# Articles

# Effectiveness of psychological interventions in prison to reduce recidivism: a systematic review and meta-analysis of randomised controlled trials

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# **Summary**

**Background** Repeat offending, also known as criminal recidivism, in people released from prison has remained high over many decades. To address this, psychological treatments have been increasingly used in criminal justice settings; however, there is little evidence about their effectiveness. We aimed to evaluate the effectiveness of interventions in prison to reduce recidivism after release.

Methods For this systematic review and meta-analysis, we searched the Cochrane Central Register of Controlled Trials, Embase, Global Health, MEDLINE, PsycINFO, and Google Scholar for articles published from database inception to Feb 17, 2021, without any language restrictions. We searched for randomised controlled trials (RCTs) that evaluated the effect of psychological interventions, delivered to adolescents and adults during incarceration, on recidivism outcomes after release. We excluded studies of solely pharmacological interventions and of participants in secure psychiatric hospitals or special residential units, or attending therapies mainly delivered outside of the prison setting. We extracted summary estimates from eligible RCTs. Data were extracted and appraised according to a prespecified protocol, with effect sizes converted to odds ratios. We used a standardised form to extract the effects of interventions on recidivism and estimated risk of bias for each RCT. Planned sensitivity analyses were done by removing studies with fewer than 50 participants. Our primary outcome was recidivism. Data from individual RCTs were combined in a random-effects meta-analysis as pooled odds ratios (ORs) and we explored sources of heterogeneity by comparing effect sizes by study size, control group, and intervention type. The protocol was pre-registered with PROSPERO, CRD42020167228.

Findings Of 6345 articles retrieved, 29 RCTs (9443 participants, 1104 [11 $\cdot$ 7%] females, 8111 [85 $\cdot$ 9%] males, and 228 [2 $\cdot$ 4%] unknown) met the inclusion criteria for the primary outcome. Mean ages were 31 $\cdot$ 4 years (SD 4 $\cdot$ 9, range 24 $\cdot$ 5–41 $\cdot$ 5) for adult participants and 17 $\cdot$ 5 years (SD 1 $\cdot$ 9; range 14 $\cdot$ 6–20 $\cdot$ 2) for adolescent participants. Race or ethnicity data were not sufficiently reported to be aggregated. If including all 29 RCTs, psychological interventions were associated with reduced reoffending outcomes (OR 0 $\cdot$ 72, 95% CI 0 $\cdot$ 56–0 $\cdot$ 92). However, after excluding smaller studies (<50 participants in the intervention group), there was no significant reduction in recidivism (OR 0 $\cdot$ 87, 95% CI 0 $\cdot$ 68–1 $\cdot$ 11). Based on two studies, therapeutic communities were associated with decreased rates of recidivism (OR 0 $\cdot$ 64, 95% CI 0 $\cdot$ 46–0 $\cdot$ 91). These risk estimates did not significantly differ by type of control group and other study characteristics.

Interpretation Widely implemented psychological interventions for people in prison to reduce offending after release need improvement. Publication bias and small-study effects appear to have overestimated the reported modest effects of such interventions, which were no longer present when only larger studies were included in analyses. Findings suggest that therapeutic communities and interventions that ensure continuity of care in community settings should be prioritised for future research. Developing new treatments should focus on addressing modifiable risk factors for reoffending.

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# Introduction

11 million people are currently held in jails or prisons worldwide and every year 30 million individuals enter and leave custody.<sup>1,2</sup> People released from jails or prisons have a higher risk of repeat offending than people given community-based sanctions, and account for nearly a fifth of all new crimes annually.<sup>3</sup> Typically, between a third and a half of people released from prison reoffend within 2 years.<sup>4</sup> The societal costs of recidivism are considerable, and include public health and associated economic effects. For example, the annual social and economic cost of reoffending is estimated at more than  $\pounds$ 18·1 billion in the UK and US\$13 billion in one US large state (Illinois) alone.<sup>5,6</sup>

Various psychological interventions have been used in custodial settings to improve outcomes for people released from prison, and to reduce reoffending in particular. Some reviews suggested that cognitive





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# **Research in context**

# Evidence before this study

We searched the Cochrane Central Register of Controlled Trials, EMBASE, Global Health, MEDLINE, PsycINFO from database inception to Feb 17, 2021, for systematic reviews and metaanalyses of the effectiveness of psychological interventions delivered in prisons, without language restrictions. We used similar keywords across databases relating to psychological interventions (eg, program\*, intervention\*, treatment\*), incarceration (eq, prison\*, incarcerat\*, custod\*), and recidivism (eq, recommit\*, reoffend\*, recidiv\*). We identified several relevant systematic reviews, but none provided a comprehensive overview of the evidence base, as their scope was limited to specific groups of individuals (eg, people with co-occurring mental illness or people in specific offence categories), or certain types of intervention (eq, CBT). Furthermore, previous reviews have included studies using non-experimental designs, which are liable to overestimate effects. Despite this limitation, these reviews stated that some psychological interventions (eq, CBT and risk-need-responsivity therapies) are effective in reducing recidivism on release from prison.

# Added value of this study

We did a comprehensive systematic review and meta-analysis of all randomised controlled trials that evaluated the

behavioural therapy (CBT) programmes are among the most effective interventions, with meta-analyses reporting recidivism risk reductions of 20-30%.7-13 Furthermore, treatment programme adherence to risk-needresponsivity principles14 is associated with reductions in reoffending; however, this link is based on predominantly quasi-experimental studies.15-17 Overall, the effectiveness of most prison-based treatments on recidivism remains unclear because the evidence is inconsistent and subject to a range of limitations.<sup>18-28</sup> Previous reviews have often focused on specific groups-eg, women,26,29 adolescents,<sup>20,23</sup> individuals who use drugs,<sup>25</sup> people living with a mental health condition,<sup>18</sup> and people with sexual<sup>21,28</sup> or other violent<sup>19,27</sup> index offences. There are considerable methodological differences between these reviews, particularly in the quality of included primary studies,20 and the sources of this heterogeneity have rarely been examined.18 Also, existing reviews have pooled estimates that combine samples from diverse settings (eg, prisons and secure psychiatric hospitals)24 or were published before 2008.<sup>19,22,23,29</sup> To address these limitations, we aimed to synthesise reoffending outcomes from all randomised controlled trials (RCTs) of psychological interventions provided in prisons.

# **Methods**

See Online for appendix

# Search strategy and selection criteria

For this systematic review and meta-analysis, we searched Cochrane Central Register of Controlled Trials,

effectiveness of psychological interventions delivered in prisons on recidivism outcomes after release. We provide an up to date systematic review, which is both broader in scope (by including all prisoners irrespective of criminal history, setting, or psychological treatment) and more precise (by including only randomised controlled trials) than previous reviews. The effects were considerably smaller than expert opinion had previously maintained, with no clear effects of CBT-based treatments.

# Implications of all the available evidence

Psychological treatments, which were developed to treat mental health conditions, need to be adapted to target modifiable risk factors that are specific to reoffending. Continued treatment after prison release should be integrated into therapeutic programmes. The evidence is inconclusive for most psychological interventions, and the findings of this systematic review could inform how different treatment modalities should be prioritised in service development and future trials.

Embase, Global Health, MEDLINE, PsycINFO, and Google Scholar for RCTs published from database inception until Feb 17, 2021. The search strategy combined terms relating to RCTs (ie, random\*, trial\*, placebo\*), psychological interventions (eg, program\*, intervention\*, treatment\*), incarceration (eg, prison\*, incarcerat\*, custod\*), and recidivism (eg, recommit\*, reoffend\*, recidiv\*). For the full list of search terms see appendix pp 3–7. We also manually searched the reference lists of included studies, and relevant articles and systematic reviews.

