

Appendix A: Scope

1 Guideline title

Delirium: diagnosis, prevention and management of delirium.

1.1 Short title

Delirium.

2 Background

a) The National Institute for Health and Clinical Excellence ('NICE' or 'the Institute') has commissioned the National Collaborating Centre for Nursing and Supportive Care to develop a clinical guideline on the diagnosis, prevention and management of delirium for use in the NHS in England and Wales. This follows referral of the topic by the Department of Health (see section 6 of this document). The guideline will provide recommendations for good practice that are based on the best available evidence of clinical and cost effectiveness.

b) The Institute's clinical guidelines support the implementation of National Service Frameworks (NSFs) in those aspects of care where a Framework has been published. The statements in each NSF reflect the evidence that was used at the time the Framework was prepared. The clinical guidelines and technology appraisals published by the Institute after an NSF has been issued have the effect of updating the Framework.

c) NICE clinical guidelines support the role of healthcare professionals in providing care in partnership with patients, taking account of their individual needs and preferences, and ensuring that patients (and their carers and families, if appropriate) can make informed decisions about their care and treatment.

1 **3 Clinical need for the guideline**

2 a) Delirium, sometimes called 'acute confusional state' is characterised by a disturbance of
3 consciousness and a change in cognition that develops over a short period of time.

4 b) Although the clinical presentation of delirium differs considerably from patient to patient,
5 there are several characteristic features that help make the diagnosis. The standard criteria for
6 delirium, are described in the 'Diagnostic and Statistical Manual of Mental Disorders' [DSM IV]
7 (1994):

- 8 • disturbance of consciousness (i.e., reduced clarity of awareness of the environment) with
9 reduced ability to focus, sustain, or shift attention.
- 10 • a change in cognition (such as memory deficit, disorientation, language disturbance) or
11 the development of a perceptual disturbance that is not better accounted for by a pre-
12 existing, established, or evolving dementia.
- 13 • the disturbance develops over a short period of time (usually hours to days) and tends
14 to fluctuate during the course of the day.
- 15 • there is evidence from the history, physical examination, and laboratory findings that:
16 (1) the disturbance is caused by the direct physiological consequences of a general
17 medical condition, (2) the symptoms in criteria (a) and (b) developed during substance
18 intoxication¹, or during or shortly after, a withdrawal syndrome, or (3) the delirium has
19 more than one aetiology”.

20
21 c) Features of delirium are recent onset of fluctuating awareness, impairment of memory and
22 attention, and disorganised thinking. Additional features may include hallucinations and
23 disturbance of sleep-wake cycle. There are three clinical subtypes of delirium: hyperactive
24 (characterised by hallucinations, delusions, agitation, and disorientation), hypoactive (sleepy
25 state, uninterested in activities of living, often unrecognised or labelled as dementia) or mixed
26 (patients can move between the two subtypes). Delirium may be present when a person is
27 admitted to hospital (prevalent delirium) or develop during an admission (incident delirium).

28 d) The prevalence of delirium in hospitalised medically ill patients ranges from 10 to 31%.
29 Most delirium occurs in the first 7 to 10 days of admission or within days of surgery. Up to 50%
30 of postoperative patients develop delirium, with patients at increased risk if they have had
31 cardiac surgery, hip surgery or transplantation. Delirium is also commonly reported to occur in
32 nursing homes, but is uncommon in community populations.

33 e) There is often difficulty in distinguishing whether a patient has delirium without dementia,
34 has dementia alone, or has delirium with pre-existing dementia. Delirium, which has acute
35 onset, is potentially preventable and treatable compared with dementia, which is ongoing. The
36 severity of delirium symptoms fluctuates over a 24-hour period; this does not occur in dementia
37 (with the exception of vascular dementia, Lewy body and Parkinson's disease dementia).
38 Duration of symptoms of delirium has been reported to range from less than 1 week to more
39 than 2 months.

¹ Refer to 4.1.2 c.

1 f) The causes of delirium can include a general medical condition and drugs (including
2 psychoactive, opioids or anticholinergic types) and surgery. Typically, delirium occurs in
3 patients who have one or more risk factors, and who then experience a precipitating factor.
4 Some groups are more at risk, for example, people with dementia, people with a severe acute
5 illness and older people (older than 65 years). Precipitating factors include acute illness (for
6 example, urinary infection, lower respiratory tract infection), unrelieved pain, sleep
7 deprivation and environmental factors. In a patient with several risk factors, a small precipitant
8 (such as a change in medication) can trigger delirium.

9 g) Diagnosis of delirium may be made by examining changes in cognitive function. In addition
10 to quick assessment methods, established instruments include the confusion assessment method,
11 used to detect delirium, and the mini-mental state examination, used to monitor the
12 development and resolution of delirium.

13 h) There is a significant burden associated with this condition. Consequences of delirium may
14 include increased length of stay in hospital or in ICU, poor functional and cognitive recovery
15 after hospital admission, earlier onset or progression of dementia, increased hospital acquired
16 complications (nosocomial infection, falls, pressure sores, and incontinence), new admission to
17 long-term care and death.

