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Abbreviations

CBT	cognitive behavioural therapy
CI	confidence interval
RR	relative risk
SOTP	sex offender treatment program

N.1 Interventions for promoting health and well being

N.1.1 Parent training for parent-child attachment for women with sub-threshold symptoms

			Quality as	ssessment			Nº of p	atients	Effect	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Parent training versus treatment as usual	promoting mental health and wellbeing in adults in contact with the criminal justice system	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Depression (0	CES-D) (Scale from	n 0 to 60; lower bette	r)			·						•
1	randomised trials	serious 1	not serious	not serious	serious ²	none	62	53	Mean 15.3 (SD 11.8)	MD 1.70 lower (5.65 lower to 2.25 higher)		CRITICAL
Number of pa	rticipants with sym	ptoms of depression	(CES-D=>16)	•	•		•					
1	randomised trials	serious 1	not serious	not serious	serious ^{2,3}	none	23/62 (37.1%)	25/53 (47.2%)	RR 0.79 (0.51 to 1.21)	99 fewer per 1,000 (from 99 more to 231 fewer)		CRITICAL
Mother-child a	attachment: Reflect	tive functioning (PDI)	(Scale from -1 to 9; h	igher better)		·						
1	randomised trials	serious 1	not serious	not serious	serious ²	none	57	52	Mean 3.15 (SD 1.33)	MD 0.39 higher (0.15 lower to 0.93 higher)		CRITICAL
Mother-child i	nteraction: Dyadic	attunement (behavio	ural observation) (sca	le from 11 to 55; high	er better)							
1	randomised trials	serious ¹	not serious	not serious	serious ²	none	51	37	-	MD 3.08 lower (6.39 lower to 0.23 higher)		IMPORTANT
Mother-child i	nteraction: Parent	positive engagement	(behavioural observa	tion; scale from 5 to 2	5; higher better)	·						·
1	randomised trials	serious 1	not serious	not serious	serious ²	none	51	37	-	MD 0.17 lower (1.44 lower to 1.10 higher)		IMPORTANT
Mother-child i	nteraction: Child in	volvement (behaviou	ral observation; scale	from 6 to 30; higher t	petter)	•						
1	randomised trials	serious ¹	not serious	not serious	serious ²	none	51	52	-	MD 0.37 lower (2.19 lower to 1.45 higher)		IMPORTANT
Maternal perc	eptions of child: W	armth (MORS)		•	•	,	•		L	I		

			Quality as	ssessment			Nº of p	atients	Effe	ct		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Parent training versus treatment as usual	promoting mental health and wellbeing in adults in contact with the criminal justice system	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
	randomised trials	serious ¹	not serious	not serious	serious ²	none	31	40	-	SMD 0.44 higher (0.04 lower to 0.91 higher)		IMPORTANT
laternal perc	ceptions of child: In	vasion (MORS)										
	randomised trials	serious ¹	not serious	not serious	serious ²	none	31	40	-	SMD 0.12 lower (0.58 lower to 0.35 higher)		IMPORTANT
Naternal perc	ceptions of child: In	tensity of problem be	ehaviour (ECBI)									
	randomised trials	serious ⁴	not serious	not serious	serious ²	none	78	25	-	SMD 0.29 lower (0.74 lower to 0.16 higher)		IMPORTANT
Aaternal perc	ceptions of child: Fi	requency of problem	behaviour (ECBI)	•				·				•
	randomised trials	serious ⁴	not serious	not serious	serious ²	none	78	25	-	SMD 0.04 higher (0.41 lower to 0.49 higher)		IMPORTANT
Naternal perc	ceptions of parentir	ng: Involvement (APC	2)				1	<u> </u>		1 1		
	randomised trials	serious 4	not serious	not serious	serious ²	none	77	25	-	SMD 0.08 lower (0.53 lower to 0.37 higher)		IMPORTANT
Naternal perc	eptions of parentir	ng: Positive parenting	j (APQ)			,		II		4 4		-1
l	randomised trials	serious ⁴	not serious	not serious	serious ²	none	78	25	-	SMD 0.66 lower (1.12 lower to 0.2 lower)		IMPORTANT
Aaternal perc	ceptions of parentir	ng: Poor monitoring/s	supervision (APQ)	•				·				
	randomised trials	serious ⁴	not serious	not serious	serious ²	none	77	25	-	SMD 0.33 higher (0.13 lower to 0.78 higher)		IMPORTANT

			Quality as	ssessment			Nº of p	atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Parent training versus treatment as usual	promoting mental health and wellbeing in adults in contact with the criminal justice system	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	serious ⁴	not serious	not serious	serious ²	none	78	25	-	MD 3.02 lower (4.72 to 1.33 lower)		IMPORTANT
Maternal perc	ceptions of parentin	ig: Corporal punishme	ent (APQ) (Scale from	n 3 to 15; lower better))							
1	randomised trials	serious ⁴	not serious	not serious	serious ²	none	78	25	-	MD 0.29 lower (1.21 lower to 0.63 higher)		IMPORTANT
Drop-out (all	cause)											
2	randomised trials	serious ^{1,4}	not serious	not serious	serious ^{2,3}	none	54/182 (29.7%)	31/126 (24.6%)	RR 1.12 (0.76 to 1.64)	30 more per 1,000 (from 59 fewer to 157 more)		IMPORTANT

CI: Confidence interval; MD: Mean difference; SMD: Standardised mean difference; RR: Risk ratio

Sleed (2013) - no blinding
 Small sample size (N<400), no sample size calculation reported
 95% CI includes both no effect and clinically significant harm or benefit

4. Menting (2014) - unclear randomisation method and no blinding

N.1.2 Yoga for promoting mental health and wellbeing

			Quality as	sessment			№ of p	patients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Yoga versus waitlist control	promoting mental health and wellbeing in adults in contact with the criminal justice system	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Positive affect	t (PANAS) (Scale f	from 10 to 50; higher	better)									
1	randomised trials	serious 1	not serious	not serious	serious ²	none	45	55	-	MD 5.94 higher (2.91 higher to 8.97 higher)		CRITICAL
Negative affect	ct (PANAS) (Scale	from 10 to 50; lower	better)	1	1			1	•			1

			Quality as	ssessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Yoga versus waitlist control	promoting mental health and wellbeing in adults in contact with the criminal justice system	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	serious 1	not serious	not serious	serious ²	none	45	55	-	MD 4.13 lower (6.80 lower to 1.46lower)		CRITICAL
Perceived str	ess (PSS) (Scale f	rom 0 to 40; lower be	etter)	•	•			·				
1	randomised trials	serious 1	not serious	not serious	serious ²	none	45	55	-	MD 4.67 lower (7.65 lower to 1.69 lower)		CRITICAL
Psychologica	l distress (BSI) (Sc	ale from 0 to 212; lov	wer better)									
1	randomised trials	serious 1	not serious	not serious	serious ²	none	45	55	-	MD 12.60 lower (22.82 lower to 2.38 lower)		CRITICAL
Drop-out (all	cause)	•		•	•	•	•	•				•
1	randomised trials	serious 1	not serious	not serious	serious 2.3	none	42/87 (48.3%)	25/80 (31.3%)	RR 1.54 (1.04 to 2.28)	169 more per 1,000 (from 13 more to 400 more)		IMPORTANT

CI: Confidence interval; MD: Mean difference; RR: Risk ratio

Bilderbeck (2013) - no blinding, attrition bias (significantly higher dropout with yoga)
 Study was an exploratory trial - without sample size calculation
 95% CI includes the possibility that the benefit is less than the minimum important difference

Meditation for promoting mental health and well-being N.1.3

			Quality as	sessment			№ of p	atients	Effect	:		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Meditation	treatment as usual	Relative (95% Cl)	Absolute (95% CI) ³	Quality	Importance
Desire to thro	ow things or hit peo	ple within past month	(study-specific meas	ure)								
1	randomised trials	serious 1	not serious	not serious	serious ²	none	17	16	-	SMD 1.01 lower (1.73 lower to 0.28 lower)		IMPORTANT

			Quality as	ssessment			№ of p	patients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Meditation	treatment as usual	Relative (95% CI)	Absolute (95% Cl) ³	Quality	Importance
Feelings of g	uilt within past mor	th (study-specific me	asure)									
1	randomised trials	serious 1	not serious	not serious	serious ²	none	17	16	-	SMD 0.42 lower (1.11 lower to 0.27 higher)	€ Low	IMPORTANT
Feelings of h	opelessness within	past month (study-sp	pecific measure)									
1	randomised trials	serious 1	not serious	not serious	serious ²	none	17	16	-	SMD 0.06 lower (0.74 lower to 0.63 higher)		IMPORTANT
Being bother	ed by nail biting wit	hin past month (study	/-specific measure)	ł	ł	<u>I</u>				1		
1	randomised trials	serious 1	not serious	not serious	serious ²	none	17	16	-	SMD 1.18 lower (1.91 lower to 0.44 lower)		IMPORTANT
Being bother	ed by sleeping diffi	culties within past mo	nth (study-specific me	easure)								
1	randomised trials	serious 1	not serious	not serious	serious ²	none	17	16	-	SMD 0.28 lower (0.96 lower to 0.41 higher)		IMPORTANT

CI: Confidence interval; SMD: Standardised mean difference

1. Sumter (2009) - no blinding, unclear allocation concealment

Small sample size (N<400), no sample size calculation reported
 It was not possible to calculate MD, so SMD is reported.

N.1.4 Physical exercise programmes versus exercise as usual for promoting mental health and well-being

			Quality as	sessment			№ of p	atients	Effect			
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Physical exercise programme	exercise as usual	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance
Change in Sy	mptom Checklist-9	0-Revised (SCL-90-F	R) Global Severity Inde	ex (GSI) - CRT or HIS	T exercise programm	ne versus exercise as usual (follow	up: 39 weeks) (Scale fro	om 0 to 4; lower better)				
1	randomised trials	very serious ¹	not serious	not serious	not serious	none	44	20	-	MD 0.17 lower (0.21 lower to 0.12 lower)		CRITICAL
Change in Sy	mptom Checklist-9	0-Revised (SCL-90-F	R) Positive Symptom 1	Total (PST) - CRT or H	HIST exercise program	nme versus exercise as usual (fol	ow up: 39 weeks) (Scale	from 0 to 90; lower bette	er)			

			Quality as	sessment			Nº of p	atients	Effect	:		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Physical exercise programme	exercise as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	very serious ¹	not serious	not serious	not serious	none	44	20	-	MD 7.08 lower (9.15 lower to 5 lower)		CRITICAL
Change in Sy	mptom Checklist-9	0-Revised (SCL-90-F	R) Positive Symptom [Distress Index (PSDI)	- CRT or HIST exerci	se programme versus exercise as	usual (follow up: 39 wee	ks) (Scale from 0 to 4; lo	wer better)			
1	randomised trials	very serious 1	not serious	not serious	not serious	none	44	20	-	MD 0.33 lower (0.41 lower to 0.25 lower)		CRITICAL

CI: Confidence interval; MD: Mean difference

1. Battaglia 2015 - unclear allocation concealment, no blinding, per-protocol analysis

N.2 Interventions for substance misuse

N.2.1 Psychological interventions

N.2.1.1 CBT versus active intervention

			Quality asse	essment			No of patient			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	CBT versus active intervention	Control	Relative (95% Cl)	Absolute		
Days usir	g cannabis (c	luring treatmo	ent) - Self-report (Better indicated	by lower values)						
1			no serious inconsistency	no serious indirectness	serious ¹	none	68	27	-	MD 10.15 days higher (6.63 lower to 26.93 higher)	⊕⊕OO LOW	CRITICAL
Days usir	g cannabis (c	luring treatmo	ent) - Urine test (E	Better indicated b	y lower values)	•	<u>.</u>					
1			no serious inconsistency	no serious indirectness	serious ²	none	68	27	-	MD 17.13 days higher (0.92 to 33.34 higher)	⊕⊕OO LOW	CRITICAL
Days with	positive urin	e test (during	treatment) (Bette	r indicated by lo	wer values)							
1	randomised trials		no serious inconsistency	no serious indirectness	very serious ²	none	38	37	-	MD 0.3 days higher (2.23 lower to 2.15 higher)	⊕OOO VERY LOW	CRITICAL
Days with	positive brea	thalyzer test	(during treatment) (Better indicate	d by lower valu	es)	•				•	•

1		2						1				
		serious ³	no serious	no serious	very serious ¹	none	38	37	-	MD 0.04 lower	$\oplus OOO$	CRITICA
	trials		inconsistency	indirectness						(0.46 lower to 0.44	VERY	
										higher)	LOW	
Days abs	stinent (during	treatment) -	Alcohol (Better i	ndicated by lowe	r values)			•				
	randomised	Very	no serious	no serious	no serious	none	36	35	-	MD 10.40 higher	⊕000	CRITICA
	trials		inconsistency	indirectness	imprecision					(1.53 to 19.27 higher)	VERY	
										ζ σ ,	LOW	
Days abs	stinent (during	treatment) -	Drugs (Better in	licated by lower	values)	- 4		1			1	
	randomised	Verv	no serious	no serious	no serious	none	36	35	-	MD 0.70 higher	⊕000	CRITICA
	trials	serious ³	inconsistency	indirectness	imprecision					(0.41 lower to 6.12	VERY	
			,							higher)	LOW	
Addictio	n Severity Inde	ex (ASI-6): ald	cohol composite	score (follow-up	26-38 weeks; B	etter indicated by lo	ower values)	4	•		Į	1
	randomised	very serious ⁴	no serious	no serious	serious ¹	none	23	21	-	MD 0.10 lower (0.22	⊕000	CRITICA
	trials		inconsistency	indirectness						lower to 0.02 higher)	VERY	
			-							Ç,	LOW	
A .I .I					OO Dett	ar indicated by low						
Addictio	n Severity Inde	ex (ASI-6): dru	ug composite sc	ore (follow-up 26	-38 weeks; Bett	er indicated by low	er values)					
Addictio			• •	ore (follow-up 26 no serious	no serious	none	er values) 23	21	-	MD 0.02 lower (0.09	⊕000	CRITICA
		very serious ⁴	• •	<u> </u>	,			21	-	MD 0.02 lower (0.09 lower to 0.05 higher)	⊕OOO VERY	CRITICA
Addictio	randomised	very serious ⁴	no serious	no serious	no serious			21	-	```		CRITICAI
1	randomised trials	very serious ⁴	no serious inconsistency	no serious	no serious imprecision			21	-	```	VERY	CRITICA
1	randomised trials bstinent (follow	very serious ⁴ w-up 26-38 w	no serious inconsistency eeks; Better indi	no serious indirectness	no serious imprecision			21	-	```	VERY LOW	
	randomised trials bstinent (follow	very serious ⁴ w-up 26-38 we very serious ⁴	no serious inconsistency eeks; Better indi	no serious indirectness cated by higher	no serious imprecision ralues)	none	23		-	lower to 0.05 higher) MD 1.30 lower (4.4	VERY	
1	randomised trials bstinent (follow randomised	very serious ⁴ w-up 26-38 we very serious ⁴	no serious inconsistency eeks; Better indi no serious	no serious indirectness cated by higher no serious	no serious imprecision ralues)	none	23		-	lower to 0.05 higher)	VERY LOW ⊕000	CRITICAI
Weeks a	randomised trials bstinent (follow randomised	very serious⁴ w-up 26-38 wo very serious⁴	no serious inconsistency eeks; Better indi no serious inconsistency	no serious indirectness cated by higher no serious	no serious imprecision ralues)	none	23		-	lower to 0.05 higher) MD 1.30 lower (4.4	VERY LOW ⊕OOO VERY	
Weeks a	randomised trials bstinent (follow randomised trials ceration (follow	very serious⁴ w-up 26-38 wo very serious⁴	no serious inconsistency eeks; Better indi no serious inconsistency eks)	no serious indirectness cated by higher no serious	no serious imprecision values) serious ¹	none	23		- - RR 0.51 (0.2	lower to 0.05 higher) MD 1.30 lower (4.4 lower to 1.8 higher)	VERY LOW ⊕OOO VERY	CRITICA
Weeks a	randomised trials bstinent (follow randomised trials ceration (follow	very serious ⁴ w-up 26-38 we very serious ⁴ /-up 26-38 we very serious ⁴	no serious inconsistency eeks; Better indi no serious inconsistency eks)	no serious indirectness cated by higher no serious indirectness	no serious imprecision ralues)	none	23	21	· · · ·	lower to 0.05 higher) MD 1.30 lower (4.4 lower to 1.8 higher)	VERY LOW ⊕000 VERY LOW ⊕000	

¹ 95% CI includes both no effect and the minimal important difference
 ² 95% CI includes the minimal important difference
 ³ high risk of performance bias. Unclear risk for allocation concealment, detection, attrition, reporting and other bias
 ⁴ high risk of concealment bias, unclear risk on all other dimensions

CBT versus control N.2.1.2

			Quality assess	sment			No of patier	nts		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	CBT versus control/TAU	Control	Relative (95% CI)			
Addiction	n Severity Inde	x (ASI-6): alco	hol composite sco	ore (Better indica	ted by lower	values)						
2	randomised trials				very serious ²	none	39	32	-	SMD 0.37 lower (0.85 lower to 0.1 higher)	⊕OOO VERY	CRITICAL

											LOW	
ddictior	n Severity Inde	ex (ASI-6): dru	ug composite sco	ore (Better indicat	ted by lower v	values)						•
	randomised trials	serious ³	no serious inconsistency	no serious indirectness	very serious ⁴	none	39	32	-	SMD 0.28 lower (0.75 lower to 0.2 higher)	⊕OOO VERY LOW	CRITICA
bstinen	t in previous 3	months (6 m	nonth follow-up)									
	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	very serious⁵	none	4/16 (25%)	2/11 (18.2%)	RR 1.38 (0.3 to 6.25)	69 more per 1000 (from 127 fewer to 955 more)	⊕⊕OO LOW	CRITICA
Veeks at	ostinent (3 mo) (Better indicate	d by lower values	s)							
	randomised trials	serious ⁶	no serious inconsistency	no serious indirectness	very serious ⁷	none	23	21	-	SMD 0.24 lower (0.84 lower to 0.35 higher)	⊕OOO VERY LOW	CRITIC
Reincarco	eration		•		•		•					•
	randomised trials	serious ³	no serious inconsistency	no serious indirectness	very serious ⁸	none	5/23 (21.7%)	9/21 (42.9%) 42.9%		210 fewer per 1000 (from 343 fewer to 116 more) 210 fewer per 1000 (from 343 fewer to 116 more)	⊕OOO VERY LOW	CRITICA
² N<100 8 ³ No expla ⁴ N<100 8 ⁵ very sma ⁵ high risk ⁷ N<100 8	CI -0.85-0.1 Anation was pro CI -0.75-0.2 All number of ev	vided vents & CI 0.3 and unclear r	isk for all other cat		ther bias' and t	_ unclear risk for all	other categories			343 tewer to 116 more)	<u> </u>	<u> </u>

N.2.1.3 ACT versus CBT

			Quality ass	essment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducation versus active intervention	Control	Relative (95% CI)	Absolute		
Addiction	Severity Ind	ex (ASI-6):	alcohol composit	e score (follow-ι	ip mean 42 wee	eks; Scale from 0	to 9; lower better)			•		
	randomised trials				no serious imprecision	none	14	16	-	MD 0.04 lower (0.07 to 0.01 lower)	⊕⊕OO LOW	CRITICAL
Addicition	n Severity Inc	dex (ASI-6):	drug composite	score (Scale fror	n 0 to 9; lower	better)	·					
1	randomised	serious	no serious	no serious	very serious ³	none	14	16	-	MD 0.01 lower	⊕000	CRITICAL

	trials	risk of bias ¹	inconsistency	indirectness						(0.05 lower to 0.03 higher)	VERY LOW	
Abstinen	t from drugs	in previous	3 months									
1			no serious inconsistency	no serious indirectness	very serious ³	none	6/14 (42.9%)	4/16 (25%)	RR 1.71 (0.6 to 4.86)	178 more per 1000 (from 100 fewer to 965 more)	⊕OOO VERY LOW	CRITICAL

1 High risk of performance and detection bias, all other domains low risk

2 optimal information size criterion not met

3 confidence interval includes both clinically significant benefit and harm

N.2.1.4 ACT versus waitlist

	sus wattist							1				
			Quality asses	ssment			No of p	oatients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	ACT	Waitlist	Relative (95% CI)	Absolute		
Addiction	Severity Index	(ASI-6): ald	cohol composite sco	ore (follow-up mea	n 42 weeks;	Better indicated by	lower v	alues)				
2	n Severity Index (ASI-6): alcohol composite randomised very very serious ² trials serious ¹ very composite sc			no serious indirectness	serious ³	none	32	24	-	SMD 0.60 lower (1.72 lower to 0.53 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Index	(ASI-6): dr	ug composite score	(follow-up mean 4	12 weeks; Be	tter indicated by lo	wer valu	ues)				
2	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	30	22	-	SMD 0.44 lower (1.19 lower to 0.3 higher)	⊕OOO VERY LOW	CRITICAL
Abstinent	from drugs in	previous 3	months (follow-up r	nean 42 weeks)								
1	randomised trials	serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	6/14 (42.9%)	2/11 (18.2%)	RR 2.36 (0.59 to 9.48)	247 more per 1000 (from 75 fewer to 1000 more)	⊕⊕OO LOW	CRITICAL

1 high risk of performance bias, unclear or mixed risk on three other facets

 $2 l^2 = 75\%$, random effects model used and outcome downgraded for inconsistency

3 confidence interval includes both clinically significant benefit and harm

4 high risk of performance bias, unclear or mixed risk on two other facets

N.2.1.5 Mindfulness-based relapse prevention versus active intervention

			Quality asse	ssment			No of patie	ents		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Mindfulness-based relapse prevention	Active intervention	Relative (95%	Absolute		

					1							
									CI)			
Drug-use	days (Better i	ndicated	by lower values)									
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	28	26	-	MD 0.46 lower (1.16 lower to 0.24 higher)	⊕OOO VERY LOW	CRITICAL
			P) follow-up (Better		· · · ·							
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	28	26	-	MD 7.30 lower (15.81 lower to 1.21 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Inde	ex: family-	social composite	score (Better ind	icated by low	ver values)						
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	28	26	-	MD 0.01 lower (0.09 lower to 0.07 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Inde	x: legal c	omposite score (B	etter indicated b	y lower value	es)				•		•
		very serious ¹	no serious inconsistency	no serious indirectness	very serious²	none	28	26	-	MD 0.31 lower (0.45 to 0.17 lower)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Inde	x: medica	al composite score	(Better indicate	d by lower va	alues)						
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	28	26	-	MD 0.20 lower (0.37 to 0.03 lower)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity inde	x: psychi	atric compose sco	re (Better indica	ted by lower	values)						•
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	28	26	-	MD 0.11 lower (0.22 lower to 0 higher)	⊕OOO VERY LOW	CRITICAL

¹ high risk of bias from blinding and other factors, unclear risk of bias on 5 other domains ² optimal information size criterion not met

N.2.1.6 Contingency management versus active intervention

			Quality asse	essment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Contingency management versus active intervention	Control	Relative (95% Cl)	Absolute		
Days usir	ng cannabis (o	during tre	atment) - Self-repo	ort (Better indica	ted by lower	values)						
2	randomised trials			no serious indirectness	serious ²	none	158	105	-	SMD 0.01 higher (0.24 lower to 0.26 higher)	⊕⊕OO LOW	CRITICAL
Days usir	ng cannabis (d	during tre	atment) - Urine tes	t (Better indicate	ed by lower	values)						
2	randomised trials	serious ³			very serious ⁴	none	67	69	-	SMD 0.23 lower (0.57 lower to 0.11 higher)	⊕OOO VERY LOW	CRITICAL

	randomised	serious⁵	no serious	no serious	very	none	37	28	_	SMD 0.18 higher (0.32	⊕000	CRITICA
	trials	3011003	inconsistency	indirectness	serious ⁶	none	51	20		lower to 0.67 higher)	VERY	
	ulais		inconsistency	indirectriess	3011003					lower to 0.07 higher)	LOW	
ddictio	on Severity Ind	ex (ASI): r	narijuana compo	site score - Follo	w-up (Better	indicated by lower	· values)				LOW	
	randomised	serious ⁷	no serious	no serious	very	none	37	28	_	SMD 0.11 higher (0.38	⊕000	CRITICA
	trials	3011003	inconsistency	indirectness	serious ⁶	none	51	20		lower to 0.6 higher)	VERY	
	thais		inconsistency		3611043					lower to 0.0 mgrier)	LOW	
ays ca	nnabis use per	r month - I	Post-treatment (E	etter indicated b	y lower value	es)						1
2	randomised	serious ⁷	no serious	no serious	very	none	37	28	-	SMD 0.5 higher (0 to 1	⊕000	CRITICA
	trials		inconsistency	indirectness	serious ⁶					higher)	VERY	
										5 - 7	LOW	
)ays ca	nnabis use per	month - I	Follow-up (Better	indicated by low	ver values)	-						
	randomised	serious ⁷	no serious	no serious	very	none	58	28	-	SMD 0.22 higher (0.24	⊕000	CRITICA
	trials		inconsistency	indirectness	serious ⁶					lower to 0.67 higher)	VERY	
											LOW	
articipa	ants still in trea	atment at	follow-up (follow-	up mean 52 wee	ks)							
	randomised	very	no serious	no serious	serious ⁶	none	18/83	22/82	RR 0.81	51 fewer per 1000	⊕000	CRITICA
	trials	serious ⁷	inconsistency	indirectness			(21.7%)	(26.8%)	(0.47 to 1.39)	(from 142 fewer to 105	VERY	
							· · · · ·	`,	``````````````````````````````````````	more)	LOW	
lo. of d	ays in treatme	nt (follow-	up mean 52 week	s; Better indicat	ed by higher	values)						
	randomised	very	no serious	no serious	serious	none	83	82	-	MD 3.00 lower (21.01	⊕000	CRITICA
	trials	serious ⁷	inconsistency	indirectness						lower to 15.01 higher)	VERY	
			,							ξ,	LOW	

¹ One study high risk for performance and attrition bias, unclear for selection and reporting bias. Other study high risk for performance and unclear for allocation concealment and reporting bias

² Optimal information size criterion not met (N<400)
 ³ high risk of bias, unclear for selection and reporting bias
 ⁴ Optimal information size criterion not met (N<200) & CI includes both clinically significant harm and no effect

⁵ performance bias is high risk, all other categories (except other) are unclear risk ⁶ CI includes both clinically significant or harm and no effect

⁷ high risk of blinding and outcome reporting bias, unclear risk of performance and concealment bias

Contingency management versus treatment as usual N.2.1.7

			Quality assessm	ient			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Contingency management	TAU	Relative (95% Cl)	Absolute		
Arrests for	public drunken	ness (Better indi	icated by lower value	es)								

1 randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	very serious ¹	none	10	10	-	MD 1.70 fewer arrests	⊕⊕OO LOW	CRITICAL
									(5.65 fewer to 2.25 more)	-	

¹ Optimal information size criterion not met (N<200); 95% CI of effect includes both clinically significant benefit and no effect