We included RCTs of psychological interventions in jails and prisons that reported on criminal recidivism occurring after release from prison as an outcome. Studies were eligible for inclusion if they met the following criteria: RCT (including pilot studies and cluster-randomised trials); all participants were incarcerated at the time of random allocation (including adolescents, people in custody awaiting trial, and people residing in immigration detention centres) and remained incarcerated for the duration of the treatment; participants assigned to control groups were exposed to the usual intervention, no intervention, or an alternative intervention to the experimental group; intervention was psychological (eg, CBT or mindfulness-based therapy) or psychoeducational (eg, vocational or educational training); interventions (both individual and group formats) were delivered in a jail or prison setting; and the recidivism outcome (eg, reconviction, reincarceration, rearrest, parole

and considered for inclusion. One author (GB) did the searches and screened the titles and abstracts of the studies identified using the search strategy and screened the full text of those matching the predetermined inclusion criteria. In cases of uncertainty, GB consulted with RY and consensus was reached about study selection. SF resolved any disagreements about inclusion and verified the eligibility of included studies. GB extracted summary estimates from eligible RCTs. This systematic review was done in accordance with the Preferred Items for Systematic Reviews and Meta-Analyses guidelines<sup>34</sup> (appendix pp 1–2). Data analysis We extracted from eligible studies information for: year of publication; geographical location; correctional setting; sample size; sex; ethnicity (Asian, Black or African American, White, Hispanic or Latinx, Indigenous, and

violation, or new charges) was reported separately for the

intervention and control groups. We included studies in

which post-prison services were offered to participants on

a voluntary basis, but were not directly part of the evaluated

intervention (eg, the Challenge to Change,<sup>30</sup> and the Amity

therapeutic community programmes<sup>31</sup>). We excluded

studies on the basis of the following criteria: trial not

randomised (eg, case studies and pretest-post-test

comparisons); participants were not in jail or prison at the

time of the study (eg, they were on parole, in a secure

psychiatric hospital, attending therapies outside of the

prison setting, or residing in community-based special

residential units formerly known as bootcamps); the

control group included primarily people who dropped out

or refused treatment altogether; the intervention was based

solely on a pharmacological approach; and the study

compared jail or prison with a community sanction

(eg, prison vs bootcamp) or involved a joint prison and

community programme for which the community

component accounted for more than half of the

intervention duration (eg, the CREST programme<sup>32,33</sup>).

There was no limit on the follow-up time period for

reoffending. Non-English language studies were translated

Other); average age of participants; follow-up period for recidivism; intervention length, type, and format; definition of recidivism; and numbers of individuals in the intervention and control groups by recidivism status (ie, having reoffended vs not having reoffended). If there were multiple assessments of recidivism in a study, we used the most serious outcome for the meta-analysis (eg, reconviction was preferred to rearrest). For samples that featured both males and females but for which the recidivism outcome was not reported separately by sex, those including at least 90% males were recorded as males, whereas those with fewer than 90% males were recorded as both. If multiple articles were available for a given study (eg, the Amity therapeutic community programme<sup>35,36</sup>), we included the article with the longest follow-up period for recidivism.32 We contacted relevant study authors if additional data or clarifications were required.

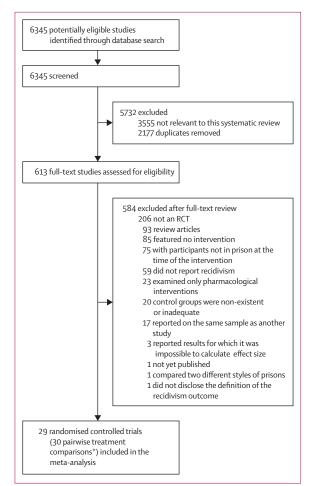
The quality of RCTs was assessed using the Cochrane Collaboration's risk-of-bias tool for randomised trials (RoB 2). Each RCT was given an overall estimation of risk of bias (ie, low risk, some concerns, or high risk) according to the following domains for risk of bias: randomisation process; deviations from intended interventions; missing outcome data; measurement of the outcome; and selection of the reported result.<sup>37</sup> Trials with a high risk of bias in at least one domain were rated as having a high risk of bias.

The primary outcome was recidivism. This measure was assessed with the summary odds ratio (OR) and corresponding 95% CI. We sought both continuous and dichotomous data on recidivism. To enable comparison across studies, when the outcome was given as continuous data, we first attempted to obtain the equivalent dichotomous data from the authors of the primary studies. If we were unable to do so, we converted the standardised mean difference to ORs (using the formula recommended by the Cochrane Handbook<sup>38</sup>). One study was excluded because of insufficient information.39 Furthermore, for multiarm trials,40,41 two distinct approaches recommended by the Cochrane Handbook<sup>38</sup> were used to avoid double-counting participants in the shared control group. For one study,40 we merged both intervention arms into a single comparison group, as they both were psychoeducational interventions. For another study,41 we included each pairwise comparison separately (one was psychoeducational and the other CBT-based) by evenly dividing the shared control group among the comparisons.

We did a random-effects meta-analysis to estimate the effect sizes, because this gives similar weights to studies with different sample sizes and substantial heterogeneity was expected between studies (eg, for type and length of interventions and follow-up periods). Pooled OR estimates were grouped into domains and summarised using forest plots. Between-study heterogeneity was estimated using Cochran's *Q* (reported with a  $\chi^2$ -value and p value) and the *I*<sup>2</sup> statistic. Amounts of heterogeneity were evaluated according to thresholds: low (0–40%), moderate (30–60%), substantial (50–90%), and considerable (75–100%).<sup>38</sup> These heterogeneity measures should be interpreted with caution if the number of studies is small (eg, in subgroup analyses).<sup>42</sup>

We first pooled all individual RCTs to calculate the summary effect size. We then stratified studies according to whether the psychological intervention group was larger than 50 participants.<sup>31,32,43–56</sup> This cutoff was determined in accordance with previous research on randomised experiments (eg, psychotherapy for adult depression<sup>57</sup>) to maximise the key beneficial effect of randomisation (ie, controlling for unknown and unmeasurable variables<sup>58,59</sup>), and rule out potential small-study effects.<sup>60</sup> Among these studies, we explored the

For more on the **Cochrane Collaboration's risk-of-bias tool for randomised trials (RoB 2)** see https://methods.cochrane. org/bias/resources/rob-2revised-cochrane-risk-bias-toolrandomized-trials



## Figure 1: Study selection

\*The 29 randomised controlled trials, included 27 RCTs that were two-arm trials and two that were three-arm trials.<sup>4041</sup> Overall, the trials described 31 psychological interventions that were combined into 30 pairwise treatment comparisons on which the statistical analyses were based.

effects of control group (ie, usual care, wait-list, and other) and intervention type (ie, CBT-based, psychoeducational, therapeutic communities, and other), and excluded two studies43,56 from the secondary analysis on the basis of considerable differences in treatment duration (eg, one session only)56 and delivery mode (eg, video feedback of previous sessions43). All interventions based on cognitive behavioural approaches were considered to be CBT-based psychological interventions.44-47,49,55 Interventions with a core vocational or educational component (eg, deterrence<sup>51</sup>) were included in the psychoeducational category.50 Interventions of therapeutic communities formed another category.<sup>30,31</sup> Both therapeutic community trials included voluntary postprison services. Most (83%) participants from the Challenge to Change trial<sup>30</sup> chose to access communitybased mental health or substance abuse services, although these were beyond the scope of that study. The Amity therapeutic community offered residential treatment to programme graduates (experimental group only) at an Amity-operated facility called Vista.<sup>31</sup> The effect of Vista on recidivism was not considered in our metaanalysis, to avoid annulling the effects of randomisation; however, we reported percentages in the Discussion. The other intervention category combined reality therapy,<sup>48</sup> social therapy,<sup>5</sup> interactive journaling,<sup>54</sup> and genderresponsive substance abuse therapy.<sup>52</sup>

Prespecified subgroup (mixed-effects) and metaregression analyses were done to examine sources of heterogeneity. The following study characteristics were assessed: year of publication (<1990  $vs \ge 1990$ ; to account for the formalisation of the risk-need-responsivity model in 1990),<sup>14</sup> study location (USA vs elsewhere), sample size (as a continuous variable), sex (sex-specific interventions vs those delivered to both males and females simultaneously), mean participant age (as a continuous variable), age group (adolescents vs adults), intervention type (CBT-based vs all other types), comparator type (usual care vs waitlist or other), follow-up time period (as a continuous variable), intervention format (individual vs group or combination), intervention aimed at substance use disorder (as a dichotomous variable) and risk of bias (high vs low or some concerns).

