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# **PSYCHOSIS WITH COEXISTING SUBSTANCE MISUSE**

**Psychosis with coexisting  
substance misuse: assessment  
and management in adults and  
young people**

**National Clinical Guideline Number X**

**National Collaborating Centre for Mental Health  
Commissioned by the  
National Institute for Health and Clinical  
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# 1 PREFACE

2 This guideline has been developed to advise on the assessment and  
3 management of adults and young people with psychosis and coexisting  
4 substance misuse. The guideline recommendations have been developed by a  
5 multidisciplinary team of healthcare professionals, a service user, a carer and  
6 guideline methodologists after careful consideration of the best available  
7 evidence. It is intended that the guideline will be useful to clinicians and  
8 service commissioners in providing and planning high-quality care for people  
9 with psychosis and coexisting substance misuse while also emphasising the  
10 importance of the experience of care for people with psychosis and coexisting  
11 substance misuse and their carers (see Appendix 1 for more details on the  
12 scope of the guideline).

13  
14 Although the evidence base is rapidly expanding, there are a number of major  
15 gaps, and future revisions of this guideline will incorporate new scientific  
16 evidence as it develops. The guideline makes a number of research  
17 recommendations specifically to address gaps in the evidence base (see  
18 Appendix 12 for the recommendations that the GDG thought were of high  
19 priority). In the meantime, it is hoped that the guideline will assist clinicians,  
20 people with psychosis and coexisting substance misuse and their carers by  
21 identifying the merits of particular treatment approaches where the evidence  
22 from research and clinical experience exists.

## 24 1.1 NATIONAL GUIDELINE

### 25 1.1.1 What are clinical practice guidelines?

26 Clinical practice guidelines are 'systematically developed statements that  
27 assist clinicians and service users in making decisions about appropriate  
28 treatment for specific conditions' (Mann, 1996). They are derived from the  
29 best available research evidence, using predetermined and systematic  
30 methods to identify and evaluate the evidence relating to the specific  
31 condition in question. Where evidence is lacking, the guidelines incorporate  
32 statements and recommendations based upon the consensus statements  
33 developed by the Guideline Development Group (GDG).

34  
35 Clinical guidelines are intended to improve the process and outcomes of  
36 healthcare in a number of different ways. They can:

37

- 1           • provide up-to-date evidence-based recommendations for the  
2           management of conditions and disorders by healthcare  
3           professionals
- 4           • be used as the basis to set standards to assess the practice of  
5           healthcare professionals
- 6           • form the basis for education and training of healthcare  
7           professionals
- 8           • assist patients and carers in making informed decisions about their  
9           treatment and care
- 10          • improve communication between healthcare professionals, service  
11          users and carers
- 12          • help identify priority areas for further research.

### 13   **1.1.2 Uses and limitation of clinical guidelines**

14   Guidelines are not a substitute for professional knowledge and clinical  
15   judgement. They can be limited in their usefulness and applicability by a  
16   number of different factors: the availability of high-quality research evidence,  
17   the quality of the methodology used in the development of the guideline, the  
18   generalisability of research findings and the uniqueness of individuals with  
19   psychosis and coexisting substance misuse.

20  
21   Although the quality of research in this field is variable, the methodology  
22   used here reflects current international understanding on the appropriate  
23   practice for guideline development ([AGREE](#) Collaboration, 2003), ensuring  
24   the collection and selection of the best research evidence available and the  
25   systematic generation of treatment recommendations applicable to the  
26   majority of people with these disorders and situations. However, there will  
27   always be some people and situations for which clinical guideline  
28   recommendations are not readily applicable. This guideline does not,  
29   therefore, override the individual responsibility of healthcare professionals to  
30   make appropriate decisions in the circumstances of the individual, in  
31   consultation with the person with psychosis and coexisting substance misuse  
32   or carer.

33  
34   In addition to the clinical evidence, cost-effectiveness information, where  
35   available, is taken into account in the generation of statements and  
36   recommendations of the clinical guidelines. While national guidelines are  
37   concerned with clinical and cost effectiveness, issues of affordability and

1 implementation costs are to be determined by the National Health Service  
2 (NHS).

3  
4 In using guidelines, it is important to remember that the absence of empirical  
5 evidence for the effectiveness of a particular intervention is not the same as  
6 evidence for ineffectiveness. In addition, of particular relevance in mental  
7 health, evidence-based treatments are often delivered within the context of an  
8 overall treatment programme including a range of activities, the purpose of  
9 which may be to help engage the person and to provide an appropriate  
10 context for the delivery of specific interventions. It is important to maintain  
11 and enhance the service context in which these interventions are delivered;  
12 otherwise the specific benefits of effective interventions will be lost. Indeed,  
13 the importance of organising care in order to support and encourage a good  
14 therapeutic relationship is at times as important as the specific treatments  
15 offered.

### 16 **1.1.3 Why develop national guidelines?**

17 The National Institute for Health and Clinical Excellence (NICE) was  
18 established as a Special Health Authority for England and Wales in 1999, with  
19 a remit to provide a single source of authoritative and reliable guidance for  
20 service users, professionals and the public. NICE guidance aims to improve  
21 standards of care, to diminish unacceptable variations in the provision and  
22 quality of care across the NHS and to ensure that the health service is person  
23 centred. All guidance is developed in a transparent and collaborative manner  
24 using the best available evidence and involving all relevant stakeholders.

25  
26 NICE generates guidance in a number of different ways, three of which are  
27 relevant here. First, national guidance is produced by the Technology  
28 Appraisal Committee to give robust advice about a particular treatment,  
29 intervention, procedure or other health technology. Second, NICE  
30 commissions public health intervention guidance focused on types of activity  
31 (interventions) that help to reduce people's risk of developing a disease or  
32 condition or help to promote or maintain a healthy lifestyle. Third, NICE  
33 commissions the production of national clinical practice guidelines focused  
34 upon the overall treatment and management of a specific condition. To enable  
35 this latter development, NICE originally established seven National  
36 Collaborating Centres in conjunction with a range of professional  
37 organisations involved in healthcare.

### 38 **1.1.4 The National Collaborating Centre for Mental Health**

39 This guideline has been commissioned by NICE and developed within the  
40 National Collaborating Centre for Mental Health (NCCMH). The NCCMH is  
41 a collaboration of the professional organisations involved in the field of

1 mental health, national service user and carer organisations, a number of  
2 academic institutions and NICE. The NCCMH is funded by NICE and is led  
3 by a partnership between the Royal College of Psychiatrists and the British  
4 Psychological Society's Centre for Outcomes Research and Effectiveness,  
5 based at University College London.

### 6 **1.1.5 From national guidelines to local protocols**

7 Once a national guideline has been published and disseminated, local  
8 healthcare groups will be expected to produce a plan and identify resources  
9 for implementation, along with appropriate timetables. Subsequently, a  
10 multidisciplinary group involving commissioners of healthcare, primary care  
11 professionals, specialist mental health and other relevant healthcare  
12 professionals, service users and carers should undertake the translation of the  
13 implementation plan into local protocols taking into account both the  
14 recommendations set out in this guideline and the priorities set in the  
15 National Service Framework for Mental Health and related documentation.  
16 The nature and pace of the local plan will reflect local healthcare needs and  
17 the nature of existing services; full implementation may take a considerable  
18 time, especially where substantial training needs are identified.

### 19 **1.1.6 Auditing the implementation of guidelines**

20 This guideline identifies key areas of clinical practice and service delivery for  
21 local and national audit. Although the generation of audit standards is an  
22 important and necessary step in the implementation of this guidance, a more  
23 broadly based implementation strategy will be developed. Nevertheless, it  
24 should be noted that the Healthcare Commission will monitor the extent to  
25 which Primary Care Trusts, trusts responsible for mental health and social  
26 care and Health Authorities have implemented these guidelines.  
27

## 28 **1.2 THE PSYCHOSIS WITH COEXISTING** 29 **SUBSTANCE MISUSE: ASSESSMENT AND** 30 **MANAGEMENT IN ADULTS AND YOUNG** 31 **PEOPLE GUIDELINE**

### 32 **1.2.1 Who has developed this guideline?**

33 The GDG was convened by the NCCMH and supported by funding from  
34 NICE. The GDG included a service user and a carer, and professionals from  
35 psychiatry, clinical psychology, general practice, nursing, pharmacy, social  
36 care, and guideline development.  
37

1 Staff from the NCCMH, who participated as full members of the GDG,  
2 provided leadership and support throughout the process of guideline  
3 development, undertaking systematic searches, information retrieval,  
4 appraisal and systematic review of the evidence. Members of the GDG  
5 received training in the process of guideline development from NCCMH staff,  
6 and the service users and carer received training and support from the NICE  
7 Patient and Public Involvement Programme. The NICE Guidelines Technical  
8 Adviser provided advice and assistance regarding aspects of the guideline  
9 development process.

10  
11 All GDG members made formal declarations of interest at the outset, which  
12 were updated at every GDG meeting. The GDG met a total of ten times  
13 throughout the process of guideline development. It met as a whole, but key  
14 topics were led by a national expert in the relevant topic. The GDG oversaw  
15 the production and synthesis of research evidence before presentation. All  
16 statements and recommendations in this guideline have been generated and  
17 agreed by the whole GDG.

### 18 **1.2.2 For whom is this guideline intended?**

19 This guideline will be relevant for adults and young people with psychosis  
20 and coexisting substance misuse.

21  
22 The guideline covers the care provided by primary, community, secondary,  
23 tertiary and other healthcare professionals who have direct contact with, and  
24 make decisions concerning the care of, adults and young people with  
25 psychosis and coexisting substance misuse.

26  
27 The guideline will also be relevant to the work, but will not cover the practice,  
28 of those in:

- 29  
30
- occupational health services
  - 31 • social services
  - 32 • the independent sector.

33 The experience of people with psychosis and coexisting substance misuse can  
34 affect the whole family and often the community. The guideline recognises  
35 the role of both in the treatment and support of people with psychosis and  
36 coexisting substance misuse.

### 1 **1.2.3 Specific aims of this guideline**

2 The guideline makes recommendations for the assessment and management  
3 of adults and young people with psychosis and coexisting substance misuse.  
4 It aims to:

- 5
- 6 • review the experience of care from the servicer user and their  
7 families' /carers' perspective
- 8 • evaluate service delivery models
- 9 • evaluate the role of psychological/ psychosocial interventions
- 10 • evaluate the role of pharmacological interventions
- 11 • integrate the above to provide best-practice advice on the  
12 assessment and care of individuals throughout the care pathway
- 13 • promote the implementation of best clinical practice through the  
14 development of recommendations tailored to the requirements of  
15 the NHS in England and Wales.

### 16 **1.2.4 The structure of this guideline**

17 The guideline is divided into chapters, each covering a set of related topics.  
18 The first three chapters provide a summary of the clinical practice and  
19 research recommendations, a general introduction to guidelines and the topic,  
20 and to the methods used to develop this guideline. Chapters 4 to 9 provide  
21 the evidence that underpins the recommendations.

22

23 Each evidence chapter begins with a general introduction to the topic that sets  
24 the recommendations in context. Depending on the nature of the evidence,  
25 narrative reviews or meta-analyses were conducted, and the structure of the  
26 chapters varies accordingly. Where appropriate, details about current practice  
27 are provided. Where meta-analyses were conducted, information is given  
28 about both the interventions included and the studies considered for review.  
29 Further sub-sections are used to present GRADE summary of findings tables,  
30 clinical summaries, and health economic evidence. A sub-section called 'from  
31 evidence to recommendations' is used to explain how the GDG moved from  
32 the evidence to the recommendations. Finally, recommendations (clinical and  
33 research) related to each topic are presented at the end of each chapter. A list  
34 of research recommendations that the GDG thought were of high priority,  
35 with the rationale for this decision, can be found in Appendix 12. On the CD-  
36 ROM, further information about the evidence and the economic plan is  
37 provided in seven appendices (see Table 1 for details).

1

**Table 1. Appendices on CD-ROM**

<b>Content</b>	<b>Appendix</b>
Clinical study characteristics tables	13
Clinical evidence forest plots	14
GRADE evidence profiles	15
Complete methodology checklists for clinical studies	16
Economic evidence profiles	17
Complete methodology checklists for economic studies	18
Economic Plan	19

2

3

4

# 1 2 PSYCHOSIS WITH COEXISTING 2 SUBSTANCE MISUSE

## 3 2.1 INTRODUCTION

4 This guideline covers the assessment and management of adults and young  
5 people (aged 14 years and older) who have a clinical diagnosis of psychosis  
6 with coexisting substance misuse.

7  
8 The term psychosis is used to describe a group of severe mental health  
9 disorders characterised by the presence of delusions and hallucinations that  
10 disrupt a person's perception, thoughts, emotions and behaviour. The main  
11 forms of psychosis are schizophrenia (including schizoaffective disorder,  
12 schizophreniform disorder and delusional disorder), bipolar disorder or other  
13 affective psychosis. Substance misuse is a broad term encompassing, in this  
14 guideline, the hazardous or harmful use of any psychotropic substance,  
15 including alcohol and either legal or illicit drugs. Such use is usually, but not  
16 always, regarded as a problem if there is evidence of dependence,  
17 characterised by psychological reinforcement of repeated substance-taking  
18 behaviour and, in some cases, a withdrawal syndrome. However, substance  
19 misuse can be harmful or hazardous without dependence, especially among  
20 people with a coexisting psychosis.

21  
22 Many people with mental health issues use substances, and for psychosis,  
23 problematic drinking and use of illicit drugs occur more frequently than in  
24 the general population (McCreadie, 2002; Regier *et al.*, 1990). For example, the  
25 Epidemiological Catchment Area (ECA) study in the USA reported a 47% and  
26 60% lifetime prevalence rate of substance misuse (drugs and alcohol) among  
27 people with schizophrenia and bipolar disorder, respectively; in the general  
28 population, the rate was 16% (Regier *et al.*, 1990). Although there is still  
29 debate as to whether there is a causal link between developing psychosis and  
30 illicit drug use, it is well established that the course of psychosis is adversely  
31 affected by substance misuse, resulting in a more prolonged and serious  
32 condition. Associated problems include non-adherence to prescribed  
33 medication, poor engagement with treatment programmes, increased risk of  
34 suicide, more inpatient stays, increased risk of violence and time spent in the  
35 criminal justice system, and poorer overall prognosis. However, many of  
36 these associations occur with substance misuse alone; the relationship  
37 between psychosis and substance misuse is complex.

38

1 Whilst an understanding of the linkage of psychosis and coexisting substance  
2 use would greatly facilitate the development of treatment approaches,  
3 knowledge to date is limited (Blanchard *et al.*, 2000). A consistency in the  
4 pattern of substance use in psychosis – alcohol being the most common  
5 substance, cannabis the most common drug, with poly substance use  
6 frequently occurring - has been established in the UK (Weaver *et al.*, 2003), the  
7 US (Blanchard *et al.*, 2000) and Australia (Kavanagh *et al.*, 2004a). This pattern  
8 of substance use in psychosis seems to be largely unrelated to service users'  
9 symptomatology (Brunette *et al.*, 1997) but rather, is associated with the same  
10 demographic correlates as for the general population (Teeson *et al.*, 2000). This  
11 suggests that in a similar way to other substance users, it is the social context  
12 and availability of substances that most often dictates substance choices in  
13 psychosis (Kavanagh *et al.*, 2004a; Patkar *et al.*, 1999). The small literature on  
14 reasons for substance use in psychosis also suggests that people with  
15 psychosis do not differ from other groups, with reasons including response to  
16 negative affective states, interpersonal conflict, and social pressures (Conrod  
17 & Stewart, 2005; Gregg *et al.*, 2009).

18  
19 Since these key dimensions of substance use are shared with the general  
20 population, the indications are that the psychological processes determining  
21 and maintaining use in people with psychosis may be similar to those found  
22 for other substance users. Hence it would seem likely that the treatment  
23 approaches developed for non – psychosis individuals will be of benefit to  
24 people with psychosis although they may need to be adapted to take account  
25 of psychosis related issues. Service user reports indicate that situations and  
26 cues triggering use may be related if not directly to psychotic symptoms then  
27 to some of the negative consequences of the illness, particularly dysphoria (an  
28 unpleasant mood state) and distress (Blanchard *et al.*, 2000). Some individuals  
29 with psychosis describe using substances to try and counteract the side effects  
30 of anti-psychotic medication; or as a preferred alternative to taking prescribed  
31 medications (Schneier & Siris, 1987). Coping motives (Mueser *et al.*, 1995), and  
32 poor problem solving abilities of this group (Carey & Carey, 1995) along with  
33 restrictive lifestyles and limitations for obtaining pleasure in other ways may  
34 then reinforce learned expectancies of the positive benefits of use.

35  
36 These vulnerability factors present considerable challenges in developing  
37 treatment programmes, and the functional aspects of substance use in  
38 psychosis may in part explain why motivation for reduction of substance use  
39 in people with psychosis is usually low (Baker *et al.*, 2006; Barrowclough *et al.*,  
40 2001; Martino *et al.*, 2002). Additionally, people with psychosis often suffer  
41 from low self esteem (Barrowclough *et al.*, 2003); thus, self efficacy may be  
42 low, which may further decrease motivation since people with psychosis may  
43 feel unable to implement changes. Moreover, psychosis is often associated

1 with a range of complex problems and within this context the contributing  
2 role of substance use may not be salient to the service user. A related issue,  
3 and again in common with substance misusers who do not have a coexisting  
4 psychosis, is that the levels of substance use may not be excessive in terms of  
5 the person's peer group, making it less likely that the person will regard their  
6 substance use as problematic.

7  
8 However, a number of psychosis-related issues increase treatment  
9 complexity. Engaging this group in treatment is often difficult and studies  
10 indicate that attrition rates are high, even for those agreeing to come into  
11 treatment (Drake *et al.*, 2004). Contributory factors may include a bias towards  
12 suspiciousness or paranoid interpretation of relationships arising from the  
13 psychotic symptoms and exacerbated by substance use; and a chaotic lifestyle  
14 along with concurrent problems making appointment scheduling and  
15 engaging in structured work more difficult. Finally, there are often  
16 medication issues that are not helpful to service user's mental state, either  
17 with service users not taking prescribed anti-psychotics (Martino *et al.*, 2002)  
18 or the non-prescription substances rendering the prescribed medication less  
19 effective.  
20

## 21 **2.2 PSYCHOSIS AND COEXISTING SUBSTANCE** 22 **MISUSE**

### 23 **2.2.1 Incidence and prevalence**

24 Reviewing the literature on comorbidity between substance misuse and  
25 psychosis presents significant challenges not least because of issues  
26 surrounding the definition of the terms involved. Substance misuse is  
27 differently defined within the diagnostic classifications (Diagnostic and  
28 Statistical Manual of Mental Disorders - *DSM-III*, *DSM-III-R*, *DSM-IV*  
29 [American Psychiatric Association, 1980, 1987, 1994] and International  
30 Statistical Classification of Diseases and Related Health Problems-10th  
31 revision [ICD-10; World Health Organization, 1992]) and operational  
32 definitions (generally scores above threshold in standardized measures of  
33 alcohol and drug misuse) employed in the contemporary literature. The  
34 literature also includes both studies relating to the comorbidity between  
35 schizophrenia (as variously defined) and substance misuse and a broader  
36 concept of psychosis that includes bipolar disorder. There is an important  
37 distinction between use of substances (which is almost ubiquitous for alcohol)  
38 on the one hand and abuse (or harmful use) and dependence on the other. In  
39 the literature by definition use of illicit substances is "abuse" and therefore  
40 problematic, although not necessarily representing harmful use or

1 dependence on the substance. Epidemiological research in this area presents  
2 many challenges and the evidence it produces must be interpreted with a  
3 degree of caution.

4  
5 Substance misuse is common in the general population: the ECA study,  
6 carried out in the USA, reported a life-time prevalence of substance misuse  
7 (including misuse of alcohol and drugs) of 16% (Regier *et al.*, 1990). In the  
8 ONS survey of psychiatric morbidity among adults living in private  
9 households in the UK, a quarter had a hazardous pattern of drinking during  
10 the year before interview, and overall, 13% of men and 8% of women aged  
11 16–74 reported using illicit drugs in the preceding 12 months (Singleton *et al.*,  
12 2000).

13  
14 Schizophrenia has a wide range of comorbidities of which substance misuse is  
15 probably the commonest (Buckley *et al.*, 2009). The ECA study in the USA  
16 found high levels of comorbidity between schizophrenia and substance  
17 misuse (47% of people with schizophrenia had a lifetime substance misuse  
18 diagnosis: odds ratio 4.6) (Regier *et al.*, 1990). Analysis of a study from  
19 Sweden that focused on the relationship between schizophrenia and  
20 offending behaviour, which found that the relationship between violent crime  
21 and schizophrenia was almost completely attenuated by coexisting substance  
22 misuse, identified comorbidity in 24.5% of service users (Fazel *et al.*, 2009a).

23  
24 Community studies of people with psychosis are challenging, but results from  
25 the US, the UK and Australia have been fairly consistent. In Australia  
26 Kavanagh and colleagues (2004a) found lifetime rates of substance misuse or  
27 dependence of 39.8% (42.1% for people with schizophrenia), with alcohol  
28 misuse (27.6%) and cannabis misuse (22.8%) the commonest. US data from the  
29 National Comorbidity Survey has provided odds ratios for coexisting  
30 substance misuse: non-affective psychosis and alcohol disorders 2.2; non-  
31 affective psychosis and drug disorders 2.7; bipolar 1 disorder and alcohol  
32 disorder 4.9; bipolar 1 and drug disorder 2.7 (Kessler *et al.*, 1994). Earlier data  
33 showed that 47% of respondents with schizophrenia met diagnostic criteria  
34 for lifetime substance misuse (including alcohol) (OR 4.6) (Regier *et al.*, 1990).

35  
36 Studies of inpatients with mixed diagnoses identify high proportions of  
37 people being admitted to a psychiatric unit with current coexisting alcohol  
38 and substance misuse – from 30% in a US sample (Huntley *et al.*, 1998) to 48%  
39 in a UK sample (Sinclair *et al.*, 2008). Similar rates are to be found in studies of  
40 service users in contact with community mental health services. Weaver and  
41 colleagues (2003) found that 44% of service users of community mental health  
42 teams in inner urban areas, where 75% of service users had a diagnosis of  
43 psychosis, had comorbid problematical use of alcohol (25%) and/or drugs

1 (31%). Alcohol and cannabis were the commonest substances to be abused  
2 and comorbidity was the norm. This was a multi-centre study and the authors  
3 noted higher levels of substance misuse in one centre (London) than the other  
4 centres (Nottingham and Sheffield). These are similar to findings from a study  
5 of the service users of a South London CMHT with “severe mental illness”  
6 where the one year prevalence of substance misuse was 36% (alcohol misuse  
7 31.6%; drug misuse 15.8%) (Menezes *et al.*, 1996).

8  
9 Margoles and colleagues (2004) reported lower rates of current substance  
10 misuse amongst a cohort of service users with schizophrenia attending an  
11 outpatient programme in Canada (15%); however they provide a telling rank  
12 order of misused substances: alcohol (10.1%); cannabis (8.2%); cocaine (2.9%);  
13 benzodiazepines (1.5%); amphetamines, stimulants and heroin (0.5% each).  
14 Substance misuse was also less common in a community cohort of service  
15 users with schizophrenia from Scotland – with 16% of service users  
16 experiencing alcohol misuse and 7% substance misuse (McCreadie, 2002). The  
17 CATIE study, which looked at drug treatment for schizophrenia, identified  
18 37% of participants as meeting diagnostic criteria for substance misuse  
19 (Swartz *et al.*, 2006).

20  
21 Studies of people with first-episode psychosis demonstrate marked  
22 differences in the prevalence of substance misuse between sites, which will  
23 plausibly reflect local patterns of substance misuse. In a German study, 23.7%  
24 of first-episode service users had a lifetime history of alcohol misuse and  
25 14.2% substance misuse (Buhler *et al.*, 2002). In contrast, 43% of a cohort of  
26 first-episode service users presenting to a service in Cambridge, UK, were  
27 diagnosed as suffering from DSM-IV alcohol misuse and 51% from cannabis  
28 misuse or dependence (Barnett *et al.*, 2007). Although the percentages of  
29 individuals with coexisting disorders are markedly different, the odds ratios  
30 between service users and age-matched controls are not. Buhler and  
31 colleagues (2002) provided an odds ratio for substance misuse against age-  
32 matched controls which for both alcohol and drugs was 2.0 – very similar to  
33 the data reported by Barnett and colleagues (2007) for all substance misuse in  
34 the previous month (OR 2.2); use of Class A drugs (OR 2.1) and use of  
35 amphetamines (OR 1.6). In addition, McCreadie (2002) reported data that  
36 showed that people with schizophrenia compared to age and gender matched  
37 general population controls, reported in the past year significantly more  
38 alcohol dependence (OR 2.7) and problem use (OR 1.80), and drug  
39 dependence (OR 7.0) and problem use (OR 4.2).

40  
41 Two recent meta-analytic studies have brought together the literature on the  
42 relationship between alcohol misuse and schizophrenia, and cannabis use and  
43 schizophrenia – cannabis being by far the commonest misused substance –

1 based on all reliable sources (Koskinen, 2009a, 2009b). These provide  
2 estimates for prevalence of comorbidity and its correlating factors. The figures  
3 are somewhat lower in absolute terms than those identified above (current  
4 alcohol use disorder 9% (IQR 4.6–19.0) – lifetime 20.6%; current cannabis use  
5 disorder 16% (IQR 8.6–28.6) – lifetime 27.1%). Cannabis use was commoner  
6 amongst first-episode service users, younger people and males rather than  
7 females (Koskinen, 2009b). Nevertheless, the prevalence and pattern of  
8 substance misuse amongst people with a psychosis will vary between  
9 geographical locations in ways that are most likely to be explained by local  
10 patterns of substance misuse in the local population; and that will be  
11 influenced by local supply and availability.

### 12 **2.2.2 Course and prognosis**

13 In some cases, the course of coexisting substance use and psychosis may be  
14 determined by the way in which it has arisen. Four main routes can be  
15 identified; (1) a primary diagnosis of psychosis with subsequent development  
16 of substance misuse, (2) a primary diagnosis of substance misuse with the  
17 secondary development of psychosis as a manifestation of the substance  
18 misuse, (3) concurrent presence of substance misuse and psychosis, the  
19 former exacerbating the latter, and (4) psychotic disorder exacerbating or  
20 altering the course of substance misuse (Lehman *et al.*, 1989). Only the second  
21 of these has a short course and good prognosis, at least in the short term, but  
22 it has been suggested that the third group, in which the substance misuse and  
23 psychosis co-occur, can be separated further into a better outcome group in  
24 which there is clearly no pre-existing psychosis, and a worse outcome group  
25 where psychosis clearly has been present in the longer term (Caton *et al.*, 2005,  
26 2007). Several drugs of misuse can lead to psychotic reactions that are  
27 unequivocally a direct consequence of the drug taken. In such cases the drug  
28 is usually taken in large or repeated doses and the psychotic reaction is  
29 manifest shortly afterwards, often after only a few hours.

30  
31 Opiates do not precipitate psychosis, but LSD (lysergic acid diethylamide) has  
32 been known to do so for many years, and perhaps is the only drug that has  
33 been incriminated in the development of long-term psychosis (Vardy & Kay,  
34 1983). True cannabis psychosis, as opposed to schizophrenia-precipitated  
35 psychosis, is a toxic state with confusion and disorientation at times as well as  
36 clearly manifest delusions and hallucinations, but this only lasts for a few  
37 hours or days (Chopra & Smith, 1974; Ghodse, 1986). Cocaine can also lead to  
38 a psychotic state with persecutory delusions and hallucinations, including the  
39 tactile hallucinations of formication (the feeling of insects crawling beneath  
40 the skin) (cocaine bug) (Ghodse *et al.*, 1998). The tropical grass, khat, although  
41 normally just acting as a mild stimulant when chewed, may also lead to brief  
42 psychotic episodes after continuous use (Alem & Shibbe, 1997). All these

1 psychotic episodes can be regarded as toxic effects of the relevant drug and,  
2 with the possible exception of LSD, resolve without any long-term  
3 consequences.

4  
5 Unfortunately, the first and fourth of these pathways to psychosis and  
6 coexisting substance misuse tend to be associated with a long course and  
7 frequent relapse. There are a series of studies that demonstrate a significantly  
8 worse outcome in terms of hospital admission (Menezes *et al.*, 1996; Zammit *et*  
9 *al.*, 2008) and bed occupancy (Menezes *et al.*, 1996; Wade *et al.*, 2006), cost  
10 (McCrone *et al.*, 2000), ceasing antipsychotic drug treatment (Wade *et al.*,  
11 2006; Zammit *et al.*, 2008), recurrence of depression and other disorders of  
12 mood (Turkington *et al.*, 2009), and the development of diabetes and early  
13 mortality (Jackson *et al.*, 2007).

### 14 **2.2.3 Morbidity and mortality**

15 People with a history of psychosis have substantially higher levels of  
16 morbidity and mortality than people without a history of psychosis. Poor  
17 physical health and premature mortality are also seen among people with  
18 drug and alcohol misuse problems. It would therefore be expected that people  
19 with psychosis plus coexisting substance misuse would have increased levels  
20 of morbidity and mortality and a large number of studies have found this to  
21 be the case.

22  
23 People with severe mental illness and substance misuse are less likely to  
24 recover from a psychotic episode and more likely to experience relapse  
25 (Dixon, 1999). Most research has focussed on the role of cannabis which  
26 appears to increase the likelihood of psychotic relapse (Linszen *et al.*, 1994).  
27 Among those admitted to hospital, symptoms of psychosis are worse among  
28 people who use cannabis and the length of stay in hospital is greater (Isaac *et*  
29 *al.*, 2005). Rates of relapse in psychosis are also higher among those who  
30 misuse other drugs, especially stimulants.

31  
32 The relationship between psychosis and coexisting substance misuse and  
33 social functioning is complex. There is evidence that, among people who  
34 develop psychosis, those with substance use have better social functioning  
35 and greater numbers of social contacts. However coexisting substance misuse  
36 can lead to social problems including impaired relationships with family  
37 members and reduced self efficacy and these may be responsible for adverse  
38 social outcomes such as housing problems and homelessness (Drake *et al.*,  
39 1991; Salyers & Museser, 2001).

40  
41 The relationship between psychosis and coexisting substance misuse and  
42 violence is more straightforward. Among people with psychosis those with

1 coexisting substance misuse are more likely be involved in violent incidents  
2 (Cuffel *et al.*, 1994). Results from a recent population-based study in Sweden  
3 suggest that the relationship between psychosis and violence may largely be  
4 the result of higher rates of substance misuse among people with severe  
5 mental illness (Fazel *et al.*, 2009b). In this study people who had schizophrenia  
6 and substance misuse were over four times more likely to be convicted of a  
7 violent crime than members of the general public. In contrast, levels of violent  
8 crime in those with schizophrenia but no substance misuse were similar to  
9 those among the general public. This study, and findings from others,  
10 provides strong evidence that any increase in levels of violence among people  
11 with psychosis is largely the result of higher levels of substance misuse in this  
12 group.

13  
14 People with psychosis and coexisting substance misuse often have poor  
15 physical health. In addition to higher rates of cardiovascular disease and  
16 other conditions that are found more frequently, those who use intravenous  
17 drugs are at far greater risk of hepatitis C, HIV and other blood borne viruses.  
18 Mortality rates are higher among people with psychosis, partly as a result of  
19 physical health problems, but also as a result of suicide. Among people with  
20 schizophrenia, coexisting substance misuse is an important risk factor for  
21 suicide with levels more than three times higher than would otherwise be  
22 expected (Hawton *et al.*, 2005).

23

## 24 **2.3 AETIOLOGY**

25 There is no single explanation for the high level of association between  
26 psychosis and substance misuse. These two disorders are usually regarded as  
27 separate diagnostic entities and therefore satisfy the strict criteria for  
28 comorbid disorders (the presence of 'any distinct clinical entity that has  
29 existed or that may occur during the clinical course of a service user who has  
30 the index disease under study (Feinstein, 1970). Although neither substance  
31 misuse nor schizophrenia are uncommon, the frequency with which they  
32 present together is many times higher than would be expected by chance (see  
33 2.2.1). It is far from clear why this is so, but several theories have been put  
34 forward for the association:

35

- 36 1. Substance misuse either precipitates the onset of, or is a direct cause of,  
37 psychosis.
- 38 2. Substance misuse is a common consequence of a psychotic disorder.
- 39 3. There is a common cause, or vulnerability, to both substance misuse  
40 and psychosis.

### 1 *Substance misuse precipitates or causes psychosis*

2 It has been known for over 40 years that substances like hallucinogens,  
3 stimulants and cannabis in high doses can be associated with or possibly  
4 cause psychotic states (Talbot & Teague, 1969). These drugs affect the  
5 dopaminergic and glutaminergic systems in the brain, which have both been  
6 associated with psychotic symptoms such as hallucinations and delusions.  
7 However, psychotic symptoms induced by substances generally tend to be  
8 short lived in comparison to psychosis in schizophrenia, and the presentation  
9 is slightly different, with predominating agitation and confusion in psychosis  
10 following drug use.

11  
12 There is a growing body of evidence showing that some substances,  
13 particularly cannabis, alcohol to a lesser extent, but not opiates, can  
14 precipitate psychosis in vulnerable people, so that the onset appears to be  
15 earlier than in those who do not take cannabis (Barnes *et al.*, 2006). Based on  
16 findings from prospective cohorts, it has been suggested that cannabis is an  
17 independent risk factor for the development of psychosis (Andreasson *et al.*,  
18 1987; Arseneault *et al.*, 2002; Van Os *et al.*, 2002), although the possibility that  
19 this association results from confounding factors or bias cannot be ruled out  
20 (Moore *et al.*, 2007). If cannabis caused schizophrenia in those who would not  
21 otherwise ever have the disease there should be an increasing prevalence of  
22 schizophrenia but this does not appear to be happening, and a very large  
23 number of cannabis consumers (1300–2700) would have to be prevented from  
24 taking cannabis to prevent just one case of schizophrenia (Hickman *et al.*,  
25 2009). The evidence to date suggests that cannabis, and to a lesser extent  
26 alcohol misuse, brings forward the onset of a psychosis that would have been  
27 likely to develop anyway.

### 28 *Psychosis causes substance misuse*

29 The most common hypothesis underlying this explanation is that people with  
30 psychosis self-medicate with substances to alleviate distressing and dysphoric  
31 symptoms of their illness. Respondents in many studies report that they use  
32 substances in order to alleviate their symptoms or negative emotional states.  
33 At the same time, it is also well documented that many service users  
34 experience exacerbation of symptoms after substance use, and there is strong  
35 evidence that the presence of substance misuse provokes relapse and  
36 generally poorer outcomes than in those with psychosis alone (Wade *et al.*,  
37 2006). Furthermore, if substances are used to alleviate symptoms, one would  
38 expect specific substances to be used to alleviate specific symptoms and  
39 substance misuse to increase with the severity of symptoms. Neither  
40 phenomenon has been demonstrated.  
41

1 However, there is some evidence to suggest that substances may be used to  
2 alleviate a more general state of dysphoria. Individuals with psychosis are  
3 more vulnerable to experiencing low mood and anxiety, not only due to  
4 symptoms of their illness, but due to social factors surrounding their situation  
5 such as stigma, social exclusion, loss of functioning ability and financial  
6 difficulties. They are therefore more likely to use substances as short term  
7 relief from the consequent unpleasant feelings (Phillips & Johnson, 2003).

8 There are further ways in which social factors may contribute to substance  
9 misuse in individuals with psychosis. This is a population in which  
10 educational and vocational failure, poverty, lack of social and recreational  
11 activity are common. Already at the margins of society, such people may feel  
12 more accepted and identify more with the drug-using population, and,  
13 because of their socio-economic position, may be housed in neighbourhoods  
14 where drug misuse is commonplace.

15  
16 It is also possible that antipsychotic medication may itself lead to an increase  
17 in substance misuse. These medications work by blocking dopamine receptors  
18 in the brain, including dopaminergic reward systems in the brain. Individuals  
19 may attempt to counteract this effect by using substances.

#### 20 *A common cause for both disorders*

21 It has been suggested that there may be a common genetic risk factor for both  
22 psychosis and substance misuse, particularly via the catechol-O-  
23 methyltransferase gene (COMT). This was initially suggested by Caspi and  
24 colleagues (2005), who postulated a gene-environment interaction as the  
25 cause of some episodes of psychosis. However, this has not been confirmed  
26 and on present evidence (Hosák, 2007; Zammit *et al.*, 2007) the relationship is  
27 too non-specific to be causal. Several studies have shown that the presence of  
28 antisocial personality disorder independently increases the incidence of both  
29 psychosis and substance misuse. Furthermore, people with antisocial  
30 personality disorder also tend to develop both psychosis and substance  
31 misuse disorder at an earlier age. More evidence is required to establish the  
32 nature of this relationship and whether there is a causative element. Further  
33 research has proposed that abnormalities in the hippocampus and frontal  
34 lobes of the brain may cause symptoms of schizophrenia and these areas also  
35 provide positive reinforcement of drug reward and reduce inhibition of drug  
36 seeking behaviour.

37  
38 A similar framework to the above three categories has been used to  
39 understand the specific group of individuals with psychosis and cannabis use.  
40 Hambrecht and Hafner (2000) describe a “vulnerability-stress-coping” model  
41 of schizophrenia and cannabis use which divides this group into three  
42 categories:

1

2 • The vulnerability group are those who use cannabis years before  
3 developing psychosis. The authors explain that cannabis may reduce  
4 their threshold of vulnerability to developing schizophrenia, either  
5 by a biological, psychological or social process, as well as reducing  
6 the service users coping resources.

7

8 • The stress group in whom the onset of cannabis misuse and  
9 psychosis occurs around the same time. This group comprises  
10 individuals already vulnerable to schizophrenia for genetic, pre- or  
11 perinatal influences and cannabis promotes the release of dopamine  
12 and this stimulation of dopamine pathways can precipitate the onset  
of disease.

13

14 • The coping group start using cannabis after the onset of psychosis  
15 and they self medicate with the drug. The theory is that they learn to  
16 counterbalance the unpleasant hypodopaminergic prefrontal state of  
schizophrenia with the dopaminergic effects of cannabis.

17

18 This model has also to accommodate the evidence of a dose-response  
19 relationship between cannabis and psychosis, as the data suggest that  
20 individuals who consume of the strongest forms of cannabis, particularly  
21 'skunk', are more prone to psychosis (Murray *et al.*, 2007; Verdoux *et al.*, 2005).

22

23 In summary, there is still some doubt as to whether cannabis precipitates the  
24 onset of psychosis in those who are vulnerable to the condition and the  
25 precise mechanism whereby such an association is generated still remains  
26 open to many explanations.

## 27 **2.4 DIAGNOSIS**

28

29 The term "dual diagnosis" is often used in both clinical practice and  
30 healthcare literature, and covers a wide spectrum of co-occurring psychiatric  
31 disorders and substance misuse with complex inter-relationships and  
32 interactions. The coexistence of psychosis with substance misuse is commonly  
33 referred to as 'dual diagnosis' when it is defined narrowly, but as this term is  
34 also used to describe other forms of comorbidity (for example, mental illness  
35 and intellectual disability), it is best avoided or, if used, the comorbidities  
36 described specifically.

37

38 People with psychosis and coexisting substance misuse may have multiple  
39 (rather than two as implied by 'dual') diagnoses both in relation to mental  
illness (for example, schizophrenia and anxiety, depression, personality

1 disorder) and substance misuse (for example, alcohol dependence, and  
2 harmful use of another substance(s)).

3  
4 In DSM-IV (American Psychiatric Association, 1994), a distinction is made  
5 between independent (primary psychiatric comorbidity) and substance-  
6 induced (organic) psychiatric comorbidity and the category of expected  
7 symptoms of substance use or withdrawal (Abou-Saleh, 2004).

8  
9 DSM-IV diagnostic criteria enable clinicians to distinguish 'primary',  
10 'substance-induced' psychiatric disorders, and the 'expected effects' of  
11 intoxication and withdrawal (Samet *et al.*, 2004). A 'primary' disorder is  
12 diagnosed if 'the symptoms are not due to the direct physiological effects of a  
13 substance'. Before diagnosing a 'substance-induced' disorder, a primary  
14 classification must first be ruled out (see Table 2 and Table 3).

15 **Table 2. Criteria for substance abuse (DSM-IV) and harmful use (ICD-10)**

DSM-IV	ICD-10
1) A maladaptive pattern of substance use leading to clinically significant impairment or distress, as manifested by one (or more) of the following occurring within a 12-month period	1) A pattern of psychoactive substance use that is causing damage to health; the damage may be to physical or mental health
2) Recurrent substance use resulting in a failure to fulfil major role obligations at work, school, or home	
3) Recurrent substance abuse in situations that are physically hazardous	
4) Recurrent substance-abuse-related legal problems	
5) Continued substance abuse despite having persistent or recurrent social or interpersonal problems caused or exacerbated by the effects of the substance	
6) Has never met the criteria for substance dependence for this class of substance	
<i>Note.</i> DSM-IV = Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (American Psychiatric Association, 1994); ICD-10 = Tenth Revision of the International Classification of Diseases and Related Health Problems (World Health Organization, 1992).	

16  
17 There are four conditions under which an episode that coexists with  
18 substance intoxication or withdrawal can be considered primary:

- 1 1. Symptoms 'are substantially in excess of what would be expected
- 2 given the type or amount of the substance used or the duration of use'.
- 3 2. A history of non-substance-related episodes.
- 4 3. The onset of symptoms precedes the onset of the substance use.
- 5 4. The symptoms persist for a substantial period of time (i.e. at least a
- 6 month) after the cessation of intoxication or acute withdrawal.

7

8 If neither 'primary' nor 'substance-induced' criteria are met, then the  
9 syndrome is considered to represent intoxication or withdrawal effects of  
10 alcohol or drugs

11

12 The ICD-10 Diagnostic Criteria for Research (World Health Organization,  
13 1992) provides specified criteria to differentiate primary disorders and  
14 disorders resulting from psychoactive substance use for psychotic disorders.  
15 As in DSM-IV, ICD-10 excludes psychotic episodes attributed to psychoactive  
16 substance use from a primary classification.

17

1

**Table 3. Criteria for dependence syndrome in DSM-IV and ICD-10**

DSM-IV	ICD-10
Diagnosis of dependence should be made if three (or more) of the following have been experienced or exhibited at any time in the same 12-month period	Diagnosis of dependence should be made if three or more of the following have been experienced or exhibited at some time during the last year
Tolerance defined by either need for markedly increased amount of substance to achieve intoxication or desired effect or markedly diminished effect with continued use of the same amount of the substance	A strong desire or sense of compulsion to take the substance
Withdrawal as evidenced by either of the following: the characteristic withdrawal syndrome for the substance <b>or</b> the same (or closely related) substance is taken to relieve or avoid withdrawal symptoms	Difficulties in controlling substance-taking behaviour in terms of its onset, termination, or levels of use
The substance is often taken in larger amounts over a longer period of time than was intended	Physiological withdrawal state when substance use has ceased or been reduced, as evidenced by either of the following: the characteristic withdrawal syndrome for the substance <b>or</b> use of the same (or closely related) substance with the intention of relieving or avoiding withdrawal symptoms
Persistent desire or repeated unsuccessful efforts to cut down or control substance use	Evidence of tolerance, such that increased doses of the psychoactive substance are required in order to achieve effects originally produced by lower doses
A great deal of time is spent in activities necessary to obtain the substance, use the substance, or recover from its effects	Progressive neglect of alternative pleasures or interests because of psychoactive substance use and increased amount of time necessary to obtain or take the substance or to recover from its effects
Important social, occupational, or recreational activities given up or reduced because of substance use	Persisting with substance use despite clear evidence of overly harmful consequences (physical or mental)
Continued substance use despite knowledge of having had a persistent or recurrent physical or psychological problem that was likely to have been caused or exacerbated by the substance	
<i>Note.</i> DSM-IV = Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (American Psychiatric Association, 1994); ICD-10 = Tenth Revision of the International Classification of Diseases and Related Health Problems (World Health Organization, 1992).	

2

1 In ICD-10, psychotic disorders can be attributed to psychoactive substance  
2 use under three conditions:

3

4 1. The onset of symptoms must occur during or within 2-weeks of  
5 substance use.

6 2. The psychotic symptoms must persist for more than 48-hours.

7 3. The duration of the disorder must not exceed 6 months.

8

9 A psychotic disorder attributed to psychoactive substance use can be  
10 specified as predominantly depressive or predominantly manic. However,  
11 unlike DSM-IV, ICD-10 does not provide a separate psychoactive substance  
12 related category for any other type of psychiatric disorder. By definition, ICD-  
13 10 'organic mental disorder' excludes alcohol or other psychoactive  
14 substance-related disorders. ICD-10 organic mood disorder and organic  
15 delusional disorder cannot be used to diagnose episodes co-occurring with  
16 heavy psychoactive substance use. Thus, the DSM-IV concept of symptoms  
17 that are greater than the expected effects of intoxication and withdrawal is not  
18 included in ICD-10. The DSM-IV concept of 'primary' and 'substance-  
19 induced' syndromes, and the ICD-10 concept of 'psychotic disorders due to  
20 psychoactive substance use,' support the notion that a psychiatric disorder  
21 warranting clinical attention can co-occur with heavy substance use.  
22 However, these categories continue to present diagnostic challenges.  
23 Differential diagnosis of categories of depression, anxiety, and psychosis often  
24 hinges on interpretation of the term 'in excess' of the 'expected' effects of  
25 substance use, including service users with chronic substance use beginning  
26 at an early age. These expected effects are not clearly defined by either system  
27 and are thus left to clinical judgment (Samet *et al.*, 2004).

28

## 29 **2.5 TREATMENT AND MANAGEMENT IN THE NHS**

30 A major problem in the treatment and management of psychosis and  
31 coexisting substance misuse is that services fail to recognise and detect both  
32 problems, hence the need for a comprehensive assessment and package of  
33 care.

### 34 **2.5.1 Pharmacological treatments**

#### 35 *Treatments for psychosis*

36 As part of a comprehensive package of care, a range of treatments can be  
37 recommended for people with psychosis and coexisting substance misuse.  
38 Most commonly, antipsychotic drugs are used to manage the symptoms of

1 psychosis. The updated NICE guidelines for the management of  
2 schizophrenia provide a helpful framework to guide the use of these drugs  
3 (NICE, 2009a). The range of treatments offered for people with psychosis and  
4 coexisting substance misuse may not be in line with treatments offered in  
5 other NICE guidelines however, as there is significant local variation in  
6 treatments offered for this population.

7  
8 With the exception of clozapine, all available antipsychotic drugs appear to be  
9 equally effective in controlling symptoms; therefore the decision to use a  
10 particular agent may be determined by the need to avoid particular side  
11 effects or other complications of treatment such as drug interactions.

12  
13 Where possible, the choice of which antipsychotic to use can be guided by the  
14 informed view of the service user. Outcomes from previous treatments may  
15 help refine the choice. Oral formulations are generally preferable, but where  
16 covert non-adherence is problematic, a long acting depot formulation may be  
17 advantageous.

18  
19 Previous guidance has stated that doses above the licensed range or  
20 combinations of antipsychotics are problematic (NICE, 2002, 2009a; Royal  
21 College of Psychiatrists, 2006), as for the majority of service users, there have  
22 been few advantages found over the licensed dose of the individual drugs. If  
23 treatment response is inadequate, despite the use of licensed doses of at least  
24 two antipsychotics over a fixed duration of time, one option which can be  
25 considered for further treatment is clozapine.

### 26 *Treatments for addiction*

27 Engagement with the service user is vital so that active treatment can then  
28 commence. There are a number of pharmacological treatments for substance  
29 problems, including replacement treatments (nicotine, opiates etc.) and  
30 others. These are commonly delivered within the context of psychosocial  
31 interventions, and the overall framework of a primary care setting and/or the  
32 specialist multidisciplinary team. Medications are available for the treatment  
33 of withdrawal, for stabilization, for substitution and maintenance regimes,  
34 and for relapse prevention. For alcohol, medications include chlordiazepoxide  
35 and diazepam for withdrawal while for opiates, methadone and  
36 buprenorphine are prescribed. Relapse prevention is achieved by the use of  
37 naltrexone and acamprosate for alcohol dependence, and naltrexone for  
38 opiate dependence.

39  
40 Additional treatment for nutritional deficiencies deficiency syndromes, or  
41 physical illness, such as diabetes or hypertension may be required as many  
42 people with psychosis and coexisting substance misuse will have physical

1 illnesses (associated with, or independent of, their psychosis and substance  
2 misuse) that will require the appropriate pharmacological interventions.  
3 There are a range of NICE guidelines and health technology assessments  
4 which are related to the treatment of addiction and mental illness (see NICE  
5 website: [www.nice.org](http://www.nice.org)).

## 6 **2.5.2 Psychological treatments**

7 Similarly, there are a range of psychological interventions that are beneficial  
8 in the treatment of psychosis and coexisting substance misuse. In general, a  
9 non-judgmental style of engagement is considered appropriate as a prelude to  
10 enhancing engagement. In the course of such a motivational approach, the  
11 individual's appreciation and attitude to their illness can be elicited and  
12 further, more intensive psychosocial interventions commenced. These may  
13 include supportive counselling, behavioural and cognitive techniques with an  
14 individual, group or family, as well as contingency management and skills  
15 training. There are a wealth of self-help mutual aid groups which provide  
16 sustained support.

### 17 *Psychological treatment approaches*

18 In both the UK and the US consensus agreements have been reached on key  
19 elements of treatment approaches for assisting service users with psychosis  
20 and coexisting substance use (Department of Health, 2002; Ziedonis *et al.*,  
21 2005). It is proposed that effective treatment for people with psychosis and  
22 coexisting substance use usually requires an integrated treatment approach.  
23 Such "integrated care" which combines elements of mental health and  
24 substance use approaches in one delivery system, was pioneered in New  
25 Hampshire, US, in the 80's, and has been well documented (Mueser & Drake,  
26 2003). The advantages of an integrated approach include ensuring that both  
27 elements of the dual problems are given attention and that interaction  
28 between mental health and substance use problems described above can be  
29 formulated and addressed. There is further consensus agreement that  
30 interventions need to take account of service users' motivation to address or  
31 reduce their substance use and there has been particular emphasis on  
32 applying motivational interventions, and in particular motivational  
33 interviewing (MI) (Miller & Rollnick, 2002). Miller and Rollnick define MI as  
34 "a client-centred, directive method for enhancing intrinsic motivation to  
35 change by exploring and resolving ambivalence". Building intrinsic  
36 motivation for change involves the therapist selectively eliciting and  
37 reinforcing 'Change Talk', that is the service user's own arguments and  
38 motivations for change. Essentially this involves engaging the service user,  
39 offering information and feedback from assessments where appropriate and  
40 exploring and resolving ambivalence in an affirming and non judgemental  
41 way.

1  
2 The additional element that has been used most commonly in recent  
3 treatment approaches for psychosis and coexisting substance misuse is  
4 cognitive behaviour therapy (CBT). CBT is one of the most commonly used  
5 therapeutic orientations in the field of substance disorders (Stewart & Conrod,  
6 2005). Moreover, in recent years CBT has been recognised to be effective in  
7 reducing the symptoms of psychosis (Pilling *et al.*, 2002). The CBT approach  
8 for individuals with psychosis and coexisting substance use problems is  
9 guided by individual formulations and by Marlatt and Gordon's (1985) model  
10 of relapse prevention. Components may include: identifying and increasing  
11 awareness of high risk situations/warning signs; developing new coping  
12 skills for handling such high risk situations/warning signs, with particular  
13 attention to psychosis symptom and mental health related problems  
14 highlighted in the formulation (for example, strategies for dealing with  
15 distressing voices or with depressed mood); coping with cravings and urges;  
16 making lifestyle changes so as to decrease need/urges for drugs and/or  
17 alcohol or to increase healthy activities/alternative options to substance use;  
18 normalising lapses in substance use and developing strategies and plans for  
19 acting in the event of lapse/relapse so that adverse consequences may be  
20 minimized; cognitive restructuring around alcohol and drug expectancies.

21  
22 Environmental factors also play an important part in the maintenance and  
23 persistence of drug misuse in psychosis. Many individuals in this group have  
24 life styles in which drug use is part of the daily fabric of existence and they  
25 cannot contemplate changes that are associated with cessation of substances  
26 that are regarded as essential requirements. Major environmental change is  
27 often regarded as desirable but very difficult to achieve. Exhortations to stop  
28 or reduce drug intake usually fail but concentration on changing the social  
29 and personal environment may be of value (Tyrer *et al.*, in press).

### 30 **2.5.3 Service level and other interventions**

31 Three models of service provision have been identified for the care and  
32 treatment of people with psychosis and coexisting substance misuse: serial,  
33 parallel and integrated. In the serial model psychosis and substance misuse  
34 disorders are treated consecutively by different services. In the parallel both  
35 are treated at the same time but by different services (mental health address  
36 the psychosis, substance misuse the drug and/or alcohol issues). In the  
37 integrated model, psychosis and substance issues are addressed at the same  
38 time, in one setting, by one team. This is the model that was advocated by the  
39 Department of Health (2002) building on work conducted in New Hampshire  
40 (US) (for example, Mueser & Drake, 2003).

41

1 In the UK service configurations, treatment philosophies and funding streams  
2 militate against integrated provision. Mental health and substance misuse  
3 services are separate. They are often provided by different organisations and  
4 even when both are provided by the same NHS Trust they usually have  
5 different organisational and managerial structures. Furthermore staff within  
6 each service often lack the knowledge and skills for working with people  
7 from the 'other' group. There has been a tendency for people to be 'bounced'  
8 between services, each requiring the service user to deal with the 'other'  
9 problem first (serial model). In some areas service provision has been  
10 enhanced by mental health and substance misuse services working together,  
11 with the mental health services focusing on care and treatment of the person's  
12 psychosis, and the substance misuse service the substance misuse issues  
13 (parallel model). This is generally considered to be an improvement on the  
14 serial model but it still has weaknesses, for example: treatment in either  
15 system may be incomplete due to a lack of attention to the co-morbid  
16 condition; each system can continue to provide standard treatment and not  
17 modify it to accommodate the co-morbid condition; there is the potential for  
18 miscommunication and contradictory recommendations and it falls to the  
19 service user to integrate the two systems (Drake *et al.*, 1993, 1995). Moreover  
20 in the current UK drug treatment system the focus is on 'problem drug users'  
21 (heroin and crack cocaine) leaving gaps in provision for those using other  
22 substances.

23  
24 The differing treatment philosophies for mental health and substance misuse  
25 services can also make it difficult for people to receive coherent treatment. If  
26 necessary mental health services can compel people to receive treatment  
27 under the provision of the Mental Health Act (2007) (HMSO, 2007). Some  
28 services are also proactive in engaging and retaining vulnerable service users  
29 with psychosis in treatment (in particular assertive outreach teams).  
30 Substance misuse services usually expect some level of readiness to change  
31 and the service user to attend a team base to receive treatment. Many people  
32 with psychosis and coexisting substance misuse do not see their substance use  
33 as problematic so are unlikely to access substance misuse services. If mental  
34 health services do not view the treatment of substance misuse as an integral  
35 part of mental health treatment, this aspect of the service users' needs is likely  
36 to be overlooked.

37  
38 Given the high prevalence of substance misuse in people with psychosis, the  
39 fact that many do not see their substance use as a problem, and the negative  
40 impact substance use can have on mental health, it is inevitable that many  
41 service users in both community and inpatient mental health services will  
42 have psychosis and coexisting substance misuse. Yet evidence suggests that  
43 substance misuse often goes undetected in people with mental illness (for

1 example, Barnaby, 2003; Noordsky *et al.*, 2003). Even when it has been  
2 identified, the lack of competence in working with substance misuse issues in  
3 general mental health settings, and the sometimes negative attitudes of staff  
4 to this group, may result in substance misuse needs not being addressed at all  
5 or, if they are, interventions not being delivered in line with best practice.

6  
7 In some areas dual diagnosis practitioners/teams have been developed to  
8 support the delivery of more integrated care. Models vary in different  
9 localities but typically their work includes delivering staff training and  
10 supervision, and engaging in joint work with mental health colleagues.

11  
12 People with psychosis and coexisting substance misuse often have multiple  
13 needs related to their psychosis and substance use, for example, physical  
14 health problems, financial difficulties, housing problems, difficulty in caring  
15 for their children and being involved in illegal activity. As a consequence they  
16 are likely to have contact with a variety of services, only some of which will  
17 be provided by the NHS. Not all the public services necessary for this  
18 desperate group of people will therefore be covered by this guidance.

#### 19 **2.5.4 Forensic/justice system**

20 Assessments for substance misuse history or problems in secure hospital units  
21 or prisons usually rely on good history taking rather than the use of research  
22 tools. Bloye and colleagues (2003) recommend a multi assessment approach to  
23 enable a more comprehensive assessment of substance use disorders within  
24 the forensic population.

25  
26 In recently established personality disorder services funded by the Dangerous  
27 and Severe Personality Disorder (DSPD) programme the Violence Risk Scale  
28 (VRS; Wong *et al.*, 2006, 2007) is routinely being used. This is designed to  
29 integrate the assessment of risk, need, responsivity and treatment change in a  
30 single tool. It assesses the service user's risk of violence, identifies treatment  
31 targets linked to violence, and assesses the service user's readiness for change  
32 and their post-treatment improvement on the treatment targets. The tool uses  
33 the stages of change model and integrates the presence of substance misuse  
34 histories and problems in the risk assessment and the formulation of  
35 treatment targets. It is important to note that some of the service users in these  
36 DSPD units have a history of comorbid psychosis and personality disorder, as  
37 well as substance misuse.

38  
39 The treatment of prisoners identified as having mental illness with or without  
40 coexisting substance misuse problems takes place in NHS or other hospitals  
41 once a prisoner has been identified as having a psychiatric disorder and been  
42 diverted. Treatment with medication can be given in prison for those

1 prisoners who can give informed consent. For those service users who are  
2 remitted back to prison following a period of treatment in hospital, there are  
3 difficulties in providing specific substance misuse treatment programmes  
4 because the mental health inreach teams are not adequately resourced  
5 (Sainsbury Centre for Mental Health, 2008).

6  
7 Most hospital secure units have treatment programmes for substance misuse  
8 based on cognitive behavioural principles (Derry, 2008). Most of these  
9 programmes are offered on a group basis and incorporate elements of  
10 motivation to change work, understanding links between substance misuse,  
11 mental health and offending, relapse prevention and skills development.  
12 These treatment programmes are not specific to forensic settings and are  
13 similar to interventions offered for generic service users in inpatient and  
14 community services. There are no good controlled evaluations with large  
15 sample sizes of these treatments, however in a recent retrospective evaluation  
16 of an inpatient drug and alcohol treatment programme, Derry and Batson  
17 (2008) found some evidence to suggest that those who had completed a  
18 treatment programme were less likely to use drugs or alcohol after discharge.  
19 In addition, those who had completed a treatment programme spent a greater  
20 proportion of time in the community compared with those who did not  
21 complete the programme. Suggestions for future research included more  
22 objective assessments of drug use, the need to control for treatment  
23 adherence, motivation to change, and incorporating a level of personal insight  
24 of mental health problems in studies using large sample sizes.

25  
26 Within secure units, there is a common practice of considering discharge into  
27 the community after service users with a history of drug or alcohol misuse  
28 have remained abstinent whilst utilising significant amounts of unescorted  
29 community leave. This practice can lead to extended detention long after  
30 abnormal mental states have been treated. Despite the significant impact this  
31 may have on length of stay, there is no good research evaluation of this  
32 practice and the impact on substance misuse post discharge has not been  
33 described. The effect of banning service users from using illicit substances or  
34 alcohol as part of the conditions of discharge has also not been evaluated.

## 36 **2.6 ECONOMIC COSTS**

37 The available epidemiological data from within the UK suggests that a  
38 significant number of individuals with psychosis, have coexisting substance  
39 misuse (Menezes *et al.*, 1996; Sinclair *et al.*, 2008; Weaver *et al.*, 2003). However,  
40 evidence on the extent to which these individuals incur extra costs in terms of  
41 health care or lost productivity is very limited both within and outside the  
42 UK.

1  
2 To date, only one UK study compared the service use and costs of individuals  
3 with a diagnosis of psychosis and coexisting substance misuse with those  
4 with a diagnosis of psychosis alone (McCrone *et al.*, 2000). Service use data,  
5 including core psychiatric services, general health care, social, education,  
6 employment and legal services, were collected over a six month period using  
7 the Client Service Receipt Interview (CSRI). Mean core health care costs  
8 (including psychiatric inpatient episodes, contacts with mental health staff  
9 and emergency and day care attendances) were significantly higher in service  
10 users with psychosis and coexisting substance misuse (£2,626 vs. £1,060;  
11  $p=0.038$ ). However, the difference in total mean costs (including supported  
12 accommodation, social and legal services) did not reach statistical significance  
13 between the two groups (£3,913 vs. £2,903;  $p=0.271$ ).  
14

15 A US-based study examined the costs of psychiatric treatment for seriously  
16 mentally ill people (diagnosed with schizophrenia; major affective disorder or  
17 other psychoses) with coexisting substance misuse in comparison with  
18 mentally ill people without substance misuse (Dickey & Azeni, 1996). Paid  
19 claims for psychiatric care, including hospital admissions, residential  
20 treatment, medical treatments and case management were collected for adult  
21 Medicaid beneficiaries in the state of Massachusetts. In this study, total  
22 annual mean costs (1992) were substantially higher in service users with  
23 coexisting substance misuse (\$22,917 vs. \$13,930). Importantly, these cost  
24 differences were largely explained by greater inpatient psychiatric treatment  
25 whilst substance misuse treatment accounted for a small proportion of the  
26 extra cost.  
27

28 Another US study compared the long-term patterns of service use and costs in  
29 service users with a dual diagnosis of psychiatric and substance misuse  
30 disorders, with those without a dual diagnosis. Of service users with  
31 psychosis and coexisting substance misuse, 46–48% had a primary diagnosis  
32 of schizophrenia or bipolar disorder (Hoff & Rosenheck, 1998). Data was  
33 analysed from longitudinal services use files that recorded all hospital and  
34 outpatient services provided by the Department of Veterans Affairs mental  
35 health system from 1990 to 1996. Costs were calculated for five types of health  
36 care: inpatient and outpatient psychiatric services, substance misuse and  
37 medical/surgical care. Separate analyses were conducted for service users  
38 who were categorised either as inpatient or outpatient at the time of case  
39 identification. Overall, there was no significant difference in mean annual  
40 costs between those with psychiatric and combined substance misuse when  
41 compared to those with a psychiatric diagnosis alone in the hospital sample.  
42 However, in the outpatient sample, service users with coexisting psychiatric  
43 and substance misuse disorders incurred substantially higher mean annual

1 costs between 1990 and 1996. Most of these extra costs incurred by people  
2 with psychosis and coexisting substance misuse in the outpatient sample  
3 were due to inpatient psychiatric and substance misuse care.

4

5 To date, no single UK study has attempted to estimate the combined total  
6 health care and societal costs of treating people with a diagnosis of psychosis  
7 and coexisting substance misuse. In 2007, the total health service costs of  
8 severe mental illness (Schizophrenia; Bipolar Disorder and related conditions)  
9 were estimated at £3.8 billion whilst the total costs of lost employment were  
10 estimated at £5.4 billion (McCrone *et al.*, 2008). Based on UK-based estimates  
11 of prevalence rates of between 36–44% for people with comorbid substance  
12 misuse (Menezes *et al.*, 1996; Weaver *et al.*, 2003), it is possible that the total  
13 annual health service and productivity costs of psychosis and substance  
14 misuse could be between £3.3 and £4 billion. However, further empirical  
15 research is required to assess the true economic burden of severe mental  
16 illness and substance misuse in the UK.

17

# 3 METHOD USED TO DEVELOP THIS GUIDELINE

## 3.1 OVERVIEW

The development of this guideline drew upon methods outlined by NICE (further information is available in *The Guidelines Manual* [NICE, 2009b]). A team of health professionals, lay representatives and technical experts known as the Guideline Development Group (GDG), with support from the NCCMH staff, undertook the development of a person centred, evidence-based guideline. There are six basic steps in the process of developing a guideline:

1. Define the scope, which sets the parameters of the guideline and provides a focus and steer for the development work.
2. Define review questions considered important for practitioners and service users.
3. Develop criteria for evidence searching and search for evidence.
4. Design validated protocols for systematic review and apply to evidence recovered by search.
5. Synthesise and (meta-) analyse data retrieved, guided by the review questions, and produce GRADE evidence profiles and summaries.
6. Answer review questions with evidence-based recommendations for clinical practice.

The clinical practice recommendations made by the GDG are therefore derived from the most up-to-date and robust evidence base for the clinical and cost effectiveness of the treatments and services used in the treatment and management of psychosis and coexisting substance misuse. In addition, to ensure a service user and carer focus, the concerns of service users and carers regarding health and social care have been highlighted and addressed by recommendations agreed by the whole GDG.

## 3.2 THE SCOPE

Guideline topics are selected by the Department of Health and the Welsh Assembly Government, which identify the main areas to be covered by the guideline in a specific remit (see *The Guidelines Manual* for further information). The NCCMH developed a scope for the guideline based on the remit. The purpose of the scope is to:

- 1       • provide an overview of what the guideline will include and exclude
- 2       • identify the key aspects of care that must be included
- 3       • set the boundaries of the development work and provide a clear
- 4       framework to enable work to stay within the priorities agreed by
- 5       NICE and the NCC and the remit from the Department of
- 6       Health/Welsh Assembly Government
- 7       • inform the development of the review questions and search strategy
- 8       • inform professionals and the public about expected content of the
- 9       guideline
- 10      • keep the guideline to a reasonable size to ensure that its
- 11      development can be carried out within the allocated period.

12 An initial draft of the scope was sent to registered stakeholders who had  
13 agreed to attend a scoping workshop. The workshop was used to:

- 14
- 15      • obtain feedback on the selected key clinical issues
- 16      • identify which patient or population subgroups should be specified
- 17      (if any)
- 18      • seek views on the composition of the GDG
- 19      • encourage applications for GDG membership.

20

21 The draft scope was subject to consultation with registered stakeholders over  
22 a 4-week period. During the consultation period, the scope was posted on the  
23 NICE website ([www.nice.org.uk](http://www.nice.org.uk)). Comments were invited from stakeholder  
24 organisations and the Guideline Review Panel (GRP). Further information  
25 about the GRP can also be found on the NICE website. The NCCMH and  
26 NICE reviewed the scope in light of comments received, and the revised  
27 scope was signed off by the GRP.

28

### 29 **3.3 THE GUIDELINE DEVELOPMENT GROUP**

30 The GDG consisted of: a service user, a representative from a service user  
31 organisation and a carer; professionals in psychiatry, clinical psychology,  
32 nursing, social work, and general practice; academic experts in psychiatry and  
33 psychology; experts in guideline development. The guideline development

1 process was supported by staff from the NCCMH, who acted as full members  
2 of the GDG, and undertook the clinical and health economics literature  
3 searches, reviewed and presented the evidence to the other members of the  
4 GDG, managed the process, and contributed to drafting the guideline.

### 5 **3.3.1 Guideline Development Group meetings**

6 Ten GDG meetings were held between May 2009 and October 2010. During  
7 each day-long GDG meeting, in a plenary session, review questions and  
8 clinical and economic evidence were reviewed and assessed, and  
9 recommendations formulated. At each meeting, all GDG members declared  
10 any potential conflicts of interest, and service user and carer concerns were  
11 routinely discussed as part of a standing agenda.

### 12 **3.3.2 Service users and carers**

13 Individuals with direct experience of services gave an integral service-user  
14 focus to the GDG and the guideline. The GDG included a service user and a  
15 representative of a service user group. They contributed as full GDG members  
16 to writing the review questions, helping to ensure that the evidence  
17 addressed their views and preferences, highlighting sensitive issues and  
18 terminology relevant to the guideline, and bringing service-user research to  
19 the attention of the GDG. In drafting the guideline, they contributed to  
20 writing the guideline's introduction and identified recommendations from the  
21 service user and carer perspective.

### 22 **3.3.3 National and international experts**

23 National and international experts in the area under review were identified  
24 through the literature search and through the experience of the GDG  
25 members. These experts were contacted to recommend unpublished or soon-  
26 to-be published studies in order to ensure up-to-date evidence was included  
27 in the development of the guideline. They informed the group about  
28 completed trials at the pre-publication stage, systematic reviews in the  
29 process of being published, studies relating to the cost effectiveness of  
30 treatment and trial data if the GDG could be provided with full access to the  
31 complete trial report. Appendix 5 lists researchers who were contacted.  
32

## 33 **3.4 REVIEW QUESTIONS**

34 Review (clinical) questions were used to guide the identification and  
35 interrogation of the evidence base relevant to the topic of the guideline. Before  
36 the first GDG meeting, an analytic framework (see Appendix 6) was prepared  
37 by NCCMH staff based on the scope and an overview of existing guidelines,  
38 and discussed with the guideline Chair. The framework was used to provide

1 a structure from which the review questions were drafted. Both the analytic  
 2 framework and the draft review questions were then discussed by the GDG at  
 3 the first few meetings and amended as necessary. Where appropriate, the  
 4 framework and questions were refined once the evidence had been searched  
 5 and, where necessary, sub-questions were generated. Questions submitted by  
 6 stakeholders were also discussed by the GDG and the rationale for not  
 7 including any questions was recorded in the minutes. The final list of review  
 8 questions can be found in Appendix 6.

9  
 10 For questions about interventions, the PICO (Patient, Intervention,  
 11 Comparison and Outcome) framework was used (see Table 4).  
 12

**Table 4: Features of a well-formulated question on effectiveness  
 intervention – the PICO guide**

Patients/ population	Which patients or population of patients are we interested in? How can they be best described? Are there subgroups that need to be considered?
Intervention	Which intervention, treatment or approach should be used?
Comparison	What is/are the main alternative/s to compare with the intervention?
Outcome	What is really important for the patient? Which outcomes should be considered: intermediate or short-term measures; mortality; morbidity and treatment complications; rates of relapse; late morbidity and readmission; return to work, physical and social functioning and other measures such as quality of life; general health status; costs?

13  
 14 In some situations, the prognosis of a particular condition is of fundamental  
 15 importance, over and above its general significance in relation to specific  
 16 interventions. Areas where this is particularly likely to occur relate to  
 17 assessment of risk, for example in terms of behaviour modification or  
 18 screening and early intervention. In addition, review questions related to  
 19 issues of service delivery are occasionally specified in the remit from the  
 20 Department of Health/Welsh Assembly Government. In these cases,  
 21 appropriate review questions were developed to be clear and concise.  
 22

23 To help facilitate the literature review, a note was made of the best study  
 24 design type to answer each question. There are four main types of review  
 25 question of relevance to NICE guidelines. These are listed in Table 5. For each  
 26 type of question, the best primary study design varies, where 'best' is  
 27 interpreted as 'least likely to give misleading answers to the question'.  
 28

29 However, in all cases, a well-conducted systematic review (of the appropriate  
 30 type of study) is likely to always yield a better answer than a single study.

1  
2 Deciding on the best design type to answer a specific review question does  
3 not mean that studies of different design types addressing the same question  
4 were discarded.

5

**Table 5: Best study design to answer each type of question**

Type of question	Best primary study design
Effectiveness or other impact of an intervention	Randomised controlled trial (RCT); other studies that may be considered in the absence of RCTs are the following: internally/externally controlled before and after trial, interrupted time-series
Accuracy of information (for example, risk factor, test, prediction rule)	Comparing the information against a valid gold standard in a randomised trial or inception cohort study
Rates (of disease, patient experience, rare side effects)	Prospective cohort, registry, cross-sectional study
Costs	Naturalistic prospective cost study

6

## 7 **3.5 SYSTEMATIC CLINICAL LITERATURE REVIEW**

8 The aim of the clinical literature review was to systematically identify and  
9 synthesise relevant evidence from the literature in order to answer the specific  
10 review questions developed by the GDG. Thus, clinical practice  
11 recommendations are evidence-based, where possible, and, if evidence is not  
12 available, informal consensus methods are used (see Section 3.5.6) and the  
13 need for future research is specified.

### 14 **3.5.1 Methodology**

15 A stepwise, hierarchical approach was taken to locating and presenting  
16 evidence to the GDG. The NCCMH developed this process based on methods  
17 set out by NICE (*The Guidelines Manual* [NICE, 2009b]), and after considering  
18 recommendations from a range of other sources. These included:

19

- 20 • Clinical Policy and Practice Program of the New South Wales  
21 Department of Health (Australia)
- 22 • *BMJ Clinical Evidence*
- 23 • Grading of Recommendations: Assessment, Development and  
24 Evaluation (GRADE) Working Group

- 1       • New Zealand Guidelines Group
- 2       • NHS Centre for Reviews and Dissemination
- 3       • Oxford Centre for Evidence-Based Medicine
- 4       • Oxford Systematic Review Development Programme
- 5       • Scottish Intercollegiate Guidelines Network (SIGN)
- 6       • The Cochrane Collaboration
- 7       • United States Agency for Healthcare Research and Quality.

### 8   **3.5.2 The review process**

#### 9   *Scoping searches*

10 A broad preliminary search of the literature was undertaken in January 2009  
11 to obtain an overview of the issues likely to be covered by the scope, and to  
12 help define key areas. Searches were restricted to clinical guidelines, health  
13 technology assessment reports, key systematic reviews and randomised  
14 controlled trials (RCTs), and conducted in the following databases and  
15 websites:

- 16
- 17       • BMJ Clinical Evidence
- 18       • Canadian Medical Association (CMA) Infobase [Canadian  
19        guidelines]
- 20       • Clinical Policy and Practice Program of the New South Wales  
21        Department of Health (Australia)
- 22       • Clinical Practice Guidelines [Australian Guidelines]
- 23       • Cochrane Central Register of Controlled Trials (CENTRAL)
- 24       • Cochrane Database of Abstracts of Reviews of Effects (DARE)
- 25       • Cochrane Database of Systematic Reviews (CDSR)
- 26       • EMBASE
- 27       • Guidelines International Network (G-I-N)
- 28       • Health Evidence Bulletin Wales

- 1       • Health Management Information Consortium [HMIC]
- 2       • Health Technology Assessment (HTA) database (technology
- 3       assessments)
- 4       • MEDLINE / MEDLINE in Process
- 5       • National Health and Medical Research Council (NHMRC)
- 6       • National Library for Health (NLH) Guidelines Finder
- 7       • New Zealand Guidelines Group
- 8       • NHS Centre for Reviews and Dissemination (CRD)
- 9       • OMNI Medical Search
- 10      • Scottish Intercollegiate Guidelines Network (SIGN)
- 11      • Turning Research Into Practice (TRIP)
- 12      • United States Agency for Healthcare Research and Quality (AHRQ)
- 13      • Websites of NICE and the National Institute for Health Research
- 14      (NIHR) HTA Programme for guidelines and HTAs in development.

15 Existing NICE guidelines were updated where necessary. Other relevant  
16 guidelines were assessed for quality using the AGREE instrument (AGREE  
17 Collaboration, 2003). The evidence base underlying high-quality existing  
18 guidelines was utilised and updated as appropriate. Further information  
19 about this process can be found in The Guidelines Manual (NICE, 2009b).

#### 20 *Systematic literature searches*

21 After the scope was finalised, a systematic search strategy was developed to  
22 locate all the relevant evidence. The balance between sensitivity (the power to  
23 identify all studies on a particular topic) and specificity (the ability to exclude  
24 irrelevant studies from the results) was carefully considered, and a decision  
25 made to develop highly sensitive strategies to identify as complete a set as  
26 possible of clinically relevant studies.

27

28 Searches were conducted in the following databases:

29

- 30       • CINAHL
- 31       • EMBASE

- 1       • MEDLINE / MEDLINE In-Process
- 2       • PsycINFO
- 3       • Cochrane Central Register of Controlled Trials (CENTRAL)

4•

5 The search strategies were initially developed for Medline before being  
6 translated for use in other databases/interfaces. Strategies were built up  
7 through a number of trial searches, and discussions of the results of the  
8 searches with the review team and GDG to ensure that all possible relevant  
9 search terms were covered. In order to assure comprehensive coverage,  
10 search terms for psychosis with substance misuse were kept purposely broad  
11 to help counter dissimilarities in database indexing practices and thesaurus  
12 terms, and imprecise reporting of study populations by authors in the titles  
13 and abstracts of records. Search terms for substance misuse were limited to  
14 the main drugs associated with the term at the advice of the GDG. The search  
15 terms for each Medline search are set out in full in Appendix 7.

16 *Reference Manager*

17 Citations from each search were downloaded into Reference Manager (a  
18 software product for managing references and formatting bibliographies) and  
19 duplicates removed. Records were then screened against the inclusion criteria  
20 of the reviews before being quality appraised (see below). The unfiltered  
21 search results were saved and retained for future potential re-analysis to help  
22 keep the process both replicable and transparent.

23 *Search filters*

24 To aid retrieval of relevant and sound studies, filters were used to limit a  
25 number of searches to randomised controlled trials, observational studies and  
26 qualitative research. The randomised controlled trial filter is an adaptation of  
27 a filter designed by the Centre for Reviews and Dissemination (CRD) and the  
28 Health Information Research Unit of McMaster University, Ontario. The  
29 observational studies filter and qualitative research filter were developed in-  
30 house. Each filter comprises index terms relating to the study type(s) and  
31 associated text words for the methodological description of the design(s).

32 *Date and language restrictions*

33 Systematic database searches were initially conducted in July 2009 up to the  
34 most recent searchable date. Search updates were generated on a 6-monthly  
35 basis, with the final re-runs carried out in May 2010 ahead of the guideline  
36 consultation. After this point, studies were only included if they were judged

1 to be exceptional by the GDG (for example, if the evidence was likely to  
2 change a recommendation).

3

4 Although no language restrictions were applied at the searching stage,  
5 foreign language papers were not requested or reviewed, unless they were of  
6 particular importance to a review question. Date restrictions were applied for  
7 searches for qualitative research for the period from 1995 onwards, and for  
8 updates of published reviews. No date restrictions were imposed for the  
9 remainder of the searches.

### 10 *Other search methods*

11 Other search methods involved: 1) scanning the reference lists of all eligible  
12 publications (systematic reviews, stakeholder evidence and included studies)  
13 for more published reports and citations of unpublished research; 2) sending  
14 lists of studies meeting the inclusion criteria to subject experts (identified  
15 through searches and the GDG) and asking them to check the lists for  
16 completeness, and to provide information of any published or unpublished  
17 research for consideration (See Appendix 5); 3) checking the tables of contents  
18 of key journals for studies that might have been missed by the database and  
19 reference list searches; 4) tracking key papers in the Science Citation Index  
20 (prospectively) over time for further useful references.

21 Full details of the Medline search strategies/filters used for the systematic  
22 review of clinical evidence are provided in Appendix 7.

### 23 *Study selection and quality assessment*

24 All primary-level studies included after the first scan of citations were  
25 acquired in full and re-evaluated for eligibility at the time they were being  
26 entered into the study information database. More specific eligibility criteria  
27 were developed for each review question and are described in the relevant  
28 clinical evidence chapters. Eligible systematic reviews and primary-level  
29 studies were critically appraised for methodological quality (see Appendix 10  
30 for methodology checklists). The eligibility of each study was confirmed by at  
31 least one member of the GDG.

32

33 For some review questions, it was necessary to prioritise the evidence with  
34 respect to the UK context (that is, external validity). To make this process  
35 explicit, the GDG took into account the following factors when assessing the  
36 evidence:

37

- 38 • participant factors (for example, gender, age and ethnicity)

- 1           • provider factors (for example, model fidelity, the conditions under  
2           which the intervention was performed and the availability of  
3           experienced staff to undertake the procedure)
  
- 4           • cultural factors (for example, differences in standard care and  
5           differences in the welfare system).

6  
7 The GDG decided which prioritisation factors were relevant to each review  
8 question in light of the UK context and then decided how to modify  
9 recommendations. In each case where this was done, further detail can be  
10 found in the relevant evidence to recommendations section.

### 11 *Unpublished evidence*

12 The GDG used a number of criteria when deciding whether or not to accept  
13 unpublished data. First, the evidence must have been accompanied by a trial  
14 report containing sufficient detail to properly assess the quality of the data.  
15 Second, the evidence must have been submitted with the understanding that  
16 data from the study and a summary of the study's characteristics would be  
17 published in the full guideline. Therefore, the GDG did not accept evidence  
18 submitted as commercial in confidence. However, the GDG recognised that  
19 unpublished evidence submitted by investigators might later be retracted by  
20 those investigators if the inclusion of such data would jeopardise publication  
21 of their research.

### 22 **3.5.3 Data extraction**

23 Study characteristics and outcome data were extracted from all eligible  
24 studies, which met the minimum quality criteria, using Review Manager 5  
25 (The Cochrane Collaboration, 2008).

26  
27 In most circumstances, for a given outcome (continuous and dichotomous),  
28 where more than 50% of the number randomised to any group were lost to  
29 follow up, the data were excluded from the analysis (except for the outcome  
30 'leaving the study early', in which case, the denominator was the number  
31 randomised). Where possible, dichotomous efficacy outcomes were calculated  
32 on an intention-to-treat basis (that is, a 'once-randomised-always-analyse'  
33 basis). Where the GDG advised that those participants who ceased to engage  
34 in the study were likely to have an unfavourable outcome, early withdrawals  
35 were included in both the numerator and denominator. For example, for the  
36 outcome of relapse of psychotic symptoms, in studies that did not use an ITT  
37 analysis, we counted participants who left the study early as relapsing..  
38 Adverse effects were entered into Review Manager as reported by the study  
39 authors because it is usually not possible to determine whether early

1 withdrawals had an unfavourable outcome. Where there was limited data for  
2 a particular review, the 50% rule was not applied. In these circumstances the  
3 evidence was downgraded due to the risk of bias.

4

5 Consultation with another reviewer or members of the GDG was used to  
6 overcome difficulties with coding. Data from studies included in existing  
7 systematic reviews were extracted independently by one reviewer and cross-  
8 checked with the existing data set. Where possible, two independent  
9 reviewers extracted data from new studies. Where double data extraction was  
10 not possible, data extracted by one reviewer was checked by the second  
11 reviewer. Disagreements were resolved through discussion. Where consensus  
12 could not be reached, a third reviewer or GDG members resolved the  
13 disagreement. Masked assessment (that is, blind to the journal from which the  
14 article comes, the authors, the institution and the magnitude of the effect) was  
15 not used since it is unclear that doing so reduces bias (Berlin, 2001; Jadad *et*  
16 *al.*, 1996).

17

## 1 3.5.4 Synthesising the evidence

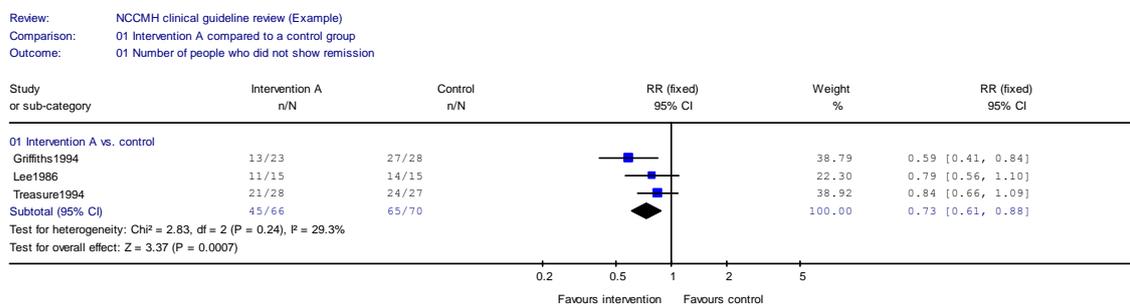
### 2 *Meta-analysis*

3 Where possible, meta-analysis based on a random-effects model  
 4 (DerSimonian & Laird, 1986) was used to synthesise the evidence using  
 5 Review Manager. If necessary, reanalyses of the data or sub-analyses were  
 6 used to answer review questions not addressed in the original studies or  
 7 reviews.

8  
 9 Dichotomous outcomes were analysed as relative risks (RR) with the  
 10 associated 95% CI (for an example, see Figure 1). A relative risk (also called a  
 11 risk ratio) is the ratio of the treatment event rate to the control event rate. An  
 12 RR of 1 indicates no difference between treatment and control. In Figure 1, the  
 13 overall RR of 0.73 indicates that the event rate (that is, non-remission rate)  
 14 associated with intervention A is about three quarters of that with the control  
 15 intervention or, in other words, the relative risk reduction is 27%.

16  
 17 The CI shows a range of values within which we are 95% confident that the  
 18 true effect will lie. If the effect size has a CI that does not cross the 'line of no  
 19 effect', then the effect is commonly interpreted as being statistically  
 20 significant.

21



22

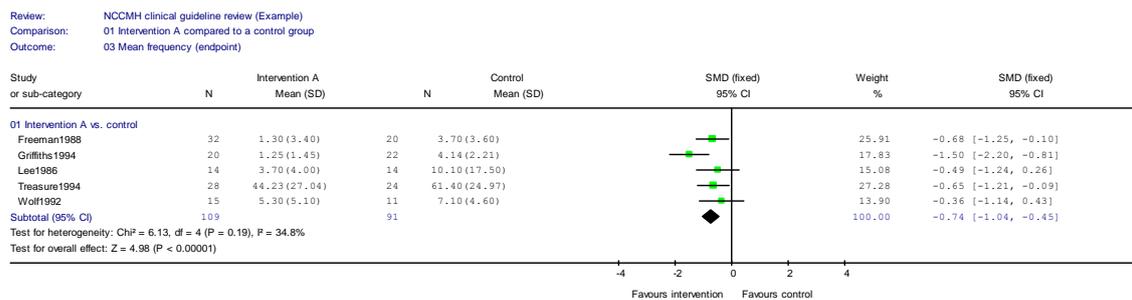
23

24 **Figure 1: Example of a forest plot displaying dichotomous data.**

25 Continuous outcomes were analysed using the mean difference (MD), or  
 26 standardised mean difference (SMD) when different measures were used in  
 27 different studies to estimate the same underlying effect (for an example, see  
 28 Figure 2). If reported by study authors, intention-to-treat data, using a valid  
 29 method for imputation of missing data, were preferred over data only from  
 30 people who completed the study.

31

32

1  
23 **Figure 2: Example of a forest plot displaying continuous data.**4 ***Heterogeneity***

5 To check for consistency of effects among studies, both the  $I^2$  statistic and the  
 6 chi-squared test of heterogeneity, as well as a visual inspection of the forest  
 7 plots were used. The  $I^2$  statistic describes the proportion of total variation in  
 8 study estimates that is due to heterogeneity (Higgins & Thompson, 2002). The  
 9  $I^2$  statistic was interpreted in the following way:

10 &gt;50%: notable heterogeneity

11  $\geq 30$  to  $\leq 50\%$ : moderate heterogeneity

12 &lt;30%: mild heterogeneity.

13 Two factors were used to make a judgement about importance of the  
 14 observed value of  $I^2$ : a) the magnitude and direction of effects, and b) the  
 15 strength of evidence for heterogeneity (for example,  $P$  value from the chi-  
 16 squared test, or a confidence interval for  $I^2$ ). Where heterogeneity was judged  
 17 to be important, an attempt was made to explain the variation by conducting  
 18 sub-analyses to examine potential moderators.

19 ***Publication bias***

20 Where there was sufficient data, we intended to use funnel plots to explore  
 21 the possibility of publication bias. Asymmetry of the plot would be taken to  
 22 indicate possible publication bias and investigated further. However, due to a  
 23 paucity of data, funnel plots could not be used.

24 **3.5.5 Presenting the data to the GDG**

25 Study characteristics tables and, where appropriate, forest plots generated  
 26 with Review Manager were presented to the GDG.

27

28 Where meta-analysis was not appropriate and/or possible, the reported  
 29 results from each primary-level study were included in the study  
 30 characteristics table (and where appropriate, in a narrative review).

## 1 *Evidence profile tables*

2 A GRADE<sup>1</sup> evidence profile was used to summarise both the quality of the  
3 evidence and the results of the evidence synthesis (see Table 6 for an example  
4 of an evidence profile). The GRADE approach is based on a sequential  
5 assessment of the quality of evidence, followed by judgment about the  
6 balance between desirable and undesirable effects, and subsequent decision  
7 about the strength of a recommendation.

8

9 For each outcome, quality may be reduced depending on the following  
10 factors:

- 11 • **study design** (randomised trial, observational study, or any other  
12 evidence)
- 13 • **limitations** (based on the quality of individual studies)
- 14 • **inconsistency** (see section 3.5.4 for how consistency was assessed)
- 15 • **indirectness** (that is, how closely the outcome measures,  
16 interventions and participants match those of interest)
- 17 • **imprecision** (based on the confidence interval around the effect  
18 size).

19 For observational studies, the quality may be increased if there is a large  
20 effect, plausible confounding would have changed the effect, or there is  
21 evidence of a dose-response gradient (details would be provided under the  
22 other considerations column). Each evidence profile also included a summary  
23 of the findings: number of service users included in each group, an estimate  
24 of the magnitude of the effect, and the overall quality of the evidence for each  
25 outcome.

---

<sup>1</sup> For further information about GRADE, see [www.gradeworkinggroup.org](http://www.gradeworkinggroup.org)

**Table 6: Example of GRADE evidence profile**

Quality assessment							Summary of findings				
No. of studies	Design	Limitations	Inconsistency	Indirectness	Imprecision	Other	No. of patients		Effect		Quality
							Intervention	Control	Relative (95% CI)	Absolute	
Outcome 1											
6	randomised trials	no serious limitations	no serious inconsistency	no serious indirectness	very serious <sup>1,2</sup>	none	8/191	7/150	RR 0.94 (0.39 to 2.23)	0 fewer per 100 (from 3 fewer to 6 more)	Low
Outcome 2											
3	randomised trials	no serious limitations	no serious inconsistency	no serious indirectness	no serious imprecision	none	120/600	220/450	RR 0.39 (0.23 to 0.65)	30 fewer per 100 (from 17 fewer to 38 fewer)	High
Outcome 3											
3	randomised trials	no serious limitations	serious inconsistency <sup>3</sup>	no serious indirectness	very serious <sup>1,2</sup>	none	83	81	-	MD -3.51 (-11.51 to 4.49)	Very low
Outcome 4											
3	randomised trials	no serious limitations	no serious inconsistency	no serious indirectness	serious <sup>1</sup>	none	88	93	-	SMD -0.26 (-0.50 to -0.03)	Moderate
Outcome 5											
4	randomised trials	no serious limitations	no serious inconsistency	no serious indirectness	very serious <sup>1,2</sup>	none	109	114	-	SMD -0.13 (-0.6 to 0.34)	Low
<sup>1</sup> Optimal information size (for dichotomous outcomes, OIS = 300 events; for continuous outcomes, OIS = 400 participants) not met. <sup>2</sup> The CI includes both 1) no effect and 2) appreciable benefit or appreciable harm. <sup>3</sup> Considerable heterogeneity.											