N.2.1.8 Motivational enhancement therapy versus active intervention

			Quality asso	essment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Motivational enhancement therapy versus active intervention	Control	Relative (95% Cl)	Absolute		
Percenta	ge of days ab	stinent fro	om alcohol (self-re	port) - 3 month f	ollow-up (Be	tter indicated by lo	ower values)	•				•
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	119	119	-	MD 9.5 % more (2.51 to 16.49 % more)	⊕000 VERY LOW	CRITICAL
Percenta	ge of days ab	stinent fro	om alcohol (self-re	port) - 6 month f	ollow-up (Be	tter indicated by lo	ower values)					
1	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	107	107	-	MD 4.8 % more (2.50 % fewer to 12.10 % more)	⊕000 VERY LOW	CRITICAL
Percenta	ge of days ab	stinent fro	om alcohol (self-re	port) - 12 month	follow-up (Be	etter indicated by	lower values)					
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	95	95	-	MD 0.8 % more (8.37 % fewer to 6.77 % more)	⊕000 VERY LOW	CRITICAL
Percenta	ge of days ab	stinent fro	m alcohol and dru	igs - 3 month fol	low-up (Bette	er indicated by low	ver values)	•				
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	119	119	-	MD 9.7 % more (0.7 % more to 18.63 % more)	⊕OOO VERY LOW	CRITICAL
Percenta	ge of days ab	stinent fro	m alcohol and dru	ugs - 6 month fol	low-up (Bette	er indicated by low	ver values)					
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	107	107	-	MD 5.2 % more (4.05 % fewer to 14.45 % more)	⊕OOO VERY LOW	CRITICAL
Percenta	ge of days ab	stinent fro	om alcohol and dru	ugs - 12 month fo	ollow-up (Bet	ter indicated by lo	wer values)					
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	95	95	-	MD 9.7 % more (0.7 % more to 18.63 % more)	⊕000 VERY LOW	CRITICAL
Drinks pe	er drinking da	ys - 3 mor	nth follow-up (Bett	er indicated by l	ower values)							
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	119	119	-	MD 1.7 drinks fewer (3.75 fewer to 0.35 more)	⊕000 VERY LOW	CRITICAL

Drinks pe	r drinking day	ys - 6 mor	th follow-up (Bett	er indicated by lo	ower values)							
1		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	107	107	-	MD 0.70 drinks more (0.93 fewer to 2.33 more)	⊕OOO VERY LOW	CRITICAL
Drinks pe	r drinking day	ys - 12 mo	onth follow-up (Bet	tter indicated by	lower values	5)						
1		very serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	97	95	-	MD 0.30 drinks fewer (1.90 fewer to 1.33 more)	⊕OOO VERY LOW	CRITICAL
Percentag	ge of days wit	h cannab	is use (during trea	tment) (Better in	dicated by lo	ower values)						
1		very serious ⁴	no serious inconsistency	no serious indirectness	very serious ³	none	69	67	-	SMD 0.1 lower (0.44 lower to 0.24 higher)	⊕OOO VERY LOW	CRITICAL
Percentag	ge of urine tes	sts positiv	e for cannabis us	e (during treatme	ent) (Better ir	dicated by lower	values)					
1		very serious ⁴	no serious inconsistency	no serious indirectness	very serious ³	none	69	67	-	SMD 0.91 lower (1.27 to 0.56 lower)	⊕OOO VERY LOW	CRITICAL
Self-repo	rted motivatio	on to take	steps to change s	ubstance abuse	scores (Bette	er indicated by hig	her values)					
1		very serious⁵	no serious inconsistency	no serious indirectness	serious ²	none	18	9	-	MD 4.10 higher (5.77 lower to 13.97 higher)	⊕OOO VERY LOW	CRITICAL

¹ High performance bias + unclear for 4 other bias types.
 ² Optimal information size criterion not met (N < 400)
 ³ Attrition bias (more than 50% of sample)
 ⁴ High performance bias + high attrition bias + unclear on 3 other types of bias.
 ⁵ High risk of performance, detection and other bias, unclear selection and attrition bias

N.2.1.9 Motivational interviewing or feedback versus active intervention

			Quality ass	essment			No of patients			Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Motivational interviewing/Motivational feedback versus control/TAU	Control	Relative (95% CI)	Absolute			
Self-repo	elf-reported drug use - 1 month follow-up												
1		- 1	no serious inconsistency	very serious ²	very serious ³	none	24/39 (61.5%)	19/40 (47.5%)	RR 1.3 (0.86 to 1.95)	142 more per 1000 (from 66 fewer to 451 more)	⊕OOO VERY LOW	CRITICAL	
Self-repo	rted days wit	h drug u	se in past 30 days	s (10 month foll	ow-up) (Bett	er indicated by lo	wer values)			•			
1	randomised	very	no serious	no serious	very	none	90	24	-	SMD 0.04 higher	⊕000	CRITICAL	

	trials	serious ⁴	inconsistency	indirectness	serious⁵					(0.41 lower to 0.49 higher)	VERY LOW	
Jrine tes	st positive for	drug use	e (during study	period)	-	-1	-			,		ļ
1	randomised trials	very serious ⁶	no serious inconsistency	very serious ²	very serious ³	none	15/39 (38.5%)	14/40 (35%)	RR 1.1 (0.62 to 1.96)	35 more per 1000 (from 133 fewer to 336 more)	⊕OOO VERY LOW	CRITICA
Self-rep	orted alcohol	use - 1 m	onth follow-up									
1	randomised trials	very serious ¹	no serious inconsistency	very serious ²	very serious ³	none	24/39 (61.5%)	19/40 (47.5%)	RR 1.3 (0.86 to 1.95)	142 more per 1000 (from 66 fewer to 451 more)	⊕OOO VERY LOW	CRITICAL
Days wit	th illegal activ	ity in pas	t 30 days (10 m	onth follow-up)	(Better indic	ated by lower val	ues)					1
1		very serious ⁴	no serious inconsistency	no serious indirectness	very serious⁵	none	80	23	-	SMD 0.07 higher (0.4 lower to 0.53 higher)	⊕000 VERY LOW	CRITICAL
Drop-ou	t from subsec	uent trea	tment - binge d	lrinking group (f	ollow-up me	an 26 weeks)						
1	randomised trials	serious ⁷	no serious inconsistency	no serious indirectness	serious ⁸	none	2/11 (18.2%)	8/12 (66.7%)	RR 0.27 (0.07 to 1.02)	487 fewer per 1000 (from 620 fewer to 13 more)	⊕⊕OO LOW	CRITICAL
Drop-ou	t from subsec	uent trea	tment - no bing	e drinking grou	p (follow-up	mean 26 weeks)						I
1	1	-	no serious inconsistency	no serious indirectness	serious ⁸	none	5/20 (25%)	4/15 (26.7%)	RR 0.94 (0.3 to 2.91)	16 fewer per 1000 (from 187 fewer to 509 more)	⊕⊕OO LOW	CRITICAL
Number	of subsequer	nt treatme	ent sessions att	ended - binge d	rinking grou	p (follow-up mear	26 weeks; Better indicated by high	er values)		· · · ·		
1	randomised trials	serious ⁷	no serious inconsistency	no serious indirectness	serious ⁸	none	10	9	-	MD 11.16 higher (3.86 to 18.46 higher)	⊕⊕OO LOW	CRITICAL
Number	of subsequer	nt treatme	ent sessions att	ended - no bing	e drinking gr	roup (follow-up m	ean 26 weeks; Better indicated by hi	gher value	es)			
1	randomised trials		inconsistency	no serious indirectness	serious ⁸	none	20	15	-	MD 1.65 lower (8.28 lower to 4.98 higher)	⊕⊕OO LOW	CRITICAL
Specialit	ty addiction c	linic atter	ndance							· · · · · · · · · · · · · · · · · · ·		
1	randomised trials	very serious ⁹	no serious inconsistency	no serious indirectness	serious ⁸	none	8/17 (47.1%)	4/13 (30.8%)	RR 1.53 (0.59 to 3.99)	163 more per 1000 (from 126 fewer to 920 more)	⊕OOO VERY LOW	CRITICAL

1 high performance bias + high other bias + 3 unclear;

2 very serious limitations (outcome)

Mental health of adults in contact with the criminal justice system Appendix N: Clinical evidence - GRADE evidence profiles

3 Optimal information size criterion not met (n = 79)

4 high performance and detection bias.

5 Optimal information size criterion not met (n = 114)

6 high performance bias + high other bias + 3 unclear

7 High risk of performance bias, unclear selection and detection bias

8 Optimal information size criterion not met

9 High risk of blinding, performance and detection bias, unclear selection and concealment bias

N.2.1.10 Group counselling versus treatment as usual

			Quality asso	essment			No of patie	ents		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Group counselling	TAU	Relative (95% CI)	Absolute		
Rearrest (*	12 month follo	w-up)	•	•	•	•		•	•		•	
1		- ,	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	46/85 (54.1%)	24/43 (55.8%)	RR 0.97 (0.7 to 1.35)	17 fewer per 1000 (from 167 fewer to 195 more)	⊕OOO VERY LOW	CRITICAL
Number of	f reconviction	s (12 mont	h follow-up) (Bette	er indicated by lov	wer values)	1		•				
1		- ,	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	99	50	-	MD 0.10 fewer reconvictions (0.68 fewer to 0.48 more)	⊕OOO VERY LOW	CRITICAL
Reincarce	ration (12 mor	th follow-	up)			•			•			•
1		· ·	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	24/85 (28.2%)	14/43 (32.6%)	RR 0.87 (0.5 to 1.5)	42 fewer per 1000 (from 163 fewer to 163 more)	⊕OOO VERY LOW	CRITICAL
Days inca	rcerated (12 m	onth follo	w-up) (Better indic	ated by lower val	ues)	I		1	1			1
1		- ,	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	99	50	-	MD 0.30 days more (28.9 fewer to 29.5 more)	⊕OOO VERY LOW	CRITICAL
Self-repor	ted drug use (12 month	follow-up) - Mariju	ana	+	ł			•		۰ 	
1		- ,	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	31/85 (36.5%)	24/43 (55.8%)		195 fewer per 1000 (from 22 fewer to 313 fewer)	⊕OOO VERY LOW	CRITICAL

Self-repor	ted drug use (12 month	follow-up) - LSD									
1	randomised trials	1	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	14/85 (16.5%)	9/43 (20.9%)	RR 0.79 (0.37 to 1.67)	44 fewer per 1000 (from 132 fewer to 140 more)	⊕OOO VERY LOW	CRITICAL
Self-repor	ted drug use (12 month	follow-up) - Speed		1				<u> </u>			l
1	randomised trials	- ,	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	14/85 (16.5%)	4/43 (9.3%)	RR 1.77 (0.62 to 5.05)	72 more per 1000 (from 35 fewer to 377 more)	⊕OOO VERY LOW	CRITICAL
Self-repor	ted drug use (12 month	follow-up) - Heroin	l								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ^{2,3}	none	7/85 (8.2%)	3/43 (7%)	RR 1.18 (0.32 to 4.34)	13 more per 1000 (from 47 fewer to 233 more)	⊕OOO VERY LOW	CRITICAL

1 high risk of performance and detection bias. Unclear risk of remaining categories (other than 'other' bias)

2 Imprecision: optimal information size criterion not met

3 Confidence interval of effect includes both clinically significant benefit and harm

N.2.1.11 Self-help versus control for substance misuse

			Quality ass	essment			No of patie	ents		Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Self-help versus control	Control	Relative (95% CI)	Absolute			
Subseque	Subsequent bookings (12 month follow-up)												
	randomised trials	serious risk of bias ¹			no serious imprecision	none	49/98 (50%)	56/85 (65.9%)		158 fewer per 1000 (from 20 fewer to 270 fewer)	⊕⊕OO LOW	CRITICAL	

¹ Sample size not reported. 183 participants were randomised but is unclear how many were assessed for eligibility

N.2.2 Pharmacological interventions

N.2.2.1 Naloxone versus placebo

			Quality asse	essment			No of patier			Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Naloxone versus placebo	Control	Relative (95% CI)	Absolute			
Discontinu	continued medication												
	randomised trials	very serious ¹	no serious inconsistency		very serious ²	none	16/55 (29.1%)	8/42 (19%)	RR 1.53 (0.72 to 3.23)	101 more per 1000 (from 53 fewer to 425 more)	⊕OOO VERY LOW	CRITICAL	
Number of	urine tests p	ositive dur	ing treatment										
	randomised trials	1	no serious inconsistency		very serious ²	none	5/73 (6.8%)	10/90 (11.1%)		42 fewer per 1000 (from 87 fewer to 80 more)	⊕OOO VERY LOW	CRITICAL	

1 unclear risk of bias for detection and performance bias.

2 optimal information size criterion not met; confidence interval for the effect includes clinically significant benefit

N.2.2.2 Naltrexone versus active intervention for drug misuse

			Quality asse	essment			No of	patients		Effect	Quality	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Naltrexone versus active intervention	Active intervention	vention (95% CI) Absolute		Quality	Importance
Retained i	in treatment											
	randomised trials	very serious ¹			very serious ²	none	17/34 (50%)	5/17 (29.4%)	RR 1.7 (0.76 to 3.82)	206 more per 1000 (from 71 fewer to 829 more)	⊕OOO VERY LOW	CRITICAL
Urine test	positive for	drugs (du	ring treatment) -	Alcohol								

1		serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1/34 (2.9%)	1/17 (5.9%)	RR 0.5 (0.03 to 7.51)	29 fewer per 1000 (from 57 fewer to 383 more)	⊕OOO VERY LOW	CRITICAL
Urine test	positive for	drugs (du	ring treatment) -	Amphetamine								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	0/34 (0%)	0/17 (0%)	not estimable	not estimable ⁷	⊕OOO VERY LOW	CRITICAL
Urine test	positive for	druas (dı	uring treatment) -	Benzodiazepine	.	_			_		I	
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	1/34 (2.9%)	1/17 (5.9%)	RR 0.5 (0.03 to 7.51)	29 fewer per 1000 (from 57 fewer to 383 more)	⊕OOO VERY LOW	CRITICAL
Urine test	t positive for	drugs (dı	uring treatment) -	Cocaine							<u> </u>	
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	11/34 (32.4%)	8/17 (47.1%)	RR 0.69 (0.34 to 1.38)	146 fewer per 1000 (from 311 fewer to 179 more)	⊕OOO VERY LOW	CRITICAL
Urine test	t positive for	drugs (du	uring treatment) -	Marijuana								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	4/34 (11.8%)	3/17 (17.6%)	RR 0.67 (0.17 to 2.65)	58 fewer per 1000 (from 146 fewer to 291 more)	⊕OOO VERY LOW	CRITICAL
Urine test	positive for	drugs (dı	uring treatment) -	Opiates								
1	randomised		no serious inconsistency	no serious indirectness	very serious ²	none	3/34 (8.8%)	5/17 (29.4%)	RR 0.3 (0.08 to 1.11)	206 fewer per 1000 (from 271 fewer to 32 more)	⊕OOO VERY LOW	CRITICAL
Cocaine ι	use (post-trea	atment)	I	•	_ I				_ I		I	1
2	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	very serious ²	none	14/47 (29.8%)	10/49 (20.4%)	RR 1.34 (0.73 to 2.48)	69 more per 1000 (from 55 fewer to 302 more)	⊕OOO VERY LOW	CRITICAL

	randomised trials	very	no serious									
		serious ^{3,4}	inconsistency	no serious indirectness	very serious ²	none	13/47 (27.7%)	24/49 (49%)	RR 0.55 (0.35 to 0.87)	220 fewer per 1000 (from 64 fewer to 318 fewer)	⊕OOO VERY LOW	CRITICAL
njection of	drug use (po	st-treatme	ent)									
	randomised trials	very serious⁴	no serious inconsistency	no serious indirectness	very serious ²	none	4/16 (25%)	1/17 (5.9%)	RR 4.25 (0.53 to 34.1)	191 more per 1000 (from 28 fewer to 1000 more)	⊕OOO VERY LOW	CRITICAL
ays of d	rug use per r	nonth (6 r	nonth follow-up)	- Amphetamine (Better indica	ated by lower valu	ies)					
	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	very serious ⁶	none	23	21	-	MD 2.50 higher (3.86 lower to 8.86 higher)	⊕OOO VERY LOW	CRITICAL
ays of d	rug use per r	month (6 r	nonth follow-up)	- Benzodiazepine	e (Better indi	cated by lower va	alues)					
	randomised trials		no serious inconsistency	no serious indirectness	very serious ⁶	none	23	21	-	MD 2.0 higher (4.49 lower to 8.49 higher)	⊕OOO VERY LOW	CRITICAL
ays of d	rug use per r	nonth (6 r	nonth follow-up)	- Heroin (Better i	ndicated by	lower values)						
		very serious⁵	no serious inconsistency	no serious indirectness	very serious ⁶	none	23	21	-	MD 4.60 lower (12.74 lower to 3.54 higher)	⊕OOO VERY LOW	CRITICAL
eincarce	eration											
	randomised trials		no serious inconsistency	no serious indirectness	very serious ⁶	none	19/73 (26%)	21/55 (38.2%)	RR 0.64 (0.39 to 1.06)	137 fewer per 1000 (from 233 fewer to 23 more)	⊕OOO VERY LOW	CRITICAL
eincarce	eration - Duri	ng treatm	ent		I	ļ			I			
	randomised		no serious inconsistency	no serious indirectness	very serious ²	none	9/34 (26.5%)	9/17 (52.9%)	RR 0.5 (0.24 to 1.02)	265 fewer per 1000 (from 402 fewer to 11 more)	⊕OOO VERY LOW	CRITICAL
eincarce	eration - Post	t-treatmen	t	•	+	, ,			۱ <u>ــــــــــــــــــــــــــــــــــــ</u>			

	randomised trials		no serious inconsistency	no serious indirectness	very serious ²	none	5/16 (31.3%)	7/17 (41.2%)	RR 0.76 (0.3 to 1.91)	99 fewer per 1000 (from 288 fewer to 375 more)	⊕OOO VERY LOW	CRITICAL
eincarce	ration - 6 m	onth follow	w-up									
	randomised trials	, ,	no serious inconsistency	no serious indirectness	very serious ²	none	5/23 (21.7%)	5/21 (23.8%)	RR 0.91 (0.31 to 2.71)	21 fewer per 1000 (from 164 fewer to 407 more)	⊕OOO VERY LOW	CRITICAL
arole vio	lations (pos	t-treatmer	nt)		•							
	randomised trials		no serious inconsistency	no serious indirectness	very serious ²	none	2/31 (6.5%)	9/32 (28.1%)	RR 0.23 (0.05 to 0.98)	217 fewer per 1000 (from 6 fewer to 267 fewer)	⊕OOO VERY LOW	CRITICAL
Drug char	ges (post-tre	eatment)			1	_	11					I
	randomised trials		no serious inconsistency	no serious indirectness	very serious ²	none	3/31 (9.7%)	1/32 (3.1%)	RR 3.1 (0.34 to 28.19)	66 more per 1000 (from 21 fewer to 850 more)	⊕OOO VERY LOW	CRITICAL
Days of cr	iminal activ	ity per mo	onth (6 month foll	ow-up) (Better i	ndicated by lo	ower values)						
	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	very serious ⁶	none	23	21	-	Mean 14.4 days (SD 13.11)	⊕000 VERY LOW	CRITICAL

1 Cornish 1997 - unclear randomisation and allocation concealment; unclear blinding; ITT analysis

2 Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome (imprecision) respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

3 Caviello 2010 - Unclear randomisation and allocation concealment; unclear blinding; available case analysis

4 Lee 2016 - Appropriate randomisation and unclear allocation concealment; No blinding to participants; ITT analysis

5 Lobmaier 2010 - appropriate randomisation and allocation concealment; no blinding; ITT analysis

6 Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome (imprecision) respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

7 No event in either arm of the trial.

N.2.2.3 Methadone versus waitlist control

Quality assessment	No of patients	Effect	Quality Importa	ince
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No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Methadone versus waitlist control	Control	Relative (95% CI)	Absolute		
Drop-out												
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	152/191 (79.6%)	123/191 (64.4%)	RR 1.24 (1.09 to 1.4)	155 more per 1000 (from 58 more to 258 more)	⊕OOO VERY LOW	CRITICAL
Positive f	or opioids - P	ost-treatme	ent									
2	randomised	very serious ^{1,3}	serious ⁴	no serious indirectness	no serious imprecision	none	82/277 (29.6%)	90/270 (33.3%)	RR 0.86 (0.61 to 1.23)	47 fewer per 1000 (from 130 fewer to 77 more)	⊕OOO VERY LOW	CRITICAL
Positive f	or opioids - 2	month foll	ow-up									
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	no serious imprecision	none	10/106 (9.4%)	12/101 (11.9%)	RR 0.79 (0.36 to 1.76)	25 fewer per 1000 (from 76 fewer to 90 more)	⊕⊕OO LOW	CRITICAL
Positive f	or opioids - 3	month foll	ow-up									
2	randomised	very serious ^{1,3}	no serious inconsistency	no serious indirectness	serious⁵	none	40/233 (17.2%)	51/211 (24.2%)	RR 0.7 (0.5 to 0.99)	73 fewer per 1000 (from 2 fewer to 121 fewer)	⊕OOO VERY LOW	CRITICAL
Positive f	or opioids - 4	month foll	ow-up							-		
2	randomised trials	very serious ^{1,3}	no serious inconsistency	no serious indirectness	no serious imprecision	none	38/280 (13.6%)	39/258 (15.1%)	RR 0.91 (0.62 to 1.35)	14 fewer per 1000 (from 57 fewer to 53 more)	⊕⊕OO LOW	CRITICAL
Reincarce	eration (4 year	r follow-up))					I	I	l		I
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	143/191 (74.9%)	137/191 (71.7%)		29 more per 1000 (from 57 fewer to 129 more)	⊕OOO VERY LOW	CRITICAL

1 Dolan 2003/2005 - appropriate randomisation and allocation concealment; unclear blinding and available case analysis

2 Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference(MID) for the outcome (imprecision) respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

3 Shearer 2006 - unclear randomisation and allocation concealment; unclear blinding; available case analysis

4 Evidence was downgraded by one level due to serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of 50%-74.99%) and by two levels due to very serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of >75%).

5 Rich 2015 - appropriate randomisation and allocation concealment; unclear blinding; ITT analysis

N.2.2.4 Alpha-adrenergic agonists versus opioid maintenance

			Quality assess	sment			No of	oatients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Alpha- adrenergics	Opioid maintenance	Relative (95% Cl)	Absolute		
Total with	drawal sympto	oms (follow-up	o mean 10 days; Be	etter indicated by	lower value	s)						
	randomised trials			no serious indirectness	very serious ¹	none	29	34	-	MD 24 higher (73.86 lower to 121.86 higher)	⊕⊕OO LOW	CRITICAL

optimal information size criterion not met; confidence interval of effect includes both appreciable benefit and harm

N.2.2.5 Opioid substitution versus active intervention or placebo

			Quality asses	sment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Opioid substitution therapy versus active intervention	Control	Relative (95% Cl)	Absolute		
Drop-out												
	randomised trials	very serious ¹	no serious inconsistency	serious ²	serious ³	none	21/102 (20.6%)	29/104 (27.9%)	RR 0.75 (0.46 to 1.22)	70 fewer per 1000 (from 151 fewer to 61 more)	⊕OOO VERY LOW	CRITICAL
Abstinend	ce - Post-treat	tment										

4		· 4			. 5		74/400	70/440	<u>DD 4 00 (0 0</u>	40 4000		
	randomised trials		no serious inconsistency	no serious indirectness	serious⁵	none	74/100 (74%)	79/113 (69.9%)	RR 1.06 (0.9 to 1.25)	42 more per 1000 (from 70 fewer to 175 more)	⊕⊕OO LOW	CRITICAL
Abstinend	ce - 1 month f	ollow-up										
			no serious	no serious	very serious ⁶	nono	45/72	64/87	RR 0.85	110 fewer per 1000	000	CRITICAL
	trials			indirectness	very senous		(62.5%)	(73.6%)	(0.68 to 1.06)	(from 235 fewer to 44 more)	⊕000 VERY LOW	CRITICAL
Abstinend	ce - 3 month f	ollow-up	<u></u>	ļ		ļ				<u> </u>		
1	randomised trials		no serious inconsistency	no serious indirectness	very serious ⁷	none	31/46 (67.4%)	27/48 (56.3%)	RR 1.2 (0.87 to 1.65)	113 more per 1000 (from 73 fewer to 366 more)	⊕OOO VERY LOW	CRITICAL
Abstinend	ce - 6 month f	ollow-up	<u> </u>									
2	randomised trials	10	no serious inconsistency	no serious indirectness	very serious ⁹	none	26/75 (34.7%)	21/75 (28%)	RR 1.08 (0.74 to 1.59)	22 more per 1000 (from 73 fewer to 165 more)	⊕OOO VERY LOW	CRITICAL
Opioid ab	ouse (3 month	follow-up)	I									
	randomised trials	,	no serious inconsistency	serious ²	very serious ¹⁰	none	32/60 (53.3%)	37/56 (66.1%)	RR 0.81 (0.6 to 1.09)	126 fewer per 1000 (from 264 fewer to 59 more)	⊕OOO VERY LOW	CRITICAL
Self-repor	rted injection	drug use - Pe	ost-treatment		I	I				I		
	randomised trials			no serious indirectness	very serious ¹¹	none	8/24 (33.3%)	7/12 (58.3%)	RR 0.57 (0.27 to 1.2)	251 fewer per 1000 (from 426 fewer to 117 more)	⊕⊕OO LOW	CRITICAL
Self-repor	rted injection	drug use - 3	month follow-up		<u> </u>							
		-										

1	randomised trials	no serious risk of bias	no serious inconsistency	no serious indirectness	very serious ¹²	none	7/24 (29.2%)	6/12 (50%)	RR 0.58 (0.25 to 1.35)	210 fewer per 1000 (from 375 fewer to 175 more)	⊕⊕OO LOW	CRITICAL
Number o	of times rearr	ested (3 mon	th follow-up) (Be	tter indicated by	lower values	s)	•					
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	very serious ¹⁰	none	60	56	-	SMD 0.02 lower (0.39 lower to 0.34 higher)	⊕000 VERY LOW	CRITICAL
Rearrest	for drug crim	es (3 month f	ollow-up)									
1	randomised trials	serious ¹²	no serious inconsistency	serious ¹²	very serious ¹³	none	8/60 (13.3%)	13/56 (23.2%)	RR 0.57 (0.26 to 1.28)	100 fewer per 1000 (from 172 fewer to 65 more)	⊕OOO VERY LOW	CRITICAL
Reincarce	eration (post-	treatment)										
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	very serious ¹⁰	none	24/60 (40%)	28/56 (50%)	RR 0.8 (0.53 to 1.2)	100 fewer per 1000 (from 235 fewer to 100 more)	⊕OOO VERY LOW	CRITICAL

1 high risk performance of bias

2 serious indirectness Maguara 2009 due to population)

3 Optimal information size criterion not met (combined n = 206)

4 high risk performance of bias

5 Optimal information size criterion not met (n = 213)

6 Optimal information size criterion not met (n = 159)

7 Optimal information size criterion not met (n = 94)

8 ROB - Sheared = high performance bias + unclear detection bias + 2 unclear bias.

9 Optimal information size criterion not met (Combined n = 150)

10 Optimal information size criterion not met (n = 116)

11 Optimal information size criterion not met (n = 36)

12 Optimal information size criterion not met (events<100) and CI of effect includes appreciable benefit and harm

N.2.3 Combined pharmacological and psychological interventions

N.2.3.1 Antidepressants plus psychological therapy versus psychological therapy alone