We did influence analysis on all studies to determine which of them disproportionately influenced the summary effect of our meta-analysis. We used the leave-one-out method and showed results using the Baujat plot.<sup>61</sup>

We examined publication bias in all studies using the Egger's test of the intercept<sup>62</sup> and funnel plot analysis. If the Egger's test reported publication bias and betweenstudy heterogeneity was not substantial,<sup>63</sup> we followed the trim and fill procedure<sup>64</sup> to correct for publication bias by imputing missing studies into a new symmetrical funnel plot.<sup>65</sup>

If the results of the publication bias analysis indicated small-study effects, we did further sensitivity analyses. First, we compared the fixed-effect and random-effect estimates of the intervention effect, because a more favourable estimate in the random-effects model might indicate that interventions were more effective in smaller studies. We did an additional analysis by only including studies with an intervention group of at least 100 participants.<sup>30,31,44,6,47,49,50,53,55</sup> We did this to reduce small-study effects, and to evaluate the robustness of the findings, as small trials are susceptible to selection bias and tend to have larger treatment effects than large trials.<sup>65,66</sup> We also investigated the effect of study quality on the pooled effect size, by removing studies at high risk of bias.

All statistical analyses were done in R version 3.6.2 and R Studio version 1.4.1717.<sup>67,68</sup> The study protocol was registered with PROSPERO, CRD42020167228.

# Role of the funding source

The funders of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report.

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Follow-up period of recidivism	Mean 9.5 months (further details not reported)	1 year	1 year	77-82 months	10 years	12-15 months	≥1 year and ≤2 years	Mean 2 years (range 411-1530 days; further details not reported)	(Table 1 continues on next page)
Detailed definition of recidivism outcome	Reinstitutionalisation in any penal institution	Incarcerated at follow-up	Subsequent arrest, or charge, or both	Marginal failure (ie, return to prison for minor crime or technical violation of parole regulations) or clear recidivism (ie, return to prison for major offence)	Reincarceration for a new felony (ie, serious criminal offence) or reincarceration on felony (ie, breach of post-release supervision conditions)	Further criminal activity (ie, being charged and sentenced to incarceration in an institution)	Parole violation	Rearrest	(Table 1 contir
Duration of intervention and number or frequency of sessions	20 weeks (80 h over 60 sessions) total; twice per week group psychotrapy (1.5 h psychotrapy (1.6 h per session; plus an average of 1 h per week individual psychotherapy	8 weeks total; mean 224 h of programme sessions (further details not reported)	3 consecutive Saturday morning sessions (3 h per session)	Not reported	Mean number of therapy sessions during the first year of imprisonment: 18.6 experimental group, 4.0 control group (further details not reported)	8 sessions (further details not reported)	12 weeks total; once a week meetings (1 h per session)	Not reported	
Comparator	Notreatment	Routine institutional care	No treatment	No treatment	Standard care (control group participants were free to out therapy [group or individual] through the usual channels)	No treatment	No treatment	Routine care (eg, assignment to the first available vocational training programme or to a prison job)	
Psychological intervention; category; format	Psychotherapy; other, combination	Awareness group (with and without video feedback); psychoeducational; group	Squires programme; psychoeducational; group	Prison educational programme: psychoeducational; combination	Group therapy; other; group	Social interaction skills programme and stress management training programme; psychoed ucational; group	Cognition mediation training plus attention control; CBT-based and psychoeducational; group	Vocation delivery system; psychoeducational; group	
Mean age, years (SD)	16.4 years (SD not reported)	24·5 years (range 18–64; SD not reported)	16·3 years (range 14–18; SD not reported)	Not reported	Not reported	Mean not reported (range 15-17 years)	17.2 years (range 15–18; SD not reported)	20.0 years (SD not reported)	
Sex	Males	Males	Males	Males	Males	Males	Both (50% females, 50% males)	Males	
Participants followed up (%)	82 (100%)	128 (85%)	108 (100%)	55 (83%)	86 (93%)	42 (93%)	83 (50%)	247 (42%)	
Participants randomly allocated	82	150	108	66	92	45	165	591	
Setting	Institution for boys	Minimum- security institution	Four camps	Two penitentiaries (maximum and medium security)	Prison	Institution for incarcerated delinquents	Juvenile correctional facility	Prison	
Country	USA	Canada	USA	Canada	USA	Canada	USA	USA	
	Persons (1967)≌	Annis (1979) <sup>43</sup>	Lewis (1983) <sup>51</sup>	Linden et al (1984) <sup>22</sup>	Homant (1986) <sup>21</sup>	Shivrattan (1988)⁴	Guerra and Slaby (1990) <sup>41</sup>	Lattimore et al (1990) <sup>ss</sup>	

20.2 years Moral reconation (range 15-22; therapy, CBT-based; SD 1-0) group 30-7 years Amity therapeutic community programme; therapeutic communities; group 34.3 years Prison modified (SD 8.8) plus afterare;		576 (81%) Males 107 (45%) Males	715 576 (81%) 236 107 (45%)	576 (81%) 107 (45%)
pe	(5D 8.8) (SD 8.8) Not reported	Males	eanon (n C+) (u.	Prison (A. CF) (A. CF) (A. CC) Males

Follow-up period of recidivism		6 months	1 year	1 year	1 year	Mean 518 days (SD 264)	Mean 3.6 years (range 0.1–5.8)	1 year	1 year	mber of post- 1 year rests
Detailed definition of recidivism outcome		Reincarceration	Reincarceration	Being booked (ie, processed after arrest) in the county jail	Reincarceration	Reconviction	Reincarceration	Reincarceration	Reincarceration	Mean number of post- release arrests
Duration of intervention and number or frequency of sessions		6–8 weeks tota!; 90 min sessions, typically 3 times per week	Helping Women Recover (17 sessions) and Beyond Trauma (11 sessions)	Not reported	Planned 6 months tenure; programme activities were provided 4 h per day, 5 days per week	4 weeks total; 10 sessions; approximately 20 h of group treatment and ≥4 h of individual support	3 months; six personalised feedback letters; letter sent twice per month	12 weeks total; 48 interactive, multimedia modules; once a week for 2 h or twice per week for 1 h (depending on laboratory availability)	20 sessions; 40 h total	12 weeks total, 2.5 h sessions, three times per week
Comparator		Treatment as usual (similar to other US state prison programmes for substance users)	Standard prison therapeutic community programme	Placebo (government booklet on substance misuse disorders and criminal behaviour)	CBT-based intervention for substance misuse	Treatment as usual	No treatment	Standard care	Treatment as usual	Services as usual
Psychological intervention; category; format		Seeking Safety plus treatment as usual; CBT-based; group	Gender responsive therapy using manualised curricula (Helping Women Recover; Beyond Trauma); other; group	Interactive journalling; other, individual	Challenge to change therapeutic community; therapeutic communities; group	Control of violence for angry, impulsive drinkers plus treatment as usual; CBT-based; group	Personalised feedback intervention; other; individual	Experimental condition therapeutic education system; CBT-based; individual	Beyond violence; other; group	Parent management training CBT-based; group
Mean age, years (SD)		34.6 years (SD 7.4)	35-9 years (SD 9-6)	36.6 years (SD 11.1)	35-1 years (SD 7-9)	24-5 years (SD 5-7)	41·5 years (SD 10·5)	36-6 years (SD 9-6)	33-7 years (SD 8-9)	31.4 years (SD not reported)
Sex		Females	Females	Males	Females	Males	Males	Both (31.4% females, 69.6% males)	Females	Both (55% females, 45% males)
Participants followed up (%)		44 (90%)	115 (100%)	183 (99%)	370 (79%)	109 (95%)	50 (100%)	482 (98%)	35 (83%)	359 (100%)
Participants randomly allocated		49	115	185	468	115	50	494	42	359
Setting	us page)	Residential substance abuse treatment programme in a minimum security wing of a women's prison	Women's prison	Jail	Women's correctional facility	Two medium- security prisons	Prison	Ten prisons	Prison for women	Four US state correctional facilities (releasing institutions)
Country	om previot	USA	USA	USA	USA	Х	Japan	USA	NSA	USA
	(Continued from previous page)	Zlotnick et al (2009) <sup>35</sup>	Messina et al (2010)≌	Proctor et al (2012) <sup>54</sup>	Sacks et al (2012) <sup>20</sup>	Bowes et al (2014) <sup>45</sup>	Yokotani and Tamura (2015) <sup>22</sup>	Chaple et al (2016) <sup>47</sup>	Kubiak et al (2016) <sup>28</sup>	Burraston and Eddy (2017) <sup>46</sup>