### 1 **3.5.6 Method used to answer a review question in the absence of** 2 **appropriately designed, high-quality research**

3 In the absence of appropriately designed, high-quality research, or where the  
4 GDG were of the opinion (on the basis of previous searches or their  
5 knowledge of the literature) that there were unlikely to be such evidence, an  
6 informal consensus process was adopted. This process focused on those  
7 questions that the GDG considered a priority.

8  
9 The starting point for the process of informal consensus was that a member of  
10 the GDG used expert opinion about good practice and any relevant papers  
11 identified by GDG members to write a narrative review.

### 12 **3.5.7 Forming the clinical summaries and recommendations**

13 Once the GRADE evidence profiles relating to a particular review question  
14 were completed, summary evidence tables were developed (these tables are  
15 presented in the evidence chapters). Finally, the systematic reviewer in  
16 conjunction with the members of the GDG produced a clinical evidence  
17 summary.

18  
19 After the GRADE profiles and clinical summaries were presented to the GDG,  
20 the associated recommendations were drafted. In making recommendations,  
21 the GDG took into account the trade-off between the benefits and downsides  
22 of treatment as well as other important factors, such as economic  
23 considerations, social value judgements<sup>2</sup>, the requirements to prevent  
24 discrimination and to promote equality<sup>3</sup>, and the group's awareness of  
25 practical issues (Eccles *et al.*, 1998; NICE, 2009b).

26  
27 Finally, to show clearly how the GDG moved from the evidence to the  
28 recommendations, each chapter has a section called 'from evidence to  
29 recommendations'. Underpinning this section is the concept of the 'strength'  
30 of a recommendation (Schunemann *et al.*, 2003). This takes into account the  
31 quality of the evidence but is conceptually different. Some recommendations  
32 are 'strong' in that the GDG believes that the vast majority of healthcare  
33 professionals and service users would choose a particular intervention if they

---

<sup>2</sup> See NICE's Social Value Judgements: Principles for the Development of  
NICE Guidance:

[www.nice.org.uk/aboutnice/howwework/socialvaluejudgements/socialvaluejudgements.jsp](http://www.nice.org.uk/aboutnice/howwework/socialvaluejudgements/socialvaluejudgements.jsp)

<sup>3</sup> See NICE's equality scheme:

[www.nice.org.uk/aboutnice/howwework/NICEEqualityScheme.jsp](http://www.nice.org.uk/aboutnice/howwework/NICEEqualityScheme.jsp)

1 considered the evidence in the same way that the GDG has. This is generally  
2 the case if the benefits clearly outweigh the harms for most people and the  
3 intervention is likely to be cost effective. However, there is often a closer  
4 balance between benefits and harms, and some service users would not  
5 choose an intervention whereas others would. This may happen, for example,  
6 if some service users are particularly averse to some side effect and others are  
7 not. In these circumstances the recommendation is generally weaker,  
8 although it may be possible to make stronger recommendations about specific  
9 groups of service users. The strength of each recommendation is reflected in  
10 the wording of the recommendation, rather than by using labels or symbols.

11  
12 Where the GDG identified areas in which there are uncertainties or where  
13 robust evidence was lacking, they developed research recommendations.  
14 Those that were identified as 'high-priority' were included in the NICE  
15 version of the guideline, and in Appendix 12.  
16

### 17 **3.6 HEALTH ECONOMICS METHODS**

18 The role of the health economist was to contribute to the guideline's  
19 development by providing evidence on the cost-effectiveness of interventions  
20 covered in this guideline. This was achieved by:

- 21 • Systematic literature review of existing economic evidence
- 22 • Economic modelling, where economic evidence was lacking or was  
23 considered inadequate to inform decisions.

24  
25 Systematic reviews of economic literature were conducted in all areas covered  
26 in the guideline. Economic modelling was planned in areas with potentially  
27 major resource implications, where the current extent of uncertainty over  
28 cost-effectiveness was significant and economic analysis was expected to  
29 reduce this uncertainty, in accordance with the NICE guidelines manual  
30 (NICE, 2009b). Prioritisation of areas for economic modelling was a joint  
31 decision between the Health Economist and the GDG. The rationale for  
32 prioritising review questions for economic modelling was set out in an  
33 economic plan agreed between NICE, the GDG, the Health Economist and  
34 other members of the technical team. The economic plan is presented in  
35 Appendix 19. The following review questions were selected as key issues that  
36 could potentially be addressed by further economic modelling:  
37

- 1           • Cost-effectiveness of integrated models of care (usually involving  
2           the model of assertive community treatment) in people with  
3           psychosis and coexisting substance misuse
  
- 4           • Cost-effectiveness of specific psychological/psychosocial  
5           interventions (delivered within an integrated service model) in  
6           people with psychosis and coexisting substance misuse including:
  - 7               - individual interventions
  - 8               - group interventions
  - 9               - family interventions
  - 10              - contingency management
  - 11              - residential treatment (with/without recovery model)
  - 12              - combined interventions.

13  
14 In addition, literature on the health-related quality of life of people with  
15 psychosis and coexisting substance misuse was systematically searched to  
16 identify studies reporting appropriate health state utility scores that could be  
17 used in potential cost-utility analysis.

18  
19 The rest of this section describes the methods adopted in the systematic  
20 literature review of health economics studies. Methods employed in any  
21 economic modelling undertaken are described in the respective sections of the  
22 guideline.

### 23 **3.6.1 Search strategy for economic evidence**

#### 24 *Scoping searches*

25 A broad preliminary search of the literature was undertaken in January 2009  
26 to obtain an overview of the issues likely to be covered by the scope, and help  
27 define key areas. Searches were restricted to economic studies and health  
28 technology assessment reports, and conducted in the following databases:

- 29
- 30           • EMBASE
- 31           • MEDLINE / MEDLINE In-Process
- 32           • Health Technology Assessment (HTA) database (technology  
33           assessments)
- 34           • NHS Economic Evaluation Database (NHS EED).

1 \* Any relevant economic evidence arising from the clinical scoping searches  
2 was also made available to the health economist during the same period.

3 *Systematic literature searches*

4 After the scope was finalised, a systematic search strategy was developed to  
5 locate all the relevant evidence. The balance between sensitivity (the power to  
6 identify all studies on a particular topic) and specificity (the ability to exclude  
7 irrelevant studies from the results) was carefully considered, and a decision  
8 made to utilise a broad approach to searching to maximise retrieval of  
9 evidence to all parts of the guideline. Searches were restricted to economic  
10 studies and health technology assessment reports, and conducted in the  
11 following databases:

12

- 13 • CINAHL
- 14 • EconLit
- 15 • EMBASE
- 16 • MEDLINE / MEDLINE In-Process
- 17 • PsycINFO
- 18 • Health Technology Assessment (HTA) database (technology  
19 assessments)
- 20 • NHS Economic Evaluation Database (NHS EED).

21

22 \* Any relevant economic evidence arising from the clinical searches was also  
23 made available to the health economist during the same period.

24

25 The search strategies were initially developed for Medline before being  
26 translated for use in other databases/interfaces. Strategies were built up  
27 through a number of trial searches, and discussions of the results of the  
28 searches with the review team and GDG to ensure that all possible relevant  
29 search terms were covered. In order to assure comprehensive coverage,  
30 search terms for psychosis with substance misuse were kept purposely broad  
31 to help counter dissimilarities in database indexing practices and thesaurus  
32 terms, and imprecise reporting of study populations by authors in the titles  
33 and abstracts of records. Search terms for substance misuse were limited to  
34 the main drugs associated with the term at the advice of the GDG.

35

1 For standard mainstream bibliographic databases (CINAHL, EMBASE,  
2 MEDLINE and PsycINFO) search terms for psychosis and substance misuse  
3 were combined with a search filter for health economic studies. For searches  
4 generated in topic-specific databases (EconLit, HTA, NHS EED) search terms  
5 for psychosis and substance abuse were used without a filter. The sensitivity  
6 of this approach was aimed at minimising the risk of overlooking relevant  
7 publications, due to potential weaknesses resulting from more focused search  
8 strategies. The Medline search terms are set out in full in Appendix 9.

### 9 *Reference Manager*

10 Citations from each search were downloaded into Reference Manager (a  
11 software product for managing references and formatting bibliographies) and  
12 duplicates removed. Records were then screened against the inclusion criteria  
13 of the reviews before being quality appraised. The unfiltered search results  
14 were saved and retained for future potential re-analysis to help keep the  
15 process both replicable and transparent.

### 16 *Search filters*

17 The search filter for health economics is an adaptation of a filter designed by  
18 Centre for Reviews and Dissemination (CRD). The filter comprises a  
19 combination of controlled vocabulary and free-text retrieval methods.

### 20 *Date and language restrictions*

21 Systematic database searches were initially conducted in July 2009 up to the  
22 most recent searchable date. Search updates were generated on a 6-monthly  
23 basis, with the final re-runs carried out in May 2010 ahead of the guideline  
24 consultation. After this point, studies were included only if they were judged  
25 by the GDG to be exceptional (for example, the evidence was likely to change  
26 a recommendation).

27  
28 Although no language restrictions were applied at the searching stage,  
29 foreign language papers were not requested or reviewed, unless they were of  
30 particular importance to an area under review. All the searches were  
31 restricted to research published from 1994 onwards in order to obtain data  
32 relevant to current healthcare settings and costs.

### 33 *Other search methods*

34 Other search methods involved scanning the reference lists of all eligible  
35 publications (systematic reviews, stakeholder evidence and included studies  
36 from the economic and clinical reviews) to identify further studies for  
37 consideration.

38

1 Full details of the Medline search strategies/filter used for the systematic  
2 review of health economic evidence are provided in Appendix 9.

### 3 **3.6.2 Inclusion criteria for economic studies**

4 The following inclusion criteria were applied to select studies identified by  
5 the economic searches for further consideration:

- 6       • No restriction was placed on language or publication status of the  
7       papers.
- 8       • Studies published from 1996 onwards were included. This date  
9       restriction was imposed in order to obtain data relevant to current  
10      healthcare settings and costs.
- 11      • Only studies from Organisation for Economic Co-operation and  
12      Development countries were included, as the aim of the review was  
13      to identify economic information transferable to the UK context.
- 14      • Selection criteria based on types of clinical conditions and service  
15      users as well as interventions assessed were identical to the clinical  
16      literature review.
- 17      • Studies were included provided that sufficient details regarding  
18      methods and results were available to enable the methodological  
19      quality of the study to be assessed, and provided that the study's  
20      data and results were extractable.
- 21      • Full economic evaluations that compared two or more relevant  
22      options and considered both costs and consequences (that is, cost-  
23      consequence analysis, cost-effectiveness analysis, cost-utility  
24      analysis or cost-benefit analysis), as well as costing analyses that  
25      compared only costs between two or more interventions, were  
26      included in the review.
- 27      • Economic studies were included if they used clinical effectiveness  
28      data from an RCT, a cohort study, or a systematic review and meta-  
29      analysis of clinical studies. Studies that had a mirror-image design  
30      were excluded from the review.
- 31      • Studies were included only if the examined interventions were  
32      clearly described. This involved the dosage and route of  
33      administration and the duration of treatment in the case of  
34      pharmacological therapies; and the types of health professionals  
35      involved as well as the frequency and duration of treatment in the

1 case of psychological interventions. Evaluations in which  
2 medications were treated as a class were excluded from further  
3 consideration.

### 4 **3.6.3 Applicability and quality criteria for economic studies**

5 All economic papers eligible for inclusion were appraised for their  
6 applicability and quality using the methodology checklist for economic  
7 evaluations recommended by NICE (NICE, 2009b), which is shown in  
8 Appendix 18 of this guideline. The methodology checklist for economic  
9 evaluations was also applied to the economic models developed specifically  
10 for this guideline. All studies that fully or partially met the applicability and  
11 quality criteria described in the methodology checklist were considered  
12 during the guideline development process, along with the results of the  
13 economic modelling conducted specifically for this guideline. The completed  
14 methodology checklists for all economic evaluations considered in the  
15 guideline are provided in Appendix 18.

### 16 **3.6.4 Presentation of economic evidence**

17 The economic evidence considered in the guideline is provided in the  
18 respective evidence chapters, following presentation of the relevant clinical  
19 evidence. The references to included studies as well as the evidence tables  
20 with the characteristics and results of economic studies included in the  
21 review, are provided in Appendix 17. Methods and results of any economic  
22 modelling undertaken alongside the guideline development process are  
23 presented in the relevant evidence chapters. Characteristics and results of all  
24 economic studies considered during the guideline development process are  
25 summarised in economic evidence profiles accompanying respective GRADE  
26 clinical evidence profiles in Appendix 17.

### 27 **3.6.5 Results of the systematic search of economic literature**

28 The titles of all studies identified by the systematic search of the literature  
29 were screened for their relevance to the topic (i.e. consideration of health  
30 economics issues and health-related quality of life in people with psychosis  
31 and coexisting substance misuse). References that were clearly not relevant  
32 were excluded first. The abstracts of all potentially relevant publications (82  
33 references) were then assessed against the inclusion criteria for economic  
34 evaluations by the health economist. Full texts of the studies potentially  
35 meeting the inclusion criteria (including those for which eligibility was not  
36 clear from the abstract) were obtained. Studies that did not meet the inclusion  
37 criteria, were duplicates, secondary publications of one study, or had been  
38 updated in more recent publications were subsequently excluded. Overall, six  
39 economic evaluations were identified as being eligible for inclusion and were

1 appraised for their applicability and quality using the methodology checklist  
2 for economic evaluations. The findings of these studies were considered when  
3 formulating the guideline recommendations.  
4

### 5 **3.7 STAKEHOLDER CONTRIBUTIONS**

6 Professionals, service users, and companies have contributed to and  
7 commented on the guideline at key stages in its development. Stakeholders  
8 for this guideline include:  
9

- 10 • patient and carer stakeholders: national patient and carer  
11 organisations that represent the interests of people whose care will  
12 be covered by the guideline
- 13 • local patient and carer organisations: but only if there is no relevant  
14 national organisation
- 15 • professional stakeholders' national organisations: that represent the  
16 healthcare professionals who provide the services described in the  
17 guideline
- 18 • commercial stakeholders: companies that manufacture drugs or  
19 devices used in treatment of the condition covered by the guideline  
20 and whose interests may be significantly affected by the guideline
- 21 • providers and commissioners of health services in England and  
22 Wales
- 23 • statutory organisations: including the Department of Health, the  
24 Welsh Assembly
- 25 • Government, NHS Quality Improvement Scotland, the Healthcare  
26 Commission and the National Patient Safety Agency
- 27 • research organisations: that have carried out nationally recognised  
28 research in the area.

29  
30 NICE clinical guidelines are produced for the NHS in England and Wales, so  
31 a 'national' organisation is defined as one that represents England and/or  
32 Wales, or has a commercial interest in England and/or Wales.  
33 Stakeholders have been involved in the guideline's development at the  
34 following points:

1  
2  
3  
4  
5  
6  
7

- commenting on the initial scope of the guideline and attending a scoping workshop held by NICE
- contributing possible review questions and lists of evidence to the GDG
- commenting on the draft of the guideline
- highlighting factual errors in the pre-publication check.

### 8 **3.8 VALIDATION OF THE GUIDELINE**

9 Registered stakeholders had an opportunity to comment on the draft  
10 guideline, which was posted on the NICE website during the consultation  
11 period. Following the consultation, all comments from stakeholders and  
12 others were responded to, and the guideline updated as appropriate. The  
13 GRP also reviewed the guideline and checked that stakeholders' comments  
14 had been addressed.

15  
16 Following the consultation period, the GDG finalised the recommendations  
17 and the NCCMH produced the final documents. These were then submitted  
18 to NICE for the pre-publication check where stakeholders are given the  
19 opportunity to highlight factual errors. Any errors are corrected by the  
20 NCCMH, then the guideline is formally approved by NICE and issued as  
21 guidance to the NHS in England and Wales.  
22

23

# 1 **4 EXPERIENCE OF CARE**

## 2 **4.1 INTRODUCTION**

3 This chapter provides an overview of the experience of people with psychosis  
4 and coexisting substance misuse, and the experience of their families/carers.  
5 The first two sections present first-hand personal accounts written by people  
6 with psychosis and coexisting substance misuse, and their families and carers.  
7 These sections provide an insight into the experience of being diagnosed,  
8 accessing services, receiving treatment and caring for someone with psychosis  
9 and coexisting substance misuse. It should be noted that these accounts of the  
10 experience of people with psychosis and coexisting substance misuse are  
11 illustrative. The third section is a qualitative analysis of transcripts of people  
12 with psychosis and coexisting substance misuse from seven online websites  
13 and a review of the qualitative literature of the experience of people with  
14 psychosis and coexisting substance misuse. Following this is a summary of  
15 the themes emerging from the personal accounts, the online transcripts and  
16 the literature review which provides a basis for the associated  
17 recommendations.  
18

## 19 **4.2 PERSONAL ACCOUNTS**

### 20 **4.2.1 Introduction**

21 The writers of the personal accounts from people with psychosis and  
22 coexisting substance misuse were contacted through representatives on the  
23 GDG and through various agencies that had access to people with psychosis  
24 and coexisting substance misuse. The people who were approached to write  
25 the accounts were asked to consider a number of questions when composing  
26 their narratives. These included:

- 27 • When did you first seek help for your psychosis and coexisting  
28 substance misuse and whom did you contact? Please describe this  
29 first contact.
- 30 • What helped or did not help you gain access to services? Did a  
31 friend or family member help you gain access to these services?
- 32 • Do you think that any life experiences led to the onset of the  
33 problem? If so, please describe if you feel able to do so.

- 1           • In what ways has psychosis and substance misuse affected your  
2 everyday life (such as education, employment and making  
3 relationships) and the lives of those close to you?
- 4           • What possible treatments were discussed with you?
- 5           • What treatment(s) did you receive? Please describe any drug  
6 treatment and/or psychological therapy.
- 7           • Was the treatment(s) helpful? Please describe what worked for you  
8 and what didn't work for you.
- 9           • How would you describe your relationship with your  
10 practitioner(s) (for example, your GP, psychologist or other)
- 11          • Did you use any other approaches to help your psychosis and  
12 substance misuse in addition to those provided by NHS services, for  
13 example private treatment? If so please describe what was helpful  
14 and not helpful.
- 15          • Do you have any language support needs, including needing help  
16 with reading or speaking English? If so, did this have an impact on  
17 your understanding of the psychosis and substance misuse or on  
18 receiving treatment?
- 19          • Did you attend a support group and was this helpful? Did family  
20 and friends close to you or people in your community help and  
21 support you?
- 22          • How has the nature of the problem changed over time?
- 23          • How do you feel now?
- 24          • If your psychosis and coexisting substance misuse has improved,  
25 do you use any strategies to help you to stay well? If so, please  
26 describe these strategies.

27 Each author signed a consent form allowing the account to be reproduced in  
28 this guideline. Two personal accounts from people (both male) with psychosis  
29 and coexisting substance misuse were received in total. They offer different  
30 perspectives of their experience of illness and treatment, but despite the  
31 differences some common themes do emerge. Each person speaks of the  
32 isolation he felt at various stages of his illness and treatment and the  
33 challenges in finding employment after a long period out of work. In terms of  
34 treatment, the service users valued staff who were 'empathic', 'helpful',

1 'motivated' and 'keen', and understood mental health and substance misuse  
2 issues. Lack of planned care, gaps in their treatment and treatment being  
3 stopped abruptly (especially for the person being released from prison) were  
4 deemed unhelpful.

5  
6 The service users identified a range of helpful and unhelpful treatments.  
7 Person A found that in prison CBT, group work, and creative and educative  
8 activities were helpful and, out of prison, his local alcohol service provided  
9 support better suited to him than Alcoholics Anonymous; self-help (delivered  
10 in prison) was deemed to be unhelpful because the service user felt it was not  
11 properly explained to him. Person B was very positive about the treatment he  
12 received from his dual diagnosis practitioner which included writing a drug  
13 diary and a feelings notebook, and identifying and managing the risks and  
14 triggers.

15  
16 Both men identified that support from assertive outreach teams and other  
17 workers to enable them to re-enter society and find employment (either paid  
18 or voluntary) was vital in building self-esteem and restoring confidence.

#### 19 **4.2.2 Personal account A**

20 I was born in 1961 in London, and my parents came from Jamaica. I had a  
21 very successful career until 2003. At this time I would go days without sleep,  
22 having detailed nightmares, hallucinations and I wouldn't go out in the day  
23 time or answer my phone. As time went on my mood swings got worse and I  
24 had no control over them. I thought the world was against me and everyone  
25 wanted to do me harm.

26  
27 I was drinking a lot and socially smoking weed. I lost my job, wife, family and  
28 home in 2004 and ended up in prison. In 2005, I was diagnosed with severe  
29 depression and personality disorder with agoraphobic, paranoid and  
30 psychotic features by a clinical psychiatrist.

31  
32 In August 2005, I was arrested and remanded in custody. My lawyer had a  
33 good understanding of the prison system and talked me through the booking  
34 in process and what was best to say and do. At my booking in, I advised them  
35 of my mental health and all of my issues. I was interviewed the next day and I  
36 was told that the services I needed would be provided as soon as possible.

37  
38 The doctor gave me four sleeping tablets (one per night) to keep me stable  
39 until I could see the CMHT. The staff that I met in the first 48 hours showed  
40 empathy and concern about my well-being, but the service provided didn't  
41 always live up to their promises. The action plan was good, and the full-time

1 staff were helpful, motivated and keen, but the specialist team of a clinical  
2 psychologist, psychiatrist and counsellor didn't keep their appointments and  
3 this led to me having relapses in my mental health. On a couple of occasions,  
4 the staff forgot to open my cell door or were late in doing so and I missed my  
5 appointment. To address this problem, I was given stronger medication or  
6 larger doses. I never missed taking my medication because if you did you  
7 were escorted to the nurse and your mouth was checked after.

8  
9 I took olanzapine and diazepam daily, and if I was having a bad night I might  
10 get temazepam to help me sleep. I was offered lots of meaningful activities to  
11 do during the day, such as focus groups, arts and crafts, games and  
12 education. This did keep my mind occupied and help me feel better. I was  
13 also taught CBT and I started self-help treatment but it didn't entirely work  
14 because it wasn't fully explained to me; however it did show me what I could  
15 do to help myself and how to handle my relationship with my family and  
16 friends, and my problems with drink and drugs .

17  
18 One of the good things that came out of my prison stay was when we got the  
19 governor to change the day centre from being located in a mental health unit  
20 to a multicultural mental health day centre. This was my first taste that  
21 service user involvement works.

22  
23 I was released on bail straight from court without any medication and  
24 ordered to stay with my family until my court date. My GP was in another  
25 town so to get treatment I had to lie and say I still lived there. The paperwork  
26 took a while to get to my GP and I was not given any antidepressants, only a  
27 referral to the CMHT and sleeping tablets.

28  
29 On my return to court, the judge gave me probation as long as I followed the  
30 guidelines without fail. These included taking my medication and attending  
31 anger management, literacy and numeracy classes, in addition to attending all  
32 sessions recommended by the CMHT and my probation officer. The CMHT  
33 and my probation officer put together an action plan for me without my  
34 input. Six specialists were assigned to me. Again, the plan was good, but the  
35 services I needed were not available to start at the same time. At first this was  
36 not a problem but as time went by my mental health and drinking issues were  
37 not dealt with – the services looked at what they could provide and not what I  
38 needed. The clinical psychiatrist I saw was very good at her job,  
39 knowledgeable and showed lots of empathy and people skills. However, after  
40 seven sessions she advised me she was going on honeymoon for 6 weeks and  
41 my treatment would be put on hold until her return. Again, as I was making  
42 progress, my treatment was put on hold. I had to rely on the CBT I had been

1 taught in prison, and on drink and pills to get though any crisis I may come  
2 across.

3

4 I had to use drink to get though the hard days; by the time, I got help for my  
5 drinking it had become a bigger problem. Alcoholics Anonymous did not  
6 work for me because it was not holistic and I was always very depressed after  
7 AA meetings. I was asked to leave because I wasn't engaging correctly.

8

9 My brother paid for me to have four private sessions with a clinical  
10 psychiatrist, but he was only willing to help develop my CBT and coping  
11 skills. I was referred to Mind for counselling by my GP but failed a risk  
12 assessment (my local Mind only had female staff, small interview rooms and  
13 no security). At this stage of my recovery journey, I got housed by an  
14 organisation for the homeless, and accessed their services. I was given a  
15 keyworker, who was very knowledgeable and showed a lot of empathy and a  
16 willingness to help me address all my issues and support me to reach my  
17 aims and goals. We drew up an action plan together with targets and rewards  
18 for hitting them. We met with my GP and had my medication reduced and  
19 sorted out some meaningful actives for me to do. I had interviews with the  
20 mental health and substance abuse team at the homeless organisation and  
21 was put on their self-help programme; the service provided was excellent and  
22 empowered me to aim higher and believe I could recover. However, just as I  
23 was feeling the benefit and moving on leaps and bounds the service came to  
24 an end due to lack of money.

25

26 I attended my local alcohol counselling services for my drinking problems;  
27 this service suited me better than AA and sorted out my drinking. The  
28 counsellor asked me keep a diary, account for my drinking and look for the  
29 triggers that caused it.

30

31 Then we worked with my keyworker and clinical psychologist to find ways  
32 for me to cope.

33

34 The service provided by the CMHT came to an end because my probation  
35 was up and not because I was ready to rejoin the community or because I had  
36 fully recovered. Ultimately I found the service patchy; it was full of great  
37 intentions but they failed to deliver what they had promised.

38

39 I also attended a programme that helped me to prepare for the moving back  
40 into the community. The homeless organisation's resettlement officer helped  
41 me sort out my housing benefit, got my gas and electricity turned on, and  
42 hired a removal van, a bed, and cooker for me. She also gave me advice on  
43 paying my bills. The system would not give me a community or crisis loan

1 because I was not on Jobseeker's Allowance or Income Support. I only had the  
2 bare minimum in my flat. This did not help my mental health or empower me  
3 to keep on going.

4  
5 Now it was time to look for full-time work. Trying to get employment with a  
6 criminal record and mental health issues was near on impossible. I had a lot  
7 of interviews but even more excuses why people were not employing me. I  
8 was appointed a floating support worker to help me with my move on from  
9 supported housing back into the community. His caseload is large and the  
10 length of time his support will be available to me relies on funding; however,  
11 the service provided was good because he worked in a holistic way, always  
12 returned my calls within 2 hours, kept all of our appointments, treated me as  
13 a person at all times, and provided a professional, honest and reliable service.

14  
15 All the services helped me in different ways but because the services  
16 provided didn't all start at the same time the process was slow and put a lot of  
17 pressure on me and my ability to cope. This led to relapse, binge drinking,  
18 and withdrawal from the community. I think my recovery journey is going  
19 well but I know my hardest tests are still to come.

### 20 **4.2.3 Personal account B**

21 I am 33 years old and have a history of paranoid schizophrenia and substance  
22 misuse.

23  
24 In 1994 after I finished my A levels I started to hang out with the 'trendy  
25 guys' who lived in my town and spent many hours smoking cannabis spliffs  
26 (rolled tobacco cigarettes laced with cannabis resin) and bongs (water pipes  
27 which would cool down the cannabis smoke). In the following autumn, I  
28 went to university. I thought that students should spend most of their time  
29 getting stoned and living the life of a 1960s hippie. That was the plan and  
30 that's what I did. I not only continued to smoke cannabis but also became  
31 experienced with other substances: speed (amphetamine), ecstasy, LSD and  
32 magic mushrooms.

33  
34 Initially, much of my university work was of a high quality. However, as the  
35 year progressed and I became more involved with drugs, I began to feel more  
36 self-conscious about my existence. I would feel uncomfortable walking to the  
37 campus and developed a dread about my course. A feeling of helplessness  
38 and a sort of isolation developed and my academic work began to suffer. I  
39 changed courses the following year – I didn't feel so anxious but I was  
40 smoking one to two ounces of cannabis resin a week – and taking a variety of  
41 other drugs.

1  
2 I finished my degree (with a third class) and found an office job. However, I  
3 found the job tedious and in 1999 decided to do a master's degree. I continued  
4 to use drugs every weekend (ecstasy and cannabis and occasionally cocaine  
5 and magic mushrooms). The amount of cannabis I was using led to lung  
6 problems.

7  
8 During the new year celebrations of 2000, I decided to take about 10 ecstasy  
9 tablets in about 45 minutes. That new year's party may have changed my  
10 whole life. During the next term my tutor was concerned that I had very dull  
11 eyes. I thought nothing of it. Then as the year went on I started thinking that a  
12 DJ was talking to me through the radio and the walls contained mini-  
13 microphones and cameras. My body felt more and more intense, and not in a  
14 good way. My behaviour became more angry and irrational. I accused people  
15 of ridiculous things (for example, I thought that my flatmate had broken into  
16 my room and removed a bit of my printer to stop it working). Nevertheless I  
17 continued to see my old university friends every weekend and my pattern of  
18 drug use continued.

19  
20 I felt uncertain as to what was happening to me. My feelings became more  
21 and more intense. My friends kept telling me that instead of the smiles which  
22 I had initially met them with, I looked angry and depressed. My mood  
23 deteriorated and I became more isolated. I thought that I should get some  
24 help, so I went to the university student services. I got to the front door, felt  
25 very self-conscious and walked away.

26  
27 Despite my continued drug use and deteriorating mental health I completed  
28 my masters degree. I found an interesting job but as I walked through the  
29 factory and heard Radio 4 talking about me, that was it. How would I be able  
30 to do a job well if I thought that a national radio station was talking about  
31 me?

32  
33 I wanted to get treatment but had heard (incorrectly) from a GP that the only  
34 way a doctor in the UK would treat me was if I posed a serious risk to myself  
35 or others and that would mean putting me on a section of the Mental Health  
36 Act.

37  
38 My parents became worried about my mental health and accessed a  
39 neurologist in the United States (which is where we come from). We were  
40 concerned that I might have more than just mental health problems and there  
41 could be some underlying physiological problem. After seeing the neurologist  
42 I was referred on to a psychologist. By the end of it they had identified that I

1 was psychotic and referred me to a psychiatrist who gave me drugs to stop  
2 those symptoms.

3  
4 I returned to England and lived with my parents for about 10 months. My GP  
5 referred me to the local psychiatrist and I accessed a community psychiatric  
6 nurse (CPN) and mental health support worker. My CPN was very helpful  
7 and the support worker helped me get out of the house and do things like  
8 play badminton and have lunch at the seaside. I was in some form of recovery  
9 at this stage but still felt that I was functioning at a much lower level than I  
10 was capable of. I would describe my mental state as 'gormless'. I did not feel  
11 very sharp in my thinking. Looking back I'm not sure if this was a reflection  
12 of my mental state, the medication I was being prescribed, or a combination  
13 of both.

14  
15 Eventually, I acquired some voluntary work, still feeling gormless, but better  
16 able to get things done. This was negotiated through an employment  
17 company for disadvantaged people who were able to persuade them that I  
18 would be an asset to the team. I was assigned a support worker, which  
19 worked out well. I was able to get out of the house and be a part of society at  
20 some level, which was better than staying in, watching telly and eating junk  
21 food on my own. Indeed, I was even provided with a reference, which helped  
22 me get work subsequently.

23  
24 I decide to move to London and find paid work. I knew a guy who was  
25 renting out cheap rooms and I managed to get a job. Initially I was socially  
26 isolated but eventually my old friends from my university days contacted me.  
27 I was glad to have friends again but we were soon back smoking skunk –  
28 about 20 to 30 joints over the weekend. I began to feel 'gormless' again and  
29 my behaviour became weird. I could no longer undertake simple tasks at  
30 work and this along with other things, such as being slightly smelly, being  
31 late to work, spending more time smoking cigarettes than doing the job, led to  
32 my dismissal.

33  
34 Still getting stoned on skunk, I went from one job to the next, each being  
35 progressively worse than the former. I just wasn't able to do my job properly.  
36 Nevertheless, I continued to smoke weed. Soon, I got to the stage where I  
37 would sit at home all day, in my smelly unwashed clothes, eat biscuits for  
38 dinner and defer bill payments.

39  
40 I needed to change my life. My main social contact was a middle-aged artist  
41 who would convince me that I should give him money to buy cannabis. Most  
42 of my friends had moved away and I did not get on very well with my family.  
43 I could not maintain any kind of employment and I had little or no money. I

1 had lost control of my own life and the people who did have control of it were  
2 mostly dealers and 'friends'. I began to get scared just walking down my  
3 road. Every year I would watch my life go no further than the previous one.  
4 And most of all, I was very vulnerable and truly out of control. I wanted my  
5 life back. Desperately.

6  
7 Throughout this period I saw my psychiatrist every 6 months and I would tell  
8 him how smoking weed ruined my chances of having a real life. After 2 or so  
9 years, he put me in touch with a dual diagnosis practitioner. For me, it was  
10 very important to stop using cannabis. I would probably not have been able  
11 do this on my own but by accessing the dual diagnosis service it was much  
12 easier.

13  
14 I met with my dual diagnosis practitioner every 3 weeks. One area of work I  
15 did with her was identify the triggers that stimulated me to smoke spliffs. The  
16 triggers would range from spending time with the artist or my old friends to  
17 watching films alone on television (strong spliffs and funny movies go  
18 together like strawberries and cream for me). We identified that the artist  
19 posed a real danger to my recovery. Every time I stopped smoking weed I  
20 would go and see him and the habit would restart.

21  
22 We also identified that the addiction to cannabis is strong and psychological,  
23 that my brain craves that 'lovely' THC (tetrahydrocannabinol - the chemical  
24 in cannabis which makes the feeling of using so pleasant) and that it would  
25 manipulate me to score by changing my thinking patterns. I would think, 'the  
26 artist has a book that I want back'; that is the THC addiction sending me to  
27 the artist to smoke that crafty spliff. A tool to combat this is to 'know your  
28 enemy'.

29  
30 My dual diagnosis worker helped me to identify and overcome the triggers  
31 and armed me with tools to fight the cravings. One tool I use is to picture  
32 traffic lights. If I want a joint I look at a picture of a traffic light on my wall..  
33 The traffic lights act like a reminder, or a prompt, challenging me to think  
34 about whether I really want this and/or how smoking cannabis affected me in  
35 the past. Red is the first warning. This alerts me to ask myself: Do I really  
36 want to get stoned? Remember your history. Do I want to be that smelly,  
37 unkempt, poor drug user again? Remember that it was hard enough coming  
38 off the weed and would be just as easy to get back onto the 'addiction wagon'.  
39 Yellow is 'well why not, life is pretty bad', like getting sacked from my job  
40 and my family disowning me. Yellow is considering the threat that using  
41 cannabis would have and the consequences which would come from smoking  
42 it. In this case, I may think that there is little else to lose and having a joint  
43 wouldn't hurt. This may be the case, but considering my history of cannabis

1 addiction the threat would be significant. And the bottom line would be 'do I  
2 really want to go through that all over again?' This would refer me back to the  
3 red traffic light. Then there is the green light, which is 'nuclear holocaust'.  
4 Everything that could possibly go wrong has and is getting worse. In that  
5 case, going out, scoring a draw and getting obliterated might not be so bad. I  
6 haven't got to green yet!

7  
8 For about 9 months, the THC addiction was still strong. I felt that by writing  
9 stories and feelings in a notebook, I could manage these very intense feelings,  
10 which included blaming everyone except me for the failures of my life (such  
11 as 'I was poor because my brother introduced me to smoking cannabis'). In  
12 real life, I could not blame anyone for my substance misuse. Often feelings of  
13 social isolation would come out in my notebook. Using cannabis had masked  
14 these feelings and would make me less lonely. Harboursing unpleasant  
15 thoughts and not being able to express them, especially during rehabilitation,  
16 could lead to mental anguish. By writing these thoughts on paper and being  
17 able to look back on them, I felt emotionally liberated. I could release the  
18 mental tension and feel better. It was like popping a blister.

19  
20 I also found that smoking tobacco in 'rollies' was a great substitute for  
21 smoking joints, in terms of the process of preparing the rollies, the act of  
22 smoking, and doing something with my hands. Over time I reduced the  
23 rollies and, recognising the harms tobacco itself can cause, I now smoke one  
24 herbal cigarette a day.

25  
26 I was spending long periods at home watching television and thinking about  
27 how much I would like to smoke a joint and feeling lonely and socially  
28 isolated, so my dual diagnosis practitioner and I identified that activity was  
29 the best way forward. I looked at every possible opportunity to get involved  
30 with as much as possible. I volunteered to do things that interested me. I  
31 considered working as a support worker with people with learning  
32 disabilities or in the office of my housing association, or befriending an old  
33 lady. None of these activities came to much, but just the 'doing' helped to stop  
34 that lonely feeling which comes with social isolation. I felt that involvement  
35 with society would be the best way ahead in terms of recovery from substance  
36 misuse. It would also help me to regain my confidence by proving that I can  
37 do jobs successfully even though I have a history of mental health issues.

38  
39 The changes I have made to my drug use and lifestyle have brought about  
40 wider benefits too. I have re-established good relationships with my family  
41 again and recently spent about a month with them. I am training to be a drugs  
42 worker through work I am involved in at a local substance misuse service. I

1 have also taken part in delivering dual diagnosis training and been a service  
2 user link worker to an acute psychiatric ward.

3  
4 I also run a social club, which is proving to be very successful. It provides hot  
5 meals to people who may have issues with substance misuse, mental health  
6 and/or learning disabilities. We aim to re integrate people with these issues  
7 back into society at their own pace, by providing opportunities such as fun  
8 classes, which may inspire them into mainstream education, or making new  
9 social networks or joining the management committee. From my own  
10 perspective, running this club has enabled me to regain a huge amount of  
11 confidence and I am keen to start these clubs more widely. My vision is for  
12 each club, under the umbrella of the wider social club organisation, to be run  
13 independently –they would choose their own activities and food (within  
14 reason). By providing this responsibility, it may help others in their recovery  
15 journeys.

16  
17 My status has improved, as well as my mental health. Since I have accessed  
18 the dual diagnosis service my medication dose has dropped by 25%. Two  
19 years ago, I was frightened of a 30-minute bus ride to visit my friends but I  
20 am not scared on buses any longer or even walking the streets of London at  
21 night. I have made new friends and these friendships are blossoming. I have  
22 found a new kind of respect for myself and am truly looking forward to a  
23 future without limits.

24  
25 From my point of view, de-stigmatising treatment for mental health is vital to  
26 promoting early diagnosis and recovery. An approachable practitioner who  
27 empathises and understands mental health and substance misuse issues is  
28 also vital. It's important for professionals to plan treatment in conjunction  
29 with the service user, taking account of the person's readiness to change.  
30 Mental health professionals need to maintain an open mind and sense of  
31 optimism about what the service user can achieve, rather than limiting  
32 options through low expectations. This can help to develop the person's self-  
33 esteem. Reducing or stopping substance misuse altogether may reduce  
34 medication doses. When a person is in recovery, social support from the NHS,  
35 family members and other social systems, is crucial. When addressing  
36 substance misuse, tools such as a drug diary, feelings notebook, and traffic  
37 lights, can be useful to enable the person to identify and manage the  
38 risks/triggers. Distraction techniques (such as volunteering and fun classes)  
39 can help them to start rebuilding their lives and returning to work is  
40 important because that is part of the person's identity. Ideally the work  
41 should be something that is suited to the person's skills and/or wishes. It's  
42 important for the service user to feel a sense of achievement and involving  
43 others can help them develop important connections and make new friends.

1

## 2 **4.3 PERSONAL ACCOUNTS – CARERS**

### 3 **4.3.1 Introduction**

4 The methods used for obtaining the carers' accounts were the same as  
5 outlined in section 4.2.1, but the questions included:

- 6 • In what way do you care for someone with psychosis and substance  
7 misuse?
- 8 • How long have you been a carer of someone with psychosis and  
9 substance misuse??
- 10 • In what ways has being a carer affected your everyday life (such as  
11 schooling, employment and making relationships) and the lives of  
12 those close to you?
- 13 • How involved are/were you in the treatment plans of the person  
14 with psychosis and substance misuse??
- 15 • Were you offered support by the person's practitioners (for  
16 example, their GP, psychologist, or other)?
- 17 • How would you describe your relationship with the person's  
18 practitioner(s)?
- 19 • Have you and your family been offered help or received  
20 assessment/treatment by a healthcare professional?
- 21 • Did you attend a support group and was this helpful?
- 22 • Did any people close to you help and support you in your role as a  
23 carer?

24 Three accounts from carers of people with psychosis and coexisting substance  
25 misuse were received, which offer different perspectives of being a carer. Two  
26 of the carers are parents (one mother, one father) and one is a grandmother.  
27 Many of the common themes from the personal accounts are echoed in the  
28 carer accounts, including the lack of continuity of care, which may impact on  
29 carers as well, who have to fill in the gap. The accounts below reveal the  
30 difficulties of caring with someone who has psychosis and coexisting  
31 substance misuse, such as challenging behaviour and, in the case of drug  
32 misuse, contending with the drugs world, including dealers and other users.  
33 All of the carers spoke of providing practical support to their family

1 members, which ranges from helping them with their shopping, taking their  
2 medication, finding appropriate housing and employment, and managing  
3 money and benefits. For carer B a significant financial burden was placed on  
4 the family. As all of the accounts below demonstrate, carers value support  
5 from healthcare professionals and other workers, and appreciate it when they  
6 recognise that they, the carers, have valuable knowledge about their family  
7 member's illness and substance problem which can help adherence to  
8 treatment and prevent relapse. What is clear from the accounts is that carers  
9 have very different individual needs: some may require more support from  
10 healthcare professionals than others, who may prefer to cope within their  
11 family environment, rather than attending support groups. However during a  
12 crisis, all of the carers expressed that they would like to know whom to  
13 contact and to be able to access help quickly.

#### 14 **4.3.2 Carer account A**

15 It is difficult to know where to begin to summarise what it has meant to see  
16 myself as the carer of my son Jack. Did it all begin 20 years ago when, aged 18,  
17 he had the first episode that could be deemed to be psychotic? Or was it much  
18 earlier when he was having difficulties at school and was labelled dyslexic,  
19 although one teacher said that she wondered whether he was a genius?  
20

21 In some ways we were fortunate in being able to pay for him to see  
22 educational psychologists and Jack went through various tests and attended  
23 special schools that were supposed to meet his needs and help to prepare him  
24 for life in the world outside the safety of his family.  
25

26 However, as I discovered much later, some of the boys at his specialist day  
27 school had access to marijuana and what began as a prank led to him self-  
28 medicating because of his worries about not 'fitting' in and not being able to  
29 keep up at school.  
30

31 Jack is the youngest of three siblings and his older brother and sister were  
32 high achievers at school and university and are both married with children.  
33 This has highlighted Jack's feelings of inadequacy and fuelled his anger at  
34 what he feels to be an unfriendly world.  
35

36 In his late teens Jack began experimenting with LSD, which led to his first  
37 admission to a private psychiatric hospital. It soon became apparent that we  
38 would not be able to afford long-term private treatment and he was  
39 transferred to an NHS hospital under the care of the same psychiatrist.  
40

1 The nightmare began. There were times when he seemed quite mad – he grew  
2 his hair and a beard and my beautiful, funny and happy little boy turned into  
3 a frightened and frightening stranger. We went through outpatients, then he  
4 was sectioned and spent a few weeks in one major teaching hospital. The  
5 psychiatrist said to me at the time that there was nothing they could do to  
6 stop people bringing in ‘ganja’, so while heavy medication (haloperidol,  
7 called the ‘liquid cosh’ by the patients) was being administered the patients  
8 were smoking dope on the patios! As I am a psychotherapist and had a lot of  
9 support, I battled the system at that time in which parents were not told  
10 which drugs were being prescribed. This meant that when one’s child was  
11 sent home, the family had no idea of the possible side effects and what to do  
12 about them. We had one terrifying Sunday when Jack went into spasms and  
13 his face and jaw locked until we managed to get the antidote pill through a  
14 private doctor.

15  
16 I became involved in what was then the National Schizophrenia Fellowship  
17 where there was some support and a bit of information for what were mostly  
18 the mothers of children with a similar diagnosis to Jack. By then he was  
19 labelled as schizophrenic, although this has now been removed and replaced  
20 by ‘possible Asperger’s’.

21  
22 As Jack became more alienated from us, things got worse. He was picked up  
23 by the police, once while wandering along the underground railway line and  
24 once while climbing on a statue in a park. He broke things in the house, and  
25 although he never attacked me or stole money I was often frightened as he  
26 crashed about upstairs.

27  
28 Things came to a head when he was sectioned for the second time and spent  
29 10 weeks in a locked ward. Although dope was still available there his  
30 medication was changed and he gradually improved. We were lucky to have  
31 an excellent and understanding social worker and for the first time I felt  
32 supported to some degree by the system.

33  
34 The next stroke of luck was that Jack was offered a place on a rehabilitation  
35 programme so that when he came out he was monitored by a team under an  
36 exceptional psychiatrist who was the first psychiatrist who appeared to see  
37 his patients as human beings. Although very overworked, this doctor took the  
38 time to consider each patient individually and agreed to gradually reduce  
39 Jack’s medication. Jack also managed to stop using dope in order to be  
40 allowed to come home from his half-way house.

41  
42 Fast forward about 10 years and Jack has been off neuroleptic drugs but still  
43 needs antidepressants and gets very bad headaches. He is not happy – he

1 leads an isolated life and has had a couple of strange, seemingly psychotic  
2 episodes, over the last year. We need support, but the services are  
3 underfunded and understaffed; only last week Jack kept an appointment with  
4 his social worker (a different one sadly to our earlier helper) and no-one told  
5 him that they had been called out on an emergency. He felt let down and  
6 angry that he was just left to wait rather than being told. Three close friends  
7 of ours have had sons of a similar age who have committed suicide, and this  
8 never leaves my mind especially when I hear Jack feeling let down and  
9 undervalued.

10  
11 I struggle with my sadness, wondering what I could have done differently in  
12 Jack's early life. Sometimes it is unbearable. Jack's father and I separated 22  
13 years ago – how much was this a factor?

14  
15 The family and my relationship with Jack's very patient step-father is  
16 affected. The ache in my heart is always there due to living with a son who  
17 wishes that he was not alive. I suffer for him and I suffer for myself. I am  
18 lucky in many ways in that Jack has a decent small flat and is able to drive his  
19 car; he also studies a lot and practises martial arts when he has the energy.  
20 But there are days when he stays in bed all day, and he is sometimes angry  
21 and unapproachable and leaves a mess in the kitchen and fills our non-  
22 smoking household with his cigarette fumes. He has not used 'recreational'  
23 drugs for many years and hardly drinks alcohol, but he is very self-  
24 deprecating and bitter and very much into the occult as a way of escaping the  
25 reality of everyday life. This can lead to some dangerous practices.

26  
27 My experience with the mental health services has been that there is no  
28 awareness of the need for continuity – the staff in our centre seem to change  
29 almost monthly. The one psychiatrist is overworked and so only crises are  
30 dealt with promptly. Most of the social workers are very friendly and well  
31 meaning, but don't seem to have much in the way of counselling or  
32 psychological training or support for themselves.

33  
34 We have been offered a consultation for a diagnosis of Asperger's, but  
35 nothing has come of this. Basically Jack is not ill enough to get real help or  
36 well enough to lead a 'normal' life. We continue to do our best to manage in a  
37 kind of limbo, but it is not a comfortable place for Jack, or those who love  
38 him.

### 39 **4.3.3 Carer account B**

40 I am the carer of my son who is 32 years old and currently has a dual  
41 diagnosis. He has been ill for 12 years, originally with the diagnosis of

1 schizoaffective disorder, but over the past few years this has changed to dual  
2 diagnosis, though his condition and substance misuse behaviour have been  
3 much the same throughout. His main drug is cannabis (skunk), but he has  
4 used most of the other commonly available recreational drugs. Initially, and  
5 before he was ill, these were mainly ecstasy, amphetamines and alcohol. He  
6 still uses these but crack, cocaine and heroin (smoked) have become regulars.

7  
8 When my son was first ill he was 200 miles away at university. The first  
9 indication of problems was a call from a friend with whom he shared student  
10 accommodation, who expressed some concern about his behaviour. I then  
11 received a call from my son about money problems. When I suggested I visit  
12 to help sort things out, my son readily agreed. I found him pleased to see me  
13 but quite agitated, and exhibiting some paranoia, but the most disturbing  
14 issue was his 'pressure of speech'. I assumed it was problems with his studies,  
15 though he denied it. I then managed to meet with his professor who said he  
16 was coping well, the only concern being a lack of actual work being  
17 submitted. He suggested I speak to student welfare. They felt that his  
18 behaviour suggested mental health problems and suggested talking to the  
19 university GP. She referred me to a visiting psychiatric nurse at the end of the  
20 week. The intervening few days convinced me that the problems were serious  
21 as my son's paranoia and pressured speech became more apparent. I also  
22 became aware of the heavy cannabis use of my son and his fellow students,  
23 almost at the level of ordinary tobacco use - my presence in the house only  
24 inhibited them slightly. The psychiatric nurse became quite alarmed and  
25 arranged an immediate meeting with a psychiatrist, who wanted to admit  
26 him to hospital but, given the distances involved for me, agreed to my request  
27 that we returned home. A consultation with our GP at home resulted in my  
28 son being admitted to hospital under a Section 3.

29  
30 Over the next 4 years my son was in hospital several times, mainly under  
31 section. For the rest of that period he lived in the family home. He was then  
32 encouraged by the assertive outreach team to move into independent  
33 accommodation on the rather spurious grounds that a young man of 24  
34 needed his independence. While he was able to live independently with only  
35 limited support, his drug use accelerated due to his lack of ability to control  
36 his social circumstances. The flat became the hangout for both his old friends,  
37 who were still living at home and therefore had their illegal activities  
38 restricted, together with, more unfortunately, members of the drug  
39 community (fellow users and suppliers), who in effect made use of him. This  
40 situation has persisted since, being relieved slightly by a period in a council  
41 hostel and other short periods when he effectively moved back home.

42

1 Approximately 7 years ago during another Section 3 enforced period in  
2 hospital he was put on depot injections of Clopixol, which has kept his illness  
3 under control but means he is quite debilitated for a few days after the  
4 fortnightly injections and generally claims that, in part, his drug use  
5 (particularly cannabis), is necessary to relieve side effects of the medication.  
6

7 My life has been affected in several ways. There is the normal disruption  
8 suffered by all carers of somebody with a serious mental health condition  
9 such as daily visits when he was in hospital, urgent calls at any time of the  
10 day or night for support during periods of paranoia or stress, and highly  
11 charged, emotionally stressful situations dealing with illogical and delusional  
12 arguments and accusations. The drug misuse adds financial and safety  
13 concerns. Encounters with drug suppliers have not only been stressful, they  
14 were also probably dangerous. In the early days I had to settle drug debts  
15 running to several hundred pounds. Currently we have a fairly stable  
16 relationship, with small loans usually being repaid the following week from  
17 benefits, though arguments still arise when it is obvious that all of the week's  
18 benefits have been spent within a few hours and I am expected to fund the  
19 whole week; it also stressful to be called in the early hours of the morning for  
20 money. I am not sure that my financial support is in my son's best interests -  
21 while it ensures he does not go without, it does not encourage him to be  
22 independent and I suspect drug suppliers have been happy to advance credit  
23 to him because he has me to bail him out when debts get too high.  
24

25 Initially treatment for my son was only offered for his mental health  
26 problems, indeed, his first consultant said that his admitted use of cannabis  
27 was not a problem so long as it was not excessive. Times have changed.  
28 Various antipsychotic drugs were tried, including clozapine, but none was  
29 really very successful until the Clopixol depots. Very little other treatment has  
30 been offered. During the second detention in hospital an assessment was  
31 carried out by a clinical psychologist and although he felt sessions could be  
32 helpful, the consultant insisted that it was too early. I did not feel I was  
33 involved in any real sense in forming treatment plans at this time but anyway  
34 they amounted to little more than prescribing medication. Just as importantly  
35 I was not asked about my views on my son's history and therefore several  
36 things were recorded as delusions that were in fact true. Although he was  
37 definitely ill, the assumption that most of his stories were untrue still rankles  
38 with my son and means he distrusts the medical team. During the central  
39 period of his illness I had a good relationship with his key worker on the  
40 assertive outreach team and was invited to CPA reviews. My son was  
41 generally uncooperative at these due to the build up of stress at the situation  
42 causing problems, but the outcome was that little was offered apart from  
43 continuation of the medication; even variation of the dosage to reduce side

1 effects was never seriously discussed. Since that particular key worker moved  
2 on 3 years ago I have had little contact with his care team, and only when  
3 initiated by me.

4  
5 Initially his drug misuse was almost ignored. He was encouraged to go to the  
6 drug and alcohol service but having eventually got him there, they decided he  
7 was not ready for treatment as his mental state was not stabilised. The main  
8 reason for this attitude was his lack of interest in stopping his drug use (he  
9 still maintains his stance on cannabis though he does accept that other drugs ,  
10 especially crack, cause him financial problems). Following a change in the  
11 structure of the drug and alcohol service and the emergence of dual diagnosis  
12 as a label, my son did start regular meetings with a counsellor. Although  
13 these went on for several months they appeared to have little effect,  
14 floundering again on the belief of my son that cannabis use is not a real  
15 problem. At the time of writing his only treatment is medication though he  
16 has been relatively stable and open to other possibilities.

17  
18 My view is that the traditional approach to substance misuse is not really  
19 suitable for dual diagnosis sufferers since it relies heavily on the premise that  
20 there is a desire to stop using drugs that needs to be supported. My  
21 experience with my son and his peers is that they have little interest in  
22 stopping their drug use and their mental health problems mean they are not  
23 open to the normal logic. This is especially true of cannabis use where there is  
24 a strong belief in the general population that use is not a problem anymore  
25 than responsible drinking is.

26  
27 At the start of my son's illness a family counsellor came to our home. She  
28 spent most of the time talking to my wife, although she did little to reassure  
29 her and offered little in the way of advice on dealing with our son's delusions.  
30 His drug use was ignored other than suggesting that we were over  
31 controlling in trying to stop it. I do not remember much about her visits,  
32 except that I was unimpressed, especially when she criticised me for putting  
33 pressure on my son to take his medication; shortly afterwards he was re-  
34 admitted after relapsing because of non-compliance. She completely ignored  
35 my daughter, who had great difficulty coming to terms with her 'big  
36 brother's' problems. My daughter still has reservations about contact with  
37 him but these are now largely over fears for her young family and his social  
38 situation.

39  
40 In an attempt to understand more about the illness and the help available we  
41 became involved with Rethink (then National Schizophrenia Fellowship).  
42 This was helpful in a social sense but only to a limited extent since nobody  
43 else appeared to have drug misuse concerns. From this I became involved

1 with the PCT advisory group, NIMHE and the National Forum for Assertive  
2 Outreach. From these I gained more insight into services but, unfortunately,  
3 what I learnt primarily was how little there was to offer someone like my son.  
4 Most interventions I have seen relate to injectors (for example, needle  
5 exchanges, substitution programmes) and are not relevant to cannabis and  
6 crack smokers. More structured activities would help as at least part of the  
7 problem is boredom and emptiness.

8  
9 Generally people I was in contact with were sympathetic but were unable to  
10 offer much help. As a civil servant my managers were quite helpful in  
11 allowing time off for visiting, consultations and meetings. Over time most  
12 non-professional support fell away including my wife, who appeared to lose  
13 hope as time went on and things did not seem to be improving. Others, such  
14 as his neighbours, have had almost no sympathy for his situation. The council  
15 housing department were particularly lacking in understanding for his  
16 condition and how it affected his ability to obey their rules. Housing has been  
17 a particular problem and the caring team seemed unprepared to engage with  
18 the issue, despite the obvious effects it had on his illness (he reacts  
19 particularly badly to stressful situations). However, the police were generally  
20 very helpful and understanding in their contact with him, largely as a victim.

#### 21 **4.3.4 Carer account C**

22 I have been the main carer of my grandson for nearly 15 years. Jim is now 30  
23 and has a diagnosis of schizophrenia and an alcohol problem. He started  
24 living with me when he was 15 after things became increasingly difficult for  
25 him while living with his stepfather and mother, who also has mental health  
26 problems.

27  
28 When Jim started living with me he was taking drugs and drinking. At that  
29 time I had no idea about the drug use but did know that he was drinking with  
30 his friends at weekends. He was unhappy and quite isolated. He got some  
31 work with his father (my son), but his behaviour started becoming a bit  
32 strange and he would say odd things. We knew there was something wrong  
33 and his father paid for him to go to a private hospital; he did not receive a  
34 diagnosis at this time.

35  
36 Not long after that first admission he was admitted to another hospital near to  
37 where his mother lived. Around 2000 Jim became increasingly unwell and we  
38 had our first contact with our local mental health services. A consultant  
39 psychiatrist and nurse came to see him at home. They thought he might have  
40 a drug-induced psychosis. They were both good: they listened, provided  
41 advice and gave us information. Jim was started on medication for the

1 psychosis but it made little, if any, difference and he got worse. He would be  
2 agitated and suspicious and think things had special meanings for him. He  
3 was not offered any help for his drug use.

4

5 Sometimes he could be very scary and on one occasion he smashed up my  
6 house and attacked me. I had to call the police. Jim ended up being taken to  
7 hospital under a section of the Mental Health Act. As well as the police, there  
8 was an ambulance, doctor, social workers. I hadn't realised that was how it  
9 would be.

10

11 Jim has had several admissions to hospital, the longest of which was for 18  
12 months. During that admission he spent a long time on the psychiatric  
13 intensive care unit as well as time on other wards. The hospital was a terrible  
14 place. Most of the staff – doctors and nurses - were awful. They were  
15 disrespectful and not interested in the patients. I wrote a letter of complaint  
16 about one of the wards but did not get any response. The one exception was  
17 the manager of the intensive care unit. He was gentle and calm and would  
18 always explain what was going on and the reason for things. Although Jim  
19 hated it there he did not want me to complain as he was afraid it would have  
20 negative consequences for him. He used to spend most of his time in his room  
21 so that he could keep out of the way of the other patients and staff.

22

23 When he was in hospital I visited Jim every day – including Christmas day. I  
24 took him food and cigarettes. After one of his admissions Jim was placed in a  
25 hostel. It was dirty and the staff were awful. It was just dreadful. I couldn't let  
26 him stay there.

27

28 Despite being tried on lots of different medications Jim didn't really get any  
29 better. When he was on the open wards he would abscond, often to go out  
30 drinking. I used to go out looking for him, but he would often end up back at  
31 my house.

32

33 It wasn't until one of his mental health review tribunals that a doctor asked  
34 why he had not been tried on clozapine. After that he was started on it and it  
35 made a difference straight away. Since being discharged from that admission  
36 he hasn't been re-admitted to hospital – that's about 6 years now. Clozapine  
37 has been a lifesaver for him.

38

39 After his discharge Jim was put under the care of the assertive outreach team.  
40 I've got nothing but praise for them. Over the years he has had a number of  
41 care co-ordinators and two support, time and recovery (STR) workers. The  
42 consultant psychiatrist responsible for his care is the one that we met during  
43 our first contact with local services. The dual diagnosis nurse specialist has

1 also been involved over quite a few years now. Having continuity, where you  
2 can build up a strong relationship with someone, has been really helpful. All  
3 the assertive outreach staff have been very good and they're always reliable.  
4 I've been given their mobile phone numbers so I can contact them if I need to.  
5 They always take any concerns I have seriously and recognise that I know Jim  
6 really well and can spot when things aren't right at an early stage. When there  
7 have been times when Jim's mental health has deteriorated they have  
8 responded quickly and, when necessary, have visited him at home every day.  
9 The STR workers have bent over backwards to get Jim out and doing more  
10 social things. They'll phone, pick him up and do things like going to the gym,  
11 meeting up for coffee or going shopping. They've all been really flexible and  
12 helpful. I always attend the CPA meetings and these have been arranged at  
13 times that are convenient for me - I still work a few hours each week.

14  
15 Over the years I've provided Jim with a lot of practical support, like doing his  
16 washing, ironing and shopping, making sure he's managing his money and  
17 not getting behind with his bills, liaising with his bank and the utility  
18 companies, and taking him up to the mental health team to have his blood  
19 taken, or to collect his medication. Although he's lived in his own flat for a  
20 long time now, he always comes to stay with me overnight once or twice a  
21 week - and sometimes has stays for longer periods. When he does that I know  
22 he's had a decent meal. I set limits on his drinking. I won't let him drink  
23 strong lagers in my house. He knows I don't like him drinking and am  
24 worried about the effect it has on him. I'm sure he would make more progress  
25 if only he could stop. I phone him everyday to remind him to take his  
26 medication - even when I'm away on holiday.

27  
28 I have been offered a carer's assessment and been given information about  
29 carers' groups but they're not my sort of thing. I get a lot of support from my  
30 partner, who gets on well with Jim, and other family members provide  
31 support too.

32  
33 Over the years Jim has gradually made changes: he can live on his own,  
34 manage his money, take his medication (with reminders from me), do some  
35 shopping, travel on public transport on his own, and visit his brothers and  
36 Mum and stay over with them. He stopped taking drugs a long time ago and  
37 has had a few periods when he has stopped drinking but he keeps going back  
38 to it. Jim has often talked about courses or getting some voluntary or paid  
39 work but hasn't been able to follow through on his ideas yet. His assertive  
40 outreach team offered to do things with him but he always declines. Left to  
41 his own devices he will often stay in bed all morning. I think he lacks  
42 confidence. If only he had a bit more self-belief he could achieve more. I think

1 it's difficult for him because his Dad and brother have been very successful. I  
2 think his Dad is a bit embarrassed and disappointed by him and he feels that.  
3  
4 I strongly believe that whatever happens to Jim it is up to me and my family  
5 to deal with it. I'll continue to keep supporting him as long as he needs me.  
6

## 7 **4.4 REVIEW OF QUALITATIVE RESEARCH**

### 8 **4.4.1 Clinical review protocol (qualitative research)**

9 The review protocol, including the review question, information about the  
10 databases searched and the eligibility criteria used for this section of the  
11 guideline can be found in Table 7.  
12

13 A systematic search for qualitative studies, observational studies and reviews  
14 of qualitative studies of people with psychosis and coexisting substance  
15 misuse was undertaken. The aim of the review was to explore the experience  
16 of care for people with psychosis and coexisting substance misuse and their  
17 families and carers in terms of the broad topics of receiving a diagnosis,  
18 accessing services and having treatment. Reviews were sought of qualitative  
19 studies that used relevant first-hand experiences of people with psychosis and  
20 coexisting substance misuse and their families/carers.  
21  
22

**Table 7: Clinical review protocol for the review of qualitative studies**

Component	Description
Review question(s)	1.5.1 For people with psychosis and coexisting substance misuse, what are their experiences of having problems with psychosis and coexisting substance misuse, of access to services and of treatment?  1.5.2 For families and carers of people who have psychosis and coexisting substance misuse, what are their experiences of caring for people with psychosis and coexisting substance misuse, and what support is available for families and carers?
Electronic databases	CINAHL, EMBASE, MEDLINE, PSYCINFO, HMIC, PsycEXTRA, PsycBOOKS
Date searched	Database inception to 25.06.2010
Study design	Systematic reviews of qualitative studies, qualitative studies
Population	People with psychosis and coexisting substance misuse
Critical outcomes	None specified - any narrative description of service user experience of psychosis and coexisting substance misuse

#### 1 4.4.2 Studies considered

2 Based on the advice of the GDG, this review was focused on qualitative  
3 research only as it was felt it was most appropriate to answer questions about  
4 the experience of care of those with psychosis and coexisting substance  
5 misuse. As good quality qualitative research exists within the literature,  
6 quantitative and survey studies were excluded.

7  
8 The search found 21 qualitative studies which met the inclusion criteria  
9 (Alvidrez *et al.*, 2004; Bradizza & Stasiewicz, 2003; Carey *et al.*, 1999; Charles &  
10 Weaver, 2010; Costain, 2008; Dinos *et al.*, 2004; Hawkins & Abrams, 2007;  
11 Healey *et al.*, 2009; Johnson, 2000; Lobban *et al.*, 2010; Loneck & Way, 1997;  
12 Padgett *et al.*, 2008a, Padgett *et al.*, 2008b; Penn *et al.*, 2002; Pollack *et al.*, 1998;  
13 Strickler *et al.*, 2009; Todd *et al.*, 2002; Turton *et al.*, 2009; Vogel *et al.*, 1998;  
14 Wagstaff, 2007; Warfa *et al.*, 2006) and 28 were considered for the review but  
15 they did not meet the inclusion criteria. The most common reasons for  
16 exclusion were because quantitative or survey methodology had been used or  
17 because the people included in the research did not have psychosis and  
18 coexisting substance misuse. The characteristics of all the studies reviewed in  
19 this section, and references to excluded studies are summarised in Appendix  
20 13.

21  
22 Once qualitative studies were assessed for methodological quality, themes  
23 from each study were extracted and synthesized in a narrative synthesis to  
24 reflect overarching themes to capture the experience of people with psychosis  
25 and coexisting substance misuse, and their carers. The studies have been

1 categorised under seven main headings: service user experience of psychosis  
2 and coexisting substance use, access and engagement, carers' perspective,  
3 service user experience of psychosis and coexisting substance use, social  
4 networks, employment, and treatment.

#### 5 **4.4.3 Experience of psychosis and coexisting substance misuse** 6 **and effects of substance use**

7 Eight studies (Alvidrez *et al.*, 2004; Bradizza & Stasiewicz, 2003; Carey *et al.*,  
8 1999; Charles & Weaver, 2010; Costain, 2008; Healey *et al.*, 2009; Lobban *et al.*,  
9 2010; Warfa *et al.*, 2006), four of which were conducted in the UK, looked at  
10 effects of substance use in a population of participants with psychosis and  
11 coexisting substance misuse. The main topics emerging within effects of  
12 substance use were management of symptoms with substances, physical and  
13 psychosocial consequences and effects of substance use, and triggers leading  
14 to substance use.

15  
16 Carey and colleagues (1999) and Alvidrez and colleagues (2004) interviewed  
17 participants about positive and negative aspects and consequences of  
18 substance misuse and abstaining. Both studies identified interpersonal  
19 problems and alienation from social networks (especially substance using  
20 social networks) as a negative aspect of abstaining from substance use.

21 Conversely, one positive aspect of substance use mentioned was improved  
22 social skills and less social inhibition. While some participants felt that their  
23 drug use was the driving force behind the development of mental disorders  
24 (*'It activates...it triggers the mental illness'*), the majority of participants  
25 expressed that drug use has both beneficial and negative effects on their  
26 psychiatric symptoms (Alvidrez *et al.*, 2004). In a more recent study by  
27 Charles and Weaver (2010), five of 14 participants perceived their substance  
28 use to directly influence development of their mental health problems, while  
29 five others felt that substance use made their psychiatric symptoms worse.  
30 Additionally, seven people acknowledged that substance use contributed to  
31 relapse and worsened their mental health after the onset of psychosis.

32  
33 Seven studies found that substances were commonly used by people with  
34 psychosis for managing their symptoms. Charles & Weaver (2010) found that  
35 participants did not self-medicate, but did use substances to prevent the  
36 effects caused by their anti-psychotic medication (for example, drowsiness).  
37 Bradizza & Stasiewicz (2003) also found that experiencing symptoms of  
38 psychosis triggered alcohol and drug urges, as such substances helped people  
39 to cope with psychotic episodes:

40

1 *'that's why I kept using heroin. I mean, my paranoia was bad. I thought everything*  
2 *and everyone was after me'.*

3  
4 For people with schizophrenia, substance use relieved negative symptoms  
5 (for example, lack of motivation and energy) but exacerbated psychotic  
6 symptoms (for example, paranoia). Participants described the cyclical nature  
7 of their mental illness and drug misuse. Psychiatric symptoms trigger  
8 substance use, which acts as a catalyst for additional symptoms that  
9 precipitate further substance use:

10  
11 *'..The worst problem in my life right now is this vicious cycle that I've been in for the*  
12 *past seven years, which is battling substance abuse and then how the substance abuse*  
13 *impacts my depression, my self-esteem and the paranoia...'* (Alvidrez et al., 2004)

14  
15 *'It's like you know something really isn't no good for you, but at the same time, you*  
16 *want the results of an escape from reality temporarily, so you go ahead and do it'.*  
17 (Alvidrez et al., 2004)

18  
19 Positive aspects of abstaining consisted of improved living skills, better  
20 physical health, getting off the streets and away from crime, regaining trust  
21 from others and engaging in social activities. Fears and negative perceptions  
22 of abstaining from substance use included anticipating the physical effects of  
23 withdrawal, loss of relationships with substance-using friends, and the cycle  
24 of relapse.

25  
26 Despite the perceived positive aspects of substance use, participants did have  
27 insight and awareness about the dangers of using substances to alleviate  
28 symptoms:

29  
30 *'[alcohol] has a tendency to make a person think that his problem is less severe than it*  
31 *might be. It kind of clouds an image of what's really going on and will cause*  
32 *continual problems.'* (Alvidrez et al., 2004)

33  
34 Cannabis was most often mentioned for helping with delusions, controlling  
35 symptoms, and 'normalising behaviour' (Costain, 2008). Participants in  
36 Costain's (2008) study also perceived improvement in cognitive functioning  
37 from cannabis, as well as increased levels of energy and reduced  
38 psychological pain. The authors point out that this may influence adherence  
39 to treatment for service users with schizophrenia, and that clinicians must be  
40 aware of the phenomenological expressions and beliefs of service users with  
41 schizophrenia. They argue that ignoring this issue may have an impact on the  
42 development of a therapeutic relationship. Additionally, service users with  
43 bipolar disorder would often use substances because they had a desire to feel

1 normal without the sedative effects of their medication, or to attempt to  
2 recapture how they felt pre-diagnosis (Healey *et al.*, 2009). Substances used to  
3 help people relax were most often alcohol or cannabis (Wagstaff, 2007). Warfa  
4 and colleagues (2006) also found cannabis was used by participants to have a  
5 'good impact' or feeling of being strong.

6  
7 Feelings of anger and loneliness were most often expressed as emotions  
8 leading to substance use. In relation to this, other participants with bipolar  
9 disorder felt that substance use was a way to control and manage mood  
10 states, particularly mania and depression (Healey *et al.*, 2009), though many  
11 realised that this was not a reliable method of controlling mania. Anxiety,  
12 depressive symptoms and relieving pressure were also cited as reasons for  
13 substance use (Alvidrez *et al.*, 2004; Carey *et al.*, 1999; Healey *et al.*, 2009). Most  
14 participants experimented with alcohol and drugs before receiving a  
15 diagnosis of psychosis or in the early course of their illness. The substance  
16 misuse then became out of control, either because they were unaware of their  
17 mental disorder, or did not understand the effects the substances had on their  
18 mood. In this experimental phase with substances, dependency is often  
19 established.

20  
21 Additional triggers leading to substance misuse were feelings of being  
22 stressed or overwhelmed by life events. These issues could stem from poor  
23 housing, unemployment, family relationships and legal problems (Bradizza &  
24 Stasiewicz, 2003; Carey *et al.*, 1999). In some instances, previous traumatic life  
25 events served as a trigger for substance use (Charles & Weaver, 2010).

#### 26 **4.4.4 Access and engagement**

27 Having a diagnosis of psychosis and coexisting substance misuse can  
28 significantly impact on a person's ability to access and engage in services and  
29 in treatment. This can be due to a myriad of factors including stigma,  
30 ethnicity, socioeconomic status, gender, and perception of services. Several  
31 themes emerged under the broad heading of 'access and engagement' to  
32 services for those with psychosis and coexisting substance misuse, including  
33 the factors that may act as barriers to accessing treatment services, such as  
34 external and internal stigma, ethnicity and gender. This review also identified  
35 'reasons for seeking help' as a theme emerging from the included studies.  
36 There were six studies from which themes of access and engagement emerged  
37 (Dinos *et al.*, 2004; Johnson, 2000; Loneck & Way, 1997; Penn *et al.*, 2002; Todd  
38 *et al.*, 2002; Warfa *et al.*, 2006).

39  
40 Dinos and colleagues (2004) interviewed service users in community and day  
41 mental health services in London in an attempt to describe the relationship of

1 stigma to mental illness and the consequences of stigma for the individual.  
2 One significant theme that emerged for participants with a psychosis and  
3 coexisting substance misuse was anxiety surrounding managing information  
4 regarding both their illnesses, and issues of disclosure (whether to disclose to  
5 friends, family and prospective employers). Overt discrimination from others  
6 was experienced by most of the participants in this study, typically in the  
7 form of verbal or physical harassment, or through actions such as damage to  
8 property. Those with a comorbid mental illness and substance misuse  
9 reported having been verbally abused and patronised more frequently than  
10 those with other diagnoses. People with psychotic disorders experienced  
11 physical violence, as well as reduced contact with others. They also felt that  
12 they had been discriminated against in that they had not been selected by  
13 educational institutions or employers due to their diagnosis. As a result, most  
14 participants felt fearful, anxious, angry, and depressed, as well as isolated,  
15 guilty and embarrassed. These feelings resulting from stigma were a  
16 significant hindrance to recovery and a barrier to seeking help:

17  
18 *'It makes you feel bad.. it makes you feel even worse... when people don't*  
19 *trust you and think you're going to do something to someone.'*  
20

21 On the other hand, many participants reported positive aspects to having a  
22 mental illness, expressing relief that they had a proper diagnosis and  
23 appreciating their treatment:

24  
25 *'I feel that if I survive it I've been through a very privileged experience and*  
26 *that I can actually make something of it...'*  
27

28 Interestingly, no participants who were drug dependent expressed this  
29 positive view of their illness. It is evident that for this study population,  
30 stigma was a pervasive concern for the majority.

### 31 ***Black and minority ethnic groups and socioeconomic status***

32 One UK study (Warfa *et al.*, 2006) looked at drug use (specifically cannabis  
33 and khat<sup>4</sup>) in black and minority ethnic (BME) groups. Whereas East African  
34 communities showed that use of khat was linked to their culture, cannabis  
35 was seen as entangled with religious uses for black Caribbean populations.  
36 Participants in the study stated that the cultural context of their substance use  
37 was not taken into account by healthcare professionals. Some participants in  
38 the study mentioned that their clinics or clinicians exhibited cultural

---

<sup>4</sup> Khat is a plant native to East Africa and the Arabian Peninsula, and when chewed, acts as a stimulant.

1 awareness, while others felt that there needed to be increased cultural and  
2 religious sensitivity within services in the UK (Warfa *et al.*, 2006).

3  
4 Johnson (2000) interviewed families in the United States caring for a family  
5 member with psychosis and coexisting substance misuse. The marked  
6 differences in socioeconomic status and its connection with access and  
7 engagement in care emerged as significant themes. Upper-middle class  
8 European-American families felt a greater sense of individual and organised  
9 support compared with families of a lower socioeconomic status. In contrast,  
10 upper middle class families from an ethnic minority were most difficult to  
11 identify as they did not access care as frequently. They were very rarely  
12 connected with an organised support group and therefore were less visible to  
13 services compared with other socioeconomic status groups. The lower middle  
14 class families were found to have a more extensive family network although  
15 this did not seem to facilitate management of family members' illnesses.

16  
17 Families of individuals with psychosis and coexisting substance misuse from  
18 all ethnic and socioeconomic status groups felt disregarded or dismissed by  
19 mental health professionals with whom they engaged, feeling that their  
20 knowledge and opinion was rarely taken into account by mental health  
21 professionals (especially staff at crisis centres, hospitals, and psychiatrists in  
22 all settings). The experience of stigma for middle-class families differed from  
23 the lower-class families, in that those in the upper-middle class were often  
24 embarrassed that a family member was ill and therefore not functioning to  
25 their own or their social network's standards, and consequently felt distanced  
26 from other families in their network. The low and lower-middle class families  
27 felt stigmatised mostly when dealing with professional mental health and  
28 legal systems. Surprisingly, only 25% of the families interviewed had been  
29 involved in an organised support network (for example, a family group or  
30 self-help group). One suggestion the authors make is that there needs to be  
31 greater knowledge of other families struggling with an ill family member and  
32 information about community groups to go to for support.

### 33 *Gender*

34 Penn and colleagues (2002) examined treatment concerns for women with  
35 coexisting mental illness and substance misuse. The women interviewed  
36 emphasised how a person-centred approach facilitates treatment, especially  
37 when the clinician embodies traits such as empathy, honesty, and being  
38 encouraging and direct. All participants identified that negative staff attitudes  
39 or changes in the service significantly hindered their treatment progress (for  
40 example high staff turnover, lack of coordination between services, feeling  
41 judged). Childcare services were mentioned as necessary for women

1 accessing treatment, as was support that specifically accounted for women's  
2 needs.

### 3 *Reasons for seeking and accessing help*

4 Many people with psychosis and coexisting substance misuse do not come to  
5 treatment until the pattern of illness is well established (Vogel *et al.*, 1998).  
6 Similarly, Padgett and colleagues (2008b) interviewed psychiatric service  
7 users with a psychosis and coexisting substance misuse who used to be  
8 homeless and found that people typically entered treatment once symptoms  
9 of mental illness became overwhelming (for example, increased  
10 hallucinations):

11

12 *'I got to a point.. I can't take it no more. I'm going to the hospital'.*

13

14 Another key reason for reducing or stopping substance misuse was a change  
15 in personal life goals, for example an increase in the perceived value of health,  
16 income, and social relationships (Lobban *et al.*, 2010). In addition, the desire to  
17 be accepted within a certain social milieu can play a part in both initiating  
18 drug use and in terminating it. A significant event can lead to a dramatic  
19 change in behaviour and lend support to wanting to become abstinent as well  
20 (Lobban *et al.*, 2010).

### 21 **4.4.5 Importance of social networks**

22 There were eight qualitative studies addressing the effect of social networks  
23 on people with psychosis and coexisting substance misuse (Bradizza &  
24 Stasiewicz, 2003; Carey *et al.*, 1999; Charles & Weaver, 2010; Hawkins &  
25 Abrams, 2007; Lobban *et al.*, 2010; Padgett *et al.*, 2008a; Turton *et al.*, 2009;  
26 Wagstaff, 2007). All the studies highlighted that individuals often feel isolated  
27 from their social networks and do not have many people with whom to  
28 socialise. Given the pervasiveness of their illness, many found it difficult to  
29 make new friends and often relied on substance-abusing friends for support  
30 (Bradizza & Stasiewicz, 2003). Other participants highlighted the need for  
31 support and having contact with others who have experienced similar mental  
32 health and substance problems (Turton *et al.*, 2009):

33

34 *'most of the counsellors there were ex-addicts themselves and I could relate to them,*  
35 *and the things they said because they've been through it'.*

36

37 Both Hawkins & Abrams (2007) and Padgett and colleagues (2008a) examined  
38 the social networks of those with a psychosis and coexisting substance misuse  
39 who were homeless. Social networks were perceived to be smaller, primarily  
40 because many members of their social networks died prematurely (homeless

1 service users with stressful environments were at a higher risk of mortality),  
2 or service users withdrew or pushed others away. Many participants had  
3 witnessed a death of a loved one; and death appeared prominently in all of  
4 the narratives in this study. When social networks diminished, some  
5 participants reacted by attempting to rebuild their network, even if this  
6 involved negative social interactions with strong substance use triggers, while  
7 others reacted by isolating themselves further to escape social pressures.  
8 Many participants adopted 'loner talk' and wanted privacy, which arose from  
9 negative life experiences or distrust of those around them.

10  
11 Social benefits were also frequently cited as reasons for substance misuse.  
12 Lobban and colleagues (2010) differentiated between internal and external  
13 attributions for ongoing drug-taking behaviour. Participants who made  
14 internal attributions for substance use described seeking out information and  
15 weighing up advantages and disadvantages of taking drugs in order to make  
16 their decisions. This was also found in Carey and colleagues' (1999) study,  
17 where participants made a 'decisional balance' before using substances.  
18 Substance use was found to have a positive effect on interpersonal  
19 relationships in helping people 'fit in' and facilitating connections with others.  
20 Furthermore, drugs were a way to reduce social anxieties.

21  
22 Social networks were seen as a way to experiment with substances in order to  
23 gain experience, providing the person with 'social currency' which further  
24 encourages substance misuse (Charles & Weaver, 2010). A study by Vogel  
25 and colleagues (1998) and by Charles and Weaver (2010) also confirms this  
26 finding, in that participants felt that using drugs and/or alcohol elicited  
27 feelings of confidence and 'belonging', which often promoted even more  
28 substance use.

29  
30 Many participants talked about how drug use in their community was the  
31 'norm' (Lobban *et al.*, 2010). Participants who attributed their substance use to  
32 those around them found that their social networks grew around drug-using  
33 communities, and also increased their level of detachment from non-drug  
34 using networks. Socialising in drug-using communities reinforced not only  
35 shared experiences, but also facilitated drug accessibility and consumption  
36 (Charles & Weaver, 2010; Lobban *et al.*, 2010). Therefore, the social aspect of  
37 belonging and acceptance plays a part in both initiating and terminating drug  
38 use, and is fundamental in increasing motivation to use substances. When the  
39 social networks are associated with drug-using behaviour or triggers, this is a  
40 hindrance to promoting and maintaining abstinence. Young people in  
41 particular identified that their social networks were very important to them,  
42 and much of their substance use was linked to social activities. Thus, they felt

1 that they would require drastic changes to their social networks and  
2 surroundings in order to reduce their substance use.

3

4 Evidently, social inclusion is important to this population in terms of building  
5 relationships (and re-building social capital post-treatment), and influencing  
6 substance use.

7

#### 8 **4.4.6 Experience of treatment**

9 The experience of treatment for people with psychosis and coexisting  
10 substance misuse varied widely. Central themes appeared to be ambivalence  
11 towards medication, ceasing medication, the importance of self-help and  
12 mutual support groups, having a key worker, and cultural sensitivity  
13 integrated within services. Eight studies highlighted the experience of  
14 treatment for people with psychosis and coexisting substance misuse  
15 (Costain, 2008; Johnson, 2000; Loneck & Way, 1997; Pollack *et al.*, 1998; Todd  
16 *et al.*, 2002; Vogel *et al.*, 1998; Wagstaff, 2007; Warfa *et al.*, 2006).

#### 17 *Experience of assessment and referral from the staff perspective*

18 Loneck and Way (1997) and Todd and colleagues (2002) looked at how to  
19 assess service users with psychosis and coexisting substance misuse from a  
20 staff perspective, how to refer them to appropriate services, and keep them  
21 engaged in the care plan. In the study by Loneck and Way (1997), healthcare  
22 professionals working in an accident and emergency ward emphasise that for  
23 service users with schizophrenia, a more supportive approach to engagement  
24 must be employed, whereas those with substance use disorders are more  
25 receptive to a style that is more directive and if necessary, confrontational.  
26 The approach advocated by these healthcare professionals for service users  
27 with psychosis and coexisting substance misuse is a combination of  
28 supportive and directive styles, and is confrontational only when necessary.  
29 Support was characterised by listening and assessing needs, whereas a  
30 directive approach meant having a structure and steps in order to move  
31 service users into appropriate services. If service users were resistant to the  
32 supportive approach and unwilling to accept referrals, persuasion and  
33 motivational techniques could be adopted to motivate service users to accept  
34 more appropriate referrals to services. Lastly, healthcare professionals  
35 identified that the therapeutic alliance is crucial to successfully engaging with  
36 service users with psychosis and coexisting substance misuse. The most  
37 important factors to ensure a strong therapeutic alliance were: agreement  
38 about goals and tasks, and strengthening the service user-clinician bond.  
39 Todd and colleagues (2002) found that the essence of optimal care was the  
40 provision of a comprehensive assessment and a care plan that addresses both

1 urgent and non-urgent issues related to both illnesses. The care plan should  
2 be integrated across services, and make sense to the service user such that it  
3 encourages engagement and motivation to change, and is readily accessible.  
4 However, staff feared that this proposed treatment service consisting of an  
5 integrated assessment and care plan would further strain the system and  
6 increase workload.

### 7 *Experience of therapeutic relationship*

8 When participants were asked about their most positive experience of  
9 services in the UK, they highlighted having a key worker (for example, a  
10 social worker) with whom they have a good relationship, in addition to  
11 accessing local counselling services or alternative treatment options (for  
12 example, spiritual services or specific cultural support groups) (Warfa *et al.*,  
13 2006). These services and options were seen as integral to their progress in  
14 treatment.

15  
16 One limitation cited by many participants was the lack of cultural awareness  
17 and sensitivity in mental health services. They also mentioned that meetings  
18 with healthcare professionals were not long enough, and there was not  
19 enough attention being paid to social activities (Warfa *et al.*, 2006).  
20 Participants emphasised that alcohol or drug dependence made service  
21 engagement extremely difficult.

22  
23 Emotional support and time investment by service providers was important  
24 across all cultural groups with psychosis and coexisting substance misuse  
25 (Warfa *et al.*, 2006). This, therefore, highlights the importance of developing  
26 an active therapeutic relationship with a service user, fostering trust and  
27 confidence and addressing all of the person's identified needs.

### 28 *Treatment options*

29 Once service users were in treatment, many were frustrated at the lack of  
30 individual 'talk' therapy to help discuss and heal the trauma incurred from  
31 having a mental illness, having a substance problem, and living on the streets.  
32 Conversely, some participants had positive views about services, particularly  
33 the atmosphere and amenities, the sense of privacy, and staff who were warm  
34 and humane (Warfa *et al.*, 2006).

### 35 *Medication adherence and effects*

36 Service users in the study by Warfa and colleagues (2006) found that  
37 medication for their psychosis works for them and generally improved their  
38 mental health. However, antipsychotic medication typically is associated with  
39 negative perceptions and, consistent with this view, the Wagstaff (2007) study

1 found that the most common reason for participants to cease taking their  
2 psychotropic medication was that they did not perceive themselves as  
3 requiring medication in the first place. Costain (2008) found that many  
4 participants had side effects from their antipsychotic medication, and when  
5 participants also had anxiety symptoms, they stopped taking their medication  
6 and increased their cannabis use. The reasons for non-adherence to  
7 medication were varied. Many felt that adherence to medication would not  
8 enable them to have control over their symptoms (for example, delusions).  
9 Others did not perceive they had a mental illness and therefore the  
10 medications were irrelevant (Costain, 2008).

11  
12 Pollack and colleagues (1998) found that participants cited symptom  
13 improvement as the bigger driver for adhering to their medication, however  
14 the side effects and potential to be stigmatised because of the need for  
15 medication were a concern:

16  
17 *'So actually, when you say you're suffering because of your side effects, it's not only*  
18 *the physical part, but how you think you're perceived by other people'.*  
19

20 Other service users suggested that therapists should address ambivalence  
21 towards medication (Warfa *et al.*, 2006).  
22

23 Relapse was also associated with discontinuing medication treatment because  
24 of wanting to avoid the stigma of 'needing medication':  
25

26 *'I've realised the medication is doing a lot for me, but at the same time, it's*  
27 *going back and grabbing that security blanket again and that feeling, or that*  
28 *high, that desire, that craving...'* (Pollack *et al.*, 1998)  
29

30 All of these factors highlight the notion that the relationship between  
31 adherence to medication and substance use is complex. In terms of improving  
32 medication adherence or aftercare attendance, participants highlighted family  
33 influences as the most positive, especially in providing support or initiative.

#### 34 ***Self-help groups***

35 Many participants interviewed by Vogel and colleagues (1998) mentioned  
36 that a mutual support programme was extremely beneficial in enabling  
37 people with psychosis and coexisting substance misuse to share similar  
38 experiences and providing a non-judgemental atmosphere in which they  
39 could discuss problems. The support group increased participants' optimism,  
40 brought them comfort and changed their attitudes towards taking their  
41 medication (Vogel *et al.*, 1998).

1  
2 Pollack and colleagues (1998) interviewed inpatients with psychosis and  
3 coexisting substance misuse about the factors that affected their attendance in  
4 an aftercare programme. Self-help meetings (for example, Alcoholics  
5 Anonymous [AA]) were easier to attend because of the flexible timing and the  
6 fact that they facilitated social activities:

7  
8 *'Just being around the other people, you know, I've pretty much alienated*  
9 *everyone due to my drug addiction and alcohol...so it provides me the*  
10 *opportunity to...generate a new relationship'.*

11  
12 *'I found that it was a joy to go and share my daily achievements with a group*  
13 *of people that knew my condition because their own condition was so*  
14 *similar'.*

15  
16 On the other hand, attending AA meetings that were not designed for those  
17 with psychosis and coexisting substance misuse was unhelpful and perceived  
18 as contributing to relapse. As the meetings were tailored to people with  
19 alcohol and drug misuse disorders, one participant felt that they were treated  
20 differently because of their other diagnosis, leading them to seek other  
21 meetings.

### 22 *Experience of treatment from the carers' perspective*

23 One prominent theme that emerged from the interviews conducted by  
24 Johnson (2000) with carers of people with psychosis and coexisting substance  
25 misuse was the benefits and marked differences due to person taking their  
26 prescribed medication. Most families had noticed a significant improvement  
27 in functioning when their family member was on medication. However, many  
28 service users replaced their prescribed medication with street drugs, leading  
29 to deterioration in functioning and to rehospitalisation. Family members who  
30 cared for people with psychosis and coexisting substance misuse felt excluded  
31 from mental health services and considered that their efforts were largely  
32 ignored by mental health practitioners.

33  
34 It was emphasised that greater knowledge of and contact with other families  
35 struggling with the same problem would be beneficial, as would more  
36 emotional support from extended social networks. Support groups, led by  
37 professionals, that were specifically for people with psychosis and coexisting  
38 substance misuse and their families and carers were also mentioned by carers  
39 as beneficial.