			Quality asso	essment			No of patie	ents		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Antidepressants + psychological therapy	Psychological therapy only	Relative (95% Cl)	Absolute		
No. partio	cipants who f	ailed to c	omplete treatmen	t (follow-up me	an 12 weeks		· · · · · · · · · · · · · · · · · · ·			·		
1	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	13/31 (41.9%)	9/29 (31%)	RR 1.35 (0.68 to 2.67)	109 more per 1000 (from 99 fewer to 518 more)	⊕⊕OO LOW	CRITICAL
Spielberg	ger state anxi	ety invent	tory score (follow	-up mean 12 we	eks; Scale f	rom 20 to 80; lowe	er better)					
1	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	31	29	-	MD 0.30 lower (6.44 lower to 5.84 higher)		CRITICAL
Hamilton	depression r	ating sca	le score (follow-u	ıp mean 12 weel	ks; Scale fro	m 0 to 52; lower b	etter)					
1	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	31	29	-	MD 3.10 lower (6.18 to 0.02 lower)	⊕⊕OO LOW	CRITICAL

¹ unclear selection, detection and attrition bias ² optimal information size criterion not met

N.2.4 Support and educational interventions

Psychoeducation versus control N.2.4.1

			Quality asse	ssment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Other considerations	Psychoeducation versus control/TAU	Control	Relative (95% Cl)	Absolute			
Number of	days with unc	ontrolled o	drinking (Better ind	icated by lower va	lues)							
	trials			indirectness	very serious ²	none	18	16	-	MD 4.85 days fewer (11.46 fewer to 1.76 more)	⊕OOO VERY LOW	CRITICAL

¹ high risk for performance, detection and selective reporting ² Optimal information size criterion not met (N<400); 95% CI of effect includes both appreciable benefit and harm

N.2.4.2 Employment workshop versus control or treatment as usual

			Quality as	sessment			No of pa	tients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Employment workshops	Control/TAU	Relative (95% Cl)	Absolute		
No. of par	ticipants emp	oloyed (fol	low-up 12-52 weel	ks)								
2		very serious ¹	very serious ²	serious ³	serious ⁴	none	220/272 (80.9%)	189/257 (73.5%)	RR 1.24 (0.84 to 1.81)	176 more per 1000 (from 118 fewer to 596 more)	⊕OOO VERY LOW	CRITICAL
Days in pa	aid employme	ent (follow	-up mean 52 week	s; Better indicat	ed by higher val	ues)		·				
1		- , _			no serious imprecision	none	244	233	-	MD 10.20 higher (11.8 lower to 32.2 higher)	⊕⊕OO LOW	CRITICAL

1 high risk of performance, detection and reporting bias, unclear bias on 3 other dimensions

2 12=73%; random effects model used; no reasons for this heterogeneity were identified; study effect estimates were RR=1.58 [1.06, 2.36] for Hall (1961) and RR = 1.06 [0.97, 1.17] for Webster (2014)

3 Hall 1981-unclear whether the population have a current drug or other mental health problem

4 Hall 1981, small sample size

5 high risk of detection and performance bias, unclear risk on 3 other domains

N.2.5 Physical interventions

N.2.5.1 Acupuncture versus active intervention

			Quality asse	essment			No of J	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Acupuncture	active intervention	Relative (95% CI)	Absolute		
Drop-out												
	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	50/82 (61%)	32/76 (42.1%)	RR 1.45 (1.06 to 1.99)	189 more per 1000 (from 25 more to 417 more)	⊕⊕OO LOW	CRITICAL
Urine test	positive for d	rug use po	ost-treatment									

randomised trials	very serious ³	very serious ⁴	very serious ⁶	none	15/46 (32.6%)	8/62 (12.9%)	RR 3.65 (0.33 to 41)	342 more per 1000 (from 86 fewer to 1000 more)	⊕000 VERY LOW	CRITICAL

1 allocation concealment, attrition and selective reporting all high risk of bias

2 Optimal information size criterion not met (N<300 events)

3 Both studies had allocation concealment, attrition and selective reporting all high risk of bias

4 1² 66% - random effects model used; large variation in effect sizes: Berman 16.39, Konefal 1.59, but no explanation for the heterogeneity was identified

5 For one study (Konefal 1995) - only 51% of participants were in contact with CJS

6 Optimal information size criterion not met (N < 300 events) and CI of effect includes both appreciable benefit and harm

N.3 Interventions for 'other' mental health disorders

N.3.1 Depression

N.3.1.1 Psychotherapy vs PSYCHOED

			Quality asse	ssment			No of pa	tients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychotherapy		Relative (95% Cl)	Absolute		
Depression	n by HRSD sca	les (Scale	from 0 to 52; lower	better)								
1		- /	no serious inconsistency	no serious indirectness	serious ²	none	19	19	-	MD 6.5 lower (12.52 to 0.48 lower)	⊕OOO VERY LOW	CRITICAL
Depression	n by HRSD sca	les (13 we	eks Follow-up) (Sca	le from 0 to 52; lov	wer better)							
1		· 1	no serious inconsistency	no serious indirectness	serious ²	none	19	19	-	MD 3.8 higher (3.83 lower to 11.43 higher)	⊕OOO VERY LOW	CRITICAL

¹ Johnson 2012 - Unclear risk of bias, unclear blinding of participants and care administrators, blinding of outcome assessors, low attrition bias, unclear selective outcome bias, low other risk of bias

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.1.2 Group therapy vs Individual therapy for depression

			Quality asso	essment			No of	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Group therapy	Individual therapy	Relative (95% CI)	Absolute		
Depressio	n by BDI (Scale	e from 0 to	20; lower better)				•	•				
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	5	5	-	MD 3.2 lower (13.56 lower to 7.16 higher)	⊕OOO VERY LOW	CRITICAL
Depressio	n by Hopeless	scale (Bett	er indicated by lowe	er values)			-	•				
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	5	5	-	MD 2.6 higher (4.98 lower to 10.18 higher)	⊕OOO VERY LOW	CRITICAL
Depressio	n by MMPI D s	cale (Better	indicated by lower	values)	*	•		•	•			•
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	5	5	-	MD 12.6 higher (3.38 lower to 28.58 higher)	⊕OOO VERY LOW	CRITICAL
Depressio	n by MMPI D se	cale (39 we	eks Follow-up) (Bet	ter indicated by lo	wer values)		•	•			•	
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	5	5	-	MD 4.8 higher (9.68 lower to 19.28 higher)	⊕OOO VERY LOW	CRITICAL
Depressio	n by Multiple a	ffect adject	ive Check list D sca	ale (Better indicate	ed by lower val	ues)						
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	5	5	-	MD 0.6 higher (4.93 lower to 6.13 higher)	⊕OOO VERY LOW	CRITICAL
² The evia	lence was dow the outcome, i	vngraded k	by one level and tw	o levels if the co	nfidence interv		hed one or	both boundar		defined minimally imp l group (if MD was use		

N.3.1.3 Arts-based therapy vs TAU for depression

Quality assessment							No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Arts-based therapy	TAU	Relative			
Change in	Change in Adult Nowicki-Strickland Locus of Control Scale (ANS) (Better indicated by lower values)											
1	randomised	very	no serious	no serious	serious ²	none	72	50	-	MD 3.88 lower (8.15 lower	⊕000	CRITICAL

	trials	serious ¹	inconsistency	indirectness						to 0.39 higher)	VERY LOW	
Change in	Adult Nowick	-Strickland	Locus of Contro	ol Scale (ANS) - Ma	le (Better ind	icated by lower va	alues)					
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	37	25	-	MD 2.26 lower (4.18 to 0.34 lower)	⊕OOO VERY LOW	CRITICAL
Change in	Adult Nowick	-Strickland	Locus of Contro	ol Scale (ANS) - Fe	male (Better i	ndicated by lower	values)			•		
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	35	25	-	MD 6.81 lower (11.97 to 1.65 lower)	⊕OOO VERY LOW	CRITICAL
Change in	Beck Depress	ion Invente	ory (BDI): Total (E	Better indicated by	lower values	.)						
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	111	45	-	MD 6.5 lower (9.33 to 3.67 lower)	⊕OOO VERY LOW	CRITICAL
Change in	Beck Depress	ion Invent	ory (BDI): Total -	Male (Better indica	ted by lower	values)						-
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	35	25	-	MD 6.81 lower (11.97 to 1.65 lower)	⊕OOO VERY LOW	CRITICAL
Change in	Formal Eleme	nts of Arts	Therapy Scale ra	ting guide(FEATS): Prominenc	e of color (Better	ndicated by low	er valu	les)			-
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	65	19	-	MD 0.81 lower (1.51 to 0.11 lower)	⊕OOO VERY LOW	LIMITEDIMPORTAN
Change in	Formal Eleme	nts of Arts	Therapy Scale ra	ting guide (FEATS	6): color fit (B	etter indicated by	lower values)			•		
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	65	19	-	MD 0.45 lower (0.84 to 0.06 lower)	⊕OOO VERY LOW	LIMITED IMPORTANT

¹ Gussak 2009 - Unclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessorsUnclear randomization and allocation, No blinding of patients and care administrators, Blinding of outcome assessors, (Gussak 2009)

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries

N.3.2 Vulnerable inmates with suicidal risks

N.3.2.1 Social problem solving group vs No treatment control

	Quality assessment						No of patients			Effect	Quality	Importance
No of studies	Design	esign Risk of Inconsistency Indirectness Imprecision Other consideration		Other considerations	Social problem solving group for vulnerable	No treatment	Relative (95%	Absoluto				

							inmates		CI)			
Depressi	on by HADS s	cale (Scal	e from 0 to 21; lov	ver better)					<u> </u>			
1	randomised	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23	23	-	MD 3.6 lower (5.76 to 1.44 lower)	⊕000 VERY LOW	CRITICAL
Anxiety b	y HADS scale	s (Scale f	rom 0 to 21; lower	better)								
1		very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	23	23	-	MD 2.9 lower (4.67 to 1.13 lower)	⊕OOO VERY LOW	CRITICAL
Depressi	on by Beck Ho	opeless so	ales (Scale from () to 20; lower bet	ter)							
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23	23	-	MD 2.5 lower (4.89 to 0.11 lower)	⊕000 VERY LOW	CRITICAL
Decision	making ability	by SPSI:	R scales (Scale fro	om 0 to 21; lower	better)							
1		very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	23	23	-	MD 5.3 higher (2.66 to 7.94 higher)	⊕⊕OO LOW	CRITICAL
Depressi	on by HADS s	cale (13 w	eeks Follow-up) (Scale from 0 to 2	0; lower better)		· · · · · · · · · · · · · · · · · · ·			••		•
1		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23	23	-	MD 3.3 lower (5.19 to 1.41 lower)	⊕000 VERY LOW	CRITICAL
Anxiety b	y HADS scale	s (13 wee	ks Follow-up) (Sca	ale from 0 to 21;	lower better)							
1		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23	23	-	MD 2.7 lower (4.61 to 0.79 lower)	⊕000 VERY LOW	CRITICAL
Depressi	on by Beck Ho	peless so	ales (13 weeks Fo	llow-up) (Scale f	rom 0 to 20; low	ver better)						
1	randomised	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23	23	-	MD 2.8 lower (5.13 to 0.47 lower)	⊕000 VERY LOW	CRITICAL

¹Biggam 2002 - Unclear risk of selection bias, No blinding, low attrition bias, unclear selective outcome reporting, low other risk of bias ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.3 **Anxiety disorders**

Self-help therapy vs Wait-list control N.3.3.1

Quality assessment No of patients Effect Quality Import

No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Self-help materials	Control	Relative (95% Cl)	Absolute		
Anxiety by	nxiety by HADS scales (Scale from 0 to 21; lower better)											
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	15	18	-	MD 1.06 lower (3.63 lower to 1.51 higher)	⊕⊕OO LOW	CRITICAL
Anxiety by	Anxiety by HADS scales (4 weeks follow-up) (Scale from 0 to 21; lower better)											
	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	15	18	-	MD 2.98 lower (5.82 to 0.14 lower)	⊕⊕OO LOW	CRITICAL

¹ Maunder 2009 - low selection risk of bias, No blinding of participants but blinding of care administrators (+), unclear outcome assessor, unclear attrition risk of bias, unclear other risk of bias (blocked randomization with single blinded trial)

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.4 PTSD

N.3.4.1 Psychotherapy vs Wait-list/No-contact control

Quality assessment								nts		Effect	Quality	Importance		
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychotherapy Control		Relative (95% Cl)	Absolute				
Trauma by ⁻	Trauma by TSI - Group Therapy (Wait-list/No-contact Control) (Scale from 0 to 300; lower better)													
		very serious ^{1,2}	,	no serious indirectness	serious ⁴	none	17	23	-	MD 11.67 lower (30.36 lower to 7.02 higher)	⊕000 VERY LOW	CRITICAL		

¹ Cole 2007 - high risks of selection bias, No blinding, Unclear attrition bias, low selective outcome bias and low other risk of bias

² Bradley 2003 - unclear risks of selection bias, No blinding, Unclear attrition, High selective outcomes bias and low other risks of bias 12=83%; studies combined by randomised model because similar population, intervention and the outcome measured by same measure.

³ Evidence was downgraded by one level due to serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of 50%-74.99%) and by two levels due to very serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of >75%).

⁴The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.4.2 TIR vs Wait-list control

			Quality asses	sment			No	of patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	TIR	Wait-list control	Relative (95% Cl)	Absolute		
Depression	by BDI - Traun	natic Incide	nt Reduction (Scale f	rom 0 to 63; lower l	better)							•
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 3.8 lower (7.52 to 0.08 lower)	⊕OOO VERY LOW	CRITICAL
Depression	by BDI total (1	3 weeks Fo	llow-up) (Scale from	0 to 63; lower bette	r))	<u>.</u>	•					
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 7.8 lower (12.64 to 2.96 lower)	⊕OOO VERY LOW	CRITICAL
PTSD by PS	SS scales at po	st-treatmen	t (Scale from 0 to 51;	lower better)								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	56	67	-	MD 4.1 lower (7.96 to 0.24 lower)	⊕OOO VERY LOW	CRITICAL
PTSD by PS	SS scales (13 w	eeks follow	-up) (Scale from 0 to	51; lower better)								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 7.3 lower (11.49 to 3.11 lower)	⊕OOO VERY LOW	CRITICAL
Generalized	Expectancy for	or Success	Scale at post-treatme	ent (Scale from 30 to	o 150; higher	better)	• •					
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 15.9 higher (5.7 to 26.1 higher)	⊕OOO VERY LOW	CRITICAL
Generalized	d Expectancy for	or Success	Scale (13 weeks follo	w-up) (Scale from 3	30 to 150; hig	her better)						
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 3.6 higher (2.69 lower to 9.89 higher)	⊕OOO VERY LOW	CRITICAL
MH outcom	es: Clinical An	xiety scale	at post-treatment (Sc	ale from 0 to 100; lo								
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	Serious ²	none	56	67	-	MD 3.3 lower (8.55 lower to 1.95 higher)	⊕OOO VERY LOW	CRITICAL
MH outcom		xiety scale	(13 weeks follow-up)	(Scale from 0 to 10		er)					-	
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	56	67	-	MD 7.8 lower (12.64 to 2.96 lower)	⊕OOO VERY LOW	CRITICAL

¹ Valentine 2001 - high risk of selection bias, No blinding, unclear attrition bias, low selective outcome bias, low other risk of bias ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference

(MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.4.3 TARGET vs SGT

			Quality as	sessment			No o patien			Effect	Quality	Importance
No of studies	Design	Imprecision	Other considerations	TARGET		Relative (95% Cl)	Absolute					
PTSD symp	otoms by CAPS	scales (Be	etter indicated by low	ver values)	•	•						
	randomised trials			no serious indirectness	no serious imprecision	none	38	34	-	MD 0.5 lower (11.01 lower to 10.01 higher)	⊕⊕⊕O MODERATE	CRITICAL
Heartland forgiveness scale (Better indicated by lower values)												
	randomised trials			no serious indirectness	serious ²	none	23	9	-	MD 4.6 higher (6.73 lower to 15.93 higher)	⊕⊕OO LOW	CRITICAL

¹ Ford 2013- low risk of selection bias, blinding of care administrators and outcome assessors but no blinding of participants, low attrition rate, low selective outcome bias, low other risk of bias Ford - low risk of selection bias, blinding of care administrators and outcome assessors but no blinding of participants, low attrition rate, low selective outcome bias, low other risk of bias ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries..

N.3.4.4 Focused group therapy vs Wait-list control

			Quality as	sessment			No of pat	ients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Focused group therapy	Wait-list control	Relative (95% Cl)	Absolute		
Symptom	checklist-90-R	: Global S	everity Index (Scale	from 0 to 90; low	er better)							
1	randomised trials	1	no serious inconsistency	no serious indirectness	no serious imprecision	none	4	5	-	MD 16.3 lower (26.23 to 6.37 lower)	⊕⊕OO LOW	CRITICAL
Symptom	Checklist-90R	Positive S	Symptom Distress I	ndex (Scale from	0 to 90; lower bet	ter)						
1	randomised trials	1	no serious inconsistency	no serious indirectness	serious ²	none	4	5	-	MD 13.9 lower (24.8 to 3 lower)	⊕OOO VERY LOW	CRITICAL
Symptom	Checklist-90R	Positive S	Symptom Total (Sca	ale from 0 to 90; lo	ower better)							
1	randomised trials	1	no serious inconsistency	no serious indirectness	no serious imprecision	none	4	5	-	MD 16.1 lower (26.67 to 5.53 lower)	⊕⊕OO LOW	CRITICAL

¹ Cole 2007 - high risks of selection bias, No blinding, Unclear attrition bias, low selective outcome bias and low other risk of bias

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.4.5 Group Therapy vs No contact control

			Quality asses	ssment			No of	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Group Therapy	No contact control	Relative (95% Cl)	Absolute		
IIP-32 scale	IIP-32 scales (Scale from 0 to 128; lower better)											
		- 1	no serious inconsistency	no serious indirectness		none	13	18	-	MD 10.1 lower (24.43 lower to 4.23 higher)		CRITICAL

¹Bradley 2003 - unclear risks of selection bias, No blinding, Unclear attrition, High selective outcomes bias and low other risks of bias

N.3.5 ADHD

N.3.6 Methylphenidate vs Placebo

			Quality ass	sessment			No of patien	ts		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Methylphenidate (MPH)	Placebo	Relative (95% Cl)	Absolute		
Conner Ad	dult ADHD rati	ing scale -	Observer: Screen	ing Version (CAA	RS-OSV) – post	-treatment (52 wee	eks) (Scale from 0 to	o 90; Iow	er better)			
	randomised trials	serious ^{1,2}		no serious indirectness	Serious⁴	none	42	42	-	MD 12.85 lower (22.5 to 3.20 lower)	⊕OOO VERY LOW	CRITICAL
Conner Ad	dult ADHD rati	ing scale -	Observer: Screen	ing Version (CAA	RS-OSV) - Follo	ow-up (3 years) (Co	py) (Better indicate	d by low	er values)			
		1			no serious imprecision	none	15	5	-	MD 16.9 lower (24.5 to 9.3 lower)	⊕⊕OO LOW	CRITICAL
Number o	f participants	with drug	negative urine									
		2		no serious indirectness	serious⁵	none	6/27 (22.2%)	4/27 (14.8%)		74 more per 1000 (from 77 fewer to 551 more)	⊕OOO VERY LOW	CRITICAL

¹Ginsberg 2012 - high risk of selection bias, No blinding, low risk of attrition, unclear selective outcome reporting and low risk of other bias ²Konstenius 2013- low risk of selection bias, Blinding of participants, care administrators and outcome detectors, unclear attrition bias and unclear selective outcome reporting, low risk of other bias ³ Evidence was downgraded by one level due to serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of 50%-74.99%) and by two levels due to very serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of >75%)

⁴ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries. If SMD was used, +0.5 and -0.5 on the SMD scale as MID boundaries.'

⁵ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference for the outcome (imprecision) respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.3.7 Antisocial personality disorders

N.3.7.1 Tiagabine vs Placebo

			Quality asses	sment			No of pa	atients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Tiagabine	Placebo	Relative (95% CI)	Absolute		
Aggressiv	gressive Response (Better indicated by lower values)											
1	randomised trials	· ·	no serious inconsistency	no serious indirectness	serious ²	none	6	6	-	MD 1.86 lower (2.7 to 1.02 lower)	⊕OOO VERY LOW	CRITICAL
Number of	subjects with ad	verse effe	cts				•				•	
		- ,		no serious indirectness	very serious ³	none	6/157* (3.8%)	6/65* (9.2%)	RR 0.41 (0.14 to 1.24)	54 fewer per 1000 (from 79 fewer to 22 more)	⊕OOO VERY LOW	CRITICAL

Gowin 2012- Unclear risk of selection bias, blinding to participants and care person involved (+), low risk of attrition, unclear selective outcome reporting, low risk of other bias.

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

³ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome (imprecision) respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

*Denominator - total number of 'Yes' reports to the side-effects at least once

N.3.8 Severe mental illness

N.3.8.1 IM Paliperidone vs Oral Antipsychotics for schizophrenia

Quality assessment	No of patients	Effect	Quality	Importance	
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No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	IM Paliperidone	Oral antipsychotics	Relative (95% CI)	Absolute		
First-time	treatment fa	ilure*										
1	randomised trials	very serious ¹		no serious indirectness	serious ²	none	90/226 (39.8%)	117/218 (53.7%)	RR 0.74 (0.61 to 0.91)	140 fewer per 1000 (from 48 fewer to 209 fewer)	⊕000 VERY LOW	CRITICAL
Incidence	of prolactin-	-related si	de-effects	•			•				·	
1	randomised trials	very serious ¹			no serious imprecision	none	53/226 (23.5%)	9/219 (4.1%)	RR 5.71 (2.89 to 11.28)	194 more per 1000 (from 78 more to 422 more)	⊕⊕OO LOW	LIMITED IMPORTANCE

¹ Alphs 2015a- Unclear risk of selection bias, no blinding, low risk of attrition bias, low risk of selective outcome bias, low risk of other bias ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

* First-time treatment failure - arrest/incarceration, psychiatric hospitalization, suicide, discontinuation of antipsychotic treatment due to inadequate efficacy, treatment supplementation with another antipsychotic due to inadequate efficacy, discontinuation of antipsychotic treatment due to safety or tolerability concerns, or an increase in the level of psychiatric services to prevent imminent psychiatric hospitalization

The Citizenship project for severe mental illness N.3.8.2

			Quality as	sessment			No of patients	5		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	The Citizenship Project	TAU	Relative (95% Cl)	Absolute		
Overall qua	ality of life (Be	tter indica	ted by lower values		•	-			•			
	randomised trials	very serious ¹		no serious indirectness	serious ²	none	73	41	-	MD 0.68 higher (0 to 1.36 higher)	⊕OOO VERY LOW	CRITICAL
Number of	all conviction	s (Better ir	ndicated by lower va	lues)								
	randomised trials			no serious indirectness	no serious imprecision	none	73	41	-	MD 0.05 higher (0.79 lower to 0.89 higher)	⊕⊕OO LOW	CRITICAL
Alcohol co	mposite ratio	(Scale from	n 0 to 9; lower bette	r)	•			•				
	randomised trials	very serious ¹		no serious indirectness	Serious ²	none	40	29	-	MD 0.29 lower (0.57 to 0.01 lower)	⊕OOO VERY LOW	IMPORTANT
Brief Psych	niatric Rating	Scale: With	hdrawal symptoms (Scale from 1 to 7;	lower better)							
	randomised trials	very serious ¹		no serious indirectness	Serious ²	none	73	41	-	MD 0.28 higher (0.01 to 0.55 higher)	⊕OOO VERY LOW	IMPORTANT
Addition se	everity index:	Drug index	(Scale from 0 to 9;	lower better)								
1	randomised	very	no serious	no serious	serious ²	none	73	41	-	MD 0.04 lower (0.08 lower	⊕000	IMPORTANT

trials	serious ¹	inconsistency	indirectness			to 0 higher)	VERY	
							LOW	

¹ Clayton 2013 - Unclear selection bias, No blinding, Unclear attrition, low risk of selective outcome reporting, low risk of other bias

²The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.8.3 Individual Placement and Support vs Peer support for severe mental illness

			Quality asse	essment			No of patient	S		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Individual Placement and Support (IPS)	Peer support	Relative (95% CI)	Absolute		
Competiti	ve job placem	ent										
1	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	13/42 (31%)	3/43 (7%)	RR 4.44 (1.36 to 14.46)	240 more per 1000 (from 25 more to 939 more)	⊕⊕OO LOW	IMPORTANT
Number o	f hospitalizati	ons (Bette	er indicated by lov	ver values)	•		•					•
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	41	43	-	MD 0.5 higher (0.07 lower to 1.07 higher)	⊕⊕OO LOW	CRITICAL
Number o	f days being l	nospitaliz	ed (Better indicate	d by lower value	es)							
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	41	43	-	MD 5.51 higher (1.91 lower to 12.93 higher)	⊕OOO VERY LOW	CRITICAL

¹ Bond 2015 - Appropriate randomization with concealed allocation, no blinding of participants and care administrators, ITT analysis, appropriate outcome report ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries..

N.3.9 Uncategorised mental health disorders

N.3.9.1 Parenting from inside vs wait-list control

Quality assessment	No of patients	Effect	Quality	Importance	
					i i

No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Parenting from the Inside (PFI)	Wait-list control	Relative (95% Cl)	Absolute		
Parenting	Stress Inde	x (Scale f	from 27 to 135; lo	wer better)								
	randomised trials	- /			no serious imprecision	none	60	76	-	MD 0.04 higher (0.17 lower to 0.25 higher)	⊕⊕OO LOW	CRITICAL
Brief Sym	ptom Invent	ory (BSI)	: Total (Scale from	m 0 to 212; lowe	r better)	•						
	randomised trials	- /		no serious indirectness	serious ²	none	60	76	-	MD 0.2 higher (0.12 lower to 0.52 higher)	⊕OOO VERY LOW	CRITICAL
Parenting	Alliance (So	cale from	20 to 100; higher	better)								
	randomised trials	1			no serious imprecision	none	60	76	-	MD 0.31 lower (6.23 lower to 5.61 higher)	⊕⊕OO LOW	IMPORTANT

¹ Loper 2011 - Unclear selection bias; No blinding; Unclear attrition bias, low risk of selective outcomes, low risk of other bias ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.3.9.2 Music therapy vs standard care for anxiety and depression disorders

			Quality as	sessment			No of I	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Music therapy	Standard care	Relative (95% Cl)	Absolute		
State and T	Frait Anxiety Ir	ventory -	State (Scale from 2	to 80; lower bette	er)							
1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	93	91	-	MD 8.05 lower (10.74 to 5.36 lower)	⊕⊕⊕O MODERATE	CRITICAL
State and 1	Frait Anxiety Ir	ventory -	Trait (Scale from 20	to 80; lower bette	r)							
1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	93	91	-	MD 8.51 lower (10.91 to 6.11 lower)	⊕⊕⊕O MODERATE	CRITICAL
Brief Symp	tom Inventory	(BSI): Tot	al (Scale from 0 to 2	212; lower better)					•			
1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	93	91	-	MD 8.81 lower (11.82 to 5.8 lower)	⊕⊕⊕O MODERATE	CRITICAL
Rosenberg	self-esteem i	nventory (S	Scale from 0 to 30;	higher better)								
1	randomised trials			no serious indirectness	no serious imprecision	none	93	91	-	MD 2.26 higher (0.98 to 3.54 higher)	⊕⊕⊕O MODERATE	CRITICAL
Texas soci	al behaviour i	nventory (S	Scale from 0 to 128	higher better)								
1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	93	91	-	MD 7.54 higher (3.24 to 11.84 higher)	⊕⊕⊕O MODERATE	CRITICAL

¹ Chen 2015 - Appropriate randomization with proper concealment; blinding of care administrators, but not participants; ITT analysis; appropriate outcome report

N.3.9.3 Music therapy vs wait-list control for antisocial personality disorders

			Quality as	sessment			No of	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Music therpy	Wait-list control	Relative (95% Cl)	Absolute		
ASP-1: Self	f-management	of psychia	atric symptoms (Sc	ale from 0 to 4; hig	her better)							
-		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	8	5	-	MD 0.44 higher (0.03 lower to 0.91 higher)	⊕000 VERY LOW	CRITICAL
ASP-4: self	f-management	of assault	ive symptoms (Scal	e from 0 to 4; high	ner better)							
-	randomised trials	1	no serious inconsistency	no serious indirectness	serious ²	none	8	5	-	MD 0.11 lower (0.67 lower to 0.45 higher)	⊕000 VERY LOW	CRITICAL
ASP-9: Inte	erpersonal skil	Is (Scale fi	rom 0 to 4; higher b	etter)	•		•					
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	8	5	-	MD 0.02 higher (0.06 lower to 0.1 higher)	⊕⊕OO LOW	CRITICAL
Social dysf	unction and a	ggression	scale (Scale from 0	to 44; lower bette	r)	·						
		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	8	5	-	MD 0.8 lower (2.73 lower to 1.13 higher)	⊕000 VERY LOW	CRITICAL
forensic ps	sychiatric prof	iles 40 (FP	40): positive coping	skills (Better indi	cated by lower va	lues)						
1		- 1	no serious inconsistency	no serious indirectness	serious ²	none	8	5	-	MD 0.43 higher (0.12 to 0.74 higher)	⊕OOO VERY LOW	CRITICAL

¹ Hakvoort 2013 - unclear randomisation and concealment; No blinding; available case analysis; appropriate outcome report

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.4 Interventions for acquired cognitive impairment

A narrative overview of indirectly relevant systematic reviews was performed for this question. The evidence was not from criminal justice system populations and was not subject to critical appraisal of quality.