	Country	Country Setting	Participants randomly allocated	Participants followed up (%)	Sex	Mean age, years (SD)	Psychological intervention; category; format	Comparator	Duration of intervention and number or frequency of sessions	Detailed definition of Follow-up recidivism outcome period of recidivism	Follow-up period of recidivism
(Continued from pre Malouf et al USA (2017) <sup>33</sup>	(Continued from previous page) Malouf et al USA Jail (2017) <sup>32</sup>	us page) Jail	49	31 (63%)	Males	37.2 years (range 18–81; SD 15.7)	Re-entry values and mindfulness programme plus treatment as usual; other, group	Treatment as usual	Treatment as usual 4 weeks total; 90 min sessions, twice per week	Rearrest	3 years
Gold et al (2020) <sup>ଛ୍ଡ</sup>	Norway	Prison	66	64 (96%)	Males	Median 26 years (range 18–53; SD not reported)	Music therapy; other; usually group but in some cases individual	Standard care	Mean 4.4 (range 0-12; SD 3-9); median 3-0), typically two to three times per week	Serious events, excluding writs	5 years
Hein et al (2020) <sup>49</sup> Data are n (%)	USA or mean (SD),	Hein et al USA Juvenile justice 289 289 (100%) Males (2020) <sup>46</sup> setting Data are n (%) or mean (5D), unless otherwise specified. CBT=cognitive behavioural therapy.	289 ecified. CBT=cogr	289 (100%) Males itive behavioural therap	Males therapy.	14-9 years (SD 1-0)	Training on solving social problems; CBT- based; group	Treatment as usual	Treatment as usual 10 sessions each lasting At least one offence 1 h during follow-up	At least one offence during follow-up	2 years
Table 1: Chara	acteristics of	Table 1: Characteristics of randomised controlled trials of psychological interventions in prison to reduce recidivism	rolled trials of <b>p</b>	sychological int	terventions in pris	on to reduce red	cidivism				

# Results

We identified 6345 articles through electronic searches and 29 eligible trials (for selection process see figure 1 and for study characteristics see table 1).<sup>30,31,40,41,43-56,69-75,77-80</sup> Most RCTs were two-arm trials (n=27); two were three-arm trials.40,41 These trials described 31 psychological interventions that were combined into 30 pairwise treatment comparisons, on which the statistical analyses were based. In total, 9443 individuals (1104 [11.7%] females, 8111 [85.9%] males, and 228 [2.4%] individuals for whom sex was not reported) participated in the trials, and 6528 (1118 [17.1%] adolescents and 5410 [82.9%] adults) had recidivism outcome data. The mean age was 31.4 years (SD 4.9, range 24.5-41.5) in adults and 17.5 years (1.9, 14.6-20.2) in adolescents. Descriptive statistics on the age of participants were calculated using the mean age from each study and the range of mean ages (if available). Race or ethnicity data from each study are summarised in the appendix (pp 8-9). Among included trials, 19 were from the USA (n=3578 [54.8%]),  $^{30,31,41,44,46-52,54,69,71,72,74,75,78,79}$ four from Canada (n=2351),<sup>40,43,55,70</sup> two from the UK (n=203);<sup>45,56</sup> and one each from Germany (n=223),<sup>53</sup> Sweden (n=59),73 Japan (n=50),77 and Norway (n=64).80 Treatment duration varied considerably between trials, ranging from one session only<sup>56</sup> to multiple interventions that lasted for 1 year.<sup>31,74</sup> The most frequent source of trial funding was government-funded research council. None of the psychological interventions was described as being mandatory and recruitment of participants was voluntary. However, it is possible that perceived coercion and other incentives could have contributed to the decision to participate.

In terms of risk of bias, most RCTs were rated as having concerns (n=18, 60%) or being at high risk (n=10, 33%), and only two studies46.54 were rated as having a low risk of bias (appendix pp 10–12). There was a low risk of bias in outcome measurement for all studies, because recidivism was ascertained from official criminal records.

Overall in the meta-analysis, psychological interventions were associated with reduced reoffending, with a pooled OR of 0.72 (95% CI 0.56-0.92) and moderate levels of heterogeneity (I2=49%; Q=57.3; p<0.01; figure 2). To prevent overestimation caused by small-study effect, as suggested by the literature65,66 and confirmed by our influence analysis, we pooled results excluding studies with fewer than 50 participants in the experimental group, as a planned sensitivity analysis. The reduction in recidivism was attenuated in the 14 trials (6446 followedup participants) with an intervention group of at least 50 participants (OR 0.87, 95% CI 0.68-1.11; I<sup>2</sup>=54%; figure 3).

Subgroup analyses are shown by comparator type in figure 4, and by intervention type in figure 5. RCTs with a control group of usual care were associated with recidivism but not significantly so (OR 0.97, 95% CI 0.70-1.34; I<sup>2</sup>=59%). If using waiting list (0.74, 0.56-0.99; 17%) or other interventions (0.64, 0.40-1.01;