#### 1 **4.4.7 Employment**

2 Strickler and colleagues (2009) interviewed people with psychosis and  
3 coexisting substance misuse about their experience with employment. Having  
4 a psychosis and coexisting substance misuse was perceived as a prominent  
5 barrier to gaining and maintaining employment; the most frequently cited  
6 barriers were the psychiatric symptoms themselves (such as manic episodes,  
7 delusions, anxiety and stress). Both Strickler and colleagues (2009) and  
8 Bradizza and Stasiewicz (2003) found that regular employment was difficult  
9 to obtain for those with psychosis and coexisting substance misuse.  
10 Furthermore, the longer the period of unemployment, the more the difficulty  
11 of finding and sustaining employment increased. As a result, there is often an  
12 extended period of unemployment with little money available to engage in  
13 additional activities. This in turn, can encourage substance use. Employment  
14 was conceptualised as a positive event which aids recovery, and adds  
15 therapeutic value to a service user's life:

16  
17 *'Work was really kind of helpful. I didn't have as many symptoms because I*  
18 *was too busy working'.*

19  
20 *'It helps my mental illness. It gives me structure'.*

21  
22 Employment helped to reduce substance use and keep participants away  
23 from drugs or alcohol. It occupied the service user and kept their daily living  
24 skills intact (for example, maintaining daily hygiene at a level suitable to  
25 attend work). The regular use or dependence on substances made consistent  
26 employment significantly more difficult.

27  
28 Employment, therefore, held a positive structural value to participants,  
29 providing them with an additional sense of belonging and contributing to  
30 society:

31  
32 *'When I am working I feel like I am contributing. I don't feel isolated.'*

#### 33 **4.4.8 Summary**

34 The evidence from the narrative synthesis of the qualitative studies provides  
35 some important insights into the experience of people with psychosis and  
36 coexisting substance misuse and their carers. Substance misuse appears to  
37 stem from a range of environmental and social factors including the  
38 management of psychiatric symptoms and/or social situations that encourage  
39 and exacerbate substance use.

40

1 Perhaps the most central theme of the reviewed literature was the importance  
2 of social networks. People with psychosis and coexisting substance misuse  
3 commonly identified interpersonal problems and alienation from social  
4 networks across all studies. This alienation and lack of a positive social  
5 support network seemed to influence their substance use, ability to seek  
6 treatment, maintain positive change, and increased vulnerability to relapse.  
7 Many negative social networks grew around drug-using communities and  
8 reinforced substance misuse.

9

10 The reasons for substance misuse were cited in nearly every qualitative study  
11 included in this review. For the most part, service users highlighted the  
12 positive and negative drawbacks to substance use and its direct effect on their  
13 psychosis.

14

15 People with psychosis and coexisting substance misuse were often  
16 stigmatised by others and faced discrimination. Many also felt internal stigma  
17 which made them hesitant to disclose their diagnosis or 'edit' it. Awareness of  
18 stigma can often be a hindrance to recovery and a barrier to seeking help in  
19 this population. People from a minority ethnic group also felt that the cultural  
20 context of their substance use was not taken into account by healthcare  
21 professionals. From the carers' perspective, families from ethnic groups and  
22 groups of lower socioeconomic status felt disregarded by mental health  
23 professionals. As a group, women felt that they faced additional barriers to  
24 treatment in the form of more social stigma, and the need for childcare while  
25 seeking and undergoing treatment. In addition, women felt that they received  
26 less support from treatment providers, and would benefit from a more  
27 empathetic and therapeutic approach. The studies focusing on women  
28 emphasise that a person-centred and non-judgemental atmosphere is  
29 necessary in order to foster openness and willingness to change. All  
30 participants highlighted that negative staff attitudes hindered their treatment  
31 progress.

32

33 An inability to access services easily, combined with negative interactions  
34 with healthcare professionals, highlights the importance of an appropriate  
35 assessment and referral process, which takes into account both the psychosis  
36 and the substance misuse. The literature indicated that a good assessment,  
37 which is direct in nature, should be employed for the substance use problem,  
38 whereas a non-judgemental, empathetic approach is preferred for assessment  
39 of psychosis. Staff however, found this comprehensive assessment  
40 problematic due to the increase in resource use and strain on time for  
41 healthcare professionals.

42

1 Regarding treatment, most participants found medication to be beneficial, but  
2 ambivalence about it was common often due to the regimen and side effects.  
3 Participants also spoke positively about having a good relationship with a key  
4 worker or participating in a self-help group. Employment was seen as  
5 providing positive structural value and a sense of belonging.

6  
7 Family and friends can have an important role to play in supporting a person  
8 with psychosis and coexisting substance misuse. They can promote and  
9 maintain change, but in order to do this they require information and support  
10 from healthcare professionals. The strain on carers, however, can be  
11 challenging and they may require a carer's assessment.

12  
13 From a staff perspective, the qualitative studies suggest that an improvement  
14 in staff training is required to facilitate access and engagement in treatment  
15 for people with psychosis and coexisting substance misuse. When  
16 interventions were successfully delivered, a thorough assessment, as well as  
17 coordination between mental health services and substance misuse services,  
18 were two components of care perceived as crucial.

19  
20 One interesting result emerging from all the studies was the realisation that it  
21 is possible to conduct qualitative research with this specific population and  
22 engage them in focus groups and interviews. This finding can hopefully  
23 facilitate further research in the future for people with psychosis and  
24 coexisting substance misuse.

25  
26 While these qualitative studies provide insight about the experience of care  
27 for service users with psychosis and coexisting substance misuse, the overall  
28 quality of the evidence was moderate. All studies were assessed for  
29 methodological quality according to a qualitative study checklist (NICE,  
30 2009), however several of the included studies could have been improved by  
31 describing methodology and data analysis further. In addition, the theoretical  
32 frameworks and approaches were variable across studies, as were the  
33 populations they focused on.

## 35 **4.5 QUALITATIVE ANALYSIS**

### 36 **4.5.1 Introduction**

37 The following section includes a qualitative analysis of transcripts available  
38 on the internet from people with psychosis and coexisting substance misuse.  
39 These were accessed from the following websites: Healthtalkonline  
40 (<http://www.healthtalkonline.org/>), Dual Recovery Anonymous

1 (<http://draonline.org/>), Meriden Family Programme  
2 (<http://www.meridenfamilyprogramme.com/>),  
3 Talktofrank(<http://www.healthtalkonline.org/>), Foundations Associates  
4 (<http://dualdiagnosis.org/>), Bipolarworld(<http://www.bipolarworld.net/>),  
5 and Rethink (<http://www.rethink.org/>). The websites all provided  
6 information and support to people with psychosis and coexisting substance  
7 misuse and included personal narratives from people with these conditions  
8 and their carers. The review team undertook their own thematic analysis of  
9 the narrative accounts to explore emergent themes that could be used to  
10 inform recommendations for the provision of care for people with psychosis  
11 and coexisting substance misuse. It should be noted that service users with  
12 diagnoses of bipolar disorder, schizophrenia, schizoaffective disorder, and  
13 psychotic disorder were all included in these transcripts, in addition to having  
14 problematic or dependent substance use.

#### 15 **4.5.2 Methods**

16 Using all the personal experiences available from seven websites, the review  
17 team analysed the accounts of 48 service users. All accounts were published  
18 on the website in their original form. The majority are written by people from  
19 the UK but there are also some from the US. Poems and letters were excluded  
20 from the analysis. Each transcript was read and re-read and sections of the  
21 text were collected under different headings using a qualitative software  
22 programme (NVivo). Initially the text from the transcripts was divided into  
23 six broad headings emerging from the data: impact and experience of  
24 psychosis and coexisting substance misuse; access and engagement;  
25 experience of treatment; carers' perspectives; and support and services. Under  
26 these broad headings, specific emergent themes that were identified  
27 separately and coded by two researchers. Three GDG members also  
28 individually coded the testimonies into emergent themes. Overlapping  
29 themes and themes with the highest frequency count across all testimonies  
30 were extracted and regrouped under the subsections below.

31  
32 There are some limitations to the qualitative analysis for this guideline. Some  
33 of the accounts are written in retrospect, whereas others are written more  
34 recently, or in the present. This may have had an impact on the way in which  
35 the experiences were recalled; moreover, the accounts cover different time  
36 periods which may affect factors such as attitudes, and information and  
37 services available.

1 **4.5.3 Impact and experience of psychosis and coexisting**  
2 **substance misuse**

3 Given the debilitating impact of having a diagnosis of psychosis or a  
4 psychotic-related disorder with coexisting substance misuse, the main themes  
5 emerging from the online accounts regarding experience of illness described  
6 the symptomatology of their disorder(s), the emotions they felt in receiving an  
7 accurate diagnosis, the use of self-medication to control psychiatric  
8 symptoms, and, lastly, gaining insight into their mental illnesses.

9 *Symptoms of psychosis and coexisting substance misuse*

10 Many service users alluded to the cyclical nature of their mental health  
11 problems (especially those with bipolar disorder), and how these symptoms  
12 were or were not affected by their substance use:

13  
14 *'When I first got sober, the manic-depressive disorder appeared even more*  
15 *pronounced than it had before. It was no longer hidden by alcohol and drugs.*  
16 *The stress of withdrawal in my early recovery triggered wild mood swings*  
17 *for me.'*

18  
19 *'At times my moods were changing from depression to manic even without*  
20 *booze or drugs. Sometimes I got so depressed I would seclude myself for*  
21 *weeks at a time with out paying attention to whether I bathed or ate.'*

22  
23 Participants also described how they would hide their symptoms from others:

24  
25 *'You can't lump everybody in together, you know, to say oh this is, these*  
26 *people are manic depressives, so their behaviour would be blah, blah, blah.*  
27 *Everybody is different...I might act different to the next manic depressive or*  
28 *whatever and, you know, perhaps I might not show my symptoms because*  
29 *there's one thing about manic depression, depressives you really are clever at*  
30 *hiding your symptoms and very good at manipulating people.'*

31  
32 *Self-medication as a reason to misuse substances*

33 Self-medicating with drugs or alcohol as a way to manage symptoms  
34 emerged as a prominent theme in the online accounts. The most common  
35 reasons for self-medication were to manage manic or depressive symptoms:

36  
37 *'The Army caught on to my problem, and tried to treat me with Lithium and*  
38 *Prozac. This helped for a little while, but I also started drinking. Eventually,*  
39 *I went off the meds and started self-medication with the alcohol.'*

40

1           *'I began to self medicate myself. Smoking weed drinking alcohol these help*  
2           *me come down from my intense moods '*

3  
4           *'I started to self-medicate. Alcohol and speed were my crutches. If I felt*  
5           *myself getting too high I would drink, if I felt I was getting too low then I*  
6           *would take a few grams of speed.'*

### 7    ***Gaining understanding***

8    Gaining an understanding of mental illness is an important step towards both  
9    engaging in treatment and promoting the recovery process. The themes that  
10   emerged centred on accepting both diagnoses of a psychotic and substance  
11   misuse disorder, and understanding how both illnesses could be treated and  
12   how their substance misuse had had an impact on their psychiatric  
13   symptoms. Understanding their conditions frequently led to positive  
14   thoughts about their illnesses and the future:

15  
16           *'Recovery from chemical dependency requires that I accept my addiction and*  
17           *abstain from mood-altering chemicals. It involves attending 12-Step*  
18           *meetings, working with my sponsor, working the 12-Steps and improving*  
19           *my physical health. Recovery from bipolar disorder..requires that I accept the*  
20           *disease. Attend dual disorder meetings; increase my activity when I'm*  
21           *depressed and decrease my activity when I'm manic, or slow down and think*  
22           *constructively.'*

23  
24           *'Believing that my mind would return to rational thinking once time healed*  
25           *it from the years of drug abuse. The entire time ignorant of [bipolar*  
26           *disorder]. As if my mind completely blocked out those years of hospitals and*  
27           *knowledge. I'm beginning to believe it was shame, fear of stigma. But still,*  
28           *why I sabotage myself is a mystery, and I still have to fight it!'*

29  
30           *' ... drugs might not be responsible for all mental illness but where, where*  
31           *people with mental illness take drugs they greatly compound the problem*  
32           *and prevent recovery. And I think that other things being equal, people do*  
33           *recover more or less but the drugs stop them recovering.'*

### 34    **4.5.4 Access and engagement**

35    Due to the additional burden of having both psychosis and a substance  
36    misuse problem, there are many barriers to accessing and/or engaging in  
37    treatment. This can stem from experience of stigma, cultural or ethnic factors,  
38    lack of coordination between services, and assessing and engaging the service  
39    user.

### 40    ***Stigma***

1 There is a significant amount of stigma attached to a severe mental illness like  
2 psychosis, and coupled with a substance misuse problem there is additional  
3 risk of stigma. Many online accounts, from both service users and carers,  
4 highlighted the experience of interacting with others in the community and  
5 the stigma that their dual diagnoses carried. The experience of stigma often  
6 elicited feelings of shame, embarrassment, and frustration:  
7  
8

9 *'When we go out there in the community people might know you have got a*  
10 *mental health problem, you might not look different to the, but they know*  
11 *you have got that. There is a stigma against it and a discrimination*  
12 *taboo..because of the label, and because of what it stands for. Which is people*  
13 *don't understand.'*  
14

15 *'I found that a lot of people disbelieve me when I say I've had schizophrenia,*  
16 *... They don't believe it because my behaviour doesn't match their stereotype*  
17 *and if there's one thing that makes me upset more than anything else is.'*  
18

19 *'So if we can get actually people on board to recognise that not all ...*  
20 *mentally ill people are violent, psychopathic or whatever that which actually*  
21 *we're just normal people trying to live our lives every day with the added*  
22 *burden of having a mental health issue then perhaps... people would get on a*  
23 *lot better.'*  
24

25 *'If anybody heard that you have a sick son, they don't want to know you.*  
26 *That's the worst part...I still hear people saying to me, "...he has two sons,*  
27 *they are sick". And when people hear that, they don't want their children to*  
28 *even come any nearer. Because they are afraid... that your son might do*  
29 *something... because they do not have enough knowledge that not all sick*  
30 *people are violent'.*  
31

32 *'When he was sectioned, we told them he had been spiked, probably with*  
33 *LSD. Bizarrely that explanation is more socially acceptable than telling*  
34 *people your son has a mental health problem. That's how far this society is*  
35 *entrenched in stigma and prejudice about mental health, but tolerates drugs*  
36 *as part of the social structure.'*  
37

### 38 ***Access for BME groups and cultural factors***

39 One theme that emerged in several testimonies was that access to care was  
40 more difficult for those coming from a BME group or a different cultural  
41 background. Factors that affected access to care for BME groups were a fear of  
42 accessing treatment due to the conceptualisation of mental illness in their  
43 home country or native culture, or fear of stigma:

1  
2  
3  
4  
5  
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7  
8  
9  
10  
11  
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23  
24  
25  
26

*'Well people look at you differently if you say you've got a mental health problem back home. They don't treat you the same. I think now it's changed but that, when I was there it was different...'*

Many felt that they were or would be treated differently by mental health professionals as a result of their ethnicity or cultural background:

*'...it wasn't so much racist it was more institutionalised racist. It's embedded within the system.'*

*'...within the mental health system it's their foreign-ness which is emphasised because it is their foreign-ness which is considered to, to shape their, their diagnosis'.*

*'...it's very hard for minority to express their views, because any time a minority express their views... "if you don't like it, what are you doing here?"'*

*'But they don't know where to go to no one. They don't go to a doctor or no GP. They want to deal with it themselves.'*

*'You know, some Black folk they don't want to go to the GP, they don't want to go, then them's not treated, because the stories they hear about the system, so we've got to find a way to make it more attractive to help them to go and get treatment before it gets worse.'*

### **Access to services**

A significant number of factors affected accessing services, including fear of contacting a healthcare professional about substance misuse, and uncertainty about how to begin accessing treatment or who to contact:

*'And I did ask somebody from my mental health team if it was possible to have like a social worker and she said no, she didn't know how I would access that. I asked my doctor the same thing she didn't know how I would access anything like that so it just leaves you vulnerable.'*

### **Coordination between services**

Another theme which emerged from the online accounts was the link between mental health services and the criminal justice system and the police. Several accounts compared how, in the UK, there needs to be more coordination between the police and mental health services in order to make the most effective referrals for people with psychosis and coexisting substance misuse.

1 In addition, information regarding mental illness was mentioned as necessary  
2 to circulate to the police:

3  
4 *'...if you're struggling with a substance misuse problem you'd be better off*  
5 *in, in the criminal justice system. People say that their lives have been saved*  
6 *by being put in the criminal justice system being forced to come off the drugs*  
7 *and then given help to stay off. And I have to tell you that at the moment*  
8 *there's no, no plan to, to give that kind of care to, to people in my trust*  
9 *[NHS].'*

10  
11 *'...if they realise that somebody is, you know, is not particularly a drunk,*  
12 *that there's something underlying with that person as well, mental health*  
13 *issues I think a mental health team should be available, a crisis team of some*  
14 *sort should be available to help that person while they're at in police custody,*  
15 *yeah. I never had any of that and so you can't, you haven't got access to your*  
16 *medication, you're off your medication, that's only going to make you worse.*  
17 *'*

18  
19 *'Like my son, the policeman came, he was so rough on him, you know*  
20 *although he has mental problem. The police are not trained. The police don't*  
21 *know what is mental health...if every community would work with the law*  
22 *enforcement, hand in hand, things might get better...'*

#### 23 **4.5.5 Support and services for people with psychosis and** 24 **coexisting substance misuse**

25 In the online accounts, people with psychosis and coexisting substance misuse  
26 frequently highlighted the positive and negative aspects of their support  
27 networks, be it personal social networks, peers accessed through mutual  
28 support groups, or mental health services. Many participants described how  
29 their social networks facilitated or impinged on accessing care or treatment.

#### 30 *Positive and negative social support networks*

31 One theme that emerged was how a lack of social support, or a social network  
32 that was based around substance misuse, hindered recovery:

33  
34 *'I had nobody there to help me with this '.*

35  
36 *'I also remember having friends who really weren't my friends if I had booze*  
37 *or drugs they were always there, if I had nothing or tried to quit they were*  
38 *always gone. It really hurt to find out who were your real friends'.*  
39

40 However, having positive social support networks actively encouraged  
41 recovery:

1  
2 *'I have the encouragement and support of my wife even though we are*  
3 *planning to separate in the near future...I also have a very close...friend, and*  
4 *although he doesn't understand bi-polar disorder, he has been very*  
5 *supportive. He makes sure that I get out of the house at least three times a*  
6 *week.'*

7  
8 *'The care and loving doesn't come from professionals. They haven't got time*  
9 *to hug me and kiss me and tell me how much they love me, and give me*  
10 *sweet things, chocolate to eat. That comes from a different source that comes*  
11 *from your friends, it comes from your family, it comes from the community.*  
12 *It comes from your spouse, your husband, your boyfriend and that happens*  
13 *after you've finished the day time treatment. So I think that is what the other*  
14 *thing is. The care and loving that we need.'*

### 15 ***The impact of key workers***

16 Another theme that emerged from the online accounts was the helpfulness of  
17 particular key workers in addressing both the psychosis and the substance  
18 misuse, acting as a positive role model and supporter, helping to encourage  
19 recovery, and referring the service user to useful community services. A key  
20 worker typically made the service user feel cared for and increased their  
21 motivation to get involved in social activities. Key workers were people to  
22 whom service users could go for help, who were separate from their personal  
23 support network and their clinicians:

24  
25 *'I have great help from [my key worker] who I see once a week and I know*  
26 *that if I have a problem I can just pick up the phone and, you know, as long*  
27 *as it's within working hours he's here.'*

28  
29 *'Because he did say to me, "The first time I met you..you were seriously ill..*  
30 *mentally," and he said, "The, the improvement over time has been great."*  
31 *And I said.. "[money adviser] that is partly because .. you've took a lot of my*  
32 *burden..and let me concentrate on getting better in myself.. putting apart*  
33 *that, the worry of all of that.'*

34  
35 *'But just that small group it makes you feel like you're being cared about and*  
36 *cared for and [my key worker] does a great job with that I think.. He can be a*  
37 *pest at times making sure that you, I've got to go out with him, "Come on*  
38 *you're coming for a cup of coffee," that's only to get, make sure that I'm*  
39 *getting out.'*

## 1 **4.5.6 Experience of treatment**

2 Due to the nature of treating both psychosis and substance misuse  
3 simultaneously, treatment for the dually diagnosed is complex and often  
4 managed across multiple services. Many online accounts highlighted  
5 experience of medication, the need for specific attributes in a therapist or  
6 mental health services, and the beneficial nature of mutual support groups  
7 addressing both of their illnesses. They also expressed the opinion that  
8 services and treatment were often disconnected.

### 9 *Interactions with healthcare professionals*

10 There were many reports within the online accounts of interactions with  
11 healthcare professionals. Some service users lacked confidence and trust in  
12 their healthcare professional:

13

14 *'And the GP, oh they have no clue about mental illness. If you go to them*  
15 *about any major problem, they look into the book, any tablets they can give*  
16 *you.'*

17

18 *'I would get very frustrated with what I felt was incompetence and*  
19 *ineptitude by my doctors. I did not feel that they were listening to me nor*  
20 *were they willing to make medication changes when my current mix of*  
21 *medications did not seem to be stopping my cycling. I had three doctors*  
22 *within that year, until I found my current doctor, who I am finally*  
23 *comfortable with.'*

24

25 *'I've seen different psychiatrists but to me they always feel, they, it's always*  
26 *felt like they're sitting on a pedestal... and I'm just there as part of their job*  
27 *really'.*

28

29 *'So the important thing is they listen to what people are saying, especially the*  
30 *people who have the illness...But they don't listen to them. They just make*  
31 *presumptions. Because of the label of they have been given. They look at a label.*  
32 *"He's paranoid schizophrenic. So we put him in that category, he must be*  
33 *saying this." Not necessarily. Things can change. Actually listen to what he's*  
34 *saying. Look at what he does. Look at his care plan. And listen ... And now*  
35 *people are beginning to listen to me and that is what makes me feel good.'*

36

37 There was a feeling among service users of having to conceal certain issues or  
38 disclose specific aspects of their illness in order to comply with their  
39 healthcare professional:

40

41 *'...make it clear that you believe what they say, very clearly that you believe*  
42 *what they say because if you show or hint that you don't believe what they*

1           *say then that's, then you've undermined your own authority in their eyes*  
2           *and therefore that makes the repair process a lot, a lot more difficult and a lot*  
3           *more long term.'*

4  
5 However some service users understood the pressures facing healthcare  
6 professionals:

7  
8           *'They've got loads to cope with. It's not their fault. Most of these things,*  
9           *people have a go about their consultant and the doctor. It's not their fault*  
10          *why these things are happening. It's the way the system is.'*

11  
12 Others highlighted the positive aspects of their healthcare professionals, such  
13 as how their doctor helped them achieve insight into their illnesses:

14  
15           *'I began to work with a new doctor, and when I told him about my continued*  
16           *marijuana smoking, he stated simply, "Do you know marijuana is bad for your*  
17           *mental health?" It was a non-judgmental statement. But, somehow it*  
18           *reverberated in me. I do not believe he judged me as good or bad for the choices I*  
19           *was making, but he just wanted to empower me by allowing me insight into*  
20           *what I was doing to myself.'*

## 21 ***Self-help***

22 Self-help groups, particularly in the online accounts from the US, emerged as  
23 a beneficial treatment option where people could openly discuss both their  
24 psychosis and substance misuse. Mutual support enabled service users to  
25 relate to someone with similar diagnoses and experiences, as well as to  
26 develop a positive social network outside of the formal group sessions. It was  
27 strongly emphasised that the support group should be focused on both  
28 illnesses, as one targeting only the substance misuse led to frustration for  
29 those who wished for their mental illness to be simultaneously addressed:

30  
31           *'I lost the zeal for AA several years ago because they didn't understand my*  
32           *bipolar condition. They felt meetings, a sponsor, and the big book along with*  
33           *a spiritual program were all you needed to obtain good sobriety.'*

34  
35           *'Dual Recovery Anonymous helps keep my whole self together so I have a*  
36           *chance to hope, cope and heal from the impact a dual disorder has had on my*  
37           *life'.*

38  
39           *'The people at the meeting really made an impression on me. I could tell they*  
40           *were sincere and serious about what they were doing, and they said they used*  
41           *to be like me until they started working this honest program. They were*  
42           *practical and realistic, yet had uncommon sense, They were humble and*

1            *unselfish, and I wanted to be as much like them as possible. I wanted what*  
2            *they had.'*

3  
4            *'I was not compliant with good mental health practices...I refused psychiatric*  
5            *medication, assuring myself that increased effort to work the 12 Steps would*  
6            *restore me to sanity... Later I would learn that my sobriety program would*  
7            *restore me to sanity from addiction and not my total mental health, but it*  
8            *went a long way in improving my quality of life.'*

9  
10           *'I met my third husband at my sponsor's house. He is also bipolar, and*  
11           *because we have worked through stabilizing his medication, then mine... we*  
12           *have learned why people in dual recovery need each other... '*

13  
14           *'I think joining a group is a big help. You'll find that you make friends, you*  
15           *make the odd friend here and there and it's up to you if you want to continue*  
16           *the friendship outside which we have done with our, when we had our black*  
17           *and ethnic group going here we all made friends and we all had each other's*  
18           *telephone numbers and we'd go out independently as well.'*

19  
20           *'My group has been a godsend... I get so much from my brothers and sisters*  
21           *in DRA [Dual Recovery Anonymous]... love, support, encouragement and*  
22           *finally, a sense of belonging. . . . I have DRA to treat my dual illnesses as a*  
23           *whole, rather than a part here, and a part there.*

24  
25           *'People show up at our meeting that I have never seen at the social club*  
26           *where it's held. They say how happy they are that they have somewhere to go,*  
27           *and they share their experience, strength and hope without reserve. They ask*  
28           *questions, and they hang around for awhile to yak and drink coffee. And we*  
29           *don't feel alone anymore. They come back the next week.'*

30  
31           *'So when you do start recognising your symptoms hopefully there will be*  
32           *somebody there, on the other end of a phone or perhaps a group you can go,*  
33           *even if it's just another mental health, mentally challenged person like*  
34           *yourself and sometimes they're better than the professionals I'm telling you,*  
35           *and give you better advice...'*

36

### 37 ***Resistance or ambivalence towards medication***

38 One of the most prominent themes that emerged from all the online accounts  
39 was a strong opinion about medication regimes for psychosis. Feelings  
40 towards medication were typically ambivalent, and side effects often  
41 outweighed the positive aspects of medication in managing symptoms. In  
42 some cases, medication had a debilitating effect and was not allowing the

1 service user to engage in other activities in their daily life (for example,  
2 holding down a job, staying awake).

3

4 Some online accounts highlighted the problematic nature of increasing and  
5 changing doses, and how this resulted in them stopping their medication  
6 altogether, or relapsing:

7

8 *'I was seeing a psychiatrist once a week and slowly I felt like my life was*  
9 *getting better. However the medication did not continue to work. So my*  
10 *doctors just put the dose up each time they saw me. I was incredibly*  
11 *frustrated with this and decided that I would take myself off all the*  
12 *medication and do it my own way.'*

13

14 *'Medications would only work for short periods of time, then we would have*  
15 *to increase dosages until we reached maximums, then we would have to*  
16 *search for something new. It was so frustrating for me, and I would often*  
17 *lose hope of ever feeling better.'*

18

19 *'However, my dosage kept increasing...even at such a high dosage, the*  
20 *medication was not showing up in my system so the doctors dropped me off*  
21 *the medication out of concern. Again, I started drinking.'*

22

23 Others were concerned about the side effects of their medication:

24

25 *'Well, lithium turned me into an emotionless zombie. I think they just had*  
26 *me on too high of a dose, but I wasn't about to live my life that way, so I*  
27 *stopped taking it. Of course, I went back on a manic high right away.'*

28

29 *'I went back to the doctors and they started me on new meds. I was exhausted*  
30 *by fatigue as a side effect of meds. I couldn't hold a job.'*

31

32 *'... most of the time you just try and dodge your medication anyway,*  
33 *everybody did it if they could.'*

34

35 *'I was in a bit of a fog with all this sedating medications so I started reducing*  
36 *it with out telling the doctors.'*

37

38 *'I soon stopped taking my prescribed medication preferring to self-medicate*  
39 *with substances that had euphoric side effects instead of the lethargy, dry*  
40 *mouth, impotence, and muscle spasms of the legitimate drugs.'*

41

42 However several online accounts expressed more positive views towards  
43 medication:

44

1           *'Coming off my meds the second i felt better..then crashing...back on my*  
2           *meds again..then crashing lower..it was a vicious cycle. I met my disability*  
3           *counselor and she explained to me everytime I came off my meds and I*  
4           *dropped to a new low it was that much harder for the medication to bring me*  
5           *back to the original me...that scared me I didn't want to lose me forever..so I*  
6           *have been faithfully taking my meds for over a year!'*

7  
8           *'Once I started taking medication for my bipolar disease, I became balanced;*  
9           *my mood swings were less severe. Medication management is critical for me,*  
10          *because any fluctuation of time or dosage can affect the purpose of the*  
11          *medication.'*

12  
13          Some service users, who were initially compliant with their medication  
14          regime, gradually stopped taking their medication without consulting anyone  
15          once they felt better, which led to relapse:

16  
17          *'For over a year I was taking my medication faithfully and feeling balanced*  
18          *and 'normal'. As with substance abuse, 'stinking thinking' started to set in,*  
19          *for my mental illness. I believed that I was 'well', so I slowly stopped taking*  
20          *my meds.'*

21  
22          *'... however I started to believe that I did not need to continue taking my*  
23          *medication because I was feeling so much better. So I stopped it all together.*  
24          *Life retuned to the rollercoaster.'*

#### 25          **4.5.7 Experience of recovery**

26          Many online accounts were positive about the future in terms of recovery and  
27          learning how to cope with their mental illness as well as maintaining  
28          abstinence from substances. The majority of the accounts expressing feelings  
29          about their recovery mentioned the tumultuous journey and the need to  
30          recognise recovery as a constant yet manageable and rewarding struggle:

31  
32          *'Life does get better and it is an enabling disability...a sort of a perceptual*  
33          *thing that never leaves you. But it is actually a gift if you can learn about it*  
34          *and manage it and get the best out of yourself. I mean it's no different from*  
35          *what anybody else is trying to do is get the best out of ourselves aren't we so,*  
36          *you know, it's pretty good.'*

37  
38          *'I still take each day as it comes. I'm always prepared for a relapse; even*  
39          *though I have five years 'under my belt' of being relatively 'episode free,' I'm*  
40          *always on alert.'*

41  
42          *'I still experience peaks and valleys, but now the cycles aren't so great or*  
43          *frequent, and they are more manageable. I know that experience teaches*

1 *expertise, help and hope replace helplessness and hopelessness, and*  
2 *weaknesses turn around to become strengths.'*

3  
4 *'Now, after a few years.....some med changes and a lot of work. I AM getting*  
5 *better! I can see the light at the end of the tunnel! I know that I have to work*  
6 *everyday to deal with my illness and I will always have to be diligent with*  
7 *my meds. But, I also know that I can feel better...'*

8  
9 *'With thanks to the Doctor's I have seen since, my condition, though present,*  
10 *is understandable now. I have greatly controlled the symptoms I have*  
11 *experienced. Gone are the days of binge drinking and marital infidelity. I*  
12 *have settled into the life of being a simple person, who get's great pleasure*  
13 *out of all the little things in life, while coping with my disability at the same*  
14 *time.'*

#### 15 **4.5.8 Carers' perspective of services**

16 Many carers held strong views on the efficacy of mental health services for  
17 people with psychosis and coexisting substance misuse. There were obvious  
18 differences between engagement in services in the US versus the UK. Carers  
19 perceived that US services outside of mental health care (for example, the  
20 police), had a better understanding of mental health care than in the UK.  
21 Others drew on the lack of communication between services in the UK. Carers  
22 perceived mental health professionals as most effective when they spent a  
23 significant amount of time with not only the service user, but the carer as  
24 well, allowing for questions to be asked about treatment and medication  
25 regimes:

26  
27 *'I can go in there and the patient and the parent, and there will be a head*  
28 *nurse or a psychiatrist or somebody there to organise the meeting. And my*  
29 *son can say anything to me and I can give a good, -and I can answer him*  
30 *back. Then a psychiatrist will say, -will tell my son he is wrong or I am*  
31 *wrong or something like that, you know. A friendly, -this thing. And to me,*  
32 *that is very, very helpful, because sometimes -you don't say things in anger,*  
33 *things go better. My son has his view, I have my view, or my son wants*  
34 *something, I will say, "I will try my best to do it". And that is very helpful.'*  
35

36 Others expressed concern about the discontinuity of care, for example in the  
37 transition to adult services:

38  
39 *'...he was eighteen...and CAMHS needed to get rid of him, but he wasn't*  
40 *having any of it. We had no idea that such a schism existed within the*  
41 *services and had assumed there would be a thread of continuity...his*

1 CAMHS doctor is a saint. But he is an overworked and under-resourced  
2 saint and he hung on to him as long as he could.'

3  
4 'The day after their eighteenth birthday they are adults and you are expected  
5 to be carers. But carers whose motives are suddenly viewed with suspicion.  
6 Carers whose agenda it is automatically opposed to theirs. You are part of the  
7 problem. You have to play by confidentiality rules and observe their  
8 conventions of procedure.'

9  
10 Some carers felt neglected by services, feeling that they received inadequate  
11 information about their family member's illness:

12  
13 'No-one told us what to expect or how to deal with anything...on a day-to-  
14 day basis; the services; medication; relapses; claiming our rightful benefits;  
15 Nothing!'

16  
17 Carers emphasised the impact of coping with their family member's illness  
18 and substance use problems on their own. Many carers provided insight into  
19 experiences and offered advice on coping and caring for someone with both  
20 illnesses:

21  
22 'Mental health needs to be handled with care and support. You have to put  
23 yourself into that person's shoes- if you are this person how would your  
24 family feel...'

25  
26 'Learning all you can is a vital part. His mood swings have many times made  
27 me want to say I give up...this isn't worth it. After I learned, and still  
28 learning each day, all that I can about bipolar disorder I now know and have  
29 some idea of what I should expect and how to handle those things.'

30  
31 Several online accounts highlighted the importance of having the right  
32 accommodation for people with psychosis and coexisting substance misuse:

33  
34 'Along with non-compliance with medication regimes and continued  
35 substance abuse, inappropriate accommodation would seem to be one of the  
36 most common causes of relapse, including remaining too long with  
37 parent/carers.'

38  
39 'Whilst there are some excellent models of supported accommodation, a huge  
40 percentage of options offer very little or no proper support, most especially if  
41 there are no family carers in the background. Service users are left vulnerable  
42 to a financially motivated system, overseen by under-resourced, underfunded  
43 and under-informed social workers, trained to feed them into what has  
44 become a multi-billion pound industry, regardless of consequences.'

1 **4.5.9 Summary of the qualitative analysis of the online accounts**

2 The online accounts highlighted the effect of substance use on psychiatric  
3 symptoms, and how many people hide their symptoms from others around  
4 them. Self-medication was frequently cited as a reason to use substances, as a  
5 way to manage or normalise psychiatric symptoms. The accounts illustrated  
6 the cycle of increased symptomatology and escalating substance use.

7  
8 The theme of social networks also ran through all of the online accounts,  
9 especially in highlighting how influential positive support can be in  
10 promoting change and optimism in the life of someone who has psychosis  
11 and coexisting substance misuse. This social support could come in the form  
12 of a carer, a key worker or advocate, or formal support through a self-help  
13 group. A number of people commented that the relationship between service  
14 user and therapist is of prime importance.

15  
16 Discontinuity of care and lack of coordination between services was also a  
17 prominent theme emerging from the accounts. A few highlighted how police  
18 and criminal justice systems could increase awareness about mental health,  
19 and promote more coordination and integration between services.

20  
21 Having a psychiatric diagnosis was often viewed as stigmatising and resulted  
22 in the service user concealing problems and symptoms from others. Many  
23 people expressed that they felt discriminated against because of their  
24 diagnosis.

25  
26 When accessing services, those from BME groups emphasised that it was  
27 difficult for minorities to express their views, and many were reluctant to  
28 approach their GP for help. Lack of information from healthcare professionals  
29 is a barrier to coming to a full understanding of psychosis and its interaction  
30 with substance misuse, the range of treatments available and the role of  
31 services.

32  
33 There were varied views about healthcare professionals emerging from the  
34 online accounts, and the main area of criticism concerned contact with the GP  
35 and maintaining a therapeutic relationship with a healthcare professional. A  
36 number expressed negative views, such as the healthcare professional being  
37 too brief and uninterested in the service user. Others felt that they had to  
38 conceal information from staff, and generally expressed a lack of confidence  
39 and trust in their healthcare practitioners. Conversely, positive interactions  
40 with healthcare professionals led to greater insight and facilitated readiness to  
41 change.

42

1 Another overarching theme emerging from the online accounts was a strong  
2 opinion about medication for psychiatric illness. There were mixed reports  
3 regarding medication; ambivalence and resistance towards medication were  
4 frequently cited due to side effects and other factors, and some people  
5 abruptly discontinued their medication once they felt better. Self-help groups  
6 (such as Dual Recovery Anonymous) were cited as beneficial in promoting  
7 change and ongoing support.

8  
9 The impact of psychosis and coexisting substance misuse on carers was a  
10 prolific theme. Some people remarked on the change of roles that occurred as  
11 a result of one person having a diagnosis of psychosis and coexisting  
12 substance misuse. Many people also commented on the supportive nature of  
13 family members and carers.

14  
15 Lastly, several online accounts explained the process of recovery, and  
16 expressed optimism and hope for the future, stemming from ongoing support  
17 from their social networks, medication and treatment, and readiness to  
18 change.

## 20 **4.6 OVERALL SUMMARY**

21 Twenty-one studies were reviewed in the narrative synthesis of the  
22 qualitative literature and 48 testimonies from seven websites were analysed in  
23 the qualitative analysis (of the websites four were UK-based and three were  
24 US-based). Many of the same themes merged from both the qualitative  
25 literature and the online accounts. Table 8 provides a list of the themes  
26 emerging from both sources of evidence.

27

**Table 8. List of themes emerging from the qualitative analysis and the narrative synthesis of the qualitative literature**

	Qualitative (thematic) analysis of online accounts	Narrative synthesis of the qualitative literature
Reasons for substance use	✓	✓
Feelings of stigma	✓	✓
Socioeconomic status as a barrier to accessing treatment	x	✓
Culture or ethnicity as a barrier to accessing treatment	✓	✓
Gender-specific barriers to care	x	✓
The importance of a comprehensive assessment and referral	x	✓
Importance of social networks	✓	✓
Positive aspects of employment		✓
Difficulty accessing and engaging in services	✓	✓
Ambivalence towards medication	✓	✓
Medication compliance and effects	✓	✓
Utility of mutual help and self-help groups	✓	✓

1

2 The literature review of qualitative studies and the qualitative analysis of  
3 online accounts revealed that many people used substances (the most  
4 common of which were alcohol, cannabis and cocaine) in an effort to control  
5 their psychiatric symptoms, such as mania or depression, although substance  
6 use was often reported as exacerbating psychotic episodes. Additional  
7 reasons for substance use with coexisting psychosis included the social  
8 benefits. Being aware of the reasons for substance misuse is important in  
9 contributing to an understanding of the relationship between psychosis and  
10 substance misuse, and how staff can better identify and help maintain  
11 positive change.

12

13 Stigma was discussed in the qualitative analysis as well as in the literature  
14 review. Those with psychosis and coexisting substance misuse concealed their  
15 feelings and thoughts, which was a barrier to getting help or support. The  
16 literature showed that few people with psychosis and coexisting substance  
17 misuse seek help until they have had a serious psychotic episode or have hit  
18 'rock bottom'. When people do present to services, typically one of their  
19 coexisting illnesses is treated while the other problem is left untreated.  
20 Furthermore, carers from BME groups of all socioeconomic statuses were  
21 difficult to engage in services. The primary study authors felt that more  
22 attention should be given to engaging this carer group and population in  
23 treatment (for example, through the provision of culturally-specific  
24 community groups). Families with a higher socioeconomic status had

1 adequate support networks and did engage more frequently in treatment. The  
2 online testimonies highlighted that an increase of support groups with a focus  
3 on recovery for both psychosis and substance misuse could be beneficial.

4  
5 Moreover, the GDG discussed that healthcare professionals in both mental  
6 health and substance misuse services could have benefitted from having more  
7 cultural sensitivity and awareness towards the linkages between culture and  
8 substance use, and provide culturally-specific services for BME groups  
9 presenting with psychosis and coexisting substance misuse. Evidence from  
10 the Warfa *et al.* (2006) study showed that BME groups were heavily accessing  
11 culturally-tailored programmes in the UK.

12  
13 Women felt additional internal stigma due to alcohol misuse being perceived  
14 largely as a male problem. They reacted positively to healthcare professionals  
15 who employed an empathic, non-judgemental approach, but were critical of a  
16 lack of childcare opportunities and rigid treatment programmes that did not  
17 allow for flexible timing to enable women to enter treatment and care for their  
18 family. Treatment could potentially be adjusted or more flexible treatment  
19 times could be provided in order to account for this.

20  
21 Both the literature and the online accounts highlighted the perceived lack of  
22 coordination and communication between services (mental health and  
23 substance use). It is important to take these findings into account and ensure a  
24 better continuity of care. Having a key worker was frequently cited in both  
25 the literature and the online accounts as providing objective support to the  
26 service user and being beneficial for facilitating recovery and referring the  
27 person on to appropriate services.

28  
29 One study highlighted the need for a comprehensive assessment to properly  
30 diagnose both the psychosis and coexisting substance misuse so that the  
31 person could be referred to appropriate services, and the need to provide a  
32 more integrated treatment where the coexisting disorders can be treated  
33 concurrently. A comprehensive assessment improves professionals'  
34 understanding of the role of substance misuse in a service user's life and  
35 provides insight into their lifestyle and social circumstances. This increases  
36 the possibility of providing effective, tailored treatment and support suited to  
37 the service user. Healthcare professionals should work collaboratively with  
38 people to agree a structured support plan and encourage and motivate service  
39 users with psychosis and coexisting substance misuse to engage in treatment.  
40 A non-judgmental attitude that will engender trust in their service users is  
41 crucial. Integrating treatment and referrals are important in establishing a  
42 therapeutic relationship with the service user, together with continuity of

1 care. The positive aspects and benefits of a therapeutic relationship both in a  
2 treatment setting and in assessment procedures were cited frequently.

3  
4 The need for more information about psychosis and substance misuse (as well  
5 as the relationship between the two) with regards to treatment modalities and  
6 options, and medication regimes were mentioned consistently in the literature  
7 and the online accounts. Lack of accessible information may be a particular  
8 issue for people from BME groups, as well as for carers.

9  
10 Social networks emerged as a prominent theme in both the literature and the  
11 the online accounts. Positive social networks were seen as helping to promote  
12 long-term recovery and maintaining positive change, whereas negative social  
13 networks pressured people to use substances, exacerbated mental illness and  
14 encouraged relapse.

15  
16 Employment and positive social activities in addition to standard treatment  
17 can help prevent relapse from substance use disorders occurring from  
18 boredom or re-engagement with substance using social networks.

19 Employment promotes empowerment in this population, as do social  
20 activities that promote autonomy and independence.

21  
22 Both reviews highlight the importance of mutual support and self-help  
23 groups so that people with psychosis and coexisting substance misuse can  
24 connect, communicate, and interact with those with similar complex needs  
25 and experiences. The literature and online accounts had a prominent theme of  
26 ambivalence and resistance towards medication regimens, due to side effects  
27 or the perceived irrelevance of drug treatment. Many cease taking their  
28 medication, leading them to relapse and causing their psychiatric symptoms  
29 to return. In order to control the onset of psychiatric symptoms, people self-  
30 medicate with more substance use, perpetuating the cycle. This results in  
31 more hospitalisations and treatment, therefore an effort should be made to  
32 promote adherence to medication, including providing as much information  
33 as possible about medication regimes to individuals and carers, and to ensure  
34 medication monitoring and follow-up.

35  
36 In the literature as well as in the online accounts, one prominent issue which  
37 emerged for carers of people with psychosis and coexisting substance misuse  
38 was a feeling of being neglected by mental health services. The GDG  
39 discussed that more attention should be paid to carers in the care plan. There  
40 should be opportunities for carers to ask questions and information about  
41 medication and treatment should be provided. Where possible carers should  
42 be encouraged to participate in family support groups so that they can share  
43 their experiences.

1  
2 Lastly, the qualitative analysis and qualitative review of the literature  
3 reflected patient/carer views on what type of treatment is considered more  
4 important.  
5 Second, treatments other than medication (for example, certain psychological  
6 interventions, alternative treatments) did not emerge as themes as expected.  
7

### 8 *Limitations*

9 There are some limitations to the qualitative analysis and qualitative review  
10 of people's experience of psychosis and coexisting substance misuse in this  
11 guideline. First, the illustrative and retrospective nature of the online accounts  
12 must be taken into account. Furthermore a large proportion of these accounts  
13 were from the United States and treatment modalities or processes may differ  
14 or not be accessible in the UK. Secondly, only certain substances were  
15 mentioned as substances of misuse in the literature and the online accounts  
16 (for example, cannabis and alcohol), whereas other substances were not  
17 mentioned frequently, or at all (for example, hallucinogens or heroin). Despite  
18 these limitations, a number of themes were identified and ran through both  
19 sources of evidence.  
20

21 Overall, the validity of the qualitative evidence needs to be mentioned,  
22 particularly regarding the triangulation of findings from different qualitative  
23 methods and its potential limitations. It may be that it is inappropriate to use  
24 data gathered from various methods and contexts to inform the experience of  
25 care of people with psychosis and coexisting substance misuse. Furthermore,  
26 the qualitative testimonies were informative and analysed in a systematic,  
27 consensus based way, however the motivation between writing these  
28 testimonies is unknown and there could be a bias in the information these  
29 testimonies provide. This needs to therefore be taken into consideration when  
30 considering the validity of the analysis.

## 31 **4.7 FROM EVIDENCE TO RECOMMENDATIONS**

32 Both the narrative synthesis of the qualitative literature and the qualitative  
33 analysis of the online accounts revealed overlapping and similar themes,  
34 which were discussed by the GDG. Both forms of evidence highlight the  
35 value of gathering information about service user experience of psychosis and  
36 coexisting substance misuse. The qualitative evidence can therefore further  
37 inform the quantitative evidence in making better informed recommendations  
38 for improving the experience of service users and their carers. Though  
39 qualitative research is largely subjective due to its narrative nature and was  
40 aimed at a specific population that may not generalise widely to the UK

1 population, a number of themes were identified that ran through both sources  
2 of evidence.

3  
4

5 The GDG thought that the evidence from both the narrative synthesis of the  
6 qualitative literature and the qualitative analysis of the online accounts  
7 suggests that those with psychosis and coexisting substance misuse should be  
8 provided information regarding comprehensive assessment, treatment  
9 decisions and options, and aftercare. This issue is important for carers as well,  
10 as many felt neglected by services and could benefit from more inclusion in  
11 the treatment progress and be provided with more information, if the service  
12 user agrees. The GDG identified that when families, carers or chosen  
13 supporters are involved in supporting the person with psychosis and  
14 coexisting substance misuse, a carer's assessment of their caring, physical,  
15 social, and mental health needs will be important. The GDG also agreed that  
16 family intervention, as recommended in the NICE schizophrenia guideline  
17 (NCCMH, 2010), was appropriate. The GDG felt that healthcare professionals  
18 could also provide information about carer support groups and voluntary  
19 organisations, including those for psychosis and substance misuse, and help  
20 families or carers to access these, as many carers felt that they would have  
21 benefited from support from other carers with similar circumstances. The  
22 GDG also discussed issues of consent, capacity and advance decisions,  
23 agreeing that advice was needed about these issues and the legal  
24 requirements under the Mental Capacity Act and Mental Health Act.

25

26 Furthermore, the GDG thought that the literature and the online accounts  
27 highlighted that healthcare professionals should be culturally competent and  
28 able to take account of the service user's cultural or ethnic background when  
29 providing information and treatment. Information about voluntary  
30 organisations and support groups in the community which may be culturally  
31 specific could benefit both service users and carers and facilitate treatment  
32 access and engagement. No evidence was found in the economic literature of  
33 the burden on carers of service users, both in terms of financial cost and  
34 quality of life. Further research would be required to provide an empirical  
35 estimate of this burden, although such costs would be considered outside of  
36 the current NICE reference case (NICE, 2008).

37

38 Although highlighted in the website testimonies and the narrative synthesis  
39 of the qualitative studies, the GDG additionally discussed the importance of  
40 having an advocate or key worker to provide ongoing support and ensure  
41 coordination between services. It was also established within the group by  
42 consensus, that a positive therapeutic relationship between the healthcare  
43 practitioner and the service user is important in facilitating service user

- 1 engagement in services and treatment and promoting change. The evidence
- 2 reviewed here supports these discussions.
- 3

1 **4.8 CLINICAL PRACTICE RECOMMENDATIONS**

2 **4.8.1 Recommendations**

3 *Working with adults and young people with psychosis and*  
4 *coexisting substance misuse*

5 **4.8.1.1** When working with adults and young people with known or  
6 suspected psychosis and coexisting substance misuse, take time to  
7 engage the person from the start, and build a respectful, trusting,  
8 non-judgmental relationship in an atmosphere of hope and optimism.  
9 Be direct in your communications, use a flexible and motivational  
10 approach, and take into account that:

- 11 • stigma and discrimination are associated with both  
12 psychosis and substance misuse
- 13 • some people will try to conceal either one or both of their  
14 conditions
- 15 • many people with psychosis and coexisting substance  
16 misuse fear being detained or imprisoned, being given  
17 psychiatric medication forcibly or having their children  
18 taken into care, and some fear that they may be 'mad'.

19 **4.8.1.2** When working with adults and young people with known or  
20 suspected psychosis and coexisting substance misuse:

- 21 • ensure that discussions take place in settings in which  
22 confidentiality, privacy and dignity can be maintained
- 23 • avoid clinical language without adequate explanation
- 24 • provide independent interpreters (who are not related to the  
25 person) if needed
- 26 • aim to preserve continuity of care and minimise changes of  
27 key workers in order to foster a therapeutic relationship.

28 **Race and culture**

29 **4.8.1.3** Healthcare professionals working with adults and young people with  
30 psychosis and coexisting substance misuse should ensure that they  
31 are competent to engage, assess, and negotiate with service users  
32 from diverse cultural and ethnic backgrounds and their families,  
33 carers or chosen supporters.

1 **4.8.1.4** Work with local black and minority ethnic organisations and groups  
2 to help support and engage adults and young people with psychosis  
3 and coexisting substance misuse. Offer organisations and groups  
4 information and training about how to recognise psychosis with  
5 coexisting substance misuse and access treatment and care locally.

6 **Providing information**

7 **4.8.1.5** Offer written and verbal information to adults and young people  
8 appropriate to their level of understanding about the nature and  
9 treatment of both their psychosis and substance misuse. Written  
10 information should:

- 11 • include the ‘Understanding NICE guidance’ booklet<sup>5</sup>, which  
12 includes a list of organisations that can provide more  
13 information
- 14 • be available in the appropriate language or, for those who  
15 cannot use written text, in an alternative format (audio or  
16 video).

17

18 **4.8.1.6** All healthcare professionals in primary, secondary or specialist  
19 substance misuse services working with adults and young people  
20 with psychosis should offer information and advice about the risks  
21 associated with substance misuse and the negative impact that it can  
22 have on the experience and management of psychosis.

23 **Working with and supporting families, carers and chosen supporters**

24 **4.8.1.7** Encourage families, carers or chosen supporters to be involved in the  
25 treatment of adults and young people with psychosis and coexisting  
26 substance misuse to help support treatment and care and promote  
27 recovery.

28 **4.8.1.8** When families, carers or chosen supporters live or are in close contact  
29 with the person with psychosis and coexisting substance misuse, offer  
30 family intervention as recommended in ‘Schizophrenia: core  
31 interventions in the treatment and management of schizophrenia in  
32 adults in primary and secondary care’ (NICE clinical guideline 82).

33 **4.8.1.9** When families, carers or chosen supporters are involved in supporting  
34 the person with psychosis and coexisting substance misuse, discuss  
35 with them any concerns about the impact of these conditions on them  
36 and other family members.

---

<sup>5</sup> Available in English and Welsh from [www.nice.org.uk/guidance/CGxx](http://www.nice.org.uk/guidance/CGxx)

- 1 **4.8.1.10** Offer families, carers or chosen supporters a carer's assessment of  
2 their caring, physical, social, and mental health needs. Where needs  
3 are identified, develop a care plan for the family member or carer.
- 4 **4.8.1.11** Offer written and verbal information to families, carers or chosen  
5 supporters appropriate to their level of understanding about the  
6 nature and treatment of psychosis and substance misuse, including  
7 how they can help to support the person. Written information should  
8 be available in the appropriate language or, for those who cannot use  
9 written text, in an accessible format (audio or video).
- 10 **4.8.1.12** Offer information to families, carers or chosen supporters about local  
11 family or carer support groups and voluntary organisations,  
12 including those for psychosis and for substance misuse, and help  
13 families, carers or chosen supporters to access these.
- 14 **4.8.1.13** Negotiate confidentiality and sharing of information between the  
15 person with psychosis and coexisting substance misuse and their  
16 family, carer or chosen supporter.
- 17 **4.8.1.14** Ensure the needs of young carers or dependent adults of the person  
18 with psychosis and coexisting substance misuse are assessed. Initiate  
19 safeguarding procedures where appropriate (see recommendations  
20 5.8.1.23–5.8.1.27).

### 21 **Support for healthcare professionals**

- 22 **4.8.1.15** Working with people with psychosis and coexisting substance misuse  
23 can be challenging and healthcare professionals should seek effective  
24 support – for example, through professional supervision or staff  
25 support groups.

### 26 **Consent, capacity and treatment decisions**

- 27 **4.8.1.16** Before undertaking any investigations for substance misuse, and  
28 before each treatment decision is taken:
- 29 • provide service users with full information appropriate to  
30 their needs about psychosis and substance misuse and the  
31 management of both conditions, to ensure informed consent
  - 32 • understand and apply the principles underpinning the  
33 Mental Capacity Act (2005), and be aware that mental  
34 capacity is decision-specific (that is, if there is doubt about  
35 mental capacity, assessment of mental capacity should be  
36 made in relation to each decision)
  - 37 • be able to assess mental capacity using the test set out in the  
38 Mental Capacity Act (2005).

1           These principles should apply whether or not people are being  
2           detained or treated under the Mental Health Act (1983; amended 1995  
3           and 2007).

4   **Advance decisions and statements**

5   **4.8.1.17** Develop advance decisions and advance statements in collaboration  
6           with adults with psychosis and coexisting substance misuse,  
7           especially if their condition is severe and they have been treated  
8           under the Mental Health Act (1983; amended 1995 and 2007). Record  
9           the decisions and statements and include copies in the care plan in  
10          primary and secondary care. Give copies to the person, their care  
11          coordinator, and their family, carer or chosen supporter if the person  
12          agrees.

13   **4.8.1.18** Take advance decisions and advance statements into account in  
14          accordance with the Mental Capacity Act (2005). Although advance  
15          decisions and advance statements can be overridden using the Mental  
16          Health Act (1983; amended 1995 and 2007), try to honour them  
17          wherever possible.

18

19

# 1 5 ASSESSMENT AND CARE 2 PATHWAYS

## 3 5.1 INTRODUCTION

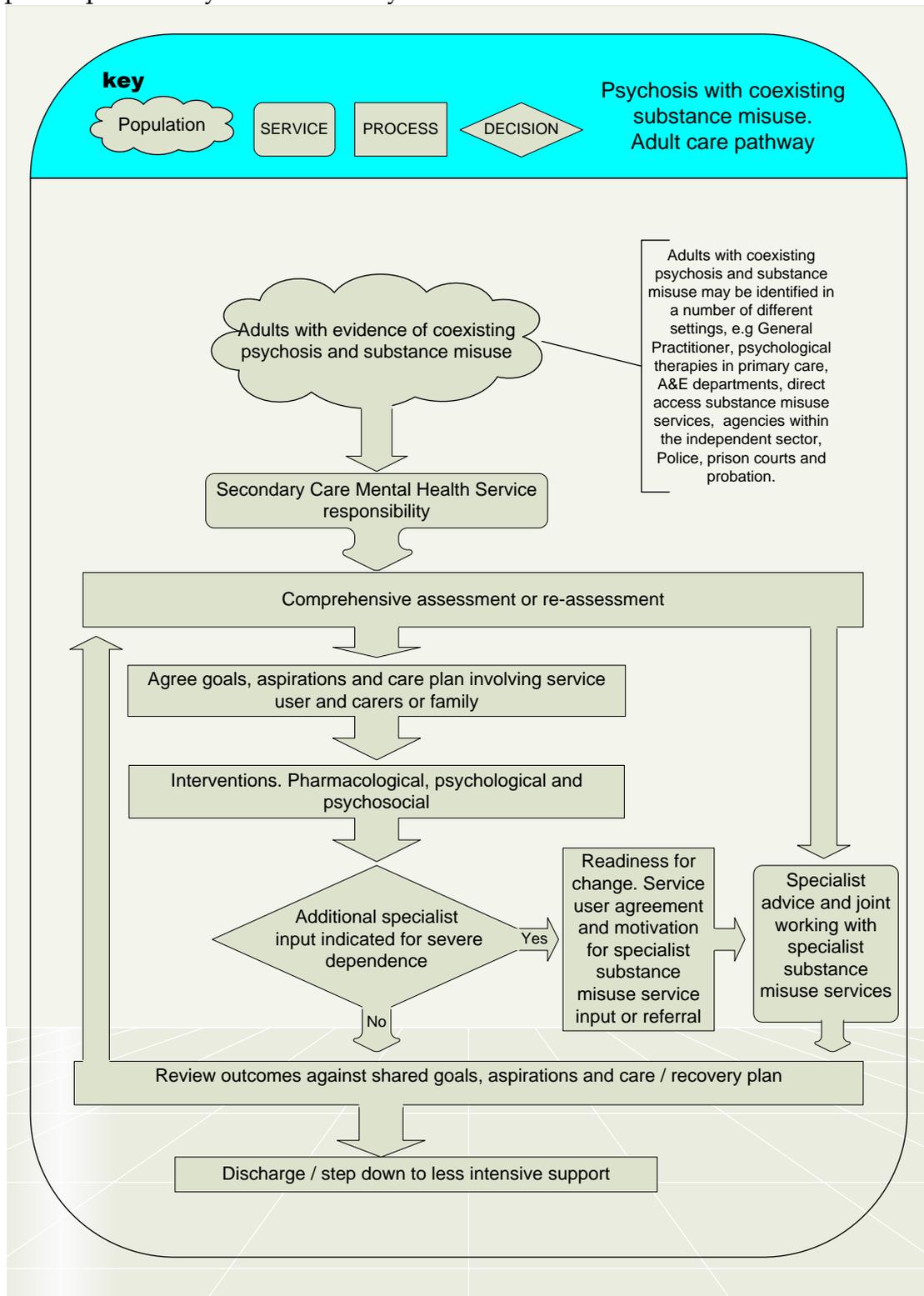
4 Because of a paucity of evidence, the GDG addressed, through expert  
5 consensus, the review questions concerning assessment (review question  
6 1.1.1) and care pathways and referral guidance (review question 1.4.1) (for  
7 further information about the methods used in this chapter, please see  
8 Chapter 3, section 3.5.6; for a list of all review questions see Appendix 6).

9  
10 The traditional problem in dealing with this group of people has been the  
11 disparity between clinical models used in different parts of the care system,  
12 particularly between addiction/substance misuse specialities and the  
13 mainstream mental health services. This has been compounded by the two  
14 services being funded and commissioned separately, and variation and  
15 confusion over which service holds clinical responsibility for people with  
16 differing relative severities of each single condition. This has, at worst, led to  
17 the exclusion of individuals with a coexisting disorder from both treatment  
18 systems, or more often, led to variable access and then attempts at parallel or  
19 sequential treatment which may become disjointed and where accountability  
20 and governance is dispersed.

21  
22 *Models of care for treatment of adult drug misusers: update 2006* (National  
23 Treatment Agency for Substance Misuse, 2006), gives a workable definition of  
24 a care pathway and the required components to be articulated: 'An integrated  
25 care pathway (ICP) describes the nature and anticipated course of treatment  
26 for a particular service user and a predetermined plan of treatment. A system  
27 of care should be dynamic and able to respond to changing individual needs  
28 over time. It should also be able to provide access to a range of services and  
29 interventions that meet an individual's needs in a comprehensive way.' The  
30 pathway therefore seeks to standardise the steps taken through access,  
31 assessment, treatment and discharge as well as provide guidance points for  
32 the thresholds and relationships between different treatment teams and  
33 services. Care pathways have been developed for drug misuse and for  
34 schizophrenia and bipolar disorder within the respective NICE guidelines  
35 (NCCMH, 2006, 2008a, 2008b, 2010).

36  
37 The care pathway is summarised in Figure 3 (Chapter 9 includes a companion  
38 care pathway for young people). Both the text and Figure 3 are designed to be

- 1 illustrative and offer some broad principles and direction, rather than to be
- 2 prescriptive. They are sufficiently broad to take into account



- 3
- 4

1 **Figure 3: Care pathway for people with psychosis and coexisting substance**  
2 **misuse.**

3 local context on the availability of services, individual need, and clinical  
4 discretion whilst providing a framework based on expert consensus.  
5

6 **5.2 PRINCIPLES UNDERPINNING CARE PATHWAYS**

7 **5.2.1 Access to mainstream services**

8 People with a psychosis and coexisting substance misuse deserve access to  
9 good quality, person-focused, and coordinated care (Department of Health,  
10 2002). The key message in the Department of Health guidance is that  
11 mainstream mental health services take responsibility for addressing the  
12 needs of people with a psychosis and substance misuse, drawing on support  
13 from substance misuse services. The rationale for this is that “substance  
14 misuse is usual rather than exceptional among people with severe mental  
15 health problems”.

16  
17 Locally agreed care pathways need to be explicit so that responsibilities are  
18 clear, and services for people with psychosis and coexisting substance misuse  
19 are delivered within mainstream mental health services with specialist  
20 support. In addition, mechanisms for resolving disagreements about team  
21 responsibility and specialist input for individuals need to be in place, such as  
22 regular care pathway meetings with executive powers.

23 **5.2.2 Right care at the right intensity**

24 Effective team working draws upon specialist skills and knowledge from  
25 within the team complemented by care pathways allowing access to further  
26 step up or step down resources ensuring that complexity is managed at the  
27 right intensity of care and that support for staff is maintained. The quadrant  
28 model (Department of Health, 2002) offers a tool for titrating the likely  
29 intensity of care and service involvement required based on the assessed  
30 relative severity of mental illness and substance misuse. Individuals who  
31 score high on both counts of need (for example, unstable schizophrenia with  
32 substance dependency) would therefore be candidates for coordinated  
33 specialist care for people with psychosis and coexisting substance misuse  
34 where available, or care from the mental health team with input from  
35 substance misuse services where required. Similarly a dependent drinker  
36 with moderate depressive symptoms would more likely be managed by  
37 substance misuse services and primary care services. The GDG decided  
38 however that we could not simply plot the service provision against the need

1 identified by each quadrant as the provision of services varies by locality and  
2 the evidence for integrated services compared to standard care is not robust  
3 (see Chapter 6).

### 4 **5.2.3 Skills and competencies**

5 Skills and competencies for working with people with psychosis and  
6 coexisting substance misuse need to be developed through training and  
7 supervision to match demand. Suitable frameworks exist for developing skills  
8 at core, generalist and specialist levels depending on the type of staff and  
9 exposure to individuals with psychosis and coexisting substance misuse  
10 (Hughes, 2006). For example, staff working in psychiatric inpatient settings,  
11 early intervention for psychosis teams and assertive outreach teams are likely  
12 to have high exposure. The competencies encompass values and attitudes,  
13 knowledge and skills, and practice development. During the review of service  
14 models reported in Chapter 6, one RCT was found that examined the  
15 effectiveness of staff training, and this is reviewed in more detail below.

#### 16 *Clinical evidence of substance misuse training*

17 Craig and colleagues (Craig *et al.*, 2008; Hughes *et al.*, 2008; Johnson *et al.*,  
18 2007) undertook a cluster-randomised trial involving brief (5 day) substance  
19 misuse training of care coordinators working within community mental  
20 health teams in South London (the COMO study). In addition to the training  
21 the care coordinators received supervision from the trainer during the follow-  
22 up period. Forty care coordinators received training and their service users  
23 with coexisting substance misuse and psychosis were followed up over  
24 eighteen months (127 service users). One hundred and five service users of  
25 thirty-nine care coordinators who did not receive the training were also  
26 followed up.

27  
28 There was no significance difference at follow-up between service users in  
29 terms of inpatient bed days, admissions and substance use at follow-up  
30 (Johnson *et al.*, 2007). Craig and colleagues (2008) reported that there were no  
31 significant differences in service costs but symptoms (as measured by the  
32 Brief Psychiatric Rating Scale [BPRS]) and needs for care were significantly  
33 lower at follow-up in the intervention group. Hughes and colleagues (2008)  
34 reported that the training course in psychosis and coexisting substance  
35 misuse interventions had a significant effect on secondary measures of staff  
36 knowledge and self-efficacy that was detectable at 18 months post-training.  
37 However improvements in attitudes towards working with drinkers and  
38 drug users in mental health settings failed to reach statistical significance.  
39 This study did not meet the eligibility criteria for the review of service  
40 delivery models but did provide some evidence that a training programme

1 for staff in substance misuse combined with supervision may have an impact  
2 on symptoms. The brief training course had only a modest impact on staff  
3 knowledge and skills in working with substance misusers.

#### 4 *Health economic evidence of substance misuse training*

5 The study by Craig and colleagues (2008) included an economic evaluation,  
6 comparing the costs and outcomes of a programme for case managers  
7 receiving substance misuse training with a waiting list control condition. A  
8 societal perspective was used for the cost analysis. The Client Service Receipt  
9 Inventory (CSRI) was used to collect resource use data over the 18 month  
10 follow-up period, including inpatient days, health care professional visits  
11 (Psychiatrist, Social worker, GP, Drug or Alcohol worker), medications and  
12 criminal justice (court, police, prison). An array of effectiveness measures  
13 were used in the study including psychiatric symptoms (BPRS), drug and  
14 alcohol consumption, quality of life (Manchester Short Assessment) and social  
15 functioning. Mean total 18-month costs were £18,672 in the intervention  
16 group and £17,639 in the control group, resulting in a difference of £1,033  
17 (95% CI, -£5,568 to £6,734). The authors did not attempt to synthesise  
18 incremental costs and outcomes, therefore the economic evaluation took the  
19 form of a simple cost-analysis. Although the results of the analysis are  
20 applicable to the UK context, it is difficult to interpret whether the training  
21 programme was cost-effective, given the variety of outcome measures used  
22 and the variability across the effectiveness measures of the training  
23 programme compared to the control group.

#### 24 **5.2.4 Choice**

25 While at times people may struggle to make informed choices about their care  
26 and treatment options, it is good practice to promote shared decision making  
27 using the assumption of competency unless assessed otherwise. Even where  
28 capacity may be limited, the active involvement of family and carers can  
29 reinforce messages from services about personal responsibility and  
30 consideration of the impact the individual's choices have upon themselves  
31 and others. Motivation and stage of readiness for change concerning  
32 substance misuse behaviour are key points determining routes on the care  
33 pathway. Sustained change comes about from engaging in a constructive  
34 alliance with the individual where the individual is supported in working  
35 through the stages of change without losing their sense of capability and self-  
36 direction towards shared goals.  
37

## 1 **5.3 PRIMARY CARE**

### 2 **5.3.1 Identification and assessment**

3 For this care pathway, primary care refers to general practice, accident and  
4 emergency departments and psychological therapy services in primary care.  
5 Services are generalist, office or department based, and offer limited intensity  
6 and frequency of contact. GPs are commonly the first resource that worried  
7 individuals or families will choose to consult and they often have a long-term  
8 perspective and relationship with people and families on their list. Frequent  
9 consultations with apparently minor ailments may signal underlying issues  
10 individuals are reluctant to disclose and the GPs' task is to elicit these hidden  
11 concerns. GPs and other primary care services play a key role in early  
12 identification and appropriate referral with full assessment of psychosis and  
13 harmful substance misuse taking place in secondary care mental health or  
14 addictions services.

#### 15 *Initial assessment in primary care*

16 Ziedonis and Brady (1997) suggested that primary care professionals should  
17 always maintain a high index of suspicion for either substance misuse in  
18 people with psychosis, or mental illness in people who misuse substances.  
19 These authors go on to suggest that when psychosis or substance misuse is  
20 detected, initial assessment for the other disorder should always take place  
21 and the findings included in referrals for secondary assessment. Alertness to  
22 and assessment for signs of current intoxication is particularly pertinent in  
23 presentations to accident and emergency departments.

24  
25 It is important for primary care practitioners to suspect and exclude physical  
26 causes for presenting symptoms, including acute intoxication, withdrawal,  
27 and side effects from medications.

28  
29 Primary care also plays a role in screening for physical co-morbidities which  
30 have a high rate of incidence in individuals with substance misuse and  
31 psychosis, including liver damage, blood borne viruses, cognitive changes,  
32 and nutritional deficiencies, particularly where dependent drinking and  
33 injecting drug use is suspected.

#### 34 *Further assessment in primary care*

35 Primary care practitioners may see individuals over a period of time and may  
36 hear the concerns of family and friends. They are therefore in an ideal  
37 position to detect the insidious decline in functioning which may be the  
38 premonitory signs of a psychotic illness. Substance misuse may present with  
39 very similar symptoms, and it is the GPs' task to establish the duration and

1 extent of substance misuse in relation to the onset of symptoms. For example,  
2 a service user may describe increasing consumption of alcohol to the point  
3 where it takes priority over other activities and results in a shortage of money,  
4 self-neglect and social withdrawal. This may clearly be distinguished from an  
5 individual who describes hearing voices and withdraws from social contact  
6 due to paranoid beliefs about others, but has a few drinks in order to sleep.

7  
8 It will usually be helpful to make an assessment of the individual's social  
9 support networks of family, friends, occupation and the degree to which the  
10 individual's networks are predicated around drinking or drug use activities.  
11 Carers may also need an assessment of their needs.

12  
13 Where significant substance use is detected in primary care, the practitioner  
14 will usually need to assess the extent to which this substance use is  
15 problematic to the individual and those they come into contact with,  
16 including children, and whether there is physical or psychological  
17 dependency on the substance.

### 18 **5.3.2 Management**

19 GPs or other primary care practitioners will normally refer a person with a  
20 first presentation of suspected psychosis for secondary assessment and not  
21 attempt to treat symptoms except to manage crisis situations until a  
22 secondary care appointment can be obtained.

23  
24 While individuals with a diagnosis of psychosis and substance misuse will  
25 normally be managed in secondary care, they remain service users of primary  
26 care and GPs may play a key role as a source of background information and  
27 may be the first to be aware of changes in individuals' physical and mental  
28 health as well as their social situations. Therefore, close liaison with the  
29 secondary care team will be necessary, and efforts should normally be made  
30 to include primary care practitioners in CPA reviews.

31  
32 People with psychosis are known to have poorer physical health than the  
33 average service user and thus will benefit from annual health checks,  
34 including monitoring of weight, blood pressure, cardiovascular risk (if  
35 indicated), respiratory symptoms and smoking cessation intervention.  
36 Regular blood test monitoring is indicated for some medications, such as  
37 lithium. These individuals will also need to be counselled regarding  
38 contraception and may need information on the safety of their medications in  
39 pregnancy.

1 The Department of Health in England and Wales has drawn up Primary Care  
2 Quality Outcomes Frameworks (QOF) (BMA & NHS Employers, 2009)<sup>6</sup>  
3 including for psychosis which detail minimum standards general practices  
4 should strive to achieve regarding the monitoring and care of these service  
5 users. The QOF for schizophrenia, bipolar disorder and other psychosis asks  
6 practices to keep a register of these service users and to record how many of  
7 them have had a review within the previous 15 months. This should evidence  
8 that the service user has been offered routine health promotion and  
9 prevention advice appropriate to their age, gender and health status. In  
10 addition, there are further indicators for the percentage of service users on  
11 lithium who have had their renal and thyroid function measured in the past  
12 15 months and a therapeutic lithium level recorded in the past 6 months.  
13  
14 Primary care physicians may also need to provide information and support to  
15 carers, families and friends, and in particular they play a vital role in  
16 monitoring and assessing the welfare of any children involved.

### 17 **5.3.3 Discharge back to primary care**

18 People with psychosis and coexisting substance misuse may be discharged  
19 back to primary care when their secondary care team is satisfied that their  
20 psychotic illness is stable and their substance use has stopped or is stable at a  
21 level at which it is unlikely to affect their mental health. Indicators of relapse  
22 need to be agreed prior to discharge including contingency plans in place to  
23 cope with a crisis.  
24

25 The GP may need to see these individuals at least for annual review and more  
26 often if indicated. They may need to ask questions to elicit symptoms of  
27 relapse of psychosis as well as gain an accurate picture of the type and  
28 quantity of substances the individual is using and the stability of their  
29 lifestyles. Prescribing records may give an indication of these service users'  
30 adherence with their prescribed medication, and individuals should normally  
31 be asked about their adherence with medication and any side effects or other  
32 problems they may be experiencing with medicines. Changes to medications  
33 would not normally be made within Primary Care but GPs may liaise with  
34 secondary care staff to gain advice about changes thought necessary and if  
35 indicated the service user may be seen for a secondary care review.  
36

---

<sup>6</sup> Further information about QOF: <http://www.qof.ic.nhs.uk/>

## 1 **5.4 SECONDARY CARE (GENERAL MENTAL** 2 **HEALTH SERVICES)**

### 3 **5.4.1 Assessment**

4 NICE Schizophrenia Clinical Guidance 82 (NCCMH, 2010) section 2.4, NICE  
5 Bipolar Disorder Clinical Guidance 38, section 4.4.4 (NICE, 2006) and NICE  
6 Drug Misuse Clinical Guidance 51 and 52 (NICE, 2007a, 2007b) sections 3.7  
7 and 6.2 respectively outline good practice core areas for comprehensive  
8 assessment and assessment questionnaires and tools. These tools have not  
9 been validated for this specific population with psychosis and coexisting  
10 substance misuse, but by consensus, the GDG considers them suitable.  
11 Assessment is also introduced in 2.4 of this guidance together with DSM-IV  
12 and ICD-10 criteria for substance misuse and harmful use and dependence  
13 syndrome.

14  
15 Assessment of substance use will normally be an integral component of  
16 mental health assessments. Some substances can trigger psychotic episodes  
17 (in use and/or withdrawal) and some can trigger relapse in pre-existing  
18 psychotic disorders. Evidence suggests that substance use is often  
19 inadequately assessed and therefore under-detected (Barnaby *et al.*, 2003;  
20 Noordsky *et al.*, 2003), resulting in potential misdiagnosis and inappropriate  
21 treatment (Carey & Correia, 1998). Even low levels of substance use by people  
22 with psychosis can worsen symptoms.

23  
24 Expert advice and assessment from substance misuse services will normally  
25 need to be sought where the service user is complex and high risk, for  
26 example injecting opiate use and dependency, or substances less commonly  
27 encountered in general mental health services. Referral thresholds for advice  
28 and subsequent interventions from substance misuse services are described in  
29 section 5.5.1.

### 30 **5.4.2 Engagement and sources of information**

31 Regardless of the circumstances at first presentation, engaging the person and  
32 working towards establishing a collaborative, respectful, trusting relationship  
33 is essential. This may require considerable sensitivity, flexibility and  
34 persistence on the part of the healthcare professional. The healthcare  
35 professional and service user may have differing views on the 'main  
36 problem', working with the person on what they see as the priority can  
37 provide a basis for working more collaboratively in the short term, and  
38 building on the relationship over the longer term.

39

1 Some people will have family members, partners or friends involved in their  
2 care. A similar collaborative relationship is also required with this support  
3 system. They can provide helpful information to contribute to the assessment  
4 process and may subsequently provide support with treatment.  
5

6 Given the multiple needs of people with psychosis and substance misuse  
7 problems a range of other service providers may be involved or have  
8 knowledge of the person (for example, GP, accident and emergency staff,  
9 housing providers, probation staff, drug/alcohol services). As well as  
10 contributing to assessment, maintaining constructive relationships and  
11 information sharing will be essential in developing effective coordinated  
12 plans.  
13

14 Confidentiality may be a particular concern for this population and their  
15 family or carers. For example, whether information about use of substances  
16 will negatively impact on treatment received, whether information about  
17 illegal activity will be passed on to the police, whether information about  
18 illness will be passed on to employers, or concerns about parenting abilities to  
19 Children and Families social services. Wherever possible the organisations'  
20 confidentiality policy should be explained at the outset. It is important to  
21 highlight that the agreed care plan is likely to involve working with other  
22 agencies and as such information sharing is an integral part of providing  
23 appropriate care. Consent to obtain and share such information should be  
24 sought at an early stage. Under some circumstances it will be necessary to  
25 break confidentiality and pass on information to relevant agencies (for  
26 example, where there is a risk to children, vulnerable adults, or others).  
27 Where possible, it will be necessary to make service user aware of the action  
28 being taken.  
29

30 Reliable systems and protocols for ensuring the safety of staff in both  
31 outpatient and community setting will normally include avoidance of  
32 attempting to assess or deliver interventions to people whilst they are  
33 severely intoxicated. A non-confrontational approach with the service user  
34 will need to be taken to agreeing to rearrange the assessment on a future  
35 occasion.  
36

37 Most assessment information is likely to be obtained by asking the person  
38 themselves unless they are floridly psychotic. Supporting self-report with  
39 observation is an important aspect of assessment and can be particularly so  
40 when people are reluctant to engage with services or to disclose feelings, what  
41 they are experiencing, or details of their substance use and funding  
42 behaviour.  
43

1 The GDG was concerned about the routine use of biological testing because of  
2 its potential to work against a collaborative approach. In typical healthcare  
3 settings a case by case approach set against a clearly explained rationale for  
4 care and treatment is preferred. NICE Drug Misuse Psychosocial  
5 Interventions Clinical Guidance 51 (NICE, 2007b) states that “urine testing for  
6 the absence or presence of drugs is an important part of assessment and  
7 monitoring”. The guidance notes that “routine screening for drug misuse is  
8 largely restricted in the UK to criminal justice settings, including police  
9 custody and prisons (Matrix Research and Consultancy & National  
10 Association for the Care and Rehabilitation of Offenders [NACRO], 2004); it is  
11 sparsely applied in health and social care settings.”  
12

13 The NICE Drug Misuse Psychosocial Interventions Clinical Guideline 51  
14 (NICE, 2007b), (section 6.2.1) provides a thorough review of biological testing,  
15 and drug misuse clinician rated and self-report identification questionnaires  
16 and their potential for identifying drug misuse in high risk populations for  
17 both adults and young people.

### 18 **5.4.3 Components of assessment**

19 Table 9 provides an overview of the assessment components for people  
20 suspected of experiencing psychosis and substance misuse (column 1) and  
21 key factors to consider when obtaining such information (column 2). This  
22 table is consistent with related NICE guidance detailed in 5.4.1.  
23

24 Having drawn together information from the assessment some consideration  
25 of the relationship between mental health and substance misuse will be  
26 possible. Knowing when the person last used particular substances may be  
27 important in determining whether their current presentation could be related  
28 to substance use alone, or whether it is a contributory factor to an underlying  
29 psychotic presentation. However, it can be difficult to distinguish symptoms  
30 and effects of mental illness from the effects of the misused substances.  
31

32 There has been a tendency to try to identify primary and secondary diagnosis  
33 however, even with careful history taking it can be impossible to disentangle  
34 symptoms, and it is recommended that both are considered primary and  
35 treated at the same time.  
36

37 It is important to obtain a picture of the person’s reasons for using substances  
38 and their understanding of the relationship between their substance use and  
39 mental health. For example, some individuals will believe that drinking  
40 alcohol lifts their low moods, while others will have insight into the fact that  
41 crack cocaine makes them more paranoid.

1  
2 When a diagnosis has been reached it will normally be fully explained and  
3 discussed with the person and their family or carers subject to consent.  
4 Information about substance use, medications being prescribed, the  
5 interaction between prescribed medication and illicit/non-prescribed  
6 substances should also be discussed and written information offered.  
7

**Table 9: Assessment - Components and considerations**

Assessment component	Key considerations
Current/recent substance use	<ul style="list-style-type: none"> <li>• Which substances is the person using? (polysubstance use is common)</li> <li>• How much they are using? (this may be expressed as weight or cost)</li> <li>• How often they are using?</li> <li>• Route(s) of administration (for example, oral, smoking, injecting)</li> <li>• When last used? (may help to explain current presentation)</li> <li>• How long they have been using at the current level?</li> <li>• Daily use: detail over past week</li> <li>• Patterns of use (for example, stable/chaotic, one substance to counteract effect of other, use following receipt of benefits followed by period of abstinence)</li> <li>• Evidence of physical dependence – past/recent experience of withdrawal symptoms in absence of substance use (may indicate need for pharmacological interventions (for example, for alcohol, opioids, benzodiazepines))</li> <li>• Whether meets diagnostic criteria DSM-IV/ICD 10</li> <li>• Severity of dependence (? Use severity of dependence questionnaire)</li> <li>• Service users' understanding of effects of use on physical and mental health</li> </ul>
Substance use history	<ul style="list-style-type: none"> <li>• Identify substances that have been used</li> <li>• Build chronology: age of first use - 'first tried', weekend, weekly, daily – pattern of use over time, whether dependent</li> <li>• Reasons for use</li> <li>• Impact on physical health, mental health, relationships, education/employment, involvement with criminal justice system,</li> <li>• Periods of abstinence – length, impact on mental health and other areas of life</li> <li>• Treatment episodes: dates, services interventions, what helped, triggers to relapse</li> </ul>
Risks	<ul style="list-style-type: none"> <li>• Consider risks associated with mental illness, substance use and inter-relationships between them</li> <li>• Consider risks to person themselves, family, carers, children, staff (on organisational premises and home visits) and wider community, for example, violence, self-harm, suicide, self-neglect, vulnerability to abuse and exploitation, accidental injury, withdrawal symptoms (for example, seizures, delirium tremens), injecting practices, blood borne viruses, accidental overdose, interactions between prescribed medication and illicit drugs and/or alcohol, unstable accommodation/homelessness, physical health problems, criminal activity</li> <li>• Risks to children</li> <li>• Risks to service users (are there vulnerable adult issues?)</li> </ul>
Social circumstances	<ul style="list-style-type: none"> <li>• Accommodation – situation and any identified needs</li> <li>• Family relationships – supportive or otherwise</li> <li>• Caring responsibilities: children, others – any safeguarding children or vulnerable adult issues?</li> <li>• Domestic violence</li> <li>• Friendships – supportive or otherwise (substance users?)</li> <li>• Education/employment (past and current) – vocational assessment required?</li> </ul>
Finances	<ul style="list-style-type: none"> <li>• Benefits/other income</li> <li>• Cost of current use</li> <li>• How substance use is being funded</li> </ul>

	<ul style="list-style-type: none"> <li>• Debts for example, rent arrears, utility arrears, to dealers</li> </ul>
Legal/forensic	<ul style="list-style-type: none"> <li>• Involvement in criminal activity to fund use (for example, shoplifting, burglary), as consequence of use (for example, drink/drug driving, violence)</li> <li>• Previous convictions, custodial sentences, any charges pending – were mental illness and/or substance use contributory factors?</li> </ul>
Medication	<ul style="list-style-type: none"> <li>• Current and past – for psychiatric, physical and substance use issues: prescribed, over the counter and homeopathic remedies – check whether prescribed medication is taken as indicated (consider non-adherence and/or abuse)</li> </ul>
Personal and family history	<ul style="list-style-type: none"> <li>• Family background</li> <li>• Early development – developmental milestones, schooling</li> <li>• Psychosocial history – physical or sexual abuse?</li> <li>• Family history of mental illness/psychological problems; substance misuse; physical health problems</li> </ul>
Physical health/ medical history	<ul style="list-style-type: none"> <li>• Physical illness(es) – past and current: consider those associated with mental illness and those associated with substance use for example, diabetes, cardiovascular disease, respiratory problems, blood borne viruses (hepatitis, HIV), liver disease, seizures, accidental injury, abscesses, bacterial endocarditis, DVT, tuberculosis, sexually transmitted diseases</li> <li>• If intravenous user, inspect injection sites</li> <li>• Hospital admissions, treatment and outcomes</li> </ul>
Psychiatric/ mental health history	<ul style="list-style-type: none"> <li>• Diagnoses, treatment, hospital admissions</li> <li>• Review of previous acute episodes, relapse signatures (taking account of substance use issues)</li> <li>• Symptoms – during acute episodes – between episodes</li> </ul>
Spiritual/cultural needs	<ul style="list-style-type: none"> <li>• Beliefs, practices</li> </ul>
Investigations	<ul style="list-style-type: none"> <li>• Biological: Urine or saliva testing can be helpful to corroborate self-reports</li> <li>• Haematological: full blood count, liver function test, hepatitis B, C, HIV</li> <li>• ECG – important for people prescribed methadone who are also prescribed other medication that can cause QT-elongation</li> </ul>
Reasons for and perceptions of use, motivation for change	<ul style="list-style-type: none"> <li>• What are the reasons for use? (for example, block out auditory hallucinations, alleviate boredom, conform with peers)</li> <li>• Does the person view their use as problematic?</li> <li>• Does s/he have want to make changes to current use (manner of use, stopping use)?</li> </ul>
Strengths and supports	<ul style="list-style-type: none"> <li>• What can the service user do well, what support do they have outside of statutory services?</li> </ul>
Involvement of other agencies	<ul style="list-style-type: none"> <li>• Identify all other agencies involved with the service user</li> <li>• Obtain collateral information</li> <li>• With consent of service user include them in future care/treatment planning and review</li> </ul>
Family/carer needs	<ul style="list-style-type: none"> <li>• Consider physical, mental health and social needs</li> <li>• Consider impact of mental illness/substance use on relationships, welfare of children, siblings, vulnerable adults</li> <li>• Assess knowledge/understanding regarding mental illness/substance use, inter-relationship, risks</li> </ul>

1  
2

1 **5.4.4 Care planning**

2 Care planning is normally a collaborative process involving the service user,  
3 and, where appropriate, his/her family/carers, and any other agencies.

4  
5 Although any substance use is likely to have detrimental effects on health,  
6 and professionals will usually think the person should work towards  
7 abstinence, many people will be unwilling or unable to do so.

8  
9 Understanding the person's perceptions of their use and motivation for  
10 change is essential for planning appropriate care/treatments. The  
11 transtheoretical model of change provides a helpful framework for informing  
12 decisions (Prochaska & Di Clemente, 1986; Prochaska *et al.*, 1992). It is  
13 important to note that the person's motivation to make changes may be  
14 different for different substances.

15  
16 Working collaboratively and accepting the person's relative autonomy is  
17 essential in maintaining a therapeutic relationship. Being non-judgemental,  
18 avoiding confrontation and maintaining optimism are likely to be associated  
19 with better long term outcomes (Miller & Rollnick, 2002; Raistrick *et al.*, 2006).

20 **5.4.5 Safeguarding**

21 Although it is essential to work collaboratively with people with psychosis  
22 and substance misuse, it is also important to recognise that those dependent  
23 upon them may also need help, and sometimes protection. When someone  
24 with psychosis and coexisting substance misuse looks after or has significant  
25 involvement with dependent children the needs and safeguarding of the  
26 child must be secured according to the Common Assessment Framework (see  
27 Chapter 9). The care co-ordinator or key worker may need to ensure that  
28 children's services are alerted to the need for assessment and possible help for  
29 the child. Similarly, when dependent or vulnerable adults are involved, the  
30 vulnerable adult may need to be assessed at home, the risks assessed and any  
31 necessary safeguarding procedures initiated.

32

## 1 **5.5 SECONDARY MENTAL HEALTH CARE** 2 **REFERRAL TO SPECIALIST SUBSTANCE MISUSE** 3 **SERVICES**

### 4 **5.5.1 Referral threshold**

5 Specialist drug and alcohol services whether hospital (inpatient units) or  
6 community-based (community drug and alcohol teams) are dedicated to  
7 providing assessment and treatment for problematic drug / alcohol users, for  
8 example, heroin and cocaine and service users with alcohol problems. There  
9 is no reason why people with psychosis and coexisting substance misuse  
10 should be excluded from access to substance misuse services because of a  
11 diagnosis of psychosis.

12  
13 Referral from mainstream mental health services for specialist advice and  
14 joint working with specialist substance misuse services will occur where  
15 individuals with psychosis are known to be (although there will be variation  
16 between services):

- 17
- 18 • Severely dependent on alcohol **or**
- 19 • dependent on both alcohol and benzodiazepines **or**
- 20 • dependent on opioids.

21 As can be seen in Figure 3, tertiary referral allows access to more specialist  
22 skills and knowledge, and resources, including opiate prescribing and  
23 inpatient detoxification, residential rehabilitation, support or treatment  
24 groups.

25  
26 Because motivation is an important element of entry criteria to specialist  
27 addiction services secondary care staff may need to help individuals toward  
28 this readiness for change.

### 29 **5.5.2 Assessment and recognition**

30 The possible coexistence of a psychosis among people who come to specialist  
31 substance misuse services is often underestimated at least in part as a result of  
32 the complex clinical picture often presented when substance misuse is severe,  
33 involves the use of multiple substances and in people with evidence of  
34 personality disorder or other mental health problems. This is further  
35 complicated by that fact that substances may well be used to combat  
36 particular psychiatric symptoms or experiences such as anxiety, depression,  
37 intrusive thoughts, difficulties sleeping or more severe and troublesome  
38 experiences such as hallucinations. Moreover, significant life events, such as

1 bereavement, divorce and trauma, are frequently associated with the  
2 emergence of mental health problems, including relapse for people with  
3 psychosis, are commonly also triggers for the beginning of, or a significant  
4 increase in substance misuse. Furthermore, substance misuse may alter the  
5 presentation of symptoms, improving some and worsening others; this is  
6 especially so when a person is either intoxicated or experiencing withdrawal.  
7 For these, and many other reasons, assessment of mental state for people with  
8 substance misuse problems can prove to be difficult and recognition of a  
9 coexisting psychosis delayed.