N.5 Interventions for paraphilic disorders

N.5.1 Medroxyprogesterone + psychological intervention compared to psychological intervention only for paraphilic disorders

			Quality as	sessment			Nº of patie	ents	Effec	:t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Medroxyprogesterone + psych intervention	psych intervention only	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance
Repetition	of anomalous	behaviour (follov	v up: range 15-52	weeks to; assess	sed with: self-rep	ort questionnaire and case	notes)					
2	randomised trials	serious	serious ¹	not serious ²	very serious ²	none	2/25 (8.0%)	6/27 (22.2%)	RR 0.58 (0.04 to 8.30)	93 fewer per 1,000 (from 213 fewer to 1,000 more)	⊕○○○ VERY LOW	CRITICAL
Reduced a	anomalous des	sires (follow up: 5	2; assessed with:	self-report quest	ionnaire)		••					
1	randomised trials	serious ³	not serious	not serious	very serious ²	none	5/10 (50.0%)	6/10 (60.0%)	RR 0.83 (0.12 to 1.55)	102 fewer per 1,000 (from 330 more to 528 fewer)	⊕⊖⊖⊖ VERY LOW	CRITICAL
Dropout (fe	ollow up: 15; a	ssessed with: nu	mber of participar	ts who did not co	omplete treatmen	t)						
1	randomised trials	serious ³	not serious	not serious	serious ²	none	10/15 (66.7%)	5/17 (29.4%)	RR 2.27 (1.00 to 5.14)	374 more per 1,000 (from 0 fewer to 1,000 more)	⊕⊕⊖⊖ LOW	IMPORTANT

CI: Confidence interval; RR: Risk ratio

1. Downgraded for inconsistency

2. Confidence interval of the effect estimate includes appreciable benefit, harm and no effect

3. High risk of selection and performance bias

N.5.2 Medroxyprogesterone compared to imaginal desensitisation for paraphilic disorders

			Quality as	sessment			№ of pa	tients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Medroxyprogesterone	imaginal desensitisation	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance
Reduced and	malous desire (fol	low up: 52; assessed	with: self-report ques	tionnaire)								
1	randomised trials	serious ¹	not serious	not serious	serious ^{2,3}	none	3/10 (30.0%)	6/10 (60.0%)	RR 0.50 (0.17 to 1.46)	300 fewer per 1,000 (from 276 more to 498 fewer)		CRITICAL
Reduced and	malous behaviour	(follow up: 52; asses	sed with: self-report q	uestionnaire)								
1	randomised trials	serious ¹	not serious	not serious	serious ^{2,3}	none	1/10 (10.0%)	3/10 (30.0%)	RR 0.33 (0.04 to 2.69)	201 fewer per 1,000 (from 288 fewer to 507 more)		CRITICAL

1. High risk of performance and attrition bias.

2. Optimal information size criterion not met (event rate less than 300)

3. Confidence interval for the effect estimate spans both MID thresholds (0.80 to 1.25).

N.5.3 Psychoeducational interventions, principally CBT-informed psychoeducation (including SOTP) versus treatment as usual, no treatment or waitlist control for paraphilic disorders.

			Quality ass	sessment			Nº of p	atients	Effect	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Cognitive dist	ortions (Abel and	Becker Cognition Scale,	ABCS; Children and	Sex Questionnaire) –	RCT (Scale from 26	to 130; higher better)						
1	randomised trials	serious 1	not serious	not serious	not serious	none	30	30	-	MD 13.43 lower (20.05 lower to 6.81 lower)		IMPORTANT
Cognitive dist	ortions (Abel and	Becker Cognition Scale,	ABCS; Children and	Sex Questionnaire) -	Controlled non-rando	mised studies						

			Quality ass	essment			№ of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
	observational studies	very serious 2	not serious	not serious	serious ³	none	38	19	-	MD 8.6 lower (14.48 lower to 2.72 lower)		IMPORTANT
ognitive dis	tortions (Abel and	Becker Cognition Scale	[ABCS]; number of pa	articipants who 'impro	ved' [pre- to post-test	t score increased by at least 10 pc	pints]) - Controlled non-ra	ndomised studies				•
	observational studies	very serious 4	not serious	not serious	very serious 5	none	4/19 (21.1%)	2/5 (40.0%)	RR 0.53 (0.13 to 2.10)	188 fewer per 1,000 (from 348 fewer to 440 more)		IMPORTANT
Cognitive dis	tortions (Multiphas	ic Sex Inventory [MSI]: (Cognitive distortions; r	number of participant	s who 'improved' [pre	- to post-test score increased by a	t least 2 points]) - Contro	olled non-randomised stu	dies			·
	observational studies	very serious 4	not serious	not serious	very serious 5	none	6/19 (31.6%)	1/5 (20.0%)	RR 1.58 (0.24 to 10.28)	116 more per 1,000 (from 152 fewer to 1,000 more)		IMPORTANT
motional co	ngruence with child	dren (Children and Sex	Questionnaire) - Cont	rolled non-randomise	d studies							
	observational studies	very serious ²	not serious	not serious	serious ⁶	none	38	19	-	MD 1.29 lower (8.8 lower to 6.22 higher)		IMPORTANT
ictim empat	hy distortions (Vict	im Empathy Distortions	scale) - Controlled no	n-randomised studie	S	•						
	observational studies	very serious ²	not serious	not serious	serious 7	none	38	19	-	MD 13 lower (21.56 lower to 4.44 lower)		IMPORTANT
cceptance of	of accountability (N	Iultiphasic Sex Inventory	y [MSI]: Justifications)	- RCT			ł			,		1
	randomised trials	serious 1	not serious	not serious	serious ⁸	none	30	30	-	MD 0.8 lower (6.13 lower to 4.53 higher)		IMPORTANT
cceptance of	of accountability (N	Iultiphasic Sex Inventory	y [MSI]: Justifications;	number of participan	ts who 'improved' [pre	e- to post-test score increased by	at least 2 points]) - Contr	olled non-randomised st	udies			
	observational studies	very serious 4	not serious	not serious	very serious 5	none	6/19 (31.6%)	2/5 (40.0%)	RR 0.79 (0.22 to 2.79)	84 fewer per 1,000 (from 312 fewer to 716		IMPORTANT

			Quality ass	sessment			Nº of p	atients	Effec	:t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
	observational studies	very serious ⁴	not serious	not serious	very serious 5	none	3/19 (15.8%)	0/5 (0.0%)	RR 2.10 (0.13 to 35.20)	0 fewer per 1,000 (from 0 fewer to 0 fewer)		IMPORTANT
enness/ho	nesty about sexua	l outlets (Multiphasic Se	ex Inventory [MSI]: Ch	ild molest; number of	participants who 'imp	proved' [pre- to post-test score inc	reased by at least 2 point	ts]) - Controlled non-rand	lomised studies			
	observational studies	very serious ⁴	not serious	not serious	very serious ⁵	none	7/19 (36.8%)	2/5 (40.0%)	RR 0.92 (0.27 to 3.13)	32 fewer per 1,000 (from 292 fewer to 852 more)		IMPORTANT
exual anxie	ty (Multiphasic Se	Inventory [MSI]: Sexua	I inadequacies) - RCI	г								
	randomised trials	serious ¹	not serious	not serious	serious 9	none	30	30	-	MD 6.2 lower (13.43 lower to 1.06 higher)		IMPORTANT
ixiety (Soc	al Anxiety and Dis	tress Scale, SADS) – R	CT (Scale from 0 to 28	8; lower better)	•		ł			· · ·		_ b
	randomised trials	serious ^{1,10}	not serious	serious 11	not serious	none	38	37	-	MD 2.19 lower (7.31 lower to 2.92 higher)		CRITICAL
arrest (CJ	S database; contro	lled non-randomised st	udies; longest follow-u	up available) - 2-year	follow-up					<u> </u>		
	randomised trials	very serious 12,13	not serious	not serious	very serious 5	none	38/197 (19.3%)	72/367 (19.6%)	RR 1.00 (0.63 to 1.59)	0 fewer per 1,000 (from 73 fewer to 116 more)		CRITICAL
arrest (CJ	S database; contro	lled non-randomised stu	udies; longest follow-u	ıp available) - 3-year	follow-up	1	1	L1		,		
	observational studies	very serious 14,15	not serious	serious ¹⁶	serious ⁵	none	436/1317 (33.1%)	1000/2118 (47.2%)	RR 0.78 (0.71 to 0.86)	104 fewer per 1,000 (from 66 fewer to 137 fewer)		CRITICAL
-	earrest (CJS data	base; controlled non-rar	idomised studies; long	gest follow-up availab	le) - 2-year follow-up		L	I		1 I		
ex offence			not serious	not serious	very serious 5	none	17/197 (8.6%)	26/367 (7.1%)	RR 1.03	2 more per	000	CRITICAL

			Quality ass	essment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
2	observational studies	very serious 14,15	not serious	serious	serious ⁵	none	105/1317 (8.0%)	199/2118 (9.4%)	RR 0.80 (0.57 to 1.12)	19 fewer per 1,000 (from 11 more to 40 fewer)		CRITICAL
Violent rearre	est (CJS database;	controlled non-randomi	sed studies; longest fo	ollow-up available) - 2	2-year follow-up							
1	observational studies	very serious 12	not serious	not serious	very serious ⁵	none	1/119 (0.8%)	5/159 (3.1%)	RR 0.27 (0.03 to 2.26)	23 fewer per 1,000 (from 31 fewer to 40 more)		CRITICAL
Violent rearre	est (CJS database;	controlled non-randomi	sed studies; longest fo	ollow-up available) - 3	-year follow-up					· · · · ·		
2	observational studies	very serious 14,15	not serious	serious ¹⁶	serious ⁵	none	197/1317 (15.0%)	485/2118 (22.9%)	RR 0.71 (0.60 to 0.83)	66 fewer per 1,000 (from 39 fewer to 92 fewer)		CRITICAL
Criminal char	ges (CJS database	e; controlled non-randon	nised studies; longest	follow-up available) -	2-year follow-up		<u>.</u>	<u>.</u>				
1	observational studies	very serious 17	not serious	serious ¹⁶	very serious 5	none	2/54 (3.7%)	1/14 (7.1%)	RR 0.52 (0.05 to 5.32)	34 fewer per 1,000 (from 68 fewer to 309 more)		CRITICAL
Sex offence of	charges (CJS datal	base; controlled non-ran	domised studies; long	gest follow-up availab	le) - 2-year follow-up							
1	observational studies	very serious 17	not serious	serious ¹⁶	very serious 5	none	0/54 (0.0%)	1/14 (7.1%)	RR 0.09 (0.00 to 2.12)	65 fewer per 1,000 (from to 80 more)		CRITICAL
Reconviction	(CJS database; co	ontrolled non-randomise	d studies; longest follo	ow-up available) - 2-y	ear follow-up		•	•	•			•
3	observational studies	very serious 13,18,19	very serious 20	not serious	very serious 5	none	37/243 (15.2%)	247/493 (50.1%)	RR 0.54 (0.16 to 1.82)	230 fewer per 1,000 (from 411 more to 421 fewer)		CRITICAL
Reconviction	(CJS database; co	ontrolled non-randomise	d studies; longest follo	ow-up available) - 3-y	ear follow-up							
1	observational studies	very serious 21	not serious	not serious	serious ⁵	none	4/94 (4.3%)	11/86 (12.8%)	RR 0.33 (0.11 to 1.01)	86 fewer per 1,000 (from 1 more to 114 fewer)		CRITICAL

			Quality ass	essment			Nº of p	atients	Effec	t l		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
econviction	(CJS database; co	ontrolled non-randomise	d studies; longest folle	ow-up available) - 4-y	ear follow-up							
	observational studies	very serious 22	not serious	serious ¹⁶	not serious	none	3/49 (6.1%)	23/74 (31.1%)	RR 0.20 (0.06 to 0.62)	249 fewer per 1,000 (from 118 fewer to 292 fewer)		CRITICAL
econviction	(CJS database; co	ontrolled non-randomise	d studies; longest follo	ow-up available) - 5-y	ear follow-up							
	observational studies	very serious 23,24,25	serious 26	serious ¹⁶	not serious	none	81/549 (14.8%)	116/484 (24.0%)	RR 0.53 (0.30 to 0.92)	113 fewer per 1,000 (from 19 fewer to 168 fewer)		CRITICAL
leconviction	(CJS database; co	ontrolled non-randomise	d studies; longest follo	ow-up available) - 7-y	ear follow-up	•	•					-
	observational studies	very serious 27	not serious	serious ¹⁶	not serious	none	199/403 (49.4%)	160/321 (49.8%)	RR 0.99 (0.85 to 1.15)	5 fewer per 1,000 (from 75 fewer to 75 more)		CRITICAL
econviction	at 2-year follow-up	o (risk of reconviction su	l b-analyses) - Low risk	(Į	I	ļ	II		ĮĮ		
	randomised trials	very serious 18	not serious	serious ¹⁶	not serious	none	15/263 (5.7%)	65/969 (6.7%)	RR 0.85 (0.49 to 1.47)	10 fewer per 1,000 (from 32 more to 34 fewer)		CRITICAL
leconviction	at 2-year follow-up	o (risk of reconviction su	b-analyses) - Medium	I-low risk		L				1		1
	randomised trials	very serious 18	not serious	serious ¹⁶	not serious	none	30/225 (13.3%)	166/655 (25.3%)	RR 0.53 (0.37 to 0.75)	119 fewer per 1,000 (from 63 fewer to 160 fewer)		CRITICAL
Reconviction	at 2-year follow-up	o (risk of reconviction su	b-analyses) - Medium	i-high risk	•		•			, ,		,
	observational studies	very serious 18	not serious	serious ¹⁶	not serious	none	23/109 (21.1%)	229/229 (100.0%)	RR 0.21 (0.15 to 0.31)	790 fewer per 1,000 (from 690 fewer to 850 fewer)		CRITICAL

			Quality ass	essment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious 18	not serious	serious ¹⁶	very serious 5	none	18/50 (36.0%)	22/57 (38.6%)	RR 0.93 (0.57 to 1.53)	27 fewer per 1,000 (from 166 fewer to 205 more)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 2-year follow-up							
2	observational studies	very serious 18,19	not serious	not serious	very serious 5	none	20/703 (2.8%)	55/1966 (2.8%)	RR 0.99 (0.59 to 1.68)	0 fewer per 1,000 (from 11 fewer to 19 more)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 3-year follow-up		•					
1	observational studies	very serious 21	not serious	not serious	very serious 5	none	1/94 (1.1%)	4/86 (4.7%)	RR 0.23 (0.03 to 2.01)	36 fewer per 1,000 (from 45 fewer to 47 more)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 4-year follow-up	1	ł			1		-1
2	observational studies	very serious 22	not serious	not serious	very serious 5	none	5/93 (5.4%)	17/118 (14.4%)	RR 0.42 (0.13 to 1.34)	84 fewer per 1,000 (from 49 more to 125 fewer)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 5-year follow-up							1
3	observational studies	very serious 24,25,28	not serious	serious ¹⁶	serious ⁵	none	23/276 (8.3%)	48/241 (19.9%)	RR 0.37 (0.16 to 0.83)	125 fewer per 1,000 (from 34 fewer to 167 fewer)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 7-year follow-up							
1	observational studies	very serious 27	not serious	serious ¹⁶	very serious 5	none	61/403 (15.1%)	46/321 (14.3%)	RR 1.06 (0.74 to 1.50)	9 more per 1,000 (from 37 fewer to 72 more)		CRITICAL
Sexual reconv	viction (CJS datab	ase; controlled non-rand	domised studies; long	est follow-up available	e) - 11-year follow-up							
1	observational studies	serious 29	not serious	serious ¹⁶	serious ⁵	none	66/616 (10.7%)	21/104 (20.2%)	RR 0.53 (0.34 to 0.83)	95 fewer per 1,000 (from 34 fewer to 133 fewer)		CRITICAL

			Quality ass	essment			№ of p	atients	Effe	ct		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Sexual recon	viction (CJS datab	ase; controlled non-ran	domised studies; long	est follow-up available	e) - Length of follow-u	up not reported						
1	observational studies	very serious 30	not serious	not serious	very serious 5	none	12/95 (12.6%)	17/145 (11.7%)	RR 1.08 (0.54 to 2.15)	9 more per 1,000 (from 54 fewer to 135 more)		CRITICAL
Sexual recon	viction at 2 or 11-y	ear follow-up (risk of ree	conviction sub-analyse	es) - Low risk	•	•	•	••				-
2	observational studies	very serious 18,29	not serious	serious ¹⁶	very serious 5	none	12/511 (2.3%)	14/994 (1.4%)	RR 0.68 (0.26 to 1.78)	5 fewer per 1,000 (from 10 fewer to 11 more)		CRITICAL
Sexual recon	viction at 2 or 11-y	ear follow-up (risk of re	conviction sub-analyse	es) - Medium-low risk						· · ·		
2	observational studies	very serious 18,29	not serious	serious ¹⁶	very serious 5	none	25/393 (6.4%)	25/683 (3.7%)	RR 0.71 (0.23 to 2.16)	11 fewer per 1,000 (from 28 fewer to 42 more)		CRITICAL
Sexual recon	viction at 2 or 11-y	ear follow-up (risk of re	conviction sub-analyse	es) - Medium-high ris	k			L				1
2	observational studies	very serious 18,29	not serious	serious ¹⁶	very serious 5	none	27/253 (10.7%)	19/260 (7.3%)	RR 0.67 (0.36 to 1.28)	24 fewer per 1,000 (from 20 more to 47 fewer)		CRITICAL
Sexual recon	viction at 2 or 11-y	ear follow-up (risk of re	conviction sub-analyse	es) - High risk		•		L1				1
2	observational studies	very serious 18,29	serious ²⁶	serious ¹⁶	very serious 5	none	19/106 (17.9%)	17/77 (22.1%)	RR 0.68 (0.26 to 1.76)	71 fewer per 1,000 (from 163 fewer to 168 more)		CRITICAL
Violent recon	viction (CJS datab	ase; controlled non-ran	domised studies; long	est follow-up available	e) - 3-year follow-up			· · · · · · ·				,
I	observational studies	very serious 21	not serious	not serious	serious 5	none	1/94 (1.1%)	7/86 (8.1%)	RR 0.13 (0.02 to 1.04)	71 fewer per 1,000 (from 3 more to 80 fewer)		CRITICAL

			Quality ass	sessment			Nº of p	atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
	observational studies	very serious 24,25	not serious	serious ¹⁶	not serious	none	16/176 (9.1%)	32/141 (22.7%)	RR 0.43 (0.25 to 0.74)	129 fewer per 1,000 (from 59 fewer to 170 fewer)		CRITICAL
olent recon	viction (CJS datab	ase; controlled non-ran	domised studies; long	est follow-up available	e) - 7-year follow-up							
	observational studies	very serious 27	not serious	serious ¹⁶	serious ⁵	none	136/403 (33.7%)	99/321 (30.8%)	RR 1.09 (0.88 to 1.35)	28 more per 1,000 (from 37 fewer to 108 more)		CRITICAL
olent recon	viction (CJS datab	ase; controlled non-ran	domised studies; long	est follow-up available	e) - Length of follow-u	up not reported						•
	observational studies	very serious 30	not serious	not serious	very serious 5	none	11/95 (11.6%)	24/145 (16.6%)	RR 0.70 (0.36 to 1.36)	50 fewer per 1,000 (from 60 more to 106 fewer)		CRITICAL
olent recon	viction (CJS datab	ase; controlled non-ran	domised studies; long	est follow-up available	e) - 11-year follow-up	1	4	<u> </u>		Į		_ I
	observational studies	very serious 29	not serious	serious ¹⁶	not serious	none	163/616 (26.5%)	46/104 (44.2%)	RR 0.60 (0.46 to 0.77)	177 fewer per 1,000 (from 102 fewer to 239 fewer)		CRITICAL
olent recon	viction at 11-year f	ollow-up (risk of reconv	iction sub-analyses) -	Low risk		•						•
	observational studies	very serious 29	not serious	serious ¹⁶	serious ⁵	none	28/248 (11.3%)	6/25 (24.0%)	RR 0.47 (0.22 to 1.03)	127 fewer per 1,000 (from 7 more to 187 fewer)		CRITICAL
olent recon	viction at 11-year t	ollow-up (risk of reconv	iction sub-analyses) -	Medium-low risk	1	1	1	II				-1
	observational studies	very serious 29	not serious	serious ¹⁶	very serious 5	none	56/168 (33.3%)	11/28 (39.3%)	RR 0.85 (0.51 to 1.41)	59 fewer per 1,000 (from 161 more to 193 fewer)		CRITICAL
iolent recon	viction at 11-year t	follow-up (risk of reconv	iction sub-analyses) -	Medium-high risk			I	I		<u> </u>		
	observational studies	very serious 29	not serious	serious ¹⁶	serious ⁵	none	53/144 (36.8%)	16/31 (51.6%)	RR 0.71 (0.48 to 1.07)	150 fewer per 1,000 (from 36 more to 268 fewer)		CRITICAL

			Quality ass	essment			Nº of p	patients	Effec	st		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
iolent recon	viction at 11-year f	ollow-up (risk of reconvi	ction sub-analyses) -	High risk						·		
I	observational studies	very serious 15	not serious	serious ¹⁶	not serious	none	26/56 (46.4%)	13/20 (65.0%)	RR 0.71 (0.47 to 1.10)	189 fewer per 1,000 (from 65 more to 345 fewer)		CRITICAL
ncarceration	(CJS database; co	ontrolled non-randomise	d studies; longest foll	ow-up available) - 3-y	ear follow-up	•		••		• • •		- ·
I	observational studies	very serious ¹⁵	not serious	serious ¹⁶	not serious	none	35/297 (11.8%)	228/1098 (20.8%)	RR 0.57 (0.41 to 0.79)	89 fewer per 1,000 (from 44 fewer to 123 fewer)		CRITICAL
ncarceration	for sexual offence	(CJS database; controll	ed non-randomised s	tudies; longest follow	-up available) - 3-yea	r follow-up				·		
I	observational studies	very serious 15	not serious	serious ¹⁶	very serious 5	none	9/297 (3.0%)	42/1098 (3.8%)	RR 0.79 (0.39 to 1.61)	8 fewer per 1,000 (from 23 fewer to 23 more)		CRITICAL
ncarceration	for violent offence	(CJS database; controll	ed non-randomised s	tudies; longest follow-	-up available) - 3-yea	r follow-up	L			1		
	observational studies	very serious 15	not serious	serious ¹⁶	serious ⁵	none	9/297 (3.0%)	74/1098 (6.7%)	RR 0.45 (0.23 to 0.89)	37 fewer per 1,000 (from 7 fewer to 52 fewer)		CRITICAL
Revocation, b	preaches of the Se	x Offender Register or p	robation violation (CJ	S database; controlle	d non-randomised st	udies; longest follow-up available)	- 2-year follow-up			1 1		
2	observational studies	very serious 13,17	very serious 20	not serious	very serious 5	none	31/132 (23.5%)	31/222 (14.0%)	RR 0.88 (0.12 to 6.74)	17 fewer per 1,000 (from 123 fewer to 802 more)		CRITICAL
Revocation, b	preaches of the Se	x Offender Register or p	robation violation (CJ	S database; controlle	d non-randomised st	ı udies; longest follow-up available)	- 5-year follow-up	ا ــــــــــــــــــــــــــــــــــــ				
	observational studies	very serious 15,24	not serious	serious ¹⁶	not serious	none	66/231 (28.6%)	643/1361 (47.2%)	RR 0.64 (0.51 to 0.80)	170 fewer per 1,000 (from 94 fewer to 231 fewer)		CRITICAL
evocation, b	preaches of the Se	x Offender Register or p	robation violation (CJ	S database; controlle	d non-randomised st	udies; longest follow-up available)	- Length of follow-up no	t reported				

			Quality ass	essment			№ of p	atients	Effect			
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious 30	not serious	not serious	not serious	none	7/95 (7.4%)	35/145 (24.1%)	RR 0.31 (0.14 to 0.66)	167 fewer per 1,000 (from 82 fewer to 208 fewer)		CRITICAL
Global treatm	ent response: Any	change (positively rated	for improvements in	denial, positive chang	ges on scales and atte	endance at therapy, and negative	ly rated for reconvictions	and breach of probation	order or parole licence)	- Controlled non-ra	ndomised studies	
1	observational studies	very serious ⁴	not serious	not serious	very serious ⁵	none	15/20 (75.0%)	3/5 (60.0%)	RR 1.25 (0.59 to 2.67)	150 more per 1,000 (from 246 fewer to 1,000 more)		IMPORTANT
Global treatm	ent response: Maj	or change (positively rate	ed for improvements i	n denial, positive cha	nges on scales and a	ttendance at therapy, and negativ	vely rated for reconviction	ns and breach of probation	on order or parole licenc	e) - Controlled non-	randomised studies	
1	observational studies	very serious 4	not serious	not serious	very serious 5	none	4/20 (20.0%)	0/5 (0.0%)	RR 2.57 (0.16 to 41.34)	0 fewer per 1,000 (from 0 fewer to 0 fewer)		IMPORTANT
Sub-analysis	by country: Recon	viction(Any)										
9	observational studies	very serious 13,18,19,21,22,23,24,25,27	serious ²⁶	serious ¹⁶	very serious 5	none	324/1338 (24.2%)	557/1458 (38.2%)	RR 0.49 (0.30 to 0.82)	195 fewer per 1,000 (from 69 fewer to 267 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction(Any) - UK										
1	observational studies	very serious 18	not serious	serious ¹⁶	not serious	none	23/109 (21.1%)	229/229 (100.0%)	RR 0.21 (0.15 to 0.31)	790 fewer per 1,000 (from 690 fewer to 850 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction(Any) - Netherland	ds									
1	observational studies	very serious 19	not serious	not serious	very serious ⁵	none	12/56 (21.4%)	14/56 (25.0%)	RR 0.86 (0.44 to 1.69)	35 fewer per 1,000 (from 140 fewer to 173 more)		CRITICAL
Sub-analysis	by country: Recon	viction(Any) - Spain										