	Intervention (n/N)	Control (n/N)	Weight	Odds ratio (95% CI)
Persons (1967) <sup>69</sup>	13/41	25/41	3.2%	0.30 (0.12-0.74)
Annis (1979)43	24/85	14/43	3.6%	0.82 (0.37–1.80)
Lewis (1983) <sup>51</sup>	43/53	37/55	3.3%	2.09 (0.86–5.09)
Linden (1984) <sup>70</sup>	20/30	20/26		0.60 (0.18–1.97)
Homant (1986) <sup>71</sup>	11/43	7/43	2.8%	1.77 (0.61–5.11)
Shivrattan (1988)40	13/27	9/15	2.3%	0.62 (0.17-2.23)
Lattimore et al (1990)50	50/138	50/109	4.6%	0.67 (0.40-1.12)
Guerra and Slaby (1990a)41	10/29	5.5/12	2.1%	0.62 (0.16-2.45)
Guerra and Slaby (1990b)41	12/28	5.5/12	2.1%	0.89 (0.23-3.46)
Leeman (1993)72	3/20	15/37	2.0%	0.26 (0.06–1.04)
Robinson (1995)55	371/1746	94/379	5.3%	0.82 (0.63–1.06)
Lindforss (1997)73	18/30	25/29	2.3%	0.24 (0.07-0.87)
Dugan and Everett (1998) <sup>48</sup>			4.1%	1.41 (0.73–2.73)
Ortmann (2000)53	67/111	76/112	4.4%	0.72 (0.42-1.25)
Armstrong (2003)44	71/110	66/102	4.4%	0.99 (0.57-1.75)
Prendergast et al (2004) <sup>31</sup>	258/341	196/235	4.9%	0.62 (0.41-0.95)
Sacks et al (2004) <sup>74</sup>	2/43	21/64	1.8%	0.10 (0.02-0.45)
Shapland (2008)56	18/52	16/42	3.4%	0.86 (0.37-2.00)
Zlotnick (2009)75	5/23	9/21	2.2%	0.37 (0.10-1.38)
Messina et al (2010)52	19/60	25/55	3.7%	0.56 (0.26–1.19)
Sacks et al (2012) <sup>30</sup>	27/207	29/163	4.4%	0.69 (0.39–1.23)
Proctor et al (2012)54	50/98	56/85	4.3%	0.54 (0.30-0.98)
Bowes et al (2014)45	27/52	41/57	3.6%	0.42 (0.19-0.93)
Yokotani (2015) <sup>77</sup>	5/20	12/30	2.3%	0.50 (0.14–1.74)
Kubiak (2016) <sup>78</sup>	3/19	8/16	1.7%	0.19 (0.04–0.91)
Chaple et al (2016)47	67/244	51/238	4.9%	1.39 (0.91–2.11)
Malouf (2017) <sup>79</sup>	10/16	12/15	1.7%	0.42 (0.08-2.11)
Burraston and Eddy (2017) <sup>46</sup>			5.0%	0.97 (0.67–1.42)
Gold (2020) <sup>80</sup>	13/32	10/32	2.9%	1.50 (0.54-4.20)
Hein et al (2020) <sup>49</sup>	98/118	126/171	4.3%	1.75 (0.97-3.15)
Random effects model			100.0%	0.72 (0.56-0.92)
Heterogeneity: I <sup>2</sup> =49%, p<0·02	L	-	0.1 0.5 1.0 2.0 10.0	
			Decreased risk of recidivism	

Figure 2: Effectiveness of psychological interventions in prison in reducing recidivism

Data are for all 29 included randomised controlled trials. Error bars show 95% CI. The number of participants in the intervention and control groups were not available for Dugan and Everett<sup>48</sup> or Burraston and Eddy<sup>46</sup> because these studies presented outcomes as continuous rather than dichotomous data.

0%), the reduction in recidivism was larger although CIs were overlapping. By treatment modality, CBT-based interventions were not associated with recidivism (1·00, 0·69–1·44; 60%) neither were psychoeducational interventions (1·11, 0·38–3·20; 79%). Other types of interventions were associated with non-significant reductions in recidivism (0·74, 0·47–1·18; 44%). However, there were reductions in reoffending risk for therapeutic community programmes (0·64, 0·46–0·91; 0%).

On univariate analyses, there was a statistically significant difference between the pooled effects of trials which included sex-specific samples compared with trails that included both males and females (Q 4·30; p=0.04). Sex-specific interventions were significantly associated with reduced recidivism (OR 0.67, 95% CI 0.50–0.90), whereas those including both males and females were not (1.09, 0.77–1.55). No other significant associations were found between prespecified study

characteristics and effect sizes in subgroup or metaregression analyses (table 2).

Two studies<sup>49,74</sup> that contributed disproportionately to the pooled effect were identified using influence analyses in all RCTs. Removal of these outliers reduced the degree of heterogeneity between studies from moderate ( $I^2$ =49%) to low (38%) but did not materially alter the pooled effect size (OR 0.73, 95% CI 0.58–0.91; appendix pp 13–15).

We found evidence of publication bias using Egger's test (t=-2·12; p=0·04) suggesting small-study effects. This finding was supported by visual inspection of the related funnel plot, which showed asymmetry (appendix pp 16–17). Seven smaller studies were identified and trimmed using the trim and fill method,<sup>40,69,72-75,78</sup> and the OR after adjusting for publication bias was 0·86 (95% CI 0·65–1·15).

The fixed-effect estimate (OR 0.81, 95% CI 0.72-0.91; *I*<sup>2</sup>=49%; appendix p 18) did not materially differ from the

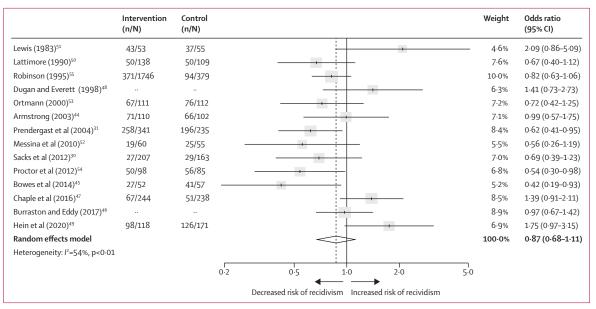


Figure 3: Effectiveness of psychological interventions in prison in reducing recidivism

Data are for the 14 randomised controlled trials with an intervention group of at least 50 participants, excluding two outlier studies.<sup>4356</sup> Error bars show 95% CI. The number of participants in the intervention and control groups were not available for Dugan and Everett<sup>48</sup> or Burraston and Eddy<sup>46</sup> because these studies presented outcomes as continuous rather than dichotomous data.

random-effects model. Repeating the meta-analysis and only including larger studies (ie,  $\geq$ 100 participants in the psychological intervention group) resulted in a decrease of the strength of the association to OR 0.90 (0.71–1.14; appendix p 19).<sup>60</sup>

# Discussion

In this meta-analysis of psychological interventions for recidivism, we identified 29 jail-based or prison-based RCTs of 9443 individuals from seven countries. Overall, there was evidence of reduced odds of reoffending. To account for small-study effects, in a planned sensitivity analysis, we excluded studies with fewer than 50 people in each experimental arm, resulting in 14 trials with 6446 followed-up participants, and the overall pooled OR 0.87 (95% CI 0.68-1.11) indicated, at most, modest effects.

We report two other main findings. First, in a sensitivity analysis, we found no strong evidence of reduced reoffending after participation in CBT-based programmes in prison (OR 1.00, 95% CI 0.69-1.44;  $I^2=60\%$ ). This is by contrast with a 2007 systematic review combining both prison-based and community-based interventions that reported reduced risks of 20-30%.<sup>13</sup> One potential explanation for no clear effectiveness of such CBT interventions found in the current systematic review is that these interventions are not linked with psychosocial support upon release. It might also be that these psychological therapies, which were developed for mental health problems, do not address the accommodation, employment, and financial difficulties after release that contribute to recidivism risk.<sup>81</sup>

A second finding, from a subgroup analysis, was that participation in a therapeutic community was associated with reduced reoffending risk. However, this finding was limited to only two studies,<sup>30,31</sup> both of which linked people released from prison to voluntary post-prison services. In support of this finding, in one of the two trials, links to community services were associated with a lower return to custody rate (33 [42%] of 79) than for participants without such links (137 [86%] of 159).31 Findings from a systematic review<sup>82</sup> of psychoeducational programmes for reducing prison violence are consistent with the potential role of therapeutic communities, as programmes tailored to specific needs (eg, substance use disorder) were associated with reduced institutional violence. Similar results were reported in a Cochrane review<sup>83</sup> of any people who offended and had co-occurring drug and mental health problems, as three35,74,76 of the four included studies<sup>35,56,74,76</sup> found therapeutic communities were associated with reductions in recidivism.