10  
11 It is important that the assessment of people with a substance misuse problem  
12 is comprehensive, and may need to take place over several meetings over an  
13 extended period. It is also important to obtain additional information and  
14 history from friends, carers, chosen supporters or indeed advocates, where  
15 this is permitted and feasible. Ideally assessment will cover not only all the  
16 information needed for a substance misuse assessment and that needed for a  
17 mental health assessment, but it should also aim to examine how the  
18 individuals' behaviour, mental state and experiences co-vary (or not) with  
19 changing patterns of substance misuse; and how patterns of substance misuse  
20 may co-vary (or not) with changes in mental state; and how both substance  
21 misuse and mental state change in the light of different life events.

22 Understanding changes in mental state when someone misusing substances  
23 becomes either relatively or completely abstinent can be crucial in making the  
24 right diagnostic formulation, not least because communicative and cognitive  
25 functions can be greatly improved at these times. In any event, for some  
26 people where the index of suspicion for the coexistence of a psychosis with  
27 known substance misuse is high, use of the mental health act (for assessment)  
28 can be necessary and decisive.

### 29 **5.5.3 Interfaces and coordination**

30 Substance misuse services will normally need to work closely with secondary  
31 mental health services, to ensure that there are agreed local protocols derived  
32 from these guidelines that set out responsibilities and processes for  
33 assessment, referral, treatment and shared care across the whole care pathway  
34 for people with psychosis and coexisting substance misuse. This includes  
35 substance misuse professionals being available for care programme meetings  
36 for individuals receiving shared care with a secondary care mental health  
37 team. Secondary care community mental health services will usually need to  
38 continue to monitor and treat psychosis, and provide care co-ordination.

39  
40 Referral and signposting options will always need to be discussed with and  
41 agreed by the service user. There may be choice of agencies and it is

1 important that the service user is informed and involved in a shared decision.  
2 A range of Tier 2 and 3 drug and alcohol services will need to be considered  
3 in this respect (see section below), in line with the principle of the right care at  
4 the right intensity outlined in 5.2.2. Tier 2 examples would be information  
5 giving and signposting to mutual aid groups such as Alcoholics Anonymous  
6 or Narcotics Anonymous, and advice and linkage to needle exchanges  
7 provided by pharmacy, and other services. Specialist liver clinics, probation  
8 services and homeless or housing agencies are also interfaces to be managed  
9 and fostered.

10 Ensure there is clarity regarding the role of each service, clearly reflected in  
11 the care plan, with regular communication and appropriate information  
12 sharing between agencies.

13 It should be noted that effective coordination between statutory health and  
14 social care, non-statutory and voluntary organisations should be taken into  
15 account. Advocates working in voluntary organisations and other third  
16 sector groups will need to be involved in care planning and care  
17 programming where this is agreed with the service user.

#### 18 **5.5.4 Responsibility for prescribing**

19 Where a treatment plan is agreed involving secondary care and specialist  
20 substance misuse services the responsibility for any opiate substitute  
21 prescribing will need to be clearly agreed between the consultants for the two  
22 teams, incorporated into the service user's written care plan, and  
23 implemented according to the prescribing guidelines. Any doctor prescribing  
24 for the service user will need to see the service user regularly.

25  
26 Advice and guidelines on prescribing for service users with substance misuse  
27 problems, for example, on home alcohol detoxification programmes should be  
28 available from substance misuse services. Mental healthcare professionals  
29 working with people with psychosis and coexisting substance misuse will  
30 need to consider having supervision, advice, consultation and/or training  
31 from those with expertise in substance misuse specialist services to aid in  
32 developing and implementing treatment plans for substance misuse within  
33 secondary care mental health services.

#### 34 **5.5.5 Care Framework differences**

35 Individuals with coexisting psychosis and significant substance misuse will  
36 need to remain under the care of secondary care, managed within the Care  
37 Programme Approach. The term Care Programme Approach describes the  
38 approach used in secondary adult mental health care to assess, plan, review  
39 and co-ordinate the range of treatment, care and support needs for people in

1 contact with secondary mental health services who have complex  
2 characteristics

3  
4 Specialist drug services operate under Models of Care for Treatment of Adult  
5 Drug Misusers: Update 2006 (National Treatment Agency for Substance  
6 Misuse, 2006), whereas specialist alcohol services operate under Models of  
7 Care for Alcohol Misuse (Department of Health & National Treatment  
8 Agency for Substance Misuse, 2006). Both models of care utilise a four-tier  
9 framework and these refer to the level of the interventions provided and not  
10 the provider organisations:

- 11  
12 • Tier 1 interventions include provision of drug-related /alcohol-  
13 related information and advice, screening and referral. For alcohol  
14 tier 1 can also involve simple brief interventions.
- 15 • Tier 2 interventions include provision of drug-related information  
16 and advice, triage assessment, referral to structured drug treatment,  
17 brief psychosocial interventions, harm reduction interventions  
18 (including needle exchange) and aftercare. For alcohol interventions  
19 include provision of open access facilities and outreach that provide:  
20 alcohol-specific advice, information and support; extended brief  
21 interventions to help alcohol misusers reduce alcohol-related harm;  
22 and assessment and referral of those with more serious alcohol-  
23 related problems for care-planned treatment.
- 24 • Tier 3 interventions include provision of community-based  
25 specialised drug/ alcohol misuse assessment and co-ordinated care  
26 planned treatment and drug specialist liaison.
- 27 • Tier 4 interventions include provision of residential specialised  
28 drug / alcohol treatment, which is care planned and care  
29 coordinated to ensure continuity of care and aftercare.

## 31 **5.6 INPATIENT AND RESIDENTIAL SERVICES**

### 32 **5.6.1 Adult mental health services**

33 Substance misuse is a major problem within adult inpatient mental health  
34 settings. It is common amongst inpatients (Barnaby *et al.*, 2003; Bonsack *et al.*,  
35 2006; Phillips & Johnson, 2003; Sinclair *et al.*, 2008), with alcohol, cannabis and  
36 cocaine being the most commonly abused substances in inner urban settings.  
37 Service users with psychosis who abuse substances spend more time as

1 inpatients and are admitted more frequently (Isaac *et al.*, 2005; Menezes *et al.*,  
2 1996). Very high rates of cannabis use were found in a study of service users  
3 admitted to an inner urban Psychiatric Intensive Care Unit and those who  
4 continued to abuse cannabis (despite the best attempts of staff to restrict  
5 access to cannabis) spent longer in hospital (Isaac *et al.*, 2005).

6  
7 Violence is also a major cause of concern on acute inpatient wards (Healthcare  
8 Commission, 2007). Substance misuse has been identified by staff as an  
9 important contributor to violence on wards (Healthcare Commission, 2007).  
10 This is consistent with the epidemiological finding that most of the excess in  
11 serious offending behaviour seen in people with a diagnosis of schizophrenia  
12 occurs where there is co-morbid substance misuse disorder (Fazel *et al.*,  
13 2009b). In the substance-abusing population as a whole, cocaine and alcohol  
14 are particularly associated with violence (Macdonald *et al.*, 2008).

15  
16 Individuals with psychosis are usually admitted to a general adult mental  
17 health inpatient bed because of deterioration in their mental state and/or  
18 evidence of increased risk either to themselves or others. Substance misuse  
19 may be a co-incidental factor or play a causal role in the circumstances  
20 surrounding admission. In either case, assessment and management of the  
21 substance misuse will follow the general principles outlined above in other  
22 settings.

23  
24 The Department of Health has issued specific guidance about the  
25 management of people with coexisting mental illness and substance misuse  
26 being cared for in day hospital and inpatient settings (Department of Health,  
27 2006). Particular potential difficulties that face healthcare professionals in  
28 inpatient services include: the place and role of routine and occasional testing  
29 of biological samples (urine, blood, hair and, for alcohol, breath) as part of an  
30 agreed treatment plan; the requirement for policies on searching; and the  
31 practical management of episodes of substance misuse occurring in  
32 inpatients. This requires the development of local policies on the management  
33 of substances found on the premises, consideration of exclusion of visitors  
34 believed to be bringing-in illicit substances and good liaison with the police.  
35 For detained service users management of ongoing substance misuse may  
36 involve a review of the leave status of the service user and the appropriate  
37 level of security for safe and effective care.

38  
39 Admission of service users with coexisting opiate misuse and psychosis to an  
40 adult psychiatric inpatient unit is uncommon; but when it does it poses  
41 particular challenges. In this context it is imperative that an appropriate  
42 assessment by an expert in substance misuse and/or advice to the adult  
43 psychiatric team is available before developing a treatment plan for the opiate

1 misuse. The treatment plan will often include prescription of substitute  
2 opiates (methadone or buprenorphine). Healthcare professionals working  
3 within adult mental health services generally, and in inpatient settings in  
4 particular, need to be aware of current guidelines on the management of  
5 substance misuse provided by the National Treatment Agency (Department  
6 of Health, 2007).

### 7 **5.6.2 Secure mental health services**

8 Although substance misuse is a very significant problem within general adult  
9 mental health services, both in the community and especially on inpatient  
10 units, a significant past history of substance misuse is even more common  
11 amongst patients in secure care (Department of Health, 2006; D'Silva &  
12 Ferriter, 2003; Isherwood & Brooke, 2001). Inpatients in medium secure units  
13 report high levels of previous substance misuse, which has commonly  
14 continued after admission (Wyte *et al.*, 2004). Historically, dedicated  
15 substance misuse programmes were lacking within secure services despite the  
16 robust epidemiological evidence that links substance abuse and misuse with  
17 offending behaviour in people with a psychotic illness (Scott *et al.*, 2004).  
18 Secure services now commonly provide structured substance misuse  
19 interventions: these are only in the early stages of evaluation (Miles *et al.*,  
20 2007).

### 21 **5.6.3 Substance misuse inpatient services**

22 There is evidence that a diagnosis of psychosis is much more prevalent in  
23 people in contact with community substance misuse services than in the  
24 general population (Weaver *et al.*, 2003). There appears to be no data on the  
25 prevalence of psychosis that is not a consequence of substance misuse  
26 amongst inpatients in substance misuse services, who are admitted for  
27 detoxification. People who become or are recognised as being acutely  
28 psychotic whilst being treated in a substance misuse inpatient setting are  
29 often appropriately referred for treatment in general adult psychiatric  
30 inpatient services (an exception here is delirium tremens in the context of  
31 alcohol withdrawal, which is a medical emergency and would not occur in a  
32 competent inpatient setting providing alcohol withdrawal). There is no  
33 evidence that a diagnosis of a psychotic illness is a contra-indication for  
34 admission for treatment of coexisting substance misuse where the psychotic  
35 illness has been effectively treated.

### 36 **5.6.4 Residential and supported housing services**

37 Residential and supported housing services for people with a diagnosis of a  
38 psychotic illness inevitably work with people who abuse substances. The  
39 general principles of assessment, treatment and care set out above are

1 relevant to staff working in these settings; which will commonly be delivered  
2 through agencies other than the housing provider. There is a lack of evidence  
3 about how residential and supported housing services should work most  
4 effectively with people with psychosis and coexisting substance misuse  
5 although some practice guidance has been developed (Turning Point, 2007).

6  
7 Residential and supported housing services for people with substance misuse  
8 have in the past commonly been reluctant to take in people with psychotic  
9 illness, despite the fact that psychosis is common amongst substance misusers  
10 (Weaver *et al.*, 2003). The National Treatment Agency has identified a need for  
11 residential programmes that take account of the specific needs of “drug  
12 misusers with severe and enduring mental health problems” (National  
13 Treatment Agency, 2006). There is no evidence that a diagnosis of a psychotic  
14 illness is a contra-indication for residential rehabilitative services for people  
15 with coexisting substance misuse where the psychotic illness has been  
16 effectively treated.

### 17 **5.6.5 Prison mental health services and criminal justice**

18 The Bradley Report (Department of Health 2009a) and the subsequent  
19 Government response and delivery plan (Department of Health 2009b)  
20 focuses on people with mental health and learning disabilities who become  
21 involved with the criminal justice system and makes wide ranging  
22 recommendations. The report recognizes the prevalence of psychosis with  
23 coexisting substance misuse in this population and makes a specific  
24 recommendation to develop improved services in prisons for these prisoners.  
25 Current problems within this system echo those outside:

26  
27 *“Mental health services and substance misuse services in prisons do not currently*  
28 *work well together; national policy is developed separately for mental health and*  
29 *for substance misuse, and this is reflected on the ground, where dual diagnosis is*  
30 *used as a reason for exclusion from services rather than supporting access”*

31 *(p16 executive summary*

32 [http://www.dh.gov.uk/prod\\_consum\\_dh/groups/dh\\_digitalassets/documents/digitalasset/dh\\_098699.pdf](http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/documents/digitalasset/dh_098699.pdf).

34  
35 In terms of the care pathway the report calls for liaison and court diversion  
36 services to reduce the need for custodial interventions and allow access to  
37 appropriate treatment at an earlier stage in their offending behaviour. The  
38 Bradley Report also calls for better links into community mental health  
39 provision when people are leaving prison with psychosis and coexisting  
40 substance misuse.

## 1 **5.7 FROM EVIDENCE TO RECOMMENDATIONS**

2 There is only a limited amount of empirical evidence about the prevalence,  
3 pattern and epidemiology of different combinations of coexisting psychosis  
4 and substance misuse. Such information is necessary to target resources at  
5 groups most at risk of very poor outcomes, to determine whether early  
6 intervention efforts might be more effective than interventions for long-  
7 standing comorbidity and to investigate whether different interventions are  
8 required for different diagnostic groups and types of substance. In addition,  
9 little research is available to determine how healthcare professionals should  
10 work together to provide the most appropriate care and treatment for people  
11 with psychosis and coexisting substance misuse. And, what evidence we  
12 have, in this and other chapters, is often collected in different countries, such  
13 as the US, where the interventions, the training and competence of  
14 professionals, the configuration of the healthcare system, and in particular,  
15 what counts as 'standard care' may be very different. The GDG, nevertheless,  
16 extrapolated where this was possible and useful. The following  
17 recommendations are, therefore, developed through an iterative process,  
18 synthesising our collective experience to develop a framework of good  
19 practice recommendations that we hope will support healthcare professionals  
20 develop services in mental health, and substance misuse services in  
21 particular, so that people with psychosis and coexisting substance misuse can  
22 receive the care and treatment most likely to bring benefit and to improve  
23 their lives and those of their carers.

24  
25 The recommendations for good practice concerned a number of topics: 1)  
26 recognition, 2) primary care, 3) secondary care mental health services, 4)  
27 substance misuse services, and 5) working with adults and young people with  
28 psychosis and coexisting substance misuse.

29  
30 With regard to recognition, given that substance misuse is usual rather than  
31 exceptional among people with psychosis, the GDG felt it was vital that  
32 healthcare professionals in all settings ask service users about substance use,  
33 and where appropriate, an assessment of dependency should be conducted  
34 using the existing NICE guidelines on drug misuse (REF) and alcohol use  
35 disorders (REF). Likewise, in people with known or suspected substance  
36 misuse, there should be an assessment for possible psychosis.

37  
38 In primary care, the GDG felt that there was a clear rationale (supported by  
39 DH guidance) to recommend that people with psychosis or suspected  
40 psychosis, including those who are suspected of having coexisting substance  
41 misuse problems, should be referred to either secondary care mental health  
42 services or CAMHS for assessment and further management. Likewise,

1 people with substance misuse or suspected substance misuse who are  
2 suspected of having coexisting psychosis, should be referred to either  
3 secondary care mental health services or CAMHS.

4  
5 In secondary care mental health services, the GDG felt there was a need to  
6 recommend that healthcare professionals should ensure they are competent in  
7 the recognition, treatment and care of people with psychosis and coexisting  
8 substance misuse. In addition, mental health professionals should consider  
9 having supervision, advice, consultation and/or training from specialists in  
10 substance misuse services. The GDG considered that this would aid in the  
11 development and implementation of treatment plans for substance misuse  
12 within CAMHS or adult community mental health services. Also, because  
13 adults and young people with psychosis and coexisting substance misuse are  
14 often excluded from age-appropriate services for no justifiable reason, the  
15 GDG felt there was a strong rationale for recommending against exclusion.  
16 Finally, the GDG made a number of recommendations covering the process of  
17 assessment and the use of biological/physical testing. With regard to the  
18 latter, the GDG felt there was a place for testing when used as part of a care  
19 plan if this is agreed to by the service user. After a great deal of discussion,  
20 the decided that biological or physical testing should not be used in routine  
21 screening for substance misuse. This applies in inpatient settings, and where  
22 mental capacity is lacking, healthcare professionals should refer to the Mental  
23 Capacity Act.

24  
25 In substance misuse services, the GDG felt there was a clear need to make a  
26 recommendation that healthcare professionals should be competent to  
27 recognise the signs and symptoms of psychosis, and undertake a mental  
28 health needs and risk assessment with sufficient ability to know how and  
29 when to refer to secondary care mental health services. The GDG also felt that  
30 recommendations for joint working needed to be made as this was not, in  
31 their experience, done well.

32  
33 When working with people with psychosis and coexisting substance misuse,  
34 the GDG thought that a number of safeguarding issues were important and  
35 needed recommendations. In addition, the GDG felt that voluntary sector  
36 organisations had an important role to play in lives of people with psychosis  
37 and coexisting substance misuse, therefore, recommendations were made  
38 about collaborative working.

39  
40 Although there is a paucity of evidence regarding all aspects of assessment  
41 and care pathways, the GDG felt that two research recommendations should  
42 be given priority. First, as described above, the prevalence, risk and protective  
43 factors, and course of illness for different combinations of psychosis and

1 coexisting substance misuse needs to be examined. Second, there are cogent  
2 reasons given the high prevalence of substance misuse amongst service users  
3 with a psychosis that staff working within psychosis services develop as part  
4 of their basic training and continuing professional development, skills and  
5 knowledge in substance misuse assessment and treatment interventions.  
6 More research is required on how this training is provided and the impact of  
7 ongoing supervision when working with people with psychosis and  
8 coexisting substance misuse. The GDG considered that the responsibility for  
9 monitoring the physical health of people with psychosis and coexisting  
10 substance misuse should remain in primary care as recommended in the  
11 NICE guideline on schizophrenia (REF).  
12

## 1 **5.8 CLINICAL PRACTICE RECOMMENDATIONS**

### 2 **5.8.1 Recommendations**

#### 3 *Recognition of psychosis with coexisting substance misuse*

4 **5.8.1.1** Healthcare professionals in all settings, including primary care,  
5 secondary care mental health services, CAMHS and accident and  
6 emergency departments, and those in prisons and criminal justice  
7 mental health liaison schemes, should routinely ask adults and young  
8 people with known or suspected psychosis about their use of alcohol  
9 and/or prescribed and non-prescribed (including illicit) drugs. If the  
10 person has used substances ask them about all of the following:

- 11 • particular substance(s) used
- 12 • quantity, frequency and pattern of use
- 13 • route of administration
- 14 • duration of current level of use.

15

16 In addition, conduct an assessment of dependency. [See 'Drug  
17 misuse: opioid detoxification' (NICE clinical guideline 52) and  
18 'Alcohol use disorders: diagnosis, assessment and management of  
19 harmful drinking and alcohol dependence' (NICE clinical guideline,  
20 forthcoming)], and also seek corroborative evidence from families,  
21 carers or chosen supporters, where this is possible and permission is  
22 given.

23 **5.8.1.2** Healthcare professionals in primary care, secondary care mental health  
24 services, CAMHS and specialist substance misuse services should  
25 routinely assess adults and young people with known or suspected  
26 substance misuse for possible psychosis. Seek corroborative evidence  
27 from families, carers or chosen supporters, where this is possible and  
28 permission is given.

#### 29 *Primary care*

### 30 **Referral from primary care**

31 **5.8.1.3** Refer all adults and young people with psychosis or suspected  
32 psychosis, including those who are suspected of coexisting substance  
33 misuse, to either secondary care mental health services or CAMHS for  
34 assessment and further management.

1 **5.8.1.4** Refer all adults and young people with substance misuse or suspected  
2 substance misuse who are suspected of having coexisting psychosis to  
3 secondary care mental health services or CAMHS for assessment and  
4 further management

5 **Physical healthcare**

6 **5.8.1.5** Monitor the physical health of adults and young people with psychosis  
7 and coexisting substance misuse, as described in the guideline on  
8 schizophrenia (NICE clinical guideline 82). Pay particular attention to  
9 the impact of alcohol and drugs (prescribed and non-prescribed) on  
10 physical health. Monitoring should be conducted at least once a year  
11 or more frequently if the person has a significant physical illness or  
12 there is a risk of physical illness because of substance misuse.

13 *Secondary care mental health services*

14 **Competence**

15 **5.8.1.6** Healthcare professionals working within secondary care mental  
16 health services should ensure they are competent in the recognition,  
17 treatment and care of adults and young people with psychosis and  
18 coexisting substance misuse.

19 **5.8.1.7** Healthcare professionals working within secondary care mental  
20 health services with adults and young people with psychosis and  
21 coexisting substance misuse should consider having supervision,  
22 advice, consultation and/or training from specialists in substance  
23 misuse services. This is to aid in the development and  
24 implementation of treatment plans for substance misuse within  
25 CAMHS or adult community mental health services.

26 **Pathways into care**

27 **5.8.1.8** Do not exclude adults and young people with psychosis and  
28 coexisting substance misuse from age-appropriate mental healthcare  
29 because of their substance misuse.

30 **5.8.1.9** Do not exclude adults and young people with psychosis and  
31 coexisting substance misuse from age-appropriate substance misuse  
32 services because of a diagnosis of psychosis.

33 **Assessment**

1 **5.8.1.10** Adults and young people with psychosis and coexisting substance  
2 misuse attending secondary care mental health services should be  
3 offered a comprehensive, multidisciplinary assessment, including  
4 assessment of **all** of the following:

- 5 • personal history
- 6 • mental, physical and sexual health
- 7 • social, family and economic situation
- 8 • accommodation, including history of homelessness and  
9 stability of current living arrangements
- 10 • current and past substance misuse and its impact upon their  
11 life, health and response to treatment
- 12 • criminal justice history and current status
- 13 • personal strengths and weaknesses and readiness to change  
14 their substance use and other aspects of their lives.

15 The assessment may need to take place over several meetings to gain  
16 a full understanding of the person and the range of problems they  
17 experience, and to promote engagement.

18 **5.8.1.11** When assessing adults and young people with psychosis and  
19 coexisting substance misuse, seek corroborative evidence from  
20 families, carers or chosen supporters where this is possible and  
21 permission is given. Summarise the findings, share this with the  
22 person and record it in their care plan.

23 **5.8.1.12** Review any changes in the person's use of substances. This should  
24 include changes in:

- 25 • the way the use of substances affects the person over time
- 26 • patterns of use
- 27 • mental and physical state
- 28 • circumstances and treatment.

29 Share the summary with the person and record it in their care plan.

30 **5.8.1.13** When assessing adults and young people with psychosis and  
31 coexisting substance misuse, be aware that low levels of substance  
32 use that would not usually be considered harmful or problematic in  
33 people without psychosis, can have a significant impact on the mental  
34 health of people with psychosis.

35 **5.8.1.14** Regularly assess and monitor risk of harm to self and/or others and  
36 develop and implement a risk management plan to be reviewed when  
37 the service users' circumstances or levels of risk change. Specifically  
38 consider additional risks associated with substance misuse, including:

- 1                   • physical health risks (for example, withdrawal seizures,  
2                   delirium tremens, blood-borne viruses, accidental overdose,  
3                   and interactions with prescribed medication) **and**  
4                   • the impact that substance use may have on other risks such  
5                   as self-harm, suicide, self-neglect, violence, abuse of or by  
6                   others, exploitation, accidental injury and offending  
7                   behaviour.

8   **5.8.1.15** When developing a care plan for an adult or young person with  
9            psychosis and coexisting substance misuse, take account of the  
10           complex and individual relationships between substance misuse,  
11           psychotic symptoms, emotional state, behaviour and the person's  
12           social context.

### 13   **Biological/physical testing**

14   **5.8.1.16** Biological or physical tests for substance use (such as blood and urine  
15            tests or hair analysis) may be useful in the assessment, treatment and  
16           management of substance misuse for adults and young people with  
17           psychosis. However, this should be agreed with the person first as  
18           part of their care plan. Do not use biological or physical tests in  
19           routine screening for substance misuse in adults and young people  
20           with psychosis.

21   **5.8.1.17** Biological or physical tests for substance use should only be  
22            considered in inpatient services as part of the assessment and  
23           treatment planning for adults and young people with psychosis and  
24           coexisting substance misuse. Obtain consent for these tests and  
25           inform the person of the results as part of an agreed treatment plan.  
26           Where mental capacity is lacking, refer to the Mental Capacity Act  
27           (2005).

### 28   *Substance misuse services*

#### 29   **Competence**

30   **5.8.1.18** Healthcare professionals in substance misuse services should be  
31            competent to:

- 32                   • recognise the signs and symptoms of psychosis  
33                   • undertake a mental health needs and risk assessment  
34                   sufficient to know how and when to refer to secondary care  
35                   mental health services.

#### 36   **Assessment**

1 **5.8.1.19** Adults and young people with psychosis and coexisting substance  
2 misuse attending substance misuse services should be offered a  
3 comprehensive, multidisciplinary mental health assessment in  
4 addition to an assessment of their substance misuse.

5 **Joint working**

1 **5.8.1.20** Healthcare professionals in substance misuse services should be  
2 present at care programme approach meetings for adults and young  
3 people with psychosis and coexisting substance misuse within their  
4 service who are also receiving treatment and support in other health  
5 services.

6 **5.8.1.21** Specialist substance misuse services should provide advice,  
7 consultation, and training for healthcare professionals in adult mental  
8 health services and CAMHS regarding the assessment and treatment  
9 of substance misuse, and of substance misuse with coexisting  
10 psychosis.

11 **5.8.1.22** Specialist substance misuse services should work closely with  
12 secondary care mental health services to develop local protocols  
13 derived from this NICE guideline for adults and young people with  
14 psychosis and coexisting substance misuse. The agreed local  
15 protocols should set out responsibilities and processes for assessment,  
16 referral, treatment and shared care across the whole care pathway.

17 *Working with adults and young people with psychosis and*  
18 *coexisting substance misuse*

#### 19 **Safeguarding issues**

20 **5.8.1.23** If people with psychosis and coexisting substance misuse are parents  
21 or carers of children or young people, ensure that the child's or young  
22 person's needs are assessed according to local safeguarding  
23 procedures<sup>7</sup>.

24 **5.8.1.24** If children or young people being cared for by people with psychosis  
25 and coexisting substance misuse are referred to CAMHS under local  
26 safeguarding procedures:

- 27 • use a multi-agency approach, including social care and  
28 education, to ensure that various perspectives on the child's  
29 life are considered
- 30 • consider using the Common Assessment Framework<sup>8</sup>;  
31 advice on this can be sought from the local named lead for  
32 safeguarding.

33 **5.8.1.25** If serious concerns are identified, health or social care professionals  
34 working with the child or young person (see 5.8.1.23) should develop  
35 a child protection plan.

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<sup>7</sup> [www.safeguardingchildren.org.uk](http://www.safeguardingchildren.org.uk)

<sup>8</sup> [www.dcsf.gov.uk/everychildmatters/strategy/deliveringservices1/caf/cafframework](http://www.dcsf.gov.uk/everychildmatters/strategy/deliveringservices1/caf/cafframework)

1 **5.8.1.26** When working with people with psychosis and coexisting substance  
2 misuse who are responsible for vulnerable adults, ensure that the  
3 home situation is risk assessed and that safeguarding procedures are  
4 in place for the vulnerable adult. Advice on safeguarding vulnerable  
5 adults can be sought from the local named lead for safeguarding.

6 **5.8.1.27** Consider adults with psychosis and coexisting substance misuse for  
7 assessment according to local safeguarding procedures for vulnerable  
8 adults if there are concerns regarding exploitation or self-care, or if  
9 they have been in contact with the criminal justice system.

## 10 **Working with the voluntary sector**

11 **5.8.1.28** Healthcare professionals in primary care and secondary care mental  
12 health services, and in specialist substance misuse services, should  
13 work collaboratively with voluntary sector organisations that provide  
14 help and support for adults and young people with psychosis and  
15 coexisting substance misuse. Ensure that advocates from such  
16 organisations are included in the care planning and care  
17 programming process wherever this is possible and agreed by the  
18 person with psychosis and coexisting substance misuse.

19 **5.8.1.29** Healthcare professionals in primary care and secondary care mental  
20 health services, and in specialist substance misuse services, should  
21 work collaboratively with voluntary sector organisations providing  
22 services for adults and young people with psychosis and coexisting  
23 substance misuse to develop agreed protocols for routine and crisis  
24 care.

## 25 **5.8.2 Research recommendations**

26 **5.8.2.1** What are the prevalence, risk and protective factors, and course of  
27 illness for different combinations of psychosis and coexisting  
28 substance misuse (for example, schizophrenia and cannabis misuse or  
29 bipolar disorder and alcohol misuse)?

30 **5.8.2.2** What and how should training be provided to healthcare professionals  
31 working with people with psychosis and substance misuse?

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# 1 **6 SERVICE DELIVERY MODELS** 2 **FOR PEOPLE WITH PSYCHOSIS** 3 **AND COEXISTING SUBSTANCE** 4 **MISUSE**

## 5 **6.1 INTRODUCTION**

6 This chapter looks at models of service delivery for people with psychosis and  
7 coexisting substance misuse. These models are means by which therapeutic  
8 interventions and supports are provided. Two broad questions are addressed  
9 in this chapter. First, is there evidence that providing therapeutic  
10 interventions and support relevant to both conditions in an integrated fashion  
11 (the same team addressing both issues), is superior to these interventions  
12 being provided separately? Second, is there evidence about the role of staffed  
13 accommodation and inpatient care in the management of coexisting substance  
14 misuse and psychosis?  
15

16 In reviewing the evidence for the effectiveness of different service delivery  
17 models, the GDG decided to focus on RCTs. By using this type of study  
18 design to evaluate service-level interventions there are specific problems  
19 relating to defining such interventions precisely; for example, the  
20 'intervention' and 'standard care' may vary between studies, between  
21 countries and over time; and experimental interventions have a tendency to  
22 overlap with standard care. Service-level interventions that claim superiority  
23 over other methods of care delivery must be able to characterise clearly what  
24 they do, how they do it, and how they differ from alternative types of service  
25 and from the standard care they hope to replace. For these reasons, it is  
26 essential for new services to be subjected to the rigour of evaluation through  
27 RCTs; services must be able to demonstrate their overall value in comparison  
28 with other interventions to remain a supportable component of care within  
29 the NHS. Other types of study design (that is, longer-term observational  
30 studies), might help to differentiate, evaluate and refine services and the ways  
31 in which they operate. For this reason, a narrative synthesis of observational  
32 studies was conducted after the review of RCTs.

## 1 **6.2 INTEGRATED SERVICE MODELS**

### 2 **6.2.1 Introduction**

3 Both in the UK, and elsewhere in the world, it has been proposed that  
4 effective treatment for people with psychosis and coexisting substance misuse  
5 usually requires an integrated treatment approach (Department of Health,  
6 2002; Ziedonis *et al.*, 2005). An integrated approach combines elements of  
7 mental health and substance misuse service models in one delivery system.  
8 This approach was originally pioneered in the US in the 1980s, and was  
9 developed in contrast with traditional treatment approaches that provided  
10 separate services either in parallel or sequentially (Mueser & Drake, 2003).  
11 Such services were felt unable to meet the needs of people with severe mental  
12 health and drug/alcohol problems; typically, service users perhaps got only  
13 one or the other component, or incompatible or inconsistent treatments from  
14 both, or worse still, fell somewhere between the two and received little care  
15 (Drake *et al.*, 2008). It was proposed that integrated care meant that both  
16 mental health and substance misuse treatments could be provided from the  
17 same team of clinicians at the same time and in an integrated manner. The  
18 potential advantages of such an integrated approach include ensuring that  
19 both elements of the dual problems are given attention, and that any  
20 interactions between mental health and substance use problems are  
21 formulated and addressed. Due to differences in service provision,  
22 organisation funding, and treatment philosophies in the UK, as compared  
23 with the US, it has been suggested that more shared care with drug and  
24 alcohol services is feasible in the UK (Graham *et al.*, 2003). Moreover, current  
25 Department of Health policy suggests that the main focus for service delivery  
26 should be within mental health services, and a key principle should be that  
27 both problems and the relationship between them are addressed  
28 simultaneously (Department of Health, 2002).

29  
30 Integrated service delivery models that have been evaluated have involved  
31 changes in the health care systems to encompass intervention components  
32 delivered in a variety of service configurations. Services have included a  
33 number of different elements delivered in different combinations and with  
34 differing intensities, including motivational interventions and various forms  
35 of group, individual, and family counselling as well as housing interventions  
36 (Mueser *et al.*, 2005). Besides differing in the components of intervention  
37 offered, integrated service delivery models have also differed in structural  
38 form: varying from different case management models in community mental  
39 health teams, to more intensive, outreach oriented services, and there have  
40 also been evaluations of staffed accommodation (usually comparisons of  
41 residential integrated treatment with non-residential treatment).

1 *Definition of intervention*

2 **Integrated service models**

3 Integrated service models were defined as those that unify services  
4 at the provider level rather than requiring service users to negotiate  
5 separate mental health and substance abuse treatment programmes  
6 (Cleary *et al.*, 2008; Drake *et al.*, 1993).

7 **Standard care**

8 This was defined as the usual treatment received from a community  
9 mental health team (which will include a care coordinator) with the  
10 potential to access separate substance misuse services.

11 **6.2.2 Clinical review protocol (integrated service models)**

12 The review protocol, including the review question, information about the  
13 databases searched and the eligibility criteria used for this section of the  
14 guideline can be found in Table 10. During the early stages of guideline  
15 development, a recent Cochrane review (Cleary *et al.*, 2008) and related peer-  
16 reviewed publication (Cleary *et al.*, 2009) were identified that addressed the  
17 review question. These systematic reviews were used as a source of evidence,  
18 and only a new systematic search for more recent primary-level studies was  
19 conducted for the guideline (further information about the search strategy can  
20 be found in Appendix 7).

21  
22 Where evidence allowed, the following two sub-questions were addressed: 1)  
23 What are the elements in an integrated service model that are most likely to  
24 be associated with better outcomes? 2) Are there any subgroups of people (for  
25 example, young people, BME groups) that benefit from some elements of the  
26 service model more than others?

27

**Table 10: Clinical review protocol for the review of integrated service models**

Component	Description
Review question	1.2.1 In people with psychosis and coexisting substance misuse, does an integrated service model (usually involving the model of assertive community treatment) when compared with an alternative management strategy lead to improved outcomes?
Electronic databases	CENTRAL, CINAHL, EMBASE, MEDLINE, PsycINFO
Date searched	01.01.2008 to 26.05.2010 <sup>1</sup>
Study design	RCTs and observational studies
Population	People with psychosis and coexisting substance misuse
Intervention(s)	Integrated service model (usually involving the model of assertive community treatment)
Comparison	Alternative management strategies
Critical outcomes	<ul style="list-style-type: none"> <li>• Reduced mortality (all causes)</li> <li>• Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</li> <li>• Reduced substance misuse (however measured)</li> <li>• Improved global and social functioning (for example, employment, accommodation)</li> <li>• Improved subjective quality of life</li> <li>• Improved satisfaction with care</li> <li>• Reduced physical morbidity.</li> </ul>
Note. RCT = Randomised controlled trial.	
<sup>1</sup> The search is an update to Cleary <i>et al.</i> (2008) and Cleary <i>et al.</i> (2009).	

1

### 2 **6.2.3 Studies considered for review (integrated service models)<sup>9</sup>**

3 Four RCTs, CHANDLER2006 (Chandler & Spicer, 2006), DRAKE1998 (Drake  
4 *et al.*, 1998), ESSOCK2006 (Essock *et al.*, 2006), MORSE2006 (Morse *et al.*, 2006),  
5 that were included in the review by Cleary *et al.* (2008), met the eligibility  
6 criteria for this review. Of these, all were published in peer-reviewed journals  
7 between 1998 and 2006. In addition, one RCT identified during the search for  
8 new evidence (Craig *et al.*, 2008), was excluded from the meta-analysis  
9 because the GDG considered this to be a trial of training that was not  
10 comparable to other trials included in the analysis. Further information about  
11 this study can be found in section 5.2.3. Full study characteristics (and any  
12 associated references), as well as a list of excluded studies can be found in  
13 Appendix 13.  
14

<sup>9</sup> Here and elsewhere in the guideline, each RCT considered for review is referred to by a study ID (primary author and date of study publication, except where a study is in press or only submitted for publication, then a date is not used).

1 Of the four included RCTs, there were two involving a comparison of an  
2 integrated service model versus standard care (CHANDLER2006,  
3 MORSE2006). MORSE2006 also included an intervention group receiving  
4 non-integrated assertive community treatment (ACT), allowing a comparison  
5 between integrated and non-integrated ACT (see Table 11 for summary  
6 information). In addition, there were two trials involving a comparison of  
7 integrated ACT versus integrated standard case management (DRAKE1998,  
8 ESSOCK2006) (see Table 12 for summary information).

9

10 In addition to the RCTs, three observational studies (Drake *et al.*, 1997; Ho *et*  
11 *al.*, 1999; Mangrum *et al.*, 2006), that were included in the review by Cleary  
12 and colleagues (2008), met eligibility criteria for this review. All studies were  
13 published in peer-reviewed journals between 1997 and 2006.

14

15 Of the three observational studies, there was one involving a comparison of  
16 an integrated service model versus a parallel service model (Mangrum *et al.*,  
17 2006), one before-and-after study of a 'dual-diagnosis treatment program' (Ho  
18 *et al.*, 1999), and one comparing an integrated service model with standard  
19 care (Drake *et al.*, 1997) (see section 6.2.5 for further information about each  
20 study and a narrative summary of results).

**Table 11. Study information table for RCTs comparing an integrated service model with a non-integrated management strategy**

	<b>Integrated service model (ACT/DDT) versus standard care</b>	<b>Integrated ACT versus non-integrated ACT</b>
Total no. of trials (N)	2 RCTs (277)	1 RCT (100)
Study ID	(1) CHANDLER2006 (2) MORSE2006	(1) MORSE2006
Number randomised	(1) 182 (2) 95	(1) 100
Diagnosis	(1) 66% DSM-IV schizophrenia, schizoaffective disorder, bipolar or psychotic disorder NOS and 100% current substance use disorder (34% alcohol dependence, 47% drug dependence) <sup>1</sup> (2) 89% DSM-IV schizophrenia, schizoaffective, atypical psychotic disorder or bipolar disorder; 9% major depression-recurrent disorder, 2% other. All had one or more substance use disorders; 46% substance dependence disorder for alcohol and/or drugs; 64% substance abuse disorder for alcohol and/or drugs, 40% an alcohol-only diagnosis, 18% drug-only diagnosis, 42% had both drug and alcohol disorders - cocaine most frequently used drug (34%) cannabis (19%)	(1) 89% DSM-IV schizophrenia, schizoaffective, atypical psychotic disorder or bipolar disorder; 9% major depression-recurrent disorder, 2% other. All had one or more substance use disorders; 46% substance dependence disorder for alcohol and/or drugs; 64% substance abuse disorder for alcohol and/or drugs, 40% an alcohol-only diagnosis, 18% drug-only diagnosis, 42% had both drug and alcohol disorders - cocaine most frequently used drug (34%) cannabis (19%)
Ethnicity	(1) 66% African American, 21% White (2) 73% African American, 25% White	(1) 73% African American, 25% White, 2% other
Treatment length	(1) 36 months (2) 24 months	(1) 24 months
Country	(1) USA (2) USA	(1) USA
Intervention (n)	(1) In-custody standard care + brief aftercare + Integrated DDT (post-custody, participants received MI, substance abuse counselling, group treatment oriented to both disorder, family psychoeducation regarding 'dual disorders', multidisciplinary team, integrated substance abuse specialists, stagewise interventions, time unlimited services, outreach etc.) (n=103) <sup>2</sup> (2) Integrated ACT (n=46)	(1) Integrated ACT (n=46)
Control (n)	(1) In-custody standard care + usual post custody services + 60 days of post release case management and housing assistance (n=79)	(1) Non-integrated ACT. Referred service users to other community providers for outpatient or individual substance abuse services and to 12-

	(2) Provided with a list of community agencies (mental health and substance abuse treatment) and staff provided linkage assistance to facilitate access (n=49)	step groups (n=54)
<p><i>Note.</i> ACT = Assertive Community Treatment; DDT = Dual Disorders Treatment; DSM-IV = Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (American Psychiatric Association, 1994); MI = motivational interviewing; N = Total number of participants; n = number of participants in each group; RCT = Randomised controlled trial.</p> <p><sup>1</sup> Some participants had more than one dependence.</p> <p><sup>2</sup> Before release from custody, all participants received an intervention including intensive assessment, medications, treatment planning in preparation for discharge, consultation with jail staff, one-to-one counselling, and crisis intervention (for more details about the intervention, see Mercer-McFadden <i>et al.</i> 1998).</p>		

- 1
- 2

**Table 12. Study information table for RCTs comparing integrated ACT with integrated standard case management**

	<b>Integrated ACT versus integrated standard case management</b>
Total no. of trials (N)	2 RCTs (421)
Study ID	(1) DRAKE1998 (2) ESSOCK2006
Number randomised	(1) 223 (2) 198
Diagnosis	(1) 53% DSM-III-R schizophrenia with active DSM-III-R substance use disorder (73% alcohol abuse, 42% drug abuse) <sup>1</sup> (2) 76% DSM-III-R schizophrenia, 17% mood disorder with co-occurring DSM-III-R substance use disorder ( 74% alcohol abuse, 81% other substances) <sup>1</sup>
Ethnicity	(1) 96% White (2) 55% African American, 27% White
Treatment length	(1) 36 months (2) 36 months
Country	(1) USA (2) USA
Intervention (n)	(1) Integrated ACT: community-based, high intensity, direct substance abuse treatment by team members, use of stage-wise 'dual-disorder' model, 'dual-disorder' treatment groups & exclusive team focus on service users for those with psychosis and coexisting substance misuse. Caseload ~ 12 (n=109) (2) Integrated ACT with a direct substance use component (n=99)
Control (n)	(1) Standard case management: community-based, team working with service user's support system & vigorously addressing co-occurring substance use. Caseload ~ 25 (n=114) (2) Standard case management: some services provided directly and teams had training from study authors in integrated treatment, including comprehensive assessment, individual motivational interviewing, group treatments, and stagewise interventions (n=99)
<i>Note.</i> ACT = Assertive Community Treatment; DSM-IV = Diagnostic and Statistical Manual of Mental Disorders, 4th Edition (American Psychiatric Association, 1994); N = Total number of participants; n = number of participants in each group; RCT = Randomised controlled trial. <sup>1</sup> Some participants had more than one dependence.	

1

2

### 3 **6.2.4 Evidence from RCTs (integrated service models)**

4 Meta-analysis was used to synthesise the evidence for each comparison. For  
5 the comparison of an integrated service model with a non-integrated  
6 management strategy, a GRADE summary of findings table is shown in Table  
7 13 and Table 14. For the comparison of integrated ACT with integrated  
8 standard case management, a GRADE summary of findings table is shown in  
9 Table 15.

10

- 1 The forest plots and full GRADE evidence profiles can be found in Appendix
- 2 14 and 15, respectively.
- 3

**Table 13. GRADE summary of findings table for RCTs comparing integrated ACT with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Substance use rating			
by 6 months	SMD 0.19 (-0.21 to 0.59)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.27 (-0.14 to 0.67)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	SMD 0.12 (-0.29 to 0.52)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.12 (-0.28 to 0.53)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 2. Days used substances			
6 months	SMD 0.08 (-0.33 to 0.48)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.11 (-0.3 to 0.51)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	SMD 0.09 (-0.31 to 0.49)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.13 (-0.28 to 0.53)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Service use: 1. Days in stable community residences (not in hospital)			
by 6 months	MD 3.17 (-0.52 to 6.86)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	MD 2.84 (-2.07 to 7.75)	95 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	MD 6.46 (1.36 to 11.56)	95 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 24 months	MD 5.70 (0.59 to 10.81)	95 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
<p><i>Note.</i> Negative SMDs favour integrated service models, positive MDs favour integrated service models; CI = confident interval; MD = mean difference; SMD = Standardised mean difference.</p> <p><sup>1</sup> Optimal information size (for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> MORSE2006</p>			

4

**Table 14. GRADE summary of findings table for RCTs comparing integrated ACT with non-integrated ACT**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Substance use rating			
by 6 months	SMD 0.14 (-0.25 to 0.53)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.18 (-0.22 to 0.57)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	SMD -0.15 (-0.54 to 0.25)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.05 (-0.34 to 0.44)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 2. Days used substances			
6 months	SMD 0.09 (-0.31 to 0.48)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.27 (-0.12 to 0.67)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	SMD 0.09 (-0.30 to 0.48)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.08 (-0.32 to 0.47)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Service use: 1. Days in stable community residences (not in hospital)			
by 6 months	MD 2.42 (-1.01 to 5.85)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	MD 0.31 (-4.42 to 5.04)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	MD -1.18 (-5.94 to 3.58)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 24 months	MD 0.51 (-4.36 to 5.38)	100 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
<p><i>Note.</i> Negative SMDs favour integrated service models, positive MDs favour integrated service models; CI = confident interval; MD = mean difference; SMD = Standardised mean difference.</p> <p><sup>1</sup> Optimal information size (for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> MORSE2006</p>			

- 1
- 2
- 3

**Table 15. GRADE summary of findings table for RCTs comparing integrated ACT with integrated standard case management**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Death - by 36 months	RR 1.18 (0.39 to 3.57)	421 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
Substance use: 1. Not in remission			
by 36 months - alcohol	RR 1.15 (0.84 to 1.56)	143 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 36 months - drugs	RR 0.89 (0.63 to 1.25)	85 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 2. Substance abuse (SATS)			
by 6 months	SMD 0.03 (-0.17 to 0.23)	379 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 12 months	SMD 0.08 (-0.23 to 0.39)	374 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 18 months	SMD -0.02 (-0.22 to 0.19)	375 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 24 months	SMD 0.11 (-0.14 to 0.37)	365 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 30 months	SMD 0.11 (-0.1 to 0.31)	358 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 36 months	SMD 0.05 (-0.15 to 0.26)	360 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
Service use: 1. Days in stable community residences (not in hospital)			
by 12 months	MD -10 (-38.61 to 18.6)	378 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
by 24 months	MD 8.54 (-4.46 to 21.55)	377 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
by 36 months	MD 5.17 (-9.2 to 19.55)	364 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
Functioning: 1. Average general score (GAS)			
by 6 months	SMD 0.13 (-0.18 to 0.43)	162 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.07 (-0.23 to 0.38)	171 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 18 months	SMD 0.11 (-0.18 to 0.41)	176 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.18 (-0.13 to 0.48)	166 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 30 months	SMD -0.06 (-0.37 to 0.24)	164 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 36 months	SMD 0.04 (-0.26 to 0.34)	170 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
Satisfaction: Average general score (QOLI)			
by 6 months	SMD 0.06 (-0.17 to 0.29)	377 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
by 12 months	SMD 0.01 (-0.2 to 0.23)	370	Low <sup>1,2</sup>

		(2 studies) <sup>3,4</sup>	
by 18 months	SMD 0.02 (-0.19 to 0.22)	366 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
by 24 months	SMD 0.07 (-0.13 to 0.27)	373 (2 studies) <sup>3,4</sup>	Low <sup>1,2</sup>
by 30 months	SMD 0.03 (-0.17 to 0.23)	379 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
by 36 months	SMD 0.08 (-0.23 to 0.39)	374 (2 studies) <sup>3,4</sup>	Moderate <sup>1</sup>
<p><i>Note.</i> A RR of &lt; 1 favours integrated ACT; Negative SMDs favour integrated ACT, positive MDs favour integrated ACT; CI = confident interval; MD = mean difference; RR = Relative Risk; SMD = Standardised mean difference.</p> <p><sup>1</sup> Optimal information size (for dichotomous outcomes, OIS = 300 events; for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> DRAKE1998.</p> <p><sup>4</sup> ESSOCK2006.</p>			

1

## 2 6.2.5 Evidence from observational studies (integrated service 3 models)

4 Mangrum and colleagues (2006) investigated hospitalisation and arrest  
5 outcomes for people with psychosis and coexisting substance misuse  
6 allocated to integrated (n=123) or parallel treatment (n=93). Of the total  
7 sample, 21% had a principal diagnosis of schizophrenia, 20% bipolar, 11%  
8 alcohol or substance use disorder. Service Users in the parallel treatment  
9 condition received substance abuse and mental health treatment by separate  
10 clinics; therefore services were not coordinated and lacked a centralised case  
11 management component. Results using weighted least squares methods  
12 revealed a significant effect favouring the integrated treatment group post-  
13 baseline on measures of any psychiatric hospitalisation,  $F(1) = 21.17$ ,  $p <$   
14  $0.0001$  and hospital days,  $F(1) = 4.28$ ,  $p = 0.04$ . Thus, a significant difference  
15 was found in number of days hospitalised favouring those in the integrated  
16 group.

17

18 Ho and colleagues (1999) prospectively looked at 6-month treatment  
19 engagement and outcome of four groups (n=179) successively enrolled in a  
20 day hospital of a 'dual-diagnosis treatment program', monitoring  
21 effectiveness changes over a 2-year period. The entire sample met criteria for  
22 psychosis (schizophrenia, schizoaffective disorder, or psychotic disorder not  
23 otherwise specified) and substance dependence (with the primary drug of use  
24 being cocaine, followed by alcohol and marijuana). Results demonstrated that  
25 all groups made sequential improvements (from group 1 to 4). Participants in  
26 group 4 had the highest engagement, attendance and retention rates, as they  
27 received the fullest spectrum of treatment (and had access to more activities

1 and therapeutic treatments) when compared with the other three groups.  
2 Furthermore, an increasing percentage of participants from group 1 to 4  
3 maintained sobriety for at least 1 to 4 months in the first six months of  
4 treatment (Cochrane-Armitage trend test statistic: 1 month, 2.16,  $p = 0.03$ ; 2  
5 months, 4.26,  $p = 0.01$ ; 3 months, 6.37,  $p = 0.001$ ; 4 months, 2.02,  $p = 0.04$ ).

6  
7 Drake and colleagues (1997) conducted a quasi-experimental study  
8 comparing integrated treatment with standard treatment on outcomes of  
9 mental health, substance abuse and housing for homeless individuals with  
10 psychosis and coexisting substance misuse. The entire sample met criteria for  
11 alcohol or drug dependence, and most had a diagnosis of schizophrenia (50%)  
12 or bipolar disorder (17%). At 18 month follow-up, service users in the  
13 integrated treatment group ( $n=158$ ) had significantly fewer days in an  
14 institution and more days in stable housing, made more progress in terms of  
15 substance abuse recovery ( $p=0.002$ ), and showed greater improvement of  
16 alcohol use disorders than those in standard treatment ( $n=59$ ) ( $p=0.05$ ). There  
17 were no significant differences between the two groups on treatment  
18 retention.

### 19 **6.2.6 Clinical evidence summary (integrated service models)**

20 There were two trials comparing an integrated service model (integrated ACT  
21 or integrated DDT [Dual Disorders Treatment]) with standard care ( $N=277$ );  
22 one of these trials also compared integrated ACT with non-integrated ACT  
23 ( $N=100$ ). However, no data from the critical outcomes could be combined  
24 using meta-analysis, so for each outcome the evidence comes from a single  
25 study. Based on these critical outcomes, the evidence (*GRADED* moderate to  
26 low quality) is inconclusive regarding the effectiveness of using an integrated  
27 approach for people with psychosis and coexisting substance misuse.

28  
29 In addition, there were two trials compared integrated ACT with integrated  
30 standard case management ( $N=421$ ), but again the evidence (*GRADED*  
31 moderate to low quality) was inconclusive.

32  
33 The three observational studies generally demonstrated support for  
34 integrated service models, but methodological issues and study setting make  
35 it difficult to generalise their results to the UK.

### 36 **6.2.7 Health economic evidence (integrated service models)**

37 The systematic search of the health economics literature identified two US-  
38 based studies (Clark *et al.*, 1998; Morse *et al.*, 2006) that considered the cost-  
39 effectiveness of integrated service models versus standard or non-integrated

1 care. Details on the methods used for the systematic search of the economics  
2 literature are described in Appendix 9.

3  
4 The study by Clark *et al.* (1998), assessing the cost-effectiveness of ACT versus  
5 standard case management (SCM), was based on the RCT described by Drake  
6 and colleagues (1998). The study sample consisted of 193 people recruited  
7 across multiple sites, diagnosed with schizophrenia, schizoaffective disorder  
8 or bipolar disorder alongside an active substance use disorder. The time  
9 horizon of the economic analysis was three years with participants  
10 interviewed at six-month intervals. A societal perspective was adopted for the  
11 cost analysis. Therefore, resource use data including mental health and  
12 general health care, legal services, community services (for example, homeless  
13 shelters) and informal care-giving, were all collected. The primary outcome  
14 measure used for the cost-effectiveness analysis was the QoL year which  
15 weighted participants' subjective quality of life (measured by the Quality of  
16 Life Interview on a 0-1 scale) over consecutive six-monthly intervals.

17  
18 Overall, mean three-year costs were similar across both groups: \$118,079 for  
19 ACT and \$124,145 for SCM. Average QoL year ratios per \$10,000 were 0.24 for  
20 integrated care participants and 0.20 for standard care participants. Overall,  
21 no significant differences in costs and effectiveness were detected between the  
22 two groups over the three-year period. There are several methodological  
23 issues with the study that limits the generalisability of the results to the UK  
24 context. First, estimates of quality of life were elicited directly from service  
25 users in the study rather than from national sample estimates. The latter  
26 approach is recommended by NICE for estimating QALYs for cost-utility  
27 analyses in the UK (NICE, 2009b). The authors did not attempt to combine  
28 total costs and outcomes by using incremental cost-effectiveness ratios,  
29 instead calculating ratios of cumulative quality of life years to total costs. No  
30 power calculations were provided in the determination of sample sizes and  
31 no formal consideration was given to study non-completers which may have  
32 biased the results.

33  
34 The study by Morse and colleagues (2006) included a cost analysis, which  
35 compared costs over 24 months between three treatment programmes:  
36 integrated ACT, non-integrated ACT, and standard care. The study was based  
37 on an RCT of 149 individuals with coexisting severe mental illness and  
38 substance use disorders who were homeless at baseline. Again a societal  
39 perspective was adopted for the cost analysis. Resource use data associated  
40 with mental health care, substance abuse treatment, physical health care and  
41 emergency shelters were collected from Medicaid claims. Over 24-months,  
42 total average costs in integrated ACT (\$48,764) and standard care (\$41,726)  
43 were significantly lower than in the non-integrated ACT programme

1 (\$71,211), while no significant cost differences were detected between the  
2 integrated ACT and standard care programmes. Most of the cost differences  
3 were explained by higher outpatient care incurred by the non-integrated ACT  
4 group, while inpatient care was similar across all three programmes. The  
5 results of the study have limited applicability to the UK setting for a number  
6 of reasons. First, the study was US based and it is unlikely that treatment  
7 patterns and associated resource use is generalisable to the UK context.  
8 Sample attrition may have biased the results of the cost analysis, although  
9 Morse and colleagues argue that attrition resulted in low statistical power, but  
10 did not affect internal validity. Finally, the study was a cost analysis and no  
11 formal attempt was made to compare the differences in total costs across the  
12 two treatment pathways with any differences in effectiveness.

### 13 *Health economics summary*

14 The literature review identified only two US-based studies that considered  
15 the cost-effectiveness of integrated care models (Clark *et al.*, 1998; Morse *et al.*,  
16 2006). Both studies suggest that integrated care models may be no more costly  
17 than non-integrated models, with no differences in health outcomes. Both  
18 studies adopted a societal perspective, including costs incurred by  
19 community services and families of service users. However, these costs  
20 accounted for a fraction of the total costs of the integrated service models  
21 considered. Both US-based studies are of limited applicability to the NHS  
22 context and limited in terms of their overall methodological quality.

23  
24 Given the uncertainty surrounding the cost-effectiveness of integrated models  
25 of care and the associated resource implications, it was anticipated that an  
26 economic model would be developed to address these issues. However, due  
27 to both the scarcity and the generally low quality of the clinical data that was  
28 identified in the guideline systematic review, the GDG agreed that it would  
29 not be possible to model the cost-effectiveness of integrated models of care.

### 30 **6.2.8 From evidence to recommendations (integrated service 31 models)**

32 Early in the development process, the GDG distinguished between outcomes  
33 that were critical to decision making and those that were important but not  
34 critical. Critical outcomes included: mortality (all causes), relapse rates  
35 (measured by exacerbation of symptoms requiring change in health care  
36 management), substance misuse (however measured), global and social  
37 functioning (for example, employment, accommodation), subjective quality of  
38 life, satisfaction with care, and physical morbidity. Only critical outcomes  
39 were included in the GRADE evidence profiles.

40

1 The review found only moderate to low quality evidence from randomised  
2 trials relating to integrated service models, and the GDG concluded that this  
3 was inconclusive. Furthermore, all of the clinical evidence and the health  
4 economic evidence included in this review were from North America, and  
5 therefore, are of questionable relevance to clinical practice in the UK.

6

7 Policy suggests that mental health services should be the lead service in  
8 working with people who are misusing substances and have a diagnosis of  
9 psychosis, and the GDG felt it was important to make a recommendation  
10 reflecting this policy.

11

12 The literature does not address the needs of people with psychosis who are  
13 severely dependent on alcohol or dependent on both alcohol and  
14 benzodiazepines or dependent on opioids and/or cocaine or crack cocaine: a  
15 small group amongst service users with psychosis. For reasons of safety in  
16 prescribing and the expertise required in monitoring the service user's  
17 requirements of substitute opiates, the GDG concluded that it would be  
18 appropriate to recommend a parallel model in which both substance misuse  
19 services and mental health services work with the service user in the overall  
20 context of the Care Programme Approach. There was no evidence that  
21 addressed the two sub-questions regarding elements of an integrated service  
22 model and subgroups of people (see section 6.2.2 for further information  
23 about these sub-questions).

1 **6.2.9 Recommendations (integrated service models)**

2 **6.2.9.1** For most adults with psychosis and coexisting substance misuse,  
3 treatment for both conditions should be provided by healthcare  
4 professionals in secondary care mental health services such as  
5 community-based mental health teams.

6 **Coordinating care**

7 **6.2.9.2** Consider seeking specialist advice and initiating joint working  
8 arrangements with specialist substance misuse services for adults and  
9 young people with psychosis being treated by community mental  
10 health teams, and known to be:

- 11 • severely dependent on alcohol **or**
- 12 • dependent on both alcohol and benzodiazepines **or**
- 13 • dependent on opioids and/or cocaine or crack cocaine.

14  
15 Adult community mental health services or CAMHS should continue  
16 to provide care coordination and treatment for the psychosis within  
17 joint working arrangements.

18 **6.2.9.3** Consider seeking specialist advice and initiate joint working  
19 arrangements with specialist substance misuse services if the person's  
20 substance misuse:

- 21 • is difficult to control **and/or**
- 22 • leads to significant impairment of functioning, family  
23 breakdown or significant social disruption such as  
24 homelessness.

25 **6.2.9.4** Delivery of care and transfer between services for adults and young  
26 people with psychosis and coexisting substance misuse should  
27 include a care coordinator and use the care programme approach.

28  
29  
30

## 1 **6.3 STAFFED ACCOMMODATION**

### 2 **6.3.1 Introduction**

3 People with severe mental health problems frequently live in staffed or  
4 supported accommodation, either as a step in a rehabilitation programme or  
5 more permanently (Macpherson *et al.*, 2004; Wolfson *et al.*, 2009). There is a  
6 wide range of accommodation providing varying degrees of support from 24-  
7 hour staffing to daytime staffing with out-of-hours telephone cover, to out-of-  
8 hours cover provided by the generic on-call service for emergencies only. The  
9 staffing can range from a full NHS multidisciplinary team to third-sector or  
10 private providers with unqualified staff. Registered care homes have to meet  
11 standards set by the Care Quality Commission in terms of the levels and  
12 experience of the care staff and will offer 24-hour staffing.

13  
14 Projects funded through Supporting People programme<sup>10</sup> will have less staff  
15 who will not be expected to provide direct care: the numbers of staff hours  
16 will depend on the nature of the project and the presumed needs of the  
17 service user group. At the lowest level people may live independently with  
18 “floating support”. Additional direct care inputs may also be provided to  
19 people in Supporting People projects.

20  
21 Other variations include housing scheme with a warden (Sheltered Housing  
22 or Special Sheltered Housing) generally for older people. In Core and Cluster  
23 housing: staff are based in the core setting that houses residents with the  
24 greatest support needs. Satellite (cluster) housing accommodates other  
25 residents grouped by needs for support.

26  
27 In Family Placements, the service user becomes part of the family. This may  
28 particularly suit people with educational under-achievement or cognitive  
29 impairment. In Adult Placement (also known as supported lodgings) a  
30 private landlord provides support to tenants renting rooms in a house. Group  
31 homes, generally for older people, provide mutual support for those who  
32 value it. Finally, dispersed intensive supported housing (Howat *et al.*, 1988)  
33 offers a specialist form of supported housing with support provided over  
34 extended hours as an alternative to residential care.

#### 35 *Current practice*

36 In the past, substance misuse was generally seen as a reason for exclusion  
37 from residential care, staffed and supported housing. Few units were

---

<sup>10</sup> Further information is available here: <http://www.communities.gov.uk>

1 prepared to tackle the challenges presented by people with coexisting mental  
2 illness and substance misuse, leading to very vulnerable individuals in  
3 housing need, being placed in extremely unsatisfactory bed and breakfast  
4 accommodation and to service users spending extended periods on acute  
5 inpatient wards in the absence of suitable alternative accommodation.

6  
7 Residential care for people with substance misuse (“rehab”) is seen as an  
8 important component in the management of people recovering from severe  
9 substance dependence. Traditionally such units were very reluctant to take in  
10 service users with a diagnosis of psychosis, even if this was effectively  
11 treated.

### 12 *Definition of intervention*

13 Any staffed accommodation or supported housing for people with a  
14 diagnosis of psychosis and coexisting substance misuse that may include an  
15 element of specific treatment for the substance misuse.

### 16 **6.3.2 Clinical review protocol (staffed accommodation)**

17 The review protocol, including the primary review question, information  
18 about the databases searched and the eligibility criteria used for this section of  
19 the guideline can be found in Table 16. During the early phase of guideline  
20 development, a recent peer-reviewed systematic review (Cleary *et al.*, 2009)  
21 was identified that addressed the review question. This systematic review  
22 was used as a source of evidence, and only a new systematic search for more  
23 recent primary-level studies was conducted for the guideline (further  
24 information about the search strategy can be found in Appendix 7).

25

**Table 16: Clinical review protocol for staffed accommodation**

Component	Description
Review question	1.2.3 In people with psychosis and coexisting substance misuse, does staffed accommodation when compared to an alternative management strategy lead to improved outcomes?
Electronic databases	CENTRAL, CINAHL, EMBASE, MEDLINE, PsycINFO
Date searched	01.01.2008 to 26.05.2010 <sup>1</sup>
Study design	RCTs and observational studies
Population	People with psychosis and coexisting substance misuse
Intervention(s)	Staffed accommodation
Comparison	Alternative management strategies
Critical outcomes	<ul style="list-style-type: none"> <li>• Reduced mortality (all causes)</li> <li>• Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</li> <li>• Reduced substance misuse (however measured)</li> <li>• Improved global and social functioning (for example, employment, accommodation)</li> <li>• Improved subjective quality of life</li> <li>• Improved satisfaction with care</li> <li>• Reduced physical morbidity.</li> </ul>
<p><i>Note.</i> RCT = Randomised controlled trial.  <sup>1</sup>The search is an update to Cleary <i>et al.</i> (2009).</p>	

1

### 2 6.3.3 Studies considered for review (staffed accommodation)

3 One RCT (N=132), BURNAM1995 (Burnam *et al.*, 1995), included in the  
4 review by Cleary and colleagues (2008), met eligibility criteria for this review.  
5 BURNAM1995 involved a comparison of a residential integrated mental  
6 health and substance use treatment programme versus standard care (see  
7 Table 17 for summary information). Full study characteristics (and any  
8 associated references), as well as a list of excluded studies can be found in  
9 Appendix 13. Forest plots and a GRADE evidence profile can be found in  
10 Appendix 14 and 15, respectively).

11

12 In addition to the RCT, five observational studies (Anderson, 1999; Blankertz  
13 & Cnaan, 1994; Brunette *et al.*, 2001; De Leon *et al.*, 2000; Nuttbrock *et al.*, 1998)  
14 met eligibility criteria for this review. Of these, all were published between  
15 1994 and 2004. Further information about each observational study and a  
16 narrative summary of results can be found in section 6.3.5.

17

**Table 17: Study information table for trials comparing staffed accommodation with standard care**

	Staffed accommodation versus standard care
Total no. of trials (N)	1 RCT (132)
Study ID	(1) BURNAM1995
Number randomised	(1) 132
Diagnosis	(1) Schizophrenia and or major affective disorder with co-occurring substance disorder <sup>1</sup>
Ethnicity	(1) 58% White
Treatment length	(1) 9 months
Country	(1) USA
Intervention (n)	(1) Residential integrated mental health and substance use treatment: educational groups, 12-step programmes including AA or NA, discussion groups, individual counselling, case-management, psychiatric consultation, ongoing medication management, general community activities (n=67)
Control (n)	(1) Routine care with no special intervention but free to access other services (shelters, mental health clinics, AA groups) (n=65)
<i>Note.</i> AA = Alcoholics Anonymous; N = Total number of participants; n = number of participants in each group; NA = Narcotics Anonymous; RCT = Randomised controlled trial. <sup>1</sup> Participants paid \$10 for each assessment interview.	

1

2 **6.3.4 Evidence from RCTs (staffed accommodation)**

3 For the comparison of staffed accommodation with standard care, a GRADE  
4 summary of findings table is shown in Table 18.

**Table 18. GRADE summary of findings table for RCTs comparing staffed accommodation with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Days used alcohol			
3 months	SMD -0.32 (-0.71 to 0.07)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD 0.00 (-0.4 to 0.4)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD -0.05 (-0.49 to 0.38)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 2. Level of alcohol use			
3 months	SMD -0.21 (-0.6 to 0.18)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD -0.06 (-0.46 to 0.33)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD -0.21 (-0.65 to 0.23)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 3. Days used drugs			
3 months	SMD -0.22 (-0.61 to 0.17)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD -0.11 (-0.51 to 0.28)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD -0.04 (-0.48 to 0.39)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 4. Severity of drug use			
3 months	SMD -0.14 (-0.52 to 0.25)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD -0.18 (-0.57 to 0.22)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD -0.16 (-0.6 to 0.28)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Functioning: 1. % time on streets			
3 months	SMD 0.04 (-0.35 to 0.42)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD -0.06 (-0.46 to 0.34)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD 0.10 (-0.34 to 0.54)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Functioning: 2. % time in independent housing			
3 months	SMD -0.16 (-0.55 to 0.23)	104 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	SMD -0.22 (-0.61 to 0.18)	97 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
9 months	SMD 0.22 (-0.22 to 0.66)	82 (1 study) <sup>3</sup>	Low <sup>1,2</sup>

Note. Negative SMDs favour staffed accommodation; CI = confident interval; SMD = Standardised mean difference.

<sup>1</sup> Optimal information size (for continuous outcomes, OIS = 400 participants) not met.

<sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.

<sup>3</sup> BURNAM1995.

### 1 **6.3.5 Evidence from observational studies (staffed** 2 **accommodation)**

3 There were five studies (Anderson, 1999; Blankertz & Cnaan, 1994; Brunette *et*  
4 *al.*, 2001; De Leon *et al.*, 2000; Nuttbrock *et al.*, 1998) which employed a non-  
5 randomised approach and examined the efficacy of residential settings for  
6 people with psychosis and coexisting substance misuse.

7  
8 Brunette and colleagues (2001) compared the effectiveness of long-term and  
9 short-term residential treatment programs. The sample consisted of  
10 participants diagnosed primarily with schizophrenia spectrum disorder (63%  
11 of the sample), in conjunction with an alcohol use disorder (32%), substance  
12 use disorder (12%) or polysubstance use (56%). Service Users in the long-term  
13 program had better engagement in treatment (Chi-square test,  $\chi^2 = 11.4$ ,  $df =$   
14  $1$ ,  $p < .001$ ) and were more likely to maintain abstinence from substance use  
15 post-discharge (Chi-square test,  $\chi^2 = 10.4$ ,  $df = 1$ ,  $p < .001$ ). There were no  
16 significant differences between short and long term residential treatment on  
17 other measures, including psychiatric hospitalisation or incarceration. It is  
18 important to note that the groups were non-equivalent however; so the data  
19 may be biased.

20  
21 Anderson (1999) explored the different impacts of an integrated approach for  
22 the treatment of psychosis and coexisting substance misuse ( $n=76$ ) and a more  
23 restrictive and traditional substance abuse model based on a therapeutic  
24 community approach ( $n=139$ ). The sample consisted of homeless participants,  
25 of whom 68.4% had a psychotic spectrum disorder (Axis 1). Fifty percent of  
26 the sample had a polysubstance abuse diagnosis (Axis 1), 22.9% had  
27 crack/cocaine problems, and 29.8% alcohol dependent. Results indicated  
28 significant differences in only five of the 33 characteristics studied. Length of  
29 stay in the program was correlated to positive treatment outcomes.  
30 Furthermore, the restrictive program was associated with twice the number of  
31 medically unadvised dropouts. It should be noted that results from this study  
32 should be interpreted with caution and cause and effect cannot be assumed,  
33 as the data analysis was based on a bivariate correlational analysis as well as a  
34 service user satisfaction survey.

35  
36 Blankertz and Cnaan (1992, 1994) compared the effectiveness of psychosocial  
37 rehabilitation versus a modified therapeutic community for homeless

1 individuals with psychosis and coexisting substance misuse. Nearly eighty  
2 percent of the overall sample had schizophrenia, and 11% had bipolar  
3 disorder. Two thirds of the sample population had a concurrent Axis III  
4 personality disorder. Substance use included alcohol (66%) cocaine, (55%),  
5 amphetamine (27%), heroin (29%), marijuana (40%), and other drugs (30%).  
6 Of the sample, 57% of the service users were polysubstance users. Results  
7 demonstrated that those receiving two years of psychosocial rehabilitation  
8 had increased abstinence (based on the ASI,  $p < 0.01$ ), improved mental state  
9 and increased treatment retention compared to the therapeutic community.

10  
11 Nuttbrock and colleagues (1998) compared a community residential treatment  
12 programme (n=87) with a therapeutic community (n=98). Of the total sample,  
13 48.8% had a primary diagnosis of a nonaffective psychotic disorder, and  
14 53.5% had a secondary diagnosis of a substance use disorder (abuse or  
15 dependence). Of those with a substance use disorder, 87.6% reported  
16 polysubstance use, 43.9% reported crack, and 21.2% reported alcohol as their  
17 primary drug of use). Service users in both programs improved on substance  
18 abuse and psychopathology outcomes, however the reductions and  
19 improvements were even greater in the therapeutic community. These results  
20 were not statistically significant after a Bonferroni correction was applied.  
21 Service users in the therapeutic community were more drug free, had more  
22 improvement in psychiatric symptoms and had improved cognitive  
23 functioning. Regression analyses indicated that improvements on  
24 psychological symptoms at 2 month follow-up and level of functioning at 12  
25 month follow-up were significantly greater among therapeutic community  
26 residents.

27  
28 More recently, De Leon and colleagues (2000) compared two types of  
29 therapeutic communities for dually diagnosed service users (medium  
30 intensity therapeutic community (n=66) and low intensity therapeutic  
31 community (n=93) versus treatment as usual (n=183). Treatment as usual  
32 consisted of the general residential programs and support services (housing,  
33 case management, day treatment) available for those with mental illness and  
34 substance use problems. In order to meet inclusion criteria, participants had  
35 to have a primary mental illness Axis 1 referral diagnosis (usually  
36 schizophrenia or major depression), a secondary Axis 1 referral diagnosis of  
37 substance abuse/dependent disorder, and a history of homelessness. Results  
38 indicated that those in the more modified, higher intensity therapeutic  
39 community (TC<sub>2</sub>) had significantly higher retention rates and did better on 12  
40 month follow-up outcomes than did those in the lower intensity (TC<sub>1</sub>) (Chi-  
41 square test,  $\chi^2 = 12.05$ ,  $p < 0.002$ ). Moreover, at two year follow-up,  
42 participants in the low intensity therapeutic community had significantly  
43 lower substance use as well as significant improved mental state (TC<sub>1</sub>). There

1 were no significant differences found on other measures, or favouring the  
2 high intensity modified therapeutic community. Those in the TC<sub>2</sub> improved  
3 statistically on 9 out of 12 outcome measures (including reduced frequency of  
4 alcohol and drug use, criminality, increased employment and improvements  
5 on the two measures of psychological functioning (SMAS and TSCS). Those in  
6 TC<sub>1</sub> and TAU improved on less outcome measures, 7 and 3 of 12, respectively.

### 7 **6.3.6 Clinical evidence summary (staffed accommodation)**

8 In one trial of residential accommodation (N=132), the evidence (*GRADED*  
9 low quality) was inconclusive to reach a decision about the effectiveness of  
10 this approach when compared to standard care for people with psychosis and  
11 coexisting substance misuse.

12  
13 Taken together, the observational studies suggest that substance use  
14 outcomes improved at follow-up, and the majority of these studies favoured  
15 longer duration integrated residential programs than shorter residential  
16 programmes. However, the substantial methodological limitations of these  
17 studies make interpretation very difficult.

### 18 **6.3.7 Health economic evidence (staffed accommodation)**

19 The systematic search of the health economics literature identified one US-  
20 based study that considered the cost-effectiveness of a staffed accommodation  
21 intervention (French *et al.*, 1999). Details on the methods used for the  
22 systematic search of the economics literature are described in Appendix 9.

23  
24 The study by French and colleagues (1999) assessed the costs and outcomes of  
25 a modified therapeutic community (TC) intervention over 12-months follow-  
26 up for homeless mentally ill chemical abusers (MICAs), compared with  
27 standard services in a treatment-as-usual (TAU) condition. This study was  
28 based on the same US service user cohort assessed by De Leon and colleagues  
29 (2000). An array of outcome measures were used in the economic analysis,  
30 including substance use, criminal activity, HIV-risk behaviour, psychological  
31 status and employment status. The perspective of the cost analysis was from  
32 the health service provider. Resource use data were collected for the modified  
33 TC intervention, hospital detoxification, A&E visits, inpatient days,  
34 residential days, non-residential day visits, outpatient visits and methadone  
35 maintenance. Over 12 months, the total mean cost per service user was  
36 \$29,255 for the modified TC group and \$29,638 for the TAU group. Overall,  
37 the higher initial cost of the modified TC intervention was offset by the higher  
38 health service utilisation in the TAU group, including residential and non-  
39 residential day visits. In terms of effectiveness, multivariate analysis showed  
40 that modified TC service users reported significantly greater reductions in

1 criminal activity and psychological dysfunction whilst no significant  
2 differences in substance use or HIV-risk behaviour were detected. No formal  
3 synthesis of costs and outcomes was carried out by the authors.

4  
5 The results of this study is of limited applicability to the UK, as it is based on  
6 a US cohort and does not attempt to synthesise costs and benefits of the two  
7 interventions being compared in the form of an incremental cost-effectiveness  
8 ratio (ICER). The authors used an array of effectiveness measures rather than  
9 a single measure such as the QALY which makes interpretation of the results  
10 difficult. Other methodological limitations relate to the cohort study design,  
11 specifically in terms of comparability between the two treatment groups in  
12 terms of subject demographic characteristics. No mention was made of how  
13 service users were allocated to both treatment groups, leading to possible  
14 selection bias, although the authors used multivariate statistical analyses to  
15 attempt to control for this. The sample sizes used for clinical outcomes and  
16 the cost analysis were different and no sensitivity analyses were performed to  
17 explore uncertainty around the base-case results.

### 18 **6.3.8 From evidence to recommendations (staffed** 19 **accommodation)**

20 Early in the development process, the GDG distinguished between outcomes  
21 that were critical to decision making and those that were important but not  
22 critical. Critical outcomes included: mortality (all causes), relapse rates  
23 (measured by exacerbation of symptoms requiring change in health care  
24 management), substance misuse (however measured), global and social  
25 functioning (for example, employment, accommodation), subjective quality of  
26 life, satisfaction with care, and physical morbidity. Only critical outcomes  
27 were included in the GRADE evidence profiles.

28  
29 Service users with coexisting substance misuse and psychosis are not ideally  
30 treated in a general ward setting, but tend to spend long periods in hospital  
31 (Menezes *et al.*, 1996). This environment is often counter-productive, where  
32 they generate great concern over the restrictions that are often imposed on  
33 them with regard to their potential to acquire illicit drugs, and in the  
34 disruption that is often created in their relationships with non-addicted  
35 service users.

36  
37 Many of the service users with combined diagnoses are too vulnerable to be  
38 discharged from hospital and yet gain little from staying in, so there have  
39 been moves to place such service users in supported staffed accommodation  
40 that may include an element of specific treatment for the substance misuse.

41

1 The evidence from randomised evidence is currently inconclusive, and  
2 positive results from observational studies could be explained by other  
3 factors, and was conducted in the United States, which makes generalisation  
4 to the UK context problematic. Nevertheless, the GDG felt that people with  
5 psychosis and coexisting substance misuse are often excluded from staffed  
6 accommodation or from treatment delivered when living in staffed  
7 accommodation, and there was no good reason for this. Therefore, in the  
8 absence of good quality evidence, the GDG decided that the main priority  
9 was to ensure people with psychosis and coexisting substance misuse were  
10 not excluded and received appropriate treatment. However, given the paucity  
11 of evidence the GDG also thought that further research was needed to decide  
12 if staffed accommodation was more cost-effective than a combination of  
13 hospital and home treatment. The GDG also thought that research was needed  
14 to decide whether there was a service delivery model that would allow  
15 people with psychosis and coexisting substance misuse to remain living  
16 outside hospital.  
17

## 18 **6.4 CLINICAL PRACTICE RECOMMENDATIONS**

### 19 **6.4.1 Recommendations (staffed accommodation)**

#### 20 *Staffed accommodation*

#### 21 **Exclusion from services**

22 **6.4.1.1** Do not exclude people with psychosis and coexisting substance misuse  
23 from staffed accommodation (such as supported or residential care)  
24 solely because of their substance misuse.

25 **6.4.1.2** Do not exclude people with psychosis and coexisting substance misuse  
26 from staffed accommodation aimed at addressing substance misuse  
27 solely because of their diagnosis of psychosis.

#### 28 **Aims of treatment**

1 **6.4.1.3** Ensure that people with psychosis and coexisting substance misuse  
2 who live in staffed accommodation receive treatment for both their  
3 psychosis and their substance misuse with the explicit aim of helping  
4 the person remain in stable accommodation.

5 **6.4.2 Research recommendations (staffed accommodation)**

6 **6.4.2.1** Is providing treatment for psychosis and substance misuse services  
7 within staffed accommodation more cost-effective than a combination  
8 of hospital and home treatment?

9 **6.4.2.2** What service delivery models allow people with psychosis and  
10 coexisting substance misuse to remain living outside hospital?

11

## 1 **6.5 INPATIENT CARE**

### 2 **6.5.1 Introduction**

3 The issues surrounding the management of inpatients with coexisting  
4 substance misuse and psychosis have been discussed in some detail in  
5 Chapter 5 (section 5.6). In brief, substance misuse is a common problem  
6 amongst people with a psychotic illness admitted to inpatient services  
7 (including secure services). Coexisting substance misuse results in longer  
8 lengths of stay in hospital and contributes substantially to incidents of  
9 violence within inpatient settings (Isaac *et al.*, 2005; Healthcare Commission,  
10 2007). Continuing substance misuse may be a reason for delay in discharge  
11 from hospital either because psychotic symptoms are exacerbated or because  
12 of concern over the future risks to themselves or others that the service user  
13 might present should they continue to abuse substances.

#### 14 *Current practice*

15 Current practice within inpatient services is not well described in the  
16 literature, although the difficulties of both staff and service users experience  
17 due to coexisting substance misuse have been very clearly documented  
18 (Healthcare Commission, 2007; Loubser *et al.*, 2009). The Department of  
19 Health has issued guidance for inpatient services about working with people  
20 with psychosis and coexisting substance misuse (Department of Health, 2006),  
21 which is focused on the need to develop policies and procedures surrounding  
22 the practicalities associated with substance misuse amongst inpatients.

#### 23 *Definition of service*

24 Any hospital-based specialist mental health service.

### 25 **6.5.2 Clinical review protocol (inpatient care)**

26 The review protocol, including the review question(s), information about the  
27 databases searched and the eligibility criteria used for this section of the  
28 guideline can be found in Table 19. During the early phase of guideline  
29 development, a recent peer-reviewed systematic review (Cleary *et al.*, 2009)  
30 was identified that addressed the review question. This systematic review  
31 was used as a source of evidence, and only a new systematic search for more  
32 recent primary-level studies was conducted for the guideline (further  
33 information about the search strategy can be found in Appendix 7). A new  
34 systematic search for systematic reviews published since 2000 was conducted  
35 in August 2009 (further information about the search strategy can be found in  
36 Appendix 7).

37

**Table 19. Clinical review protocol for inpatient care**

Component	Description
Review question	1.3.1 When a person with psychosis and coexisting substance misuse is admitted to an inpatient mental health setting (including forensic settings), should treatment follow the same principles as interventions delivered in a community setting?
Electronic databases	CENTRAL, CINAHL, EMBASE, MEDLINE, PsycINFO
Date searched	01.01.2008 to 26.05.2010
Study design	RCTs and observational studies
Population	People with psychosis and coexisting substance misuse
Intervention(s)	Inpatient care
Comparison	Community care
Critical outcomes	<ul style="list-style-type: none"> <li>• Reduced mortality (all causes)</li> <li>• Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</li> <li>• Reduced substance misuse (however measured)</li> <li>• Improved global and social functioning (for example, employment, accommodation)</li> <li>• Improved subjective quality of life</li> <li>• Improved satisfaction with care</li> <li>• Reduced physical morbidity.</li> </ul>
<p><i>Note.</i> RCT = Randomised controlled trial.  <sup>1</sup>The search is an update to Cleary <i>et al.</i> (2008) and Cleary <i>et al.</i> (2009).</p>	

1

### 2 **6.5.3 Studies considered for review (inpatient care)**

3 Two studies included in the psychological interventions chapter were  
4 conducted in inpatient settings, KAVANAGH2004 (Kavanagh *et al.*, 2004b)  
5 and LYKKE2010 (Lykke *et al.*, 2010).

6

7 Of the included studies, one was a RCT examining motivational interviewing  
8 (MI) versus standard care (KAVANAGH2004), and one was an observational  
9 study of 'cognitive milieu therapy' (LYKKE2010).

10

11 A number of other studies were also conducted in inpatient settings, but these  
12 were excluded from the review because only a small proportion of the sample  
13 were diagnosed with psychosis (for example, Moos *et al.*, 2000; Rosenheck &  
14 Fontana, 2001; Timko *et al.*, 2006).

### 15 **6.5.4 Clinical evidence summary (inpatient care)**

16 Evidence from two studies included in the psychological interventions  
17 chapter was of low quality and difficult to interpret, but suggested possible  
18 benefit of using psychological interventions to reduce substance misuse.

### 1 **6.5.5 Health economic evidence (inpatient care)**

2 No studies assessing the cost-effectiveness of inpatient care for people with  
3 psychosis and coexisting substance misuse were identified by the systematic  
4 search of the economic literature undertaken for this guideline. Details on the  
5 methods used for the systematic search of the economics literature are  
6 described in Appendix 9.

### 7 **6.5.6 From evidence to recommendations (inpatient care)**

8 The empirical literature does not at present provide good evidence to support  
9 clinical practice in this field. There are very few examples of evaluations of  
10 approaches to the management of substance misuse or specific substance  
11 misuse programmes within inpatient mental health settings. Two studies  
12 have evaluated psychological therapies delivered in the inpatient setting, but  
13 provide little evidence to reach conclusions about the effectiveness of  
14 treatment (in addition, Miles *et al.*, 2007, report the results of a non-controlled  
15 study evaluating an integrated treatment for inpatients). In the absence of  
16 good quality evidence, the GDG felt that it was appropriate to ensure that any  
17 interventions that have proven efficacy in community settings in working  
18 with this population be deployed when a person with psychosis and  
19 coexisting substance misuse is in an inpatient setting, wherever this is  
20 practicable. The GDG also felt that it was appropriate to make several  
21 recommendations for good practice concerning policies and procedures,  
22 assessment, and discharge. In particular, the GDG thought it was important  
23 that people with psychosis and coexisting substance misuse are not  
24 discharged from an inpatient mental health service solely because of their  
25 substance misuse.  
26

## 27 **6.6 CLINICAL PRACTICE RECOMMENDATIONS**

### 28 **6.6.1 Recommendations (inpatient care)**

29 *Inpatient mental health services*

30 **Substance misuse**

1 **6.6.1.1** All inpatient mental health services should ensure that they have  
2 policies and procedures for promoting a therapeutic environment free  
3 from drugs and alcohol that have been developed together with  
4 service users and their families, carers or chosen supporters. These  
5 should include: search procedures, visiting arrangements, planning  
6 and reviewing leave, drug and alcohol testing, disposal of legal and  
7 illicit substances, and other security measures. Soon after admission,  
8 provide all service users, and their families, carers or chosen  
9 supporters, with information about the policies and procedures.

10 **6.6.1.2** When carrying out a comprehensive assessment for all adults and  
11 young people admitted to inpatient mental health services, ensure  
12 that they are assessed for current substance misuse and evidence of  
13 withdrawal symptoms at the point of admission.

14 **6.6.1.3** Ensure that planned detoxification from either drugs or alcohol is  
15 undertaken only:

- 16 • with the involvement and advice of substance misuse  
17 services
- 18 • in an inpatient setting, preferably in specialist detoxification  
19 units, or designated detoxification beds within inpatient  
20 mental health services **and**
- 21 • as part of an overall treatment plan.

22  
23 For the further management of opioid detoxification see the guideline  
24 on opioid detoxification (NICE clinical guideline 52). For the further  
25 management of assisted alcohol withdrawal see the guideline on  
26 alcohol dependence and harmful alcohol use' (NICE clinical guideline,  
27 forthcoming).

## 28 **Discharge**

29 **6.6.1.4** Do not discharge adults and young people with psychosis and  
30 coexisting substance misuse from an inpatient mental health service  
31 solely because of their substance misuse.

32 **6.6.1.5** When adults and young people with psychosis and coexisting  
33 substance misuse are discharged from an inpatient mental health  
34 service, ensure that they have:

- 35 • an identified care coordinator **and**
- 36 • a care plan that includes a consideration of needs associated  
37 with both their psychosis and their substance misuse.

1  
2  
3  
4  
5  
6  
7

- been informed of the risks of overdose if they start reusing substances, especially opioids, that have been discontinued during the inpatient stay.

# 1 7 PSYCHOLOGICAL AND 2 PSYCHOSOCIAL 3 INTERVENTIONS FOR PEOPLE 4 WITH PSYCHOSIS AND 5 COEXISTING SUBSTANCE 6 MISUSE

## 7 7.1 INTRODUCTION

### 8 7.1.1 Factors related to the development of psychological 9 treatment approaches

10 There is limited understanding of just how the problems of psychosis and  
11 substance use tend to be linked together (Blanchard *et al.*, 2000). Whilst people  
12 with psychosis give many different reasons for substance use, the research  
13 consistently shows that drugs and alcohol are used by this group for many of  
14 the same reasons as those reported by the general population: to increase  
15 pleasure, to fit in with others and to alleviate negative affective states,  
16 including boredom and depression (Gregg *et al.*, 2009). However, compared  
17 with the rest of the population, these reasons may be more prominent for  
18 people with psychosis. Many people with psychosis experience negative  
19 affective symptoms (Blanchard *et al.*, 2000), and Gregg and colleagues (2009)  
20 found that reports of drug and alcohol use to cope with distressing emotions  
21 and symptoms were common, with more than half of the large sample of  
22 people with psychosis and substance use reporting they used to cope with or  
23 reduce hallucinations or feelings of suspiciousness. Some individuals with  
24 psychosis describe using substances to try and counteract the side effects of  
25 antipsychotic medication (for example, Gregg *et al.*, 2007; Spencer *et al.*, 2002);  
26 or as a preferred alternative to taking prescribed medications (Schneier &  
27 Siris, 1987). Restrictive lifestyles and limitations for obtaining pleasure in  
28 other ways may also play a part (Barrowclough *et al.*, 2006); along with a  
29 desire to fit in and be accepted by others, especially since psychosis is  
30 characterised by high levels of interpersonal difficulties (Penn *et al.*, 2004).

31  
32 Alcohol is the substance most frequently used by people with psychosis. As  
33 regards illicit drugs, cannabis is most common, although rates of poly  
34 substance use are high. This pattern of use is seen in the UK (Weaver *et al.*,

1 2003), the US (see review by Blanchard *et al.*, 2000) and Australia (Kavanagh *et*  
2 *al.*, 2004a) and is associated with the same demographic correlates as for the  
3 general population (Teeson *et al.*, 2000). It would seem that the social context  
4 and availability of substances most often influence substance choices in  
5 psychosis (Kavanagh *et al.*, 2004a; Patkar *et al.*, 1999) rather than any  
6 relationship to service users' symptomatology (Brunette *et al.*, 1997).

7  
8 Since the patterns and key motives of substance use are shared with the  
9 general population, the indications are that the psychological processes  
10 determining and maintaining use in people with psychosis may be similar to  
11 those found in non psychosis populations (Barrowclough *et al.*, 2006).

12 Therefore it would seem likely that people with psychosis may benefit from  
13 treatment approaches developed for non - psychosis service users, although  
14 treatment may need to be modified to take account of issues specific to their  
15 mental health problems and associated circumstances.