			Quality ass	sessment			Nº of p	atients	Effec	:t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious 22	not serious	serious ¹⁶	not serious	none	3/49 (6.1%)	23/74 (31.1%)	RR 0.20 (0.06 to 0.62)	249 fewer per 1,000 (from 118 fewer to 292 fewer)		CRITICAL
Sub-analysis	by country: Recor	viction(Any) - US	•	•	•		•	•		•		-
4	observational studies	very serious 13,23,24,25	serious 26	serious ¹⁶	serious 5	none	83/627 (13.2%)	120/692 (17.3%)	RR 0.57 (0.34 to 0.96)	75 fewer per 1,000 (from 7 fewer to 114 fewer)		CRITICAL
Sub-analysis	by country: Recor	viction(Any) - Canada										
2	observational studies	very serious 21,27	serious 26	serious ¹⁶	very serious 5	none	203/497 (40.8%)	171/407 (42.0%)	RR 0.66 (0.23 to 1.88)	143 fewer per 1,000 (from 324 fewer to 370 more)		CRITICAL
Sub-analysis	by country: Sexua	I reconviction	Į	Į	Į	1	Į	<u> </u>		ĮĮ		_ ļ
11	observational studies	very serious 18,19,21,22,24,25,27,28,29,30,31	serious ²⁶	serious 16	serious ⁵	none	188/2280 (8.2%)	208/2981 (7.0%)	RR 0.66 (0.47 to 0.93)	24 fewer per 1,000 (from 5 fewer to 37 fewer)		CRITICAL
Sub-analysis	by country: Sexua	al reconviction - UK				•	I	l		1 1		
3	observational studies	very serious 18,30,31	not serious	not serious	very serious 5	none	32/786 (4.1%)	75/2099 (3.6%)	RR 0.96 (0.64 to 1.44)	1 fewer per 1,000 (from 13 fewer to 16 more)		CRITICAL
Sub-analysis	by country: Sexua	al reconviction - US										
3	observational studies	very serious ^{24,25,28}	not serious	serious 16	serious ⁵	none	23/276 (8.3%)	48/241 (19.9%)	RR 0.37 (0.16 to 0.83)	125 fewer per 1,000 (from 34 fewer to 167 fewer)		CRITICAL
Sub-analysis	by country: Sexua	I reconviction - Netherla	nds							· ·		
1	observational studies	very serious 19	not serious	not serious	very serious 5	none	3/56 (5.4%)	1/56 (1.8%)	RR 3.00 (0.32 to 27.97)	36 more per 1,000 (from 12 fewer to 482 more)		CRITICAL

Model Model Sub- syndes Rask of bias Instantian Informations Instantian Informations Other considerations (PR - DATE) (PR - DATE) (PR - DATE) Instantian Instantian (PR - DATE) Retail (PR - DATE) Absolute (PR - DATE) Obter (PR - DATE) Sub-arrian Sub-arrian Instantin Instantian				Quality ass	sessment			Nº of p	atients	Effec	t		
1 cbservaloral itudies very serious ¹² not serious serious ¹³ none 2/49 (4.1%) 13/74 (17.6%) RR 0.23 (0.5 fb 0.38) 135 fewer per 1.06 (0.5 fb 0.38) ····································		Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	intervention: principally CBT- informed psychoeducation	usual, no treatment			Quality	Importance
Budies Incl	ub-analysis	by country: Sexua	I reconviction - Spain										
a observational studies very serious *127.19 serious *1 not serious serious *1 none 128/1113 (11.5%) 71/511 (13.9%) RR 0.69 (0.36 to 1.33) 43 fewer per (1.00 (tom flower) b 89 fewer) Sub-analysis by country: Wolent reconviction 5 observational very serious *1 not serious serious *1 none 327/1384 (23.6%) 208/797 (26.1%) RR 0.69 (0.40 to 0.96) 99 fewer per 1.000 (from flower) vERY LOW Sub-analysis by country: Wolent reconviction not serious serious *1 none 327/1384 (23.6%) 208/797 (26.1%) RR 0.61 (0.40 to 0.96) 99 fewer per 1.000 (from flower) VERY LOW Sub-analysis by country: Wolent reconviction - UK observational studies very serious *1.2% not serious very serious *1 none 1195 (11.6%) 24/145 (16.6%) RR 0.70 (0.36 to 1.36) Serieur per 1.000 (from 60 more) VERY LOW Sub-analysis by country: Wolent reconviction - US not serious none 16/176 (9.1%) 32/141 (22.7%) RR 0.71 (0.25 to 0.74) 129 fewer per 1.000 (from 80 fewer) VERY LOW Sub-analysis by country: Wolent reconviction - Canada serious *1.2% none 16/176 (9.1%) 32/141 (22.7%) RR 0.71 (0.25 to 0.74)	1		very serious 22	not serious	serious ¹⁶	serious ⁵	none	2/49 (4.1%)	13/74 (17.6%)		1,000 (from 4 fewer		CRITICAL
$\frac{1}{100} \frac{1}{100} \frac{1}$	Sub-analysis	by country: Sexua	l reconviction - Canada		•	•	•	•			, , ,		•
6 beervational studies very serious ³⁰ not serious not serious serious ⁵ none 327/1384 (23.6%) 208/797 (26.1%) RR 0.62 (0.40 b 0.96) 99 fewer per from 10 fewer for 100 fewer in 107 fewer) Sub-analysis by country: Violent reconviction - UK 1 observational studies very serious ^{24.29} not serious not serious very serious ⁵ none 11/195 (11.6%) 24/145 (16.6%) RR 0.70 (0.36 to 1.36) 50 fewer per to 106 fewer 0 Sub-analysis by country: Violent reconviction - UK 1 observational studies very serious ^{24.29} not serious not serious none 11/195 (11.6%) 24/145 (16.6%) RR 0.70 (0.36 to 1.36) 50 fewer to 106 fewer Sub-analysis by country: Violent reconviction - US 2 observational studies very serious ^{24.29} not serious serious ¹⁰ not serious none 16/176 (8.1%) 32/141 (22.7%) RR 0.43 (0.25 to 0.74) 129 fewer per 1.000 (from 59 fewer 1.000 (from 59 fewer 1.000 (3		very serious 21,27,29	serious ²⁶	not serious	serious 16	none	128/1113 (11.5%)	71/511 (13.9%)		1,000 (from 46 more		CRITICAL
studies Image: Studi	Sub-analysis	by country: Violen	t reconviction					·			· · · ·		
Image: studies very serious ^{24,26} not serious not serious very serious ³ none 11/95 (11.6%) 24/145 (16.6%) RR 0.70 (0.36 b 1.36) 50 fewer per 1.000 (from 60 more to 16 lewer) Sub-analysis by country: Violent reconviction - US 2 observational very serious ^{24,26} not serious serious ¹⁶ none 16/176 (9.1%) 32/141 (22.7%) RR 0.43 (0.25 b 0.74) 129 fewer per 1.000 (from 69 more to 16 fewer) 2 observational very serious ^{24,26} not serious serious ¹⁶ not serious none 16/176 (9.1%) 32/141 (22.7%) RR 0.43 (0.25 b 0.74) 129 fewer per 1.000 (from 59 fewer to 170 fewer) VERY LOW Sub-analysis by country: Violent reconviction - Canada serious ¹⁶ not serious ⁶ none 300/1113 (27.0%) 152/511 (29.7%) RR 0.71 (0.39 b 0.131) 86 fewer per 1.000 (from 92 more to 170 fower) VERY LOW Sub-analysis by country: Violent reconviction - Canada serious ¹⁶ very serious ⁶ none 300/1113 (27.0%) 152/511 (29.7%) RR 0.71 (0.39 b 0.131) 86 fewer per 1.000 (from 92 more to 181 fewer) VERY LOW Sub-analysis by country: Recoration serious ¹⁶ very serious ⁶ none 104/458 (22.7%) 709/1728 (41.0%) RR 0.	i		very serious 30	not serious	not serious	serious 5	none	327/1384 (23.6%)	208/797 (26.1%)		1,000 (from 10 fewer		CRITICAL
studies i </td <td>Sub-analysis</td> <td>by country: Violen</td> <td>t reconviction - UK</td> <td></td> <td></td> <td></td> <td>•</td> <td>•</td> <td>L</td> <td></td> <td></td> <td></td> <td></td>	Sub-analysis	by country: Violen	t reconviction - UK				•	•	L				
Productional studies very serious ^{24,25} not serious serious ¹⁶ not serious none 16/176 (9.1%) $32/141 (22.7%)$ RR 0.43 (0.25 to 0.74) 129 fewer per (0.025 to 0.74) $\Theta \odot \odot \odot \odot \nabla \Theta$ Sub-analysis by country: Violent reconviction - Canada serious ^{21,27,29} very serious ²⁰ serious ¹⁶ very serious ⁵ none $300/113 (27.0%)$ $152/511 (29.7%)$ RR 0.71 (0.39 to 1.31) 86 fewer per 1,000 (from 92 more to 181 fewer) $\Theta \odot \odot \odot \nabla \Theta$ Sub-analysis by country: Revocation very serious ^{21,27,29} very serious ²⁰ serious ¹⁶ very serious ⁵ none $300/113 (27.0%)$ $152/511 (29.7%)$ RR 0.71 (0.39 to 1.31) 86 fewer per 1,000 (from 92 more to 181 fewer) $\Theta \odot \odot \odot \odot \nabla \Theta$ Sub-analysis by country: Revocation very serious 10.15, 17, 24.30 serious ²⁰ serious ⁵ none $104/458 (22.7\%)$ $709/1728 (41.0\%)$ RR 0.66 (0.35 to 1.23) 140 fewer per 1,000 $\Theta \odot \odot \odot \odot$			very serious ^{24,25}	not serious	not serious	very serious 5	none	11/95 (11.6%)	24/145 (16.6%)		1,000 (from 60 more		CRITICAL
studies i </td <td>Sub-analysis</td> <td>by country: Violen</td> <td>t reconviction - US</td> <td></td> <td></td> <td></td> <td>•</td> <td>•</td> <td>L</td> <td></td> <td></td> <td></td> <td></td>	Sub-analysis	by country: Violen	t reconviction - US				•	•	L				
$\frac{1}{3} \frac{1}{3} \frac{1}$	2		very serious 24,25	not serious	serious 16	not serious	none	16/176 (9.1%)	32/141 (22.7%)		1,000 (from 59 fewer		CRITICAL
studies indication	Sub-analysis	by country: Violen	t reconviction - Canada										
observational studies very serious 13,15,17.24,30 serious 20 serious 16 serious 5 none 104/458 (22.7%) 709/1728 (41.0%) RR 0.66 (0.35 to 1.23) 140 fewer per 1,000	3		very serious ^{21,27,29}	very serious 20	serious ¹⁶	very serious ⁵	none	300/1113 (27.0%)	152/511 (29.7%)		1,000 (from 92 more		CRITICAL
studies 13.15.17.24.30 (0.35 to 1.23) 1,000	Sub-analysis	by country: Revoo	ation										
to 267 fewer)	5			serious 20	serious 16	serious ⁵	none	104/458 (22.7%)	709/1728 (41.0%)		1,000 (from 94 more		CRITICAL

			Quality ass	essment			Nº of p	oatients	Effec	:t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychoeducational intervention: principally CBT- informed psychoeducation (including SOTP)	Treatment as usual, no treatment or waitlist control	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Sub-analysis	by country: Revoo	ation - UK										
1	observational studies	very serious 30	not serious	not serious	not serious	none	7/95 (7.4%)	35/145 (24.1%)	RR 0.31 (0.14 to 0.66)	167 fewer per 1,000 (from 82 fewer to 208 fewer)		CRITICAL
Sub-analysis	by country: Revoo	ation - US										
4	observational studies	very serious 13,15,17,24	very serious 20	serious ¹⁶	very serious 5	none	97/363 (26.7%)	674/1583 (42.6%)	RR 0.77 (0.39 to 1.55)	98 fewer per 1,000 (from 234 more to 260 fewer)		CRITICAL
Sub-analysis	by setting: Any rea	conviction	ł	I	4		ł	ĮI				-1
9	observational studies	very serious 13,18,19,21,22,24,25,27,32	very serious 20	serious ¹⁶	serious ⁵	none	349/1054 (33.1%)	582/1204 (48.3%)	RR 0.52 (0.33 to 0.80)	232 fewer per 1,000 (from 97 fewer to 324 fewer)		CRITICAL
Sub-analysis	by setting: Any re	conviction - Inpatient				1	<u> </u>	<u> </u>		, <u>,</u>		
1	observational studies	very serious 32	very serious 20	serious ¹⁶	serious ⁵	none	55/89 (61.8%)	66/89 (74.2%)	RR 0.83 (0.68 to 1.02)	126 fewer per 1,000 (from 15 more to 237 fewer)		CRITICAL
Sub-analysis	by setting: Any rea	conviction - Prison				•				<u> </u>		-
4	observational studies	very serious 18.21,22,25	serious 26	serious ¹⁶	serious ⁵	none	74/357 (20.7%)	315/479 (65.8%)	RR 0.33 (0.13 to 0.81)	441 fewer per 1,000 (from 125 fewer to 572 fewer)		CRITICAL
Sub-analysis	by setting: Any rea	conviction - Community								· ·		
4	observational studies	very serious 13,19,24,27	serious ²⁶	serious ¹⁶	very serious 5	none	220/608 (36.2%)	201/636 (31.6%)	RR 0.67 (0.32 to 1.40)	104 fewer per 1,000 (from 126 more to 215 fewer)		CRITICAL

CI: Confidence interval; MD: Mean difference; RR: Risk ratio; SOTP, sex offender treatment programme

Mental health of adults in contact with the criminal justice system Appendix N: Clinical evidence - GRADE evidence profiles

- 1. Anderson-Varney 1991 unclear risk of selection bias; no blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 2. O'Reilly 2010 Controlled Non-RCT; high risk of selection bias (significant difference in age between groups); No blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 3. The MID calculated from SD of control was +/-6.26.
- 4. Craissati 1997 Controlled Non-RCT; at baseline, men in the group condition was more likely to have abused children within the family; Increased loss of data in individual treatment programme (68%) than group treatment (38%); no selective outcome bias, no other risk of bias
- 5. The 95% CI considered for imprecision was 0.8 to 1.25.
- 6. The MID calculated from SD of control was +/-6.39.
- 7. The MID calculated from SD of control was +/-9.11.
- The MID calculated from SD of control was +/-5.41.
- 9. The MID calculated from SD of control was +/-6.01.
- 10. Hopkins 1991 Unclear selection bias; No blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias.
- 11. Hopkins 1991 Participants involved roughly equal numbers of incarcerated paedophile and rapists.
- 12. Song 1995 Controlled Non-RCT; significant group differences at baseline in race, prior sex offences and type of offence; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias;
- 13. Stalans 2001 Controlled Non-RCT; significant group differences at baseline in current offence and on prior criminal history; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 14. Duwe 2009 Controlled Non-RCT; high risk of selection bias; no blinding; low risk of attrition bias; high risk of selective outcome bias; low risk of other bias;
- 15. Lowden 2003 Controlled Non-RCT; significant group differences at baseline on age, marital status and criminal history; high risk of selection bias; no blinding; low risk of attrition bias; high risk of selective outcome bias; low risk of other bias
- 16. Unclear proportion of paraphilia participants
- 17. McGuire 2000 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 18. Friendship 2003 Controlled Non-RCT; confounders controlled in analysis; no blinding; unclear risk of attrition bias; high risk of selective outcome bias; low risk of other bias
- 19. Ruddijs 2000 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 20. 12>80%
- 21. Marshall 2008 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 22. Illescas 2008 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 23. Aytes 2001 Controlled Non-RCT; significant group differences at baseline in prior incarceration and prior felony conviction; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 24. McGrath 1998 Controlled Non-RCT; significant group differences at baseline in prior convictions; average time incarcerated and type of sexual offence committed; no blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 25. McGrath 2003 Controlled Non-RCT; significant group differences at baseline on prior convictions and time at risk in the community; no blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 26. 50%<12<80%
- 27. Hanson 2004 Controlled Non-RCT; higher proportion of prior sexual offences in intervention group compared with control group; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 28. Turner 2000 McGrath 1998 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 29. Olver 2013a Controlled Non-RCT; low risk of selection bias (profounders properly controlled); no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 30. Craissati 2009 Controlled Non-RCT; high risk of selection bias; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 31. Procter 1996 Controlled Non-RCT; no blinding; low risk of attrition bias; low risk of selective outcome bias; low risk of other bias
- 32. Looman 2000 Controlled Non-RCT; no blinding; unclear risk of attrition bias; low risk of selective outcome bias; unclear risk of other bias

N.5.4 Good Lives Model (GLM) versus Relapse Prevention (RP) for paraphilic disorders

			Quality as	sessment			Nº of p	atients	Effect	t			
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Good Lives Model (GLM)	Relapse Prevention (RP)	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance	
Cognitive dist	Cognitive distortions (Children and Sex Questionnaire) (Scale from 0 to 75; lower better) - Controlled non-randomised studies												
1	observational studies	very serious ¹	not serious	not serious	serious ²	none	207	294	-	MD 7.15 lower (9.06 lower to 5.25 lower)		IMPORTANT	
Emotional cor	ngruence with child	Iren (Children and Se	x Questionnaire) (Sca	le from 0 to 75; lower	better) - Controlled r	non-randomised studies							

			Quality as	sessment			№ of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Good Lives Model (GLM)	Relapse Prevention (RP)	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious 1	not serious	not serious	serious ³	none	207	294	-	MD 7.72 lower (10.13 lower to 5.3 lower)		IMPORTANT
Victim empat	hy distortions (Victi	m Empathy Distortion	ns scale) (Scale from (0 to 120; lower better)	- Controlled non-rand	domised studies						
1	observational studies	very serious 1	not serious	not serious	not serious ⁴	none	207	294	-	MD 0.44 higher (2.56 lower to 3.44 higher)		IMPORTANT
Treatment res studies	sponse for pro-offe	nding attitudes (using	g algorithm based on p	pre-post change and c	comparison with score	s of a non-offender on Beliefs abo	ut Children Scale [cognit	ive distortions and emoti	onal congruence with ch	nildren subscales] a	nd Victim Empathy Scale) - Co	ntrolled non-randomised
1	randomised trials	very serious 5	not serious	not serious	not serious	none	46/67 (68.7%)	366/520 (70.4%)	RR 0.98 (0.82 to 1.16)	14 fewer per 1,000 (from 113 more to 127 fewer)		CRITICAL
Drop-out (any	(cause) - Controlle	ed non-randomised st	udies									
1	observational studies	very serious 5	not serious	not serious	very serious 6	none	2/87 (2.3%)	2/182 (1.1%)	RR 2.09 (0.30 to 14.60)	12 more per 1,000 (from 8 fewer to 149 more)		IMPORTANT

CI: Confidence interval; MD: Mean difference; RR: Risk ratio

- 1. Barnett 2014 Controlled Non-RCT; no blinding; data on drop-out was not available for some outcomes; low risk of other bias.
- 2. The MID calculated from SD of control was +/-6.79.
- 3. The MID calculated from SD of control was +/-7.95.
- 4. The MID calculated from SD of control was +/-8.48.
- 5. Harkins 2012 Controlled Non-RCT; No blinding; data for individual scales were not reported; low other risk of bias.
- 6. The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.5 Reintegration programmes versus treatment as usual for paraphilic disorders

			Quality as	ssessment			Nº of p	atients	Effect	:		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Reintegration programme (COSA)	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance

			Quality a	ssessment			Nº of p	oatients	Effe	ct		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Reintegration programme (COSA)	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Rearrest at 2	-year follow-up (C.	IS database) - RCT								·		
1	randomised trials	very serious 1	not serious	serious ²	serious ³	none	12/31 (38.7%)	20/31 (64.5%)	RR 0.60 (0.36 to 1.00)	258 fewer per 1,000 (from 0 fewer to 413 fewer)		CRITICAL
Sex offence	rearrest at 2-year fo	ollow-up (CJS databa	ase) - RCT							·		
1	randomised trials	very serious 1	not serious	serious ²	very serious ³	none	0/31 (0.0%)	1/31 (3.2%)	RR 0.33 (0.01 to 7.88)	22 fewer per 1,000 (from 32 fewer to 222 more)		CRITICAL
Reconviction	at 2- to 4-year follo	ow-up (CJS database	e) - RCT (2-year follov	v-up)			•			• •		•
1	randomised trials	very serious 1	not serious	serious ²	serious ³	none	8/31 (25.8%)	14/31 (45.2%)	RR 0.57 (0.28 to 1.16)	194 fewer per 1,000 (from 72 more to 325 fewer)		CRITICAL
Reconviction	at 2- to 4-year follo	ow-up (CJS database	e) - Controlled non-rar	ndomised studies (3- d	or 4-year follow-up)		•					
3	observational studies	very serious 4	not serious	serious 5	serious ³	none	29/175 (16.6%)	57/175 (32.6%)	RR 0.52 (0.33 to 0.81)	156 fewer per 1,000 (from 62 fewer to 218 fewer)		CRITICAL
Sex offence	reconviction at 3- o	r 4-year follow-up (C	JS database) - Contro	olled non-randomised	studies							- 1
3	observational studies	very serious 4	not serious	serious ⁵	serious ³	none	8/175 (4.6%)	21/175 (12.0%)	RR 0.41 (0.18 to 0.94)	71 fewer per 1,000 (from 7 fewer to 98 fewer)		CRITICAL
Sex offence	reconviction (conta	ct) at 4-year follow-u	p (CJS database) - Co	ontrolled non-randomi	sed studies		•					•
1	observational studies	very serious 6	not serious	not serious	very serious 3	none	0/71 (0.0%)	3/71 (4.2%)	RR 0.14 (0.01 to 2.72)	36 fewer per 1,000 (from 42 fewer to 73 more)		CRITICAL
Violent recor	viction at 3- or 4-ye	ear follow-up (CJS da	atabase) - Controlled I	non-randomised studi	es	1		ıl		-I		-+
3	observational studies	very serious 4	not serious	serious ⁵	not serious	none	13/175 (7.4%)	43/175 (24.6%)	RR 0.34 (0.19 to 0.61)	162 fewer per 1,000 (from 96 fewer		CRITICAL

			Quality as	sessment			Nº of p	atients	Effec	:t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Reintegration programme (COSA)	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
	randomised trials	very serious 1	not serious	serious ²	very serious 3	none	3/31 (9.7%)	8/31 (25.8%)	RR 0.38 (0.11 to 1.28)	160 fewer per 1,000 (from 72 more to 230 fewer)		CRITICAL
ny reincarce	eration at 2-year fol	llow-up (CJS databas	se) - RCT									
1	randomised trials	very serious 1	not serious	serious ²	very serious ³	none	15/31 (48.4%)	19/31 (61.3%)	RR 0.79 (0.50 to 1.25)	129 fewer per 1,000 (from 153 more to 306 fewer)		CRITICAL
Reincarcerati	on for a technical v	violation revocation o	r failure to comply with	Sex Offender's Regis	ster (SOR) requireme	ents at 2- or 4-year follow-up (CJS	database) - RCT (reinca	ceration for revocation; 2	-year follow-up)			
1	randomised trials	very serious 1	not serious	serious ²	serious ³	none	13/27 (48.1%)	17/25 (68.0%)	RR 0.71 (0.44 to 1.14)	197 fewer per 1,000 (from 95 more to 381 fewer)		CRITICAL
Reincarcerati	on for a technical v	violation revocation o	r failure to comply with	Sex Offender's Regis	ster (SOR) requireme	nts at 2- or 4-year follow-up (CJS	database) - Controlled ne	on-randomised studies (fa	ailure to comly with SOF	R requirements; 4-yea	ar follow-up)	
1	observational studies	very serious 6	not serious	not serious	very serious ³	none	4/71 (5.6%)	6/71 (8.5%)	RR 0.67 (0.20 to 2.26)	28 fewer per 1,000 (from 68 fewer to 106 more)		CRITICAL
Sub-analysis	by country: Recon	viction (Any) - UK (co	ontrolled non-randomis	sed)								
1	observational studies	very serious 6	not serious	not serious	serious ³	none	7/71 (9.9%)	14/71 (19.7%)	RR 0.50 (0.21 to 1.16)	99 fewer per 1,000 (from 32 more to 156 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction (Any) - US (R	CT)	L	•							
	randomised trials	very serious 1	not serious	serious ²	serious ³	none	8/31 (25.8%)	14/31 (45.2%)	RR 0.57 (0.28 to 1.16)	194 fewer per 1,000 (from 72 more to 325 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction (Any) - Canac	a (controlled non-rand	lomised)	+			· · · · · · · · · · · · · · · · · · ·				•
	observational studies	very serious 7	serious ⁸	serious ⁵	serious ³	none	22/104 (21.2%)	43/104 (41.3%)	RR 0.48 (0.22 to 1.04)	215 fewer per 1,000 (from 17 more to 323 fewer)		CRITICAL
ub-analysis	by country: Recon	viction (sexual)					•			· · ·		•

			Quality as	ssessment			№ of p	atients	Effe	ct		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Reintegration programme (COSA)	Treatment as usual	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
	observational studies	very serious ⁴	not serious	serious ⁵	serious ³	none	8/175 (4.6%)	21/175 (12.0%)	RR 0.41 (0.18 to 0.94)	71 fewer per 1,000 (from 7 fewer to 98 fewer)		CRITICAL
ub-analysis	by country: Recon	viction (sexual) - UK										
	observational studies	very serious 6	not serious	not serious	very serious 3	none	4/71 (5.6%)	5/71 (7.0%)	RR 0.80 (0.22 to 2.86)	14 fewer per 1,000 (from 55 fewer to 131 more)		CRITICAL
Sub-analysis	by country: Recon	viction (sexual) - Car	nada									
2	observational studies	very serious 7	not serious	serious ⁵	not serious	none	4/104 (3.8%)	16/104 (15.4%)	RR 0.26 (0.09 to 0.75)	114 fewer per 1,000 (from 38 fewer to 140 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction (violent)										
3	observational studies	very serious 4	not serious	serious 5	not serious	none	13/175 (7.4%)	43/175 (24.6%)	RR 0.34 (0.19 to 0.61)	162 fewer per 1,000 (from 96 fewer to 199 fewer)		CRITICAL
Sub-analysis	by country: Recon	viction (violent) - UK								<u> </u>		_
	observational studies	very serious 6	not serious	not serious	serious ³	none	0/71 (0.0%)	7/71 (9.9%)	RR 0.07 (0.00 to 1.15)	92 fewer per 1,000 (from to 15 more)		CRITICAL
Sub-analysis	by country: Recon	viction (violent) - Car	nada	•	•					<u> </u>		
2	observational studies	very serious 7	not serious	serious ⁵	not serious	none	13/104 (12.5%)	36/104 (34.6%)	RR 0.37 (0.21 to 0.65)	218 fewer per 1,000 (from 121 fewer to 273 fewer)		CRITICAL
Sub-analysis	by country: Revoc	ation - UK (failure to	comply with SOR requ	uirements: 4-year follo	w-up; controlled non	-randomised studies)	<u> </u>	·		+ +		4
	observational studies	very serious 4	not serious	not serious	very serious ³	none	4/71 (5.6%)	6/71 (8.5%)	RR 0.67 (0.20 to 2.26)	28 fewer per 1,000 (from 68 fewer to 106 more)		CRITICAL