There are several implications for treatments offered in prison. First, in-prison interventions might not be effective unless they are linked with interventions that target the psychosocial needs of released individuals. For example, two therapeutic community trials<sup>30,31</sup> highlighted the potential importance of community aftercare to maintain the therapeutic gains delivered in prison. Hence, psychological interventions that combine prisonbased and community-based services should be prioritised for future research. It should be noted that UK efforts to implement the Through the Gate service for resettling people released from prison have been widely criticised for inadequate communication between

	Intervention (n/N)	Control (n/N)	Weight	Odds ratio (95% CI)
Other				
Messina et al (2010)52	19/60	25/55		0.56 (0.26–1.19)
Sacks et al (2012) <sup>30</sup>	27/207	29/163	7.0%	0.69 (0.39-1.23)
Random effects model			12.5%	0.64 (0.40–1.01
Heterogeneity: /²=0%, $\chi_1^2$ =0·21	(p<0·01)			
Usual care				
Lewis (1983) <sup>51</sup>	43/53	37/55	+ 4.6%	2.09 (0.86–5.09)
Lattimore et al (1990) <sup>50</sup>	50/138	50/109	7.6%	0.67 (0.40–1.12)
Dugan and Everett (1998) <sup>48</sup>				1.41 (0.73–2.73)
Ortmann (2000)53	67/111	76/112	7.2%	0.72 (0.42–1.25)
Armstrong (2003) <sup>44</sup>	71/110	66/102	7.1%	0.99 (0.57–1.75)
Proctor et al (2012)54	50/98	56/85		0.54 (0.30–0.98)
Bowes et al (2014)45	27/52	41/57	5.2%	0.42 (0.19–0.93)
Chaple et al (2016)47	67/244	51/238	8.5%	1.39 (0.91–2.11)
Burraston and Eddy (2017)46			8.9%	0.97 (0.67–1.42)
Hein et al (2020) <sup>49</sup>	98/118	126/171	, 6.9%	1.75 (0.97-3.15)
Random effects model			69.1%	0.97 (0.70-1.34)
Heterogeneity: $l^2=59\%$ , $\chi_9^2=21$	89 (p=0·27)			
Waiting list				
Robinson (1995) <sup>55</sup>	371/1746	94/379	10.0%	0.82 (0.63–1.06)
Prendergast et al (2004) <sup>31</sup>	258/341	196/235	* 8.4%	0.62 (0.41–0.95)
Random effects model			18.4%	0.74 (0.56–0.99)
Heterogeneity: $l^2=17\%$ , $\chi_1^2=1.21$	L (p=0·65)			
Random effects model			100-0%	0.87 (0.68–1.11)
Heterogeneity: $l^2=54\%$ , $\chi^2_{13}=28$	·33 (p<0·01)		0.5 1.0 2.0 5.0	
			Decreased risk of recidivism Increased risk of recividism	

Figure 4: Effectiveness of psychological interventions in prison for reducing recidivism, by comparator type

Data are for randomised controlled trials with an intervention group of at least 50 participants, excluding two outlier studies.<sup>43,6</sup> Error bars show 95% Cl. The number of participants in the intervention and control groups were not available for Dugan and Everett<sup>48</sup> or Burraston and Eddy<sup>46</sup> because these studies presented outcomes as continuous rather than dichotomous data.

prisons and community services, and for poor assessment of resettlement needs, which should occur early in the sentence of a person in prison.<sup>84</sup>

Second, most of the tested interventions were developed in the community or in clinical populations for other outcomes, and hence might not address risk factors specific to reoffending. Such risk factors need to be identified by high quality assessment, and then linked to interventions for reducing recidivism. Risk assessments should be informed by scalable and transparent clinical prediction tools, such as the Oxford Risk of Recidivism tool (also known as OxRec),85 which includes assessment of modifiable risk factors for recidivism (eg, substance misuse and mental health status), supplemented by detailed assessments that consider additional dynamic factors. Considering that the resources allocated for interventions in prison populations are limited,<sup>86</sup> stratification of risk is necessary to guide risk management and the treatment of people on release from prison.

A third implication regards CBT. The absence of effect that we reported is different to evidence from some

reviews (including one published by the Campbell Collaboration<sup>13</sup>), which have suggested that CBT is one of the most effective forms of treatment for people in prison.<sup>7-12</sup> However, these previous reviews combined RCTs with less than rigorous study designs and the current new findings question the widespread roll-out of these treatment approaches in prisons. Only one<sup>45</sup> of the six CBT studies<sup>44–47,49,55</sup> in our systematic review reported significant reductions in reoffending. Other research, in selected populations of all people who have offended and also use drugs, also found little support for CBT.<sup>83,87</sup>

Another implication of our review is that the effects of in-prison psychological interventions on recidivism appear to be smaller than those reported in previous meta-analyses, which have been estimated to be around 0.65 (95% CI 0.57-0.75).<sup>24</sup> This difference is probably because the previous reviews included studies using weak research designs, such as quasiexperimental studies.<sup>88</sup> A similar difference has been noted for psychotherapy effectiveness in depression, whereby overall effectiveness was overestimated in