16  
17 Some of these issues present considerable challenges to treatment  
18 programmes. The functional aspects of substance use in psychosis may in part  
19 explain why motivation for reduction of substance use in service users with  
20 psychosis is usually low (Baker *et al.*, 2002; Barrowclough *et al.*, 2001; Martino  
21 *et al.*, 2002), and for many of this service user group, attempting to facilitate  
22 motivation to reduce or abstain from substances may need to be the primary  
23 focus of therapy. Importantly, people with psychosis often suffer from low  
24 self esteem (Barrowclough *et al.*, 2003); thus, self efficacy may be low, which  
25 may further decrease motivation since people may feel unable to make  
26 change. Additionally, psychosis is commonly associated with a range of  
27 complex problems, making the problematic aspects of drug and alcohol use  
28 less obvious to the individual. This may be especially so when others in the  
29 same peer group are using at the same level, so use is not seen as unusual or  
30 particularly harmful. Added to these motivational issues, the nature of the  
31 mental health problems may lead to further treatment challenges. Studies  
32 indicate that engagement in treatment is often difficult and attrition rates are  
33 high (Drake *et al.*, 2004). Reasons why this might be the case include  
34 suspiciousness or paranoid symptoms, exacerbated by substance use; chaotic  
35 lifestyles making appointment scheduling difficult; and medication issues  
36 such as poor adherence to anti-psychotics (Martino *et al.*, 2002) or the  
37 substances rendering the medications less effective.

### 38 **7.1.2 Current Practice**

39 In both the UK and the US there has been agreement by consensus that a key  
40 element of treatment approaches for coexisting substance use and psychosis is  
41 the need to take account of individuals' motivation to address or reduce their

1 substance use (Department of Health, 2002; Ziedonis *et al.*, 2005). Since  
2 motivation to change is often low, motivational techniques including  
3 motivational interviewing (MI, Miller & Rollnick, 2002) have been  
4 emphasised. Motivational interviewing is “a person-centred, directive method  
5 for enhancing intrinsic motivation to change by exploring and resolving  
6 ambivalence” (Miller & Rollnick, 2002). It aims to build intrinsic motivation  
7 for change and involves engaging the service user, offering information and  
8 feedback from assessments, where appropriate, and exploring and resolving  
9 ambivalence in an affirming and non judgemental way. It is reported that the  
10 approach can successfully be employed with people with psychosis, although  
11 the process is likely to be lengthier and some of the strategies may need  
12 adaptation to take account of issues such as thought disorder, psychotic  
13 symptoms and impaired cognitive ability (Barrowclough *et al.*, 2005;  
14 Handmaker *et al.*, 2002; Martino *et al.*, 2002).

15  
16 The additional element that has been used most commonly in recent  
17 treatment approaches for people with psychosis and coexisting substance  
18 misuse is cognitive behaviour therapy (CBT). CBT is one of the most  
19 commonly used therapeutic orientations in the field of substance use  
20 disorders (Stewart & Conrad, 2005). Moreover, CBT is recommended for all  
21 people with schizophrenia (NCCMH, 2010), and for depression in pregnant  
22 women with bipolar disorder (NCCMH, 2006). The CBT approach for  
23 individuals with psychosis and coexisting substance misuse is guided by  
24 individual formulations and by Marlatt and Gordon’s (1985) model of relapse  
25 prevention. Components may include: identifying and increasing awareness  
26 of high risk situations/warning signs; developing new coping skills for  
27 handling such situations and signs, with particular attention to psychotic  
28 symptoms and mental health related problems identified as contributing to  
29 risk of use (for example, CBT strategies for dealing with distressing voices,  
30 paranoia or depressed mood); coping with cravings and urges; making  
31 lifestyle changes so as to decrease need/urges for drugs and/or alcohol or to  
32 increase healthy activities/alternative options to substance use; normalising  
33 lapses in substance use and developing strategies and plans for acting in the  
34 event of lapse/relapse so that adverse consequences may be minimised;  
35 cognitive restructuring around alcohol and drug expectancies.

36

## 37 **7.2 EVIDENCE REVIEW**

### 38 **7.2.1 Introduction**

39 A number of existing NICE guidelines have reviewed the evidence for  
40 psychological and psychosocial interventions, and provided

1 recommendations, both for people with psychosis without substance misuse  
2 (that is, bipolar disorder; schizophrenia), and for people with substance  
3 misuse without psychosis (that is, alcohol; drug misuse: psychosocial  
4 interventions) (see Table 20).

5

6 For the purposes of the current guideline, two main issues were addressed.

7 First, in people with psychosis and coexisting substance misuse, is there  
8 evidence that any psychological/ psychosocial intervention, or combination  
9 of interventions, improve outcomes such as substance misuse, global and

10 social functioning, and quality of life? Second, should interventions  
11 recommended for a single diagnosis (either psychosis or substance misuse) be  
12 modified as a result of the presence of the coexisting diagnosis and treatment  
13 provided? For example, in people with psychosis and coexisting substance  
14 misuse, should family intervention for treatment of their psychosis be  
15 modified as a result of the substance misuse problem and the treatment  
16 provided (for example, methadone)? In addition to the main issues, the GDG  
17 were also interested in whether there was any evidence that sub-groups of  
18 people (for example, young people, people with a particular type of  
19 psychosis, people from BME groups) may benefit from alternative treatment  
20 strategies?

21

22 Where no evidence existed for a particular intervention in people with  
23 psychosis and coexisting substance misuse, the GDG used informal consensus  
24 to reach a conclusion about whether it was appropriate to use interventions  
25 recommended by existing NICE guidance.

**Table 20: Relevant interventions included in current NICE guidelines**

<b>Intervention name</b>	<b>Existing NICE guideline<sup>1</sup></b>
<b>Opportunistic brief interventions</b>	
Brief interventions for people not in contact with services	Substance misuse: DMP
Brief interventions for people in contact with services	Substance misuse: DMP
<b>Self-help based interventions</b>	
Self-help interventions (including guided self-help/bibliotherapy, 12-step based interventions)	Substance misuse: Alcohol <sup>2</sup> DMP
<b>Behavioural therapies</b>	
Cue exposure	Substance misuse: Alcohol <sup>2</sup>
Behavioural self-control training	Substance misuse: Alcohol <sup>2</sup>
Contingency management	Substance misuse: Alcohol <sup>2</sup> DMP
<b>Cognitive and behavioural based therapies</b>	
CBT	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Bipolar disorder Schizophrenia (update)
Coping and Social skills training	Substance misuse: Alcohol <sup>2</sup>
Relapse prevention	Substance misuse: Alcohol <sup>2</sup>
<b>Family-based interventions</b>	
Family intervention	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Bipolar disorder Schizophrenia (update)
<b>Motivational techniques</b>	
Motivational interviewing/ Motivational Enhancement Therapy	Substance misuse: Alcohol <sup>2</sup> DMP
<b>Social Network and Environment Based Therapies</b>	
Social Behaviour and Network Therapy	Substance misuse: Alcohol <sup>2</sup>
The Community Reinforcement Approach	Substance misuse: Alcohol <sup>2</sup>
Social-systems interventions	Substance misuse: DMD

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	DMP
<b>Other interventions</b>	
Adherence therapy	Psychosis: Schizophrenia (update)
Arts therapies	Psychosis: Schizophrenia (update)
Cognitive remediation	Psychosis: Schizophrenia (update)
Counselling and supportive psychotherapy	Substance misuse: Alcohol <sup>2</sup> Psychosis: Schizophrenia (update)
Couples-based interventions (including behavioural couples therapy)	Substance misuse: Alcohol <sup>2</sup> DMD DMP
Individual drug counselling	Substance misuse: DMD
Interpersonal and social rhythm therapy (IPSRT)	Psychosis: Bipolar disorder
Interpersonal therapy	Substance misuse: DMD DMP
Multi-modal care programmes	Substance misuse: Alcohol <sup>2</sup> DMP
Psychoeducational interventions	Substance misuse: Alcohol <sup>2</sup> Psychosis: Bipolar disorder Schizophrenia (update)
Psychodynamic psychotherapy and psychoanalysis	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Schizophrenia (update)
Social skills training	Psychosis: Schizophrenia (update)
Vocational interventions	Substance misuse: DMP
<p><i>Note.</i> DMD = Drug misuse: opioid detoxification; DMP = Drug misuse: psychosocial interventions.</p> <p><sup>1</sup> Available from <a href="http://www.nice.org.uk">www.nice.org.uk</a></p> <p><sup>2</sup> Management of alcohol dependence guideline.</p>	

1

1

## 2 **7.2.2 Definitions**

### 3 *Brief interventions*

4 In the NICE Drug Misuse Psychosocial interventions guideline (NCCMH,  
5 2008b), brief interventions were defined as interventions with a maximum  
6 duration of two sessions. The main aim of the intervention is to enhance the  
7 possibility of change in terms of abstinence or the reduction of harmful  
8 behaviours associated with drug misuse. The principles of brief interventions  
9 include expressing empathy with the service user, not opposing resistance  
10 and offering feedback, with a focus on reducing ambivalence about drug  
11 misuse and possible treatment. A number of brief interventions are based on  
12 principles drawn from motivational interviewing. Brief interventions can be  
13 conducted in a variety of settings, including non-medical settings, and can be  
14 given opportunistically to people not in formal drug treatment or as an  
15 adjunct to formal structured drug treatment (Ashton, 2005).

### 16 *Self-help based interventions*

#### 17 **Self-help intervention**

18 In the NICE alcohol guideline (NCCMH, in press), a self-help intervention  
19 was defined as an intervention where a healthcare professional (or para-  
20 professional) would facilitate the use of the self-help material by introducing,  
21 monitoring and reviewing the outcome of such treatment. The intervention is  
22 limited in nature, usually no more than three to five sessions some of which  
23 may be delivered by telephone. Self-administered intervention is designed to  
24 modify drinking behaviour and makes use of a range of books, web pages,  
25 CD-ROMs or a self-help manual that is based on an evidence-based  
26 intervention and designed specifically for the purpose. An example is Guided  
27 Self Change (GSC) (Sobell & Sobell, 1993). This treatment is manual-based  
28 and uses the principles of cognitive behavioural therapy and motivational  
29 enhancement therapy. The service user has an initial assessment followed by  
30 four treatment sessions and two follow-up telephone calls.

31

#### 32 **Self-help group**

33 In the NICE DMP guideline (NCCMH, 2008b), a self-help group was defined  
34 as a group of people who misuse drugs who meet regularly to provide help  
35 and support for one another. The group is typically community based, peer  
36 led and non-professional.

37

#### 38 **12-step self-help group**

39 In the NICE DMP guideline (NCCMH, 2008b), a 12-step self-help group was  
40 defined as a non-profit fellowship of people who meet regularly to help each

1 other remain abstinent. The core of the 12-step programme is a series of 12  
2 steps that include admitting to a drug problem, seeking help, self-appraisal,  
3 confidential self-disclosure, making amends – when possible – where harm  
4 has been done, achieving a spiritual awakening and supporting other drug-  
5 dependent people who want to recover.

6

### 7 **Twelve-Step Facilitation (TSF)**

8 In the NICE alcohol guideline (NCCMH, in press), Twelve-Step Facilitation  
9 was defined as an intervention based on the twelve-step or Alcoholics  
10 Anonymous (AA) concept that alcoholism is a spiritual and medical disease.  
11 As well as a goal of abstinence, this intervention aims to actively encourage  
12 commitment to and participation in AA meeting. Participants are asked to  
13 keep a journal of AA attendance and participation and are given AA literature  
14 relevant to the ‘step’ of the programme the service user has reached. Twelve-  
15 Step Facilitation is highly structured and manualised (Nowinski *et al.*, 1992)  
16 and involves a weekly session in which the service user is asked about their  
17 drinking, AA attendance and participation, given an explanation of the  
18 themes of the current sessions, and goals for AA attendance are set.

### 19 *Behavioural therapies*

#### 20 **Cue exposure**

21 In the NICE alcohol guideline (NCCMH, in press), cue exposure was defined  
22 as a treatment for alcohol misuse that is based on both learning theory models  
23 and social learning theory and suggests that environmental cues associated  
24 with drinking can elicit conditioned responses which can in turn lead to a  
25 relapse (Niaura *et al.* 1988). The first case study using cue exposure treatment  
26 for excessive alcohol consumption was reported by Hodgson & Rankin (1976).  
27 Treatment is designed to reduce craving for alcohol by repeatedly exposing  
28 the service user to alcohol related cues until the service user ‘habituates’ to the  
29 cues and can hence maintain self-control in a real-life situation where these  
30 cues are present.

31

#### 32 **Behavioural self-control training**

33 In the NICE alcohol guideline (NCCMH, in press), behavioural self-control  
34 training (also referred to as ‘behavioural self-management training’) was  
35 defined as approach based on the techniques described by Miller and Muñoz  
36 (1976). Service users are taught to set limits for drinking and self-monitor  
37 drinking episodes and are offered refusal skills training and training for  
38 coping behaviours in high-risk relapse situations. Behavioural self-control  
39 training is focused on a moderation goal rather than abstinence.

40

#### 41 **Contingency management**

1 In the NICE DMP guideline (NCCMH, 2008b) contingency management was  
2 defined as an approach that considers drug use as an example of operant  
3 behaviour that is maintained partly by the pharmacological effects of the drug  
4 in combination with other social and non-drug reinforcement provided by the  
5 drug using lifestyle (Petry, 2006). In the Alcohol guideline, contingency  
6 management was described as a system of reinforcement designed to make  
7 continual alcohol use less attractive and abstinence more attractive.

8  
9 Contingency management seeks to provide alternative incentives contingent  
10 on abstinence from a particular target drug. There are four primary methods  
11 of providing incentives:

- 12 • Voucher-based reinforcement: People who misuse drugs or alcohol  
13 receive vouchers with various monetary values (usually increasing  
14 in value after successive periods of abstinence) for providing  
15 biological samples (usually urine) that are negative for the tested  
16 substances. These vouchers are withheld when the biological sample  
17 indicates recent substance use. Once earned, vouchers are  
18 exchanged for goods or services that are compatible with a  
19 substance-free lifestyle.
  
- 20 • Prize-based reinforcement: This is more formally referred to as the  
21 'variable magnitude of reinforcement procedure' (Prendergast *et al.*,  
22 2006). Participants receive draws, often from a number of slips of  
23 paper kept in a fishbowl, for providing a negative biological  
24 specimen. Provision of a specimen indicating recent substance use  
25 results in the withholding of draws. Each draw has a chance of  
26 winning a 'prize', the value of which varies. Typically, about half the  
27 draws say 'Good job!'. The other half results in the earning of a  
28 prize, which may range in value from £1 to £100 (Prendergast *et al.*,  
29 2006).
  
- 30 • Clinic privileges: Participants receive clinic privileges for  
31 performing the target behaviour, for example, providing a negative  
32 biological sample. But these privileges are withheld when the target  
33 behaviour is not performed. An example of a clinic privilege is a  
34 take-home methadone dose (for example, Stitzer *et al.*, 1992).
  
- 35 • Cash incentives: People who misuse drugs receive cash (usually of a  
36 relatively low value, for example, £1.50–£10) for performing the  
37 target behaviour, such as submitting a urine sample negative for  
38 drugs or adherence with particular interventions. Cash incentives  
39 are withheld when the target behaviour is not performed.

1 *Cognitive and behavioural based therapies*

2 **Standard Cognitive Behavioural Therapy (CBT)**

3 In the NICE alcohol guideline (NCCMH, in press) and DMP guideline  
4 (NCCMH, 2008b), standard CBT was defined as a discrete, time-limited,  
5 structured psychological intervention, derived from a cognitive model of  
6 drug misuse (Beck *et al.*, 1993). There is an emphasis on identifying and  
7 modifying irrational thoughts, managing negative mood and intervening  
8 after a lapse to prevent a full-blown relapse.

9

10 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010)<sup>11</sup>,  
11 CBT was defined as a discrete psychological intervention where service users:

- 12 • establish links between their thoughts, feelings or actions with  
13 respect to the current or past symptoms, and/or functioning, and
- 14 • re-evaluate their perceptions, beliefs or reasoning in relation to the  
15 target symptoms.

16

17 In addition, a further component of the intervention should involve the  
18 following:

- 19 • service users monitoring their own thoughts, feelings or behaviours  
20 with respect to the symptom or recurrence of symptoms, and/or
- 21 • promotion of alternative ways of coping with the target symptom,  
22 and/or
- 23 • reduction of distress, and/or
- 24 • improvement of functioning.

25

26 **Coping and Social Skills Training**

27 In the NICE alcohol guideline (NCCMH, in press), coping and social skills  
28 training was defined as a variant of CBT that is based on social learning  
29 theory of addiction and the relationship between drinking behaviour and life  
30 problems (Kadden *et al.*, 1992; Marlatt & Gordon, 1985). Treatment is manual-  
31 based (Marlatt & Gordon, 1985) and involves increasing the individual's  
32 ability to cope with high-risk social situations and inter-personal difficulties.

33

34 **Relapse-prevention**

---

<sup>11</sup> A similar definition was provided in the NICE bipolar guideline.

1 In the NICE alcohol guideline (NCCMH, in press), relapse prevention was  
2 defined as a CBT adaptation based on the work of Marlatt (Marlatt & Gordon,  
3 1985), this incorporates a range of cognitive and behavioural therapeutic  
4 techniques to identify high risk situations, alter expectancies and increase self-  
5 efficacy. This differs from standard CBT in the emphasis on training people  
6 who misuse alcohol to develop skills to identify situations or states where  
7 they are most vulnerable to alcohol use, to avoid high-risk situations, and to  
8 use a range of cognitive and behavioural strategies to cope effectively with  
9 these situations (Annis, 1986; Marlatt & Gordon, 1985).

## 10 *Family-based interventions*

### 11 **Family intervention**

12 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
13 family intervention was defined as discrete psychological interventions  
14 where:

- 15 • family sessions have a specific supportive, educational or treatment  
16 function and contain at least one of the following components:
  - 17 - problem solving/crisis management work, or
  - 18 - intervention with the identified service user.

## 19 *Motivational techniques*

### 20 **Motivational interviewing**

21 For the purposes of the current guideline, MI was defined as “a client-centred,  
22 directive method for enhancing intrinsic motivation to change by exploring  
23 and resolving ambivalence” (Miller & Rollnick, 2002). It aims to build intrinsic  
24 motivation for change and involves engaging the service user, offering  
25 information and feedback from assessments, where appropriate, and  
26 exploring and resolving ambivalence in an affirming and non judgemental  
27 way. In people with psychosis, the process is likely to be lengthier and some  
28 of the strategies may need adaptation to take account of issues such as  
29 thought disorder, psychotic symptoms and impaired cognitive ability  
30 (Barrowclough *et al.*, 2005; Handmaker *et al.*, 2002, Martino *et al.*, 2002).

### 32 **Motivational Enhancement Therapy**

33 In the NICE alcohol guideline (NCCMH, in press), Motivational Enhancement  
34 Therapy (MET) was defined as an approach based on the methods and  
35 principles of MI (Miller *et al.*, 1992). It is person-centred and aims to result in  
36 rapid internally motivated changes by exploring and resolving ambivalence  
37 towards behaviour. The treatment strategy of motivational interviewing is not  
38 to guide the service user through recovery step by step, but to use  
39 motivational methods and strategies to utilise the service user’s resources. A

1 more specific manualised and structured form of motivational interviewing  
2 based on the work of Project MATCH is usually utilised (Project MATCH  
3 Research Group, 1993).

#### 4 *Social Network and Environment Based Therapies*

##### 5 **Social Behaviour and Network Therapy**

6 In the NICE alcohol guideline (NCCMH, in press), Social Behaviour and  
7 Network Therapy (SBNT) was defined as comprising of a range of cognitive  
8 and behavioural strategies to help service users build social networks  
9 supportive of change which involve the service user and members of the  
10 service user's networks (for example, friends and family) (Copello, 2002). The  
11 integration of these strategies has the aim of helping the service user to build  
12 'positive social support for a change in drinking'.  
13

##### 14 **The Community Reinforcement Approach**

15 In the NICE alcohol guideline (NCCMH, in press), the community  
16 reinforcement approach (Hunt & Azrin, 1973; Meyers & Miller, 2001; Sisson &  
17 Azrin, 1986), was defined as an approach where emphasis is placed on  
18 maintaining abstinence through the development of activities that do not  
19 promote alcohol use, for example, recreational and social activities,  
20 employment and family involvement.  
21

##### 22 **Social-systems interventions**

23 In the NICE DMP guideline (NCCMH, 2008b), it was suggested that social-  
24 systems interventions were developed primarily (but not exclusively) for  
25 young people. These interventions aim to address a range of risk and  
26 protective factors for drug misuse within the service user's wider social  
27 network. Family members, partners, close friends and other significant  
28 individuals (such as teachers or probation officers) may be involved in joint  
29 treatment sessions with the service user in a range of settings (for example,  
30 Henggeler *et al.*, 1999).

#### 31 *Other interventions*

##### 32 **Adherence therapy**

33 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
34 adherence therapy was defined as any programme involving interaction  
35 between service provider and service user, during which service users are  
36 provided with support, information and management strategies to improve  
37 their adherence to medication and/or with the specific aim of improving  
38 symptoms, quality of life and preventing relapse.  
39

##### 40 **Arts therapies**

1 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
2 arts therapies were defined as complex interventions that combine  
3 psychotherapeutic techniques with activities aimed at promoting creative  
4 expression. In all arts therapies:

- 5 • the creative process is used to facilitate self-expression within a  
6 specific therapeutic framework
- 7 • the aesthetic form is used to 'contain' and give meaning to the  
8 service user's experience
- 9 • the artistic medium is used as a bridge to verbal dialogue and  
10 insight-based
- 11 • psychological development if appropriate
- 12 • the aim is to enable the service user to experience him/herself  
13 differently and develop new ways of relating to others.

14 Arts therapies currently provided in the UK comprise: art therapy or art  
15 psychotherapy, dance movement therapy, body psychotherapy,  
16 dramatherapy and music therapy.

### 17 **Cognitive remediation**

18 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
19 cognitive remediation was defined as:

- 20 • an identified procedure that is specifically focused on basic  
21 cognitive processes, such as attention, working memory or executive  
22 functioning, and  
23
- 24 • having the specific intention of bringing about an improvement in  
25 the level of performance on that specified cognitive function or other  
26 functions, including daily living, social or vocational skills.

### 27 **Counselling and supportive psychotherapy**

28 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
29 counselling and supportive therapy were defined as discrete psychological  
30 interventions that:  
31

- 32 • are facilitative, non-directive and/or relationship focused, with the  
33 content largely determined by the service user, and
- 34 • do not fulfil the criteria for any other psychological intervention.

### 35 **Couples-based interventions**

1 In the NICE alcohol guideline (NCCMH, in press), it is suggested that the  
2 content and definition of couples therapy can vary and reflect different  
3 approaches, for example, cognitive behavioural or psychodynamic. Couples-  
4 based interventions (including behavioural couple's therapy [BCT]) involve  
5 the spouse or partner expressing active support for the person who misuses  
6 alcohol in reducing alcohol use, including via the use of behavioural  
7 contracts. Couples are helped to improve their relationship through more  
8 effective communication skills, and encouraged to increase positive  
9 behavioural exchanges through acknowledgement of pleasing behaviours and  
10 engagement in shared recreational activities (Fals-Stewart *et al.*, 2005).  
11 Standard BCT is manual based and structured (Fals-Stewart *et al.*, 2004) and  
12 combines cognitive-behaviour treatment strategies with methods that address  
13 relationship issues arising from alcohol misuse as well as more general  
14 relationship problems with the aim of reducing distress.

15

### 16 **Individual drug counselling**

17 In the NICE DMD guideline (NCCMH, 2008a), individual drug counselling  
18 was defined as the assessment of an individual's needs, provision of  
19 information and referral to services to meet these needs (including  
20 psychosocial interventions, methadone and residential rehabilitation). No  
21 attempt is made to engage in any specific formal psychological intervention.  
22 Sessions are normally weekly and last 15–20 minutes (Rawson *et al.*, 1983).  
23 This to some extent resembles keyworking as used in the UK drug treatment  
24 field.

25

### 26 **Interpersonal and social rhythm therapy (IPSRT)**

27 In the NICE guideline on bipolar disorder (NCCMH, 2006), IPSRT was  
28 defined as discrete, time limited, structured psychological intervention  
29 derived from an interpersonal model of affective disorders that focuses on:

- 30 • working collaboratively with the therapist to identify the effects of  
31 key problematic areas related to interpersonal conflicts, role  
32 transitions, grief and loss, and social skills, and their effects on  
33 current symptoms, feelings states and/or problems
  
- 34 • seeking to reduce symptoms by learning to cope with or resolve  
35 these interpersonal problem areas
  
- 36 • seeking to improve the regularity of daily life in order to minimise  
37 relapse.

38

### 39 **Interpersonal therapy**

1 In the NICE DMP guideline (NCCMH, 2008b), interpersonal therapy (IPT)  
2 was defined as a discrete, time-limited, structured psychological intervention,  
3 originally developed for the treatment of depression, which focuses on  
4 interpersonal issues and where therapist and service user:

- 5 • work collaboratively to identify the effects of key problematic areas  
6 related to interpersonal conflicts, role transitions, grief and loss, and  
7 social skills, and their effects on current drug misuse, feelings states  
8 and/or problems; and
- 9 • seek to reduce drug misuse problems by learning to cope with or  
10 resolve interpersonal problem areas (Weissman *et al.*, 2000).

### 11 **Multi-modal care programmes**

12 In the NICE DMP guideline (NCCMH, 2008b), multi-modal care programmes  
13 were defined as those including a combination of therapy activities delivered  
14 in intensive schedules of 10 hours per week or more. Content of these  
15 programmes varies but would usually include education, daily living skills  
16 and other psychologically based interventions (for example, CBT, relapse  
17 prevention and reinforcement-based approaches), mostly delivered in group  
18 format. Such programmes are not common in generic drug treatment services  
19 in the UK, although they are available in some areas. They are more  
20 commonly used within drug services linked to the criminal justice system as a  
21 way of providing more intensive programmes for those referred. The current  
22 use of these interventions in the UK is limited and their distribution is not  
23 well understood.  
24

### 25 **Psychoeducational interventions**

26 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
27 psychoeducational interventions were defined as:

- 29 • any programme involving interaction between an information  
30 provider and service users or their carers, which has the primary  
31 aim of offering information about the condition; and
- 32 • the provision of support and management strategies to service  
33 users and carers.

34 To be considered as well defined, the educational strategy should be tailored  
35 to the need of individuals or carers.

### 36 **Psychodynamic and psychoanalytic therapies**

37 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
38 psychodynamic interventions were defined as having:  
39

- 1           • regular therapy sessions based on a psychodynamic or  
2           psychoanalytic model; and
- 3           • sessions that could rely on a variety of strategies (including  
4           explorative insight-orientated, supportive or directive activity),  
5           applied flexibly.

6 To be considered as well-defined psychodynamic psychotherapy, the  
7 intervention needed to include working with transference and unconscious  
8 processes.

9  
10 Psychoanalytic interventions were defined as having:

- 11           • regular individual sessions planned to continue for at least 1 year;  
12           and
- 13           • analysts required to adhere to a strict definition of psychoanalytic  
14           technique.

15 To be considered as well-defined psychoanalysis, the intervention needed to  
16 involve working with the unconscious and early child/adult relationships.

17  
18 **Social skills training**

19 In the NICE guideline on schizophrenia (updated edition; NCCMH, 2010),  
20 social skills training was defined as a structured psychosocial intervention  
21 (group or individual) that aims to enhance social performance, and reduce  
22 distress and difficulty in social situations. The intervention must:

- 23           • include behaviourally-based assessments of a range of social and  
24           interpersonal skills, and
- 25           • place importance on both verbal and non-verbal communication,  
26           the individual's ability to perceive and process relevant social cues,  
27           and respond to and provide appropriate social reinforcement.

28  
29 **Vocational interventions**

30 In the NICE DMP guideline (NCCMH, 2008b), pre-vocational training was  
31 defined as any approach to vocational rehabilitation in which participants are  
32 expected to undergo a period of preparation before being encouraged to seek  
33 competitive employment. This preparation could involve either work in a  
34 sheltered environment (such as a workshop or work unit), or some form of  
35 pre-employment training or transitional employment (Crowther *et al.*, 2001).  
36 Supported employment was defined as any approach to vocational  
37 rehabilitation that attempts to place service users immediately in competitive

1 employment. It is acceptable for supported employment to begin with a short  
2 period of preparation, but this has to be of less than one month's duration and  
3 not involve work placement in a sheltered setting, or training, or transitional  
4 employment (Crowther *et al.*, 2001).

### 5 **7.2.3 Clinical review protocol (psychological/ psychosocial** 6 **interventions)**

7 The review protocol, including the review questions, information about the  
8 databases searched, and the eligibility criteria used for this section of the  
9 guideline, can be found in Table 21. During the early stages of guideline  
10 development, a recent Cochrane review (Cleary *et al.*, 2008) and related peer-  
11 reviewed publication (Cleary *et al.*, 2009) were identified that addressed the  
12 review question. These systematic reviews were used as a source of evidence,  
13 and only a new systematic search for more recent primary-level studies was  
14 conducted for the guideline (further information about the search strategy can  
15 be found in Appendix 7).

16  
17 If the evidence allowed, the following sub-question was asked for review  
18 question 2.2.1 and 2.4.1: Are there sub-groups of people (for example, young  
19 people, people with a particular type of psychosis, BME groups) that may  
20 benefit from alternative strategies? In addition, the following sub-question  
21 was asked for review question 2.4.1: Should interventions be matched to  
22 stages of the treatment process (i.e. engagement, persuasion, active treatment,  
23 relapse prevention)?  
24

**Table 21: Clinical review protocol for the review of psychological/ psychosocial interventions**

Component	Description
Review question	<p>1.2.2 In people with psychosis and coexisting substance misuse, do psychological/ psychosocial interventions when compared to an alternative management strategy lead to improved outcomes?</p> <p>2.2.1 For people with psychosis and coexisting substance misuse, should the psychological/ psychosocial (family interventions, CBT, arts therapies) treatment of their psychosis be modified as a result of the substance misuse problem and the treatment provided (for example, methadone, buprenorphine, psychological treatment etc)?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>2.4.1 For people with psychosis and coexisting substance misuse, should psychological/ psychosocial treatment for substance misuse be modified as a result of the presence of psychosis and the treatment provided?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p>
Electronic databases	CENTRAL, CINAHL, EMBASE, MEDLINE, PsycINFO
Date searched	01.01.2008 to 26.05.2010 <sup>1</sup>
Study design	RCTs and observational studies
Population	People with psychosis and coexisting substance misuse
Intervention(s)	Individual psychological/ psychosocial interventions for people with psychosis and coexisting substance misuse
Comparison	An alternative management strategy
Critical outcomes	<p>Reduced mortality (all causes)</p> <p>Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</p> <p>Reduced substance misuse (however measured)</p> <p>Improved global and social functioning (for example, employment, accommodation)</p> <p>Improved subjective quality of life</p> <p>Improved satisfaction with care</p> <p>Reduced physical morbidity.</p>
<sup>1</sup> The search is an update to Cleary <i>et al.</i> (2008) and Cleary <i>et al.</i> (2009).	

1

## 1 7.2.4 Studies considered for review (psychological/psychosocial 2 interventions)<sup>12</sup>

3 12 RCTs, BAKER2006 (Baker *et al.*, 2006), BARROWCLOUGH2001  
4 (Barrowclough *et al.*, 2001), BARROWCLOUGH2010 (Barrowclough *et al.*, in  
5 press), EDWARDS2006 (Edwards *et al.*, 2006), GRAEBER2003 (Graeber *et al.*,  
6 2003), HELLERSTEIN1995 (Hellerstein *et al.*, 1995), JERRELL1995 (Jerrell &  
7 Ridgely, 1995), KAVANAGH2004 (Kavanagh *et al.*, 2004b), RIES2004 (Ries *et*  
8 *al.*, 2004), SCHMITZ2002 (Schmitz *et al.*, 2002), TRACY2007 (Tracy *et al.*, 2007),  
9 WEISS2007 (Weiss *et al.*, 2007), that were included in the review by Cleary and  
10 colleagues (2008), met the eligibility criteria for this review. In addition, one  
11 further trial was identified during the search for evidence, WEISS2009 (Weiss  
12 *et al.*, 2009). Full study characteristics (and any associated references), as well  
13 as a list of excluded studies can be found in Appendix 13.

14  
15 Of the 13 included RCTs, there were four involving a comparison of CBT  
16 versus standard care (EDWARDS2006, SCHMITZ2002, WEISS2007,  
17 WEISS2009), two of MI versus standard care (GRAEBER2003,  
18 KAVANAGH2004), two of a group therapy (social skills training/  
19 psychoeducation) versus standard care (HELLERSTEIN1995, JERRELL1995),  
20 two of contingency management versus standard care (RIES2004,  
21 TRACY2007), and three of CBT combined with MI versus standard care  
22 (BAKER2006, BARROWCLOUGH2001, BARROWCLOUGH2010) (see Table  
23 22 and Table 23 for summary information about each trial).

24  
25 In addition to the RCTs, three observational studies (James *et al.*, 2004; Santa  
26 Ana *et al.*, 2007; Weiss *et al.*, 2000), that were included in the review by Cleary  
27 and colleagues (2008), met the eligibility criteria for review. A further three  
28 studies (Helmus *et al.*, 2003; Lykke *et al.*, 2010; Tyrer *et al.*, in press) were  
29 found during the search for evidence.

30  
31 Of the six observational studies, one involved a comparison of CBT versus  
32 standard care (Weiss *et al.*, 2000), one of motivational interviewing versus  
33 therapist attention activity control (Santa Ana *et al.*, 2007), one of group  
34 psychotherapy versus standard care (single educational session) (James *et al.*,  
35 2004), one of a contingency management program (Helmus *et al.*, 2003), one of  
36 cognitive milieu therapy (Lykke *et al.*, 2010), and one of nidotherapy (Tyrer *et*  
37 *al.*, in press) (see section 7.2.6 for further information about each study and a  
38 narrative summary of results).

39

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<sup>12</sup> Here and elsewhere in the guideline, each RCT considered for review is referred to by a study ID (primary author and date of study publication).

1 After the consultation period was complete, the GDG received the  
2 BARROWCLOUGH2010 trial report pre-publication. Having been accepted  
3 for publication in the BMJ, and the quality of the study having been judged as  
4 acceptable, a fresh meta-analysis of now three trials of CBT combined with  
5 MI, compared to standard care, was undertaken. This analysis is presented in  
6 the results, but readers should be aware that this small part of the guideline  
7 has not been consulted upon. As the fresh meta-analysis did not lead to any  
8 changes in the recommendations, the GDG, following consultation with NICE  
9 deemed the lack of consultation to be acceptable.  
10

**Table 22: Study information table for trials comparing CBT, MI, or CBT plus MI with standard care**

	CBT versus standard care	MI versus standard care	CBT + MI versus standard care
Total no. of trials (N)	4 RCTs (216)	2 RCTs (56)	3 RCTs (493)
Study ID	(1) EDWARDS2006 (2) SCHMITZ2002 (3) WEISS2007 (4) WEISS2009	(1) GRAEBER2003 (2) KAVANAGH2004	(1) BAKER2006 (2) BARROWCLOUGH2001 (3) BARROWCLOUGH2010
Number randomised	(1) 47 (2) 46 (3) 62 (4) 61	(1) 30 (2) 25	(1) 130 (2) 36 (3) 327
Diagnosis	(1) 72% DSM-IV schizophrenia/schizophreniform, 11% affective psychosis, 17% NOS/delusional /other and all actively using cannabis. (2) 100% DSM-IV bipolar disorder and substance use disorder (72% alcohol, 61% cocaine, 26% marijuana, 59% were dependent on more than 1 drug). (3) 100% DSM-IV bipolar disorder and substance dependence (most common; 27% alcohol, 26% marijuana). (4) 100% DSM-IV bipolar disorder with dependence (26.2% had alcohol dependence only, 8.2% had drug dependence only, and 65.6% had both).	(1) 100% DSM-IV schizophrenia and met criteria for an alcohol use disorder within the 3- month period prior to study enrolment; service users with additional non-alcohol substance use (except active intravenous drug abuse) were eligible for protocol enrolment. (2) 100% DSM-IV psychotic disorder with a current DSM-IV substance use disorder (88% alcohol, 76% cannabis, 12% inhalants, 8% cocaine or heroin).	(1) 75% ICD-10 schizophrenia or schizoaffective disorder with SCID-1 diagnosis of abuse or dependence past 12 months (alcohol 69%, cannabis 74%, amphetamine 42%) <sup>1</sup> (2) ICD-10 & DSM-IV schizophrenia or schizoaffective disorder with DSM-IV substance abuse or dependence. (3) ICD-10 & DSM-IV schizophrenia, schizophreniform or schizoaffective disorder with DSM-IV substance abuse or dependence.
Ethnicity	(1) NR (2) 80% White (3) 94% White (4) 92% White	(1) 40% White, 40% Hispanic, 20% African American (2) 84% White	(1) NR (2) White European (3) 81% White, 11% Black
Treatment length	(1) 6 months (2) 3 months (3) 8 months	(1) 6 months (2) 12 months	(1) 15 weeks (FU at 6 and 12 months) (2) 9 months (FU at 12

	(4) 6 months		and 18 months) (3) 12 months (FU at 24 months)
Country	(1) Australia (2) USA (3) USA (4) USA	(1) USA (2) Australia	(1) Australia (2) UK (3) UK
Intervention (n)	(1) Cannabis-focused CBT (weekly over 3 months) (n=23) (2) Medication monitoring and CBT (16 sessions) (n=25) (3) Integrated group CBT (20 weekly 1 hour sessions) (n=31) (4) Integrated group CBT (12 weekly 1 hour sessions) (n=31)	(1) Motivational interviewing (3 sessions) (n=15) (2) Brief motivational intervention (6-9 sessions) (n=13)	(1) Motivational interviewing and CBT (10 weekly one hour sessions) + routine care (n=65) (2) Family support worker plus motivational interviewing, manualised individual CBT for the participant and CBT for family / caregiver (a total of 29 individual sessions) + routine care (n=18) (3) Motivational interviewing and CBT (26 individual sessions delivered over 12 months) + routine care (n=164)
Control (n)	(1) Psychoeducation + standard EPPIC care (n=24) (2) Standard care (includes medication monitoring) (n=21) (3) Group drug counselling (n=31) (4) Group drug counselling (n=30)	(1) Three-session educational intervention (n=15) (2) Standard care (n=12)	(1) Routine care plus self-help books (n=65) (2) Routine care plus family support worker (n=18) (3) Routine care (n=163)
<p><i>Note.</i> CBT = cognitive behavioural therapy; FU = follow up; MI = motivational interviewing; N = total number of participants; n = number of participants in each group.</p>			

1

**Table 23: Study information table for trials comparing group approaches or contingency management with standard care**

	<b>Group psychotherapy/ behavioural skills programme versus standard care</b>	<b>Contingency management versus standard care</b>
Total no. of trials (N)	2 RCTs (94)	2 RCTs (71)
Study ID	(1) HELLERSTEIN1995 (2) JERRELL1995	(1) RIES2004 (2) TRACY2007
Number randomised	(1) 47 (2) 47	(1) 41 (2) 30
Diagnosis	(1) RDC schizophrenia with 74% DSM-III-R psychoactive substance abuse/ dependence. (2) 62% DSM-III-R schizophrenia with coexisting substance disorder.	(1) 73% schizophrenia or schizoaffective disorder, 24% major recurrent depression or bipolar disorder, 2% other, and DSM-IV substance misuse disorder with active substance use in the previous 6 months. (2) 100% current or lifetime DSM-IV diagnosis of an Axis I psychiatric disorder and current diagnosis of cocaine or alcohol abuse or dependence.
Ethnicity	(1) 43% African American, 32% Hispanic (2) 64% White	(1) NR (2) NR
Treatment length	(1) 8 months (2) 18 months	(1) 6.5 months (2) 1 month
Country	(1) USA (2) USA	(1) USA (2) USA
Intervention (n)	(1) Group outpatient psychotherapy & psychoeducation plus drug treatment all at same site (twice weekly) (n=23) (2) Behavioural skills programme: psychoeducational approach with self-management skills, repeated practice & reinforcement (weekly group sessions with two licensed clinicians) (n=22)	(1) Contingency management of supplementary social security income/food vouchers and motivational message (n=22) (2) Petry's low-cost contingency management with variable ratio reinforcement (n=15)
Control (n)	(1) Comparable levels of psychiatric care and substance abuse treatment from separate sites without formal case-coordination (n=24) (2) Twelve step recovery programme: clinical staff (some 'recoverers') offered mock AA	(1) Non-contingency management of benefits (n=19) (2) Assessment-only treatment (n=15)

	meetings within the Mental Health Centre, took or referred service users to community AA meetings, facilitated a sponsor relationship & provided counselling (n=25)	
<p><i>Note.</i> N = total number of participants; n = number of participants in each group; NR = not reported; RCT = randomised controlled trial.  <sup>1</sup>Some participants were dependent on more than one of these.</p>		

1

2 **7.2.5 Evidence from RCTs (psychological/psychosocial**  
 3 **interventions)**

4 Meta-analysis was used to synthesise the evidence for each comparison  
 5 (GRADE summary of findings tables are shown in Table 24, Table 25, Table  
 6 26, Table 27, and Table 28).

7

8 The forest plots and full GRADE evidence profiles can be found in Appendix  
 9 14 and 15, respectively.

10

11

**Table 24. GRADE summary of findings table for RCTs comparing CBT with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Using substances			
by 1 month - alcohol or drugs	RR 0.48 (0.26 to 0.9)	61 (1 study) <sup>4</sup>	Moderate <sup>1</sup>
Substance use: 2. Using substances			
by 3 months - alcohol	RR 5.88 (0.79 to 44.03)	46 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
by 3 months - drugs	RR 2.02 (0.85 to 4.8)	46 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
by 3 months - alcohol or drugs	RR 0.74 (0.55 to 1)	61 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
Substance use: 3. Any substance (skewed data) - average score (ASI)			
by 3 months	MD -0.07 (-0.16 to 0.02)	62 (1 study) <sup>6</sup>	Low <sup>1,3</sup>
by 6-9 months	MD -0.06 (-0.16 to 0.04)	62 (1 study) <sup>6</sup>	Low <sup>1,3</sup>
Substance use: 4. Any substance (skewed data) - days reporting any substance use (ASI)			
by 3 months	MD -2.1 (-5.9 to 1.7)	61 (1 study) <sup>4</sup>	Low <sup>1,2,3</sup>
by 6 months	MD -2.7 (7.25 to 1.85)	61 (1 study) <sup>4</sup>	Low <sup>1,2,3</sup>
Substance use: 5. Drugs use (skewed data)			
by 3 months	MD 0.05 (-1.55 to 1.66)	103 (2 studies) <sup>4,5</sup>	Low <sup>1,3</sup>
by 6 months	MD -3.7 (-7.99 to 0.59)	57 (1 study) <sup>4</sup>	Low <sup>1,2,3</sup>
Substance use: 6. Alcohol use (skewed data)			
by 3 months	MD -1.95 (-4.48 to 0.58)	103 (2 studies) <sup>4,5</sup>	Low <sup>1,2,3</sup>
by 6 months	MD 0.00 (-3.66 to 3.66)	57 (1 study) <sup>4</sup>	Low <sup>1,2,3</sup>
<p><i>Note.</i> A RR of &lt; 1 favours the intervention, negative MDs favour the intervention; CI = confidence interval; MD = mean difference; RR = Relative Risk.</p> <p><sup>1</sup> Optimal information size (for dichotomous outcomes, OIS = 300 events; for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> Skewed data.</p> <p><sup>4</sup> WEISS2009.</p> <p><sup>5</sup> SCHMITZ2002.</p> <p><sup>6</sup> WEISS2007.</p>			

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**Table 25. GRADE summary of findings table for RCTs comparing MI with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Not abstinent or not improved on all substances			
by 12 months	RR 0.51 (0.24 to 1.10)	25 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
Substance use: 2. Not abstaining from alcohol			
by 3 months	RR 0.52 (0.26 to 1.03)	28 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
by 6 months	RR 0.36 (0.17 to 0.75)	28 (1 study) <sup>5</sup>	Moderate <sup>1</sup>
Substance use: 3. Other measures of alcohol use (skewed data) - drinking days			
by 6 months	SMD -1.29 (-2.12 to -0.46)	28 (1 study) <sup>5</sup>	Low <sup>1,3</sup>
<p><i>Note.</i> A RR of &lt; 1 favours the intervention, negative SMDs favour the intervention; CI = confidence interval; MI = motivational interviewing; RCT = randomised controlled trial; RR = Relative Risk; SMD = Standardised mean difference.</p> <p><sup>1</sup> Optimal information size (for dichotomous outcomes, OIS = 300 events; for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> Skewed data.</p> <p><sup>4</sup> KAVANAGH2004.</p> <p><sup>5</sup> GRAEBER2003.</p>			

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**Table 26. GRADE summary of findings table for RCTs comparing CBT plus MI with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Death - by about 1 year	RR 0.73 (0.22 to 2.41)	492 (3 studies) <sup>3,4,5</sup>	Low <sup>1,2</sup>
Substance use: 1. Average number of different drugs used during the past month (OTI)			
by 3 months	MD 0.37 (-0.01, 0.75)	119 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 6 months	MD 0.19 (-0.22, 0.60)	119 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Substance use: 2. Average score - alcohol (skewed data) - alcohol - estimated daily consumption - past month			
3 months	MD 1.57 (-0.90, 4.04)	52 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
6 months	MD 1.21 (-1.07, 3.49)	52 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
12 months	MD 1.39 (-1.10, 3.88)	46 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Substance use: 3. Average score - amphetamine (skewed data) - amphetamine- estimated daily consumption - past month			
3 months	MD 0.09 (-0.40, 0.58)	20 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
6 months	MD -1.28 (-2.79, 0.23)	20 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
12 months	MD 0.13 (-0.11, 0.37)	17 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Substance use: 4. Average score - cannabis (skewed data) - cannabis- estimated daily consumption - past month			
3 months	MD -0.57 (-4.27, 3.13)	73 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	MD 0.70 (-4.00, 5.40)	73 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
12 months	MD 4.41 (-1.40, 10.22)	58 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Substance use: 7. TLFB: % days abstinent main substance (skewed data)			
12 months	MD 6.81 (-2.07 to 15.69)	275 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
18 months	MD -1.21 (-10.74 to 8.32)	258 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
24 months	MD 2.52 (-7.42 to 12.46)	246 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
Substance use: 8. TLFB: % days abstinent all substance (skewed data)			
12 months	MD 5.73 (-2.62 to 14.08)	273 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
18 months	MD -0.30 (-9.14 to 8.54)	256 (1 study) <sup>5</sup>	Low <sup>1,2</sup>
24 months	MD 7.07 (-2.32 to	247	Low <sup>1,2</sup>

	16.46)	(1 study) <sup>5</sup>	
Functioning: 1. Average global functioning score (GAF)			
3 months	MD -2.70* (-7.05, 1.65)	119 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
6 months	MD -0.09* (-3.70, 3.52)	119 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
9 months	MD 8.44* (0.48, 16.40)	32 (1 study) <sup>4</sup>	Moderate <sup>1</sup>
12 months	MD 1.87* (-2.36, 6.11)	398 (3 studies) <sup>3,4,5</sup>	Low <sup>1,2</sup>
18-24 months	MD 0.69* (-3.86, 5.25)	262 (2 study) <sup>4,5</sup>	Low <sup>1,2</sup>
Functioning: 2. Average social functioning score (SFS)			
by end of 9 month treatment	MD 5.01* (-0.55, 10.57)	32 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
by 12 months (3 months following treatment end)	MD 7.27* (0.86, 13.68)	32 (1 study) <sup>4</sup>	Moderate <sup>1</sup>
<p><i>Note.</i> A RR of &lt; 1 favours the intervention, negative MDs favour the intervention (except if marked with *, then positive MDs favour the intervention); CI = confidence interval; MD = mean difference; MI = motivational interviewing; RR = Relative Risk.</p> <p><sup>1</sup> Optimal information size (for dichotomous outcomes, OIS = 300 events; for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> BAKER2006.</p> <p><sup>4</sup> BARROWCLOUGH2001.</p> <p><sup>5</sup> BARROWCLOUGH2010.</p>			

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**Table 27. GRADE summary of findings table for RCTs comparing group psychotherapy with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. Average score - C-DIS-R Drugs (skewed data) - C-DIS-R DRUGS			
by 6 months	MD -2.99 (-5.51 to -0.47)	46 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 12 months	MD -2.47 (-5.76 to 0.82)	46 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
By 18 months	MD -0.79 (-3.35 to 1.77)	25 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Substance use: 2. Average score - C-DIS-R Alcohol (skewed data) - C-DIS-R Alcohol			
by 6 months	MD -1.81 (-3.41 to -0.21)	46 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 12 months	MD -0.71 (-2.54 to 1.12)	46 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 18 months	MD 0.04 (-2.27 to 2.35)	25 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Functioning: 1. Average role functioning score (RFS)			
by 6 months	MD 0.61* (-1.63 to 2.85)	47 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 12 months	MD 1.07* (-1.15 to 3.29)	47 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
by 18 months	MD -2.55* (-6.24 to 1.14)	25 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Functioning: 2. Average social adjustment score (SAS)			
by 6 months	MD -0.92* (-6.58 to 4.74)	47 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 12 months	MD 2.58* (-3.39 to 8.55)	47 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
by 18 months	MD -4.66* (-15.29 to 5.97)	25 (1 study) <sup>3</sup>	Low <sup>1,2</sup>
Service use: Days in hospital (skewed data)	MD 1.80 (-4.46 to 8.06)	29 (1 study) <sup>4</sup>	Low <sup>1,2</sup>
<p><i>Note.</i> Negative MDs favour the intervention (except if marked with *, then positive MDs favour the intervention); CI = confidence interval; MD = mean difference.</p> <p><sup>1</sup> Optimal information size (for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> CI includes both 1) no effect and 2) appreciable benefit or appreciable harm.</p> <p><sup>3</sup> JERRELL1995.</p> <p><sup>4</sup> HELLERSTEIN1995.</p>			

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**Table 28. GRADE summary of findings table for RCTs comparing contingency management with standard care**

Outcomes	Effect size (95% CI)	No of participants (studies)	Quality of the evidence (GRADE)
Substance use: 1. No. of days/weeks of drug use (confirmation by urine drug screen) - Days of cocaine use	SMD -1.04 (-1.8 to -0.28)	30 (1 study) <sup>2</sup>	Moderate <sup>1</sup>
Substance use: 2. No. of days/weeks of alcohol use (confirmation by breathalyzer)	SMD -1.16 (-1.83 to -0.49)	71 (2 studies) <sup>2,3</sup>	Moderate <sup>1</sup>
Substance use: 3. No. of days/weeks using both drugs and alcohol (confirmation by urine or breathalyzer) - weeks	SMD -0.82 (-1.47 to -0.17)	41 (1 study) <sup>3</sup>	Moderate <sup>1</sup>
Substance use: 4. Alcohol positive breathalyzer samples	SMD -0.82 (-1.47 to -0.17)	30 (1 study) <sup>2</sup>	Moderate <sup>1</sup>
<p><i>Note.</i> Negative SMDs favour the intervention; CI = confidence interval; SMD = Standardised mean difference.</p> <p><sup>1</sup> Optimal information size (for continuous outcomes, OIS = 400 participants) not met.</p> <p><sup>2</sup> TRACY2007.</p> <p><sup>3</sup> RIES2004.</p>			

2

### 3 7.2.6 Observational studies (psychological/ psychosocial 4 interventions)

5 Cleary and colleagues (2009) included three observational studies that met the  
6 guideline eligibility criteria. Of the three, one US study (Weiss *et al.*, 2000) of  
7 people with coexisting bipolar disorder and substance dependence was  
8 classified as examining integrated group sessions (12–20 weekly 1 hour) using  
9 a CBT relapse prevention model (n=21) versus standard care (n=24). After 6  
10 months follow up, there were statistically significant treatment group  
11 differences favouring CBT on a number of substance misuse outcomes and a  
12 measure of mania. However, assessment was not blind, although the  
13 substance misuse outcomes were verified by urine toxicology screens and  
14 breath alcohol assessments.

15

16 One US study (Santa Ana *et al.*, 2007), was described by Cleary and colleagues  
17 (2009) as a comparison of group motivational interviewing (two, 2-hour  
18 sessions; n=50) versus a control group (group discussion, two, 2-hour  
19 sessions; n=51). Participants were psychiatric inpatients with coexisting  
20 substance dependence. At 1- and 3-months follow-up there was a statistically

1 significant difference between groups favouring the motivational  
2 interviewing group on rates of alcohol use and binge drinking, and drug use  
3 days. There were no significant differences between groups on measures of  
4 abstinence or on aftercare treatment attendance.

5  
6 Cleary and colleagues (2009) included one Australian study (James *et al.*,  
7 2004), that compared the effectiveness of a 6 week manualised group-based  
8 intervention (incorporating both substance use and mental health  
9 interventions; n=32) versus standard care (consisting of a single educational  
10 session; n=31). Participants were diagnosed with schizophrenia or bipolar  
11 disorder and coexisting substance dependence or harmful use. At 3-months  
12 follow-up, there were statistically significant differences between the two  
13 groups, favouring group therapy in terms of reduced drug use and symptoms  
14 of psychosis, but not severity of dependence or alcohol use.

15  
16 One non-randomised study (Helmus *et al.*, 2003), not included by Cleary and  
17 colleagues (2009), examined the effectiveness of a community based  
18 contingency management program. The sample consisted of 20 participants  
19 diagnosed with schizophrenia (15%), schizoaffective disorder (20%), bipolar  
20 disorder (30%), or MDD (35%) and a coexisting substance use disorder  
21 (alcohol dependence, 70%; cocaine abuse, 5%; polysubstance dependence,  
22 5%). Using an A-B-A within-subjects reversal design, participants had a 4-  
23 week baseline phase, followed by 12 weeks of contingency management  
24 reinforcing their psychosis and coexisting substance misuse group  
25 counselling attendance and alcohol abstinence (based on breath alcohol  
26 levels), and then a 4 week return to baseline phase. Group counselling was  
27 provided twice weekly with alcohol breath tests given before each session.  
28 The results demonstrated that contingency management attendance was  
29 significant higher than at baseline, and remained elevated in the return to  
30 baseline phase. There were no significant effects found on alcohol use,  
31 however, as the breath tests remained negative throughout the entire study.

32  
33 Lykke and colleagues (2010) conducted a pragmatic clinical trial evaluating  
34 cognitive milieu therapy in a convenient sample of 136 inpatients in Denmark,  
35 using a pre-post intervention design. Of the 136 participants, 53 to 65% had an  
36 ICD-10 diagnosis of schizophrenia, with a coexisting diagnosis of substance  
37 abuse (29-41% alcohol only, 5-6% cannabis only, 50-59% polysubstance  
38 abuse). Cognitive milieu therapy is carried out within a structured inpatient  
39 environment, and incorporates both motivational and cognitive behavioural  
40 strategies in an effort to address both mental health and substance misuse  
41 problems simultaneously. Results revealed that the most significant changes  
42 post-treatment were in functioning (Global Assessment of Functioning scale,  
43  $p=.0001$ ), global symptomatology as assessed by the Global Assessment Scale

1 ( $p=.0001$ ), and levels of anxiety/ depression on the Brief Psychiatric Rating  
2 Scale (BPRS) ( $p=.0001$ ). In addition, participants displayed significant  
3 improvement on anxiety levels (Beck Anxiety Inventory,  $p=.0001$ ), depressive  
4 symptoms (Beck Depression Inventory,  $p=.0001$ ), and self-esteem (Robson  
5 Self-Concept Questionnaire,  $p=.0022$ ) at post-treatment follow-up. A  
6 regression analysis did not identify any predictors associated with treatment  
7 completion, although reduced chance of completion of treatment was  
8 associated with a higher BPRS score. Regression analysis for achieving  
9 sustained abstinence was associated with the absence of a polysubstance  
10 abuse diagnosis (OR = 0.19;  $p=.018$ ) and lower BPRS score (OR= 0.80, 1 per  
11 point,  $p < .01$ ).

12  
13 One further study (Tyrer *et al.*, in press), was a secondary sub-group analysis  
14 of an RCT conducted in the UK, which looked at the impact of nidotherapy  
15 for people with psychosis, a significant proportion of whom had coexisting  
16 substance misuse problems (Ranger *et al.*, 2009). Nidotherapy is a  
17 “collaborative treatment involving the systematic assessment and  
18 modification of the environment to minimise the impact of any form of  
19 mental disorder on the individual or on society” (Tyrer *et al.*, 2003). The sub-  
20 group analysis of the people with psychosis and coexisting substance misuse  
21 suggested that participants referred to nidotherapy had a 63% reduction in  
22 hospital bed use after one year compared to those referred to a standard  
23 assertive outreach team ( $p = .03$ ). There was also some evidence that  
24 nidotherapy improved social functioning (MD -2.0, 95% CI -4.0 to -0.1),  
25 without any detrimental effect on psychiatric symptoms (MD -2.6, 95% CI -8.0  
26 to 2.8) or engagement with services (MD .23, 95% CI -1.6 to 2.1).

### 27 **7.2.7 Clinical evidence summary (psychological/ psychosocial** 28 **interventions)**

29 For the majority of interventions included in related NICE guidance, the  
30 current systematic review found no direct evidence for people with psychosis  
31 and coexisting substance misuse (Table 29). With regard to the evidence that  
32 was available, it should be interpreted with some caution because the  
33 research was not conducted in the UK and methodological issues limit the  
34 quality of the evidence.

35  
36 There were two small RCTs (N=56) of MI compared to standard care.  
37 However, data could not be combined using meta-analysis, so for each  
38 outcome, the evidence comes from a single study. Nevertheless, the evidence  
39 (GRADED moderate to low quality) suggests that for people with psychosis  
40 and coexisting substance misuse this approach may reduce substance misuse

1 at up to 12 months follow-up. These results were supported by one  
2 observational study.

3

4 In two small RCTs (N=71) of contingency management compared to standard  
5 care, there was evidence (*GRADED* low quality) suggesting benefit in terms  
6 of reduced substance misuse at up to 6 months follow-up. One small  
7 observational study demonstrated improved attendance after contingency  
8 management, but no effect on alcohol use.

9

10 In four small RCTs of CBT (N=216), three trials of CBT plus MI (N=493), and  
11 two small trials of group psychotherapy (social skills training/  
12 psychoeducation) (N=94), the evidence (*GRADED* moderate to low quality) is  
13 inconclusive with regard to the effectiveness of these approaches when  
14 compared to standard care for people with psychosis and coexisting  
15 substance misuse. Two small observational studies favoured CBT and group  
16 psychotherapy in terms of reduced substance misuse and improved  
17 symptoms of psychosis.

18

19 The study of nidotherapy, suggests that collaborative psychosocial  
20 interventions involving the systematic assessment and modification of the  
21 environment may be worth studying further.

22

**Table 29: Relevant interventions included in current NICE guidelines and summary of evidence of effectiveness for people with psychosis and coexisting substance misuse**

Intervention name	Existing NICE guideline <sup>1</sup>	Recommended	Evidence relevant to people with psychosis and substance misuse
<b>Opportunistic brief interventions</b>			
Brief interventions for people not in contact with services	Substance misuse: DMP	Yes <sup>3</sup>	–
Brief interventions for people in contact with services	Substance misuse: DMP	Yes <sup>3</sup>	–
<b>Self-help based interventions</b>			
Self-help intervention (including self-help groups, 12-step self-help groups)	Substance misuse: Alcohol <sup>2</sup> DMP Psychosis: Bipolar disorder	Yes Yes Yes <sup>3</sup>	–
Twelve-step facilitation	Substance misuse: Alcohol <sup>2</sup>	Yes <sup>4</sup>	
<b>Behavioural therapies</b>			
Cue exposure	Substance misuse: Alcohol <sup>2</sup>	Yes (BT in general recommended)	–
Behavioural self-control training	Substance misuse: Alcohol <sup>2</sup>	Yes (BT in general recommended)	–
Contingency management	Substance misuse: Alcohol <sup>2</sup> DMD DMP	Research rec Yes Yes	Low quality evidence in favour of contingency management.
<b>Cognitive and behavioural based therapies</b>			
CBT	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Bipolar disorder Schizophrenia (update)	Yes No Yes <sup>3</sup> Yes <sup>3</sup> Yes	Moderate to low quality evidence available, but insufficient to reach conclusion about direction of effect.
Coping and social skills training	Substance misuse: Alcohol <sup>2</sup>	No	Moderate to low quality evidence available, but insufficient to reach conclusion about direction of effect.

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Relapse prevention	Substance misuse: Alcohol <sup>2</sup> DMD Psychosis: Bipolar disorder	Not specifically <sup>5</sup> No Yes <sup>3</sup>	-
<b>Family-based interventions</b>			
Family intervention	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Bipolar disorder Schizophrenia (update)	Yes <sup>3</sup> No Yes <sup>3</sup> Yes <sup>3</sup> Yes <sup>3</sup>	-
<b>Motivational techniques</b>			
Motivational interviewing/ Motivational Enhancement Therapy	Substance misuse: Alcohol <sup>2</sup> DMP	Yes <sup>4</sup> No	Moderate to low quality evidence in favour of motivational interviewing.
<b>Social Network and Environment Based Therapies</b>			
Social Behaviour and Network Therapy	Substance misuse: Alcohol <sup>2</sup>	Not specifically <sup>6</sup>	-
The Community Reinforcement Approach	Substance misuse: Alcohol <sup>2</sup> DMD	Not specifically <sup>6</sup> No	-
Social-systems interventions	Substance misuse: DMD DMP	No No	-
<b>Other interventions</b>			
Adherence therapy	Psychosis: Schizophrenia (update)	No	-
Arts therapies	Psychosis: Schizophrenia (update)	Yes	-
Cognitive remediation	Psychosis: Schizophrenia (update)	No	-
Counselling and supportive psychotherapy	Substance misuse: Alcohol <sup>2</sup> Psychosis: Schizophrenia (update)	No No	-
Couples-based interventions (including behavioural couples therapy)	Substance misuse: Alcohol <sup>2</sup> DMD DMP	Yes Yes <sup>3</sup>	-
Individual drug counselling	Substance misuse: DMD	No	-
Interpersonal and social rhythm therapy (IPSRT)	Psychosis: Bipolar disorder	Yes <sup>3</sup>	-
Interpersonal therapy	Substance misuse: DMD	No	-

	DMP	No	
Multi-modal care programmes	Substance misuse: Alcohol <sup>2</sup> DMP	Yes <sup>3</sup> No	-
Psychoeducational interventions	Substance misuse: Alcohol <sup>2</sup> DMP Psychosis: Bipolar disorder Schizophrenia (update)	No No Yes <sup>3</sup> No	-
Psychodynamic psychotherapy and psychoanalysis	Substance misuse: Alcohol <sup>2</sup> DMD DMP Psychosis: Schizophrenia (update)	No No No No	-
Social skills training	Psychosis: Schizophrenia (update)	No	-
Vocational interventions	Substance misuse: DMP	No	-
<p><i>Note.</i> DMD = Drug misuse: opioid detoxification; DMP = Drug misuse: psychosocial interventions; Research rec = Research recommendation (from NICE guideline).</p> <p><sup>1</sup> Available from <a href="http://www.nice.org.uk">www.nice.org.uk</a>.</p> <p><sup>2</sup> Management of alcohol dependence guideline.</p> <p><sup>3</sup> For specific groups and/or in certain circumstances (see relevant guideline for further information).</p> <p><sup>4</sup> These interventions were seen as components of any effective psychosocial intervention delivered in alcohol services with the assessment and enhancing of motivation forming a key element of the assessment process.</p> <p><sup>5</sup> Interventions that promote abstinence and prevent relapse recommended.</p> <p><sup>6</sup> But social network therapies recommended.</p>			

1

## 2 7.2.8 Health economic evidence (psychological/ psychosocial 3 interventions)

4 The systematic search of the health economics literature identified two  
5 relevant papers: one comparing the cost-effectiveness of CBT combined with  
6 MI versus standard care (Haddock *et al.*, 2003) and one comparing a group  
7 behavioural skills programme or case management with a twelve-step control  
8 condition (Jerrell & Ridgley, 1997). Details on the methods used for the  
9 systematic search of the economics literature are described in Appendix 9.

10

11 One UK study (Haddock *et al.* 2003), based on the RCT conducted by  
12 Barrowclough and colleagues (2001), evaluated the cost-effectiveness of an  
13 integrated programme of CBT combined with MI plus standard care versus  
14 standard care alone. The study sample consisted of 36 people diagnosed with  
15 psychosis and coexisting substance dependence or misuse along with their

1 carers, recruited from the mental health units of three UK NHS hospital  
2 trusts. Resource use and outcome data were collected over 18 months follow-  
3 up. The study adopted a societal perspective, with data on hospital care,  
4 primary care, community and domiciliary services, medications, service user  
5 travel and out-of-pocket expenses and productivity losses all collected from  
6 the Client Service Receipt Inventory (CSRI). The primary measure of  
7 effectiveness was change in the Global Assessment of Functioning Scale  
8 (GAF).

9  
10 Over 18 months follow-up, the intervention group was on average £1,260 ( $p =$   
11 0.25) less costly, while experiencing an average of 22.5% improvement in GAF  
12 scores in comparison to routine care. Incremental cost-effectiveness ratios  
13 were calculated by the authors but not reported in the paper. Cost-  
14 effectiveness acceptability curves (CEACs) were used to measure uncertainty  
15 around the sample estimates of mean costs and outcomes. The probability of  
16 the intervention being less costly than standard care (at a willingness-to-pay  
17 of 0) was 69.3%. Overall, the authors concluded that the integrated  
18 programme of CBT combined with MI was no more costly than standard care,  
19 and there was a high probability of it being cost-effective. The results of the  
20 study are relevant to the UK setting, although the major limitations are the  
21 small sample size (which may not have been representative of the study  
22 population) and the measure of effectiveness used in the analysis (which  
23 limits comparability across health care interventions). Furthermore, the study  
24 adopted a societal rather than an NHS and PSS perspective as recommended  
25 by NICE (NICE, 2008). However, differences between the two treatment  
26 groups, in terms of societal costs including patient travel and out-of-pocket  
27 expenditure and productivity losses, were not significant. Therefore, inclusion  
28 of these costs did not significantly alter the overall results of the cost-  
29 effectiveness analysis.

30  
31 One US-based study was identified that assessed the cost-effectiveness of two  
32 outpatient programmes (behavioural skills training, case management) with a  
33 twelve-step control condition (Jerrell & Ridgely, 1997). The study population  
34 included 132 people with an axis I DSM-III-R diagnosis of psychosis or major  
35 affective disorder with a coexisting substance disorder and previous  
36 psychiatric treatment. The primary measures of effectiveness in the study  
37 were psychological functioning, psychiatric and substance abuse symptoms.  
38 As no significant differences in clinical effectiveness were detected across the  
39 three treatment groups, the economic analysis was based on differences in  
40 costs only. A societal perspective was taken for the cost analysis, with data on  
41 mental health and general health care resource use, criminal justice and social  
42 services, family and caregiver resources and any other transfer payments,  
43 collected over an 18-month period. Total costs were reported separately for

1 intensive mental health care (inpatient days, residential treatment, emergency  
2 visits) and supportive mental health care (outpatient visits, medication visits,  
3 and supported housing visits).

4  
5 For intensive mental health care costs, the total cost in the twelve-step group  
6 was \$10,275, in the behavioural skills group was \$4,276 and in the case  
7 management group was \$7,643. For supportive mental health care costs, the  
8 total cost in the twelve -step group was \$7,798, in the behavioural skills group  
9 was \$6,112 and in the case management group was \$5,970. No formal  
10 statistical tests were conducted to quantify the significance of any cost  
11 differences between the three treatment groups. Overall, the authors  
12 concluded that no differences in outcomes were detected between the three  
13 groups, but the twelve -step group incurred the highest intensive and  
14 supportive costs over the 18-month period. The study is of limited relevance  
15 to the UK context as it was based in the US and has a number of  
16 methodological limitations. The partial randomised study design and lack of  
17 information about the power of the study, in terms of detecting differences  
18 between the three treatment groups, limits the internal validity of the  
19 effectiveness results. Resource use components were not described separately  
20 from costs and it is not possible to ascertain whether the cost analysis was  
21 based on actual costs or service charges.

## 22 *Health Economics Summary*

23 In summary, there was limited evidence of the cost-effectiveness of specific  
24 psychological/psychosocial interventions for people with psychosis and  
25 coexisting substance misuse. The UK-based study by Haddock and colleagues  
26 (2003) suggested that a combination of CBT and MI plus standard care was  
27 cost-effective compared with standard care alone. The US based study by  
28 Jerrell and Ridgely (1997) showed that a behavioural skills training was more  
29 costly in terms of intensive and supportive mental health care, when  
30 compared with 12-step recovery or case management programmes.

31  
32 Given the uncertainty surrounding the cost-effectiveness of  
33 psychological/psychosocial interventions and the associated resource  
34 implications, it was anticipated that further economic modelling would be  
35 developed to address these issues. However, due to both the scarcity and the  
36 generally low quality of the clinical data that was identified in the guideline  
37 systematic review, the GDG agreed that it would not be possible to model the  
38 cost-effectiveness of specific psychological/psychosocial interventions in  
39 people with psychosis and coexisting substance misuse.

## 1 **7.2.9 From evidence to recommendations (psychological/ 2 psychosocial interventions)**

3 Early in the development process, the GDG distinguished between outcomes  
4 that were critical to decision making and those that were important but not  
5 critical. Critical outcomes included: mortality (all causes), relapse rates  
6 (measured by exacerbation of symptoms requiring change in health care  
7 management), substance misuse (however measured), global and social  
8 functioning (for example, employment and accommodation), subjective  
9 quality of life, satisfaction with care, and physical morbidity. Only critical  
10 outcomes were included in the GRADE evidence profiles and considered  
11 when making recommendations.

12  
13 There was little direct evidence relating to most psychological interventions  
14 for people with psychosis and coexisting substance misuse. The evidence that  
15 was available was generally difficult to interpret because of the context the  
16 research was conducted in and/or methodological issues. As a result, the  
17 GDG decided that it was not possible to recommend any specific  
18 psychological or psychosocial intervention or combination of interventions to  
19 people with psychosis and coexisting substance misuse. Nevertheless, the  
20 GDG thought that given the positive evidence in favour of contingency  
21 management (even if poor quality), a recommendation should be made that  
22 people with psychosis and coexisting substance misuse should not be  
23 excluded from contingency management programmes because of their  
24 psychosis. In general though, as no good quality evidence was found relating  
25 to the modification of interventions recommended for people with a single  
26 diagnosis, the GDG concluded that people with psychosis and coexisting  
27 substance misuse should be offered the same range of evidence-based  
28 interventions recommended for people with a single diagnosis.

29  
30 However, the GDG felt it was important to emphasise that low levels of  
31 substance use that would not usually be considered harmful or problematic in  
32 people without psychosis, can have a significant impact on the mental health  
33 of people with psychosis.

34  
35 In addition, the GDG, whilst unwilling to make specific recommendations  
36 about environmental modifications such as nidotherapy, thought it would be  
37 important that research is undertaken to assess the potential for such  
38 modifications for people with psychosis and coexisting substance misuse.

39  
40 There was no evidence that addressed the two sub-questions regarding  
41 elements of an integrated service model and subgroups of people (see section  
42 7.2.3 for further information about these sub-questions). In addition, the GDG

1 noted that valuable information about the potential benefits of  
2 pharmacological and psychosocial interventions for people with psychosis  
3 and substance misuse could be obtained from trials of treatments for people  
4 with either of these two different types of problems. However, to date, most  
5 trials conducted among people with psychosis have excluded those who have  
6 coexisting substance misuse and nearly all trials among people with  
7 substance misuse have excluded those with coexisting psychosis. In some  
8 instances, it may be necessary to exclude people with coexisting problems  
9 from future studies. However, very often, this important and prevalent group  
10 of patients have been excluded from intervention trials with no clear reason  
11 being offered. Therefore, future research should not routinely exclude people  
12 with psychosis and coexisting substance misuse.

## 1 7.3 CLINICAL PRACTICE RECOMMENDATIONS

### 2 7.3.1 Recommendations (psychological/ psychosocial 3 interventions)

#### 4 *Secondary care mental health services*

#### 5 **Treatment**

6 **7.3.1.1** Before starting treatment for adults and young people with psychosis  
7 and coexisting substance misuse, review:

- 8 • the diagnosis of psychosis and of the coexisting substance  
9 misuse, especially if either diagnosis has been made during  
10 a crisis or emergency presentation **and**
- 11 • the effectiveness of previous and current treatments and the  
12 person's tolerance of them; discontinue ineffective  
13 treatments.<sup>13</sup>

14 **7.3.1.2** Ensure that adults and young people with psychosis and coexisting  
15 substance misuse are offered evidence-based treatments for both  
16 conditions (see 7.3.1.3 and 7.3.1.4).<sup>14</sup>

17 **7.3.1.3** For the treatment of psychosis, see 'Bipolar disorder: the management  
18 of bipolar disorder in adults, children and adolescents, in primary  
19 and secondary care' (NICE clinical guideline 38) or the guideline on  
20 schizophrenia (NICE clinical guideline 82).<sup>15</sup>

21 **7.3.1.4** For the treatment of substance misuse, see:

- 22 • 'Alcohol-use disorders: diagnosis and clinical management  
23 of alcohol-related physical complications' and the guideline  
24 on alcohol dependence and harmful alcohol use (NICE  
25 clinical guidelines 100 and CGXX) **and/or**
- 26 • 'Drug misuse: psychosocial interventions' and the guideline  
27 on opioid detoxification (NICE clinical guidelines 51 and  
28 52).<sup>16</sup>

---

<sup>13</sup> This recommendation also appears in section 8.3.1 where the pharmacological data is presented.

<sup>14</sup> This recommendation also appears in section 8.3.1 where the pharmacological data is presented.

<sup>15</sup> This recommendation also appears in section 8.3.1 where the pharmacological data is presented.

<sup>16</sup> This recommendation also appears in section 8.3.1 where the pharmacological data is presented.

- 1 **7.3.1.5** When developing a treatment plan for a person with psychosis and  
2 coexisting substance misuse, tailor the plan and the sequencing of  
3 treatments to the person and take account of:
- 4 • the relative severity of both the psychosis and the substance  
5 misuse at different times
  - 6 • the person's social and treatment context and
  - 7 • the person's readiness for change.
- 8 **7.3.1.6** Do not exclude adults and young people with psychosis and coexisting  
9 substance misuse from contingency management programmes  
10 because of their psychosis.

11 **7.3.2 Research recommendations (psychological/ psychosocial**  
12 **interventions)**

- 13 **7.3.2.1** Are interventions for psychosis or substance misuse clinically and cost  
14 effective when compared with standard care for people with  
15 psychosis and coexisting substance misuse?<sup>17</sup>
- 16 **7.3.2.2** Are psychosocial interventions clinically and cost effective when  
17 compared with standard care for people with psychosis and  
18 coexisting substance misuse?
- 19 **7.3.2.3** Are environmental interventions clinically and cost effective when  
20 compared with standard care for people with psychosis and  
21 coexisting substance misuse?

22

---

<sup>17</sup> This recommendation also appears in section 8.3.2 where the pharmacological data is presented.

# 1 8 PHARMACOLOGICAL AND 2 PHYSICAL INTERVENTIONS 3 FOR PEOPLE WITH PSYCHOSIS 4 AND COEXISTING SUBSTANCE 5 MISUSE

## 6 8.1 INTRODUCTION

7 There are many pharmacological treatments for both psychotic disorders and  
8 substance misuse, but there is very little overlap between the treatments for  
9 each group of disorders. The pharmacological treatments for each of the  
10 substance misuse disorders are generally specific ones for each substance of  
11 dependence, for example, disulfiram and acamprosate for alcohol  
12 dependence; methadone for opioid addiction. In the treatment of psychoses,  
13 however, there is much greater overlap, with lithium salts and other mood  
14 stabilisers, antipsychotics of all types, and anticonvulsants being used; these  
15 medications show little commonality with the treatments for substance  
16 misuse. It might be expected that with a large number of drugs being used to  
17 treat each group of disorders, there could be important interactions between  
18 them, both pharmacodynamic and pharmacokinetic. In practice, interactions  
19 appear to be rare and generally unimportant. It might also be expected that  
20 polypharmacy would be a problem for these dual disorders but the data here  
21 are conflicting with no clear evidence of greater use of drug treatment in  
22 people with psychosis and coexisting substance misuse (Centorrino *et al.*,  
23 2008; Goldberg *et al.*, 2009; Kreyenbuhl *et al.*, 2007).

24  
25 To date, few specific recommendations for pharmacological treatment of both  
26 groups of disorders have been made that are not covered by previous  
27 published NICE guidelines for substance misuse and the psychoses  
28 separately. The purpose of this chapter is to examine whether there is any  
29 evidence that pharmacological/physical treatment of each disorder should be  
30 modified as result of having a coexisting disorder.

### 31 8.1.1 Current practice

32 The pharmacological management of service users with psychosis and  
33 substance misuse is primarily concerned with treating the individual  
34 disorders. Nevertheless, special attention needs to be paid to treatment  
35 adherence in this group, not least as the risk of adverse outcomes, including

1 significant societal violence, is so much greater in this population (Kooyman  
2 *et al.*, 2007).  
3

## 4 **8.2 EVIDENCE REVIEW**

### 5 **8.2.1 Introduction**

6 A number of existing NICE guidelines have reviewed the evidence for  
7 pharmacological and physical interventions used to treat people with  
8 psychosis without substance misuse (that is, bipolar disorder and  
9 schizophrenia), and for people with substance misuse without psychosis (that  
10 is, alcohol and drug misuse: opioid detoxification).  
11

12 For the purposes of the current guideline, three main issues were addressed  
13 for people with psychosis and coexisting substance misuse. First, modification  
14 of the pharmacological treatment of psychosis as a result of substance misuse  
15 and the treatment provided (for example, methadone, buprenorphine *etc.*).  
16 Second, modification of the pharmacological/physical treatment of substance  
17 misuse as a result of the presence of psychosis and the treatment provided  
18 (for example, antipsychotic drugs, lithium). Third, management of drug  
19 interactions or adverse effects from pharmacological interventions.  
20

21 Where no evidence existed for a particular intervention in people with  
22 psychosis and coexisting substance misuse, the GDG used informal consensus  
23 to reach a conclusion about whether it was appropriate to cross-reference to  
24 existing NICE guidance.

#### 25 *Interventions and licensing in the UK*

26 Table 30 lists the interventions included in current NICE guidelines together  
27 with their licensed indications in the UK (those relevant to this guideline).  
28  
29

**Table 30: Relevant interventions included in current NICE guidelines and current licence status of medication**

Intervention type/use	Name	UK licence (only relevant indications listed)	Reviewed by existing NICE guideline
<b>MEDICATION</b>			
Alcohol dependence	Acamprosate calcium	Maintenance of abstinence in alcohol dependence (it should be combined with counseling)	Alcohol (management of alcohol dependence guideline)
Alcohol deterrent compounds	Disulfiram	Adjuvant in the treatment of carefully selected and co-operative service users with drinking problems (?15+)  Its use must be accompanied by appropriate supportive treatment	Alcohol (management of alcohol dependence guideline)
Alpha-adrenergic agonists	Clonidine	Hypertension; migraine (13+)	DMD
Alpha-adrenergic agonists	Lofexidine	Management of symptoms of opioid withdrawal (18+)	DMD
Antiepileptic drugs	Phenytoin	All forms of epilepsy except absence seizures; status epilepticus	Alcohol (clinical management guideline)
Antiepileptic drugs	Topiramate	Generalised tonic-clonic seizures or partial seizures	Alcohol (management of alcohol dependence guideline)
Antimanic drugs	Lithium	Bipolar disorder (12+)	Bipolar
Antimanic drugs	Valproic acid	Manic episodes associated with bipolar disorder (18+)  Treatment of generalised, partial or other epilepsy. No mention of manic episodes	Bipolar
Antimanic drugs/	Benzodiazepine: Diazepam	Adjunct in acute alcohol withdrawal; short-term use	Bipolar/ Alcohol <sup>1</sup>

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Anxiolytics		in anxiety or insomnia	
Antimanic drugs/ Anxiolytics	Benzodiazepine: Lorazepam	Short-term use in anxiety or insomnia, acute excitement and acute mania	Bipolar
Antimanic drugs/ Hypnotics	Benzodiazepine: Chlordiazepoxide	Adjunct in acute alcohol withdrawal; Short-term treatment of severe anxiety that is severe with or without insomnia/short-term psychosomatic/organic or psychotic illness	Alcohol <sup>1</sup>
Antimanic drugs/ Hypnotics	Chlormethiazole	Alcohol withdrawal	Alcohol <sup>1</sup>
Antimanic/ Control of epilepsy	Carbamazepine	Prophylaxis of bipolar disorder unresponsive to lithium	Bipolar
Antipsychotic drugs (first-generation)	For example: Chlorpromazine Haloperidol	Schizophrenia; mania	Bipolar/Schizophrenia (update)
Antipsychotic drugs (second-generation)	For example: Clozapine Olanzapine Risperidone	Schizophrenia; some individual drugs also indicated for mania. Note, clozapine only indicated for schizophrenia in service users unresponsive to, or intolerant of, first-generation antipsychotic drugs	Bipolar/Schizophrenia (update)
Opioid agonists & partial agonists	Buprenorphine	Treatment for opioid drug dependence (subutex) (16+)	DMD
Opioid agonists & partial agonists	Methadone	Treatment of opioid drug addictions (?15+)	DMD
Opioid antagonists	Nalmefene	Unlicensed	Alcohol <sup>1</sup> / DMD
Opioid antagonists	Naltrexone	Adjunctive prophylactic therapy in the maintenance of detoxified formerly opioid dependent service users (18+)	Alcohol <sup>1</sup> / DMD
Serotogenic agents	Ondansetron	Prevention and treatment of postoperative nausea and vomiting	Alcohol

Serotogenic agents	SSRIs	Depression	Alcohol <sup>1</sup> / Depression
Skeletal muscle relaxants	Baclofen	Chronic severe spasticity	Alcohol <sup>1</sup>
<b>PHYSICAL AND COMPLEMENTARY INTERVENTIONS</b>			
Physical	Acupuncture	-	DMD
Physical	Electrical transcranial stimulation	-	Alcohol <sup>1</sup>
Complementary	Kudzu root	-	Alcohol <sup>1</sup>
Complementary	Vipassana meditation	-	Alcohol <sup>1</sup>
<i>Note.</i> DMD = drug misuse: opioid detoxification. <sup>1</sup> Management of alcohol dependence guideline.			

1

## 2 **8.2.2 Clinical review protocol (pharmacological/ physical** 3 **interventions)**

4 The review protocol, including the primary review question, information  
5 about the databases searched and the eligibility criteria used for this section of  
6 the guideline can be found in Table 31. Initially a search for systematic  
7 reviews and existing guidelines that addressed the review question was  
8 conducted. Good quality systematic reviews were then used as a source of  
9 evidence, and only a new systematic search for more recent primary-level  
10 studies was conducted for the guideline (further information about the search  
11 strategy can be found in Appendix 7).

12

13 If the evidence allowed, the following sub-question was asked for review  
14 question 2.1.1 and 2.3.1: Are there sub-groups of people (for example, young  
15 people, people with a particular type of psychosis, people from BME groups)  
16 that may benefit from alternative strategies than those recommended for  
17 people with a single disorder?

18

**Table 31: Databases searched and eligibility criteria for clinical evidence**

Component	Description
Review questions	<p>2.1.1 For people with psychosis and coexisting substance misuse, should the medical treatment of their psychosis be modified as a result of substance misuse and the treatment provided (for example, methadone, buprenorphine etc)?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>2.3.1 For people with psychosis and coexisting substance misuse, should the medical/physical treatment of substance misuse be modified as a result of the presence of psychosis and the treatment provided (for example, antipsychotics, lithium)?</p> <p>A) During the acute phase? B) During non-acute phase?</p> <p>If so, how should treatment be modified?</p> <p>2.5.1 In people with psychosis and substance misuse, is there any evidence that the management of drug interactions or adverse effects from pharmacological treatments should be different from those people without coexisting disorders?</p> <p>If so, how should management of drug interactions be modified?</p>
Electronic databases	CENTRAL, CINAHL, EMBASE, MEDLINE, PsycINFO
Date searched	Inception to 26.05.2010
Study design	Reviews, clinical guidelines, primary-level studies
Population	People with psychosis and coexisting substance misuse
Intervention(s)	Pharmacological/physical interventions
Comparison	Any relevant treatment
Critical outcomes	<p>Reduced mortality (all causes)</p> <p>Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</p> <p>Reduced substance misuse (however measured)</p> <p>Improved global and social functioning (for example, employment, accommodation)</p> <p>Improved subjective quality of life</p> <p>Improved satisfaction with care</p> <p>Reduced physical morbidity.</p>
<i>Note.</i> BME = Black and minority ethnic.	

1  
2

### 1 **8.2.3 Studies considered for review (pharmacological/ physical** 2 **interventions)**<sup>18</sup>

3 Thirteen clinical evidence reviews and guidelines met the eligibility criteria  
4 for this section of the guideline (Buchanan *et al.*, 2009 [Schizophrenia Patient  
5 Outcomes Research Team, PORT]; Casas *et al.*, 2008; Center for Substance  
6 Abuse Treatment, 2005a [Treatment Improvement Protocol series 42]; Center  
7 for Substance Abuse Treatment, 2005b [Treatment Improvement Protocol  
8 series 43]; Center for Substance Abuse Treatment, 2006 [Treatment  
9 Improvement Protocol series 45]; Green *et al.*, 2008; Hjorthoj *et al.*, 2009; Mills  
10 *et al.*, 2009 [Australian guideline]; San *et al.*, 2007; Smelson *et al.*, 2008; Tiet &  
11 Mausbach, 2007; Vornick & Brown, 2006; Wobrock & Soyka, 2008). All were  
12 published in peer-reviewed journals between 2006 and 2009. In addition, a  
13 number of reviews were excluded as they had either been superseded by  
14 more recent reviews (for example, Brunette *et al.*, 2005; Goldstein *et al.*, 2006;  
15 Green, 2005), or are currently under review (that is, Lingford-Hughes *et al.*,  
16 2004).

17  
18 In addition, a search was conducted for RCT evidence that may have been  
19 published too recently to be included in existing reviews. From this, four  
20 RCTs were found: BROWN2009 (Brown *et al.*, 2009), KEMP2009 (Kemp *et al.*,  
21 2009), NEJTEK2008 (Nejtek *et al.*, 2008), VANNIMWEGEN2008 (Van  
22 Nimwegen *et al.*, 2008). A summary of study characteristics is given in Table  
23 32 and the results are described in the text below. Additionally, a secondary  
24 analysis from the Clinical Antipsychotic Trials of Intervention Effectiveness  
25 project was reviewed (CATIE2008; Swartz *et al.*, 2006).

---

<sup>18</sup> Here and elsewhere in the guideline, each RCT considered for review is referred to by a study ID (primary author and date of study publication, except where a study is in press or only submitted for publication, then a date is not used).

1

**Table 32: Study information table for RCTs of pharmacological interventions**

	Pharmacological interventions versus any control
Total no. of trials (N)	4 RCTs (216)
Study ID	(1) BROWN2009 (2) KEMP2009 (3) NEJTEK2008 (4) VANNIMWEGEN2008
Number randomised	(1) 50 (2) 31 (3) 94 (4) 41
Diagnosis	(1) Bipolar disorder I or II and alcohol dependence (2) Rapid cycling bipolar disorder I or II and substance abuse and/or dependence (3) Bipolar disorder I or II with and without psychotic features and stimulant dependence, currently in manic or hypomanic episode (4) Schizophrenia or schizophrreniform disorder and cannabis misuse
Treatment (mean dose) (n)	(1) Naltrexone (50 mg/day) + CBT (n=23) (2) Lithium (1440 mg/day; range 900-2400 mg) (n=16) (3) Risperidone (3.1 mg/day +/- 1.2 mg) (n=46) (4) Olanzapine (11.1mg) (n=20)
Control (mean dose) (n)	(1) Placebo + CBT (all with usual medication) (n=27) (2) Lithium (1400 mg/day; range 600-2100 mg) + divalproex (1583 mg/day; range 1000-3250 mg) (n=15) (3) Quetiapine (303.6 mg/day +/- 151.9 mg) (n=48) (4) Risperidone (3mg) (n=21)
Treatment length/design	(1) 12 weeks, double-blind RCT (2) 25 weeks, double-blind RCT (3) 20 weeks, double-blind RCT (5) 6 weeks; double-blind RCT
Country	(1) US (2) US (3) US (5) The Netherlands
<i>Note.</i> N = Total number of participants; n = number of participants in each group.	

2

### 3 **8.2.4 Evidence from existing reviews and guidelines for the use** 4 **of pharmacological interventions to treat people with** 5 **schizophrenia and coexisting substance misuse**

6 Eleven recent existing reviews and/or guidelines included evidence for the  
7 pharmacological treatment of people with coexisting schizophrenia (or related  
8 disorders) and substance misuse (Buchanan *et al.*, 2009 [Schizophrenia Patient

1 Outcomes Research Team, PORT]; Center for Substance Abuse Treatment,  
2 2005a [Treatment Improvement Protocol series 42]; Center for Substance  
3 Abuse Treatment, 2005b [Treatment Improvement Protocol series 43]; Center  
4 for Substance Abuse Treatment, 2006 [Treatment Improvement Protocol series  
5 45]; Green *et al.*, 2008; Hjorthoj *et al.*, 2009; Mills *et al.*, 2009 [Australian  
6 guideline]; San *et al.*, 2007; Smelson *et al.*, 2008; Tiet & Mausbach, 2007;  
7 Wobrock & Soyka, 2008). They review a range of evidence, from case studies  
8 to RCTs.