			Quality as	ssessment			Nº of p	atients	Effect	1		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Reintegration programme (COSA)	Treatment as usual	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
1	randomised trials	very serious 1	not serious	not serious	serious ³	none	13/27 (48.1%)	17/25 (68.0%)	RR 0.71 (0.44 to 1.14)	197 fewer per 1,000 (from 95 more to 381 fewer)		CRITICAL

- 1. Duwe 2013 high risk of selection bias (Prior sex crime conviction was 32% in intervention group compared with 10% in control group); No blinding; low attrition risks; low selective outcome bias; low risk of other bias.
- 2. 'Sex offender' unclear proportion of participants with a paraphilic disorder

3. The 95% CI considered for imprecision was 0.8 to 1.25.

- 4. Bates 2014 Controlled Non-RCT; high risk of selection bias; no blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias; Wilson 2007, Wilson 2009 Controlled Non-RCT; high risk of selection bias; significant differences in baseline risk factors between groups; no blinding; unclear attrition risk of bias; low risk of other bias; low risk of other bias
- 5. Proportion of participants with paraphilia was unclear (Wilson 2009); only over half (Wilson 2007); majority (86%) of sample (Bates 2014).
- 6. Bates 2014 Controlled Non-RCT; high risk of selection bias; no blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias;
- 7. Wilson 2007, Wilson 2009 Controlled Non-RCT; high risk of selection bias; significant differences in baseline risk factors between groups; no blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias

8. I2>50%.

N.5.6 Therapeutic communities versus no treatment for paraphilic disorders

			Quality as	ssessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Therapeutic communities	No treatment	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance
Rearrest (CJS	S database; contro	led non-randomised	studies; longest follow	v-up available)								
1	observational studies	serious ¹	not serious	not serious	not serious	none	41/119 (34.5%)	607/1098 (55.3%)	RR 0.62 (0.48 to 0.80)	210 fewer per 1,000 (from 111 fewer to 287 fewer)		CRITICAL
Rearrest (CJS	S database; contro	led non-randomised	studies; longest follow	- v-up available) - 3-yea	r follow-up	•				••		•
1	observational studies	serious ¹	not serious	not serious	not serious	none	41/119 (34.5%)	607/1098 (55.3%)	RR 0.62 (0.48 to 0.80)	210 fewer per 1,000 (from 111 fewer to 287 fewer)		CRITICAL
Sex offence r	earrest (CJS datab	ase; controlled non-r	andomised studies; lo	ngest follow-up availa	able)		l	L				
1	observational studies	serious 1	not serious	not serious	serious ²	none	8/119 (6.7%)	81/1098 (7.4%)	RR 0.91 (0.45 to 1.84)	7 fewer per 1,000 (from 41 fewer to 62 more)		CRITICAL

			Quality as	ssessment			Nº of p	patients	Effe	rt -		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Therapeutic communities	No treatment	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Sex offence	rearrest (CJS datab	base; controlled non-	randomised studies; lo	ongest follow-up avail	able) - 3-year follow-u	p						
1	observational studies	serious 1	not serious	not serious	serious ²	none	8/119 (6.7%)	81/1098 (7.4%)	RR 0.91 (0.45 to 1.84)	7 fewer per 1,000 (from 41 fewer to 62 more)		CRITICAL
Violent rearr	est (CJS database;	controlled non-rande	omised studies; longes	st follow-up available)			•			1 1		1
1	observational studies	serious 1	not serious	not serious	serious ²	none	26/119 (21.8%)	288/1098 (26.2%)	RR 0.83 (0.58 to 1.19)	45 fewer per 1,000 (from 50 more to 110 fewer)		CRITICAL
Violent rearr	est (CJS database;	controlled non-rande	omised studies; longes	st follow-up available)	- 3-year follow-up	•	•			• •		-
1	observational studies	serious 1	not serious	not serious	serious ²	none	26/119 (21.8%)	288/1098 (26.2%)	RR 0.83 (0.58 to 1.19)	45 fewer per 1,000 (from 50 more to 110 fewer)		CRITICAL
Incarceration	n (CJS database; co	ontrolled non-random	nised studies; longest f	ollow-up available)			•					
1	observational studies	serious 1	not serious	not serious	serious ²	none	12/119 (10.1%)	228/1098 (20.8%)	RR 0.49 (0.28 to 0.84)	106 fewer per 1,000 (from 33 fewer to 150 fewer)		CRITICAL
Incarceration	n (CJS database; co	ntrolled non-random	ised studies; longest f	ollow-up available) - 3	3-year follow-up			Į		4 I		-1
1	observational studies	serious ¹	not serious	not serious	serious ²	none	12/119 (10.1%)	228/1098 (20.8%)	RR 0.49 (0.28 to 0.84)	106 fewer per 1,000 (from 33 fewer to 150 fewer)		CRITICAL
Incarceration	n for sexual offence	(CJS database; con	trolled non-randomise	d studies; longest follo	ow-up available)		•					
1	observational studies	serious ¹	not serious	not serious	serious ²	none	6/119 (5.0%)	42/1098 (3.8%)	RR 1.32 (0.57 to 3.04)	12 more per 1,000 (from 16 fewer to 78 more)		CRITICAL
Incarceration	for sexual offence	(CJS database; con	trolled non-randomise	d studies; longest follo	ow-up available) - 3-y	ear follow-up	•			+ +		+
1	observational studies	serious ¹	not serious	not serious	serious ²	none	6/119 (5.0%)	42/1098 (3.8%)	RR 1.32 (0.57 to 3.04)	12 more per 1,000 (from 16 fewer to 78 more)		CRITICAL

			Quality as	sessment			Nº of p	oatients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Therapeutic communities	No treatment	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance
1	observational studies	serious ¹	not serious	not serious	serious ²	none	3/119 (2.5%)	74/1098 (6.7%)	RR 0.37 (0.12 to 1.17)	42 fewer per 1,000 (from 11 more to 59 fewer)		CRITICAL
Incarceration	for violent offence	(CJS database; contr	olled non-randomised	l studies; longest follo	w-up available) - 3-ye	ear follow-up						
1	observational studies	serious 1	not serious	not serious	serious ²	none	3/119 (2.5%)	74/1098 (6.7%)	RR 0.37 (0.12 to 1.17)	42 fewer per 1,000 (from 11 more to 59 fewer)		CRITICAL
Revocation (C	CJS database; con	trolled non-randomise	ed studies; longest foll	ow-up available)	•					•		•
1	observational studies	serious ¹	not serious	not serious	serious ²	none	18/115 (15.7%)	625/1310 (47.7%)	RR 0.33 (0.21 to 0.50)	320 fewer per 1,000 (from 239 fewer to 377 fewer)		CRITICAL
Revocation (0	CJS database; con	trolled non-randomise	ed studies; longest foll	ow-up available) - 5-y	ear follow-up							
1	observational studies	serious ¹	not serious	not serious	serious ²	none	18/115 (15.7%)	625/1310 (47.7%)	RR 0.33 (0.21 to 0.50)	320 fewer per 1,000 (from 239 fewer to 377 fewer)		CRITICAL

Lowden 2003 - Controlled Non-RCT; significant group differences at baseline on age, marital status and criminal history; high risk of selection bias; no blinding; low risk of attrition bias; high risk of selective outcome bias; low risk of other bias
 The 95% C.I. considered for imprecision was 0.80 to 1.25

Cognitive behavioural therapy (CBT) versus treatment as usual for paraphilic disorders N.5.7

			Quality as	ssessment			Nº of p	atients	Effect	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	СВТ	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Sexual recon	aual reconviction (CJS database; controlled non-randomised studies; longest follow-up available) - 4-year follow-up (exhibitionists)											
1	al reconviction (CJS database; controlled non-randomised studies; longest follow-up available) - 4-year follow-up (exhibitionists) observational studies very serious ¹ not serious not serious ² none						4/17 (23.5%)	12/21 (57.1%)	RR 0.41 (0.16 to 1.05)	337 fewer per 1,000 (from 29 more to 480 fewer)		CRITICAL

- Marshall 1988a/b/1991 Controlled Non-RCT with 4 and 9-year follow-up; No baseline risk differences; No blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias 1.
- 2. The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.8 Behavioural therapies versus treatment as usual for paraphilic disorders

			Quality as	ssessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Behavioural therapies	Treatment as usual	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
Sexual recon	viction (CJS datab	ase; controlled non-ra	ndomised studies; lor	ngest follow-up availa	ble) - 4-year follow-up			•				
1	observational studies	very serious 1	not serious	not serious	serious ²	none	6/24 (25.0%)	12/20 (60.0%)	RR 0.42 (0.19 to 0.91)	348 fewer per 1,000 (from 54 fewer to 486 fewer)		CRITICAL
Sexual recon	viction (CJS datab	ase; controlled non-ra	indomised studies; lor	ngest follow-up availa	ble) - 9-year follow-up	(exhibitionists)						
1	observational studies	very serious 1	not serious	not serious	very serious ²	none	9/23 (39.1%)	12/21 (57.1%)	RR 0.68 (0.36 to 1.29)	183 fewer per 1,000 (from 166 more to 366 fewer)		CRITICAL

CI: Confidence interval; RR: Risk ratio

Marshall 1988a/b/1991 - Controlled Non-RCT with 4 and 9-year follow-up; No baseline risk differences; No blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias
 The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.9 Imaginal desensitization plus MPA versus MPA for paraphilic disorders

			Quality as	sessment			Nº of p	atients	Effect	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Imaginal desensitization + medroxyprogesterone	Medroxyprogesterone only	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Number of pe	eople who had a re	eduction in anomalou	s behaviours (26 wee	ks follow-up)								
1	randomised trials	very serious 1	not serious	serious ²	very serious ³	none	9/10 (90.0%)	8/10 (80.0%)	RR 1.12 (0.78 to 1.63)	96 more per 1,000 (from 176 fewer to 504 more)		CRITICAL
Number of pe	eople who had a re	eduction in anomalou	s desires (26 weeks f	ollow-up)								

			Quality a	ssessment			Nº of p	atients	Effect	t		
Nº of studies			Other considerations	Imaginal desensitization + medroxyprogesterone	Medroxyprogesterone only	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance			
1	randomised trials	very serious ¹	not serious	serious ²	very serious ³	none	5/10 (50.0%)	3/10 (30.0%)	RR 1.67 (0.54 to 5.17)	201 more per 1,000 (from 138 fewer to 1,000 more)		CRITICAL

- McConaghy 1988 unclear risk of selection bias, no blinding, low risk of attrition bias, high risk of selective outcome bias, low risk of other bias. 1.
- Unclear what percentage are currently in contact with the criminal justice system
 The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.10 Imaginal desensitization versus covert sensitization for paraphilic disorders

			Quality asses	ssment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Imaginal desensitization versus Covert sensitization only (Inpatient)	Control	Relative (95% Cl)	Absolute		
Number o	of people who	had a rec	duction in anomal	lous behavio	urs							
	trials	serious ¹	no serious inconsistency		very serious ³	none	7/10 (70%)	4/10 (40%)	⊕OOO VERY LOW	CRITICAL		
			duction in anomal		lioni	2020	2/10	E/10		200 fower par 1000		CDITICAL
	randomised trials	1	no serious inconsistency		very serious ³	none	3/10 (30%)	5/10 (50%)	RR 0.6 (0.19 to 1.86)	200 fewer per 1000 (from 405 fewer to 430 more)	⊕OOO VERY LOW	CRITICAL

¹ McConaghy 1985 - unclear selection bias, no blinding, high risk of attrition bias, high risk of selective outcome bias, low other risk of bias,

² 13/20 had previously received convictions but unclear what percentage of the sample were currently in contact with the criminal justice system. Also 5 individuals requested treatment due to being homosexual, which would no longer be considered a paraphilia. ³ The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.11 Aversive conditioning and milieu therapy versus treatment as usual for paraphilic disorders

			Quality as	ssessment			Nº of p	patients	Effect	i		
№ of studies	Study design	lesign Risk of bias Inconsistency Indirectness		Imprecision	Other considerations	Aversive conditioning training and milieu therapy	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance	
Sexual and/o	r violent reconvictio	ons at 21-year follow-	up (CJS database) - 0	Controlled non-randon	nised studies							
1	observational studies	very serious 1	not serious	not serious	serious ²	none	47/106 (44.3%)	35/91 (38.5%)	RR 1.15 (0.82 to 1.61)	58 more per 1,000 (from 69 fewer to 235 more)		CRITICAL

CI: Confidence interval; RR: Risk ratio

- Hanson 1993 Controlled Non-RCT; significant baseline risk differences (+); no blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias.
 The 95% CI considered for imprecision was 0.8 to 1.25.

N.5.12 Psychotherapy versus no treatment or treatment as usual for paraphilic disorders

			Quality as	sessment			Nº of p	oatients	Effec	t					
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychotherapy	No treatment or treatment as usual	Relative (95% Cl)	Absolute (95% CI)	Quality	Importance			
Rearrest (sou	rce of data not rep	orted; controlled non-	randomised studies; I	ongest follow-up avail	able) - 2-year follow-	up						·			
1	observational studies very serious 1 not serious serious 2 not serious none 3/92 (3.3%) 20/75 (26.7%) RR 0.12 (0.04 to 0.40) 235 fewer per 1,000 (from 160 fewer) QCRITICAL x offence rearrest (source of data not reported; controlled non-randomised studies; longest follow-up available) - 2-year follow-up Description Description Description CRITICAL														
Sex offence r	earrest (source of c	data not reported; cor	trolled non-randomise	ed studies; longest fol	low-up available) - 2-	year follow-up		••		••		•			
1	observational studies	very serious 1	not serious	serious ²	serious ³	none	1/92 (1.1%)	6/75 (8.0%)	RR 0.14 (0.02 to 1.10)	69 fewer per 1,000 (from 8 more to 78 fewer)		CRITICAL			
Sexual reconv	viction (CJS databa	ase; controlled non-ra	indomised studies; lor	ngest follow-up availat	ble) - Length of follow	-up not reported									
1	observational studies	very serious 4	not serious	not serious	very serious ³	none	5/23 (21.7%)	17/145 (11.7%)	RR 1.85 (0.76 to 4.54)	100 more per 1,000 (from 28 fewer to 415 more)		CRITICAL			
iolent recon	viction (CJS databa	ase; controlled non-ra	ndomised studies; lor	ngest follow-up availat	ble) - Length of follow	-up not reported	ļ			· ·		ł			

			Quality as	ssessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Psychotherapy	No treatment or treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious ⁴	not serious	not serious	very serious ³	none	3/23 (13.0%)	24/145 (16.6%)	RR 0.79 (0.26 to 2.41)	35 fewer per 1,000 (from 122 fewer to 233 more)		CRITICAL
Breaches of t	he Sex Offender R	egister (CJS databas	e; controlled non-rand	domised studies; longe	est follow-up available	e) - Length of follow-up not reporte	d					
1	observational studies	very serious 4	not serious	not serious	very serious ³	none	8/23 (34.8%)	35/145 (24.1%)	RR 1.44 (0.77 to 2.70)	106 more per 1,000 (from 56 fewer to 410 more)		CRITICAL

- 1. Peters 1968 Controlled Non-RCT; group differences at baseline; no blinding; unclear attrition risk of bias; low risk of selective outcome bias; low risk of other bias.
- 2. 'Sex offender' unclear proportion of participants with a paraphilic disorder; also an unknown proportion of participants in the intervention group had treatment delivered in a psychiatric inpatient unit
- 3. The 95% CI considered for imprecision was 0.8 to 1.25.
- 4. Craissati 2009 Controlled Non-RCT; there might have selection bias issues such as unequal baseline risks between 2 groups and the individual psychoeducation group was also offered to to those who had already attempted group work; No blinding; only participants with available follow-up data were included; low risk of selective outcome bias; low risk of other bias

N.5.13 Polygraph testing versus treatment as usual for paraphilic disorders

			Quality as	sessment			Nº of p	atients	Effec	t		
№ of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Polygraph testing	Treatment as usual	Relative (95% Cl)	Absolute (95% Cl)	Quality	Importance
Reconviction	(CJS database; co	ntrolled non-randomi	sed studies; longest fo	ollow-up available) - 5	-year follow-up							
1	observational studies	very serious 1	not serious	not serious	serious ²	none	41/104 (39.4%)	36/104 (34.6%)	RR 1.14 (0.80 to 1.63)	48 more per 1,000 (from 69 fewer to 218 more)		CRITICAL
Sexual recon	viction (CJS databa	ase; controlled non-ra	andomised studies; lor	ngest follow-up availa	ble) - 5-year follow-up							
1	observational studies	very serious 1	not serious	not serious	serious ²	none	6/104 (5.8%)	7/104 (6.7%)	RR 0.86 (0.30 to 2.46)	9 fewer per 1,000 (from 47 fewer to 98 more)		CRITICAL
Violent recon	viction (CJS databa	ase; controlled non-ra	andomised studies; lor	ngest follow-up availa	ble) - 5-year follow-up							

			Quality as	ssessment			№ of p	atients	Effec	t		
Nº of studies	Study design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Polygraph testing	Treatment as usual	Relative (95% CI)	Absolute (95% Cl)	Quality	Importance
1	observational studies	very serious 1	not serious	not serious	serious ²	none	3/104 (2.9%)	12/104 (11.5%)	RR 0.25 (0.07 to 0.86)	87 fewer per 1,000 (from 16 fewer to 107 fewer)		CRITICAL
Incarceration	Incarceration (CJS database; controlled non-randomised studies; longest follow-up available) - 5-year follow-up											
1	observational studies	very serious 1	not serious	not serious	serious ²	none	49/104 (47.1%)	40/104 (38.5%)	RR 1.23 (0.89 to 1.68)	88 more per 1,000 (from 42 fewer to 262 more)		CRITICAL
Violation of su	upervision condition	ns (CJS database; co	ntrolled non-randomi	sed studies; longest fo	ollow-up available) - 5	-year follow-up						
1	observational studies	very serious 1	not serious	not serious	serious ²	none	54/104 (51.9%)	47/104 (45.2%)	RR 1.15 (0.87 to 1.52)	68 more per 1,000 (from 59 fewer to 235 more)		CRITICAL

- 1. McGrath 2007 Controlled Non-RCT; baseline characters were similar between the groups; no blinding; low risk of detection bias; low attrition bias; low selective outcome bias; low risk of other bias
- 2. The 95% CI considered for imprecision was 0.80 to 1.25.

N.6 Service delivery models

N.6.1 Street Triage (Before and After)

			Quality ass	essment			No of p	patients		Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Street triage	Control	Relative (95% CI)	Absolute			
Total s13	Total s136 detentions per 100,000												
	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	89/100000* (0.09%)	107/100000* (0.11%)	RR 0.83 (0.63 to 1.1)	18.2 fewer per 100,000 (from 39.6 fewer to 10.7 more)	⊕OOO VERY LOW	CRITICAL	
Number o	of s136 detention	ons in cus	stody	•	•	•		<u>.</u>		•			
	observational studies	serious ³	no serious inconsistency	no serious indirectness	no serious imprecision	strong association ⁴	6085/24687 (24.6%)		RR 0.68 (0.67 to 0.7)	115 fewer per 1000 (from 108 fewer to 119 fewer)	⊕⊕OO LOW	CRITICAL	

Number o	of s136 detentio	ons in hos	spital									
3	observational	serious⁵	no serious	no serious	no serious	none	18613/24703	16139/25250	RR 1.18	115 more per 1000	⊕000	CRITICAL
	studies		inconsistency	indirectness	imprecision		(75.3%)	(63.9%)	(1.16 to	(from 102 more to 121	VERY	
									1.19)	more)	LOW	

¹ Reveruzzi 2016 – before and after study; low risk of selection bias as the groups were formed by before and after implementation of street triage; high risk of performance bias as there was no blinding involved; high rate of missing data and complete case analysis

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ Hywel Dda 2015 – before and after study; low risk of selection bias as the groups were formed by before and after implementation of street triage; high risk of performance bias as there was no blinding involved; high rate of missing data and complete case analysis.

⁴ Evidence was upgraded if the effect estimate was considered to be large(I.e. 95% CI of RR <0.75 or RR>1.25).

⁵ Powys 2015 – before and after study; low risk of selection bias as the groups were formed by before and after implementation of street triage; high risk of performance bias as there was no blinding involved; high rate of missing data and complete case analysis

*The total population being looked at was not provided and the data was calculated per 100,000.

N.6.2 Diversion services

N.6.2.1 Before and After Diversion services

			Quality asse	essment		No of patients			Quality	Importance		
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Court Diversion	Control	Relative (95% CI)	Absolute		
Duration b	between remand	and asses	sment (days) (Bett	er indicated by lo	wer values)		•					
2	observational studies	Serious ^{1,2}		no serious indirectness	no serious imprecision	none	294	317	-	MD 31.76 lower (69.55 lower to 6.03 higher)	⊕OOO VERY LOW	CRITICAL
Days of to	tal time on rema	and (Better	indicated by lower	values)		•						
1	observational studies	Serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	280	285	-	MD 17.6 lower (28.64 to 6.56 lower)	⊕OOO VERY LOW	IMPORTANT
Proportion	ns of prisoners of	on bail	-	•	•		•	•	· · · · · · ·			
1	observational studies	Serious ³	no serious inconsistency	no serious indirectness	very serious ⁴	none	31/122 (25.4%)	20/98 (20.4%)	RR 1.25 (0.76 to 2.04)	51 more per 1000 (from 49 fewer to 212 more)	⊕OOO VERY LOW	CRITICAL
Attendanc	e at alcohol and	drug treat	ment programmes									
1	observational studies	Serious ³	no serious inconsistency	no serious indirectness	very serious ⁴	none	13/41 (31.7%)	9/29 (31%)	RR 1.02 (0.51 to 2.07)	6 more per 1000 (from 152 fewer to 332 more)	⊕OOO VERY LOW	CRITICAL

OPD atten	PD attendance rate for those release on bail													
	observational studies	Serious ³		no serious indirectness	serious ⁴	none	11/23 (47.8%)	7/13 (53.8%)	```	59 fewer per 1000 (from 291 fewer to 388 more)	⊕OOO VERY LOW	CRITICAL		
Registratio	on of care progr	ammes (Cl	PA) and supervisio	n registration (SF	R)									
	observational studies	Serious ³		no serious indirectness	very serious ⁴	none	10/122 (8.2%)	4/98 (4.1%)	RR 2.01 (0.65 to 6.21)	41 more per 1000 (from 14 fewer to 213 more)	⊕OOO VERY LOW	CRITICAL		

¹ Exworthy 1997- before and after study with no confounder being controlled; no blinding; unclear drop out and available case analysis ² Weaver 1997 – before and after study with no confounder being controlled; no blinding; unclear dropout with available case analysis ³ Chambers 1999 – controlled cohort study with no confounder being controlled; no blinding; unclear drop out and available case analysis

⁴ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

Court diversion vs Community diversion services N.6.2.2

	Quality assessment									Effect	Quality	Importance	
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Court	Community Diversion	Relative (95% CI)	Absolute			
Rate of re	Rate of re-incarceration in two years after index discharge												
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	strong association ²	60/214 (28%)	11/214 (5.1%)	RR 5.45 (2.95 to 10.08)	229 more per 1000 (from 100 more to 467 more)	⊕⊕OO LOW	IMPORTANT	
100% atte	ndance rate of	appointm	ents						•				
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	47/214 (22%)	79/214 (36.9%)	RR 0.59 (0.44 to 0.81)	151 fewer per 1000 (from 70 fewer to 207 fewer)	⊕OOO VERY LOW	CRITICAL	
Number o	f days in hospi	tal (Better	r indicated by low	er values)		•			•				
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	214	214	-	MD 17 lower (64.44 lower to 30.44 higher)	⊕OOO VERY LOW	CRITICAL	
Number o	of diverted parti	cipants w	ith no mental hea	Ith disorders									
1	observational studies	serious ¹	no serious inconsistency	no serious indirectness	very serious ³	none	6/214 (2.8%)	0/214 (0%)	RR 13 (0.74 to 229.33)	-	⊕OOO VERY LOW	CRITICAL	

¹ James 2002 - controlled cohort study; No blinding; Few missing cases and available case data analysis ² The effect size is considered large if 95% of RR<0.8 or RR>1.25. ³ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

Patient Navigation Intervention (PNI): Motivational feedback vs Control for substance misuse disorders (26 weeks follow-up) N.6.3

			Quality asse	essment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Patient navigation intervention (at 26 weeks follow-up)	Control	Relative (95% Cl)	Absolute		
Number o	of participants	who use	d drugs		•							
1	randomised trials				very serious ²	none	1/8 (12.5%)	2/10 (20%)	RR 0.62 (0.07 to 5.72)	76 fewer per 1000 (from 186 fewer to 944 more)	⊕OOO VERY LOW	CRITICAL
Number o	of participants	who use	d alcohol to intoxi	ication	•			•				
1	randomised trials			no serious indirectness	very serious ²	none	1/8 (12.5%)	3/10 (30%)	OR 0.33 (0.03 to 4.04)	176 fewer per 1000 (from 287 fewer to 334 more)	⊕OOO VERY LOW	CRITICAL
Average days when mental health was not good in the last 30 days (Better indicated by lower values)												
1	randomised trials			no serious indirectness	very serious ³	none	8	10	-	MD 1.1 lower (9.74 lower to 7.54 higher)	⊕OOO VERY LOW	CRITICAL

¹ Binswanger 2015 - unclear randomization with appropriate allocation concealment, no blinding and appropriate attrition rate

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

Neighbourhood outreach (Before and After) N.6.4

			Quality asses	sment			No of patient	s		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Neighbourhood outreach	Control	Relative (95% CI)	Absolute		
Proportion	n of crime conta	cts with p	olicing team escal	ated to court								
	observational studies			no serious indirectness	serious ²	none		149/308 (48.4%)		155 fewer per 1000 (from 73 fewer to 223 fewer)	⊕OOO VERY LOW	CRITICAL

¹ Earl 2015 – before and after study; available case analysis; high risk of selective outcome report ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25..

N.6.5 Drug Rehabilitation Requirement (DRR) (formerly known as Drug Testing Treatment Order (DTTO) vs TAU for substance misuse disorders

			Quality asso	essment				lo of tients		Effect	Quality	Importance
No of studies	Design Inconsistency Indirectness Imprecision						DRR	Control	Relative (95% Cl)	Absolute		
MAP total s	scores (Better indi	cated by lo	wer values)									
1	observational studies	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	25	27	-	MD 20.2 lower (52 lower to 11.6 higher)	⊕OOO VERY LOW	CRITICAL
Overall sat	isfaction (Better in	ndicated by	lower values)	•	•	•						
1	observational studies	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	25	27	-	MD 2.1 higher (1.16 to 3.04 higher)	⊕OOO VERY LOW	CRITICAL
HoNOS total score (Better indicated by lower values)												
1	observational studies	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	25	27	-	MD 0.2 lower (2.44 lower to 2.04 higher)	⊕OOO VERY LOW	CRITICAL

¹ Naeem 2007 – controlled cohort study; missing data imputed by regression ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries..