18 i) There is a need for guidance to improve methods of appropriate identification, diagnosis,
19 prevention and management of delirium. Failure to diagnose delirium, or misdiagnosis (mainly
20 as dementia), can lead to inappropriate treatment being given. Delirium is often preventable
21 and improvements in care practices and other treatments are needed. The improved
22 management of delirium has the potential to generate cost savings.

23 4 The guideline

24 a) The guideline development process is described in detail in two publications that are
25 available from the NICE website (see 'Further information'). 'The guideline development
26 process: an overview for stakeholders, the public and the NHS' describes how organisations
27 can become involved in the development of a guideline. 'The guidelines manual' provides
28 advice on the technical aspects of guideline development.

29 b) This document is the scope. It defines exactly what this guideline will (and will not) examine,
30 and what the guideline developers will consider. The scope is based on the referral from the
31 Department of Health (see section 6 of this document).

32 c) The areas that will be addressed by the guideline are described in the following sections.

33 4.1 Population

34 4.1.1 Groups that will be covered

35 a) Adult patients (18 years and older) in a hospital setting

36 b) Adults (18 years and older) in long-term residential care.

37 4.1.2 Groups that will not be covered

38 a) Children and young people (younger than 18 years).

1 b) People receiving end-of-life care.

2 c) People with intoxication and/or withdrawing from drugs or alcohol, and people with
3 delirium associated with these states.

4 **4.2 Healthcare setting**

5 a) The guideline will be relevant to NHS staff responsible for patients in hospital (including
6 critical care) and long term residential care settings (including primary care health care
7 professionals).

8 **4.3 Clinical management**

9 a) Assessment of risk factors and consideration of precipitants to identify people at high risk of
10 developing delirium.

11 b) Diagnosis of delirium in acute, critical and long term care. This will take into consideration
12 people presenting with learning disability, linguistic or communication problems.

13 c) Pharmacological and non pharmacological interventions to reduce the risk of delirium and its
14 consequences.

15 d) Pharmacological and non pharmacological interventions for people with delirium to reduce
16 the severity and duration and to prevent deterioration and major consequences of delirium
17 including dementia.

18 e) Guideline recommendations will normally fall within licensed indications; Use outside a
19 licensed indication may be recommended, including those licensed for other conditions, if
20 clearly supported by evidence. The guideline will assume that prescribers will use a drug's
21 summary of product characteristics to inform their decisions for individual patients.

22 f) The Guideline Development Group will take reasonable steps to identify ineffective
23 interventions and approaches to care. If robust and credible recommendations for re-
24 positioning the intervention for optimal use, or changing the approach to care to make more
25 efficient use of resources can be made, they will be clearly stated. If the resources released
26 are substantial, consideration will be given to listing such recommendations in the 'Key priorities
27 for implementation' section of the guideline.

28 **4.4 Status**

29 **4.4.1 Scope**

30 This is the final version of the scope. The consultation period is 16 April to 14 May 2008.

31 NICE has published the following related guidance which may be referred to in this guideline:

- 32
- 33 • Acutely ill patients in hospital: recognition of and response to acute illness in adults in hospital. NICE clinical guideline 50 (2007). Available from www.nice.org.uk/CG050.
 - 34 • Infection Control: prevention of healthcare-associated infection in primary and
35 community care NICE clinical guideline 2 (2003). Available from
36 www.nice.org.uk/CG2.

- 1 • Nutrition Support in adults: Nutrition support in adults: oral nutrition support, enteral
2 tube feeding and parenteral nutrition. NICE clinical guideline 32 (2006). Available
3 from www.nice.org.uk/CG032.
- 4 • Dementia: supporting people with dementia and their carers in health and social care.
5 NICE clinical guideline 42 (2006). Available from www.nice.org.uk/CG042.
- 6 • Drug Misuse: opioid detoxification. NICE clinical guideline 52 (2007). Available from
7 www.nice.org.uk/CG0452.
- 8 • Alcohol Use Disorders. Two guidelines are currently in preparation (exact titles to be
9 confirmed). One will cover acute alcohol withdrawal and the treatment of physiological
10 diseases associated with alcohol. The other will cover the management of planned
11 detoxification and the psychological treatment of dependency.
- 12 • Surgical Site Infection. NICE clinical guideline in development.
- 13 • Schizophrenia (update). NICE clinical guideline in development
- 14 • Alzheimer's disease - donepezil, galantamine, rivastigmine (review) and memantine for
15 the treatment of Alzheimer's disease. NICE technology appraisal 111 (2007).
16 Available from www.nice.org.uk/TA111.
- 17 • Schizophrenia - the clinical effectiveness and cost effectiveness of newer atypical
18 antipsychotic drugs for schizophrenia. NICE technology appraisal 43 (2002). Available
19 from www.nice.org.uk/TA43.

20 **5 Further information**

21 Information on the guideline development process is provided in:

- 22 • 'The guideline development process: an overview for stakeholders, the public and the
23 NHS'.
- 24 • 'The guidelines manual'.

25 These booklets are available as PDF files from the NICE website
26 (www.nice.org.uk/guidelinesmanual). Information on the progress of the guideline will also
27 be available from the website.

28 **6 Referral from the Department of Health**

29 "Remit: To prepare a clinical guideline on the diagnosis, prevention and management of
30 delirium"