9

10 Buchanan and colleagues (2009) updated the PORT psychopharmacological  
11 treatment recommendations last published in 2004 (Lehman *et al.*, 2004). The  
12 authors conducted a systematic review of evidence sourced from quarterly  
13 searches of MEDLINE (January 2002 to March 2008) to supplement searches  
14 undertaken for their previous guideline. No other electronic database was  
15 used. The guideline covers pharmacological treatments for schizophrenia,  
16 with a subsection on the treatment of coexisting substance misuse. It mostly  
17 focuses on double-blind RCTs. It included studies provided at least 50% of  
18 participants had a schizophrenia spectrum disorder diagnosis and where  
19 study drugs had US Food and Drug Administration (FDA) approval. Studies  
20 involving people with coexisting schizophrenia and cocaine abuse or  
21 dependence included two double-blind RCTs comparing olanzapine to  
22 haloperidol, and one double-blind RCT comparing olanzapine to risperidone.  
23 Also included was one double-blind RCT comparing naltrexone to placebo in  
24 people with coexisting schizophrenia and alcohol use disorders. Finally, the  
25 authors mention a sub-analysis of a larger RCT that examined naltrexone,  
26 disulfiram, and naltrexone plus disulfiram compared to placebo in people  
27 with psychosis and coexisting substance misuse. The guideline development  
28 group concluded that based on the research examined there was insufficient  
29 evidence to support a specific recommendation for a pharmacological  
30 intervention to treat people with coexisting schizophrenia and substance  
31 misuse.

32

33 Green and colleagues (2008) conducted a narrative review of evidence, but  
34 did not describe their methodology for identifying relevant research. The  
35 authors focus on antipsychotic drugs for the treatment of coexisting  
36 schizophrenia and substance misuse, but also cover medications for substance  
37 disorders. They report a range of evidence (mostly low level evidence such as  
38 case reports and open-label non-comparative studies) suggesting that  
39 “atypical” antipsychotics may be helpful in reducing substance misuse in  
40 people with coexisting schizophrenia and substance misuse. The evidence  
41 reviewed covered a range of drugs of abuse, including alcohol, cocaine and  
42 marijuana. They found the most consistent evidence (from non-randomised  
43 studies) suggesting that clozapine treatment may reduce substance use. There

1 was 'less substantial' evidence for quetiapine and aripiprazole, while that for  
2 olanzapine and risperidone is unclear, with some studies showing a benefit  
3 and others not. Overall they concluded that RCT evidence is required before  
4 firmer conclusions can be drawn.

5  
6 With regard to evidence for drugs specifically used to treat substance misuse,  
7 Green and colleagues found preliminary evidence to support the use of  
8 naltrexone and disulfiram in people with coexisting schizophrenia and  
9 alcohol dependence. They found no relevant studies of acamprosate. They  
10 report case studies indicating the potential benefit of valproic acid in people  
11 with coexisting schizophrenia and alcohol abuse or dependence.

12  
13 However, Green and colleagues conclude that "despite numerous suggestive  
14 reports, the questions of whether and to what degree antipsychotic  
15 medications and other medications for substance use disorders are effective in  
16 reducing substance use among people with [schizophrenia and] co-occurring  
17 disorders are not yet answered."

18  
19 Hjorthoj and colleagues (2009) conducted a systematic review focusing on the  
20 treatment of cannabis use disorder in schizophrenia spectrum disorders,  
21 covering all types of intervention including psychosocial. The evidence was  
22 sourced from searches of four electronic databases searched to September  
23 2008. The authors focused on studies which provided outcomes for cannabis  
24 use separately from outcomes for other substance misuse, although also  
25 looked at studies which reported cannabis use as part of a grouped outcome.  
26 With regard to pharmacological interventions for reducing cannabis use, they  
27 found evidence from non-randomised studies of benefit from using clozapine  
28 and quetiapine.

29  
30 The Australian Government Department of Health and Ageing funded the  
31 National Drug and Alcohol Research Centre (Mills *et al.*, 2009) to develop a  
32 guideline covering the management of people with mental health conditions  
33 with coexisting alcohol and other drug abuse. The guideline, designed for  
34 alcohol and other drug workers, was based on a comprehensive review of the  
35 available evidence together with the experience of an expert panel. However,  
36 no details of the methodology used to undertake the review work were  
37 provided. For people with psychosis, Mills and colleagues found evidence  
38 that clozapine may be useful, but that evidence of benefit for second-  
39 generation antipsychotics is not yet clear. The guideline authors also suggest  
40 that pharmacological interventions may be more effective than psychosocial  
41 interventions, because negative symptoms associated with psychosis may  
42 restrict involvement and outcomes from psychosocial interventions. In

1 addition, this group of people may have greater tolerance to medication  
2 regimes.

3

4 Mills and colleagues conclude that treatments which work for mental health  
5 disorders without coexisting substance misuse will also work for those with a  
6 coexisting disorder. They raise the issue of adherence and also the importance  
7 of an awareness of possible interactions and side effects.

8

9 San and colleagues (2007) produced a systematic review of treatment with  
10 antipsychotic drugs for people with coexisting schizophrenia and substance  
11 misuse. The evidence was sourced from searches of three electronic databases  
12 searched to November 2006. The authors found three RCTs comparing  
13 olanzapine with haloperidol, plus other non-RCT evidence. From this they  
14 concluded that there was preliminary evidence that compared with  
15 haloperidol, olanzapine is more effective in reducing cravings whilst retaining  
16 antipsychotic action, and that clozapine showed similar potential. They also  
17 concluded that older antipsychotics (first-generation) were not as appropriate  
18 in this population compared with newer drugs (second-generation) since they  
19 were more likely to increase EPS symptoms. Based on case reports, open and  
20 retrospective studies, they found that newer antipsychotics may be of use,  
21 although the evidence is generally weak. The authors point out the limitations  
22 of the evidence base, including small sample sizes, short follow-up periods,  
23 and high dropout rates, as well as the paucity of RCTs and blinded studies.

24

25 Smelson and colleagues (2008) conducted a review of FDA-approved  
26 medications for people with schizophrenia with coexisting substance misuse.  
27 There are no details of the methods used, including how evidence was  
28 sourced. However, they provide reasonably comprehensive tables of evidence  
29 found (compared with other reviews). They cover both medication for the  
30 treatment of schizophrenia (antipsychotics) and that for the treatment of  
31 substance misuse disorders. They conclude that there is very little evidence to  
32 support specific treatment recommendations and, therefore, that clinicians  
33 should base treatment decisions on what suits the service user in terms of  
34 efficacy and side effects. They found the most evidence suggesting benefit for  
35 clozapine, olanzapine and risperidone, although this evidence is not strong.  
36 They suggest that second-generation antipsychotics may be better for  
37 controlling drug craving in those with cocaine dependence. The authors make  
38 the point that non-adherence is a bigger threat to effective treatment rather  
39 than poor efficacy and, therefore, advocate clinicians should consider depot  
40 medication. The authors found evidence to support the use of disulfiram and  
41 naltrexone.

42

1 Tiet and Mausbach (2007) report a systematic review of studies of treatment  
2 for people with mental disorders, including schizophrenia and bipolar  
3 disorder, with coexisting substance abuse. Studies were sourced from a search  
4 of two electronic databases. The search date is unclear, but is probably no  
5 later than 2006. The authors estimated effect sizes using Cohen's *d* but they do  
6 not give confidence intervals. It is unclear whether, or how, they applied  
7 diagnostic criteria when assessing studies. The authors concluded that  
8 treatments which are effective in reducing psychiatric symptoms in those  
9 with mental disorder without coexisting substance abuse, also work with  
10 coexisting substance abuse, and those treatments that are effective for  
11 improving substance abuse also work in those with a mental disorder.  
12 Specifically, they found that naltrexone may reduce coexisting alcohol-related  
13 disorders. They found no evidence of enhanced efficacy with higher doses.

14

15 The Treatment Improvement Protocol (TIP) series 42, 43 and 45 published by  
16 the Center for Substance Abuse Treatment are based on systematic reviews  
17 and reviews of published meta-analyses together with the views of an expert  
18 consensus panel for the treatment of substance abuse in those with coexisting  
19 disorders (TIP series 42), medication treatment of opioid addiction –  
20 treatment of coexisting disorders (TIP series 43) and detoxification and  
21 substance misuse (TIP series 45). The methods for evidence review are not  
22 available, but the guidelines were drafted by expert panels.

23

24 Treatment Improvement Protocol series 42 (Center for Substance Abuse  
25 Treatment, 2005a) does not focus on specific pharmacological treatments, but  
26 on general management and care by clinicians, and special considerations  
27 (such as for pregnant women). It is not considered further here.

28

29 Treatment Improvement Protocol series 43 (Center for Substance Abuse  
30 Treatment, 2005b), which focuses specifically on opioid addiction,  
31 recommends stabilisation of addiction symptoms with methadone, and using  
32 newer antipsychotics as either initial or second-line treatment. This is based  
33 on the supposed lower side effect profile and increased effectiveness of many  
34 newer antipsychotics compared with older medications.

35

36 Treatment Improvement Protocol series 45 (Center for Substance Abuse  
37 Treatment, 2006), which focuses on detoxification, recommends avoiding  
38 abrupt withdrawal of existing medication because of the risk of withdrawal  
39 symptoms or precipitating a psychiatric episode. It recommends maintenance  
40 on existing medications, unless the person has been abusing the medication or  
41 the psychiatric symptoms were caused by the medication. It also recommends  
42 giving consideration to withdrawal of medications which lower seizure  
43 threshold during acute alcohol withdrawal, or at least using a loading dose or

1 schedule taper of a benzodiazepine. The authors point out the importance of  
2 balancing risks and benefits of medication for people with mental disorder  
3 and coexisting substance misuse. These include the tension between the  
4 tendency for some medications to ‘impair cognition and blunt feelings’ which  
5 may hinder people from addressing problems in their lives which they need  
6 to change in order to abstain from misused substances successfully. However,  
7 untreated mental disorders “can be powerful relapse triggers, especially for  
8 people with a long-standing pattern of relying on alcohol or other drugs to  
9 manage their symptoms”.

10  
11 With regard to psychotic disorders, TIP series 45 has no specific  
12 recommendations for treatment in the presence of coexisting substance abuse  
13 apart from usual care.

14  
15 Wobrock and Soyka (2008) conducted a systematic review of pharmacological  
16 treatment of people with schizophrenia or psychosis and coexisting substance  
17 misuse based on searches of five electronic databases searched to November  
18 2007. They report a range of evidence including other reviews, RCTs and case  
19 studies. With regard to first-generation antipsychotics, Wobrock and Soyka  
20 found that 'most studies reported that service users with the psychosis and  
21 coexisting substance misuse showed a generally poorer response to  
22 treatment'. Whether the authors are using studies with both substance abuse  
23 and substance non-abuse populations, or whether they are comparing studies  
24 with substance abuse populations with studies with non-abusing populations  
25 is unclear. They include a range of substances including alcohol. They found  
26 some evidence that switching to flupenthixol improves outcomes in alcohol or  
27 cocaine abuse.

28  
29 With regard to second-generation antipsychotics, Wobrock and Soyka found  
30 little high quality evidence, but concluded a theoretical case for the use of  
31 second-generation antipsychotics based on limited evidence that second-  
32 generation antipsychotics, particularly aripiprazole, clozapine, olanzapine,  
33 quetiapine and risperidone may be more effective than older antipsychotics  
34 for both psychotic symptoms and for reducing craving and drug  
35 consumption. They found some evidence for the use of naltrexone in  
36 controlling alcohol abuse, as well as for the use of disulfiram, but did not  
37 consider this to be appropriate because of the risk of inducing psychosis.

### 38 *Summary of evidence from reviews and guidelines*

39 Although some of the reviews and guidelines described above, either did not  
40 search widely for relevant studies, or did not describe the source of the  
41 evidence reviewed, they all came to the conclusion that there is poor evidence  
42 for the effectiveness of pharmacological interventions for people with

1 coexisting schizophrenia and substance misuse. Some authors concluded that  
2 no specific drugs can be recommended and that treatment should follow that  
3 used for schizophrenia alone, while others suggest that the limited evidence  
4 for several second-generation antipsychotics, including clozapine, quetiapine,  
5 risperidone and olanzapine should be interpreted as an indication for use of  
6 these drugs. All call for better quality research to be undertaken.

### 7 **8.2.5 Evidence from new RCTs for the use of pharmacological** 8 **interventions to treat people with coexisting schizophrenia** 9 **and substance misuse (pharmacological interventions)**

10 One additional RCT (VANNIMWEGEN2008) and a secondary analysis from  
11 an earlier RCT (CATIE2008) were found that were not included in the  
12 published reviews and guidelines.

13  
14 The VANNIMWEGEN2008 trial was a 6-week double-blind RCT comparing  
15 olanzapine with risperidone in people with schizophrenia, schizoaffective  
16 disorder or schizophreniform disorder with coexisting cannabis use.  
17 Participants were a subsample (N=41) of 138 in service users or outpatients  
18 from four mental health centres aged 18 to 30. The authors report no  
19 differences between the study drugs in terms of cannabis use or cravings.

20  
21 CATIE2008 was a secondary analysis of a large pragmatic trial that included  
22 1432 participants (643 substance users and 789 non-users). People with  
23 schizophrenia were recruited at 57 US sites and randomly assigned to  
24 olanzapine, perphenazine, quetiapine, risperidone or ziprasidone for up to 18  
25 months. Among the substance users, there were no significant differences  
26 between treatment groups in time to all-cause discontinuation. The authors  
27 also report that substance users and non-users were generally similar in terms  
28 of improvement of symptoms of psychosis and side-effects. An analysis of the  
29 effective of treatment on substance misuse outcomes has not yet been  
30 published.

#### 31 *Summary of evidence from new RCTs*

32 There is no new evidence showing increased effectiveness of any particular  
33 antipsychotic in reducing substance misuse in people with coexisting  
34 schizophrenia and substance misuse.

1 **8.2.6 Evidence from existing reviews and guidelines for the use**  
2 **of pharmacological interventions to treat people with**  
3 **coexisting bipolar disorder and substance misuse**  
4 **(pharmacological interventions)**

5 Two reviews focus solely on the treatment of people with coexisting bipolar  
6 disorder and substance misuse (Casas *et al.*, 2008; Vornik & Brown, 2006). In  
7 addition, three reviews and guidelines discussed above also cover bipolar  
8 disorder (Mills *et al.*, 2009; Tiet & Mausbach, 2007; Centre for Substance Abuse  
9 Treatment, 2006 [TIP series 45]).

10  
11 Casas *et al.* (2008) developed a guideline based on a systematic review of  
12 published evidence together with expert consensus and surveys of expert  
13 practice. Evidence was sourced from a search of MEDLINE (to 2005). How the  
14 evidence was assessed, or what outcomes were used, is unclear. Similarly the  
15 diagnostic criteria used to include or exclude studies are unclear.

16 Nevertheless, recommendations are made for the treatment of different  
17 episode types. With regard to mania, Casas and colleagues recommend that  
18 treatment for “concomitant substance use disorder ... should be initiated at  
19 the same time [as treatment for mania] without giving priority to one over the  
20 other. However, if substance abuse presents as an acute intoxication or  
21 abstinence syndrome, then the treatment of the manic episode must be  
22 adapted.” They recommend second-generation antipsychotics, as well as,  
23 carbamazepine and valproate, but not antidepressants. For rapid cycling  
24 bipolar disorder, Casas and colleagues recommend that treatment should be  
25 adapted if substance abuse presents as acute intoxication or abstinence  
26 syndrome, using the same drugs as are recommended for use in a manic  
27 episode; otherwise treat as for mania. The authors found that lithium was  
28 shown to be effective in young people with coexisting substance abuse, and  
29 that valproate was helpful in reducing alcohol consumption. They found no  
30 RCT evidence for carbamazepine, gabapentin, lamotrigine, or  
31 benzodiazepines.

32  
33 With regard to bipolar disorder, Mills and colleagues (2009) found evidence  
34 to suggest that alcohol use outcomes improved with the use of valproate; that  
35 carbamazepine and lithium may help to reduce substance misuse; and that  
36 quetiapine and lamotrigine may also be of value in those with cocaine  
37 dependence.

38  
39 In addition to the findings described above, Tiet and Mausbach (2007) found  
40 that the combination of valproate and lithium may reduce coexisting alcohol  
41 use in bipolar disorder.

1 With regard to TIP series 45 (Center for Substance Abuse Treatment, 2006),  
2 the general advice covered above, can also be applied to the treatment of  
3 bipolar disorder and coexisting substance abuse, the TIP series 45 guideline  
4 authors looked at drugs commonly prescribed for bipolar disorder. With  
5 regard to lithium, they concluded that “studies [...] have shown that lithium  
6 has no conclusively positive effect on rates of abstinence in either depressed  
7 or nondepressed patients.” They also state that “anticonvulsant mood  
8 stabilizers, such as divalproex sodium and carbamazepine, can be effective in  
9 controlling mania and, some evidence suggests, in coexisting addictive  
10 conditions as well. Carbamazepine is known to be as effective as some  
11 benzodiazepines in inpatient treatment of alcohol withdrawal and, because of  
12 its anticonvulsant properties, it may be a good choice for treating those  
13 service users at high risk of withdrawal seizures.”

14  
15 Vornik and Brown (2006) reviewed pharmacological interventions for bipolar  
16 disorder and coexisting substance abuse. There is no description of how  
17 evidence was sourced or of any criteria by which evidence was assessed,  
18 which makes it difficult to assess the overall quality of the conclusions drawn.  
19 The authors report some evidence from RCTs for the effectiveness of mood  
20 stabilisers, including carbamazepine for reducing depressive symptoms in  
21 bipolar disorder (depressed phase) and coexisting cocaine abuse; major  
22 depressive disorder and coexisting substance use; and valproate in reducing  
23 alcohol use. They report non-randomised evidence for lamotrigine in  
24 reducing psychiatric symptoms and cocaine use. They also found evidence for  
25 the effectiveness of antipsychotics, including quetiapine (randomised open-  
26 label) and aripiprazole (open-label, non-randomised) for reducing psychiatric  
27 symptoms and drug craving.

### 28 *Summary of evidence from reviews and guidelines*

29 As with schizophrenia, not all the reviews searched more than one electronic  
30 database or gave full details of their methodology, which makes it hard to  
31 judge their quality. However, the reviews and guidelines largely came to  
32 similar conclusions, other than concerning the use of lithium. Some used the  
33 Geller and colleagues (1998) trial in young people (see Chapter 9) as evidence  
34 for lithium’s effectiveness (for example, Casas *et al.*, 2008), but others found no  
35 particular effect (for example, TIP series 45). With regard to other drugs used  
36 as mood stabilisers, most reviewers found evidence for the use of  
37 carbamazepine, valproate for improving alcohol-related outcomes, and  
38 antipsychotics. One found low-level evidence for the use of lamotrigine.

1 **8.2.7 Evidence from new RCTs for the use of pharmacological**  
2 **interventions to treat people with coexisting bipolar**  
3 **disorder and substance misuse (pharmacological**  
4 **interventions)**

5 Three relevant RCTs were found which were not included in the published  
6 reviews and guidelines (BROWN2009, KEMP2009, NEJTEK2008).

7  
8 BROWN2009 reported results from a 12-week placebo-controlled double-  
9 blind RCT of naltrexone plus CBT in 50 people with bipolar disorder I or II  
10 (currently depressed or mixed phase) with coexisting alcohol dependence. All  
11 participants continued to take their usual medication throughout the trial. The  
12 authors report that although the decline in alcohol consumption was  
13 numerically greater in the naltrexone group, there was no significant  
14 difference between groups on the primary outcome (percentage of drinking  
15 days) or any secondary outcome.

16  
17 KEMP2009 reported results from a 6-month, double-blind, maintenance trial  
18 of lithium monotherapy versus the combination of lithium and divalproex in  
19 people with coexisting rapid-cycling bipolar disorder and substance abuse  
20 and/or dependence. Of 149 participants enrolled into an open-label acute  
21 stabilisation phase, 31 were randomised to the maintenance phase. The  
22 authors report no statistically significant advantage in using combination  
23 therapy in terms of the primary outcome measure (time to relapse; defined as  
24 treatment for a mood disorder) or any secondary outcome.

25  
26 NEJTEK2008 report results from a 20-week, double-blind, RCT comparing  
27 risperidone to quetiapine in people with coexisting bipolar disorder I or II  
28 and stimulant dependence. Of 96 participants who consented and were  
29 randomly assigned, 80 attended at least one follow up visit. The results  
30 suggested little difference between study medication in terms of drug use or  
31 craving, or mood.

32 *Summary of evidence from new RCTs*

33 When tested in an RCT, there was insufficient evidence to reach a conclusion  
34 about the effectiveness of using naltrexone or a combination of lithium with  
35 divalproex to improve alcohol-related outcomes in people with coexisting  
36 bipolar disorder and alcohol dependence. In terms of antipsychotic  
37 medication, evidence from one trial suggests little difference between  
38 risperidone and quetiapine, but a lack of placebo control makes it difficult to  
39 determine if these medications may be effective.

1 **8.2.8 Clinical evidence for the management of drug interactions**  
2 **or adverse events from pharmacological interventions in**  
3 **people with psychosis and coexisting substance misuse**  
4 **(pharmacological interventions)**

5 None of the reviews focus substantially on interactions between treatment  
6 medication and substances of misuse, or on adverse events which are specific  
7 to, or especially elevated in, those with psychosis and coexisting substance  
8 misuse compared with those with psychosis alone.

9  
10 Adverse events associated with most psychotropic drugs are well  
11 documented. For antipsychotics, these include extrapyramidal symptoms  
12 (notably with first-generation drugs), weight gain, and increased glucose and  
13 lipid levels, leading to increased risk of diabetes (notably with second-  
14 generation drugs). Clozapine, which is used in several of the trials discussed  
15 above, tends to be associated with more reports of side effects than other  
16 antipsychotic medication. However, as Green and colleagues (2008) state,  
17 interactions between psychotropic medications and drugs of abuse are rare.  
18 These authors also point out that some newer medication can be sedating  
19 which can be problematic with some drugs of abuse. In addition, Farren and  
20 colleagues (2000) reported near syncopal episode following cocaine use in a  
21 service user treated with clozapine.

22  
23 Meanwhile, pharmacological treatments for alcohol abuse, such as naltrexone  
24 and acamprosate, are not contraindicated in schizophrenia, and disulfiram  
25 also seems to be well tolerated, although it has been suggested that symptoms  
26 of psychosis and liver toxicity should be closely monitored (Green *et al.*, 2008).

27  
28 Treatment Improvement Protocol series 43 covers problems with treatments  
29 for opioid dependence, such as methadone and buprenorphine. These drugs  
30 can precipitate withdrawal in people also taking drugs to treat HIV infection,  
31 such as nelfinavir, efavirenz, and nevirapine. There is a similar problem with  
32 these opioid treatments and carbamazepine, phenytoin and phenobarbital.

33  
34 With antidepressants, some SSRIs which inhibit the isoenzymes that  
35 metabolise methadone (particularly, CYP3A4, CYP1A and CYP2D6) could  
36 lead to increased serum methadone levels. Fluvoxamine is the most likely to  
37 cause excessive serum methadone levels due to inhibition of CYP1A2 and has  
38 been implicated in over-sedation and respiratory depression when combined  
39 with methadone. Also, there is some indication that methadone increases  
40 serum levels of tricyclic antidepressants, so lower doses may be needed.  
41 Rifampin, carbamazepine, phenobarbital and some HIV infection medications

1 may induce liver enzymes that alter the transformation of methadone. So  
2 clinicians may need to adjust the dose of methadone accordingly.

3

4 Treatment Improvement Protocol series 45 warns that benzodiazepines,  
5 which are known to be addictive, are particularly so in those already addicted  
6 to other substances. Because of their reduced side effect profile and lower risk  
7 of dangerous drug interactions, SSRIs may be considered as the  
8 antidepressants of choice for those with addiction and coexisting psychiatric  
9 conditions. However, the potential for different SSRIs to cause drug  
10 interactions should be considered in individual cases.

### 11 **8.2.9 Clinical evidence summary (pharmacological** 12 **interventions)**

13 There is limited evidence from well conducted RCTs for the relative  
14 effectiveness of pharmacological treatments for people with psychosis and  
15 coexisting substance misuse, either of treatments for psychosis symptoms or  
16 of treatments aimed at improving substance misuse. There is also little data  
17 on interactions between drugs given as medication and drugs of abuse. See  
18 Table 33 for a summary for each medication.

**Table 33: Relevant interventions included in current NICE guidelines and summary of evidence of effectiveness**

Intervention type/use	Name	Recommended in existing NICE guideline? <sup>1</sup>	Evidence found from existing reviews and new RCTs	Notes from Summary of Product Characteristics
<b>MEDICATION</b>				
Alcohol dependence	Acamprosate calcium	Alcohol <sup>2</sup> : Yes <sup>3</sup>	No evidence, but no known contraindication in those with schizophrenia.	
Alcohol deterrent compounds	Disulfiram	Alcohol <sup>2</sup> : Yes <sup>3</sup>	At best, there is preliminary evidence of effectiveness in people with coexisting schizophrenia and alcohol dependence, but some reviewers consider that using this medication risks inducing psychosis.	Chlordiazepoxide and diazepam toxic effect may be enhanced. Very rare reports of potentiation of organic brain syndrome and choreoathetosis with pimozide. The intensity of the Disulfiram-alcohol reaction may be increased by amitriptyline and chlorpromazine and decreased by diazepam.  Avoid lithim liquid (contains 5% ethanol).
Alpha-adrenergic agonists	Clonidine	DMD: Not routinely	No evidence.	Anntipsychotics and tricyclic antidepressants may provoke orthostatic hypotension. CNS depressants may be potentiated and cause excessive drowsiness. Increased risk of rebound hypertension if clonidine is withdrawn in service users taking tricyclic antidepressants.

Alpha-adrenergic agonists	Lofexidine	DMD: Yes <sup>3</sup>	No evidence.	Efficacy may be reduced by tricyclic antidepressants. Concomitant use of drugs which prolong the QT interval should be avoided.
Antiepileptic drugs	Phenytoin	Alcohol <sup>2</sup> : No	No evidence.	Class warning for anticonvulsants. A small increased risk of suicidal ideation and behaviour reported. Potential for drug interactions is complex and includes a range of psychotropic drugs
Antiepileptic drugs	Topiramate	Alcohol <sup>2</sup> : No	No evidence.	SPC Class warning for anticonvulsants. A small increased risk of suicidal ideation and behaviour reported. Inhibits the enzyme CYP 2C19.
Antimanic drugs	Lithium	Bipolar: Yes	There is limited evidence of effectiveness in reducing substance misuse in those with bipolar disorder; of combined use with valproate in reducing coexisting alcohol use.	Avoid lithium liquid with metronidazole or in service users with alcohol misuse.
Antimanic drugs	Valproic acid	Bipolar: Yes	Case study evidence of benefit in coexisting schizophrenia and alcohol dependence; recommended by one author for mania but evidence is unclear; evidence of usefulness in reducing alcohol consumption.	Class warning for anticonvulsants. A small increased risk of suicidal ideation and behaviour reported.  Combination with olanzapine may significantly increase the risk of certain olanzapine associated adverse events.
Antimanic drugs/ Anxiolytics	Benzodiazepine (for example, diazepam,	Bipolar: Yes <sup>3</sup>  Alcohol <sup>2</sup> : Yes	No evidence, but potentially addictive.	

FINAL CONSULTATION

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	lorazepam, chlordiazepoxide):			
Antimanic drugs/ Hypnotics	Clomethiazole (Chlormethiazole)	Alcohol <sup>2</sup> : No	No evidence, but potentially addictive.	Fatal cardiorespiratory collapse reported when combined with other CNS depressant drugs.
Antimanic/ Control of epilepsy	Carbamazepine	Bipolar: Not routinely	Evidence that it may reduce substance misuse in bipolar disorder, and control mania and depressive symptoms.	<p>Class warning for anticonvulsants. A small increased risk of suicidal ideation and behaviour reported.</p> <p>Avoid with MAOI's and individuals of Han Chinese and Thai origin with positive HLA-B*1502 allele, due to increased risk of developing carbamazepine-associated Stevens-Johnson syndrome.</p> <p>Principal iso enzyme responsible for metabolism is CYP 3A4, therefore use caution with inhibitors or inducers of this isoenzyme.</p> <p>Levels of carbamazepine and its principal active metabolite may be increased by concomitant use of a range of drugs including fluoxetine, fluvoxamine, paroxetine, trazodone and olanzapine, quetiapine and valproic acid.</p> <p>Carbamazepine is also a potent inducer of</p>

				CYP3A4 and may therefore reduce the plasma concentrations of concomitant pharmacotherapy which is metabolized by CYP3A4.
Antipsychotic drugs	For example: Clozapine Haloperidol Olanzapine Risperidone	Bipolar: Yes  Schizophrenia (update): Yes	Inconsistent findings on substance misuse outcomes. More frequent reports suggest clozapine may be of benefit.	Principal isoenzyme responsible for metabolism is CYP1A2.  Clozapine is contraindicated in alcoholic and other toxic psychoses, drug intoxication and comatose conditions. Principal iso enzyme responsible for metabolism is CYP 1A2 . Sudden smoking cessation may significantly increase clozapine plasma levels, concomitant benzodiazepine use may increase risk of circulatory collapse.  Consult the SPC of individual agents for information about other drugs.
Opioid agonists & partial agonists	Buprenorphine	DMD: Yes	No evidence.	Principal isoenzyme responsible for metabolism is CYP3A4.
Opioid agonists & partial agonists	Methadone	DMD: Yes	No evidence. Some suggestion of interactions with other medications.	Principal isoenzyme responsible for metabolism is CYP3A4.  Concomitant use with MAOI's and drugs which prolong the QT interval should be avoided
Opioid	Nalmefene	Alcohol?: No	No evidence.	No UK licence.

antagonists		DMD: No		
Opioid antagonists	Naltrexone	Alcohol <sup>2</sup> : Yes <sup>3</sup> DMD: Yes <sup>3</sup>	Some evidence of effectiveness in schizophrenia with coexisting alcohol dependence.	
Serotogenic agents	Ondansetron	Alcohol <sup>2</sup> : No	No evidence.	Metabolised by multiple hepatic isoenzymes: CYP3A4, CYP2D6 and CYP1A2. Therefore enzyme inhibition or reduced activity of one enzyme is normally compensated by other enzymes and should result in little or no significant change in overall ondansetron clearance or dose requirement.
Serotogenic agents	SSRIs	Alcohol <sup>2</sup> : Not routinely for alcohol misuse Depression: Yes <sup>3</sup>	No evidence in psychosis. Some suggestion of interactions with methadone, leading to increased serum methadone levels (SSRIs).	Individual SSRIs vary in their propensity to affect Cytochrome p450 isoenzymes.  Consult current SPC for details.
Skeletal muscle relaxants	Baclofen	Alcohol <sup>2</sup> : No	No evidence.	Tricyclic antidepressants may potentiate effects, resulting in pronounced muscular hypotonia. Concomitant use of CNS drugs may lead to increased sedation.
<b>PHYSICAL AND COMPLEMENTARY INTERVENTIONS</b>				
Physical	Acupuncture	DMD: No	No evidence.	
Complementary	Mindfulness meditation	Alcohol <sup>1</sup> : No	No evidence.	

*Note.* DMD = Drug misuse: opioid detoxification; DMP = Drug misuse: psychosocial interventions.

<sup>1</sup> Available from [www.nice.org.uk](http://www.nice.org.uk).

<sup>2</sup> Management of alcohol dependence guideline.

<sup>3</sup> For specific groups and/or in certain circumstances (see relevant guideline for further information).

1 **8.2.10 Health economic evidence (pharmacological/ physical**  
2 **interventions)**

3 No studies assessing the cost-effectiveness of pharmacological/physical  
4 interventions for people with psychosis and coexisting substance misuse were  
5 identified by the systematic search of the economic literature undertaken for  
6 this guideline. Details on the methods used for the systematic search of the  
7 economic literature are described in Appendix 9.

8 **8.2.11 From evidence to recommendations (pharmacological/**  
9 **physical interventions)**

10 There is little robust evidence to guide the use of specific pharmacological  
11 treatments for people with psychosis and coexisting substance misuse in the  
12 UK. On the basis of the evidence reviewed, it is not possible to identify  
13 specific drugs which should be considered as agents of first choice.

14  
15 The GDG felt that the use of depot formulations may be expected to increase  
16 the opportunity to identify episodes of non-adherence to prescribed  
17 treatment. Whilst this may be an important consideration in individual cases  
18 there is, overall, insufficient evidence to recommend depot preparations as  
19 routine first line treatment.

20  
21 Clozapine is frequently cited as having a particular role in this population,  
22 although there is no RCT evidence to support this view. In addition, its use  
23 may increase the risk of adverse effects, and due to the possibility of a  
24 syncopal episode, the GDG felt that particular care should be exercised where  
25 the drug of misuse is cocaine.

26  
27 In general though, as no good quality evidence was found relating to the  
28 modification of interventions recommended for people with a single  
29 diagnosis, the GDG concluded that people with psychosis and coexisting  
30 substance misuse should be offered the same range of evidence-based  
31 interventions recommended for people with a single diagnosis. In addition,  
32 the GDG felt it important to make a number of recommendations for good  
33 practice concerning the initiation and use of medication.

34  
35 There was no evidence that addressed the sub-question regarding subgroups  
36 of people (see section 8.2.2 for further information about the sub-question). In  
37 addition, the GDG noted that valuable information about the potential  
38 benefits of pharmacological and psychosocial interventions for people with  
39 psychosis and substance misuse could be obtained from trials of treatments  
40 for people with either of these two different types of problems. However, to

1 date, most trials conducted among people with psychosis have excluded  
2 those who have coexisting substance misuse and nearly all trials among  
3 people with substance misuse have excluded those with coexisting psychosis.  
4 In some instances, it may be necessary to exclude people with coexisting  
5 problems from future studies. However, very often, this important and  
6 prevalent group of patients have been excluded from intervention trials with  
7 no clear reason being offered. Therefore, future research should not routinely  
8 exclude people with psychosis and coexisting substance misuse.  
9

## 10 **8.3 CLINICAL PRACTICE RECOMMENDATIONS**

### 11 **8.3.1 Recommendations (pharmacological/ physical** 12 **interventions)**

#### 13 *Secondary care mental health services*

#### 14 **Treatment**

15 **8.3.1.1** Before starting treatment for adults and young people with psychosis  
16 and coexisting substance misuse, review:

- 17 • the diagnosis of psychosis and of the coexisting substance  
18 misuse, especially if either diagnosis has been made during  
19 a crisis or emergency presentation **and**
- 20 • the effectiveness of previous and current treatments and the  
21 person's tolerance of them; discontinue ineffective  
22 treatments.<sup>19</sup>

---

<sup>19</sup> This recommendation also appears in section 7.3.1 where the psychological/psychosocial data is presented.

- 1 **8.3.1.2** Ensure that adults and young people with psychosis and coexisting  
2 substance misuse are offered evidence-based treatments for both  
3 conditions (see 8.3.1.3 and 8.3.1.4).<sup>20</sup>
- 4 **8.3.1.3** For the treatment of psychosis, see ‘Bipolar disorder: the management  
5 of bipolar disorder in adults, children and adolescents, in primary  
6 and secondary care’ (NICE clinical guideline 38) or the guideline on  
7 schizophrenia (NICE clinical guideline 82).<sup>21</sup>
- 8 **8.3.1.4** For the treatment of substance misuse, see:
- 9                   • ‘Alcohol-use disorders: diagnosis and clinical management  
10 of alcohol-related physical complications’ and the guideline  
11 on alcohol dependence and harmful alcohol use (NICE  
12 clinical guidelines 100 and CGXX) **and/or**
- 13                   • ‘Drug misuse: psychosocial interventions’ and the guideline  
14 on opioid detoxification (NICE clinical guidelines 51 and  
15 52).<sup>22</sup>
- 16 **8.3.1.5** Use antipsychotics according to the guideline on schizophrenia (NICE  
17 clinical guideline 82) or bipolar disorder (NICE clinical guideline 38)  
18 because there is no evidence for any differential benefit for one  
19 antipsychotic over another for people with psychosis and coexisting  
20 substance misuse.
- 21 **8.3.1.6** Use depot/long-acting injectable antipsychotics according to the  
22 guideline on schizophrenia (NICE clinical guideline 82) in managing  
23 covert non-adherence with treatment for psychosis and not as a  
24 specific treatment for psychosis and coexisting substance misuse.
- 25 **8.3.1.7** When prescribing medication for adults and young people with  
26 psychosis and coexisting substance misuse:
- 27                   • take into account the level and type of substance misuse,  
28 especially of alcohol, as this may alter the metabolism of  
29 prescribed medication, decrease its effectiveness and/or  
30 increase the risk of side effects
- 31                   • warn the person about potential interactions between  
32 substances of misuse and prescribed medication

---

<sup>20</sup> This recommendation also appears in section 7.3.1 where the psychological/psychosocial data is presented.

<sup>21</sup> This recommendation also appears in section 7.3.1 where the psychological/psychosocial data is presented.

<sup>22</sup> This recommendation also appears in section 7.3.1 where the psychological/psychosocial data is presented.

- 1                           • discuss the problems and potential dangers of using non-  
2                           prescribed substances and alcohol to counteract the effects  
3                           or side effects of prescribed medication.

4   **8.3.2 Research recommendations (pharmacological**  
5           **interventions)**

6   **8.3.2.1** ?Are interventions for psychosis or substance misuse clinically and cost  
7           effective when compared with standard care for people with  
8           psychosis and coexisting substance misuse?<sup>23</sup>

9   **8.3.2.2** Is clozapine clinically and cost effective when compared with other  
10          pharmacological interventions for people with psychosis and  
11          coexisting substance misuse?

12  
13  
14  
15  
16

---

<sup>23</sup> This recommendation also appears in section 7.3.2 where the psychological/psychosocial data is presented.

1

## 2 **9 YOUNG PEOPLE WITH** 3 **PSYCHOSIS AND COEXISTING** 4 **SUBSTANCE MISUSE**

### 5 **9.1 INTRODUCTION**

6 As described in Chapter 5, there is a paucity of evidence relating to  
7 assessment and care pathways specifically for people with psychosis and  
8 coexisting substance misuse. For young people, this is true of all review  
9 questions, therefore, the GDG developed through expert consensus, specific  
10 recommendations for young people (for further information about the  
11 methods used in this chapter, please see Chapter 3, section 3.5.6; for a list of  
12 all review questions see Appendix 6). The care pathway is summarised in  
13 Figure 1. As with Chapter 5, the text and are designed to be illustrative and  
14 offer some broad principles and direction, rather than to be prescriptive. They  
15 are sufficiently broad to take into account.

16

17 Adolescence is a period of major developmental transitions - physically,  
18 psychologically and socially. During this period young people experience  
19 emotional distress, frequent interpersonal disruptions and challenges in  
20 establishing a sense of identity. These factors can act as both stressors for  
21 those vulnerable to a psychotic illness and as difficulties that can lead to  
22 substance misuse as a form of escape or self-treatment.

23

24 Little research has been carried out on the specific factors that lead young  
25 people to be vulnerable to both substance misuse and psychosis. Furthermore,  
26 little is known about the effectiveness of interventions specific to this age  
27 group. This chapter, therefore, covers what is known about prevalence,  
28 outcomes and service configuration for young people. In the absence of more  
29 specific evidence, the principles of intervention will be drawn from and  
30 adapted from the adult literature.

31

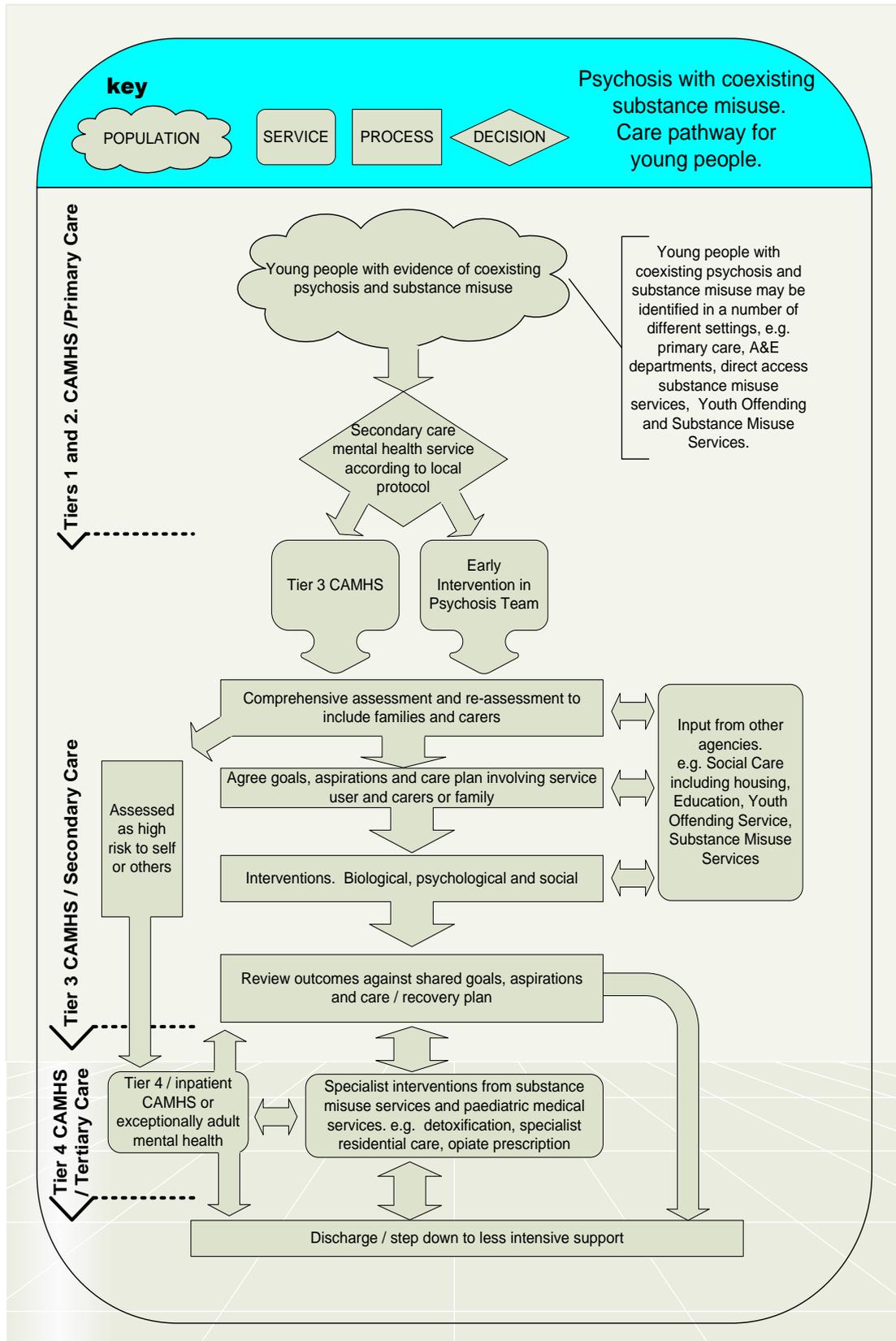
32 This guideline uses the term 'young people' to refer to people aged between  
33 their 14<sup>th</sup> and 18<sup>th</sup> birthdays, as people of this age generally prefer this  
34 descriptor to the term 'adolescent'.

35

1 **9.2 PREVALENCE**

2 It is not simple to identify the prevalence of substance misuse and psychosis  
3 in young people. Studies which explore the age range might include a  
4 discussion about each of the disorders, but rarely combine them. Studies  
5 which do investigate combined disorders usually do not focus on the under  
6 18 year olds.  
7

- 1 **Figure 4: Care pathway for young people with psychosis and coexisting**
- 2 **substance misuse.**



1

2

1 A systematic review of coexisting substance use in people with psychosis  
2 carried out by Carra and Johnson (2009) pointed to wide variations in  
3 prevalence rates. Most recent UK studies reported rates of between 20 to 37%  
4 in mental health settings, and 6 to 15% in addiction settings (Carra & Johnson,  
5 2009). Inpatient, crisis and forensic settings are, not surprisingly, higher, that  
6 is, 38 to 50% (Carra & Johnson, 2009). People from inner cities and some  
7 ethnic groups are over represented (Carra & Johnson, 2009). It should be  
8 emphasised that there are varying age ranges in these studies and few  
9 specifically focused on young people.

### 10 **9.2.1 General practice**

11 A study undertaken from 1993 to 1998 estimated that there were at least  
12 195,000 comorbid service users and 3.5 million GP consultations involving  
13 comorbid service users of all ages in England and Wales (Frisher *et al.*, 2004).  
14 An unanticipated finding was that each year 80-90% of comorbid service  
15 users were newly diagnosed, although existing service users may continue to  
16 receive treatment. Thus, there is a significant problem in terms of primary  
17 care workload. The number of people newly developing comorbidity in  
18 primary care increased year-on-year. The impact on health services is far in  
19 excess of that for mono-morbid service users; comorbid individuals have an  
20 extra consultation frequency for all problems, estimated as an excess of  
21 1,115,751 consultations in England and Wales in 1998.

22  
23 During the six year study period, the annual comorbidity rate increased by  
24 62%, but rates of comorbid schizophrenia, paranoia and psychoses increased  
25 by 128%, 144% and 147%, respectively (Frisher *et al.*, 2004). In this study, the  
26 level of comorbidity increased at a higher rate among younger service users,  
27 which indicates that comorbidity may increase, perhaps at a faster rate than  
28 observed in the study period, in future years. All comorbid diagnoses –  
29 including schizophrenia and psychosis – peaked at ages 16-24 or 25-34. In  
30 1998, it was estimated that there were about 20,000 comorbid cases between  
31 ages 16 to 34 (7773 in age 16 to 24 and 12949 in 25 to 34 age range) in primary  
32 care.

33  
34 The data reported by Frisher and colleagues indicate that substance abuse  
35 may be precipitating more serious forms of comorbidity, although it is by no  
36 means clear that this is the case. For example, nearly all diagnoses of  
37 comorbid schizophrenia precede substance abuse. In this study (Frisher *et al.*,  
38 2004), the majority (54%) of service users had a psychiatric diagnosis first, and  
39 half become comorbid within 6 months of the first diagnosis.

40

1 The findings on transition from mono to comorbidity have major implications  
2 for understanding and preventing comorbidity. Perhaps individuals with  
3 comorbidity may be qualitatively different in the form of their mono-  
4 morbidity than those who remain mono-morbid. Early development of  
5 comorbidity suggests that there may be characteristics already present at the  
6 mono-morbid stage which may predict the likelihood of developing  
7 comorbidity. Identifying such characteristics in future research might  
8 contribute to the early management or prevention of comorbidity in primary  
9 care.

## 10 **9.2.2 Community substance misuse and mental health services**

11 Weaver and colleagues (2003) conducted a multicentre study that derived  
12 estimates of psychosis and coexisting substance misuse (of whom, 76% were  
13 diagnosed with schizophrenia), in the age range of 16 to 30 years old. They  
14 found that one third of their sample was misusing substances. Although the  
15 age range looked at in this study exceeds the range considered for young  
16 people, it is helpful in providing a figure on substance misuse in the  
17 community.

## 18 **9.2.3 First-episode psychosis**

19 Donoghue and colleagues (2009) utilised data from two epidemiological  
20 studies of first-episode psychosis (the Schizophrenia in Nottingham study  
21 and the Aetiology and Ethnicity of Schizophrenia and Other Psychoses  
22 study), demonstrating that for 16 to 29 year olds, there was a significant  
23 increase from 14.9% to 30.1% in all substance use disorders between 1992-1994  
24 and 1997-1999 (Donoghue *et al.*, 2009). Similarly, for cannabis-specific  
25 substance use disorder, there was a significant increase from 3.2% to 10.6%.  
26 These increases were seen in both males and females.

## 27 **9.3 IMPACT OF SUBSTANCE MISUSE ON** 28 **OUTCOME IN PSYCHOSIS**

29 In a group of first episode service users treated with psychological therapy,  
30 33% of the under 21 year olds had self reported substance misuse (Haddock *et*  
31 *al.*, 2006). Of relevance is the finding that young people may have differing  
32 needs with regard to engagement. Counselling appeared to be more beneficial  
33 for the younger age group.

34  
35 An Australian study (Wade *et al.*, 2006), in a 15-30 year old age group (mean  
36 age 21.6 years), reported that substance misuse (53% at follow up) was an  
37 independent risk factor for problematic recovery in first-episode psychosis  
38 (for example, increased risk of admission, relapse of positive symptoms and

1 shorter time to relapse). However, substance misuse was not associated with  
2 longer time to remission.

3  
4 Hides and colleagues (2006) has pointed to a bidirectional relationship  
5 between substance misuse and cannabis relapse in that a higher frequency of  
6 cannabis use was predictive of psychotic relapse (if medication adherence,  
7 other substance use and duration of untreated psychosis were controlled for),  
8 while an increase in psychotic symptoms was predictive of relapse to  
9 cannabis use. In this study, only 15% of service users had not used any illicit  
10 substance in the previous 12 months.

## 11 **9.4 ASSESSMENT AND DIAGNOSIS**

12 Many aspects of the assessment and diagnosis of young people with  
13 psychosis and coexisting substance misuse will be the same or similar as for  
14 adults. This is covered in detail in Chapter 5.

15  
16 As is the case for adults, healthcare professionals in all settings should  
17 routinely ask young people with known or suspected psychosis about their  
18 use of substances. This may include questions about type and method of  
19 administration, quantities and frequency. It is important for healthcare  
20 professionals in all settings to routinely assess young people with known or  
21 suspected substance misuse for possible psychosis.

22  
23 For young people with psychosis and coexisting substance misuse presenting  
24 to mental health services, a comprehensive assessment of a young person's  
25 psychosis and substance misuse is crucial. This includes an assessment of  
26 psychiatric, psychological and physical health, home and family environment,  
27 educational or employment status, medication, risk to self and others,  
28 relationships and social networks, forensic and criminal justice history,  
29 strengths, and aspirations. Assessing the relationship between substance use,  
30 emotional state and reasons for substance use is also important. In addition,  
31 gaining corroborative evidence where possible is helpful in order to assess the  
32 impact of substance misuse on mental state and behaviour.

33  
34 The assessment of young people may take time and involve multiple sessions  
35 due to difficulty with concentration, ambivalence, lack of clarity about the  
36 purpose of the assessment(s), and the need to gradually gain trust and  
37 confidence in the practitioners and service. There are three crucial goals of an  
38 assessment. The first is to conduct the assessment in such a manner that  
39 fosters and promotes continuing engagement. The second is to ensure safety  
40 of the young person, and the third is to determine which substance(s) the  
41 young person is dependent on in order to determine whether administration

1 of a pharmacological agent – possibly for detoxification – is appropriate. It is  
2 important to note that even if the young person is not dependent on a  
3 substance, serious harm may result from drug misuse.

4  
5 The comprehensive assessment of a young person presenting with psychosis  
6 and coexisting substance misuse is similar to what is described for adults in  
7 Chapter 5. The issues brought up for adults however, apply even more  
8 strongly for young people, as they are more complex to engage, are more  
9 vulnerable, and can suffer from serious problems as a result of substance  
10 misuse, without having substance dependence. Additional differences  
11 between adults and young people relate to service delivery, as services for  
12 young people are usually provided separately from those for adults.

## 13 **9.5 SERVICE CONFIGURATION AND CARE** 14 **PATHWAYS**

### 15 **9.5.1 Configuration of CAMHS Services**

16 Interventions for young people with psychosis and coexisting substance  
17 misuse may be provided by a range of agencies and services within each  
18 agency. Agencies will include Children’s Services, which may be involved  
19 around social care/housing issues, education or safeguarding. Youth  
20 Offending Services may be involved. However, once a diagnosis of psychosis  
21 with substance misuse has been made, mental health services will usually be  
22 provided by specialist CAMHS or Early Intervention in Psychosis Services  
23 (EIS). Specialist substance misuse interventions for young people may be  
24 available from within core mental health services or from specialist substance  
25 misuse services.

26  
27 In order to recognise the different levels of interventions for many child  
28 mental health problems, CAMHS has been organised into four main levels, or  
29 tiers, of delivery (Department of Health, 2004; Health Advisory Service, 1995)  
30 (see Text Box 1).

### 31 **9.5.2 CAMH Services**

#### 32 *Tier 1 CAMHS*

33 Professionals at Tier 1 are most likely to encounter young people with  
34 psychosis and coexisting substance misuse when a change in their behaviour  
35 is noticed. This could be unusual behaviour or otherwise out-of-character  
36 behaviour, a decline in academic performance or increasing social isolation.  
37 Tier 1 professionals are unlikely to be involved in diagnosing psychosis, but  
38 may become aware of substance misuse difficulties. They could also become

1 involved in providing for the young person's physical healthcare, social and  
2 educational needs when the young persons mental health needs are being  
3 met. Awareness of psychosis and substance misuse in young people may  
4 prevent inappropriate dismissal of the difficulties presented by the young  
5 person and encourage them to refer on to appropriate services. For Tier 1  
6 professionals to be able to fulfil these roles for young people with psychosis  
7 and coexisting substance misuse they will need appropriate training. Training  
8 programmes for Tier 1 staff may require modification to cover psychosis with  
9 substance misuse or behaviours suggestive of the diagnosis. This training  
10 may be most effectively targeted at services that have young people with  
11 higher rates of mental health concerns for example Key Stage 4 Pupil Referral  
12 Units. Following appropriate training Tier 1 professionals may be involved in  
13 the sensitive detection of psychosis and substance misuse difficulties. When  
14 identified such concerns should lead to referral to or consultation with Tier 2  
15 professionals.

### 16 *Tier 2 CAMHS*

17 Tier 2 professionals provide consultation and training to Tier 1 professionals  
18 in regard to all mental health problems. Tier 2 professionals therefore require  
19 an awareness of the problems of young people with psychosis and coexisting  
20 substance misuse and competence to detect psychotic symptoms in young  
21 people or the early features of psychosis. If a diagnosis of psychosis or early  
22 features of psychosis is suspected, a referral to Tier 3 CAMHS or Early  
23 intervention services (EIS) teams can be made according to local protocols.

### 24 *Tier 3 CAMHS*

25 Tier 3 services can provide a comprehensive assessment of the young person  
26 with psychosis and coexisting substance misuse. When a diagnosis of  
27 psychosis is made, it is important for Tier 3 professionals to consider the  
28 possibility of substance misuse.

29  
30

1

<b>Text Box 1: Child and adolescent mental health services (CAMHS) tiers structure</b>	
<b>Tier 1</b>	<ul style="list-style-type: none"> <li>• Provide primary or direct contact with young people, primarily for reasons other than mental health, including primary care/general practice, counselling and psychotherapy, general paediatrics, social services, health visitors and schools</li> <li>• First point of contact with the child/family with mental health problems</li> <li>• Draw on specialist CAMHS personnel who can consult and advise them about working with children and young people in their care who either have, or are at risk of developing, a mental health problem</li> </ul>
<b>Tier 2</b>	<ul style="list-style-type: none"> <li>• Specialist CAMHS professionals working in a community-based setting alongside Tier 1 workers, working in primary care, schools and other relevant community settings such as social services</li> <li>• Work as a part of a team, with Tier 1 staff, built around the individual child</li> <li>• Able to provide fairly rapid assessment and treatment to children within Tier 1 settings, as well as consultation/support to Tier 1 workers</li> <li>• Able to help identify those children needing referral to more specialist services</li> <li>• Ideally organised into multidisciplinary teams, with good links to Tier 3 services, thereby facilitating a more seamless transition across tiers</li> <li>• Sometimes, Tier 2 services are provided by the voluntary sector (for example, some but not all adolescent counselling and psychotherapy services)</li> </ul>
<b>Tier 3</b>	<ul style="list-style-type: none"> <li>• Comprise multidisciplinary teams of specialist CAMHS professionals working in (secondary care) specialist CAMHS facilities (for example, Child and Family Consultation Services or Hospital Liaison Teams)</li> <li>• The National Service Framework for Children's Services states that all PCT / LHB areas should have at least one (or access to one) comprehensive Tier 3 multidisciplinary CAMHS team providing specialist co-ordinated assessments and interventions, and offering the full range of appropriate psychological and pharmacological treatments</li> <li>• Offer outreach services to those young people who are housebound or otherwise unable to access Tier 3 services based in secondary care facilities, or to work in conjunction with outpatient treatment plans (for example, monitoring of medication). Emergency services, with 24-hour availability should also be in place in all localities</li> <li>• Provide consultation and training to Tier 1 workers and refer when necessary to Tier 4 services</li> </ul>
<b>Tier 4</b>	<ul style="list-style-type: none"> <li>• Highly specialised tertiary CAMHS that provide multidisciplinary services for very severe mental health problems, or for those who need very intensive treatment or supervision. These services vary in how they are organised.</li> <li>• Includes highly specialist outpatient treatment, crisis intervention and intensive home-based therapies.</li> <li>• Referrals to Tier 4 services usually come from Tier 3 CAMHS professionals, and service users are usually discharged back to Tier 3 services or outreach services after the Tier 4 intervention</li> </ul>

2

1 When a diagnosis of psychosis and coexisting substance misuse has been  
2 made, priority should be given to both treatment of the psychosis and  
3 substance misuse. Constant review of risk is of key importance, and if the  
4 young person presents with a high risk to themselves or others due to their  
5 psychosis, then inpatient admission is important to consider.

6  
7 All the mainstays of treatment, including prescribing medication, monitoring  
8 mental state and providing psychosocial intervention can be offered in Tier 3  
9 CAMHS, by EIS teams or by a collaboration between the two.

10  
11 Given that most young people with psychosis and coexisting substance  
12 misuse live with their families, with foster parents, or in social services  
13 residential placements, involving carers in treatment is helpful. Carers can be  
14 involved in relapse prevention work as well working with professionals in  
15 supporting the young person with their substance misuse. Supporting  
16 parents, including family therapy, should be offered to all families and  
17 include a focus on high levels of criticism and intrusiveness (expressed  
18 emotion) when identified.

19  
20 As many young people with psychosis and coexisting substance misuse  
21 require a multi-agency response, clarity about the responsibilities of each  
22 agency facilitates the delivery of care. As well as their mental health and  
23 substance misuse needs, young people with psychosis and coexisting  
24 substance misuse will often have housing, employment or educational needs.  
25 Agencies must strive to collaborate to provide coordinated care. Different  
26 thresholds for entry into services can compromise this objective. For example,  
27 Tier 3 professionals may have concerns about a young person's social care  
28 that may not meet social service thresholds for intervention. This can reduce  
29 the effectiveness of therapeutic interventions as Tier 3 staff become involved  
30 in trying to coordinate or meet social care needs. Likewise social services may  
31 find accessing specialist therapy services for some of the young people they  
32 care for difficult because, for example, despite on-going substance misuse,  
33 Tier 3 staff may consider that the young person's mental health difficulties are  
34 in remission and therefore sub-threshold for active involvement. Failure to  
35 engage at all with the young person in these circumstances may prevent the  
36 success of social services interventions to improve the young person's social  
37 care and increase likelihood of relapse. Professionals need to work flexibly  
38 and creatively around these tensions over service thresholds. Respecting the  
39 validity of the principles leading to the development of thresholds whilst  
40 trying to meet the needs of the young person is required in these  
41 circumstances.

42

1 It is important for Tier 3 teams to develop sub-teams of professionals with  
2 expertise in the management of young people with psychosis and coexisting  
3 substance misuse either separately or in collaboration with EIS teams. One  
4 model of collaboration widely adopted is for CAMHS to provide psychiatric  
5 input whilst EIS provide care co-ordination and psychosocial interventions. In  
6 some areas, stand alone CAMHS psychosis services have been set up. Tier 3  
7 CAMHS professionals must also have the capacity to provide consultation  
8 and training to Tier 2 staff.

9  
10 Healthcare professionals working in Tier 3 can also follow the  
11 recommendation for adults in other chapters.

#### 12 *Tier 4 CAMHS*

13 For young people with psychosis and coexisting substance misuse, Tier 4  
14 CAMH services principally comprise inpatient services. There is usually a  
15 limited role for other Tier 4 CAMH services such as specialist outpatient  
16 services and home-based treatment teams, as most non-bed based treatments  
17 can be picked up by other services such as Tier 3 CAMHS or EIS teams.

18  
19 *Inpatient services* – Admission to an inpatient unit will usually be indicated  
20 due to the level of risk identified in managing the young person in the  
21 community. This can often present in an acute crisis. Admissions for the  
22 management of acute risk should be clearly linked to an acute exacerbation of  
23 risk, time-limited, and with clear goals in mind. Such admissions may also be  
24 required when risk is high and the motivation of the service user to  
25 collaborate in community treatment is very low or non-existent. The aim of  
26 such admissions is usually to ensure that the service user is ‘just community  
27 ready’. Transfer back to the community is clearly facilitated in circumstances  
28 where the young person is effectively engaged in a structured outpatient  
29 programme.

30  
31 Other factors warranting consideration for admission by a Tier 4 team for  
32 treatment of psychosis and coexisting substance misuse include other Axis I  
33 difficulties combined with a significant deterioration in functioning and a  
34 reduced capacity of either the family or community team to manage the  
35 young person.

36  
37 Exceptionally, if a young person’s needs are thought to be best met by and  
38 adult ward and they choose this (for example if they are almost 18 years and  
39 adult services are much closer to home), then it is acceptable for them to be  
40 admitted to an adult mental health ward. It is also acceptable for a young  
41 person aged 16 or 17 years to spend a short time on an adult ward if an age  
42 appropriate bed is not available. In both these examples safeguarding

1 measures need to be in place whilst the young person is on the adult ward. It  
2 is never acceptable for a young person under the age of 16 years to be  
3 admitted to an adult ward (See MHA 1983 revision 2007, section 31 and MHA  
4 Code of Practice [Department of Health, 2008]).

5  
6 *Specialist home-based treatment teams* for young people are in the early stages of  
7 development in the UK and consequently their place in the treatment of  
8 psychosis and coexisting substance misuse has yet to be established. Like  
9 inpatient services, existing teams frequently manage acute risk and attempt to  
10 address chronic risk and/or low functioning service users.

11  
12 Services are likely to take different forms dependent on their focus on acute or  
13 chronic issues. When focused on acute risk, services usually combine  
14 characteristics of assertive outreach and crisis intervention with intensive case  
15 management. These services have proved effective both when Tier 3  
16 treatment has been disrupted and as a mechanism for organising an effective  
17 outpatient intervention plan. Typically services have a capacity for rapid and  
18 intensive engagement lasting no more than a few weeks, followed by service  
19 user/family centred intensive case management.

20  
21 Services focused on chronic risk and/or low functioning are characterised by  
22 a stronger psychotherapy focus, a longer duration of treatment and an active  
23 engagement phase pre-treatment. These services have also been used as step-  
24 down from inpatient, when inpatient stays have become ineffective or for  
25 community rehabilitation. This type of intervention might be considered  
26 when parenting has become distorted by the service user's presentation and  
27 family relationships are undermining individually focused treatment plans.

28  
29 In most cases, psychoeducational work with parents is required prior to  
30 implementing more intensive interventions that may often be experienced as  
31 intrusive. These forms of home-based treatment are best avoided where there  
32 are longstanding concerns about parental capacity.

33  
34 Home-based treatment services, regardless of whether they focus on the  
35 treatment of acute or chronic issues, share a number of characteristics: they  
36 require experienced staff with expertise in psychosis and coexisting substance  
37 misuse and a team structure that allows a high level of supervision and the  
38 effective management of risk in the community; each is likely to offer time-  
39 limited treatment but of different durations; and each is likely to balance limit  
40 setting with developing autonomy. Services need to effectively differentiate  
41 young person, parents, family, and wider system interventions and to focus  
42 primarily on the management of risk and the promotion of functioning.

## 1 **9.6 EARLY INTERVENTION IN PSYCHOSIS** 2 **SERVICES**

3 Early intervention services (EIS) are assertive community-based  
4 multidisciplinary teams that provide care for people aged between 14 and 35  
5 years with a first presentation of psychotic symptoms during the first 3 years  
6 of psychotic illness (Department of Health, 2001) and are primarily concerned  
7 with the early identification and treatment of the early phase of psychotic  
8 illness. For young people (aged 14 to 18), EIS often work according to locally  
9 agreed protocols with Tier 3 and 4 CAMHs.

10  
11 Often, the initial focus of the EIS is on engagement in order to develop a  
12 shared, individualised recovery focussed treatment plan that incorporates a  
13 range of interventions including antipsychotic drugs, CBT, family  
14 intervention, vocational activity and reduction of substance misuse. As  
15 substance use and misuse is so common in people presenting with a first  
16 episode of psychotic illness, there are sound clinical reasons why EIS staff  
17 would consider the possibility of substance misuse in a young person  
18 presenting with psychotic symptoms, and if a diagnosis of psychosis and  
19 coexisting substance misuse is made, ensure that treatment for both  
20 conditions is offered.

21  
22 Interventions for substance misuse may be complicated if the young persons  
23 peer group are also using substances and so there is a strong rationale for  
24 why staff in EIS need to develop strategies to help enable the young person to  
25 recognise the impact of their own substance use on their psychotic symptoms.  
26 In order to do this, EIS staff will need to fully assess substance use including  
27 type, amount and frequency of use of each substance used as well as  
28 understanding the context in which the young person uses each substance  
29 and its function.

## 31 **9.7 SPECIALIST SUBSTANCE MISUSE SERVICES** 32 **FOR YOUNG PEOPLE**

33 The Health Advisory Service reports (1996, 2001) identified a four-tier  
34 framework similar to that described above for CAMHS. The functions of each  
35 tier, rather than the professional discipline involved, are the focus. Different  
36 models and configurations have developed in different regions due to a  
37 variety of factors including the prevalence of substance misuse, the general  
38 level of affluence or deprivation, existing services, and leadership in service  
39 development and innovation. A key issue is that interventions for those  
40 young people whose substance misuse is serious enough to require specialist

1 help are not isolated, but integrated with other medical and social services so  
2 that continuity is established and maintained.

3 ***Tier 1 Universal, generic and primary services***

4 This tier is aimed at all young people. It provides information and advice,  
5 health promotion and support to all young people, parents, families and  
6 carers. At this level, vulnerable individuals with risk factors including child  
7 protection issues may be identified. It is important for staff in such generic  
8 and mainstream services to be aware of the need for a destigmatising non-  
9 confrontational empathic approach to substance use and be equipped to  
10 identify where more complex interventions may be required.

11 ***Tier 2 Specialist services***

12 This tier is directed at vulnerable children who are in contact with children's  
13 services such as CAMHS, YOT, paediatrics, child psychology and voluntary  
14 services and who are potentially vulnerable to the use of substances. Staff  
15 should be skilled in the comprehensive assessment of children and young  
16 people and appreciate the context of developmental issues. Implementation of  
17 advice and counselling, crisis management, outreach, interventions with  
18 family, as well as competence in 'brief interventions' or motivational  
19 enhancement treatments for substance misuse is part of the role.  
20 Collaboration with agencies in the formulation of care planning so that  
21 interventions are integrated – and substance misuse interventions are not  
22 delivered in isolation – is a key component.

23 ***Tier 3 Specialist addiction services***

24 This tier comprises a multidisciplinary team to deliver a complex range of  
25 interventions for young people who have harmful and potentially serious  
26 substance misuse problems and dependence on substances. Close  
27 collaboration with CAMHS, youth justice, voluntary agencies and medical  
28 services is needed in the delivery of these complex care plans. These services  
29 should be integrated with children's services and should cater for the needs of  
30 young people and not be based on adult models. Staff should be competent in  
31 the delivery of the range of pharmacological and individual, group and  
32 family psychological treatments that are available for the treatment of  
33 dependent substance use. Training can be provided to staff to understand the  
34 intricacies of the relationship between mental, physical and social problems  
35 and substance misuse in this age group so that appropriate links can be  
36 forged between the diverse agencies in the locality or region.

37 ***Tier 4 Very specialised services***

1 These are intensely focused interventions of a pharmacological and  
2 psychological nature that require implementation in a residential or inpatient  
3 setting or in a structured day programme, due to the severity of the problems.  
4 Since there are no residential units for adolescent substance misusers at  
5 present, units such as inpatient CAMHS, forensic or paediatric units might be  
6 appropriate for different stages of the care plan. Inpatient detoxification for  
7 alcohol dependence or titration of opiate substitution treatment are examples  
8 of medical interventions requiring inpatient treatment. Intense daily  
9 psychological support may only be achieved in an inpatient CAMHS unit or a  
10 structured day programme. Coordination of support for accommodation,  
11 education and other social needs may also require crisis and fostering  
12 placements in order to achieve stability and safety in critical situations, rather  
13 than the professional groups involved in provision of care.

14  
15 Children and young people may need a range of services from a number of  
16 tiers at different times. Tiers 3 and 4 should not be involved without support  
17 from Tiers 1 and 2. Tiers 1 and 2 are key to the development of a broader base,  
18 a more comprehensive approach and the establishment of credibility and  
19 trust. Continuity of care from Tier 1, particularly in health and education is  
20 crucial. Where possible, coordination and management of the intervention  
21 can be done within Tier 1. This would reduce the stigmatisation and attempt  
22 to 'normalise' the child and his/her family. For those young people not  
23 connected with Tier 1, any other services involved may want to ensure re-  
24 integration and provision of services at Tier 1. Tiers 3 and 4 act as a base for  
25 specialist opinion and focussed interventions.

### 26 **9.7.1 Transition to adult services**

27 The transition to adult services for young people is often marked by a series  
28 of discontinuities in terms of personnel, frequency of treatment (often less  
29 intense in adult services) and treatment approach, and often a failure to  
30 recognise and adapt treatment to developmental stage. Parents who are used  
31 to being intensively involved with CAMH services may feel disengaged with  
32 adult services. In such circumstances the Care Programme Approach (CPA)  
33 and joint working between adult mental health services and CAMHS may  
34 facilitate the transition. A period of engagement with adult services before  
35 handover is preferable. Flexible working around age-limit cut-offs is also  
36 likely to be helpful in promoting smooth transitions.

37  
38 If the young person is primarily being managed in CAMHS, protocols with  
39 adult mental health services need to be in place to ensure the smooth  
40 transition of young people to adult services when they turn 18 years old (or in  
41 some localities 16 years). It is preferable that such protocols ensure that access

1 criteria to adult services are consistent with young people who have been  
2 previously treated by CAMHS, and involve EIS in this process.

3

4 In exceptional circumstances where no age appropriate services are available  
5 for young people, establishing protocols in place for adult services for young  
6 people admitted to adult wards is important. These protocols should include  
7 liaison with and involvement of CAMHS.

8

## 9 **9.8 INTERVENTIONS**

### 10 **9.8.1 Clinical evidence review**

11 A number of existing NICE guidelines have reviewed the evidence for  
12 interventions used to treat young people with psychosis without substance  
13 misuse (that is, bipolar disorder), and interventions used to treat young  
14 people with substance misuse without psychosis (that is, alcohol; drug  
15 misuse: opioid detoxification; drug misuse: psychosocial interventions).

16

17 For the purposes of the guideline, the review questions relating to young  
18 people with psychosis and coexisting substance misuse were sub-questions of  
19 those for adults and, therefore, the review protocols are not repeated here (see  
20 Chapter 6, 7 and 8).

21

22 Where no evidence existed for a particular intervention in young people with  
23 psychosis and coexisting substance misuse, the GDG used informal consensus  
24 to reach a conclusion about whether it was appropriate to cross-reference to  
25 existing NICE guidance.

### 26 **9.8.2 Studies considered for review**

27 Based on the searches conducted for Chapters 6, 7 and 8, only one RCT (Geller  
28 *et al.*, 1998) focusing specifically on young people with psychosis and  
29 coexisting substance misuse, met eligibility criteria. Several further RCTs  
30 (Edwards *et al.*, 2006; Green *et al.*, 2004; Kemp *et al.*, 2007) included young  
31 people, but interpretation of the evidence is difficult as the majority of  
32 participants were over 17 years old. One review (Crome & Bloor, 2005), which  
33 examined interventions for “substance misuse and psychiatric comorbidity in  
34 adolescents,” included the study by Green and colleagues, but no other  
35 research specifically about psychosis. In addition, one review (Bender, *et al.*,  
36 2006) systematically searched for studies of interventions for “dually  
37 diagnosed adolescents”. However, all of the evidence reviewed was for  
38 young people with common mental health disorders, not psychosis.

### 1 **9.8.3 Evidence for the use of pharmacological interventions**

2 One RCT (Geller *et al.*, 1998) randomised 25 young people aged 12 to 18 years  
3 old who had coexisting bipolar and substance dependency disorder to  
4 treatment with lithium or placebo. The results suggested that lithium may be  
5 effective in terms of numbers of participants screening positive for drug use  
6 after 6 weeks of treatment. This study was also reviewed for the NICE bipolar  
7 guideline (NICE, 2006), in which the evidence for psychiatric outcomes was  
8 judged to be inconclusive and of overall low quality. Substance misuse  
9 outcomes were not examined. The participants had less than two months'  
10 history of substance misuse, and the lithium serum levels achieved were high  
11 (0.9 to 1.3 meq/l – the guideline recommended 0.6 to 0.8 meq/l).

### 12 **9.8.4 Guiding principles of treatment**

13 Given the paucity of evidence relating to interventions for young people with  
14 psychosis and coexisting substance misuse, the GDG developed a set of  
15 guiding principles of treatment.

16  
17 First, mental health services are the preferred service to lead the treatment of  
18 a young person with psychosis and coexisting substance misuse. At the same  
19 time, it is necessary for specialist substance misuse services to be involved in  
20 the management of young people with opiate misuse and may advise or offer  
21 a service to those with cannabis misuse, stimulant misuse, or severe alcohol  
22 misuse or dependence. A collaborative coordinated approach is likely to be  
23 the most helpful.

#### 24 *Engagement*

25 Engagement is an essential precursor to treatment. Without it, treatments,  
26 especially psychosocial and environmental, are less likely to be effective. It is  
27 important to take time to engage the young person by adopting a  
28 straightforward, non-confrontational, non-judgemental and optimistic  
29 approach. Assessing readiness to change can help inform care planning and  
30 treatment options.

#### 31 *Risk Management*

32 Young people with psychosis and substance misuse can at times present with  
33 high risk to either themselves or others due to their psychosis, their substance  
34 misuse or a combination of the two. Careful and thorough risk assessments  
35 are needed at initial presentation and whilst ill, with risk management plans  
36 put in place to address any risks identified.

#### 37 *Medication for psychosis*

1 Medication for the treatment of bipolar disorder should follow the NICE  
2 Bipolar Guideline (NICE, 2006). There is currently no NICE guideline for the  
3 treatment of young people with schizophrenia, but guiding principles can be  
4 adopted from the adult schizophrenia guideline (NICE, 2009a).

5

6 In the UK, licensing of antipsychotic drugs for the treatment of schizophrenia  
7 and bipolar disorder in under 18 year olds is variable, with some  
8 manufacturers not recommending these drugs in those under the age of 18  
9 years and the drugs themselves not licensed for this use in this age group.

10 However despite this, considerable clinical experience of their use in young  
11 people has been developed from open trials and from some controlled  
12 evaluations of drug treatments.

13

14 In 2000, the Royal College of Paediatrics and Child Health issued a policy  
15 statement on the use of unlicensed medicines or the use of licensed medicines  
16 for unlicensed applications, in children and young people. This states clearly  
17 that such use is necessary in paediatric practice and that doctors are legally  
18 allowed to prescribe unlicensed medicines where there are no suitable  
19 alternatives and where the use is justified by a responsible body of  
20 professional opinion (Joint Royal College of Paediatrics and Child  
21 Health/Neonatal and Paediatric Pharmacists Group Standing Committee on  
22 Medicines, 2000).

23

24 Caution should be taken with possible drug interactions with substances of  
25 misuse. Dosage should be adjusted according to age and weight/body mass  
26 index.

### 27 *Psychological/ Psychosocial interventions*

28 As for adults, the following psychosocial interventions are used with young  
29 people either on their own or in combination:

- 30 • Motivational interviewing
- 31 • CBT
- 32 • Relapse prevention work
- 33 • Psychoeducation
- 34 • Family work/therapy
- 35 • Contingency management.

36

1 The choice of intervention depends on the nature of the problem and which  
2 approach may appear more appropriate and suitable for a particularly  
3 substance misuse. Motivational enhancement therapy has becoming  
4 increasingly used and evidence is accumulating about its benefits and cost-  
5 effectiveness. Some young people may feel more comfortable concentrating  
6 on behavioural methods rather than treatments that use abstract forms of  
7 reasoning. The 'treatment' needs to focus not only on the substance misuse  
8 but also the psychiatric disorders such as depression, anxiety, ADHD, and  
9 conduct disorders (Chan *et al.*, 2008; Rowe *et al.*, 2004).

10  
11 In the UK, there is also emphasis on harm reduction, including needle  
12 exchange, prevention of drug-related deaths, and treatment for physical  
13 illness and injury. Active support for families, and developing social skills  
14 and competence in parents and children is a recent focus. The Iowa  
15 Strengthening Families Program (Molgaard *et al.*, 1994) and Preparing for the  
16 Drug Free Years (Spoth *et al.*, 2004) and Community Reinforcement and  
17 Family Training (CRAFT) (Waldron *et al.*, 2007) are examples.

### 18 *Treatment of substance misuse*

19 Where available, relevant NICE guidelines can be used to inform treatment of  
20 substance misuse. In addition, it should be noted that young substance  
21 misusers who are referred to Tier 3/4 services are likely to have some  
22 psychological and physical comorbidities as well as be polysubstance  
23 misusers. Thus, treatment of substance misuse should take account of these  
24 possibilities. Constant and consistent review of a young person's clinical state  
25 is crucial, as unpredictability is a feature of young substance misusers.

26  
27 For relevant pharmacological treatments, section 9.8.3 can be consulted in  
28 addition to relevant NICE guidelines. It is crucial that dependence is  
29 diagnosed if medications for withdrawal or substitution are going to be  
30 prescribed. Medications should be prescribed by experienced practitioners  
31 who are aware of the risks in young people. Medications - apart from  
32 buprenorphine - are not licensed for use for under 18 year olds. For  
33 detoxification of alcohol dependence and management of opiate dependence  
34 by detoxification or substitution specialist substance misuse services should  
35 be involved.

### 36 *Input from other agencies*

37 Young people with psychosis and substance misuse often have a range of  
38 social needs. These should be fully assessed and the following services may  
39 need to be involved to address these needs:

- 40 • Housing

- 1           • Education
- 2           • Employment
- 3           • Youth Offending Services (YOS).

4  
5 There are several key elements which contribute to the quality and  
6 effectiveness of young people's substance misuse services. These include  
7 having a comprehensive assessment, an integrated approach, family  
8 involvement, developmental appropriateness, engagement and retention,  
9 qualified staff, gender and cultural competence and evaluation of outcomes  
10 (Knudsen, 2009). Of note was the finding that treatment quality was  
11 significantly greater in programs offering intensive levels of care.

### 12 **9.8.5 Issues of consent to treatment for young people**

13 It is desirable to gain informed consent from both the young person and their  
14 parents, not least because the success of any treatment approach significantly  
15 depends upon the development of a positive therapeutic alliance between the  
16 young person, the family and the professionals. In most outpatient settings,  
17 consent is usually straight forward, as the young person will generally have a  
18 choice to, at least, accept or decline treatment. Nevertheless, it is important to  
19 provide information about the potential risks and benefits of the intervention  
20 being offered, and where appropriate, a choice given between different  
21 treatment options.

22  
23 There may be times when professionals consider inpatient admission to be  
24 necessary, but either the young person or the family do not consent. Under  
25 the Mental Health Act 1983 (HMSO, 1983), there have been some changes to  
26 the law regarding young people under the age of 18 years.

27  
28 If a young person aged 16 or 17 years old has capacity to give or refuse  
29 consent for treatment, it is no longer possible for the person with parental  
30 authority to over-rule the young person's wishes. However, for those under  
31 the age of 16 years a 'Gillick-competent' young person can still be admitted  
32 against his or her wishes with the consent of someone with parental  
33 authority. Whilst the use of parental consent is legal, the Code of Practice for  
34 the Mental Health Act (HMSO, 2007) advises against this, suggesting it is  
35 good practice to consider the use of other appropriate legislation, usually the  
36 Mental Health Act (HMSO, 2007). This includes safeguards such as the  
37 involvement of other professionals, a time limit and a straightforward  
38 procedure for appeals and regular reviews. It also avoids a possible conflict  
39 with the Human Rights Act, 1998 (HMSO, 1998a).

1  
2 On the other hand, a 'Gillick competent' young person below the age of 16  
3 years has the right to consent to treatment. If the person with parental  
4 authority objects, these objections must be considered but will not necessarily  
5 prevail.

6  
7 Alternative legislation includes using a care order (Section 31) under the  
8 Children Act 1989 (HMSO, 1989) or a specific issue order (Section 8). Both of  
9 these options normally involve social services and can be time consuming.  
10 Another, more rapid alternative to the Children Act (HMSO, 1989), is to apply  
11 for a Wardship Order, which in an emergency can be organised by telephone.

### 12 **9.8.6 Clinical evidence summary**

13 In one small trial (N=25) assessing pharmacological interventions for young  
14 people, lithium was compared with placebo. Based on this evidence  
15 (*GRADED* low quality), it was not possible to reach a decision about the  
16 effectiveness of pharmacological interventions for young people with  
17 psychosis and coexisting substance misuse.

18  
19 There was no evidence for psychological or psychosocial interventions for  
20 young people with psychosis and coexisting substance misuse.

## 21 **9.9 FROM EVIDENCE TO RECOMMENDATIONS**

22 Based on the limited evidence base, the GDG were required to extrapolate  
23 from data which may not accurately address treatment effectiveness for  
24 young people with psychosis and coexisting substance misuse. The GDG  
25 therefore developed guiding principles of treatment and recommendations  
26 based on consensus. The GDG recognises that as new evidence emerges on  
27 treatment for young people with psychosis and coexisting substance misuse,  
28 the recommendations in this guideline will be revised and updated  
29 accordingly. The recommendations cover competency, identification and  
30 referral, and assessment and treatment.

31  
32 The GDG felt that professionals in Tier 1 CAMHS should be competent to  
33 recognise early signs of psychosis and substance misuse, while Tier 3 and 4  
34 CAMHS, and early intervention service healthcare professionals should be  
35 competent with regard to management of psychosis and coexisting substance  
36 misuse. With regard to identification and referral, the GDG felt that  
37 professionals in Tier 1 should seek advice from Tier 2 staff when signs of  
38 psychosis are detected in young people. In Tier 2 services, referral should be  
39 made according to local protocols. The GDG also thought that it was  
40 important that all young people with psychosis or suspected psychosis seen

1 by professionals in Tier 3 or 4 services, or early intervention services, should  
2 be asked about substance misuse. Referral to Tier 4 CAMHS should be done  
3 directly when a comprehensive assessment reveals a high risk of harm to self  
4 or others. In terms of assessment, the GDG thought that there needed to be a  
5 recommendation to ensure that healthcare professionals are familiar with the  
6 legal framework that applies to young people. In terms of treatment, the GDG  
7 felt that recommendations for the treatment of adults should be followed, but  
8 adapted for young people if necessary. It was also recognised that other  
9 agencies, including children's services should be involved to ensure that the  
10 young person's educational, employment, family and housing needs are met.  
11 Finally, the GDG thought that a recommendation directed at commissioners  
12 was needed to ensure that age-appropriate mental health services are  
13 available for young people with psychosis and coexisting substance misuse,  
14 and that transition arrangements to adult mental health services are in place  
15 where appropriate.

16  
17 In addition, the GDG discussed that because onset of psychosis at a younger  
18 age is also an indicator of poor prognosis, people with a combination of  
19 younger age of onset and coexisting substance misuse may have a particularly  
20 poor prognosis. A clearer understanding of the risk and protective factors for  
21 substance misuse in young people with psychosis, and the interrelationship of  
22 the two conditions over time, may facilitate the development of treatment  
23 approaches for the coexisting conditions in this group. This may then improve  
24 the longer term outcome for a group of people who tend to have a poor  
25 prognosis.  
26

1 **9.10 CLINICAL PRACTICE RECOMMENDATIONS**

2 **9.10.1 Recommendations (Specific issues for young people with**  
3 **psychosis and coexisting substance misuse)**

4 **Competence**

5 **9.10.1.1** Professionals in Tier 1 (primary care and educational settings) should  
6 be competent to recognise early signs of psychosis and substance  
7 misuse in young people.

8 **9.10.1.2** Healthcare professionals in Tier 3 (community mental health teams)  
9 and Tier 4 (specialist inpatient and regional services) CAMHS, and in  
10 early intervention in psychosis services, should be competent in the  
11 management of psychosis and substance misuse in young people.

12 **Identification and referral**

13 **9.10.1.3** Professionals in Tier 1 (primary care and educational settings) should  
14 seek advice or consultation from Tier 2 CAMHS (primary care) when  
15 signs of psychosis are detected in young people. If healthcare  
16 professionals in Tier 2 CAMHS detect signs of psychosis in young  
17 people, a referral to Tier 3 CAMHS or early intervention in psychosis  
18 services for young people should be made according to local  
19 protocols.

20 **9.10.1.4** Ask all young people seen in Tier 3 and Tier 4 CAMHS and in early  
21 intervention in psychosis services who have psychosis or suspected  
22 psychosis about substance misuse (see 5.8.1.1).

23 **9.10.1.5** Children and young people who, after comprehensive assessment,  
24 are considered to be at high risk of harm to themselves or others,  
25 should be referred directly to Tier 4 CAMHS including inpatient  
26 services where necessary.

27 **Assessment and treatment**

28 **9.10.1.6** Healthcare professionals working with young people with psychosis  
29 and coexisting substance misuse should ensure they are familiar with  
30 the legal framework that applies to young people including the  
31 Mental Health Act (1983; amended 1995 and 2007), the Mental  
32 Capacity Act (2005), and the Children Act (2004).

1 **9.10.1.7** For psychological, psychosocial, family and medical interventions for  
2 young people, follow the recommendations for adults in this  
3 guideline; they may need to be adapted according to the young  
4 person's circumstances and age. In addition, other agencies, including  
5 children's services, should be involved to ensure that the young  
6 person's educational, employment, family and housing needs are met.

7 **9.10.1.8** When prescribing medication, take into account the young person's  
8 age and weight when determining the dose. If it is appropriate to  
9 prescribe unlicensed medication, explain to the young person and/or  
10 their parents or carers the reasons for doing this.

11 **9.10.1.9** Those providing and commissioning services should ensure that:

- 12 • age-appropriate mental health services are available for young people  
13 with psychosis and coexisting substance misuse **and**
- 14 • transition arrangements to adult mental health services are in place  
15 where appropriate.

16

## 17 **9.10.2 Research Recommendations**

18 **9.10.2.1** What risk factors predict the onset of substance misuse in young  
19 people with psychosis?

20

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19	Appendix 14: Clinical evidence - forest plots.....	On website
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21	Appendix 16: Clinical evidence - completed methodology checklists.....	On website
22	Appendix 17: Economic evidence - GRADE profiles.....	On website
23	Appendix 18: Economic evidence - completed methodology checklists..	On website
24	Appendix 19: Economic plan.....	On website
25		

1 **APPENDIX 1: SCOPE FOR THE DEVELOPMENT OF THE**  
2 **CLINICAL GUIDELINE**

3 **1** *Guideline title*

4 Psychosis in conjunction with substance misuse: the assessment and  
5 management of psychosis with substance misuse

7 **1.1** *Short title*

8 Psychosis with substance misuse

10 **2** *The remit*

11 The Department of Health has asked NICE: "To develop a clinical guideline  
12 for the assessment and management of severe mental illness in conjunction  
13 with problematic substance misuse."

14 **3** *Clinical need for the guideline*

15 **3.1** *Epidemiology*

16 a) The term psychosis is used to describe a major group of severe  
17 disorders of mental health characterised by the presence of delusions  
18 and hallucinations that disrupt a person's perception, thoughts,  
19 emotions and behaviour. The two main forms of this are schizophrenia  
20 and bipolar disorder. Substance misuse is a broad term encompassing  
21 the use of any psychotropic medication or substance, whether illicit or  
22 not, or taken for pleasure or not, if the use is considered hazardous or  
23 harmful. It includes, for example, alcohol, and prescribed medications  
24 used for purposes other than those prescribed. Such use is usually, but  
25 not always, regarded as a problem if there is evidence of dependence,  
26 characterised by psychological reinforcement of repeated drug-taking  
27 behaviour and, in some cases, a withdrawal syndrome.

28  
29 b) In the UK, the annual prevalence for probable psychotic disorder  
30 among adults living in private households is about 5 per 1000. This  
31 figure is 9 per 1000 in adults aged 30–44 years and 18 per 1000 in adults  
32 with an African-Caribbean family background. Among those  
33 diagnosed with a psychotic disorder, studies show that prevalence for  
34 any substance misuse ranges from 24–36% (7–20% for alcohol misuse  
35 only, 5–9% for drug misuse only, 8% for drug and alcohol misuse). In

1 one study of people with a psychotic disorder, 35% of the sample had a  
2 lifetime history of any illicit drug use. Prevalence rates for substance  
3 misuse are even higher in forensic (50–70%) and inpatient (30–49%)  
4 mental health services. In addition, service users with comorbid drug  
5 misuse spend twice as long in hospital, on average, and have higher  
6 levels of unmet needs, compared with other inpatients with psychosis.  
7

- 8 c) Substance misuse among individuals with psychiatric disorders is  
9 associated with significantly poorer outcomes than for individuals  
10 with a single disorder. These outcomes include worsening psychiatric  
11 symptoms, poorer physical health, increased use of institutional  
12 services, poor medication adherence, homelessness and increased risk  
13 of HIV infection, as well as poor social outcomes including impact on  
14 carers and family and contact with the criminal justice system.  
15
- 16 d) There is a substantial link between substance misuse and crime. Hence  
17 the provision in the Crime and Disorder Act 1998 (HMSO, 1998b) for  
18 drug treatment and testing orders and in the Criminal Justice and  
19 Court Services Act 2000 drug abstinence orders and drug abstinence  
20 requirements.  
21
- 22 e) Compared to people with psychosis only, people with psychosis and  
23 substance misuse have greater levels of inpatient mental health service  
24 use, higher overall treatment costs, and lower concordance with  
25 community care and medication.

### 26 3.2 *Current practice*

- 27 a) The National Service Framework for Mental Health (Department of  
28 Health, 1999), sets out how services will be planned, delivered and  
29 monitored. Several areas are relevant to this guideline including  
30 mental health promotion, primary care and specialist services. The  
31 following are also relevant:
- 32 • The Care Programme Approach (CPA). This is a framework for  
33 interagency working. It seeks to ensure that service users have a  
34 proper assessment and that services are coordinated in line with  
35 service user need.
  - 36 • Assertive outreach and crisis resolution services. These are  
37 proactive approaches to engaging with service users and managing  
38 problems.