N.6.6 **Case Management**

Case Management vs TAU for substance misuse disorders N.6.6.1

			Quality as	sessment			No of patie	ents		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Case management	TAU	Relative (95% CI)	Absolute		
Rearrest ·	arrest - Post-treatment											
1	randomised trials	- /	no serious inconsistency	serious ²	serious ³	none		56/135 (41.5%)		41 fewer per 1000 (from 124 fewer to 58 more)	⊕OOO VERY LOW	IMPORTANT
Rearrest ·	· 3 month follo	ow-up										
1	randomised trials		no serious inconsistency	no serious indirectness ²	serious ³	none	56/224 (25%)	48/238 (20.2%)		48 more per 1000 (from 24 fewer to 149 more)	⊕⊕OO LOW	IMPORTANT

Reconvic	tion –Post-tre	atment										
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	serious ³	none	58/369 (15.7%)	28/135 (20.7%)	RR 0.76 (0.51 to 1.14)	50 fewer per 1000 (from 102 fewer to 29 more)	⊕OOO VERY LOW	IMPORTAN ⁻
Reincarce	eration - Post-	treatment										
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	serious ³	none	99/369 (26.8%)	44/135 (32.6%)	RR 0.82 (0.61 to 1.11)	59 fewer per 1000 (from 127 fewer to 36 more)	⊕OOO VERY LOW	IMPORTAN
Reincarce	eration - 3 mo	nth follow	-up			•		•				
1	randomised trials	serious ⁴	no serious inconsistency	no serious indirectness	very serious ³	none	54/224 (24.1%)	55/238 (23.1%)	RR 1.04 (0.75 to 1.45)	9 more per 1000 (from 58 fewer to 104 more)	⊕OOO VERY LOW	IMPORTAN ⁻
Reincarce	eration - 12 m	onth follo	w-up: female sam	ple	•	-						
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	very serious	none	16/77 (20.8%)	22/77 (28.6%)	RR 0.73 (0.41 to 1.27)	77 fewer per 1000 (from 169 fewer to 77 more)	⊕OOO VERY LOW	IMPORTAN
Reincarce	eration - 12 m	onth follo	w-up: male samp	e				•				
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	serious ³	none	120/354 (33.9%)	127/354 (35.9%)	RR 0.94 (0.77 to 1.16)	22 fewer per 1000 (from 83 fewer to 57 more)	⊕⊕OO LOW	IMPORTAN
Number o	of days jailed i	n past 6 r	nonths (12 month	follow-up) (Bette	er indicated by lo	ower values)						
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	no serious imprecision	none	207	204	-	MD 0.47 higher (6.65 lower to 7.59 higher)	⊕⊕⊕O MODERATE	IMPORTAN ⁻
Drug rela	ted crimes in	past 6 mo	nths (12 month fe	ollow-up) (Better	indicated by low	er values)						
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	serious ³	none	207	204	-	MD 25.6 lower (235.88 lower to 184.68 higher)	⊕⊕OO LOW	IMPORTAN ⁻
Drug rela	ted criminal a	ctivity du	ring treatment (12	months follow-u	p)	-		•				
1	randomised trials	serious ⁶	no serious inconsistency	no serious indirectness	very serious ³	none	32/147 (21.8%)	33/137 (24.1%)		24 fewer per 1000 (from 99 fewer to 94 more)	⊕OOO VERY LOW	IMPORTAN
Self-repo	rted alcohol u	se - Durin	g treatment	•		•	•	•				
1	randomised trials	serious ⁶	no serious inconsistency	no serious indirectness	serious ³	none	85/151 (56.3%)	93/137 (67.9%)		115 fewer per 1000 (from 7 fewer to 210 fewer)	⊕⊕OO LOW	CRITICAL
Self-repo	rted alcohol u	se - Post-	treatment			-						
1	randomised trials	serious ⁷	no serious inconsistency	no serious indirectness	serious ³	none	102/346 (29.5%)	90/334 (26.9%)	RR 1.09 (0.86 to 1.39)	24 more per 1000 (from 38 fewer to 105 more)	⊕⊕OO LOW	CRITICAL
Self-repo	rted alcohol u	se - 12 m	onth follow-up: fe	male sample at p	ost-treatment							
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	no serious imprecision	none	4/77 (5.2%)	22/77 (28.6%)	RR 0.18 (0.07 to 0.5)	234 fewer per 1000 (from 143 fewer to 266 fewer)	⊕⊕⊕O MODERATE	CRITICAL
Self-repo	rted alcohol u	se - 12 m	onth follow-up: m	ale sample at pos	st-treatment							
1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	serious ³	none	138/354 (39%)	166/354 (46.9%)	RR 0.83 (0.7 to 0.99)	80 fewer per 1000 (from 5 fewer to 141 fewer)	⊕⊕OO LOW	CRITICAL
Self-repo	rted drug use	- During t	reatment (marijua	ana)	·							
1	randomised trials	serious ⁶	no serious inconsistency	no serious indirectness	serious ³	none	44/151 (29.1%)	49/137 (35.8%)	RR 0.81 (0.58 to 1.14)	68 fewer per 1000 (from 150 fewer to 50 more)	⊕⊕OO LOW	CRITICAL
Self-repo	rted drug use	- During t	reatment (hard d	ugs)								

1	randomised	serious ⁶	no serious	no serious	very serious ³	none	76/151	69/137	RR 1 (0.79 to	0 fewer per 1000 (from	$\oplus OOO$	CRITICAL	
	trials		inconsistency	indirectness			(50.3%)	(50.4%)	1.26)	106 fewer to 131 more)	VERY LOW		
Self-repor	ted drug use	- Post-trea	atment	<u>.</u>									
1	randomised	serious ⁷	no serious	no serious	serious ³	none	100/346	90/334	RR 1.07 (0.84	19 more per 1000 (from	⊕⊕OO	CRITICAL	
	trials		inconsistency	indirectness			(28.9%)	(26.9%)	to 1.37)	43 fewer to 100 more)	LOW		
Self-repor	ted drug use	- 12 mont	h follow-up: femal	e sample at post-	treatment	,,		•					
1	randomised	serious⁵	no serious	no serious	very serious ³	none	8/77	13/77	RR 0.62 (0.27	64 fewer per 1000 (from	⊕000	CRITICAL	
	trials		inconsistency	indirectness	-		(10.4%)	(16.9%)	to 1.4)	123 fewer to 68 more)	VERY LOW		
Self-reported drug use - 12 month follow-up: male sample at post-treatment													
1	randomised	serious⁵	no serious	no serious	serious ³	none	74/354	95/354	RR 0.78 (0.6	59 fewer per 1000 (from	⊕⊕OO	CRITICAL	
	trials		inconsistency	indirectness			(20.9%)	(26.8%)	to 1.02)	107 fewer to 5 more)	LOW		
Injection of	drug use (pos	t-treatme	nt)	<u>.</u>									
1	randomised	serious ⁴	no serious	no serious	very serious ³	none	9/224	12/238	RR 0.8 (0.34	10 fewer per 1000 (from	⊕000	CRITICAL	
	trials		inconsistency	indirectness			(4%)	(5%)	to 1.85)	33 fewer to 43 more)	VERY LOW		
Abstinent	- During treat	tment (at '	12 months)	<u>.</u>									
1	randomised	serious ⁶	no serious	no serious	serious ³	none	42/147	30/136	RR 1.3 (0.86	66 more per 1000 (from	⊕⊕OO	CRITICAL	
	trials		inconsistency	indirectness			(28.6%)	(22.1%)	to 1.94)	31 fewer to 207 more)	LOW		
Abstinent	- Post-treatm	ent			•	• • •		•	•				
1	randomised	serious4	no serious	no serious	very serious ³	none	54/224	55/238	RR 1.04 (0.75	9 more per 1000 (from 58	⊕000	CRITICAL	
	trials		inconsistency	indirectness	-		(24.1%)	(23.1%)		fewer to 104 more)	VERY LOW		
			tion. No blinding of 1					•					

¹ Hanlon 1999 - Unclear randomisation; No blinding; Unclear attrition

² Evidence was downgraded by one level because study population of one study (Hanlon 1999) differed from the review question in that the study included unclear proportion of ex-herion/cocaine users.
 ³ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome

respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

⁴ Scott 2012 - appropriate randomisation with concealment; No blinding; Unclear attrition bias; No selective outcomes report

⁵ Johnson 2011/Friedmann 2012 - Unclear randomisation with unclear allocation concealment; No blinding; ITT analysis; Appropriate outcome report

⁶ Rossman 1999 - Appropriate randomisation with allocation concealment; No blinding; Unclear drop-out; Appropriate selective outcome report

⁷ Prendergast 2011 - Unclear randomisation with unclear allocation concealment; No blinding; Unclear attrition risk; high risk of selective outcome report

Case management vs active intervention for substance misuse disorders N.6.6.2

			Quality as	sessment			No of p					Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Case management			Absolute		
Remained	nained in treatment for 6 months											
1		- ,	no serious inconsistency	serious ²	no serious imprecision	none				258 more per 1000 (from 106 more to 457 more)	⊕OOO VERY LOW	CRITICAL
Rearrest -	arrest - Post-treatment											
1	randomised	very	no serious	serious ²	serious ³	none	93/270	44/99	RR 0.78	98 fewer per 1000	⊕000	IMPORTANT

	trials	serious ¹	inconsistency				(34.4%)	(44.4%)	(0.59 to 1.02)	(from 182 fewer to 9 more)	VERY LOW	
Rearrest -	3 month folle	ow-up	•	•	•							
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	96/247 (38.9%)	93/264 (35.2%)	RR 1.1 (0.88 to 1.38)	35 more per 1000 (from 42 fewer to 134 more)	⊕OOO VERY LOW	IMPORTANT
Rearrest -	12 month fol	low-up	•	•	•				•	•		-
1	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	serious ³	none	41/69 (59.4%)	33/64 (51.6%)		77 more per 1000 (from 77 fewer to 294 more)	⊕OOO VERY LOW	IMPORTANT
Rearrest f	for drug crime	e (3 month	n follow-up)									
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	very serious ⁶	none	48/247 (19.4%)	49/264 (18.6%)	RR 1.05 (0.73 to 1.5)	9 more per 1000 (from 50 fewer to 93 more)	⊕OOO VERY LOW	IMPORTANT
Reconvic	tion - Post-tre	atment										
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	serious ³	none	37/270 (13.7%)	21/99 (21.2%)	RR 0.65 (0.4 to 1.05)	74 fewer per 1000 (from 127 fewer to 11 more)	⊕OOO VERY LOW	IMPORTANT
Reconvic	tion - 3 month	n follow-u	p							· · ·		•
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	67/247 (27.1%)	54/264 (20.5%)	RR 1.33 (0.97 to 1.81)	68 more per 1000 (from 6 fewer to 166 more)	⊕000 VERY LOW	IMPORTANT
Reincarce	eration - Post-	treatment	t i	4	-1				-1			4
1	randomised trials	very serious ¹	no serious inconsistency	serious ²	very serious ³	none	71/270 (26.3%)	28/99 (28.3%)	RR 0.93 (0.64 to 1.35)	20 fewer per 1000 (from 102 fewer to 99 more)	⊕000 VERY LOW	IMPORTANT
Reincarce	eration - 3 mo	nth follow	/-up	1	-1	-1			-1	<u> </u>		•
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	88/247 (35.6%)	86/264 (32.6%)		29 more per 1000 (from 46 fewer to 127 more)	⊕OOO VERY LOW	IMPORTANT
Any self-r	eported drug	use (3 m	onth follow-up)									
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	100/247 (40.5%)	100/264 (37.9%)		27 more per 1000 (from 53 fewer to 125 more)	⊕OOO VERY LOW	CRITICAL
Positive h	nair test (3 mo	onth follow	v-up) - Crack/Coc	aine								
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	97/247 (39.3%)	99/264 (37.5%)	RR 1.05 (0.84 to 1.3)	19 more per 1000 (from 60 fewer to 112 more)	⊕OOO VERY LOW	CRITICAL
Positive h	nair test (3 mo	onth follow	v-up) - Marijuana									
1	randomised trials	very serious ⁴	no serious inconsistency	no serious indirectness	serious ³	none	50/247 (20.2%)	71/264 (26.9%)	RR 0.75 (0.55 to 1.03)	67 fewer per 1000 (from 121 fewer to 8 more)	⊕OOO VERY LOW	CRITICAL

¹ Hanlon 1999 - Unclear randomisation; No blinding; Unclear attrition

² Evidence was downgraded by one level because study population of one study (Hanlon 1999) differed from the review question in that the study included unclear proportion of ex-herion/cocaine users. ³ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

⁴ Needels 2005 - Unclear randomisation and allocation concealment; No blinding; Available case analysis with unclear drop-out; appropriate outcome report

⁵ Kinlock 2007/Kinlock 2009/ Gordon 2008 - Permuted block randomisation with unclear allocation concealment; No blinding; ITT analysis with uncomparable drop-out rates

⁶ Evidence was downgraded by one level due to serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of 50%-74.99%) and by two levels due to very serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of >75%). Random Effect Model was used if I-squared inconsistency statistic was more than or equal to 50%.

N.6.6.3 Assertive community treatment vs TAU for substance misuse disorders

			Quality asse	ssment			No of patient	S		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Assertive Community Treatment	TAU	Relative (95% Cl)	Absolute		
Urine test	positive for d	rug use di	uring treatment						•			
			no serious inconsistency	no serious indirectness	serious ²	none	14/45 (31.1%)	6/45 (13.3%)	RR 2.33 (0.98 to 5.53)	177 more per 1000 (from 3 fewer to 604 more)	⊕OOO VERY LOW	CRITICAL
Injection d	rug use durin	g treatme	nt (self-report)									
		- ,	no serious inconsistency	no serious indirectness	very serious ²	none	10/56 (17.9%)	14/63 (22.2%)	RR 0.8 (0.39 to 1.66)	44 fewer per 1000 (from 136 fewer to 147 more)	⊕OOO VERY LOW	CRITICAL
Drug use d	luring treatme	ent (self-re	eport)									
		- ,	no serious inconsistency	no serious indirectness	serious ²	none	40/56 (71.4%)	40/63 (63.5%)	RR 1.13 (0.88 to 1.44)	83 more per 1000 (from 76 fewer to 279 more)	⊕OOO VERY LOW	CRITICAL
Reincarcerated during treatment												
		- 1	no serious inconsistency	no serious indirectness	very serious ²	none	26/56 (46.4%)	32/63 (50.8%)	RR 0.91 (0.63 to 1.33)	46 fewer per 1000 (from 188 fewer to 168 more)	⊕OOO VERY LOW	IMPORTANT

¹Martin 1993 - Unclear randomisation and allocation concealment; no blinding; Available case analysis with unclear drop-out; appropriate outcome report

² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.6.6.4 Case management vs TAU for mental health disorders other than substance misuse

No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Case management	TAU	Relative (95% CI)	Absolute		
Service ut	ilization				•		•					
	randomised trials	very serious ^{1,2}	serious ³	no serious indirectness	serious⁴	none	48/113 (42.5%)	52/110 (47.3%)	```	9 fewer per 1000 (from 208 fewer to 340 more)	⊕OOO VERY LOW	CRITICAL
Rate of re-	offending	•			•							•
	randomised trials		no serious inconsistency	no serious indirectness	serious⁴	none	122/236 (51.7%)	99/196 (50.5%)		15 more per 1000 (from 81 fewer to 136 more)	⊕OOO VERY LOW	CRITICAL
No of days	s in jail(up to 2	24 mths follo	w-up) (Better indic	ated by lower val	ues)							•
	randomised trials	5 5 6	no serious inconsistency	no serious indirectness	serious ⁴	none	209	160	-	MD 12.24 higher (21.87 to 2.61 lower)	⊕OOO VERY LOW	CRITICAL
Quality of	life (Better ind	dicated by lov	wer values)									
	randomised trials	, <u> </u>	no serious inconsistency	no serious indirectness	serious⁴	none	53	39	-	MD 0.09 higher (0.51 lower to 0.69 higher)	⊕OOO VERY LOW	CRITICAL

¹ Jarrett 2012 – Unclear randomisation and allocation concealment; No blinding; Available case analysis

² Wang 2012 – Appropriate randomisation and allocation concealment; Unclear blinding; ITT analysis

³Evidence was downgraded by one level due to serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of 50%-74.99%) and by two levels due to very serious heterogeneity (chi-squared p<0.1, I-squared inconsistency statistic of >75%). Random Effect Model was used if I-squared inconsistency statistic was more than or equal to 50%.

⁴ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

⁵ Cosden 2003 – Unclear randomisation and allocation concealment; Unclear blinding; Available case analysis

⁶ Solomon 1994 – Unclear randomisation and allocation concealment; No blinding; Unclear risk of attrition bias

⁷ Cusack 2010 – Unclear randomisation and allocation concealment; ITT .

N.6.7 Drug court

N.6.7.1 Drug court vs TAU for substance misuse disorders

	Linconsistency Indirectness Imprecision									Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Drug court	TAU	Relative (95% CI)	Absolute		
Days of su	ibstance use (12 month	follow-up) - Alcoho	ol (Better indicated	by lower values	5)						
1 randomised trials serious ¹ no serious inconsistency no serious indirectness none 86 71 - MD 43.10 lower (46.8 to ⊕⊕⊕O 39.4 lower)											CRITICAL	
Days of su	Days of substance use (12 month follow-up) - Cocaine (Better indicated by lower values)											

	randomised trials	serious ¹	no serious inconsistency		no serious imprecision	none	86	71	-	MD 43.70 lower (48.16 to 39.24 lower)	⊕⊕⊕O MODERATE	CRITICAL	
Days of su	ıbstance use (12 month	follow-up) - Heroin	e (Better indicate	d by lower values	s)							
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	86	71	-	MD 54.50 lower (59.42 to 49.58 lower)	⊕⊕⊕O MODERATE	CRITICAL	
Rearrest (*	arrest (12 month follow-up)												
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	37/86 (43%)	46/71 (64.8%)	RR 0.66 (0.49 to 0.89)	220 fewer per 1000 (from 71 fewer to 330 fewer)	⊕⊕OO LOW	IMPORTANT	
Maximum	Crime Serious	sness Sca	le (Better indicated	by lower values)									
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	86	71	-	MD 1.12 lower (1.18 to 1.06 lower)	⊕⊕⊕O MODERATE	IMPORTANT	

¹ Gottfredson 2005 - Unclear randomisation and allocation concealment; No blinding; Unclear analysis; Insufficient outcome report ² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25

Drug court vs active intervention for substance misuse disorders N.6.7.2

			Quality asse	essment			No of patients	5		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Drug court versus active intervention	Control	Relative (95% CI)	Absolute		
Removed	from treatme	nt due to	unsatisfactory pro	gress								
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	11/85 (12.9%)	10/65 (15.4%)		25 fewer per 1000 (from 95 fewer to 132 more)	⊕000 VERY LOW	CRITICAL
Addiction	Severity Inde	x (ASI): a	cohol composite	score (Scale from	n 0 to 9; lowe	er better)						
	randomised trials	serious ³	no serious inconsistency	no serious indirectness	very serious ⁴	none	31	31	-	MD 0.02 lower (0.04 lower to 0 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Inde	x (ASI): d	rug composite sco	re (Scale from 0	to 9; lower b	etter)						
	randomised trials	serious ³	no serious inconsistency	no serious indirectness	very serious	none	31	31	-	MD 0.01 lower (0.04 lower to 0.02 higher)	⊕OOO VERY LOW	CRITICAL
Number of	f sanctions d	uring treat	tment (Better indic	ated by lower va	lues)							
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ⁴	none	85	65	-	MD 0.90 lower (1.99 lower to 0.19 higher)	⊕⊕OO LOW	CRITICAL
Number of	f sanctions d	uring treat	tment resulting in	jail detention (Be	etter indicate	d by lower values)						
	randomised trials	serious1		no serious indirectness	serious⁴	none	67	54	-	MD 0.5 lower (0.99 to 0.01 lower)	⊕⊕OO LOW	IMPORTANT
Reincarce	ration during	treatment	ł									

1	randomised trials	serious⁵	no serious inconsistency	no serious indirectness	very serious ⁶	none		25/68 (36.8%)	```	81 fewer per 1000 (from 195 fewer to 103 more)	 IMPORTANT
Urine test	positive for c	lrugs (pos	st-treatment)								
1	randomised trials	serious ³	no serious inconsistency	no serious indirectness	very serious ⁶	none	2/31 (6.5%)	5/31 (16.1%)		97 fewer per 1000 (from 148 fewer to 147 more)	CRITICAL

¹ Messina 2012 - Inappropriate randomisation with adequate allocation concealment; No blinding; low risk of attrition bias; appropriate selective outcomes

² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ Dakof 2010 - Unclear randomisation and allocation concealment; No blinding; ITT analysis; insufficient outcome report

⁴ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

⁵ Jones 2013 - Permuted block randomisation with unclear allocation concealment; No blinding; low risk of attrition bias; insufficient outcome report

⁶ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.6.8 Opioid substitution therapy

N.6.8.1 Opioid substitution therapy + case management vs active intervention

			Quality as	sessment			No of patier	nts		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Opioid substitution therapy + case management	Active intervention	Relative (95% CI)	Absolute		
Complete	d jail treatme	ent										
1	randomised trials	very serious ¹			no serious imprecision	none	64/104 (61.5%)	68/107 (63.6%)	RR 0.96 (0.81 to 1.14)	25 fewer per 1000 (from 121 fewer to 89 more)	⊕⊕OO LOW	CRITICAL
Complete	d jail treatme	ent - Fema	ale sample		•							
1	randomised trials	- /		no serious indirectness	serious ²	none	27/32 (84.4%)	27/31 (87.1%)	RR 0.97 (0.79 to 1.18)	26 fewer per 1000 (from 183 fewer to 157 more)	⊕OOO VERY LOW	CRITICAL
Complete	d jail treatme	ent - Male	sample	•	•							
1	randomised trials	1		no serious indirectness	very serious ²	none	37/72 (51.4%)	41/76 (53.9%)	RR 0.95 (0.7 to 1.29)	27 fewer per 1000 (from 162 fewer to 156 more)	⊕OOO VERY LOW	CRITICAL
Urine test	t positive for	cocaine -	1 month follow-u	ıp								
1	randomised trials	very serious ³		no serious indirectness	serious ²	none	31/70 (44.3%)	73/130 (56.2%)	RR 0.79 (0.58 to 1.07)	118 fewer per 1000 (from 236 fewer to 39 more)	⊕OOO VERY	CRITICAL

											LOW	
Jrine tes	st positive for	cocaine ·	- 6 month follow	-up				•				
1	randomised trials	very serious ²	no serious inconsistency	no serious indirectness	very serious ²	none	15/25 (60%)	34/51 (66.7%)	RR 0.9 (0.62 to 1.31)	67 fewer per 1000 (from 253 fewer to 207 more)	⊕000 VERY LOW	CRITICAL
Urine tes	-		- 12 month follow	w-up								
1	randomised trials	very serious ²	no serious inconsistency	no serious indirectness	serious ²	none	19/44 (43.2%)	49/71 (69%)	RR 0.63 (0.43 to 0.91)	255 fewer per 1000 (from 62 fewer to 393 fewer)	⊕000 VERY LOW	CRITICAL
Urine tes	st positive for	opioids -	1 month follow	-up				•				
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	19/70 (27.1%)	67/130 (51.5%)	RR 0.53 (0.35 to 0.8)	242 fewer per 1000 (from 103 fewer to 335 fewer)	⊕000 VERY LOW	CRITICAL
Urine tes	st positive for	opioids -	6 month follow	-up	•	•		•				
1	randomised trials	very serious	no serious inconsistency	no serious indirectness	serious ²	none	3/12 (25%)	26/45 (57.8%)	RR 0.43 (0.16 to 1.19)	329 fewer per 1000 (from 485 fewer to 110 more)	⊕000 VERY LOW	CRITICAL
Urine tes	st positive for	opioids -	12 month follow	w-up								
1	randomised trials	very serious	no serious inconsistency	no serious indirectness	no serious imprecision	none	11/44 (25%)	40/71 (56.3%)	RR 0.44 (0.26 to 0.77)	315 fewer per 1000 (from 130 fewer to 417 fewer)	⊕⊕OO LOW	CRITICAL
Days of a	substance us	e (12 mor	nth follow-up) - 0	Cocaine (Better	indicated by lov	ver values)						
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious⁴	none	71	133	-	MD 27.40 lower (47.25 to 7.55 lower)	⊕000 VERY LOW	CRITICAL
Days of a	substance us	e (12 mor	nth follow-up) - H	leroin (Better in	dicated by lowe	er values)						
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious⁴	none	71	133	-	MD 36.80 lower (74.3 lower to 0.7 higher)	⊕000 VERY LOW	CRITICAL
Self-repo	orted days wit	th drug us	se in past 30 day	s (6 month follo	w-up) - Crack/C	Cocaine						
1	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	serious ²	none	4/21 (19%)	19/41 (46.3%)	RR 0.41 (0.16 to 1.05)	273 fewer per 1000 (from 389 fewer to 23 more)	⊕000 VERY LOW	CRITICAL
Self-repo	orted days wit	th drug us	se in past 30 day	s (6 month follo	ow-up) - Heroin				·			
1	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	no serious imprecision	none	3/21 (14.3%)	22/41 (53.7%)	RR 0.27 (0.09 to 0.79)	392 fewer per 1000 (from 113 fewer to 488 fewer)	⊕⊕OO LOW	CRITICAL
Self-repo	orted days wit	th drug us	se in past 30 day	s (6 month follo	ow-up) - Marijua	na						
1	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	very serious ²	none	2/21 (9.5%)	9/41 (22%)	RR 0.43 (0.1 to 1.83)	125 fewer per 1000 (from 198 fewer to 182 more)	⊕000 VERY LOW	CRITICAL

	randomised trials	very serious⁵	no serious inconsistency	no serious indirectness	very serious ²	none	2/21 (9.5%)	15/41 (36.6%)	RR 0.26 (0.07 to 1.03)	271 fewer per 1000 (from 340 fewer to 11 more)	⊕000 VERY LOW	CRITICAL
Drug ove	rdose - 6 moi	nth follow	v-up									
1		very serious⁵	no serious inconsistency	no serious indirectness	very serious ²	none	3/21 (14.3%)	7/41 (17.1%)	RR 0.84 (0.24 to 2.91)	27 fewer per 1000 (from 130 fewer to 326 more)	⊕OOO VERY LOW	CRITICAL
Drug ove	rdose - 12 mo	onth follo	w-up		•							
1		very serious ³	no serious inconsistency	no serious indirectness	very serious ²	none	0/71 (0%)	6/133 (4.5%)	RR 0.14 (0.01 to 2.51)	39 fewer per 1000 (from 45 fewer to 68 more)	⊕000 VERY LOW	CRITICAL
Rearrest	- 6 month fol	low-up	-		•					•		
1		very serious⁵	no serious inconsistency	no serious indirectness	very serious ²	none	7/21 (33.3%)	11/41 (26.8%)	RR 1.24 (0.56 to 2.73)	64 more per 1000 (from 118 fewer to 464 more)	⊕OOO VERY LOW	IMPORTAN ⁻
Rearrest	- 12 month fo	llow-up			-							
1		very serious ³	no serious inconsistency	no serious indirectness	very serious ²	none	38/71 (53.5%)	74/133 (55.6%)	RR 0.96 (0.74 to 1.25)	22 fewer per 1000 (from 145 fewer to 139 more)	⊕000 VERY LOW	IMPORTAN
Self-repo	rted days of o	criminal a	activity (12 month	s follow-up) (Be	etter indicated I	by lower values)						
1		very serious ³	no serious inconsistency	no serious indirectness	no serious imprecision	none	71	133	-	MD 3.37 lower (35.27 lower to 28.53 higher)	⊕⊕OO LOW	IMPORTANT