Armstrong (2003) <sup>44</sup> 71/110 66/102 71/52 41/57 52% 0-42 (0.19) Bows et al (2014) <sup>45</sup> 27/52 41/57 52% 0-42 (0.19) Chaple et al (2016) <sup>69</sup> 67/244 51/238 85% 139 (0.91) Burraston and Eddy (2017) <sup>46</sup>		Intervention (n/N)	Control (n/N)		Weight	Odds ratio (95% CI)
Amstrong (2003) <sup>44</sup> 71/10       66/102       71%       0.99 (0.57)         Bows et al (2014) <sup>45</sup> 27/52       41/57       52%       0.42 (0.19)         Chaple et al (2016) <sup>47</sup> 67/744       51/28       85%       1.39 (0.91)         Burraston and Eddy (2017) <sup>46</sup> -       89%       0.97 (0.67)         Random effects model       46.6%       1.00 (0.69         Heterogeneity: P=60%, $\chi_2^+$ =12.6 (p=0.03)       -       63%       1.41 (0.73-         Other       -       -       63%       1.41 (0.73-         Dugan and Everett (1998) <sup>46</sup> -       -       63%       1.41 (0.73-         Ortharn (2000) <sup>53</sup> 67/111       76/112       -       7.2%       0.72 (0.42-         Messina et al (2012) <sup>54</sup> 50/98       56/85       -       6.8%       0.54 (0.30)         Random effects model       -       -       -       6.8%       0.54 (0.30)         Psychoeducational       -       -       -       -       6.6%       0.670 4.0-         Random effects model       -       -       -       10.038       0.670 4.0-         Heterogeneity: P=44%, $\chi_2^+=5.32$ (p=0.15)       -       -       7.6%       0.670 4.0-	CBT-based					
Bowe set al (2014) <sup>5</sup> 27/52 41/57 T 52% 0.42 (0.19) Chaple et al (2016) <sup>17</sup> 67/244 51/338 85% 139 (0.91) Buraston and Eddy (2017) <sup>46</sup> 88% 0.97 (0.67) Random effects model Heterogeneity: $P = 60\%$ , $\chi_2^{h} = 12.6$ (p=0.03) Other Dugan and Everett (1998) <sup>46</sup> 63% 1.41 (0.73- Ortmann (2000) <sup>13</sup> 67/111 76/112 63% 0.42 (0.19) Other Dugan and Everett (1998) <sup>46</sup> 63% 0.42 (0.19) Other Dugan and Everett (1998) <sup>46</sup> 63% 0.42 (0.19) Chere Dugan and Everett (1998) <sup>46</sup> 63% 0.42 (0.19) Pychoeducational Lewis (1983) <sup>17</sup> 43/53 37/55 46/85 46.8% 0.54 (0.30) Random effects model Heterogeneity: $P = 44\%$ , $\chi_3^{h} = 5.32$ (p=0.03) Pychoeducational Lewis (1983) <sup>17</sup> 43/53 37/55 46/85 76% 0.670 40- Random effects model Heterogeneity: $P = 79\%$ , $\chi_4^{h} = 4.73$ (p=0.15) Therapeutic community Prendergast et al (2004) <sup>11</sup> 258/341 196/235 77% 0.69 (0.38) Random effects model Heterogeneity: $P = 79\%$ , $\chi_4^{h} = 4.73$ (p=0.15) Therapeutic smodel Heterogeneity: $P = 79\%$ , $\chi_4^{h} = 4.73$ (p=0.15) Therapeutic smodel Heterogeneity: $P = 0\%$ , $\chi_2^{h} = 0.1$ (p=0.76) Random effects model Heterogeneity: $P = 0.5$ , $\chi_2^{h} = 0.1$ (p=0.76) Random effects model Heterogeneity: $P = 0.5$ , $\chi_2^{h} = 0.1$ (p=0.76) Random effects model Heterogeneity: $P = 0.5$ , $\chi_2^{h} = 0.1$ (p=0.76) Random effects model	Robinson (1995)55	371/1746	94/379		10.0%	0.82 (0.63–1.06
Chaple et al (2016) <sup>17</sup> 67/244 51/238 Buraston and Eddy (2017) <sup>46</sup> Hein et al (2020) <sup>49</sup> 98/118 126/171 69% 175 097 <b>Random effects model</b> Heterogeneity: $P = 60\%$ , $\chi_5^2 = 12.6$ (p=0-03) <b>Other</b> Dugan and Everett (1998) <sup>48</sup> Chapter 41 (2012) <sup>45</sup> 67/111 76/112 72% 63% 141 (073- 72% 072 (042) Heterogeneity: $P = 60\%$ , $\chi_5^2 = 12.6$ (p=0-03) <b>Other</b> Dugan and Everett (1998) <sup>48</sup> Chapter 41 (2012) <sup>45</sup> 50/98 56/85 546% 054 (0-30 <b>Random effects model</b> Heterogeneity: $P = 44\%$ , $\chi_5^2 = 52$ (p=0-03) <b>Psychoeducational</b> Lewis (1983) <sup>12</sup> 43/53 37/55 46% 057 040- <b>Random effects model</b> Heterogeneity: $P = 79\%$ , $\chi_1^2 = 473$ (p=0-15) <b>Therapeutic community</b> Prendergast et al (2001) <sup>12</sup> 25/8/341 196/235 70% 0667 049 <b>Random effects model</b> Heterogeneity: $P = 79\%$ , $\chi_1^2 = 473$ (p=0-15) <b>Therapeutic smodel</b> Heterogeneity: $P = 79\%$ , $\chi_1^2 = 473$ (p=0-15) <b>Therapeutic smodel</b> Heterogeneity: $P = 79\%$ , $\chi_1^2 = 473$ (p=0-15) <b>Therapeutic smodel</b> Heterogeneity: $P = 0\%$ , $\chi_1^2 = 0.1$ (p=0-76) <b>Random effects model</b> Heterogeneity: $P = 0\%$ , $\chi_1^2 = 0.1$ (p=0-76) <b>Random effects model</b> Heterogeneity: $P = 0\%$ , $\chi_1^2 = 0.1$ (p=0-76)	Armstrong (2003)44	71/110	66/102		7.1%	0.99 (0.57–1.75
Burraston and Eddy (2017) <sup>46</sup>	Bowes et al (2014) <sup>45</sup>	27/52	41/57 —		5.2%	0.42 (0.19-0.93
Hein et al (2020) <sup>49</sup> 98/118 126/171 Random effects model Heterogeneity: $l^{2}=60\%$ , $\chi_{2}^{2}=12.6$ (p=0-03) Other Digan and Everett (1998) <sup>46</sup>	Chaple et al (2016)47	67/244	51/238	· · · · · · · · · · · · · · · · · · ·	8.5%	1.39 (0.91–2.11
Random effects model       46.6% $1.00 (0.69)$ Heterogeneity: $l^{2}=60\%$ , $\chi_{2}^{2}=12.6 (p=0.03)$ 63% $1.41 (0.73)$ Other       72% $0.72 (0.42)$ Messina et al (2000) <sup>13</sup> $67/111$ $76/112$ 72%         Messina et al (2010) <sup>12</sup> $19/60$ $25/55$ $5.5\%$ $0.56 (0.26)$ Proctor et al (2012) <sup>24</sup> $50/98$ $56/85$ $6.8\%$ $0.54 (0.30)$ Random effects model       25.8% $0.74 (0.47)$ Heterogeneity: $l^{2}=44\%$ , $\chi_{2}^{2}=5.32 (p=0.03)$ $7.6\%$ $0.67 0.40-7$ Psychoeducational       25.8% $0.74 (0.47)$ $1.00.0\%$ $8.4\%$ $0.62 (0.41)$ Heterogeneity: $l^{2}=79\%$ , $\chi_{1}^{2}=4.73 (p=0.15)$ $1.00.0\%$ $8.4\%$ $0.62 (0.41)$ Therapeutic community       7.70% $0.69 (0.39)$ $7.6\%$ $0.60 (0.39)$ Random effects model       15.5% $0.64 (0.46)$ $1.55\%$ $0.64 (0.46)$ Heterogeneity: $l^{2}=0.0(p=0.76)$ 100.0% $0.87 (0.68)$ $0.00.0\%$ $0.87 (0.68)$	Burraston and Eddy (2017) <sup>46</sup>				8.9%	0.97 (0.67–1.42
Heterogeneity: $l^2$ =60%, $\chi_2^2$ =12.6 (p=0.03) Other Dugan and Everett (1998) <sup>46</sup> 6-3% 1.41 (0.73- Ortmann (2000) <sup>53</sup> 67/111 76/112 7.2% 0.72 (0.42- Messina et al (2010) <sup>52</sup> 19/60 25/55 6-8% 0.56 (0.26 Proctor et al (2012) <sup>54</sup> 50/98 56/85 6-8% 0.54 (0.30) Random effects model 25.8% 0.74 (0.47 Heterogeneity: $l^2$ =44%, $\chi_3^2$ =5-32 (p=0.03) Psychoeducational Lewis (1983) <sup>53</sup> 43/53 37/55 4-6% 2.09 (0.86 Lattimore et al (1990) <sup>50</sup> 50/138 50/109 7-6% 0.67 0.40- Random effects model 12.2% 1.11 (0.38 Heterogeneity: $l^2$ =79%, $\chi_1^2$ =4.73 (p=0.15) Therapeutic community Prendergast et al (2004) <sup>31</sup> 258/341 196/235	Hein et al (2020) <sup>49</sup>	98/118	126/171	· · · · · · · · · · · · · · · · · · ·	6.9%	1.75 (0.97–3.15