39

- 1        b) Less than a fifth of people who have co-existing psychosis and  
2        substance misuse receive substance misuse interventions, and there is  
3        clearly uneven distribution of services with regard to ethnicity. In  
4        substance misuse services those with a severe mental illness and co-  
5        existing substance misuse are generally white; assertive outreach teams  
6        have a much higher proportion of service users classified as African-  
7        Caribbean than all other teams.  
8  
9        c) There are no uniformly agreed screening or assessment tools.  
10  
11       d) The following three treatment models have been described in the  
12       literature, but there is currently little guidance about which is the most  
13       effective or cost effective:  
14       • Serial treatment – one treatment, either psychiatric or substance  
15       misuse is followed by the other  
16       • Parallel treatment – the concurrent but separate treatment of both  
17       the psychiatric disorder and the substance misuse disorder  
18       • Integrated treatment – substance misuse and psychiatric treatment  
19       are provided concurrently by the same personnel.

20

## 21    4        *The guideline*

22    The guideline development process is described in detail on the NICE website  
23    (see section 6, 'Further information').

24    This scope defines what the guideline will (and will not) examine, and what  
25    the guideline developers will consider. The scope is based on the referral from  
26    the Department of Health.

27    The areas that will be addressed by the guideline are described in the  
28    following sections.

29

### 30    4.1    *Population*

#### 31    4.1.1    **Groups that will be covered**

32

- 33        a) Adults and young people (14 and older) who have a clinical working  
34        diagnosis of schizophrenia<sup>24</sup>, bipolar or other affective psychosis, in  
35        conjunction with substance misuse.  
36

---

<sup>24</sup> This includes schizoaffective disorder and delusional disorder.

- 1           b) This will include specific consideration of the needs of people with  
2           coexisting learning difficulties or significant physical or sensory  
3           difficulties, and the needs of people from black and minority ethnic  
4           groups.  
5

6   **4.1.2 Groups that will not be covered**  
7

- 8           a) People with very late onset psychosis (onset after age 60) and  
9           coexisting substance misuse.  
10

11   **4.2 Healthcare setting**

- 12           a) Care that is received from healthcare professionals in primary and  
13           secondary care, including standard inpatient and forensic settings, who  
14           have direct contact with, and make decisions concerning, the care of  
15           people with severe mental illness and substance misuse.  
16

- 17           b) Whilst the guideline will not provide specific recommendations for  
18           accident and emergency departments, paramedic services, prison  
19           medical services, the police and those who work in the criminal justice  
20           and education sectors, the guideline will be relevant to their work. The  
21           evidence considered in this guideline will not be derived from these  
22           settings.  
23

24   **4.3 Clinical management**

25   **4.3.1 Key clinical issues that will be covered**

- 26           a) Identification and assessment.  
27  
28           b) Sequencing of treatment, and integrated versus non-integrated models  
29           of care.  
30  
31           c) The use of antipsychotic medication and/or psychological or  
32           psychosocial interventions (for example, family intervention) for the  
33           treatment of people with co-existing psychosis, and substance misuse.  
34  
35           d) Psychosocial interventions for the management of substance misuse  
36           (for example, cognitive behavioural therapy [CBT], motivational  
37           interviewing and contingency management) in people with coexisting  
38           psychosis.  
39

- 1 e) Pharmacological (for example, opioid antagonists) and physical  
2 interventions for the management of substance misuse in people with  
3 coexisting psychosis.  
4
- 5 f) Residential rehabilitation and inpatient mental health care of people  
6 with coexisting psychosis and substance misuse (including in a  
7 forensic setting).  
8
- 9 g) Working with non-NHS services (for example, the police and those  
10 who work in the criminal justice and education sectors).  
11
- 12 h) Ways to improve access to mental health services for people from black  
13 and minority ethnic communities (this will include issues concerned  
14 with engagement with services).  
15
- 16 i) Interactions between prescribed medication and substances misused.  
17
- 18 j) Ways to improve insight (that is, an individual's awareness of mental  
19 disorder and substance misuse, awareness of the social consequences  
20 of disorder/substance misuse, awareness of the need for treatment,  
21 awareness of symptoms and attribution of symptoms to  
22 disorder/substance misuse).  
23
- 24 k) Ways to improve and manage non-adherence to treatment. This  
25 guideline will cross refer to the NICE clinical guideline on medicines  
26 adherence where appropriate.  
27
- 28 l) Note that guideline recommendations for pharmacological  
29 interventions will normally fall within licensed indications;  
30 exceptionally, and only if clearly supported by evidence, use outside a  
31 licensed indication may be recommended. The guideline will assume  
32 that prescribers will use a drug's summary of product characteristics to  
33 support joint clinical decision-making between service users and  
34 prescribers.  
35

#### 36 **4.3.2 Clinical issues that will not be covered**

- 37 a) Primary prevention.  
38
- 39 b) Diagnosis.  
40
- 41 c) Management of violence in people with severe mental illness.  
42

1 **4.4 Economic aspects**

2 Developers will take into account both clinical and cost effectiveness when  
3 making recommendations involving a choice between alternative  
4 interventions. A review of the economic evidence will be conducted and  
5 analyses will be carried out as appropriate. The preferred unit of effectiveness  
6 is the quality-adjusted life year (QALY), and the costs considered will usually  
7 only be from an NHS and personal social services (PSS) perspective. Further  
8 detail on the methods can be found in 'The guidelines manual' (see 'Further  
9 information').

10

11 **4.5 Status**

12 **4.5.1 Scope**

13

14 This is the final scope.

15

16 **4.5.2 Timing**

17

18 The development of the guideline recommendations will begin in May 2009.

19

20 **5 Related NICE guidance**

21 **5.1 Published guidance**

22 • Schizophrenia. NICE clinical guideline 82 (2009 [NICE, 2009a]).  
23 Available from [www.nice.org.uk/CG82](http://www.nice.org.uk/CG82)

24 • Medicines adherence. NICE clinical guideline 76 (2009 [NICE,  
25 2009c]). Available from [www.nice.org.uk/CG76](http://www.nice.org.uk/CG76)

26 • Drug misuse: opioid detoxification. NICE clinical guideline 52 (2007  
27 [NICE, 2007a]). Available from [www.nice.org.uk/CG52](http://www.nice.org.uk/CG52)

28 • Drug misuse: psychosocial interventions. NICE clinical guideline 51  
29 (2007 [NICE, 2007b]). Available from [www.nice.org.uk/CG51](http://www.nice.org.uk/CG51)

30 • Interventions to reduce substance misuse among vulnerable young  
31 people. NICE public health guidance 4 (2007 [NICE, 2007c]).  
32 Available from [www.nice.org.uk/PH4](http://www.nice.org.uk/PH4)

33 • Naltrexone for the management of opioid dependence. NICE  
34 technology appraisal guidance 115 (2007 [NICE, 2007d]). Available  
35 from [www.nice.org.uk/TA115](http://www.nice.org.uk/TA115)

- 1           • Methadone and buprenorphine for managing opioid dependence.  
2           NICE technology appraisal guidance 114 (2007 [NICE, 2007e]).  
3           Available from [www.nice.org.uk/TA114](http://www.nice.org.uk/TA114)
- 4           • Bipolar disorder. NICE clinical guideline 38 (2006 [NICE, 2006]).  
5           Available from [www.nice.org.uk/CG38](http://www.nice.org.uk/CG38)
- 6           • Violence. NICE clinical guideline 25 (2005 [NICE, 2005]). Available  
7           from [www.nice.org.uk/CG25](http://www.nice.org.uk/CG25)
- 8           • Schizophrenia. NICE clinical guideline 1 (2002 [NICE, 2002]).  
9           Available from [www.nice.org.uk/CG1](http://www.nice.org.uk/CG1)

10

## 11   **5.2   Guidance under development**

- 12           • NICE is currently developing the following related guidance  
13           (details available from the NICE website).
- 14           • Alcohol use disorders (prevention). NICE public health guidance.  
15           Publication expected March 2010.
- 16           • Alcohol use disorders (clinical management). NICE clinical  
17           guideline. Publication expected May 2010.
- 18           • Alcohol dependence and harmful alcohol use. NICE clinical  
19           guideline. Publication expected January 2011.

20

## 21   **6       Further information**

22   Information on the guideline development process is provided in:

- 23           • 'How NICE clinical guidelines are developed: an overview for  
24           stakeholders' the public and the NHS'
- 25           • 'The guidelines manual'.

26

27   These are available from the NICE website  
28   ([www.nice.org.uk/guidelinesmanual](http://www.nice.org.uk/guidelinesmanual)). Information on the progress of the  
29   guideline will also be available from the NICE website ([www.nice.org.uk](http://www.nice.org.uk)).

30

31

## 1 APPENDIX 2: DECLARATIONS OF INTERESTS BY GDG

### 2 MEMBERS

3 With a range of practical experience relevant to the treatment and  
4 management of psychosis in conjunction with substance misuse in the GDG,  
5 members were appointed because of their understanding and expertise in  
6 healthcare for people with psychosis and substance misuse and support for  
7 their families/carers, including: scientific issues; health research; the delivery  
8 and receipt of healthcare, along with the work of the healthcare industry; and  
9 the role of professional organisations and organisations for people with  
10 psychosis and substance misuse and their families/carers.

11  
12 To minimise and manage any potential conflicts of interest, and to avoid any  
13 public concern that commercial or other financial interests have affected the  
14 work of the GDG and influenced guidance, members of the GDG must  
15 declare as a matter of public record any interests held by themselves or their  
16 families which fall under specified categories (see below). These categories  
17 include any relationships they have with the healthcare industries,  
18 professional organisations and organisations for people with psychosis and  
19 substance misuse and their families/carers.

20  
21 Individuals invited to join the GDG were asked to declare their interests  
22 before being appointed. To allow the management of any potential conflicts of  
23 interest that might arise during the development of the guideline, GDG  
24 members were also asked to declare their interests at each GDG meeting  
25 throughout the guideline development process. The interests of all the  
26 members of the GDG are listed below, including interests declared prior to  
27 appointment and during the guideline development process.

#### 28 *Categories of interest*

##### 29 **Paid employment**

30  
31 **Personal pecuniary interest:** financial payments or other benefits from either  
32 the manufacturer or the owner of the product or service under consideration  
33 in this guideline, or the industry or sector from which the product or service  
34 comes. This includes holding a directorship, or other paid position; carrying  
35 out consultancy or fee paid work; having shareholdings or other beneficial  
36 interests; receiving expenses and hospitality over and above what would be  
37 reasonably expected to attend meetings and conferences.

38

1 **Personal family interest:** financial payments or other benefits from the  
2 healthcare industry that were received by a member of your family.

3  
4 **Non-personal pecuniary interest:** financial payments or other benefits  
5 received by the GDG member's organisation or department, but where the  
6 GDG member has not personally received payment, including fellowships  
7 and other support provided by the healthcare industry. This includes a grant  
8 or fellowship or other payment to sponsor a post, or contribute to the running  
9 costs of the department; commissioning of research or other work; contracts  
10 with, or grants from, NICE.

11  
12 **Personal non-pecuniary interest:** these include, but are not limited to, clear  
13 opinions or public statements you have made about individuals with  
14 psychosis and substance misuse problems, holding office in a professional  
15 organisation or advocacy group with a direct interest in psychosis and  
16 substance misuse, other reputational risks relevant to psychosis and substance  
17 misuse.

18

<b>Guideline Development Group - Declarations of interest</b>	
Professor Peter Tyrer - Chair, Guideline Development Group	
Employment	Professor of Community Psychiatry Department of Psychological Medicine, Imperial College
Personal pecuniary interest	The originator of the treatment called nidotherapy which may be used in the population considered in this guideline, and conducted a study looking at Nidotherapy.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	Published books and articles on Nidotherapy
Non-personal non-pecuniary interest	A contingency management study is being conducted within my department.
Action Taken	Nidotherapy was discussed by the GDG on 2 March 2010. It was decided that it was not appropriate for the Chair to be present and Peter Tyrer left the room for this discussion. All members were asked individually if they felt this approach was acceptable and all agreed.
Professor Mohammed T. Abou-Saleh	
Employment	Professor of Psychiatry, St George's, University of London and Honorary Consultant in Addiction Psychiatry, South West London and St George's Mental Health NHS Trust, London
Personal pecuniary interest	None
Personal family interest	None

## FINAL CONSULTATION

Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Non-personal non-pecuniary interest	Asked to chair a presentation at an event sponsored by a pharmaceutical company, although he did not receive any money for this.
Action Taken	None
<b>Professor Christine Barrowclough</b>	
Employment	Prof of Clinical Psychology, University of Manchester
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	Currently Chief Investigator for two major studies evaluating psychological therapy for people with psychosis with substance misuse.
Action Taken	None
<b>Ms. Tina Braithwaite</b>	
Employment	Service User/Carer Representative. Director of Service User Involvement, Revolving Doors Agency. Also I'm a Member of the lived experience advisory panel, REFOCUS Recovery Research Project.
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr Andy Cotgrove</b>	
Employment	Young people (CAMHS level 4), Pine Lodge Young People's Centre
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr. Mike Crawford</b>	
Employment	Reader in Mental Health Services Research, Imperial College London / CNWL Mental Health NHS Trust
Personal pecuniary interest	Involved in a study on Nidotherapy.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	Nidotherapy was discussed by the GDG on 2 March 2010. It was decided that Mike Crawford could be present to answer any queries, but not be involved in

	the discussion. All members were asked individually if they felt this approach was acceptable and all agreed.
<b>Professor Ilana Crome</b>	
Employment	Professor of Addiction Psychiatry, Keele University November 2009 – ongoing Honorary Consultant Addiction Psychiatrist, South Staffordshire and Shropshire Foundation Trust. Prior to November 2009 – Honorary Consultant Addiction Psychiatrist, North Staffordshire Combined Healthcare NHS Trust.
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	The Academic Psychiatry Unit, Keele University receives funding from pharmaceutical companies which covers speakers' expenses for regular departmental seminar series.  Keele University has received funding from DH, Home Office, SCIE (Social Care Institute for Excellence, for research on drug misuse and mental illness.  Policy roles for DH, Scottish Executive and Welsh Assembly
Personal non-pecuniary interest	Member, Advisory Council on the Misuse of Drugs. Specific roles in Cannabis and Schizophrenia research which informed recommendation on Cannabis re- classification; Pathways to Problems report. ACMD, Chair Working Group on Treatment Effectiveness  Member, Faculty of Academic Psychiatry, Royal College of Psychiatrists Member, Young People's Working Group, Royal College of Psychiatrists Honorary Secretary, Professors of Psychiatry Club Chair, WG Older people and substance misuse, Royal College of Psychiatrists  Member, British Association of Psychopharmacology, Consensus group on Addiction and Comorbidity  Trustee, Society for the Study of Addiction  Chair, Steering Committee Assertive Community Treatment of Alcohol Dependence Trial, MRC funded trial led by Institute of Psychiatry  Member, Young people and drugs and alcohol study DIPEX Research Group (Youthtalk)

	<p>Member, Young people and depression study DIPEX Research Group (Youthtalk)</p> <p>Consultant, PaRticipation of the ElDerly In Clinical Trials(PREDICT) 2007-2009 European Union Project developed and recently launched a charter for evaluation of medicines in older people.</p> <p>Steering Group Advisory Panel, National Undergraduate Substance Misuse Curriculum Implementation Group</p> <p>Advisor, Turning Point</p> <p>Editorial responsibilities for several journals e.g. International editor, American Journal of Addiction Editor, Drugs Education Prevention and Policy International Advisory Board: British Journal of Psychiatry. British Journal of Psychiatry, Addiction, Journal of Mental Illness and Substance, Misuse, J of Psychopharmacology Member, International Society of Addiction Journal Editors</p>
Action Taken	None
<b>Mr. Mike Firn</b>	
Employment	Clinical Service Development Lead
Personal pecuniary interest	Non-guideline specific interest: specifically I am Chair of a mutual trading organisation (National Forum for Assertive Outreach) that has educational grants from Janssen-Cilag pharmaceuticals covering venue and catering costs of 2 regional network events in Manchester within the last year. There has been no product information or talks given at either of these events beyond acknowledgement of the room and catering costs.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr. Frank Holloway</b>	
Employment	Consultant Psychiatrist and Clinical Director, Bethlem Royal Hospital
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None

## FINAL CONSULTATION

<b>Dr. Cheryl Kipping</b>	
Employment	Nurse Consultant, South London And Maudsley NHS Foundation Trust
Personal pecuniary interest	Member of independent review team into SUIs in a PCT area.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	<ul style="list-style-type: none"> <li>• Member of PROGRESS (dual diagnosis nurse consultant group). Co-ordinated group's response to consultation on scope of PSM guideline.</li> <li>• Member of DH steering group that developed DH (2002) Mental Health Policy Implementation Guide: Dual Diagnosis Good Practice Guide</li> <li>• Co-editor of Advances in Dual Diagnosis journal</li> <li>• Provide specialist dual diagnosis advice to National Mental Health Development Unit (NMHDU) dual diagnosis and acute programmes. Involved in development of dual diagnosis elearning packages for NMHDU Dual Diagnosis programme and National Acute Project Board.</li> </ul>
Action Taken	None
<b>Dr. Kate McKinnell</b>	
Employment	Senior Medical Officer (Addictions) Sefton Integrated Recovery Team (Crime Reduction Initiatives)
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr. Jonathan Mitchell</b>	
Employment	Consultant Psychiatrist - Early Intervention, East Glade Centre
Personal pecuniary interest	In 2006 I chaired an educational meeting sponsored by Eli Lilly for which I received a payment of £250. In 2007 I chaired an educational meeting sponsored by Jansen for which I was offered, but did not accept payment. I have no current or ongoing personal pecuniary interests.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr. David Ndegwa</b>	
Employment	Consultant Forensic Psychiatrist / Strategy Director

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	South London & Maudsley NHS Foundation Trust
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Mr. Peter Pratt</b>	
Employment	Chief Pharmacist, Sheffield Health & Social Care Trust And Rotherham Doncaster & South Humber NHS Trust
Personal pecuniary interest	Gave a presentation regarding payment by results in mental health at an event sponsored by Janssen-Cilag. Executive member of NAPICU committee (National Association of Psychiatric Intensive Care Units) Received payment for market research about schizophrenia.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Ms. Theresa Renwick</b>	
Employment	Social care lead for mental health, Royal Borough of Kensington and Chelsea
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Mr. Leroy Simpson</b>	
Employment	Service User/Carer Representative. Board Member, Salvation Army Housing Association.
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Mrs. Penelope Wigram</b>	
Employment	Service User/Carer Representative
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None

## FINAL CONSULTATION

<b>Professor. Tim Kendall</b>	
Employment	Director, NCCMH Medical Director, Sheffield Health and Social Care Trust Consultant Adult Psychiatrist
Personal pecuniary interest	Grant holder for £1.44 million per year (approx) from NICE for guidelines work. Work with NICE International. Undertake some research into mental health, and the mental health workforce for DH, Royal College of Psychiatrists and the academy of medical royal colleges.
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Dr. Craig Whittington</b>	
Employment	Senior Systematic Reviewer, NCCMH
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	Conducted a sub-analysis on the nidotherapy study for publication and subsequent use by GDG
<b>Mr. Matthew Dyer</b>	
Employment	Health Economist, NCCMH
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Ms. Sarah Stockton</b>	
Employment	Senior Information Scientist, NCCMH
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Ms. Laura Shields</b>	
Employment	Research Assistant, NCCMH
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None
<b>Ms. Katherine Leggett</b>	

## FINAL CONSULTATION

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Employment	Guideline Development Manager
Personal pecuniary interest	None
Personal family interest	None
Non-personal pecuniary interest	None
Personal non-pecuniary interest	None
Action Taken	None

1  
2  
3

1 **APPENDIX 3: EXPERT REVIEWERS TO THE GDG**

2 Dr Michelle Cleary, Research Unit, Rozelle Hospital, Sydney South West Area  
3 Health Service.

4

5

6

1 **APPENDIX 4: STAKEHOLDERS AND EXPERTS WHO**  
2 **SUBMITTED COMMENTS IN RESPONSE TO THE**  
3 **CONSULTATION DRAFT OF THE GUIDELINE**

4 *Stakeholders*

Alder Hey Children's NHS Foundation Trust  
British Association for Psychopharmacology

Central and North West London NHS Trust  
College of Mental Health Pharmacy  
Department of Health  
Faculty of Forensic and Legal Medicine  
Huntercombe Group  
International Society for the Psychological Treatment of the Schizophrenias  
and Other Psychoses  
Lancashire Care NHS Foundation Trust  
Manchester Mental Health and Social Care Trust  
Mental Health Nurses Association  
MIDAS Therapists  
National Institute for Health and Clinical Excellence  
National Mental Health Development Unit  
National Treatment Agency for Substance Misuse  
NETSCC - Referee 1  
NETSCC - Referee 2  
NHS Direct  
Nottinghamshire Healthcare NHS Trust  
National Consortium of Consultant Nurses in Dual Diagnosis  
Royal College of Nursing  
Royal College of Paediatrics and Child Health  
Royal College of Psychiatrists  
Royal College of Psychiatrists (Wales)  
Royal Pharmaceutical Society  
Specialist Clinical Addiction Network  
Welsh Assembly Government  
West London Mental Health NHS Trust  
Yorkshire and the Humber LSA

5

6 *Experts*

7 Dr Carol Caton, University of Columbia, New York, USA.

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1 **APPENDIX 5: RESEARCHERS CONTACTED TO REQUEST**  
2 **FURTHER INFORMATION ABOUT PUBLISHED OR**  
3 **UNPUBLISHED EVIDENCE**

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5 Dr. Alan Bellack, University of Maryland School of Medicine

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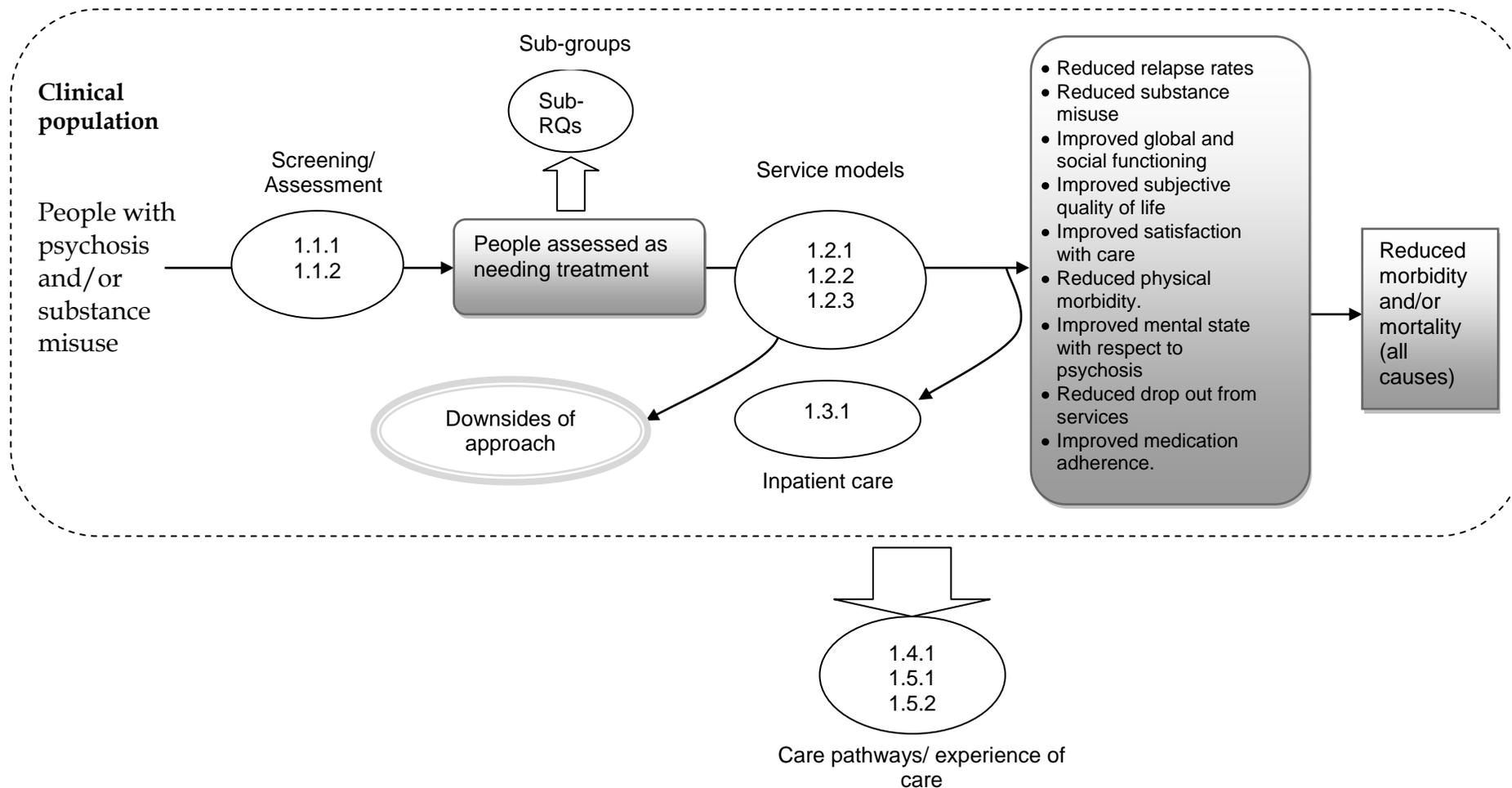
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## APPENDIX 6: ANALYTIC FRAMEWORK AND REVIEW QUESTIONS

### *Assessment/service models/ inpatient care/care pathways/experience of care*



### Assessment

No.	Primary review questions
1.1.1	<p>In people with psychosis and coexisting substance misuse, what are the key elements for a comprehensive assessment (of needs and risks)?</p> <p>Sub-question 1: should the assessment be the same in primary and secondary care?</p> <p>Sub-question 2: should the assessment be modified for sub-groups of people (for example, young people, women, people from BME groups, homeless people, offenders, type of psychosis, type of substance misuse)?</p> <p>Sub-question 3: what factors should trigger a reassessment?</p>

### Service models

No.	Primary review questions
1.2.1	<p>In people with psychosis and coexisting substance misuse, does an integrated service model (usually involving the model of assertive community treatment) when compared with an alternative management strategy lead to:</p> <p>Critical outcomes:</p> <ul style="list-style-type: none"> <li>• Reduced mortality (all causes)</li> <li>• Reduced relapse rates (measured by exacerbation of symptoms requiring change in health care management)</li> <li>• Reduced substance misuse (however measured)</li> <li>• Improved global and social functioning (for example, employment, accommodation)</li> <li>• Improved subjective quality of life</li> <li>• Improved satisfaction with care</li> <li>• Reduced physical morbidity.</li> </ul> <p>Secondary outcomes:</p> <ul style="list-style-type: none"> <li>• Insight</li> <li>• Improved medication adherence</li> <li>• Improved access to services (reduced drop out)</li> <li>• Reduced relapse rates (measured by admission to hospital; number of bed days)</li> </ul>

	<ul style="list-style-type: none"> <li>• Improved mental state with respect to psychosis (for example, PANSS)</li> <li>• Reduced offending behavior.</li> </ul> <p>Sub-question 1: What are the elements in an integrated service model that are most likely to be associated with better outcomes?</p> <p>Sub-question 2: Are there any subgroups of people (for example, young people, BME groups) that benefit from some elements of the service model more than others?</p> <p>Sub-question 3: Are there subgroups of people (for example, based on severity of substance misuse and severity of psychosis; young people, BME groups) that may benefit from alternatives strategies (non-integrated service models – serial treatment, for example)</p>
1.2.2	<p>In people with psychosis and coexisting substance misuse, do the psychological/psychosocial interventions listed below (delivered within an integrated service model) when compared to an alternative management strategy lead to improved outcomes? (for outcomes see 1.2.1)</p> <ul style="list-style-type: none"> <li>• Individual interventions</li> <li>• Group interventions</li> <li>• Family intervention</li> <li>• Contingency management</li> <li>• Combined interventions</li> </ul>
1.2.3	<p>In people with psychosis and coexisting substance misuse, does staffed accommodation when compared to an alternative management strategy lead to improved outcomes? (for outcomes see 1.2.1)</p>

### Inpatient care

No.	Primary review questions
1.3.1	<p>When a person with psychosis and coexisting substance misuse is admitted to an inpatient mental health setting (including forensic settings), should treatment follow the same principles as interventions delivered in a community setting?</p> <p>Sub-question: Are there subgroups of people for whom we would alter our approach to treatment?</p>

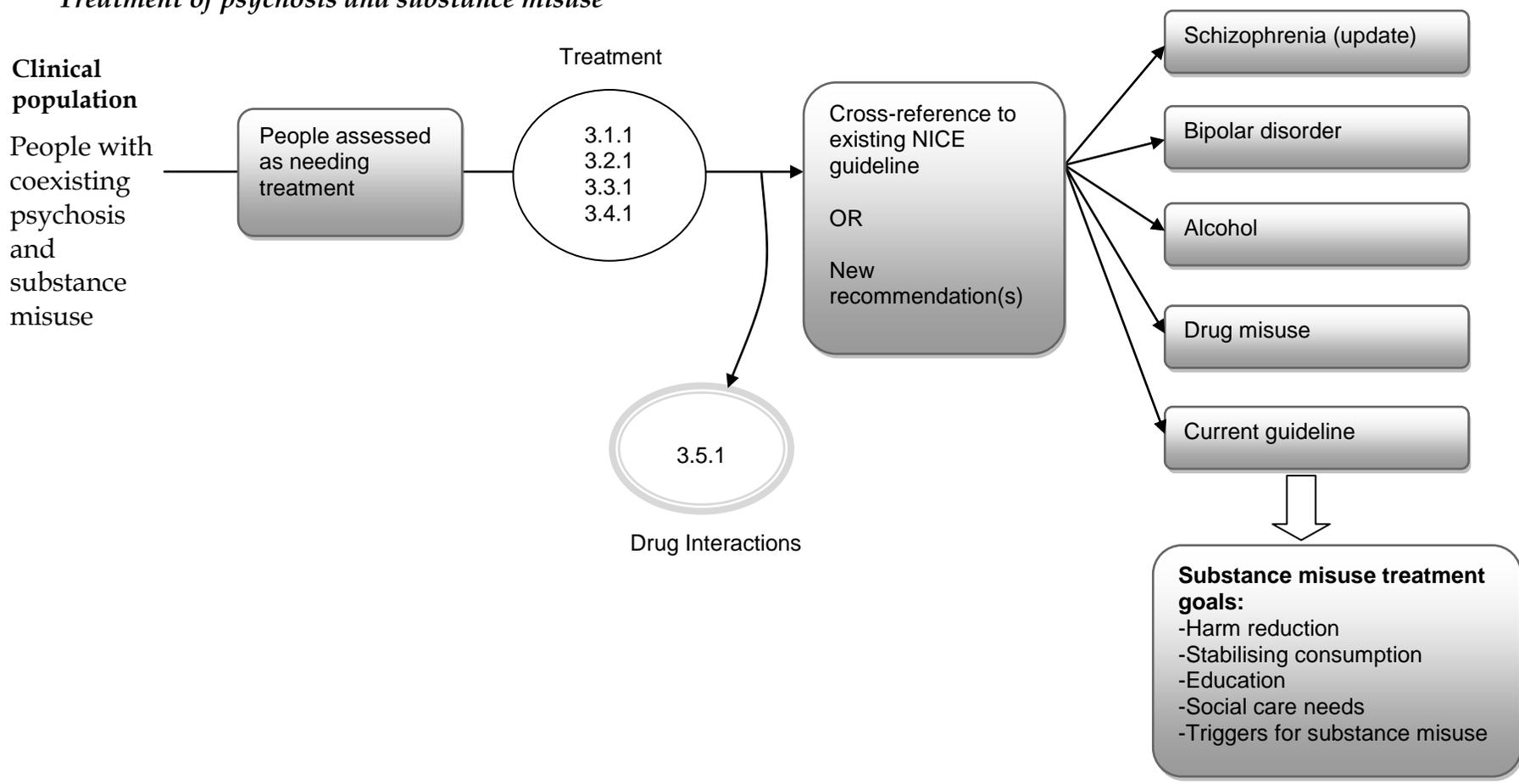
### Care pathways

No.	Primary review questions
1.4.1	In people with psychosis and coexisting substance misuse, what is the most appropriate care pathway (involving all NHS and non-NHS providers) and referral guidance at each transition?

### Experience of care

No.	Primary review questions
1.5.1	For people with psychosis and coexisting substance misuse, what is the experience of diagnosis, access to services, and treatment?
1.5.2	For families and carers of people who have psychosis and coexisting substance misuse, what is the experience of caring for people with psychosis and coexisting substance misuse, and what support is available for families and carers?

*Treatment of psychosis and substance misuse*



### Medication for psychosis

No.	Primary review question
2.1.1	<p>For people with psychosis and coexisting substance misuse, should the medical treatment of their psychosis be modified as a result of substance misuse and the treatment provided (for example, methadone, buprenorphine etc)?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>Sub-question 1: Are there sub-groups of people (for example, young people, people with a particular type of psychosis, BME groups) that may benefit from alternative strategies?</p>

### Psychological/ psychosocial interventions for psychosis

No.	Primary review question
2.2.1	<p>For people with psychosis and coexisting substance misuse, should the psychological/ psychosocial (family interventions, CBT, arts therapies) treatment of their psychosis be modified as a result of the substance misuse problem and the treatment provided (for example, methadone, Buprenorphine, psychological treatment etc)?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>Sub-question 1: Are there sub-groups of people (for example, young people, people with a particular type of psychosis, BME groups) that may benefit from alternative strategies?</p>

### Medication/physical interventions for substance misuse

No.	Primary review question
2.3.1	<p>For people with psychosis and coexisting substance misuse, should the medical/physical treatment of substance misuse be modified as a result of the presence of psychosis and the treatment provided (for example, antipsychotics, lithium)?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>Sub-question 1: Are there sub-groups of people (for example, young people, people with a particular type of psychosis, BME groups) that may benefit from alternative strategies?</p>

### Psychological/ psychosocial interventions for substance misuse

No.	Primary review question
2.4.1	<p>For people with psychosis and coexisting substance misuse, should psychological/ psychosocial treatment for substance misuse be modified as a result of the presence of psychosis and the treatment provided?</p> <p>A) During the acute phase B) During non-acute phase</p> <p>If so, how should treatment be modified?</p> <p>Sub-question 1: Are there sub-groups of people (for example, young people, people with a particular type of psychosis, BME groups) that may benefit from alternative strategies?</p> <p>Sub-question 2: Should interventions be matched to stages of the treatment process (i.e. engagement, persuasion, active treatment, relapse prevention)?</p>

### Drug interactions

No.	Primary review question
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2.5.1	In people with psychosis and substance misuse, is there any evidence that the management of drug interactions or adverse effects from pharmacological treatments should be different from those people without coexisting disorders?  If so, how should management of drug interactions be modified?
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## APPENDIX 7: SEARCH STRATEGIES FOR THE IDENTIFICATION OF CLINICAL STUDIES

### 7.1 Search strategies

The search strategies should be referred to in conjunction with information set out in Section 3.5.2. Each search was constructed using the groups of terms as set out in Table 34. The full set of search terms is documented in sections 7.1.1 to 7.1.3 in Appendix 7. Each search was initially developed for Medline before being translated for use in other databases/interfaces.

**Table 34: Summary of systematic search strategies**

#### Search strategy construction

##### Psychological/psychosocial interventions

Updates to Cleary *et al.* (2008) and Cleary *et al.* (2009). Searches were limited to updating the reviews, covering the time period since the searches for the published reviews were last conducted.

*Search dates: 2008 onwards*

- i) (Psychosis with substance misuse terms) AND (RCT filter OR Observational study filter)

##### Service delivery models

[As above]

##### Pharmacological/physical interventions

Search results covering comprising all the above (psychological, service delivery and pharmacological) were merged into one dataset for the period from 2008 onwards to cut back on unnecessary duplication of effort at the sifting stage.

*Search dates: inception of database onwards*

- i) (Psychosis with substance misuse terms) AND (pharmacological terms)

##### Experience of care

*Search dates: 1995 onwards*

- i) (Psychosis with substance misuse terms) AND (experience of care terms) AND (qualitative filter)

ii) (Psychosis with substance misuse terms) AND (experience of care terms - modified to be more precise)

### 7.1.1 Population Search terms

MEDLINE – Ovid SP interface

*\* Search terms for substance misuse were limited to the main drugs associated with the term at the advice of the GDG.*

1. exp psychotic disorders/ or exp affective disorders, psychotic/
2. exp schizophrenia/or "schizophrenia and disorders with psychotic features"/or schizophrenic psychology/
3. ((mental disorders or mentally ill persons) and chronic disease).sh.
4. exp movement disorders/ or (dyskinesias or psychomotor agitation or neuroleptic malignant syndrome).sh.
5. (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$) adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1).ti,ab.
6. (bipolar\$ or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or psychosi\$ or psychotic\$ or schizo\$).ti,ab.
7. (((tardiv\$ and dyskine\$) or akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or (neuroleptic and movement and disorder) or parkinsoni\$ or neuroleptic-induc\$) not (parkinson\$ and disease)).ti,ab.
8. (emergency services, psychiatric or hospitals, psychiatric or psychiatric department, hospital or (mentally ill persons and (inpatients or hospitalization))).sh. or (psychiatric adj2 (admission\$ or admitted or emerg\$ or hospitali\$ or inpatient\$ or in patient\$)).ti,ab.
9. or/1-8
10. comorbidity/ or "diagnosis, dual (psychiatry)"/
11. (comorbid\$ or co morbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$).ti,ab.
12. or/10-11
13. (designer drugs or needle exchange programs or needle sharing or overdose or street drugs or substance abuse detection or substance abuse, intravenous or substance abuse treatment centers or substance-related disorders or substance withdrawal syndrome).sh.

14. (((drug\$1 or polydrug\$ or psychotropic\$ or substance\$) adj3 (abstain\$ or abstinen\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1).ti,ab.
15. ((club or designer or street) adj2 (drug\$ or substance\$)).ti,ab.
16. or/13-15
17. (amphetamine or amphetamine-related disorders).sh.
18. (dextroamphetamine or methamphetamine).sh.
19. (((amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers) adj2 use\$1)).ti,ab.
20. (amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers).ti,ab.
21. or/17-19
22. 20
23. exp cocaine/ or cocaine-related disorders.sh.
24. (((benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine) adj2 use\$1)).ti,ab.
25. (benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine).ti,ab.
26. or/23-24

27. 25
28. (heroin or heroin dependence or opioid-related disorders).sh.
29. (((heroin or diacetylmorphin\$ or diagesil or diamorf or diamorphin\$) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((diamorphin\$ or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan) adj2 use\$1)).ti,ab.
30. (heroin or diacetylmorphin\$ or diagesil or diamorf or diamorphin\$).ti,ab.
31. or/28-29
32. 30
33. (cannabis or marijuana abuse or marijuana smoking).sh.
34. (((bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk) adj2 use\$1)).ti,ab.
35. (bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk).ti,ab.
36. or/33-34
37. 35
38. 9 and 12 and (or/22,27,32,37)
39. 9 and (or/16,21,26,31,36)
40. or/38-39

### 7.1.2 Question specific search strategies

#### a) Psychological/psychosocial interventions

See Table 34 for information for the strategy used to identify psychological/psychosocial evidence.

#### b) Service delivery models

See Table 34 for information for the strategy used to identify evidence for service delivery models.

*c) Pharmacological/physical interventions*

MEDLINE – Ovid SP interface

1. exp antipsychotic agents/
2. (antipsychotic\$ or anti psychotic\$ or (major adj2 (butyrophenon\$ or phenothiazin\$ or tranquil\$)) or neuroleptic\$).ti,ab.
3. (amisulprid\$1 or aminosultoprid\$1 or amisulpirid\$1 or sertol\$1 or socian or solian).ti,ab.
4. (aripiprazol\$1 or abilify or abilitat).ti,ab.
5. (benperidol\$1 or anquil or benperidon\$1 or benzoperidol\$1 or benzperidol\$1 or frenactil\$1 or frenactyl or glianimon\$1 or phenactil\$1).ti,ab.
6. chlorpromazine.sh. or (chlorpromazin\$1 or aminazin\$1 or chlorazin\$1 or chlorderazin\$1 or contomin\$1 or fenactil\$1 or largactil\$1 or propaphenin\$1 or thorazin\$1).ti,ab.
7. chlorprothixene.sh. or (chlorprothixen\$1 or aminasin\$1 or aminasin\$1 or aminazin\$1 or aminazin\$1 or ampliactil\$1 or amplictil\$1 or ancholactil\$1 or chlopromazin\$1 or chlor pz or chlorbromasin\$1 or chlorderazin\$1 or chlorderazin\$1 or chloropromazin\$1 or chlorpromanyl or chlorpromazin\$1 or chlorprotixen\$1 or clorderazin\$1 or clorpromazin\$1 or cloxan or contomin\$1 or elmarin\$1 or fenactil\$1 or hibanil\$1 or hibernal\$1 or hibernol\$1 or klorpromex or largactil\$1 or largactyl or megaphen\$1 or neurazin\$1 or novomazin\$1 or phenathyl or plegomazin\$1 or plegomazin\$1 or proma or promacid\$1 or promactil\$1 or promapar or promazil\$1 or propaphen\$1 or propaphenin\$1 or prozil or psychozin\$1 or sanopron\$1 or solidon\$1 or sonazin\$1 or taractan\$1 or taroctil\$1 or thor prom or thorazen\$1 or thorazin\$1 or torazin\$1 or truxal or vegetamin a or vegetamin b or wintamin\$1 or wintermin\$1 or zuledin\$1).ti,ab.
8. clozapine.sh. or (clozapin\$1 or alemoxan\$1 or azaleptin\$1 or clopine or clozaril\$1 or denzapin\$1 or dorval or dozapin\$1 or fazaclo or froidir or klozapol or lapenax or leponex or wander compound or zaponex).ti,ab.
9. flupenthixol.sh. or (flupentixol\$1 or flupenthixol\$1 or depixol\$1 or emergil\$1 or fluanxol\$1 or flupentixol\$1 or emergil\$1 or fluanxol\$1 or piperazineethanol\$1 or viscoleo).ti,ab.
10. fluphenazine.sh. or (anatensil or anatensol or antasol or dapotum or elinol or flufenazin\$ or flumezin or fluorfenazine or fluphenacin or

- fluphenazin or fluphenazin\$ or fluphenzine or ftorphenazine or luogen depot or lyogen or lyorodin or moditen or moditin or omca or pacinol or permitil or phthorphenazine or prolixan 300 or prolixene or prolixin\$ or sevinal or sevinol or squaline or squalon\$ or siquoline or tensofin or trancin or valamina or vespazin\$).ti,ab.
11. fluspirilene.sh. or (fluspirilen\$1 or fluspi or imap or kivat or redeptin\$1 or spirodiflamin\$1).ti,ab.
  12. haloperidol.sh. or (haloperidol\$1 or aloperidin\$1 or bioperidolo or brotopon or celenase or cerenace or dozic or duraperidol or einalon s or eukystol or fortunans\$1 or haldol or halidol or haloneural\$1 or haloperitol\$1 or halosten or keselan or linton or peluces or serenace or serenase or siegoperidol\$1 or sigaperidol\$1).ti,ab.
  13. methotrimeprazine.sh. or (levomepromazin\$1 or 2 methoxytrimeprazin\$1 or hirnamin\$1 or levo promazin\$1 or levomeprazin\$1 or levopromazin\$1 or levoprom\$1 or mepromazin\$1 or methotrimeprazin\$1 or methotrimperazin\$1 or milezin\$1 or minozinan\$1 or nozin\$1 or neuractil\$1 or neurocil\$1 or nirvan or nosinan\$1 or nozinan\$1 or sinogan or tiserцин\$1 or tizercin\$1 or tizertsin\$1 or veractil\$1).ti,ab.
  14. (olanzapin\$1 or lanzac or midax or olansek or olzapin or rexapin or zalasta or zolafren or zydis or zypadhera or zyprex\$1).ti,ab.
  15. (paliperidon\$1 or 9 hydroxyrisperidon\$1 or invega).ti,ab.
  16. paroxetine.sh. or (paroxetin\$1 or aropax or deroxat or motivan or paxil\$1 or pexeva or seroxat or tagonis).ti,ab.
  17. (pericyazin\$1 or aolect or neulactil\$1 or neuleptil\$1 or periciazin\$1 or properciazin\$1 or propericiazin\$1).ti,ab.
  18. perphenazine.sh. or (perphenazin\$1 or chlorperphenazin\$1 or chlorpiprazin\$1 or chlorpiprozin\$1 or decentan\$1 or etaperazin\$1 or ethaperazin\$1 or etrafon or fentazin\$1 or perfenazin\$1 or perfenazin\$1 or perferazin\$1 or perphenan\$1 or perphenezin\$1 or thilatazin\$1 or tranquisan\$1 or triavail or trifalon\$1 or trilafan\$1 or trilafon\$1 or trilifan\$1 or triliphan\$1).ti,ab.
  19. pimozide.sh. or (pimozid\$1 or antalon\$1 or opiran\$1 or orap or pimocid\$1 or pimorid\$1 or pinozid\$1).ti,ab.
  20. prochlorperazine.sh. or (prochlorperazin\$1 or buccastem or capazin\$1 or chlormeprazin\$1 or chlorpeazin\$1 or chlorperazin\$1 or compazin\$1 or dicopal\$1 or emelent or kronocin\$1 or meterazin\$1 or metherazin\$1 or nipodal\$1 or phenotil or prochlor perazin\$1 or prochlorpemazin\$1 or prochlorperacin\$1 or prochlorperzin\$1 or prochlorpromazin\$1 or proclorperazin\$1 or stemetil or stemzine or tementil\$1 or temetil\$1).ti,ab.

21. promazine.sh. or (promazin\$1 or alofen\$1 or alophen\$1 or ampazin\$1 or amprazim\$1 or centractyl or delazin\$1 or esparin\$1 or lete or liranol\$1 or neo hibernex or neuroplegil\$1 or piarin\$1 or prazin\$1 or pro tan or promantin\$1 or promanyl\$1 or promilen\$1 or promwill or protactil\$1 or protactyl\$1 or romthiazin\$1 or romtiazin\$1 or sediston\$1 or sinophenin\$1 or sparin\$1 or tomil or varophen\$1 or verophen\$1).ti,ab.
22. (quetiapin\$1 or ketipinor or quepin or seroquel or tienapin\$1).ti,ab.
23. risperidone.sh. or (risperidon\$1 or belivon\$1 or ridal or riscalin or risolept or rispen or risperdal\$1 or sizodon).ti,ab.
24. (sertindol\$1 or indole or serdolect or serlect).ti,ab.
25. sulpiride.sh. or (sulpirid\$1 or abilit or aiglonyl\$1 or arminol\$1 or bosnyl or deponerton\$1 or desisulpid\$1 or digton or dobren or dogmatil\$1 or dogmatyl or dolmatil\$1 or eglonyl or ekilid or equilid or guastil\$1 or isnamid\$1 or leboprid\$1 or levopraid or levosulpirid\$1 or meresa or miradol\$1 or modal or neogama or pontirid\$1 or psicocen\$1 or sulfirid\$1 or sulp\$1 or sulperid\$1 or sulpitil\$1 or sulpivert or sulpor or sulpyride or synedil\$1 or tepavil\$1 or vertigo meresa or vertigo neogama or vipral).ti,ab.
26. trifluoperazine.sh. or (trifluoperazin\$1 or apotrifluoperazine\$1 or calmazin\$1 or dihydrochlorid\$1 or eskazin\$1 or eskazin\$1 or eskazinyl or fluoperazin\$1 or flupazin\$1 or jatroneural\$1 or modalina or stelazin\$1 or terfluzin\$1 or terfluzin\$1 or trifluoperazid\$1 or trifluoperazin\$1 or trifluoperzin\$1 or trifluoroperazin\$1 or trifluorperacin\$1 or trifluperazin\$1 or triflurin\$1 or triftazin\$1 or triftazinum or triphthazin\$1 or triphthasin\$1 or triphthazin\$1).ti,ab.
27. (zotepin\$1 or lodopin\$1 or losizopilon or nipolept or setous or zoleptil).ti,ab.
28. clopenthixol.sh. or (zuclopenthixol\$1 or acuphase or clopenthixol\$1 or clopixol or cisordinol\$1 or sedanaxol\$1).ti,ab.
29. or/1-28
30. exp serotonin uptake inhibitors/
31. (ssri\$ or ((serotonin or 5 ht or 5 hydroxytryptamine) adj (uptake or reuptake or re uptake) adj inhibit\$)).ti,ab.
32. citalopram.sh. or (celexa or cipramil\$1 or cytalopram or elopram or escitalopram or lexapro or nitalapram or sepram or seropram).ti,ab.
33. (escitalopram or cipralax or lexapro or seroplex).ti,ab.
34. fluoxetine.sh. or (fluoxetin\$1 or fluctin\$1 or flunirin\$1 or fluoxifar or prosac or prozac or prozamin\$1 or sarafem or symbyax).ti,ab.
35. fluvoxamine.sh. or (fluvoxamin\$1 or depromel\$1 or desiflu or dumirox or faverin\$1 or fevarin\$1 or floxyfral\$1 or fluoxamin\$1 or fluoxamin\$1 or fluvoxadura or luvox).ti,ab.

36. (nefazadon\$1 or dutonin\$1 or nefadar or reseril\$1 or serzon\$1).ti,ab.
37. paroxetine.sh. or (paroxetin\$1 or aropax or deroxat or motivan\$1 or paxil or pexeva or seroxat or tagonis).ti,ab.
38. sertraline.sh. or (sertralin\$1 or altrulin\$1 or aremis or besitran\$1 or gladem or lustral\$1 or naphthylamin\$1 or sealdin\$1 or serad or serlain\$1 or tresleen or zoloft).ti,ab.
39. or/30-38
40. benzodiazepines.sh.
41. (benzo\$1 or benzodiazepin\$).ti,ab.
42. diazepam.sh. or (diazepam or alupram or ansiolin\$1 or antenex or apaurin\$1 or apaurin\$1 or apozepam or assival\$1 or audium\$1 or bialzepam or bialzegan\$1 or calmpos\$1 or cercin\$1 or cersin\$1 or chlordiazepam or dialar or diastat or diazeliium or diazemuls or diazidem or ducen\$1 or duxen\$1 or eridan or eurosan\$1 or evacalm\$1 or fanstan\$1 or faustan\$1 or gewacalm\$1 or lamra or lembrol\$1 or lipodiazepam or lorinon\$1 or methyl Diazepinon\$1 or methyl Diazepinon\$1 or morosan\$1 or neocalm\$1 or neurolytril\$1 or noan or novazam or paceum or plidan or psychopax or relanium or rimapam or sedapam or seduxen\$1 or serendin\$1 or setonil\$1 or sibazon\$1 or sonacon\$1 or stesolid\$1 or stesolin\$1 or tanquo tablinen\$1 or tensium or tranimul\$1 or tranquo puren or umbrium\$1 or valaxon\$1 or valclair or valiquid\$1 or valium or valpam or valreleas\$1 or vatran\$1 or vival\$1 or vivol4 or zetran\$1).ti,ab.
43. lorazepam.sh. or (lorazepam or almazin\$1 or alzapam or apolorazepam or ativan or bonatranquan\$1 or donix or duralozam or durazolam or idalprem or kendol\$1 or laubeel or lorabenz or loranas\$1 or loranz\$1 or lorans or lorax or lorazep von ct or lorigem\$1 or lorivan\$1 or mesmerin\$1 or novo lorazem\$1 or novolorazem\$1 or novo lorazem\$1 or nu loraz or nuloraz or orfidal or orifadal\$1 or pro dorm or quait or securit or sedicepan\$1 or sinestron\$1 or somagerol\$1 or tavor or temesta or tolid or wypax).ti,ab.
44. narcotic antagonists.sh.
45. ((narcotic\$ or opiate\$ or opioid\$) adj antagonist\$).ti,ab.
46. naltrexone.sh. or (antaxone or celupan or depade or nalorex or naltrel or naltrexone\$ or nemexin or opizone or revia or trexan or vivitrex or vivitrol).ti,ab.
47. (arthene or cervene or cessalor incystene or nalmefene or nalmetrene or revex or soberal).ti,ab.
48. or/40-47
49. (analgesics, opioid or opiate agonist or partial agonist).sh.
50. ((narcotic\$ or opiate\$ or opioid\$ or partial\$) adj2 (agonist\$ or analg?esi\$)).ti,ab.

51. exp methadone/ or (adanon or algidon or algolysin or algoxale or althose or amidon or amidone or amidosan or anadon or biodone or butalgin or deamin or depridol or diaminion or dianone or dolafin or dolamid or dolesone or dolophine or dorex or dorexol or fenadon or heptadon or heptanon or ketalgin or linctus or mecodin or mepecton or mephenon or metadol or metasedin or methaddict or methadon or methadone or methadose or methex or miadone or moheptan or phenadon or phenadone or phymet or physepton or physeptone or physeptone or pinadone or polamidon or polamivet or polamivit or sinalgin or symoron).ti,ab.
52. buprenorphine.sh. or (buprenex or buprenorphin\$ or buprex or finibron or lepetan or prefin or suboxone or subutex or temgesic or transtec).ti,ab.
53. or/49-52
54. adrenergic alpha-agonists.sh.
55. ((adrenergic alpha or alpha adrenergic) adj2 agonist\$).ti,ab.
56. (lofexidin\$ or britlofex or lofetensin or loxacor).ti,ab.
57. clonidine.sh. or (arkamin\$1 or caprysin\$1 or catapres or catapresan\$1 or catapressan or catapressant or catasan\$1 or chlofazolin\$1 or chlophazolin\$1 or chlophelin\$1 or clinidin\$1 or clofelin\$1 or clofelin\$1 or clofenil\$1 or clomidin\$1 or clondin\$1 or clonidin\$1 or clonistada or clonnirit or clophelin\$1 or clopheline or dcai or dichlorophenylaminoimidazoline or dixarit or duraclon or gemiton or haemiton or hemiton or isoglaucan or klofelin or klofenil or normopresan or paracefan or tenso timelets).ti,ab.
58. or/54-57
59. disulfiram.sh. or (abstensil\$1 or abstynyl or alcophobin\$1 or antabus or antabuse or antadix or antaethan\$1 or antaethyl or antiaethan\$1 or anticol\$1 or antietanol\$1 or aversan or contralin\$1 or contrapot or cronetal\$1 or dicupral or disulfid\$1 or disulfiram or disulfizam or disulphiram or espenal or esperal or etabus or ethyl thiurad or exhorran or hoca or stopethyl or stopetyl or teraetil or tetra ethyl thiuramdisulfide or tetradin\$1 or tetraethylthiuram or tetraetil\$1 or teturam or teturamin or thiuram or thiuranide or tiuram or ttd).ti,ab.
60. (acamprosate or aotal or calcium acetylhomotaurinate or campral or n acetylhomotaurine calcium).ti,ab.
61. chlormethiazole.sh. or (chlomethiazol\$ or chlorethiazol\$ or chlormethiazol\$ or clomethiazol\$ or distraneurin or distraneurin\$ or hemineurin\$1 or heminevrin\$1 or hemithiamin\$ or zendra).ti,ab.
62. chlordiazepoxide.sh. or (a poxide or ansiacal or benzodiapin\$1 or cebrum or chlordiazepoxid\$ or chlordiazepoxyd\$1 or chlorodiazepoxid\$1 or chlozepid\$1 or clopoxid\$1 or contol or

- decacil\$1 or defobin\$1 or disarim or dizepin\$1 or dopoxid\$1 or droxol\$1 or eden psich or elenium or elenum or equibral or kalmocaps or labican or librelease or libritabs or librium or lipoxide or mesural or metaminodiazepoxide or methaminodiazepoxide or mildmen or mitran or multum or murcil or napoton\$1 or novosed or o c m or ocm 505 or psichial or psicosan or psicoterina or radepur or reliberan or repositans 10 or risolid or seren vita or servium or silibrin or sk lygen or sonimen or timosin or tropium or viansin or viopsicol).ti,ab.
63. or/59-62
64. anticonvulsants.sh.
65. (anticonvuls\$ or anti convuls\$ or antiepilep\$ or anti epilep\$).ti,ab.
66. (epitomax or topamax or topamax or sprinkle or topamax or topimax or topirimate or topiramate).ti,ab.
67. valproic acid.sh. or (2 propylpentanoate or 2 propylpentanoic acid or 2 propylvalerate sodium or 2 propylvaleric acid or alpha propylvalerate or alpha propylvaleric acid or apilepsin\$1 or convulex or convulsofin\$1 or depacon\$1 or depaken\$1 or depakin\$1 or depakot\$1 or deprakin\$1 or di n propylacetate or di n propylacetate sodium or di n propylacetic acid or dipropyl acetic acid or dipropylacetate or dipropylacetatic acid or dipropylacetic acid or diprosin\$1 or divalproex or epilim or ergenyl or everiden\$1 or goilim or labazen\$1 or leptilan\$1 or leptilanil or mylproin or myproic acid or n dipropylacetic acid or orfiril or orlept or propymal or sodium 2 propylpentanoate or sodium 2 propylvalerate or sodium di n propyl acetate or sodium di n propylacetate or sodium dipropyl acetate or sodium dipropylacetate or sodium n dipropylacetate or stavzor or valerin\$1 or valparin\$1 or valpro or valproate or valproic acid or vupral\$1).ti,ab.
68. carbamazepine.sh. or (amizepin\$1 or atretol\$1 or biston or calepsin\$1 or carbagen\$1 or carbama or carbamaze or carbamazepin\$1 or carbategral or carbatrol or convuline or degranol or epimaz or epimax or epitol or equetro or finlepsin\$1 or hermolepsin\$1 or lexin or mazepin\$1 or neurotol or neurotop or servimazepin\$1 or sirtal or stazepin\$1 or tegral or tegretal or tegretol or tegrital or telesmin\$1 or teril or timonil or trimonil).ti,ab.
69. or/64-68
70. neuromuscular agents.sh.
71. ((neuromuscular or skeletal muscle) adj (agent\$ or drug\$ or relaxant\$)).ti,ab.
72. baclofen.sh. or (apobaclofen\$1 or atrofен\$1 or baclofen\$ or baclofeneirex or baclofene-irex or baclophen or baclospas or beta 4 chlorophenyl 4 aminobutanoic acid or beta amino methyl

chlorohydrocinnamic acid or beta aminomethyl para  
 chlorohydrocinnamic acid or beta para chlorophenyl gamma  
 aminobutyric acid or chlorophenyl gaba or clofen or genbaclofen or  
 genpharm or kemstro or lioresal or intralcal or lebic or lioresal or  
 lioresal or lioresyl or lyflex or nu baclo or nubaclo or pcp-gaba or  
 pmsbaclofen).ti,ab.

73. or/70-72

74. lithium\$.sh. or (lithium or camcolit or candamid\$1 or carbolith or  
 carbolitium or cibalith s or contemnol\$1 or dilithium or eskalith or  
 hypnorex or li salt or limas or linthane or liskonium or liskonum or  
 litarex or lithane or lithiofor or lithionit or lithiophor or lithobid or  
 lithocarb or lithonate or lithotabs or maniprex or mesin or micalith or  
 neurolepsin or neurolithium or plenur or priadel or quilinormretard or  
 quilonorm or quilonum or teralithe or theralite or theralithe).ti,ab.

75. or/1-74

*d) Experience of care*

*Due to the difficulties of identifying qualitative research with precision from  
 bibliographic databases, search request #15 was generated without the use of a  
 qualitative filter.*

MEDLINE – Ovid SP interface

1. (consumer participation or consumer satisfaction or health behavior or  
 hospital patient relations or medication adherence or nurse patient  
 relations or patient acceptance of health care or patient advocacy or  
 patient compliance or patient participation or patient preference or  
 physician patient relations or professional patient relations or public  
 opinion or treatment refusal).sh.
2. (attitude or attitude to health or knowledge, attitudes, practice or  
 patient satisfaction).sh.
3. (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$)  
 adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1 or bipolar\$  
 or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or  
 hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or  
 psychosi\$ or psychotic\$ or schizo\$ or (tardiv\$ and dyskine\$) or  
 akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or  
 (neuroleptic and movement and disorder) or parkinsoni\$ or  
 neuroleptic-induc\$ or (psychiatric adj2 (admission or admitted or  
 emerg\$ or hospitali\$ or inpatient\$ or in patient\$)) or ((mental\$ or

psych\$) adj (disease\$ or disorder\$ or illness\$) or comorbid\$ or comorbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$ or ((drug\$1 or polydrug\$ or substance\$) adj3 (abstain\$ or abstinen\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or overdos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1 or ((club or designer or street) adj2 (drug\$ or substance\$)) or amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers or benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine or diacetylmorphin\$ or diamorphin\$ or diagesil or diamorf or heroin or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan or bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk or polydrug\$) adj8 (acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) adj8 (adult\$1 or attender\$ or client\$ or consumer\$ or individuals or inpatient\$ or men or minorities or outpatient\$ or participant\$ or patient\$ or people or population or public or subjects or survivor\$ or women or user\$ or care giver\$ or caregiver\$ or carer\$ or (care adj (manager\$ or worker\$)) or family or families)).ti,ab.

4. (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$) adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1 or bipolar\$ or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or psychosi\$ or psychotic\$ or schizo\$ or (tardiv\$ and dyskine\$) or akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or (neuroleptic and movement and disorder) or parkinsoni\$ or neuroleptic-induc\$ or (psychiatric adj2 (admission or admitted or emerg\$ or hospitali\$ or inpatient\$ or in patient\$)) or ((mental\$ or psych\$) adj (disease\$ or disorder\$ or illness\$)) or comorbid\$ or co morbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$ or ((drug\$1 or polydrug\$ or substance\$) adj3 (abstain\$ or abstinens\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or overdos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1 or ((club or designer or street) adj2 (drug\$ or substance\$)) or amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers or benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine or diacetylmorphin\$ or diamorphin\$ or diagesil or diamorf or heroin or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan or bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk or polydrug\$) and (acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or literacy or narrat\$ or knowledge or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism

- or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) and (adult\$1 or attender\$ or client\$ or consumer\$ or individuals or inpatient\$ or men or minorities or outpatient\$ or participant\$ or patient\$ or people or population or public or subjects or survivor\$ or women or user\$ or care giver\$ or caregiver\$ or carer\$ or (care adj (manager\$ or worker\$)) or family or families)).ti.
5. (((mental\$ or psych\$ or psychiatric) adj2 (disease\$ or disorder\$ or distress or health or ill or problem\$)) and (acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) and (adult\$1 or attender\$ or client\$ or consumer\$ or individuals or inpatient\$ or men or minorities or outpatient\$ or participant\$ or patient\$ or people or population or public or subjects or survivor\$ or women or user\$ or care giver\$ or caregiver\$ or carer\$ or (care adj (manager\$ or worker\$)) or family or families)).ti.
6. (((mental\$ or psych\$ or psychiatric) adj2 (disease\$ or disorder\$ or distress or health or ill or problem\$)) and (acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) and (care or healthcare or health care or medication or service\$ or therap\$ or treatment\$)).ti.

7. ((acceptance or account\$1 or adhere\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) adj3 (adult\$1 or attender\$ or client\$ or consumer\$ or individuals or inpatient\$ or men or minorities or outpatient\$ or participant\$ or patient\$ or people or population or public or subjects or survivor\$ or women or user\$ or care giver\$ or caregiver\$ or carer\$ or (care adj (manager\$ or worker\$)) or family or families)).ti.
8. ((acceptance or account\$1 or adhere\$ or aspiration\$ or attitude\$ or aversion\$ or barrier\$ or belief\$ or centredness or communicat\$ or complianc\$ or conception\$ or concern\$1 or content\$ or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or experience\$1 or engaging or engage\$1 or happy or help\$ or idea\$1 or incentive\$ or interview\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or preference\$ or refus\$ or research or satisf\$ or scepticism or service\$ use\$ or stigma or story or stories or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) adj2 (client\$ or consumer\$ or inpatient\$ or minorities or outpatient\$ or patient\$ or people or public or survivor\$ or user\$)).ti,ab.
9. ((acceptance or account\$1 or adhere\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$)

adj4 (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$) adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1 or bipolar\$ or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or psychosi\$ or psychotic\$ or schizo\$ or (tardiv\$ and dyskine\$) or akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or (neuroleptic and movement and disorder) or parkinsoni\$ or neuroleptic-induc\$ or (psychiatric adj2 (admission or admitted or emerg\$ or hospitali\$ or inpatient\$ or in patient\$)) or ((mental\$ or psych\$) adj (disease\$ or disorder\$ or illness\$)) or comorbid\$ or co morbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$ or ((drug\$1 or polydrug\$ or substance\$) adj3 (abstain\$ or abstinens\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or overdos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1 or ((club or designer or street) adj2 (drug\$ or substance\$)) or amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers or benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine or diacetylmorphin\$ or diamorphin\$ or diagesil or diamorf or heroin or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan or bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk or polydrug\$)).ti.

10. (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$) adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1 or bipolar\$ or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or psychosi\$ or psychotic\$ or schizo\$ or (tardiv\$ and dyskine\$) or akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or (neuroleptic and movement and disorder) or parkinsoni\$ or neuroleptic-induc\$ or (psychiatric adj2 (admission or admitted or emerg\$ or hospitali\$ or inpatient\$ or in patient\$)) or ((mental\$ or

psych\$) adj (disease\$ or disorder\$ or illness\$)) or comorbid\$ or co morbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$ or ((drug\$1 or polydrug\$ or substance\$) adj3 (abstain\$ or abstinen\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1 or ((club or designer or street) adj2 (drug\$ or substance\$)) or amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers or benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine or diacetylmorphin\$ or diamorphin\$ or diagesil or diamorf or heroin or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan or bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk or polydrug\$) and (acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$) and (care or healthcare or health care or medication or service\$ or therap\$ or treatment\$)).ti.

11. ((acceptance or account\$1 or adher\$ or aspiration\$ or attitude\$ or aversion\$ or awareness or barrier\$ or belief\$ or centredness or choice\$ or cognitions or complianc\$ or conception\$1 or concern\$1 or confus\$ or content\$ or diary or diaries or demand\$ or disatisf\$ or disclos\$ or discontent\$ or disgruntle\$ or engaging or engage\$1 or experienc\$ or

feeling or happy or help\$ or incentive\$ or involv\$ or knowledge or literacy or narrat\$ or need or needs or nonadher\$ or obstacle\$ or opinion\$ or participa\$ or perception\$ or perceived or perspective\$ or position\$ or prefer or preferred or preference\$ or persistence or refus\$ or satisf\$ or scepticism or selfobservat\$ or self observat\$ or (service\$ adj2 use\$) or stigma\$ or story or stories or support\$ or tolerance or understand\$ or unhappy or utili?ation or view\$ or willing\$ or voice\$ adj4 (care or healthcare or medication\$ or psychotherapy\$ or service\$ or therap\$ or treatment\$ or ((perceived or perception\$ or unmet\$) adj need\$) )) .ti,ab.

12. caregivers/or exp disabled persons/or mentally ill persons/ or inpatients/or outpatients/or survivors/or (consumer\$ or patient\$).hw.
13. (adult\$1 or attender\$ or client\$ or consumer\$ or individuals or inpatient\$ or men or minorities or outpatient\$ or participant\$ or patient\$ or people or public or subjects or survivor\$ or women or user\$ or care giver\$ or caregiver\$ or carer\$ or (care adj (manager\$ or worker\$)) or family or families).ti,ab. or (population or sample).ti.
14. or/12-13
15. or/2,4,5,7 or (or/6,9,10 and 14)
16. or/1,3,8 or (11 and 14)

### 7.1.3 Search filters

*a) Randomised controlled trial search filter – this is an adaptation of a filter designed by the Health Information Research Unit of the McMaster University, Ontario.*

MEDLINE – Ovid SP interface

1. exp clinical trial/ or cross over studies/ or double blind method/ or random allocation/ or randomized controlled trials as topic/ or single blind method/
2. (clinical adj2 trial\$).ti,ab.
3. (crossover or cross over).ti,ab.
4. (((single\$ or doubl\$ or trebl\$ or tripl\$) adj5 blind\$) or mask\$ or dummy or doubleblind\$ or singleblind\$ or trebleblind\$ or tripleblind\$).ti,ab.
5. (placebo\$ or random\$).mp.
6. (clinical trial\$ or controlled clinical trial\$ or random\$).pt.
7. animals/ not (humans/or human\$.ti,ab.)
8. (or/1-6) not 7

*b) Observational studies filter – developed in-house*

MEDLINE – Ovid SP interface

1. exp case control studies/ or exp cohort studies/ or cross sectional studies/ or epidemiologic study characteristic as topic/ or epidemiologic studies/
2. case reports.pt.
3. ((cross sectional or epidemiologic\$ or observational) adj (study or studies)).ti,ab.
4. (case control\$ or cohort\$1 or cross sectional or followup\$ or follow up\$ or followed or longitudinal or prospective\$ or retrospective\$).ti,ab.
5. or/1-4

*c) Qualitative filter – this is an adaptation of filters designed by the Health Information Research Unit of McMaster University, Ontario, and the University of Alberta.*

MEDLINE – Ovid SP interface

1. qualitative research/
2. interview/ or personal narratives/ or exp interviews as topic/ or interview, psychological/
3. narration/
4. exp tape recording/ or videodisc recording/
5. sampling studies/ or cluster analysis/
6. anthropology, cultural/
7. nursing methodology research/
8. observation/
9. (qualitative or ethno\$ or emic or etic or heuristic or semiotics or phenomenolog\$).ti,ab.
10. interview\$.ti,ab.
11. (((audio or tape or video\$) adj5 record\$) or audiorecord\$ or taperecord\$ or videorecord\$ or videotap\$).ti,ab.
12. (story or stories or storytell\$ or story tell\$).ti,ab.
13. testimon\$.ti,ab.
14. ((focus adj4 (group\$ or sampl\$)) or narrat\$ or ((life or lived) adj experience\$)).ti,ab.
15. ((participant\$ or nonparticipant\$) adj3 observ\$).ti,ab.
16. (constant adj (comparative or comparison)).ti,ab.

17. (content analy\$ or (field adj (note\$ or record\$ or stud\$ or research)) or fieldnote\$).ti,ab.
18. (data adj1 saturat\$).ti,ab.
19. discourse analys?s.ti,ab.
20. (grounded adj (theor\$ or study or studies or research)).ti,ab.
21. (hermeneutic\$ or heidegger\$ or husserl\$ or colaizzi\$ or giorgi\$ or glaser or spiegelberg\$ or strauss).ti,ab.
22. (maximum variation or snowball).ti,ab.
23. (cross case analys\$ or metaethno\$ or meta ethno\$ or metanarrative\$ or meta narrative\$ or metasynthes\$ or meta synthes\$ or metasummar\$ or meta summar\$ or metastud\$ or meta stud\$ or qualitative synthes\$ or qualitative overview or metaoverview or meta overview).ti,ab.
24. purpos\$ sampl\$.ti,ab.
25. ((structured or unstructured) adj1 categor\$).ti,ab.
26. ((thematic\$ adj3 analys\$) or themes).ti,ab.
27. (theoretical sampl\$ or ricoeur or spiegelberg\$ or merleau).ti,ab.
28. (van kaam\$ or van manen or constant compar\$).ti,ab.
29. action research.ti,ab.
30. human science.ti,ab.
31. (critical social\$ or ethical enquiry or (pilot testing and survey) or shadowing or ((philosophical or social) adj research\$)).ti,ab.
32. or/1-31

## APPENDIX 8: METHODOLOGY CHECKLIST TEMPLATE FOR CLINICAL STUDIES AND REVIEWS

The methodological quality of each study was evaluated using NICE checklists (NICE, 2009b). The checklists for systematic reviews and for RCTs are reproduced below (for other checklists and further information about how to complete each checklist, see *The Guidelines Manual* [NICE, 2009b]).

### Methodology checklist: systematic reviews and meta-analyses

Study identification <i>Include author, title, reference, year of publication</i>	
Guideline topic:	Review question no:
Checklist completed by:	
<b>SCREENING QUESTIONS</b>	
In a well-conducted, relevant systematic review:	<i>Circle one option for each question</i>
The review addresses an appropriate and clearly focused question that is relevant to the guideline review question	Yes      No      Unclear
The review collects the type of studies you consider relevant to the guideline review question	Yes      No      Unclear
The literature search is sufficiently rigorous to identify all the relevant studies	Yes      No      Unclear
Study quality is assessed and reported	Yes      No      Unclear
An adequate description of the methodology used is included, and the methods used are appropriate to the question	Yes      No      Unclear

**Methodology checklist: randomised controlled trials**

Study identification Include author, title, reference, year of publication		
Guideline topic:		Review question no:
Checklist completed by:		
		<i>Circle one option for each question</i>
<b>A. Selection bias (systematic differences between the comparison groups)</b>		
A1	An appropriate method of randomisation was used to allocate participants to treatment groups (which would have balanced any confounding factors equally across groups)	Yes No Unclear N/A
A2	There was adequate concealment of allocation (such that investigators, clinicians and participants cannot influence enrolment or treatment allocation)	Yes No Unclear N/A
A3	The groups were comparable at baseline, including all major confounding and prognostic factors	Yes No Unclear N/A
Based on your answers to the above, in your opinion was selection bias present? If so, what is the likely direction of its effect?		
<p style="text-align: center;"> <span style="margin-right: 100px;">Low risk of bias</span> <span style="margin-right: 100px;">Unclear/unknown risk</span> <span>High risk of bias</span> </p>		
Likely direction of effect:		
<b>B. Performance bias (systematic differences between groups in the care provided, apart from the intervention under investigation)</b>		
B1	The comparison groups received the same care apart from the intervention(s) studied	Yes No Unclear N/A
B2	Participants receiving care were kept 'blind' to treatment allocation	Yes No Unclear N/A
B3	Individuals administering care were kept 'blind' to treatment allocation	Yes No Unclear N/A
Based on your answers to the above, in your opinion was performance bias present? If so, what is the likely direction of its effect?		

Low risk of bias		Unclear/unknown risk	High risk of bias	
Likely direction of effect:				
C. Attrition bias (systematic differences between the comparison groups with respect to loss of participants)				
C1	All groups were followed up for an equal length of time (or analysis was adjusted to allow for differences in length of follow-up)	Yes	No	Unclear N/A
C2	a. How many participants did not complete treatment in each group?			
	b. The groups were comparable for treatment completion (that is, there were no important or systematic differences between groups in terms of those who did not complete treatment)	Yes	No	Unclear N/A
C3	a. For how many participants in each group were no outcome data available?			
	b. The groups were comparable with respect to the availability of outcome data (that is, there were no important or systematic differences between groups in terms of those for whom outcome data were not available).	Yes	No	Unclear N/A
Based on your answers to the above, in your opinion was attrition bias present? If so, what is the likely direction of its effect?				
Low risk of bias		Unclear/unknown risk	High risk of bias	
Likely direction of effect:				
D. Detection bias (bias in how outcomes are ascertained, diagnosed or verified)				
D1	The study had an appropriate length of follow-up	Yes	No	Unclear N/A
D2	The study used a precise definition of outcome	Yes	No	Unclear N/A
D3	A valid and reliable method was used to determine the outcome	Yes	No	Unclear N/A
D4	Investigators were kept 'blind' to participants' exposure to the intervention	Yes	No	Unclear N/A

D5	Investigators were kept 'blind' to other important confounding and prognostic factors	Yes No Unclear N/A
Based on your answers to the above, in your opinion was detection bias present? If so, what is the likely direction of its effect?		
Low risk of bias	Unclear/unknown risk	High risk of bias
Likely direction of effect:		

## APPENDIX 9: SEARCH STRATEGIES FOR THE IDENTIFICATION OF HEALTH ECONOMICS EVIDENCE

### 10.1 Search strategies

The search strategies should be referred to in conjunction with information set out in Section 3.6.1.

For standard mainstream bibliographic databases (CINAHL, EMBASE, MEDLINE and PsycINFO) search terms for psychosis with substance abuse were combined with a search filter for health economic studies. For searches generated in topic-specific databases (HTA, NHS EED) search terms on psychosis with substance abuse were used without a filter. The search strategies were initially developed for Medline before being translated for use in other databases/interfaces.

#### 10.1.1 Population Search terms

MEDLINE – Ovid SP interface

*\* Search terms for substance misuse were limited to the main drugs associated with the term at the advice of the GDG.*

1. exp psychotic disorders/ or exp affective disorders, psychotic/
2. exp schizophrenia/or "schizophrenia and disorders with psychotic features"/or schizophrenic psychology/
3. ((mental disorders or mentally ill persons) and chronic disease).sh.
4. exp movement disorders/ or (dyskinesias or psychomotor agitation or neuroleptic malignant syndrome).sh.
5. (((acute or chronic\$ or serious\$ or sever\$) adj3 (mental\$ or psych\$) adj3 (disease\$ or disorder\$ or disturbanc\$ or ill\$)) or smi\$1).ti,ab.
6. (bipolar\$ or ((cyclothymi\$ or rapid or ultradian) adj5 cycl\$) or rcbd or hebephreni\$ or mania\$ or manic\$ or oligophreni\$ or psychose\$ or psychosi\$ or psychotic\$ or schizo\$).ti,ab.
7. (((tardiv\$ and dyskine\$) or akathisi\$ or acathisi\$ or (neuroleptic\$ and malignant and syndrome) or (neuroleptic and movement and disorder) or parkinsoni\$ or neuroleptic-induc\$) not (parkinson\$ and disease)).ti,ab.
8. (emergency services, psychiatric or hospitals, psychiatric or psychiatric department, hospital or (mentally ill persons and (inpatients or

- hospitalization))).sh. or (psychiatric adj2 (admission\$ or admitted or emerg\$ or hospitali\$ or inpatient\$ or in patient\$)).ti,ab.
9. or/1-8
  10. comorbidity/ or "diagnosis, dual (psychiatry)"/
  11. (comorbid\$ or co morbid\$ or ((dual\$ or tripl\$) adj2 (diagnos\$ or disease\$ or disorder\$ or illness\$ or mental or problem\$ or psych\$ or syndrome\$)) or coexist\$ or co exist\$ or concur\$ or con cur\$ or cooccur\$ or co occur\$).ti,ab.
  12. or/10-11
  13. (designer drugs or needle exchange programs or needle sharing or overdose or street drugs or substance abuse detection or substance abuse, intravenous or substance abuse treatment centers or substance-related disorders or substance withdrawal syndrome).sh.
  14. (((drug\$1 or polydrug\$ or psychotropic\$ or substance\$) adj3 (abstain\$ or abstinen\$ or abus\$ or addict\$ or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxicat\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or unlawful\$ or withdraw\$)) or ((drug\$1 or polydrug\$ or recreation\$ or substance\$) adj use\$1) or ((drug\$1 or polydrug\$ or substance\$) adj rehab\$) or abusable product\$ or (crave\$ adj2 inject\$) or hard drug\$ or needle fixation or soft drug\$ or vsa\$1).ti,ab.
  15. ((club or designer or street) adj2 (drug\$ or substance\$)).ti,ab.
  16. or/13-15
  17. (amphetamine or amphetamine-related disorders).sh.
  18. (dextroamphetamine or methamphetamine).sh.
  19. (((amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers) adj2 use\$1)).ti,ab.
  20. (amphetamin\$ or crystal meth\$ or desoxyn or dexamfetamin\$ or dexedrine or dextroamphetamin\$ or methamphetamin\$ or psychostimulant\$ or stimulant\$ or uppers).ti,ab.
  21. or/17-19
  22. 20
  23. exp cocaine/ or cocaine-related disorders.sh.

24. (((benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine) adj2 use\$1)).ti,ab.
25. (benzoylmethyl ecgonine or cocain\$ or crack\$1 or codrenine or ecgonine methyl ester benzoate or erythroxylin or locosthetic or neurocaine or sterilocaine).ti,ab.
26. or/23-24
27. 25
28. (heroin or heroin dependence or opioid-related disorders).sh.
29. (((heroin or diacetylmorphin\$ or diagesil or diamorf or diamorphin\$) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((diamorphin\$ or acetomorphine or anpec or diacephine or diacetylmorphine\$ or diagesil or diamorf or diaphorin or duromorph or epimorph or heroin or morfin\$ or morphacetin or morphia or morphian\$ or morphin\$ or morphium or opso\$1 or skenan) adj2 use\$1)).ti,ab.
30. (heroin or diacetylmorphin\$ or diagesil or diamorf or diamorphin\$).ti,ab.
31. or/28-29
32. 30
33. (cannabis or marijuana abuse or marijuana smoking).sh.
34. (((bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk) adj5 (abstain\$ or abstinen\$ or abus\$ or addict\$ or banned or excessive use\$ or criminal or depend\$ or habit\$ or illegal\$ or illicit\$ or intoxica\$ or misus\$ or nonprescri\$ or non prescri\$ or over dos\$ or overdos\$ or recreation\$ or rehab\$ or unlawful\$ or using or utilis\$ or utiliz\$ or withdraw\$)) or ((bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk) adj2 use\$1)).ti,ab.
35. (bhang or cannador or cannabis or ganja or ganjah or hashish or hemp or marihuana or marijuana or sativex or skunk).ti,ab.
36. or/33-34

- 37. 35
- 38. 9 and 12 and (or/22,27,32,37)
- 39. 9 and (or/16,21,26,31,36)
- 40. or/38-39

### 10.1.2 Search filters

*Health economics and quality of life search filter – this is an adaptation of a filter designed by the NHS Centre for Reviews and Dissemination at the University of York.*

MEDLINE - Ovid SP interface

1. exp “costs and cost analysis” / or health priorities/ or health resources/ or exp resource allocation/
2. budgets/ or socioeconomic factors/ or (economi\$ or fee or fees or financ\$).hw.
3. quality adjusted life years/ or "quality of life"/ or "value of life"/
4. exp models, economic/ or models, statistical/ or monte carlo method/
5. health status indicators/
6. decision trees/
7. (budget\$ or cost\$ or econom\$ or expenditure\$ or financ\$ or fiscal or funding or pharmaco-economic\$ or socioeconomic\$ or price or prices or pricing or (value adj3 money) or (burden adj3 (disease\$ or illness\$))).ti,ab.
8. (daly or qol or hql or hqol or hrqol or hr ql or hrql or (quality adj2 life) or (adjusted adj2 life) or qaly\$ or (health adj2 stat\$) or well being or wellbeing or qald\$ or qale\$ or qtime\$ or eq5d or eq 5d or qw b or ((quality or value\$) adj3 (life or survival or well\$)) or hui\$1 or (utilit\$ adj1 (health or score\$ or weigh\$)) or (life adj2 year\$) or health year equivalent\$ or ((disability or quality) adj adjusted) or utility value\$ or (weight\$ adj3 preference\$) or euroqol or euro qol or visual analog\$ or standard gamble or time trade or qtwist or q twist or (valu\$ adj2 quality)).ti,ab.
9. decision tree/ or decision trees/
10. (decision analy\$ or monte carlo or markov or simulation model\$ or rosser or disutili\$ or willingness to pay or tto or hye or hyes or (resource adj (allocat\$ or use\$ or utilit\$))).ti,ab.
11. (sf36 or sf 36 or short form 36 or shortform 36 or sf thirtysix or sf thirty six or shortform thirtysix or shortform thirty six or short form thirtysix or short form thirty six).ti,ab.

- 12 (sf6 or sf 6 or short form 6 or shortform 6 or sf six or sfsix or shortform six or short form six).ti,ab.
- 13 (sf12 or sf 12 or short form 12 or shortform 12 or sf twelve or sftwelve or shortform twelve or short form twelve).ti,ab.
- 14 (sf16 or sf 16 or short form 16 or shortform 16 or sf sixteen or sfsixteen or shortform sixteen or short form sixteen).ti,ab.
- 15 (sf20 or sf 20 or short form 20 or shortform 20 or sf twenty or sftwenty or shortform twenty or short form twenty).ti,ab.
- 16 ec.fs. [*ANDed with subject heading searches for the main population/topic*]
- 17 or/1-16

## APPENDIX 10: METHODOLOGY CHECKLIST TEMPLATE FOR ECONOMIC STUDIES

The methodological quality of each study was evaluated using the NICE checklists for economic evaluations, reproduced below (for information about how to complete the checklist, see *The Guidelines Manual* [NICE, 2009b]).