¹ Gordon 2014 - Permuted blocks with adequate allocation concealment, No blinding with potential of effect size bigger in intervention group; available case analysis; appropriate outcome report ² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ Kinlock 2007/Kinlock 2009/ Gordon 2008 - Permuted block randomisation with unclear allocation concealment; No blinding; ITT analysis with incomparable drop-out rates

⁴ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries. ⁵ McKenzie 2012 - Unclear randomisation and allocation concealment; No blinding with potential increased effect size in intervention arm; per protocol analysis; appropriate outcome report

N.6.9 Automated telephony with feedback vs Automated telephony alone

			Quality as	sessment			No of pa	tients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Automated telephony with feedback	Automated telephony alone	Relative (95% Cl)	Absolute		
Improven	nent in Arnetz	and Hase	son stress question	onnaire (AHSS) (Better indicated	d by lower values)						
1	randomised trials				no serious imprecision	none	52	56		MD 2.5 higher (1.13 lower to 6.13 higher)		CRITICAL
Improven	nent in sympt	om check	list-8D (SCL-8D) (Better indicated	by lower values	s)						

1 r.	randomised	serious ¹	no serious	no serious	serious ²	none	52	56	-	MD 4.5 higher (0.22	$\oplus \oplus OO$	CRITICAL
t	rials		inconsistency	indirectness						to 8.78 higher)	LOW	
Improveme	ent in daily s	tressor a	ssessment (Bette	r indicated by lo	wer values)							
1 r	randomised	serious ¹	no serious	no serious	serious ²	none	52	56	-	MD 1.91 higher	⊕⊕OO	CRITICAL
t	trials		inconsistency	indirectness						(1.11 to 2.71 higher)	LOW	
Alcohol Ur	rge Question	naires: re	duction in alcoho	ol urge (Better in	dicated by lowe	er values)						
1 r	randomised	serious ¹	no serious	no serious	no serious	none	52	56	-	MD 0.20 higher	$\oplus \oplus \oplus \Theta$	CRITICAL
t	rials		inconsistency	indirectness	imprecision					(0.35 lower to 0.75	/ODERATE	
										higher)		
Alcohol Ur	rge Question	naires: re	eduction in alcoho	ol use (Better inc	licated by lower	r values)						
1 r	randomised	serious ¹	no serious	no serious	serious ²	none	52	56	-	MD 0.8 higher (0.11	$\oplus \oplus OO$	CRITICAL
t	rials		inconsistency	indirectness						to 1.49 higher)	LOW	
Alcohol Ur	rge Question	naires: re	duction in drug u	ise (Better indica	ated by lower va	alues)						
1 r	randomised	serious ¹	no serious	no serious	serious ²	none	52	56	-	MD 1 higher (0.41 to	$\oplus \oplus OO$	CRITICAL
t	rials		inconsistency	indirectness						1.59 higher)	LOW	
Alcohol Ur	rge Question	naires: re	eduction in drug u	irge (Better indic	ated by lower v	values)						
1 r	randomised	serious ¹	no serious	no serious	serious ²	none	52	56	-	MD 0.3 higher (0.25	$\oplus \oplus OO$	CRITICAL
t	trials		inconsistency	indirectness						lower to 0.85 higher)	LOW	
4 1	0014 II			1 11 1	1	1 11 11 7 1				1		

⁷ Andersson 2014 - Unclear randomisation with unclear allocation concealment; No blinding; Low drop-out rate with available rate analysis

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.6.10 **IDDT vs TAU**

			Quality asse	ssment			No o	f patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	IDDT	Service as usual	Relative (95% CI)	Absolute		
Rate of out	tpatient medic	ation servi	ces									
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	83/103 (80.6%)	51/79 (64.6%)	RR 1.25 (1.03 to 1.51)	161 more per 1000 (from 19 more to 329 more)	⊕⊕OO LOW	CRITICAL
No of days	in hospital (B	etter indica	ated by lower values	s)								
-	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	103	79	-	MD 5.63 lower (9.59 to 1.67 lower)	⊕⊕OO LOW	CRITICAL
Rate of cris	sis visits (Bette	er indicate	d by lower values)									
	randomised trials	serious ¹	no serious inconsistency	no serious indirectness	serious ³	none	103	79	-	MD 2.26 lower (3.82 to 0.7 lower)	⊕⊕OO LOW	CRITICAL

 Inconsistency
 Inconsistency
 Inconsistency

 ¹ Chandler 2006 - Unclear randomization with unclear allocation concealment; Blinding was not reported; Analysis by imputation

 ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome

 ³ The evidence was downgraded by one
 ³ The evidence was downgraded by one

level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.6.11 Housing first vs TAU

			Quality as	sessment			No of pat	tients		Effect	Quality	Importance
No of studies	tudies Design bias Inconsistency Indirectness Imprecision cor						Housing First	TAU	Relative (95% Cl)	Absolute		
Any offend	ce											
	randomised trials		no serious inconsistency	no serious indirectness	serious ²	none	16/197 (8.1%)	19/100 (19%)	RR 0.43 (0.23 to 0.82)	108 fewer per 1000 (from 34 fewer to 146 fewer)	⊕⊕OO LOW	IMPORTANT
Any offend	e - Scattered	HF+ACT	•	•	•	•		•	•		•	
	randomised trials		no serious inconsistency		no serious imprecision	none	6/90 (6.7%)	11/50 (22%)	RR 0.3 (0.12 to 0.77)	154 fewer per 1000 (from 51 fewer to 194 fewer)	⊕⊕⊕O MODERATE	IMPORTANT
Any offend	y offence - Congregate HF											
	randomised trials		no serious inconsistency	no serious indirectness	very serious ²	none	10/107 (9.3%)	8/50 (16%)	RR 0.58 (0.25 to 1.39)	67 fewer per 1000 (from 120 fewer to 62 more)	⊕OOO VERY LOW	IMPORTANT

¹ Somers 2013 - Unclear randomisation with unclear concealment; no blinding of participants and care administrators; ITT analysis

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.6.12 TIMA vs service as usual

			Quality asses	sment			No	of patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	ТІМА	Service as usual	Relative (95% Cl)	Absolute		
Bipolar Disc	order Symptom	Scale (BDS	SS) (Scale from 7 to 7	0; lower better)								
	randomised trials			no serious indirectness	serious ²	none	30	30	-	MD 0.27 lower (0.75 lower to 0.21 higher)	⊕⊕OO LOW	CRITICAL
Brief Psychi	atric Rating Sc	ale (BPRS)	(Scale from 18 to 12	6; lower better)								
	randomised trials			no serious indirectness	serious ²	none	30	30	-	MD 0.97 higher (1.78 lower to 3.72 higher)	⊕⊕OO LOW	CRITICAL

¹ Ehret 2013 - inappropriate randomization with unclear concealment; no blinding; available case analysis

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.6.13 Service Brokerage Intervention vs Control

			Quality asse	ssment			No of patients	6		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Service brokerage intervention	Control	Relative (95% CI)	Absolute		
Number of	f participants i	in contact	with MH service									
1	randomised trials	serious ¹		no serious indirectness	very serious ²	none	55/665 (8.3%)	47/660 (7.1%)	RR 1.16 (0.8 to 1.69)	11 more per 1000 (from 14 fewer to 49 more)	⊕OOO VERY LOW	CRITICAL
Number of	f participants	who have	seen GP									
1	randomised trials	serious ¹		no serious indirectness	very serious ²	none	21/665 (3.2%)	13/660 (2%)	RR 1.6 (0.81 to 3.17)	12 more per 1000 (from 4 fewer to 43 more)	⊕OOO VERY LOW	CRITICAL
Number of participants who attended alcohol or drug service												
1	randomised trials			no serious indirectness	serious ²	none	18/665 (2.7%)	17/660 (2.6%)	RR 1.05 (0.55 to 2.02)	1 more per 1000 (from 12 fewer to 26 more)	⊕OOO VERY LOW	CRITICAL

 1 Kinner 2013/2014a/2014b - RCT with unclear allocation concealment; Blinding of care administrators; ITT analysis

² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.6.14 Therapeutic communities

N.6.14.1 Therapeutic community versus waitlist control

			Quality asse	ssment			No of patients Effect			Effect	Quality	Importance
No of studies			Indirectness Imprecision		Other considerations	Therapeutic waitlist community control		Relative (95% Cl)	Absolute			
Days until	reincarceratio	n (Better i	ndicated by lower v	alues)								
	randomised trials			no serious indirectness	serious ²	none	199	142	-	MD 83.58 higher (32.69 to 134.47 higher)	⊕OOO VERY LOW	IMPORTANT

¹ Wexler 1999 - Unclear randomisation and allocation concealment; No blinding with potential of effect size bigger in intervention group; ITT analysis; appropriate outcome report ² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

N.6.14.2 Modified therapeutic communities versus CBT informed psychoeducation

			Quality as	sessment			No of patients		Effect		Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Modified therapeutic community	CBT informed psychoeducation	Relative (95% CI)	Absolute		
Substanc	e use (12 mo	onth follow	w-up)							•		
		very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	23/75 (30.7%)	35/64 (54.7%)	RR 0.56 (0.37 to 0.84)	241 fewer per 1000 (from 88 fewer to 345 fewer)	⊕000 VERY LOW	CRITICAL
Alcohol u	se (12 monti	n follow-u	p)					•				-
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	15/75 (20%)	24/64 (37.5%)	RR 0.53 (0.31 to 0.93)	176 fewer per 1000 (from 26 fewer to 259 fewer)	⊕OOO VERY LOW	CRITICAL
Drug use	(12 month fo	ollow-up)								•	•	•
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	18/75 (24%)	28/64 (43.8%)	RR 0.55 (0.34 to 0.89)	197 fewer per 1000 (from 48 fewer to 289 fewer)	⊕OOO VERY LOW	CRITICAL
Reincarce	eration (12 m	onth follo	ow-up)	•	•					•		
	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	no serious imprecision	none	7/75 (9.3%)	21/64 (32.8%)	RR 0.28 (0.13 to 0.63)	236 fewer per 1000 (from 121 fewer to 285 fewer)	⊕⊕OO LOW	IMPORTANT
Alcohol/d	rug offence	(12 montł	n follow-up)									
	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	27/75 (36%)	37/64 (57.8%)	RR 0.62 (0.43 to 0.9)	220 fewer per 1000 (from 58 fewer to 330 fewer)	⊕000 VERY LOW	IMPORTANT
Criminal a	activity (12 m	onth foll	ow-up)		·	••		•	•	•	•	•
	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	35/75 (46.7%)	45/64 (70.3%)	RR 0.66 (0.5 to 0.89)	239 fewer per 1000 (from 77 fewer to 352 fewer)	⊕OOO VERY LOW	IMPORTANT

¹ Sullivan 2007 - unclear randomisation and allocation concealment; No blinding; unclear analysis; self-reported data

² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

³ Sacks 2004 - Unclear randomisation and allocation concealment; unclear blinding; Available case analysis; inadequate outcome report

N.6.14.3 Enhanced therapeutic community versus standard therapeutic community

Quality assessment	No of patients	Effect	Quality In	mportance	
Quality assessment	No of patients	Effect	Quality in	mportance	

No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Enhanced therapeutic community	Standard therapeutic community	Relative (95% Cl)	Absolute		
Engagem	ent with treat	ment (Bet	ter indicated by lo	ower values)								
1		1			no serious imprecision	none	232	219		MD 0.03 higher (0.01 lower to 0.07 higher)		CRITICAL
Negative	mood (as rate	d by cour	sellor) (Better ind	dicated by lower	values)							
1		- 1	no serious inconsistency	serious ²	no serious imprecision	none	230	219	-	MD 1.79 lower (2.09 to 1.49 lower)	⊕OOO VERY LOW	CRITICAL

¹ Czuchry 2003 – unclear randomisation and allocation concealment; no blinding; unclear attrition ² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

Gender-responsive therapeutic community versus standard therapeutic community N.6.14.4

			Quality ass	essment			No of patients Effect Gender-responsive standard				Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	therapeutic	standard therapeutic community	Relative (95% CI)	Absolute		
Addiction	Severity Ind	ex (ASI): a	alcohol composit	te score (Bette	er indicated by	lower values)						
	randomised trials		no serious inconsistency	serious ²	very serious ³	none	60	55	-	MD 0.04 lower (0.08 lower to 0 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Ind	ex (ASI):	psychological co	mposite score	e (Better indica	ated by lower value	es)					
1	randomised trials	1	no serious inconsistency		no serious imprecision	none	60	55	-	MD 0.01 lower (0.1 lower to 0.08 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Ind	ex (ASI): d	drug composite s	score (Better i	ndicated by lov	wer values)		•		•		
1	randomised trials	1	no serious inconsistency	serious ²	very serious ³	none	60	55	-	MD 0.02 higher (0.0 lower to 0.04 higher)	⊕OOO VERY LOW	CRITICAL
Addiction	Severity Ind	ex (ASI): f	family composite	score (Better	indicated by l	ower values)						
1	randomised trials		no serious inconsistency	Serious ²	very serious ³	none	60	55	-	MD 0.04 lower (0.12 lower to 0.04 higher)	⊕OOO VERY LOW	CRITICAL
Participat	ted in afterca	re upon re	elease					•		•		

				. 2			/	/				
	randomised	Very	no serious	serious ²	very serious⁴	none	28/60	30/55	RR 0.86 (0.6	76 fewer per 1000	⊕000	CRITICAL
	trials	serious	inconsistency				(46.7%)	(54.5%)	to 1.23)	(from 218 fewer to	VERY	
							. ,	. ,		125 more)	LOW	
Nonths	spent in aftero	are (Bette	er indicated by lo	ower values)						,		
1	randomised	Verv	no serious	Serious ²	very serious ³	none	60	55	_	MD 1.50 higher (0.29	⊕000	CRITICAL
		serious ¹	inconsistency	Conous	very serious	none	00	00		to 2.71 higher)	VERY	
	lilais	senous	inconsistency							to 2.7 i higher)		
											LOW	
Discipliı	nary removal f	rom first i	residential treatr	nent post-rel	ease							
1	randomised	Very	no serious	Serious ²	very serious ⁴	none	8/60	8/55	RR 0.92	12 fewer per 1000	⊕000	CRITICAL
	trials	serious ¹	inconsistency		,		(13.3%)	(14.5%)	(0.37 to	(from 92 fewer to 186	VERY	
		0011040					(101070)	(1.11070)	2.28)	more)	LOW	
									2.20)	more)	LOW	
Reincar	ceration (12 m	onth follo	w-up)									
1	randomised	Very	no serious	Serious ²	very serious ⁴	none	18/60	25/55	RR 0.66	155 fewer per 1000	⊕000	IMPORTANT
	trials	serious1	inconsistency				(30%)	(45.5%)	(0.41 to	(from 268 fewer to 32	VERY	
							(((())))	(1010,0)	1.07)	more)	LOW	
	<u> </u>				<u> </u>				1.07)	morey	LOW	
Volunta	rily dropped-o	ut from fi	rst residential tre	eatment post								
1	randomised	Very	no serious	serious ²	very serious ⁴	none	10/60	17/55	RR 0.54	142 fewer per 1000	⊕000	CRITICAL
	trials	serious ¹	inconsistency				(16.7%)	(30.9%)	(0.27 to	(from 226 fewer to 25	VERY	
			,				((00000)	1.08)	more)	LOW	
			I	<u> </u>					1.00)	inore)	1011	
Months		· · ·	etter indicated b									
1	randomised	Very	no serious	Serious ²	very serious ³	none	60	55	-	MD 1.90 higher (0.5	$\oplus 000$	IMPORTANT
	trials	serious ¹	inconsistency		-					to 3.3 higher)	VERY	
											LOW	
											LOW	

¹ Messina 2010 - high risk of selection bias; No blinding; available case analysis; unclear selective outcome report

² Evidence was downgraded by one level because study population of one study (Messina 2010) differed from the review question in that not all the participants met the proxy measure criteria for substance misuse disorder.

³ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or both boundaries of the defined minimally important difference (MID) for the outcome, respectively. For continuous outcomes, +/-0.5 (mean for 2 studies and median for 3 or more studies) times SD of the control group (if MD was used) were considered as MID boundaries.

⁴ Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.6.14.5 Gender-specific therapeutic community versus CBT informed psychoeducation

			Quality ass	sessment			No of	patients		Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Gender-specific therapeutic community	CBT informed psychoeducation	Relative (95% CI)	Absolute		
Self-repo	elf-reported criminal activity (sexual)											
1	randomised	very	no serious	no serious	very serious ²	none	3/163	8/151	RR 0.35	34 fewer per 1000	⊕000	IMPORTANT

					- 1						1	1
	trials	serious ¹	inconsistency	indirectness			(1.8%)	(5.3%)	(0.09 to 1.29)	(from 48 fewer to 15 more)	VERY LOW	
Receivir	ng mental hea	Ith treatmo	ent at follow-up			-			•	•		•
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	very serious ²	none	65/163 (39.9%)	63/151 (41.7%)	RR 0.96 (0.73 to 1.25)	17 fewer per 1000 (from 113 fewer to 104 more)	⊕OOO VERY LOW	CRITICAL
Alcohol	use (follow-u	p NR)	•		-							
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	41/163 (25.2%)	29/151 (19.2%)	RR 1.31 (0.86 to 2)	60 more per 1000 (from 27 fewer to 192 more)	⊕OOO VERY LOW	CRITICAL
Frequen	cy of alcohol	use (follo	w-up NR) (Bette	r indicated by lo	ower values)							
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	87	75	-	MD 0.25 higher (0.42 lower to 0.92 higher)	⊕⊕OO LOW	CRITICAL
Frequen	cy of drug us	e (follow-u	up NR) (Better in	dicated by lowe	er values)							
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	111	95	-	MD 0.42 lower (1.14 lower to 0.3 higher)	⊕⊕OO LOW	CRITICAL
Self-rep	orted drug us	e - 6 mont	h follow-up			•			-			
2	randomised trials	1 10	no serious inconsistency	no serious indirectness	serious ²	none	76/374 (20.3%)	87/328 (26.5%)	RR 0.77 (0.59 to 1.01)	61 fewer per 1000 (from 109 fewer to 3 more)	⊕OOO VERY LOW	CRITICAL
Self-rep	orted drug us	e - 12 mor	th follow-up	L	-1		- · ·		- i	I		Į
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	50/207 (24.2%)	54/163 (33.1%)	RR 0.73 (0.53 to 1.01)	89 fewer per 1000 (from 156 fewer to 3 more)	⊕OOO VERY LOW	CRITICAL
Rearres	t - 6 month fo	llow-up	•			•						
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	19/211 (9%)	32/177 (18.1%)	RR 0.5 (0.29 to 0.85)	90 fewer per 1000 (from 27 fewer to 128 fewer)	⊕000 VERY LOW	IMPORTANT
Rearres	t - 12 month f	ollow-up										
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	23/207 (11.1%)	11/163 (6.7%)	RR 1.65 (0.83 to 3.28)	44 more per 1000 (from 11 fewer to 154 more)	⊕000 VERY LOW	IMPORTAN
Rearres	t - Follow-up l	NR	•		•	•			1	•		•
1	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	42/163 (25.8%)	53/151 (35.1%)	RR 0.73 (0.52 to 1.03)	95 fewer per 1000 (from 168 fewer to 11 more)	⊕OOO VERY LOW	IMPORTANT
Reincar	ceration	•			·		·					•
1	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	59/257 (23%)	59/211 (28%)	RR 0.82 (0.6 to 1.12)	50 fewer per 1000 (from 112 fewer to 34 more)	⊕OOO VERY LOW	IMPORTANT

Self-rep	orted criminal	l activity (a	iny) - 6 month fo	llow-up								
	randomised trials	very serious ^{1,3}	no serious inconsistency	no serious indirectness	serious ²	none	130/374 (34.8%)	149/328 (45.4%)	RR 0.77 (0.64 to 0.92)	104 fewer per 1000 (from 36 fewer to 164 fewer)	⊕OOO VERY LOW	IMPORTAI
Self-rep	orted criminal	activity (a	iny) - 12 month f	ollow-up								
	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	72/207 (34.8%)	67/163 (41.1%)	RR 0.85 (0.65 to 1.1)	62 fewer per 1000 (from 144 fewer to 41 more)	⊕OOO VERY LOW	IMPORTAN
Self-rep	orted criminal	l activity (c	lrugs) - 6 month	follow-up								
2	randomised trials	very serious ^{1,3}	no serious inconsistency	no serious indirectness	serious ²	none	106/374 (28.3%)	108/328 (32.9%)	RR 0.86 (0.69 to 1.08)	46 fewer per 1000 (from 102 fewer to 26 more)	⊕000 VERY LOW	IMPORTAN
Self-rep	orted criminal	l activity (c	Irugs) - 12 mont	h follow-up	<u>.</u>							
	randomised trials	very serious ³	no serious inconsistency	no serious indirectness	serious ²	none	62/207 (30%)	60/163 (36.8%)	RR 0.81 (0.61 to 1.09)	70 fewer per 1000 (from 144 fewer to 33 more)	⊕000 VERY LOW	IMPORTAN
Receivir	ng substance	abuse trea	tment at follow-	up	<u>.</u>		<u>.</u>			· · · ·		
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	serious ²	none	109/163 (66.9%)	118/151 (78.1%)	RR 0.86 (0.75 to 0.98)	109 fewer per 1000 (from 16 fewer to 195 fewer)	⊕OOO VERY LOW	CRITICAL
Beck De	pression Inve	entory (BD) total score (Be	tter indicated b	y lower values)	-			•		•
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	163	151	-	MD 2.64 lower (5.26 to 0.02 lower)	⊕⊕OO LOW	CRITICAL
Brief Sy	mptom Invent	tory (BSI) t	otal score (Bette	er indicated by	ower values)				•			
	randomised trials	very serious ¹	no serious inconsistency	no serious indirectness	no serious imprecision	none	163	151	-	MD 1.63 lower (4.45 lower to 1.19 higher)	⊕⊕OO LOW	CRITICAL
Post-tra	umatic Sympt	om Severi	ty Scale (PSS) (Better indicated	by lower value	es)						
	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	163	151	-	MD 2.90 lower (5.68 to 0.12 lower)	⊕⊕OO LOW	CRITICAL

¹Sacks 2008 - unclear randomisation and allocation concealment; No blinding; analysis by regression technique; appropriate outcome report ² Evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or more boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25. ³ Sacks 2012a - unclear randomisation and allocation concealment; No blinding with potential of effect size bigger in intervention group; available case analysis

Re-entry modified therapeutic community versus treatment as usual N.6.14.6

	Quality assessment		No of patients		Effect	Quality	Importance
No of Design Ris	sk of Inconsistency Indirectness Imprecision	Other	Re-entry modified treatment as	Relative	Absolute		

studies		bias				considerations	therapeutic community	usual	(95% CI)			
Reincarce	eration (12 mo	onth post	prison release)		•							-
	randomised trials				very serious ²	none	14/71 (19.7%)	21/56 (37.5%)	RR 0.53 (0.29 to 0.94)	176 fewer per 1000 (from 23 fewer to 266 fewer)	⊕000 VERY LOW	IMPORTANT
Criminal a	riminal activity											
	randomised trials				very serious²	none	25/63 (39.7%)	29/47 (61.7%)	RR 0.64 (0.44 to 0.94)	222 fewer per 1000 (from 37 fewer to 346 fewer)	⊕OOO VERY LOW	IMPORTANT
Alcohol/D	rug offence											
	randomised trials				very serious²	none	23/63 (36.5%)	27/47 (57.4%)	RR 0.64 (0.42 to 0.96)	207 fewer per 1000 (from 23 fewer to 333 fewer)	⊕000 VERY LOW	IMPORTANT

¹ Sacks 2012b – inappropriate randomisation without allocation concealment; no blinding; ITT analysis; lack of outcome report on percentages of therapeutic community in prison ² The evidence was downgraded by one level and two levels if the confidence interval crossed or touched one or two boundaries of the defined minimally important difference (MID) for the outcome respectively. The MID boundaries for dichotomous outcomes (RR) were 0.8 to 1.25.

N.7 Staff Training

N.7.1 Organisational linkage intervention (OLI) plus medication-assisted training (MAT) vs Training alone for substance misuse disorders

			Quality as	sessment			No of patients			Effect	Quality	Importance
No of studies	Design	Risk of bias	Inconsistency	Indirectness	Imprecision	Other considerations	Organisational Linkage Intervention (OLI) plus training (RQ 5.1)	Training alone	Relative (95% CI)	Absolute		
MAT-Met	hadone: Fam	iliarity wi	th medication (C	hange from bas	eline to post in	tervention; range	-4 to 4; higher is better)					
1	randomised trials	serious ¹	no serious inconsistency		no serious imprecision	none	383	464	Mean 0.26 (SD 1.01)	MD 0.14 higher (0.03 lower to 0.31 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT-Met	hadone: Refe	rral know	vledge (Change fi	om baseline to	post intervent	ion; range -4 to 4;	higher is better)		,		•	
1	randomised trials		no serious inconsistency		no serious imprecision	none	383	464	Mean 0.24 (SD 1.23)	MD 0.04 higher (0.11 lower to 0.19 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT-Met	hadone:Inten	t to refer	clients to MAT (C	Change from bas	seline to post i	ntervention; range	e -4 to 4; higher is better)					
1	randomised trials		no serious inconsistency		no serious imprecision	none	383	464	Mean 0.05 (SD 1.24)	MD 0.38 higher (0.19 to 0.57 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT-Met	hadone: Ove	rall perce	ption and knowle	dge (Change fr	om baseline to	post intervention	; range -4 to 4; higher is bet	ter)				

1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	383	464	Mean 0.01 (SD 0.04)	MD 0.2 higher (0.13 to 0.27 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT-Bup	renorphine:	Familiarit	y with the Medica	ation (Change f	om baseline to	post intervention	n; range -4 to 4; higher is bet	ter)				•
1	randomised trials		no serious inconsistency	no serious indirectness	no serious imprecision	none	383	464	Mean 0.39 (SD 1.52)	MD 0.01 higher (0.19 lower to 0.21 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT-Bup	renorphine:	Referral I	Knowledge (Chan	ge from baselir	ne to post inter	vention; range -4	to 4; higher is better)				•	•
1	randomised trials			no serious indirectness	no serious imprecision	none	383	464	Mean 0.34 (SD 1.33)	MD 0.07 higher (0.12 lower to 0.26 higher)	⊕⊕⊕O MODERATE	IMPORTAN
MAT-Bup	renorphine:	Intent to	refer clients to M	AT (Change from	n baseline to p	ost intervention;	range -4 to 4; higher is better	r)				
1	randomised trials			no serious indirectness	no serious imprecision	none	383	464	Mean 0.15 (SD 1.35)	MD 0.15 higher (0.02 lower to 0.32 higher)	⊕⊕⊕O MODERATE	IMPORTANT
MAT- Bup	prenorphine:	Overall p	erception and kn	owledge (Chan	ge from baseli	ne to post interve	ntion; range -4 to 4; higher is	better)				
1	randomised trials			no serious indirectness	no serious imprecision	none	383	464	Mean 0.03 (SD 0.66)	MD 0.13 higher (0.05 to 0.21 higher)	⊕⊕⊕O MODERATE	IMPORTANT

1. Friedmann 2015 - unclear randomisation and concealment; comparable management of experimental and control group; appropriate outcome report