<b>Study identification</b> <i>Including author, title, reference, year of publication</i>			
Guideline topic:			Question no:
Checklist completed by:			
<b>Section 1: Applicability (relevance to specific guideline review question(s) and the NICE reference case). This checklist should be used first to filter out irrelevant studies.</b>		<b>Yes/ Partly/ No/Unclear /NA</b>	<b>Comments</b>
1.1	Is the study population appropriate for the guideline?		
1.2	Are the interventions appropriate for the guideline?		
1.3	Is the healthcare system in which the study was conducted sufficiently similar to the current UK NHS context?		
1.4	Are costs measured from the NHS and personal social services (PSS) perspective?		
1.5	Are all direct health effects on individuals included?		
1.6	Are both costs and health effects discounted at an annual rate of 3.5%?		
1.7	Is the value of health effects expressed in terms of quality-adjusted life years (QALYs)?		
1.8	Are changes in health-related quality of life (HRQoL) reported directly from patients and/or carers?		
1.9	Is the valuation of changes in HRQoL (utilities) obtained from a representative sample of the general public?		
1.10	Overall judgement: Directly applicable/Partially applicable/Not applicable There is no need to use section 2 of the checklist if the study is considered 'not applicable'.		
Other comments:			

<b>Section 2: Study limitations (the level of methodological quality)</b> <b>This checklist should be used once it has been decided that the study is sufficiently applicable to the context of the clinical guideline.</b>		<b>Yes/ Partly /No/ Unclear/ NA</b>	<b>Comments</b>
2.1	Does the model structure adequately reflect the nature of the health condition under evaluation?		
2.2	Is the time horizon sufficiently long to reflect all important differences in costs and outcomes?		
2.3	Are all important and relevant health outcomes included?		
2.4	Are the estimates of baseline health outcomes from the best available source?		
2.5	Are the estimates of relative treatment effects from the best available source?		
2.6	Are all important and relevant costs included?		
2.7	Are the estimates of resource use from the best available source?		
2.8	Are the unit costs of resources from the best available source?		
2.9	Is an appropriate incremental analysis presented or can it be calculated from the data?		
2.10	Are all important parameters whose values are uncertain subjected to appropriate sensitivity analysis?		
2.11	Is there no potential conflict of interest?		
2.12	Overall assessment: Minor limitations/Potentially serious limitations/Very serious limitations		
Other comments:			

## APPENDIX 11: EVIDENCE TABLES FOR ECONOMIC STUDIES

Study ID Country Study type	Intervention details	Study population Study design Data sources	Costs: description and values Outcomes: description and values	Results: Cost-effectiveness	Comments
Clark <i>et al.</i> , 1998 USA Cost-effectiveness analysis (CEA)	Compared assertive community treatment (ACT) and standard case management (SCM) for patient with severe mental illness and substance misuse disorders	Study population: Patients with DSM-III-R diagnosis of schizophrenia, schizoaffective disorder, or bipolar disorder and; an active substance use disorder. Patients randomised to ACT (n=100) or PSM (n=93) Ave age: 34 Years; 74% Male Time-frame: 3 Years Study design: RCT (multi-centre) Data source(s): 7 mental health catchment areas in the US	<b>Costs</b> Resource use: Mental health treatment; General health care; legal system; community services (homeless shelters/soup kitchens); administration; informal care (family members' input)  <b>Outcomes</b> Subjective QoL year details provided from patients' perspective using Quality of Life Interview instrument. A modified range from 0 (terrible) to 1 (delighted) was used and weighted (cumulative) scores were derived based on the time spent on each rating	<b>Costs</b> ACT: \$118,078 per patient SCM: \$124,145 per patient  <b>Outcomes (QoL improvement from baseline)</b> ACT: 0.10 SCM: 0.04  <b>Cost-effectiveness</b> Ratios of cumulative quality of life years to total societal costs rather than of incremental cost-effectiveness were computed. Average QoL ratios per \$10,000 in societal costs were 0.24 (ACT) and 0.20 (SCM).	Perspective: Societal Currency: US \$ Cost Year: 1995 Time horizon: 3 years Discounting: Yes (3% costs; 5% outcomes) Funded by: National Institute of Mental Health, National Institute on Alcohol Abuse and Alcoholism/ New Hampshire Division of Mental Health and Developmental Services

<b>Study ID Country Study type</b>	<b>Intervention details</b>	<b>Study population Study design Data sources</b>	<b>Costs: description and values Outcomes: description and values</b>	<b>Results: Cost-effectiveness</b>	<b>Comments</b>
Craig <i>et al.</i> , 2008 UK Cost-Analysis (CA)	Programme for case managers that trained them to manage substance use disorders among persons with severe mental illness compared with waiting list control	Study population: Patients with clinical diagnosis of schizophrenia, schizoaffective disorder, or other non-affective psychotic illnesses or bipolar disorder with psychotic symptoms plus abuse or dependence on at least one substance Intervention (n=124) Control (n=104) Time-frame: 18 months Study Design: RCT (Cluster) Data source(s): Community mental health services in four London boroughs	<b>Costs</b> Resource use: Hospital inpatient days; Day Care; Medication; HC professional appointments (Psychiatrist, Community Nurse, Social Worker, Psychologist, Drug or Alcohol worker, Counsellor, GP); Criminal Justice (Court/Police/Prison)	<b>Total Mean Costs</b>  Intervention: 18,672 Control: 17,639	Perspective: Societal Currency: UK £ Cost Year: 2003/04 Time horizon: 18 months Discounting: No Funded by: Bethlem and Maudsley NHS Trust

Study ID Country Study type	Intervention details	Study population Study design Data sources	Costs: description and values Outcomes: description and values	Results: Cost-effectiveness	Comments
French <i>et al.</i> , 1999 USA Cost-Consequences Analysis (CCA)	Modified therapeutic community (TC) intervention compared with standard services in a treatment-as-usual (TAU) condition	Study population: Homeless mentally ill chemical abusers (MICAs) – axis I diagnoses of schizophrenia, major depression, mania and who also use drugs or alcohol Modified TC (n=228); TAU (n=53) Study Design: Cohort Study Data source(s): Homeless facilities and psychiatric hospitals located in New York City	<b>Costs</b> Perspective: Health service Intervention, hospital detox, emergency room visits, short-term residential stays, non-residential stays, outpatient visits, methadone maintenance, inpatient days  <b>Outcomes</b> Substance use, criminal activity, HIV-risk behaviour, psychological status, employment status	<b>Costs</b> Modified TC: \$29,255 TAU: \$29,638  <b>Outcomes</b> Modified TC patients reported significantly greater reductions in criminal activity and psychological dysfunction; no significant differences in substance use or HIV-risk behaviour  No formal synthesis of costs and outcomes	Perspective: Health service Currency: US \$ Cost Year: 1994 Time horizon: 12 months Discounting: NA Funded by: National Institute on Drug Abuse, Public Health Service, US Department of Health and Human Services

Study ID Country Study type	Intervention details	Study population Study design Data sources	Costs: description and values Outcomes: description and values	Results: Cost-effectiveness	Comments
Haddock <i>et al.</i> , 2003 UK CEA	Integrated programme of cognitive-behavioural (CBT) combined with motivational intervention (MI) plus routine care (RC) versus RC alone	Study population: Patients (entered as patient and carer pairs) with ICD-10 diagnosis of schizophrenia, schizoaffective disorder or delusional disorder and DSM-IV diagnosis of substance dependence or misuse. Intervention (n=18) Control (n=18) Study Design: RCT Data source(s): Mental health units of 3 UK NHS hospital trusts	<b>Costs</b> Resource use: Intervention; hospital services; primary care services (GPs/practice nurses); community or domiciliary services (social workers/occupational therapists); day services; medication; patient costs (travel/out-of-pocket payments); productivity losses  <b>Outcomes</b> Change in the Global Assessment of Functioning Scale (GAF) over 18 months	<b>Costs</b> Intervention: 8,753 (SD 4,804) Control: 10,013 (SD 10,717)  <b>Outcomes</b> Intervention: 60.12 (SD 18.96) Control: 53.44 (SD 13.00)  <b>Cost-effectiveness</b> Not reported Probability of intervention being less costly than routine care (at WTP of 0) was 69.3%	Perspective: Societal Currency: UK £ Cost Year: 1998/99 Time horizon: 18 months Discounting: No Funded by: West Pennine, Manchester and Stockport Health Authorities, Tameside and Glossop NHS Trust R&D support

Study ID Country Study type	Intervention details	Study population Study design Data sources	Costs: description and values Outcomes: description and values	Results: Cost-effectiveness	Comments
Jerrell <i>et al.</i> , 1997 USA CEA	Comparison of three primary interventions (with emphasis on any ethnic differences): 12-Step recovery, case management and behavioural skills training	Study population: Patients with Axis I DSM-III-R diagnosis of psychosis or major depression with a co-occurring substance disorder 12-Step (n=39) Behavioural skills (n=48) Case Management (n=45) Study Design: RCT Data source (s): 5 community mental health centres in the US	<p><b>Costs</b> Perspective: Societal Resource use: 2 categories: Intensive mental health (inpatient days, nursing days, residential treatment, emergency days); Supportive mental health (case management hours, outpatient visits, supporting housing days, service days)</p> <p><b>Outcomes</b> Psychological functioning (Social Adjustment Scale-II; Role Functioning Scale), mental health and substance abuse (Diagnosis Interview Schedule used by C-DIS_R programme)</p>	<p><b>Total Costs</b></p> <p><b>Intensive mental health costs</b> 12-Step: \$10,275 Behavioural skills: \$4,276 Case Management: \$7,643</p> <p><b>Supportive mental health costs</b> 12-Step: \$7,798 Behavioural skills: \$6,112 Case Management: \$5,970</p> <p>No differences between three treatment approaches in psychological functioning or psychiatric or substance abuse symptoms. Analysis was therefore based on cost differences</p>	Perspective: US Health service Currency: US \$ Cost Year: Not reported Time horizon: 18 months Discounting: No Funded by: National Institute of Mental Health

Study ID Country Study type	Intervention details	Study population Study design Data sources	Costs: description and values Outcomes: description and values	Results: Cost-effectiveness	Comments
Morse <i>et al.</i> , 2006 USA CA	Three treatments: Integrated Assertive Community Treatment (IACT); Assertive Community Treatment Only (ACTO) and Standard Care (Control)	Study Population: Individuals (homeless at baseline) with co- occurring SMI and substance use disorders IACT (n=54); ACTO (n=54); Control (n=49) Mean Age: 40 yrs; 80% Male Study design: RCT Data source(s): US-based community mental health agencies	<p><b>Costs</b> Perspective: Societal Outpatient care (Direct treatments for IACT and ACTO; other mental health, other substance abuse treatment, physical health care, psychosocial rehabilitation); Inpatient care; Emergency Shelter; Social security; Transfer payments and maintenance benefits</p> <p><b>Outcomes</b> Client Satisfaction; BPRS scale; Substance use (Interviewer rating)</p>	<p><b>Costs</b> IACT: \$48,764 ACTO: \$71,211 Control: \$41,726</p> <p>IACT and Control groups had significantly lower total mean costs than ACTO but no significant differences between IACT and Control</p> <p><b>Outcomes</b> IACT and ACTO participants significantly more satisfied with their treatment than control; no significant differences between IACT and ACTO. There was no significant effect of treatment group on BPRS scale (p=0.1) or substance use levels (p=0.72)</p>	Perspective: Societal Currency: US \$ Cost Year: 2001 Time horizon: 24 months Discounting: No Funded by: National Institute of Mental Health

## 1 APPENDIX 12: HIGH PRIORITY RECOMMENDATIONS

2 The Guideline Development Group has made the following recommendations  
3 for research, based on its review of evidence, to improve NICE guidance and  
4 patient care in the future.  
5

### 6 **10.1.1 Determining prevalence, risk and protective factors, and** 7 **course of illness**

8 What are the prevalence, risk and protective factors, and course of illness for  
9 different combinations of psychosis and coexisting substance misuse (for  
10 example, schizophrenia and cannabis misuse or bipolar disorder and alcohol  
11 misuse)?

#### 12 *Why this is important*

13 Studies vary in terms of the definitions and diagnosis of psychosis and  
14 substance misuse, and how they are conducted. This makes it difficult to  
15 draw conclusions about prevalence and patterns in patient groups  
16 differentiated by diagnosis, ethnicity and other demographics. Additionally,  
17 most studies are cross-sectional, so little is known about how both conditions  
18 change over time. Moreover, there is little guidance about which levels and  
19 patterns of substance misuse in which patient groups are associated with the  
20 worst clinical and social outcomes. Such information is necessary to target  
21 resources at groups most at risk of very poor outcomes.

22 This question should be answered using a longitudinal study design with a  
23 representative sample large enough to establish the prevalence, pattern, and  
24 epidemiology of different combinations of psychosis and coexisting substance  
25 misuse, associated social determinants, treatment and outcome. The study  
26 should also collect information that could inform the development of new  
27 interventions or the modification of existing interventions to improve  
28 prognosis.

1 **10.1.2 Predicting the onset of substance misuse in young people**  
2 **with psychosis**

3 What risk factors predict the onset of substance misuse in young people with  
4 psychosis?

5 *Why this is important*

6 People with psychosis and coexisting substance misuse are more likely to be  
7 non-adherent to prescribed medication, and have poor engagement with  
8 treatment programmes, increased risk of suicide, more and longer inpatient  
9 stays, increased risk of violence and time spent in the criminal justice system,  
10 and poorer overall prognosis. Because the onset of psychosis at a younger age  
11 is also an indicator of poor prognosis, people with a combination of younger  
12 age of onset and coexisting substance misuse may have a particularly poor  
13 prognosis. A clearer understanding of the risk and protective factors for  
14 substance misuse in young people with psychosis, and the interrelationship of  
15 the two conditions over time, may facilitate the development of treatment  
16 approaches for the coexisting conditions in this group. This may then improve  
17 the longer term outcome for a group of people who tend to have a poor  
18 prognosis.

19 This question should be answered using a prospective cohort study design.

20 **10.1.3 Future trials of interventions for people with psychosis or**  
21 **interventions for people with substance misuse.**

22 Are interventions for psychosis or substance misuse clinically and cost  
23 effective when compared with standard care for people with psychosis and  
24 coexisting substance misuse?

25 *Why this is important*

26 Whilst there is substantial evidence for the clinical and cost effectiveness of  
27 interventions for psychosis (for example, NICE guidelines for Schizophrenia  
28 and Bipolar Disorder), these interventions have not been adequately tested in  
29 people with coexisting substance misuse. Similarly, interventions that have  
30 been shown to be effective and cost-effective in substance misuse (for  
31 example, see NICE guidelines for Substance Misuse) have not been  
32 adequately evaluated in people with coexisting psychosis. For  
33 pharmacological interventions, these gaps in evidence are partly related to the  
34 requirements of the regulatory authorities for the licensing and marketing  
35 approval of new medicines that have been tested in specific clinical  
36 populations under ideal circumstances. However more recently pragmatic  
37 RCTs which attempt to examine the effectiveness of interventions in 'real

1 world' clinical practice are increasingly being conducted. Such studies should  
2 include people with the types of complex problems that services routinely  
3 work with - including those with coexisting psychosis and substance misuse.  
4 While numbers of people with coexisting conditions in individual studies are  
5 likely to be too small to be able to draw conclusions about their effectiveness  
6 in this group, collection of such data could facilitate future systematic reviews  
7 and help increase the evidence base for the management of people with this  
8 complex combination of problems.

9

10 This question should be answered by not routinely excluding people with  
11 psychosis and coexisting substance misuse from future trials of interventions  
12 for people with psychosis or interventions for people with substance misuse.

### 13 **10.1.4 Psychosocial interventions versus standard care**

14 Are psychosocial interventions clinically and cost effective when compared  
15 with standard care for people with psychosis and coexisting substance  
16 misuse?

#### 17 *Why this is important*

18 Psychosocial interventions are recommended for the treatment of substance  
19 misuse, with contingency management showing particular promise.

20 However, they have not been adequately tested in people who also have  
21 psychosis.

22 This question should be answered using a randomised controlled trial that  
23 examines the short- and medium-term outcomes over at least 18 months.

24 Studies should focus on people whose misuse of substances is most often  
25 encountered in clinical practice and has the greatest impact on mental health  
26 (such as cannabis and polysubstance misuse), and on those interventions –  
27 such as contingency management, cognitive therapy and relapse prevention –  
28 that show most promise in people with substance misuse without psychosis.

29 Those providing the intervention should be trained and supervised to ensure  
30 that the results are robust and generalisable. Outcomes should reflect both  
31 observer and service user-rated assessments of improvement (including  
32 mental health and social functioning) and the intervention's acceptability.

33 Studies need to be large enough to determine the intervention's costs and cost  
34 effectiveness.

### 35 **10.1.5 Environmental interventions versus standard care**

36 Are environmental interventions clinically and cost effective when compared  
37 with standard care for people with psychosis and coexisting substance  
38 misuse?

1 *Why this is important*

2 Social and other environmental factors can play an important role in  
3 triggering and maintaining substance misuse in people with psychosis, and in  
4 reducing the likelihood of progress and recovery. Evidence suggests that  
5 when the primary focus of management involves improving the environment,  
6 both conditions may improve.

7 This question should be answered using a randomised controlled trial that  
8 examines short- and medium-term outcomes over at least 12 months. Studies  
9 should focus on people with psychosis whose misuse of substances is most  
10 often encountered in clinical practice and has the greatest impact on mental  
11 health (such as cannabis and polysubstance misuse, and on interventions that  
12 take a collaborative approach to identifying and modifying social and  
13 environmental factors that may trigger substance misuse. Those providing the  
14 intervention should be trained and supervised to ensure that the results are  
15 robust and generalisable. Outcomes should reflect both observer and service  
16 user-rated assessments of improvement (including mental health and social  
17 functioning) and the intervention's acceptability. Studies need to be large  
18 enough to determine the intervention's costs and cost effectiveness.

19 **10.1.6 Clozapine versus other pharmacological interventions**

20 Is clozapine clinically and cost effective when compared with other  
21 pharmacological interventions for people with psychosis and coexisting  
22 substance misuse?

23 *Why this is important*

24 The NICE guideline on schizophrenia (NICE clinical guideline 82) states that  
25 clozapine should be offered to people with schizophrenia whose illness has  
26 not responded adequately to treatment despite the sequential use of adequate  
27 doses of at least two different antipsychotic drugs. However, there is  
28 insufficient evidence to guide healthcare professionals about the use of  
29 clozapine in people with psychosis and coexisting substance misuse. Expert  
30 opinion often advocates clozapine as having a particular role in this  
31 population, but the evidence to support such statements is lacking. Clozapine  
32 is expensive and has a wide range of side effects, some of which may be life-  
33 threatening if not monitored correctly.

34 This question should be answered using a randomised controlled trial in  
35 which participants are stratified for presenting problem. It should report short  
36 and longer-term outcomes (including substance misuse, acceptability of the  
37 intervention, and cost effectiveness) of at least 12 months' duration.  
38

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# 1 11 REFERENCES

- 2 Abou-Saleh, M. T. (2004) Dual diagnosis: management within a psychosocial  
3 context. *Advances in Psychiatric Treatment*, 10, 352-360.  
4
- 5 AGREE Collaboration (2003) Development and validation of an international  
6 appraisal instrument for assessing the quality of clinical practice guidelines:  
7 the AGREE project. *Quality and Safety in Health Care*, 12, 18-23.  
8
- 9 Alem, A. & Shibbe, T. (1997) Khat induced psychosis and its medico-legal  
10 implications: a case report. *Ethiopian Medical Journal*, 35, 137-141.  
11
- 12 Alvidrez, J., Kaiser, D. & Havassy, B. E. (2004) Severely mentally ill  
13 consumers' perspectives on drug use. *Journal of Psychoactive Drugs*, 36, 347-  
14 355.  
15
- 16 American Psychiatric Association (1980) Diagnostic and Statistical Manual of  
17 Mental Disorders (3<sup>rd</sup> edn) DSM III. Washington, DC: American Psychiatric  
18 Association.  
19
- 20 American Psychiatric Association (1987) Diagnostic and Statistical Manual of  
21 Mental Disorders (3<sup>rd</sup> edn, revised) DSM III-R. Washington, DC: American  
22 Psychiatric Association.  
23
- 24 American Psychiatric Association (1994) *Diagnostic and Statistical Manual of*  
25 *Mental Disorders (4th edn) DSM IV*. Washington, DC: American Psychiatric  
26 Association.  
27
- 28 Anderson, A. J. (1999) Comparative impact evaluation of two therapeutic  
29 programs for mentally ill chemical abusers. *The International Journal of*  
30 *Psychosocial rehabilitation*, 4, 11-26.  
31
- 32 Andreasson, S., Allebeck, P., Engstrom, A., et al. (1987) Cannabis and  
33 schizophrenia: a longitudinal study of Swedish conscripts. *Lancet*, 330, 1483-  
34 1486.  
35
- 36 Annis, H. M. (1986) A relapse prevention model for treatment of alcoholics. In  
37 *Treating Addictive Behaviors: Processes of Change (Applied Clinical Psychology)*  
38 (eds W. R. Miller & N. Heather). New York: Plenum Press.  
39

- 1 Arseneault, L., Cannon, M., Poulton, R., *et al.* (2002) Cannabis use in  
2 adolescence and risk for adult psychosis: longitudinal prospective study.  
3 *British Medical Journal*, 325, 1212-1213.  
4
- 5 Ashton, M. (2005) The motivational halo. *Drug and Alcohol Findings*, 13, 23-30.  
6
- 7 Baker, A., Bucci, S., Lewin, T. J., *et al.* (2006) Cognitive-behavioural therapy for  
8 substance use disorders in people with psychotic disorders: randomised  
9 controlled trial. *British Journal of Psychiatry*, 188, 439-448.  
10
- 11 Baker, A., Lewin, T., Reichler, H., *et al.* (2002) Motivational interviewing  
12 among psychiatric in-patients with substance use disorders. *Acta Psychiatrica*  
13 *Scandinavica*, 106, 233-240.  
14
- 15 Barnaby, B., Drummond, C., McLeod, A., *et al.* (2003) Substance misuse in  
16 psychiatric inpatients: a comparison of a screening questionnaire survey with  
17 case notes. *British Medical Journal*, 327, 783-784.  
18
- 19 Barnes, T. R., Mutsatsa, S. H., Hutton, S. B., *et al.* (2006). Comorbid substance  
20 use and age at onset of schizophrenia. *British Journal of Psychiatry*, 188, 237-  
21 242.  
22
- 23 Barnett, J., Werners, U., Secher, S. M., *et al.* (2007) Substance use in a  
24 population-based clinic sample of people with first- episode psychosis. *British*  
25 *Journal of Psychiatry*, 190, 515-520.  
26
- 27 Barrowclough, C., Haddock, G., Fitzsimmons, M., *et al.* (2006) Treatment  
28 development for psychosis and co-occurring substance misuse: a descriptive  
29 review. *Journal of Mental Health*, 15, 619-632.  
30
- 31 Barrowclough, C., Haddock, G., Lowens, I., *et al.* (2005) Psychosis and drug  
32 and alcohol problems. In *Clinical Handbook of Co-existing Mental Health and*  
33 *Drug and Alcohol Problems* (eds A. Baker & R. Velleman). London: Routledge.  
34
- 35 Barrowclough, C., Haddock, G., Tarrier, N., *et al.* (2001) Randomised  
36 controlled trial of cognitive behavioural therapy plus motivational  
37 intervention for schizophrenia and substance use. *American Journal of*  
38 *Psychiatry*, 158, 1706-1713.  
39
- 40 Barrowclough, C., Haddock, G., Wykes, T., *et al.* (in press) A randomised  
41 controlled trial of integrated motivational interviewing and cognitive

- 1 behaviour therapy for people with psychosis and co-morbid substance misuse  
2 – the MIDAS trial. *BMJ*.
- 3 Barrowclough, C., Tarrier, N., Humphreys, L., *et al.* (2003). Self esteem in  
4 schizophrenia: the relationship between self evaluation, family attitudes and  
5 symptomatology. *Journal of Abnormal Psychology*, 112, 92-99.
- 6
- 7 Beck, A. T., Wright, F. D., Newman, C. F., *et al.* (1993) *Cognitive Therapy of*  
8 *Substance Abuse*. New York: Guilford Press.
- 9 Bender, K., Springer, D. W. & Kim, J. S. (2006) Treatment effectiveness with  
10 dually diagnosed adolescents: a systematic review. *Brief Treatment and Crisis*  
11 *Intervention*, 6, 177-205.
- 12
- 13 Berlin, J. A. (2001) Does blinding of readers affect the results of meta-  
14 analyses? *Lancet*, 350, 185-186.
- 15
- 16 Blanchard, J. J., Brown, S. A., Horan, W. P., *et al.* (2000) Substance use  
17 disorders in schizophrenia: review, integration and a proposed model. *Clinical*  
18 *Psychology Review*, 20, 207-234.
- 19
- 20 Blankertz, L. E. & Cnaan, R. A. (1992) Principles of care for dually diagnosed  
21 homeless persons: Findings from a demonstration project. *Research on Social*  
22 *Work Practice*, 2, 448-464.
- 23
- 24 Blankertz, L. E. & Cnaan, R. A. (1994) Assessing the impact of two residential  
25 programs for dually diagnosed homeless individuals. *Social Service Review*, 68,  
26 536-560.
- 27
- 28 Bloye, D., Ramzan, A., Leach, C., *et al.* (2003) Substance use disorders in  
29 patients admitted to a medium secure unit: a comparison of three assessment  
30 measures. *Journal of Forensic Psychiatry & Psychology*, 14, 585-599.
- 31
- 32 BMA & NHS Employers (2009) *Quality and Outcomes Framework Guidance for*  
33 *GMS Contract 2009/10: Delivering Investment in General Practice*. London: NHS  
34 Employers & General Practitioners Committee. Available at:  
35 [http://www.bma.org.uk/images/qof0309\\_tcm41-184025.pdf](http://www.bma.org.uk/images/qof0309_tcm41-184025.pdf)
- 36
- 37 Bonsack, C., Camus, D., Kaufmann, N., *et al.* (2006) Prevalence of substance  
38 use in a Swiss psychiatric hospital: interview reports and urine screening.  
39 *Addictive Behaviors*, 31, 1252-1258.
- 40

- 1 Bradizza, C. M. & Stasiewicz, P. R. (2003) Qualitative analysis of high-risk  
2 drug and alcohol use situations among severely mentally ill substance  
3 abusers. *Addictive Behaviours*, 28, 157-169.  
4
- 5 Brown, S. E., Carmody, T. J., Schmitz, J. M., *et al.* (2009) A randomized,  
6 double-blind, placebo-controlled pilot study of naltrexone in outpatients with  
7 bipolar disorder and alcohol dependence. *Alcoholism: Clinical and Experimental*  
8 *Research*, 33, 1863-1869.  
9
- 10 Brunette, M. F., Drake, R. E., Woods, M., *et al.* (2001) A comparison of long-  
11 term and short-term residential treatment programs for dual diagnosis  
12 patients. *Psychiatric Services*, 52, 526-528.  
13
- 14 Brunette, M., Mueser, K., Xie, H., *et al.* (1997) Relationships between  
15 symptoms of schizophrenia and substance abuse. *Journal of Nervous and*  
16 *Mental Disease*, 185, 251-257.  
17
- 18 Brunette, M. F., Noordsy, D. L., Buckley, P. F., *et al.* (2005) Pharmacologic  
19 treatments for co-occurring substance use disorders in patients with  
20 schizophrenia. *Journal of Dual Diagnosis*, 1, 41-55.  
21
- 22 Buchanan, R. W., Kreyenbuhl, J., Kelly, D. L., *et al.* (2009) The 2009  
23 schizophrenia PORT psychopharmacological treatment recommendations and  
24 summary statements. *Schizophrenia Bulletin*, 36, 71-93.
- 25 Burnam, M. A., Morton, S. C., McGlynn, E. A., *et al.* (1995) An experimental  
26 evaluation of residential and non-residential treatment for dually diagnosed  
27 homeless adults. *Journal of Addictive Diseases*, 14, 111-134.  
28
- 29 Buckley, P. F., Miller, B. J., Lehrer, D. S., *et al.* (2009) Psychiatric comorbidities  
30 and schizophrenia. *Schizophrenia Bulletin*, 35, 383-402.  
31
- 32 Buhler, B., Hambrecht, M., Loffler, W., *et al.* (2002) Precipitation and  
33 determination of the onset and course of schizophrenia by substance abuse: a  
34 retrospective and prospective study of 232 population-based first illness  
35 episodes. *Schizophrenia Research*, 54, 243-251.  
36
- 37 Carey, K. B. & Carey, M. P. (1995) Reasons for drinking among psychiatric  
38 outpatients: relationship to drinking patterns. *Psychology of Addictive*  
39 *Behaviours*, 9, 251-257.  
40

- 1 Carey, K. B. & Correia, C. J. (1998) Severe mental illness and addictions:  
2 assessment considerations. *Addictive Behaviours*, 23, 735-748.  
3
- 4 Carey, K. B., Purnine, D. M., Maisto, S. A., *et al.* (1999) Decisional balance  
5 regarding substance use among persons with schizophrenia. *Community*  
6 *Mental Health Journal*, 35, 289-299.
- 7 Carra, G. & Johnson, S. (2009) Variations in rates of comorbid substance use in  
8 psychosis between mental health settings and geographical areas in the UK.  
9 *Social Psychiatry and Psychiatric Epidemiology*, 44, 429-447.  
10
- 11 Casas, M., Franco, M. D., Goikolea, J. M., *et al.* (2008) Bipolar disorder  
12 associated to substance use disorders (dual diagnosis): systematic review of  
13 the scientific evidence and expert consensus. *Actas Españolas de Psiquiatría*, 36,  
14 350-361.  
15
- 16 Caspi, A., Moffitt, T. E., Cannon, M., *et al.* (2005) Moderation of the effect of  
17 adolescent-onset cannabis use on adult psychosis by a functional  
18 polymorphism in the catechol-O-methyltransferase gene: longitudinal  
19 evidence of a gene x environment interaction. *Biological Psychiatry*, 57, 1117-  
20 1127.  
21
- 22 Caton, C. L. M., Drake, R. E., Hasin, D. S., *et al.* (2005) Differences between  
23 early phase primary psychotic disorders with concurrent substance use and  
24 substance-induced psychosis. *Archives of General Psychiatry*, 62, 137-145.  
25
- 26 Caton, C. L. M., Hasin, D. S., Shrout, P. E., *et al.* (2007) Stability of early-phase  
27 primary psychotic disorders with concurrent substance use and substance-  
28 induced psychosis. *British Journal of Psychiatry*, 190, 105-111.  
29
- 30 Center for Substance Abuse Treatment (2005a) *Substance Abuse Treatment for*  
31 *Persons with Coexisting Disorders*. Treatment Improvement Protocol (TIP)  
32 Series 42. DHHS Publication No. (SMA) 05-3992. Rockville, MD: Substance  
33 Abuse and Mental Health Services Administration.  
34
- 35 Center for Substance Abuse Treatment (2005b) *Medication-Assisted Treatment*  
36 *for Opioid Addiction in Opioid Treatment Programs*. Treatment Improvement  
37 Protocol (TIP) Series 43. DHHS Publication No. (SMA) 05-4048. Rockville,  
38 MD: Substance Abuse and Mental Health Services Administration.  
39
- 40 Center for Substance Abuse Treatment (2006) *Detoxification and Substance*  
41 *Abuse Treatment*. Treatment Improvement Protocol (TIP) Series 45. DHHS

- 1 Publication No. (SMA) 06-4131. Rockville, MD: Substance Abuse and Mental  
2 Health Services Administration.  
3
- 4 Centorrino, F., Cincotta, S. L., Talamo, A., *et al.* (2008) Hospital use of  
5 antipsychotic drugs: polytherapy. *Comprehensive Psychiatry*, 49, 65-69.  
6
- 7 Chan, Y., Dennis, M. L. & Funk, R. R. (2008) Prevalence and mental health-  
8 substance use of major internalizing and externalizing problems among  
9 adolescents and adults presenting to substance abuse treatment. *Journal of*  
10 *Substance Abuse Treatment*, 34, 14-24.  
11
- 12 Chandler, D. W. & Spicer, G. (2006) Integrated treatment for jail recidivists  
13 with co-occurring psychiatric and substance use disorders. *Community Mental*  
14 *Health Journal*, 42, 405-425.  
15
- 16 Charles, V. & Weaver, T. (2010) A qualitative study of illicit and non-  
17 prescribed drug use among people with psychotic disorders. *Journal of Mental*  
18 *Health*, 19, 99-106.  
19
- 20 Chopra, G. & Smith, J. (1974) Psychotic reactions following cannabis in East  
21 Indians. *Archives of General Psychiatry*, 30, 24-27.  
22
- 23 Clark, R. E., Teague, G. B., Ricketts, S. K., *et al.* (1998) Cost-effectiveness of  
24 assertive community treatment versus standard case management for persons  
25 with co-occurring severe mental illness and substance use disorders. *Health*  
26 *Services Research*, 33, 1285-1308.  
27
- 28 Cleary, M., Hunt, G. E., Matheson, S., *et al.* (2008) Psychosocial treatment  
29 programs for people with both severe mental illness and substance misuse.  
30 *Schizophrenia Bulletin*, 34, 226-228.  
31
- 32 Cleary, M., Hunt, G. E., Matheson, S., *et al.* (2009) Psychosocial treatments for  
33 people with co-occurring severe mental illness and substance misuse:  
34 systematic review. *Journal of Advanced Nursing*, 65, 238-258.  
35
- 36 Cochrane Collaboration (2008) *Review Manager (RevMan)*. Version 5.0. Oxford:  
37 The Cochrane Collaboration.  
38
- 39 Conrod, P. J. & Stewart, S. H. (2005) A critical look at dual-focused cognitive-  
40 behavioral treatments for comorbid substance use and psychiatric disorders:  
41 strengths, limitations, and future directions. *Journal of Cognitive Psychotherapy:*  
42 *An International Quarterly*, 19, 265-289.

- 1  
2 Copello, A., Orford, J., Hodgson, R., *et al.* (2002) Social behaviour and network  
3 therapy basic principles and early experiences. *Addictive Behaviours*, 27, 345-  
4 366.
- 5  
6 Costain, W. (2008) The effects of cannabis abuse on the symptoms of  
7 schizophrenia: patient perspectives. *International Journal of Mental Health*  
8 *Nursing*, 17, 227-235.
- 9  
10 Craig, T., Johnson, S., McCrone, P., *et al.* (2008) Integrated care for co-  
11 occurring disorders: psychiatric symptoms, social functioning and service  
12 costs at 18 months. *Psychiatric Services*, 59, 276-282.
- 13  
14 Crome, I. & Bloor, R. (2005) Substance misuse and psychiatric comorbidity in  
15 adolescents. *Child and Adolescent Psychiatry*, 18, 435-439.
- 16  
17 Crowther, R., Marshall, M., Bond, G., *et al.* (2001) Vocational rehabilitation for  
18 people with severe mental illness. *The Cochrane Database of Systematic Reviews*,  
19 2, CD003080.
- 20  
21 Cuffel, B. J., Shumway, M., Chouljilan, T. A., *et al.* (1994) A longitudinal study  
22 of substance use and community violence in schizophrenia. *Journal of Nervous*  
23 *and Mental Disease*, 182, 704-708.
- 24  
25 De Leon, G., Sacks, S., Staines, G., *et al.* (2000) Modified therapeutic  
26 community for homeless mentally ill chemical abusers: treatment outcomes.  
27 *The American Journal of Drug and Alcohol Abuse*, 26, 461-480.
- 28  
29 Department of Health (1999) *National Service Framework for Mental Health:*  
30 *Modern Standards and Service Models*. London: Department of Health.  
31 Available at:  
32 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_4009598](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_4009598)  
33
- 34  
35 Department of Health (2001) *The Mental Health Policy Implementation Guide*.  
36 London: Department of Health. Available at:  
37 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_4009350](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_4009350)  
38
- 39  
40 Department of Health (2002) *Mental Health Policy Implementation Guide: Dual*  
41 *Diagnosis Good Practice Guide*. London: Department of Health. Available at:

1 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_4009058](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_4009058)

2  
3

4 Department of Health, (2004) *National Service Framework for Children, Young People and Maternity Services*. London: Department of Health. Available at:  
5 <http://www.dh.gov.uk/en/Policyandguidance/Healthandsocialcaretopics/ChildrenServices/Childrenservicesinformation/index.htm>

6  
7  
8

9 Department of Health (2006) *Dual Diagnosis in Mental Health Inpatient and Day Hospital Settings*. London: Department of Health. Available at:

10 <http://www.nmhd.org.uk/silo/files/dual-diagnosis-in-mental-health--inpatient-and-day-hospital-settings.pdf>

11  
12  
13

14 Department of Health (2007) *Drug Misuse and Dependence: UK Guidelines on Clinical Management*. London: Department of Health (England), the Scottish Government, Welsh Assembly Government and Northern Ireland Executive. Available at: [http://www.nta.nhs.uk/uploads/clinical\\_guidelines\\_2007.pdf](http://www.nta.nhs.uk/uploads/clinical_guidelines_2007.pdf)

15  
16  
17  
18

19 Department of Health (2008) *Code of Practice: Mental Health Act 1983*. London: The Stationery Office. Available at:

20 [http://www.dh.gov.uk/prod\\_consum\\_dh/groups/dh\\_digitalassets/@dh/@en/documents/digitalasset/dh\\_087073.pdf](http://www.dh.gov.uk/prod_consum_dh/groups/dh_digitalassets/@dh/@en/documents/digitalasset/dh_087073.pdf)

21  
22  
23

24 Department of Health (2009a) *The Bradley Report: Lord Bradley's Review of People with Mental Health Problems or Learning Disabilities in the Criminal Justice System*. London: Department of Health. Available at:

25 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_098694](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_098694)

26  
27  
28  
29

30 Department of Health (2009b) *Improving Health, Supporting Justice: The National Delivery Plan of the Health and Criminal Justice Programme Board*. London: Department of Health. Available at:

31 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_108606](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_108606)

32  
33  
34  
35

36 Department of Health & National Treatment Agency for Substance Misuse (2006) *Models of Care for Treatment of Alcohol Misusers (MoCAM)*. London: National Treatment Agency. Available at:

37 [http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH\\_4136806](http://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_4136806)

38  
39  
40  
41

- 1 Derry, A. (2008) The clinical response to substance use problems in forensic  
2 mental health services. *The British Journal of Forensic Practice*, 10, 20-23.  
3
- 4 Derry, D. & Batson, A. (2008) Getting out and staying out: does substance use  
5 treatment have an effect on outcome of mentally disordered offenders after  
6 discharge from medium secure service? *The British Journal of Forensic Practice*,  
7 10, 13-17.  
8
- 9 DerSimonian, R. & Laird, N. (1986) Meta-analysis in clinical trials. *Controlled*  
10 *Clinical Trials*, 7, 177-187.  
11
- 12 Dickey, B. & Azeni, H. (1996) Persons with dual diagnoses of substance abuse  
13 and major mental illness: their excess costs of psychiatric care. *American*  
14 *Journal of Public Health*, 86, 973-977.
- 15 Dinos, S., Stevens, S., Serfaty, M., *et al.* (2004) Stigma: the feelings of  
16 experiences of 46 people with mental illness. *British Journal of Psychiatry*, 184,  
17 176-181.  
18
- 19 Dixon, L. (1999) Dual diagnosis of substance abuse in schizophrenia:  
20 prevalence and impact on outcomes. *Schizophrenia Research*, 35, 93-100.  
21
- 22 Donoghue, K., Medley, I., Brewin, J., *et al.* (2009) The association between  
23 substance misuse and first episode psychosis in a defined UK geographical  
24 area during the 1990s. *Social Psychiatry and Psychiatric Epidemiology*, DOI:  
25 10.1007/s00127-009-0175-5.  
26
- 27 Drake, R. E., Bartels, S. J., Teague, G. M., *et al.* (1993). Treatment of substance  
28 abuse in severely mentally ill patients. *Journal of Nervous and Mental Disease*,  
29 181, 606-611.  
30
- 31 Drake, R. E., McHugo, G. J., Clark, R. E., *et al.* (1998) Assertive community  
32 treatment for patients with co-occurring severe mental illness and substance  
33 use disorder: a clinical trial. *American Journal of Orthopsychiatry*, 68, 201-215.  
34
- 35 Drake, R. E., Mueser, K. T., Brunette, M. F., *et al.* (2004) A review of treatments  
36 for people with severe mental illnesses and co-occurring substance use  
37 disorders. *Psychiatric Rehabilitation Journal*, 27, 360-374.  
38
- 39 Drake, R. E., Noordsky, D. L. & Ackerson, T. (1995) Integrating mental health  
40 and substance abuse treatments for persons with chronic mental disorders: a  
41 model. In *Double Jeopardy: Chronic Mental Illness and Substance Use Disorders*

- 1 (eds A. Lehman & L. B. Dixon), vol. 3. Langhorne, PA: Harwood Academic  
2 Press.
- 3
- 4 Drake, R. E., O'Neal, E. L. & Wallach, M. A. (2008) A systematic review of  
5 psychosocial research on psychosocial interventions for people with co-  
6 occurring severe mental health and substance use disorders. *Journal of*  
7 *Substance Use Treatment*, 34, 123-138.
- 8
- 9 Drake, R. E., Osher, F. C. & Wallach, M. A. (1991) Homelessness and dual  
10 diagnosis. *American Psychologist*, 46, 1149-1158.
- 11
- 12 Drake, R. E., Yovetich, N. A., Bebout, R. R., et al. (1997) Integrated treatment  
13 for dually diagnosed homeless adults. *The Journal of Nervous and Mental*  
14 *Disease*, 185, 298-305.
- 15
- 16 D'Silva, K. & Ferriter, M. (2003) Substance use by the mentally disordered  
17 committing serious offences: a high-security hospital study. *Journal of Forensic*  
18 *Psychiatry & Psychology*, 14, 178-193.
- 19
- 20 Eccles, M., Freemantle, N. & Mason, J. (1998) North of England evidence  
21 based guideline development project: methods of developing guidelines for  
22 efficient drug use in primary care. *British Medical Journal*, 316, 1232-1235.
- 23
- 24 Edwards, J., Elkins, K., Hinton, M., et al. (2006) Randomized controlled trial of  
25 a cannabis-focused intervention for young people with first-episode  
26 psychosis. *Acta Psychiatrica Scandinavica*, 114, 109-117.
- 27
- 28 Essock, S. M., Mueser, J. K. T., Drake, R. E., et al. (2006) Comparison of ACT  
29 and standard case management for delivering integrated treatment for co-  
30 occurring disorders. *Psychiatric Services*, 57, 185-196.
- 31
- 32 Fals-Stewart, W., Klosterman, K., Yates, B. T., et al. (2005) Brief relationship  
33 therapy for alcoholism: a randomized clinical trial examining clinical efficacy  
34 and cost-effectiveness. *Psychology of Addictive Behaviors*, 19, 363-371.
- 35
- 36 Fals-Stewart, W., O'Farrell, T. J., Birchler, G. R., et al. (2004) *Behavioral Couples*  
37 *Therapy for Drug Abuse and Alcoholism: A 12-session Manual*. Buffalo, NY:  
38 Addiction and Family Research Group.
- 39
- 40 Farren, C. K., Hameedi, F. A., Rosen, M. A., et al. (2000) Significant interaction  
41 between clozapine and cocaine in cocaine addicts. *Drug & Alcohol Dependency*,  
42 59, 153-163.

- 1  
2 Fazel, S., Gulati, G., Linsell, L., *et al.* (2009a) Schizophrenia and violence:  
3 systematic review and meta-analysis. *Public Library of Science Medicine*, 6,  
4 e1000120.  
5  
6 Fazel, S., Långström, N., Hjern, A., *et al.* (2009b) Schizophrenia, substance  
7 misuse, and violent crime. *Journal of the American Medical Association*, 301, 2016-  
8 2023.  
9  
10 Feinstein, A. (1970) The pre-therapeutic classification of comorbidity in  
11 chronic disease. *Journal of Chronic Disease*, 23, 455-462.  
12  
13 French, M. T., Sacks, S., De Leon, G., *et al.* (1999) Modified therapeutic  
14 community for mentally ill chemical abusers: outcomes and costs. *Evaluation*  
15 *and the Health Professions*, 1, 60-85.  
16  
17 Frisher, M., Collins, J., Millson, D., *et al.* (2004) Prevalence of comorbid  
18 psychiatric illness and substance misuse in primary care in England and  
19 Wales. *Journal of Epidemiology and Community Health*, 58, 1036-1041.  
20  
21 Geller, B., Cooper, T. B., Sun, K., *et al.* (1998) Double-blind and placebo-  
22 controlled study of lithium for adolescent bipolar disorders with secondary  
23 substance dependency. *Journal of the American Academy of Child and Adolescent*  
24 *Psychiatry*, 37, 171-178.  
25  
26 Ghodse, H. (1986). Cannabis psychosis. *British Journal of Addiction*, 81, 473-478.  
27  
28 Ghodse, H., Oyefeso, A. & Kilpatrick, B. (1998) Mortality of drug addicts in  
29 the United Kingdom (1967-1993). *International Journal of Epidemiology*, 27, 473-  
30 478.  
31  
32 Goldberg, J. F., Brooks, J. O., Kurita, K., *et al.* (2009) Depressive illness burden  
33 associated with complex polypharmacy. *Journal of Clinical Psychiatry*, 70, 155-  
34 162.  
35  
36 Goldstein, B. I., Diamantouros, A., Schaffer, A., *et al.* (2006) Pharmacotherapy  
of alcoholism in patients with co-morbid psychiatric disorders. *Drugs*, 66,  
1229-1237.  
37  
38 Graeber, D. A., Moyers, T. B., Griffith, G., *et al.* (2003) A pilot study comparing  
motivational interviewing and an educational intervention in patients with

- 1 schizophrenia and alcohol use disorders. *Community Mental Health Journal*, 39,  
2 189-202.
- 3
- 4 Graham, H. L., Copello, A., Birchwood, M. J., *et al.* (2003) The combined  
5 psychosis and substance use (COMPASS) programme: an integrated, shared  
6 care approach. In *Substance Misuse in Psychosis: Approaches to Treatment and*  
7 *Service Delivery* (eds H. L. Graham, A. Copello, M. J. Birchwood, *et al.*).  
8 Chichester: John Wiley & Sons.
- 9
- 10 Green, A. I. (2005) Schizophrenia and comorbid substance use disorder:  
11 effects of antipsychotics. *Journal of Clinical Psychiatry*, 66, 21-26.
- 12 Green, A. I., Noordsy, D. L., Brunette, M. F., *et al.* (2008) Substance abuse and  
13 schizophrenia: pharmacotherapeutic intervention. *Journal of Substance Abuse*  
14 *Treatment*, 34, 61-71.
- 15 Green, A., Tohenc, M., Hamer, R. M., *et al.* (2004) First episode schizophrenia-  
16 related psychosis and substance use disorders: acute response to olanzapine  
17 and haloperidol. *Schizophrenia Research*, 66, 125-135.
- 18
- 19 Gregg, L., Barrowclough, C., & Haddock, G. (2007) Reasons for increased  
20 substance use in psychosis. *Clinical Psychology Review*, 27, 494-510.
- 21
- 22 Gregg, L., Barrowclough, C. & Haddock, G. (2009) Development and  
23 validation of a scale for assessment of reasons for substance use in  
24 schizophrenia: the ReSUS scale. *Addictive Behaviours*, 34, 830-837.
- 25
- 26 Haddock, G., Barrowclough, C., Tarrier, N., *et al.* (2003) Cognitive-behavioural  
27 therapy and motivational intervention for schizophrenia and substance  
28 misuse: 18-month outcomes of a randomised controlled trial. *British Journal of*  
29 *Psychiatry*, 183, 418-426.
- 30 Haddock, G., Lewis, S., Bentall, R., *et al.* (2006) Influence of age on outcome of  
31 psychological treatment in first episode psychosis. *British Journal of Psychiatry*,  
32 188, 250-254.
- 33
- 34 Hambrecht, M. & Hafner, H. (2000) Cannabis, vulnerability, and the onset of  
35 schizophrenia: an epidemiological perspective. *Australian and New Zealand*  
36 *Journal of Psychiatry*, 34, 468-475.
- 37
- 38 Handmaker, N., Packard, M. & Conforti, K. (2002) Motivational interviewing  
39 in the treatment of dual disorders. In *Motivational Interviewing: Preparing*

- 1 *People for Change* (2nd edn) (eds W. R. Miller & S. Rollnick). New York:  
2 Guildford Press.
- 3
- 4 Hawkins, R. L. & Abrams, C. (2007) Disappearing acts: the social networks of  
5 formerly homeless individuals with co-occurring disorders. *Social Science &*  
6 *Medicine*, 65, 2031-2042.
- 7
- 8 Hawton, K., Sutton, L., Haw, C., *et al.* (2005) Schizophrenia and suicide:  
9 systematic review of risk factors. *British Journal of Psychiatry*, 187, 9-20.
- 10
- 11 Healey, C., Peters, S., Kinderman, P., *et al.* (2009) Reasons for substance use in  
12 dual diagnosis bipolar disorder and substance use disorders: a qualitative  
13 study. *Journal of Affective Disorders*, 113, 118-126.
- 14
- 15 Health Advisory Service (1995) *Together We Stand: The Commissioning, Role and*  
16 *Management of Child and Adolescent Mental Health Services*. London: The  
17 Stationery Office.
- 18 Health Advisory Service (1996) *Children and Young People Substance Misuse*  
19 *Services: The Substance of Young Needs*. Norwich: HMSO.
- 20
- 21 Health Advisory Service (2001) *Children and Young People Substance Misuse*  
22 *Services: The Substance of Young Needs Review 2001*. London: Drug Prevention  
23 Advisory Service.
- 24
- 25 Healthcare Commission (2007) *National Audit of Violence 2006-7 Final Report –*  
26 *Working Age Adult Services*. London: Royal College of Psychiatrists. Available  
27 at:  
28 [http://www.rcpsych.ac.uk/pdf/WAA%20Nat%20Report%20final%20with%](http://www.rcpsych.ac.uk/pdf/WAA%20Nat%20Report%20final%20with%20all%20appendices.pdf)  
29 [20all%20appendices.pdf](http://www.rcpsych.ac.uk/pdf/WAA%20Nat%20Report%20final%20with%20all%20appendices.pdf)
- 30
- 31 Hellerstein, D. J., Rosenthal, R. N. & Miner, C. R. (1995) A prospective study  
32 of integrated outpatient treatment for substance-abusing schizophrenic  
33 patients. *American Journal on Addictions*, 4, 33-42.
- 34
- 35 Helmus, T. C., Saules, K. K., Shoener, E. P., *et al.* (2003) Reinforcement of  
36 counselling attendance and alcohol abstinence in a community-based dual-  
37 diagnosis treatment program: a feasibility study. *Psychology of Addictive*  
38 *Behaviors*, 17, 249-251.
- 39
- 40 Henggeler, S. W., Pickrel, S. G. & Brondino, M. J. (1999) Multi-systemic  
41 treatment of substance-abusing and dependent delinquents: outcomes,

- 1 treatment fidelity, and transportability. *Mental Health Services Research*, 1, 171-  
2 184.
- 3
- 4 Hickman, M., Vickerman, P., Macleod, J., *et al.* (2009). If cannabis caused  
5 schizophrenia - how many cannabis users may need to be prevented in order  
6 to prevent one case of schizophrenia? England and Wales calculations.  
7 *Addiction*, 104, 1856-1861.
- 8
- 9 Hides, L., Dawe, S., Kavanagh, D. J., *et al.* (2006) Psychotic symptom and  
10 cannabis relapse in recent onset psychosis. *British Journal of Psychiatry*, 189,  
11 137-143.
- 12
- 13 Higgins, J. P. T. & Green, S. (eds) (2009) *Cochrane Handbook for Systematic*  
14 *Reviews of Interventions*. Version 5.0.2. The Cochrane Collaboration. Available  
15 at: [www.cochrane-handbook.org](http://www.cochrane-handbook.org)
- 16
- 17 Higgins, J. P. T. & Thompson, S. G. (2002) Quantifying heterogeneity in a  
18 meta-analysis. *Statistics in Medicine*, 21, 1539-1558.
- 19
- 20 Hjorthoj, C., Fohlmann, A. & Norentoft, M. (2009) Treatment of cannabis use  
21 disorders in people with schizophrenia spectrum disorders: a systematic  
22 review. *Addictive Behaviours*, 34, 846-851.
- 23
- 24 HMSO (1983) *The Mental Health Act 1983*. London: The Stationery Office.  
25 Available at:  
26 [http://www.cqc.org.uk/\\_db/\\_documents/Mental\\_Health\\_Act\\_1983\\_201005](http://www.cqc.org.uk/_db/_documents/Mental_Health_Act_1983_201005272747.pdf)  
27 [272747.pdf](http://www.cqc.org.uk/_db/_documents/Mental_Health_Act_1983_201005272747.pdf)
- 28
- 29 HMSO (1989) *The Children Act 1989*. London: The Stationery Office. Available  
30 at: [http://www.opsi.gov.uk/acts/acts1989/Ukpga\\_19890041\\_en\\_1.htm](http://www.opsi.gov.uk/acts/acts1989/Ukpga_19890041_en_1.htm)
- 31
- 32 HMSO (1995) *The Mental Capacity Act 1995*. London: The Stationery Office.  
33 Available at: <http://www.legislation.gov.uk/ukpga/2005/9/contents>
- 34
- 35 HMSO (1998a) *The Human Rights Act 1998*. London: The Stationery Office.  
36 Available at:  
37 [http://www.opsi.gov.uk/acts/acts1998/ukpga\\_19980042\\_en\\_1](http://www.opsi.gov.uk/acts/acts1998/ukpga_19980042_en_1)
- 38
- 39 HMSO (1998b) *Crime and Disorder Act 1998*. London: The Stationery Office.  
40 Available at: <http://www.legislation.gov.uk/ukpga/1998/37/contents>
- 41

- 1 HMSO (2004) *The Children Act 2004*. London: The Stationery Office. Available  
2 at: <http://www.legislation.gov.uk/ukpga/2004/31/contents>  
3
- 4 HMSO (2007) *The Mental Health Act 2007*. London: The Stationery Office.  
5 Available  
6 at: [http://www.opsi.gov.uk/acts/acts2007/pdf/ukpga\\_20070012\\_en.pdf](http://www.opsi.gov.uk/acts/acts2007/pdf/ukpga_20070012_en.pdf)  
7
- 8 Ho, A. P., Tsuang, J. W., Liberman, R. P., *et al.* (1999) Achieving effective  
9 treatment of patients with chronic psychotic illness and comorbid substance  
10 dependence. *American Journal of Psychiatry*, 156, 1765-1770.  
11
- 12 Hodgson, R. J. & Rankin, H. J. (1976) Modification of excessive drinking by  
13 cue exposure. *Behaviour Research and Therapy*, 14, 305-307.  
14
- 15 Hoff, R. & Rosenheck, R. (1998) Long-term patterns of service use and cost  
16 among patients with both psychiatric and substance abuse disorders. *Medical  
17 Care*, 36, 98-843.  
18
- 19 Hosák, L. (2007) Role of the COMT gene Val158Met polymorphism in mental  
20 disorders: a review. *European Psychiatry*, 22, 276-281.  
21
- 22 Howat, J., Bates, P., Pidgeon, J., *et al.* (1988) The development of residential  
23 care in the community. In *Community Care in Practice: Services for the  
24 Continuing Care Client* (eds A. Lavender & F. Holloway), pp. 275-293.  
25 Chichester: John Wiley & Sons.  
26
- 27 Hughes, E. (2006). *Closing the Gap: A Capability Framework for Effectively  
28 Working with People with Combined Mental Health and Substance Use Problems  
29 (Dual Diagnosis)*. Mansfield: University of Lincoln.  
30
- 31 Hughes, E., Wanigaratne, S., Gournay, K., *et al.* (2008) Training in dual  
32 diagnosis interventions (the COMO study): randomised controlled trial. *BMC  
33 Psychiatry*, 8, 1-9.  
34
- 35 Hunt, G. M., & Azrin, N. H. (1973) A community-reinforcement approach to  
36 alcoholism. *Behaviour Research and Therapy*, 11, 91-104.  
37
- 38 Huntley, D. A., Cho, D. W., Christman, J., *et al.* (1998) Predicting length of stay  
39 in an acute psychiatric hospital. *Psychiatric Services*, 49, 1049-1053.  
40

- 1 Isaac, M., Isaac, M., & Holloway, F. (2005) Is cannabis an anti-antipsychotic?  
2 The experience in psychiatric intensive care. *Human Psychopharmacology:*  
3 *Clinical and Experimental*, 20, 207-210.  
4
- 5 Isherwood, S. & Brooke, D. (2001) Prevalence and severity of substance  
6 misuse among referrals to a local forensic service. *Journal of Forensic Psychiatry*,  
7 12, 446-454.  
8
- 9 Jackson, C. T., Covell, N. H., Drake, R. E., *et al.* (2007) Relationship between  
10 diabetes and mortality among persons with co-occurring psychotic and  
11 substance use disorders. *Psychiatric Services*, 58, 270-272.  
12
- 13 Jadad, A. R., Moore, R. A., Carroll, D., *et al.* (1996) Assessing the quality of  
14 reports of randomised clinical trials: is blinding necessary? *Controlled Clinical*  
15 *Trials*, 17, 1-12.  
16
- 17 James, W., Preston, N. J., Koh, G., *et al.* (2004) A group intervention which  
18 assist patients with dual diagnosis reduce their drug use: a randomized  
19 controlled trial. *Psychological Medicine*, 34, 983-990.  
20
- 21 Jerrell, J. M., & Ridgely, M. S. (1995) Comparative effectiveness of three  
22 approaches to serving people with severe mental illness and substance abuse  
23 disorders. *The Journal of Nervous and Mental Disease*, 183, 566-576.  
24
- 25 Jerrell, J. M. & Ridgely, M. S. (1997) Dual diagnosis care for severe and  
26 persistent disorders: a comparison of three methods. *Behavioural Healthcare*  
27 *Tomorrow*, 6, 26-33.  
28
- 29 Johnson, E. D. (2000) Differences among families coping with serious mental  
30 illness: a qualitative analysis. *American Journal of Orthopsychiatry*, 70, 126-134.  
31
- 32 Johnson, S., Thornicroft, G., Afuwape, S., *et al.* (2007) Effects of training  
33 community staff for in interventions for substance misuse in dual diagnosis  
34 patients with psychosis (COMO study): cluster randomised trial. *British*  
35 *Journal of Psychiatry*, 191, 451-452.  
36
- 37 Joint RCPCH/NPPG Standing Committee on Medicines (2000) *The Use of*  
38 *Unlicensed Medicines for Unlicensed Applications in Paediatric Practice: Policy*  
39 *Statement*. London: Royal College of Paediatrics and Child Health.  
40

- 1 Kadden, R., Litt, M. D., Cooney, N. L., *et al.* (1992) Relationship between role-  
2 play measures of coping skills and alcoholism treatment outcome. *Addictive*  
3 *Behaviors*, 17, 425-437.  
4
- 5 Kavanagh, D. J., Waghorn, G., Jenner, L., *et al.* (2004a) Demographic and  
6 clinical correlates of comorbid substance use disorders in psychosis:  
7 multivariate analyses from an epidemiological sample. *Schizophrenia Research*,  
8 66, 115-124.  
9
- 10 Kavanagh, D. J., Young, R., White, A., *et al.* (2004b) A brief motivational  
11 intervention for substance misuse in recent-onset psychosis. *Drug and Alcohol*  
12 *Review*, 23, 151-155.  
13
- 14 Kemp, D. E., Gao, K., Ganocy, S. J., *et al.* (2009) A 6-month, double-blind,  
15 maintenance trial of lithium monotherapy versus the combination of lithium  
16 and divalproex for rapid-cycling bipolar disorder and co-occurring substance  
17 abuse or dependence. *Journal of Clinical Psychiatry*, 70, 113-121.  
18
- 19 Kemp, R., Harris, A., Vurel, E., *et al.* (2007) Stop Using Stuff: trial of a drug  
20 and alcohol intervention for young people with comorbid mental illness and  
21 drug and alcohol problems. *Australasian Psychiatry*, 15, 490-493.  
22
- 23 Kessler, R. C., McGongale, K. A., Zhao, S., *et al.* (1994) Lifetime and 12-  
24 month prevalence of DSM-III-R psychiatric disorders in the United States:  
25 results from the National Comorbidity Survey. *Archives of General Psychiatry*,  
26 51, 8-19.  
27
- 28 Knudsen, H.K. (2009) Adolescent-only substance treatment: Availability and  
29 adoption of components of quality. *Journal of Substance Abuse Treatment*, 36,  
30 195-204.  
31
- 32 Kooyman, I., Dean, K., Harvey, S., *et al.* (2007) Outcomes of public concern in  
33 schizophrenia. *British Journal of Psychiatry*, 191, 29-36.  
34
- 35 Koskinen, J., Lohonen, J., Koponen, H., *et al.* (2009a) Prevalence of alcohol use  
36 disorders in schizophrenia: a systematic review and meta-analysis. *Acta*  
37 *Psychiatrica Scandinavica*, 120, 85-96.  
38
- 39 Koskinen, J., Lohonen, J., Koponen, H., *et al.* (2009b) Rate of cannabis use  
40 disorders in clinical samples of patients with schizophrenia: a meta-analysis.  
41 *Schizophrenia Bulletin*, 36, 1115-1130.  
42

- 1 Kreyenbuhl, J. A., Valenstein, M., McCarthy, J. F., *et al.* (2007) Long-term  
2 antipsychotic polypharmacy in the VA health system: patient characteristics  
3 and treatment patterns. *Psychiatric Services*, 58, 489-495.  
4
- 5 Lehman, A., Kreyenbuhl, J., Buchanan, R., *et al.* (2004) The Schizophrenia  
6 Patient Outcomes Research Team (PORT): updated treatment  
7 recommendations 2003. *Schizophrenia Bulletin*, 30, 193-217.  
8
- 9 Lehman, A. F., Meyers, C. P. & Corty, E. (1989) Classification of patients with  
10 psychiatric and substance misuse syndromes. *Hospital and Community*  
11 *Psychiatry*, 40, 1019-1025.  
12
- 13 Lingford-Hughes, A. R., Welch, S. & Nutt, D. J. (2004) Evidence-based  
14 guidelines for the pharmacological management of substance misuse,  
15 addiction and comorbidity: recommendations from the British Association for  
16 Psychopharmacology. *Journal of Psychopharmacology*, 18, 293-335.
- 17 Linszen, D. H., Dingerms, P. M. & Lenior, M.E. (1994) Cannabis abuse and  
18 the course of recent-onset schizophrenic disorders. *Archives of General*  
19 *Psychiatry*, 51, 706-712.  
20
- 21 Lobban, F., Barrowclough, C., Jeffery, S., *et al.* (2010) Understanding factors  
22 influencing substance use in people with recent onset psychosis: a qualitative  
23 study. *Social Science & Medicine*, 70, 1141-1147.  
24
- 25 Loneck, B. & Way, B. (1997) Using a focus group of clinicians to develop a  
26 research project on therapeutic process with clients with dual diagnoses.  
27 *Social Work*, 42, 107-111.
- 28 Loubser, I., Chaplin, R. & Quirk, A. (2009) Violence, alcohol and drugs: the  
29 views of nurses and patients on psychiatric intensive care units, acute adult  
30 wards and forensic wards. *Journal of Psychiatric Intensive Care*, 5, 33-39.  
31
- 32 Lykke, J., Oestrich, I., Austin, S. F., *et al.* (2010) The implementation and  
33 evaluation of cognitive milieu therapy for dual diagnosis inpatients: a  
34 pragmatic clinical trial. *Journal of Dual Diagnosis*, 6, 58-72.  
35
- 36 Macdonald, S., Erickson, P., Wells, S., *et al.* (2008) Predicting violence among  
37 cocaine, cannabis, and alcohol treatment clients. *Addictive Behaviors*, 33, 201  
38 -205.  
39

- 1 Macpherson, R., Shepherd, G. & Edwards, T. (2004) Supported  
2 accommodation for people with severe mental illness: a review. *Advances in*  
3 *Psychiatric Treatment*, 10, 180-188.  
4
- 5 Mangrum, L. F., Spence, R. T. & Lopez, M. (2006) Integrated versus parallel  
6 treatment of co-occurring psychiatric and substance use disorders. *Journal of*  
7 *Substance Abuse Treatment*, 30, 79-84.  
8
- 9 Mann, T. (1996) *Clinical Guidelines: Using Clinical Guidelines to Improve Patient*  
10 *Care Within the NHS*. London: Department of Health NHS Executive.  
11
- 12 Margoles, H. C., Malchy, L., Negrete, J. C., *et al.* (2004) Drug and alcohol use  
13 among patients with schizophrenia and related psychoses: levels and  
14 consequences. *Schizophrenia Research*, 67, 157-166.  
15
- 16 Marlatt, G. A. & Gordon, J. R. (1985) *Relapse Prevention: Maintenance Strategies*  
17 *in the Treatment of Addictive Behaviours*. New York: Guildford Press.  
18
- 19 Martino, S., Carroll, K., Kostas, D., *et al.* (2002) Dual diagnosis motivational  
20 interviewing: a modification of motivational interviewing for substance  
21 abusing patients with psychotic disorders. *Journal of Substance Abuse*  
22 *Treatment*, 23, 297-308.  
23
- 24 Matrix Research and Consultancy & NACRO (2004) *Home Office Research*  
25 *Study 286: Evaluation of Drug Testing in the Criminal Justice System*. London:  
26 Home Office. Available at:  
27 <http://rds.homeoffice.gov.uk/rds/pdfs04/hors286.pdf>  
28
- 29 McCreadie, R. G. (2002) Use of drugs, alcohol and tobacco by people with  
30 schizophrenia: case-control study. *British Journal of Psychiatry*, 181, 321-325.  
31
- 32 McCrone, P., Dhanasiri, S., Patel, A., *et al.* (2008). *Paying the Price: The Cost of*  
33 *Mental Health Care in England to 2026*. London: King's Fund.  
34
- 35 McCrone, P., Menezes, P. R., Johnson, S., *et al.* (2000) Service use and costs of  
36 people with dual diagnosis in South London. *Acta Psychiatrica Scandinavica*  
37 101, 464-472.  
38
- 39 Menezes, P. O. R., Johnson, S., Thornicroft, G., *et al.* (1996) Drug and alcohol  
40 problems among individuals with severe mental illnesses in south London.  
41 *British Journal of Psychiatry*, 168, 612-619.  
42

- 1 Mercer-McFadden, C., Drake, R., Clark, R. E., *et al.* (1998) *Substance Abuse*  
2 *Treatment for People with Severe Mental Disorders: A Program Manager's Guide.*  
3 New Hampshire: New Hampshire-Dartmouth Psychiatric Research Center.  
4
- 5 Meyers, R. J., & Miller, W. R. (eds) (2001) *A Community Reinforcement Approach*  
6 *to Addiction Treatment.* Cambridge: Cambridge University Press.  
7
- 8 Miles, H., Duthell, L., Welsby, I., *et al.* (2007) "Just say no": a preliminary  
9 evaluation of a three stage model of integrated treatment for substance abuse  
10 problems in conditions of medium security. *Journal of Forensic Psychiatry and*  
11 *Psychology, 18, 141-159.*  
12
- 13 Miller, W. R. & Munoz, R. F. (1976) *How to Control Your Drinking* (1<sup>st</sup> edn).  
14 Albuquerque: University of New Mexico Press.  
15
- 16 Miller, W. R. & Rollnick, S. (2002) *Motivational Interviewing: Preparing People for*  
17 *Change* (2nd edn). New York: Guildford Press.  
18
- 19 Miller, W. R., Zweben, A., DiClemente, C. C., *et al.* (1992) *Motivational*  
20 *Enhancement Therapy Manual: A Clinical Research Guide for Therapists Treating*  
21 *Individuals with Alcohol Abuse and Dependence.* Rockville, MD: National  
22 Institute on Alcohol Abuse and Alcoholism.  
23
- 24 Mills, K. L., Deady, M., Proudfoot, H., *et al.* (2009) *Guidelines on the*  
25 *Management of Co-occurring Alcohol and Other Drug and Mental Health*  
26 *Conditions in Alcohol and Other Drug Treatment Settings.* Sydney: University of  
27 New South Wales.  
28
- 29 Molgaard, V., Kumpfer, K. L. & Spoth, R. (1994) *The Iowa Strengthening*  
30 *Families Program for Pre- and Early Teens.* Ames, IA: Iowa State University.  
31
- 32 Moore, T., Zammit, S., Lingford-Hughes, A., *et al.* (2007) Cannabis use and  
33 risk of psychotic or affective mental health outcomes: a systematic review. *The*  
34 *Lancet, 370, 319-328.*  
35
- 36 Moos, R. H., Finney, J. W. & Moos, B. S. (2000) Inpatient substance abuse care  
37 and the outcome of subsequent community residential and outpatient care.  
38 *Addiction, 95, 833-846.*  
39
- 40 Morse, G. A., Calsyn, R. J., Klinkenberg, W. D., *et al.* (2006) Treating homeless  
41 clients with severe mental illness and substance use disorders: costs and  
42 outcomes. *Community Mental Health Journal, 42, 377-404.*

- 1  
2 Mueser, K. T., Bennett, M. & Kushner, M. G. (1995) Epidemiology of  
3 substance use disorders among persons with chronic mental illnesses. In  
4 *DoubleJeopardy: Chronic Mental Illness and Substance Use Disorders* (eds A. F.  
5 Lehman & L. B. Dixon), vol. 3, pp. 9-25. Langhorne, PA: Harwood Academic  
6 Press.  
7  
8 Mueser, K. T. & Drake, R. E. (2003) Integrated dual diagnosis treatment in  
9 New Hampshire (USA). In *Substance Misuse in Psychosis: Approaches to*  
10 *Treatment and Service Delivery* (eds H. L. Graham, A. Copello, M. J. Birchwood,  
11 *et al.*), pp. 179-206. Chichester: John Wiley & Sons.  
12  
13 Mueser, K. T., Drake, R. E., Sigmon, S. C., *et al.* (2005) Psychosocial  
14 interventions for adults with severe mental illnesses and co-occurring  
15 substance use disorders: a review of specific interventions. *Journal of Dual*  
16 *Diagnosis, 1*, 57-82.  
17  
18 Murray, R. M., Morrison, P. D., Henquet, C., *et al.* (2007) Cannabis, the mind  
19 and society: the hash realities. *Nature Review Neuroscience, 8*, 885-895.  
20  
21 National Treatment Agency for Substance Misuse (2006) *Models of Care for*  
22 *Treatment of Adult Drug Misusers: Update 2006*. London: National Treatment  
23 Agency Publications.  
24  
25 NCCMH (2006) *Bipolar Disorder: The Management of Bipolar Disorder in Adults,*  
26 *Children and Adolescents, in Primary and Secondary Care*. Leicester: The British  
27 Psychological Society and the Royal College of Psychiatrists.  
28  
29 NCCMH (2008a) *Drug Misuse: Opioid Detoxification*. Leicester: The British  
30 Psychological Society and the Royal College of Psychiatrists.  
31  
32 NCCMH (2008b) *Drug Misuse: Psychosocial Interventions*. Leicester: The British  
33 Psychological Society and the Royal College of Psychiatrists.  
34 NCCMH (2010) *Schizophrenia: Core Interventions in the Treatment and*  
35 *Management of Schizophrenia in Adults in Primary and Secondary Care (Update)*.  
36 Leicester: The British Psychological Society & the Royal College of  
37 Psychiatrists.  
38  
39 NCCMH (in press) *Alcohol Use Disorders: Diagnosis, Assessment and*  
40 *Management of Harmful Drinking and Alcohol Dependence*. Leicester: The British  
41 Psychological Society and the Royal College of Psychiatrists.  
42

- 1 NICE (2002) *Schizophrenia: Core Interventions in the Treatment and Management*  
2 *of Schizophrenia in Primary and Secondary Care (Update)*. Clinical guideline 1.  
3 London: NICE. Available at:  
4 <http://www.nice.org.uk/guidance/index.jsp?action=byID&o=10916>  
5
- 6 NICE (2005) *Violence*. Clinical guideline 25. London: NICE. Available at:  
7 <http://www.nice.org.uk/CG25>  
8
- 9 NICE (2006) *Bipolar Disorder: The Management of Bipolar Disorder in Adults,*  
10 *Children and Adolescents, in Primary and Secondary Care*. Clinical guideline 38.  
11 London: NICE. Available at: [www.nice.org.uk/CG38](http://www.nice.org.uk/CG38)  
12
- 13 NICE (2007a) *Drug Misuse: Opioid Detoxification*. Clinical guideline 52.  
14 London: NICE. Available at: [www.nice.org.uk/CG52](http://www.nice.org.uk/CG52)  
15
- 16 NICE (2007b) *Drug Misuse: Psychosocial Interventions*. Clinical guideline 51.  
17 London: NICE. Available at: [www.nice.org.uk/CG51](http://www.nice.org.uk/CG51)  
18
- 19 NICE (2007c) *Interventions to Reduce Substance Misuse Among Vulnerable Young*  
20 *People*. Public health guidance 4. London: NICE. Available at:  
21 [www.nice.org.uk/PH4](http://www.nice.org.uk/PH4)  
22
- 23 NICE (2007d) *Drug Misuse: Naltrexone*. Technology appraisals 115. London:  
24 NICE. Available at: [www.nice.org.uk/TA115](http://www.nice.org.uk/TA115)  
25
- 26 NICE (2007e) *Drug Misuse: Methadone and Buprenorphine*. Technology  
27 appraisals 114. London: NICE. Available at: [www.nice.org.uk/TA114](http://www.nice.org.uk/TA114)  
28
- 29 NICE (2008) *Guide to the Methods of Technology Appraisal*. London: NICE.  
30 Available at: [www.nice.org.uk](http://www.nice.org.uk)  
31
- 32 NICE (2009a) *Schizophrenia: Core Interventions in the Treatment and Management*  
33 *of Schizophrenia in Primary and Secondary Care (Update)*. Clinical guideline 82.  
34 London: NICE. Available at: [www.nice.org.uk/CG82](http://www.nice.org.uk/CG82)NICE (2009b) *The*  
35 *Guidelines Manual*. London: NICE. Available at: [www.nice.org.uk](http://www.nice.org.uk)  
36
- 37 NICE (2009c) *Medicines Adherence*. Clinical guideline 76. London: NICE.  
38 Available at: [www.nice.org.uk/CG76](http://www.nice.org.uk/CG76)  
39
- 40 NICE (2010) *Alcohol-Use Disorders: Physical Complications*. Clinical guideline  
41 100. London: NICE. Available at: [www.nice.org.uk/CG100](http://www.nice.org.uk/CG100)  
42

- 1 NICE (in press) Alcohol Dependence and Harmful Alcohol Use.  
2
- 3 NICE (in press) Alcohol Use Disorders (Clinical Management).  
4
- 5 NICE (in press) Alcohol Use Disorders (Prevention).  
6
- 7 Nejtcek, V. A., Avila, M., Chen, L. A., *et al.* (2008) Do atypical antipsychotics  
8 effectively treat co-occurring bipolar disorder and stimulant dependence? A  
9 randomized, double-blind trial. *Journal of Clinical Psychiatry*, 69, 1257-1266.  
10
- 11 Niaura, R. S., Rohsenow, D. J., Binkoff, J. A., *et al.* (1988) Relevance of cue  
12 reactivity to understanding alcohol and smoking relapse. *Journal of Abnormal*  
13 *Psychology*, 97, 133-152.  
14
- 15 Noordsky, D. L., McQuade, D. V. & Mueser, K. (2003) Assessment  
16 considerations. In *Substance Misuse in Psychosis: Approaches to Treatment and*  
17 *Service Delivery* (eds H. L. Graham, A. Copello, M. J. Birchwood, *et al.*).  
18 Chichester: John Wiley & Sons.  
19
- 20 Nowinski, J., Baker, S. & Carroll, K. (1992) *12-Step Facilitation Therapist Manual:*  
21 *A Clinical Research Guide for Therapists Treating Individuals with Alcohol Abuse*  
22 *and Dependence* (vol. 1, Project MATCH Monograph Series). Rockville, MD:  
23 National Institute on Alcohol Abuse and Alcoholism.  
24
- 25 Nuttbrock, L. A., Rahav, M., Rivera, J. J., *et al.* (1998) Outcomes of homeless  
26 mentally ill chemical abusers in community residences and a therapeutic  
27 community. *Psychiatric Services*, 49, 68-76.  
28
- 29 Padgett, D. K., Henwood, B., Abrams, C., *et al.* (2008a) Social relationships  
30 among persons who have experienced serious mental illness, substance abuse,  
31 and homelessness: implications for recovery. *American Journal of*  
32 *Orthopsychiatry*, 78, 333-339.  
33
- 34 Padgett, D. K., Henwood, B., Abrams, C., *et al.* (2008b) Engagement and  
35 retention in services among formerly homeless adults with co-occurring  
36 mental illness and substance abuse: voices from the margins. *Psychiatric*  
37 *Rehabilitation Journal*, 31, 226-233.  
38
- 39 Patkar, A. A., Alexander, R. C., Lundy, A., *et al.* (1999) Changing patterns of  
40 illicit substance use among schizophrenic patients. *American Journal of*  
41 *Addiction*, 8, 65-71.  
42

- 1 Penn, D. L., Mueser, K. T., Tarrier, N., *et al.* (2004). Supportive therapy for  
2 schizophrenia: possible mechanisms and implications for adjunctive  
3 psychosocial treatments. *Schizophrenia Bulletin*, 30, 101-112.  
4
- 5 Penn, P. E., Brooks, A. J. & Worsham, B. D. (2002) Treatment concerns of  
6 women with co-occurring serious mental illness and substance abuse  
7 disorders. *Journal of Psychoactive Drugs*, 34, 355-362.  
8
- 9 Petry, N. M., Alessi, S. M., Carroll, K. M., *et al.* (2006) Contingency  
10 management treatments: reinforcing abstinence versus adherence with goal-  
11 related activities. *Journal of Consulting and Clinical Psychology*, 74, 592-601.  
12
- 13 Phillips, P. & Johnson, S. (2001) How does drug and alcohol misuse develop  
14 among people with psychotic illness? A literature review. *Social Psychiatry and*  
15 *Psychiatric Epidemiology*, 36, 269-276.  
16
- 17 Pilling, S., Bebbington, P., Kuipers, E., *et al.* (2002) Psychological treatments in  
18 schizophrenia: I. Meta-analysis of family intervention and cognitive  
19 behaviour therapy. *Psychological Medicine*, 32, 763-782.  
20
- 21 Pollack, L. E., Stuebben, G., Kouzekanani, K., *et al.* (1998) Aftercare  
22 compliance: perceptions of people with dual diagnoses. *Substance Abuse*, 19,  
23 33-44.  
24
- 25 Phillips, P. & Johnson, S. (2003) Drug and alcohol misuse among in-patients  
26 with psychotic illness in three inner London psychiatric units. *Psychiatric*  
27 *Bulletin* 27, 217-220.  
28
- 29 Prendergast, M., Podus, D., Finney, J., *et al.* (2006) Contingency management  
30 for treatment of substance use disorders: a meta-analysis. *Addiction*, 101, 1546-  
31 1560.  
32
- 33 Prochaska, J. & DiClemente, C. (1986) Towards a comprehensive model of  
34 change. In *Treating Addictive Behaviours: Processes of Change (Applied Clinical*  
35 *Psychology)* (eds W. Miller & N. Heather). New York: Plenum.  
36
- 37 Prochaska, J., DiClemente, C. & Nocross, J. (1992) In search of how people  
38 change: applications to addictive behaviours. *American Psychologist*, 47, 1102-  
39 1012.  
40

- 1 Project MATCH Research Group (1993) Project MATCH: rationale and  
2 methods for a multisite clinical trial matching patients to alcoholism  
3 treatment. *Alcoholism: Clinical and Experimental Research*, 17, 1130-1145.  
4
- 5 Raistrick, D., Heather, N., Godfrey, C., *et al.* (2006) *Review of the Effectiveness of*  
6 *Treatment for Alcohol Problems*. London: National Treatment Agency.  
7
- 8 Ranger, M., Tyrer, P., Milošeska, K., *et al.* (2009) Cost effectiveness  
9 of nidothrapy for comorbid personality disorder and severe mental  
10 illness: randomized controlled trial. *Epidemiologia e Psichiatria Sociale*, 18, 128-  
11 136.  
12
- 13 Rawson, R. A., Mann, A. J., Tennant, F. S., *et al.* (1983) Efficacy of  
14 psychotherapeutic counselling during 21-day ambulatory heroin  
15 detoxification. *Drug and Alcohol Dependence*, 12, 197-200.  
16
- 17 Regier, D. A., Farmer, M. E., Rae, D. S., *et al.* (1990) Comorbidity of mental  
18 disorders with alcohol and other drug abuse. results from the Epidemiologic  
19 Catchment Area (ECA) Study. *Journal of the American Medical Association*, 264,  
20 2511-2518.  
21
- 22 Ries, R. K., Dyck, D. G., Short, R., *et al.* (2004) Outcomes of managing  
23 disability benefits among patients with substance dependence and severe  
24 mental illness. *Psychiatric Services*, 55, 445-447.  
25
- 26 Rosenheck, R. & Fontana, A. (2001) Impact of efforts to reduce inpatient costs  
27 on clinical effectiveness: treatment of posttraumatic stress disorder in the  
28 Department of Veterans Affairs. *Medical Care*, 39, 168-80.  
29
- 30 Rowe, C. L., Liddle, H. A., Greenbaum, P. E., *et al.* (2004) Impact of psychiatric  
31 mental health substance use on treatment of adolescent drug users. *Journal of*  
32 *Substance Abuse Treatment*, 26, 129-140.  
33
- 34 Royal College of Psychiatrists (2006) *Consensus Statement on High-Dose*  
35 *Antipsychotic Medication*. College Report, vol. 138. London: Royal College of  
36 Psychiatrists.  
37
- 38 Sainsbury Centre for Mental Health (2008) *Short-Changed: Spending on Prison*  
39 *Mental Health Care*. London: Sainsbury Centre for Mental Health.  
40

- 1 Salyers, M. P. & Mueser, K. T. (2001) Social functioning, psychopathology,  
2 and medication side effects in relation to substance use and abuse in  
3 schizophrenia. *Schizophrenia Research*, 48, 109-123.  
4
- 5 Samet, S., Nunes, E. V. & Hasin, D. (2004) Diagnosing comorbidity: concepts,  
6 criteria, and methods. *Acta Neuropsychiatrica*, 16, 9-18.  
7
- 8 San, L., Arranz, B., & Martinez-Raga, J. (2007) Antipsychotic drug treatment of  
9 schizophrenia patients with substance abuse disorder. *European Addiction*  
10 *Research*, 13, 230-243.  
11
- 12 Santa Ana, E. J., Wulfert, E., & Nietert, P. K. (2007) Efficacy of group  
13 motivational interviewing (GMI) for psychiatric inpatients with chemical  
14 dependence. *Journal of Consulting and Clinical Psychology*, 75, 816-822.  
15
- 16 Schmitz, J. M., Averill, P., Sayre, S., *et al.* (2002) Cognitive-behavioural  
17 treatment of bipolar disorder and substance abuse: a preliminary randomized  
18 study. *Addictive Disorders and their Treatment*, 1, 17-24.  
19
- 20 Schneier, F. R., & Siris, S. G. (1987) A review of psychoactive substance use  
21 and abuse in schizophrenia: patterns of drug choice. *Journal of Nervous and*  
22 *Mental Disease*, 175, 641-652.  
23
- 24 Schunemann, H. J., Best, D., Vist, G., *et al.* (2003) Letters, numbers, symbols  
25 and words: how to communicate grades of evidence and recommendations.  
26 *Canadian Medical Association Journal*, 169, 677-680.  
27
- 28 Scott, F., Whyte, S., Burnett, R., *et al.* (2004) A National survey of substance  
29 misuse and treatment outcome in psychiatric patients in medium security.  
30 *Journal of Forensic Psychiatry and Psychology*, 15, 595-605.  
31
- 32 Sinclair, J. M. A., Latifi, A. H. & Latifi, A. W. (2008) Comorbid substance  
33 misuse in psychiatric patients: prevalence and association with length of  
34 inpatient stay. *Journal of Psychopharmacology*, 22, 92-99.  
35
- 36 Singleton, N., Bumpstead, R., O'Brien, M., *et al.* (2000) *Psychiatric Morbidity*  
37 *Among Adults Living in Private Households, 2000*. Report of a survey carried out  
38 by the Social Survey Division of the Office for National Statistics on behalf of  
39 the Department of Health, the Scottish Executive and the National Assembly  
40 for Wales. London: HMSO.  
41

- 1 Sisson, R. W., & Azrin, N. H. (1986) Family-member involvement to initiate  
2 and promote treatment of problem drinkers. *Behavior Therapy and Experimental*  
3 *Psychiatry*, 17, 15-21.  
4
- 5 Smelson, D. A., Dixon, K., Craig, T., *et al.* (2008) Pharmacological treatment of  
6 schizophrenia and co-occurring substance use disorders. *CNS Drugs*, 22, 903-  
7 916.  
8
- 9 Sobell, M. B., & Sobell, L. C. (1993) *Problem Drinkers: Guided self-change*  
10 *Treatment*. New York: Guilford Press.  
11
- 12 Spencer, C., Castle, D. & Michie, P. T. (2002) Motivations that Maintain  
13 substance use among individuals with psychotic disorders. *Schizophrenia*  
14 *Bulletin*, 28, 233-247.  
15
- 16 Spoth, R., Redmond, C., Shin, C., *et al.* (2004) Brief family intervention effects  
17 on adolescent substance initiation: school-level growth curve analyses 6 years  
18 following baseline. *Journal of Consulting and Clinical Psychology* 72, 535-542.  
19
- 20 Stewart, S. H., & Conrod, P. J. (2005) Introduction to the special issue on state-  
21 of-the-art in cognitive-behavioural interventions for substance use disorders.  
22 *Journal of Cognitive Psychotherapy: An International Quarterly*, 19, 195-198.  
23
- 24 Stitzer, M. L., Iguchi, M. Y. & Felch, L. J. (1992) Contingent take-home  
25 incentive: Effects on drug use of methadone maintenance patients. *Journal of*  
26 *Consulting and Clinical Psychology*, 60, 927-934.  
27
- 28 Strickler, D. C., Whitley, R., Becker, D. R., *et al.* (2009) First person accounts of  
29 long-term employment activity among people with dual diagnosis. *Psychiatric*  
30 *Rehabilitation Journal*, 32, 261-268.  
31
- 32 Swartz, M. S., Wagner, H. R., Swanson, J. W., *et al.* (2006) Substance use in  
33 persons with Schizophrenia: baseline prevalence and correlates from the  
34 NIMH CATIE study. *The Journal of Nervous and Mental Disease*, 194, 164-172.  
35
- 36 Talbott, J. A. & Teague, J. W. (1969) Marihuanna psychosis: Acute toxic  
37 psychosis associated with the use of cannabis derivatives. *Journal of the*  
38 *American Medical Association*, 210, 299-302.  
39
- 40 Teeson, M., Hall, W., Lynskey, M., *et al.* (2000) Alcohol and drug use disorders  
41 in Australia: implications of the National Survey of Mental Health and Well  
42 Being. *Australian and New Zealand Journal of Psychiatry*, 34, 206-213.

- 1  
2 Tiet, Q. Q. & Mausbach, B. (2007) Treatments for patients with dual diagnosis:  
3 a review. *Alcoholism: Clinical and Experimental Research*, 31, 513-536.  
4  
5 Timko, C., Chen, S., Sempel, J., *et al.* (2006) Dual diagnosis patients in  
6 community or hospital care: one-year outcomes and health care utilization  
7 and costs. *Journal of Mental Health*, 15, 163-177.  
8  
9 Todd, F. C., Sellman, D. & Robertson, P. (2002) Barriers to optimal care for  
10 patients with coexisting substance use and mental health disorders. *Australian  
11 and New Zealand Journal of Psychiatry*, 36, 792-799.  
12  
13 Tracy, K., Babuscio, T., Nich, C., *et al.* (2007) Contingency management to  
14 reduce substance use in individuals who are homeless with co-occurring  
15 psychiatric disorders. *The American Journal of Drug and Alcohol Abuse*, 33, 253-  
16 258.  
17  
18 Turkington, A., Mulholland, C. C., Rushe, T. M., *et al.* (2009) Impact of  
19 persistent substance misuse on 1-year outcome in first-episode psychosis.  
20 *British Journal of Psychiatry*, 195, 242-248.  
21  
22 Turning Point (2007) *Dual Diagnosis: Good Practice Handbook*. London: Turning  
23 Point. Available at: [www.turning-point.co.uk](http://www.turning-point.co.uk)  
24  
25 Turton, P., Demetriou, A., Boland, W., *et al.* (2009) One size fits all: or horses  
26 for courses? Recovery based care in specialist mental health services. *Social  
27 Psychiatry and Psychiatric Epidemiology*, DOI: 10.1007/s00127-009-0174-6.  
28  
29 Tyrer, P., Milošeska, K., Whittington, C., *et al.* (in press) Nidotherapy in the  
30 treatment of substance misuse, psychosis and personality disorder: secondary  
31 analysis of a controlled trial. *The Psychiatrist*.  
32  
33 Tyrer, P., Sensky, T. & Mitchard, S. (2003) Principles of nidotherapy in the  
34 treatment of persistent mental and personality disorders. *Psychotherapy and  
35 Psychosomatics*, 72, 350-356.  
36  
37 Van Nimwegen, L. J., de Haan, L., van Beveren, N. J., *et al.* (2008) Effect of  
38 olanzapine and risperidone on subjective well-being and craving for cannabis  
39 in patients with schizophrenia or related disorders: a double-blind  
40 randomized controlled trial. *Canadian Journal of Psychiatry*, 53, 400-405.  
41

- 1 Van Os, J., Bak, M., Hanssen, M., *et al.* (2002) Cannabis use and psychosis: a  
2 longitudinal population-based study. *American Journal of Epidemiology*, 156,  
3 319-327.  
4
- 5 Vardy, M. M., & Kay, S. R. (1983) LSD psychosis or LSD-induced  
6 schizophrenia? A multimethod inquiry. *Archives of General Psychiatry*, 40, 877-  
7 83.  
8
- 9 Verdoux, H., Tournier, M. & Cougnard, A. (2005) Impact of substance use on  
10 the onset and course of early psychosis. *Schizophrenia Research*, 79, 69-75.  
11
- 12 Vogel, H. S., Knight, E., Laudet, A. B., *et al.* (1998) Double trouble in recovery:  
13 self-help for people with dual diagnoses. *Psychiatric Rehabilitation Journal*, 21,  
14 356-364.  
15
- 16 Vornik, L. A. & Brown, E. S. (2006) Management of comorbid bipolar disorder  
17 and substance abuse. *Journal of Clinical Psychiatry*, 67, 24-30.
- 18 Wade, D., Harrigan, S., Edwards, J., *et al.* (2006) Substance misuse in first-  
19 episode psychosis: 15-month prospective follow-up study. *British Journal of*  
20 *Psychiatry*, 186, 229-234.  
21
- 22 Wagstaff, C. (2007) Towards understanding the self-perception of people with  
23 a psychotic illness who use illicit substances and have a history of  
24 disengagement from mental health services: qualitative research. *The*  
25 *International Journal of Psychiatric Nursing Research*, 12, 1503-1520.  
26
- 27 Waldron, H. B., Kern-Jones, S., Turner C. W., *et al.* (2007) Engaging resistant  
28 adolescents in drug abuse treatment *Journal of Substance Abuse Treatment*, 32,  
29 133-142.  
30
- 31 Warfa, N., Bhui, K., Phillips, K., *et al.* (2006) Comparison of life events,  
32 substance misuse, service use and mental illness among African-Caribbean,  
33 black Africa and white British men in east London: a qualitative Study.  
34 *Diversity in Health and Social Care*, 3, 111-121.  
35
- 36 Weaver, T., Maden, P., Charles, V., *et al.* (2003) Comorbidity of substance  
37 misuse and mental illness in community mental health and substance misuse  
38 services. *British Journal of Psychiatry*, 183, 304-313.  
39

- 1 Weiss, R. D., Griffin, M. L., Greenfield, S.F. *et al.* (2000) Group therapy for  
2 patients with bipolar disorder and substance dependence: results of a pilot  
3 study. *Journal of Clinical Psychiatry*, 61, 361-367.  
4
- 5 Weiss, R. D., Griffin, M. L., Jaffee, W. B., *et al.* (2009) A “community friendly”  
6 version of integrated group therapy for patients with bipolar disorder and  
7 substance dependence: a randomized controlled trial. *Drug and Alcohol*  
8 *Dependence*, 104, 212-219.  
9
- 10 Weiss, R. D., Griffin, M. L., Kolodziej, M. E., *et al.* (2007) A randomized trial of  
11 integrated group therapy versus group drug counselling for patients with  
12 bipolar disorder and substance dependence. *American Journal of Psychiatry*,  
13 164, 100-107.  
14
- 15 Weissman, M. M., Markowitz, J. C. & Klerman, G. L. (2000) *Comprehensive*  
16 *Guide to Interpersonal Therapy*. New York: Basic Books.  
17
- 18 Wobrock, T., & Soyka, M. (2008) Pharmacotherapy of schizophrenia with  
19 comorbid substance use disorder: reviewing the evidence and clinical  
20 recommendations. *Progress in Neuro-Psychopharmacology and Biological*  
21 *Psychiatry*, 32, 1375-1385.  
22
- 23 Wolfson, P., Holloway, F. & Killaspy, H. (2009) *Enabling Recovery for*  
24 *People with Complex Mental Health Needs: A Template for Rehabilitation*  
25 *Services in England*. London: Executive Committee of the Faculty of  
26 Rehabilitation and Social Psychiatry, Royal College of Psychiatrists. Available  
27 at: [http://www.rcpsych.ac.uk/pdf/fr\\_rs\\_1\\_forwebsite.pdf](http://www.rcpsych.ac.uk/pdf/fr_rs_1_forwebsite.pdf)  
28
- 29 Wong, S. C. P. & Gordon, A. (2006) The validity and reliability of the Violence  
30 Risk Scale: a treatment friendly violence risk assessment tool. *Psychology,*  
31 *Public Policy and Law*, 12, 279-309.  
32
- 33 Wong, S. C. P., Gordon, A. & Gu, D. (2007) Assessment and treatment of  
34 violence-prone forensic clients: an integrated approach. *British Journal of*  
35 *Psychiatry*, 190, 24-31.  
36
- 37 World Health Organization (1992) *ICD-10 Classification of Mental and*  
38 *Behavioural Disorders*. Geneva: World Health Organization.  
39
- 40 Wyte, S., Scott, F. & Maden, T. (2004) Substance misuse in secure psychiatric  
41 hospitals. *Journal of Forensic Psychiatry and Psychology*, 15, 591-594.  
42

1 Zammit, S., Moore, T. H. M., Lingford-Hughes, A., *et al.* (2008) Effects of  
2 cannabis use on outcomes of psychotic disorders: systematic review. *British*  
3 *Journal of Psychiatry*, 193, 357-363.  
4  
5 Zammit, S., Spurlock, G., Williams, H., *et al.* (2007) Genotype effects of  
6 CHRNA7, CNR1 and COMT in schizophrenia: interactions with tobacco and  
7 cannabis use. *British Journal of Psychiatry*, 191, 402-407.  
8  
9 Ziedonis, D. & Brady, K. (1997). Dual diagnosis in primary care: detecting and  
10 treating both the addiction and mental illness. *Medical Clinics of North America*,  
11 81, 1017-1036.  
12  
13 Ziedonis, D. M., Smelson, D., Rosenthal, R. N., *et al.* (2005) Improving the care  
14 of individuals with schizophrenia and substance use disorders: consensus  
15 recommendations. *Journal of Psychiatric Practice*, 11, 315-406.  
16  
17  
18  
19  
20