Other       63% $1.41(0.73-10.72\%)^{-1}$ Dugan and Everett (1998) <sup>48</sup> 72% $0.72(0.42-10.72\%)^{-1}$ Ottmann (2000) <sup>53</sup> $67/111$ $76/112$ 72%         Messina et al (2010) <sup>52</sup> $19/60$ $25/55$ $5.\%$ $0.56(0.26)$ Proctor et al (2012) <sup>54</sup> $50/98$ $56/85$ $6.8\%$ $0.54(0.30)$ Random effects model       25.8% $0.74(0.47)$ Heterogeneity: $l^2 = 44\%$ , $\chi_3^2 = 5.32(p=0.03)$ $25.8\%$ $0.74(0.47)$ Psychoeducational       25.8% $0.74(0.47)$ Lewis (1983) <sup>51</sup> $43/53$ $37/55$ $4.6\%$ $2.09(0.86)$ Lattimore et al (1990) <sup>50</sup> $50/138$ $50/109$ $7.6\%$ $0.670.40-7$ Random effects model       12.2% $1.11(0.38)$ $1.22\%$ $1.11(0.38)$ Heterogeneity: $l^2 = 79\%$ , $\chi_1^2 = 4.73$ (p=0.15) $7.0\%$ $0.69(0.39)$ $7.0\%$ $0.69(0.39)$ Random effects model       196/235 $7.0\%$ $0.69(0.39)$ $7.0\%$ $0.69(0.39)$ Random effects model       15.5% $0.64(0.46)$ $15.5\%$ $0.64(0.46)$ Heterogeneity: $l^2 = 5.4\%$ , $y^2 = 0.1(p =$	Random effects model				46.6%	1.00 (0.69–1.4
Dugan and Everett (1998) <sup>44</sup>	Heterogeneity: $l^2=60\%$ , $\chi_5^2=12$	·6 (p=0·03)				
Ortmann (2000) <sup>53</sup> $67/111$ $76/112$ $72\%$ $0.72$ (0.42-         Messina et al (2010) <sup>52</sup> $19/60$ $25/55$ $55\%$ $0.56$ (0.26         Proctor et al (2012) <sup>54</sup> $50/98$ $56/85$ $6.8\%$ $0.54$ (0.30-         Random effects model       25.8% $0.74$ (0.47         Heterogeneity: $l^2=44\%$ , $\chi_2^2=5.32$ (p=0-03) $25.8\%$ $0.74$ (0.47         Psychoeducational       25.8% $0.74$ (0.47         Lewis (1983) <sup>51</sup> 43/53 $37/55$ $4.6\%$ $2.09$ (0.86         Lattimore et al (1990) <sup>50</sup> $50/138$ $50/109$ $7.6\%$ $0.67$ 0.40-         Random effects model       12.2%       1.11 (0.38         Heterogeneity: $l^2=79\%$ , $\chi_1^2=4.73$ (p=0-15) $7.6\%$ $0.62$ (0.41-         Sacks et al (2012) <sup>30</sup> 27/207       29/163 $7.0\%$ $0.69$ (0.39         Random effects model       15.5% $0.64$ (0.46 $15.5\%$ $0.64$ (0.46         Heterogeneity: $l^2=54\%$ , $\chi_1^2=23.3$ (p=0-01) $100.0\%$ $0.87$ (0.68	Other					
Messina et al $(2010)^{52}$ 19/60       25/55         Proctor et al $(2012)^{54}$ 50/98       56/85         Random effects model       25.8%       0.74 (0.47)         Heterogeneity: $l^2$ =44%, $\chi_3^2$ =5.32 (p=0.03)       25.8%       0.74 (0.47)         Psychoeducational       25.8%       0.74 (0.47)         Lewis (1983) <sup>51</sup> 43/53       37/55         Lattimore et al (1990) <sup>50</sup> 50/138       50/109 <b>Random effects model</b> 12.2%       1.11 (0.38)         Heterogeneity: $l^2$ =79%, $\chi_1^2$ =4.73 (p=0.15)       8.4%       0.62 (0.41-         Therapeutic community       70%       0.69 (0.39)         Random effects model       15.5%       0.64 (0.46)         Heterogeneity: $l^2$ =0.1 (p=0.76)       100.0%       0.87 (0.68)         Random effects model       100.0%       0.87 (0.68)	Dugan and Everett (1998)48				6.3%	1.41 (0.73–2.73
Proctor et al $(2012)^{54}$ 50/98 56/85 Random effects model Heterogeneity: $l^2 = 44\%$ , $\chi_2^2 = 5.32$ (p=0-03) Psychoeducational Lewis (1983) <sup>51</sup> 43/53 37/55 Lattimore et al (1990) <sup>50</sup> 50/138 50/109 Random effects model Heterogeneity: $l^2 = 79\%$ , $\chi_1^2 = 4.73$ (p=0-15) Therapeutic community Prendergast et al (2004) <sup>31</sup> 258/341 196/235 Sacks et al (2012) <sup>30</sup> 27/207 29/163 Random effects model Heterogeneity: $l^2 = 5\%$ , $d^2 = 0.1$ (p=0-76) Random effects model Heterogeneity: $l^2 = 5\%$ , $\chi_1^2 = 0.1$ (p=0-76) Random effects model Heterogeneity: $l^2 = 5\%$ , $\chi_1^2 = 0.1$ (p=0-76) Random effects model Heterogeneity: $l^2 = 5\%$ , $\chi_1^2 = 0.1$ (p=0-76)	Ortmann (2000)53	67/111	76/112		7.2%	0.72 (0.42–1.25
Random effects model       25.8% $0.74$ ( $0.47$ Heterogeneity: $l^2 = 44\%$ , $\chi_1^2 = 5.32$ (p=0.03)       9       4.6%       2.09 ( $0.86$ Lewis (1983) <sup>53</sup> 43/53       37/55       4.6%       2.09 ( $0.86$ Lattimore et al (1990) <sup>50</sup> 50/138       50/109       7.6%       0.67 0.40-         Random effects model       12.2%       1.11 ( $0.38$ 1.11 ( $0.38$ Heterogeneity: $l^2 = 79\%$ , $\chi_1^2 = 4.73$ (p=0.15)       7.0%       0.69 ( $0.39$ Therapeutic community       70%       0.69 ( $0.39$ Prendergast et al (2012) <sup>30</sup> 27/207       29/163         Random effects model       15.5%       0.64 (0.46         Heterogeneity: $l^2 = 0\%$ , $\chi_1^2 = 0.1$ (p=0.76)       100.0%       0.87 (0.68	Messina et al (2010)52	19/60	25/55	· · · · · · · · · · · · · · · · · · ·	5.5%	0.56 (0.26–1.19
Heterogeneity: $l^2$ =44%, $\chi_3^2$ =5:32 (p=0-03) Psychoeducational Lewis (1983) <sup>53</sup> 43/53 37/55 Lattimore et al (1990) <sup>50</sup> 50/138 50/109 Random effects model Heterogeneity: $l^2$ =79%, $\chi_1^2$ =4.73 (p=0-15) Therapeutic community Prendergast et al (2004) <sup>31</sup> 258/341 196/235 Sacks et al (2012) <sup>30</sup> 27/207 29/163 Random effects model Heterogeneity: $l^2$ =0%, $\chi_1^2$ =0.1 (p=0.76) Random effects model Heterogeneity: $l^2$ =54%, $\chi_1^2$ =28-33 (p<0.01)	Proctor et al (2012) <sup>54</sup>	50/98	56/85		6.8%	0.54 (0.30-0.98
Psychoeducational         Lewis $(1983)^{31}$ $43/53$ $37/55$ Lattimore et al $(1990)^{50}$ $50/138$ $50/109$ Random effects model $12\cdot2\%$ $1\cdot11$ (0·38         Heterogeneity: $l^2=79\%$ , $\chi_1^2=4.73$ (p=0·15) $8\cdot4\%$ $0.62$ (0·41-         Therapeutic community $7\cdot0\%$ $0.69$ (0·39         Random effects model $15\cdot5\%$ $0.64$ (0·46         Heterogeneity: $l^2=0\%$ , $\chi_1^2=0.1$ (p=0·76) $100\cdot0\%$ $0.87$ (0·68         Random effects model $100\cdot0\%$ $0.87$ (0·68	Random effects model				25.8%	0.74 (0.47-1.1
Lewis (1983) <sup>51</sup> 43/53 37/55 Lattimore et al (1990) <sup>50</sup> 50/138 50/109 Random effects model Heterogeneity: $l^2$ =79%, $\chi_1^2$ =4.73 (p=0-15) Therapeutic community Prendergast et al (2004) <sup>31</sup> 258/341 196/235 Sacks et al (2012) <sup>30</sup> 27/207 29/163 Random effects model Heterogeneity: $l^2$ =0.1 (p=0.76) Random effects model Heterogeneity: $l^2$ =54%, $\chi_1^2$ =0.1 (p=0.76) Random effects model Heterogeneity: $l^2$ =54%, $\chi_1^2$ =0.28-33 (p<0.01)	Heterogeneity: $l^2=44\%$ , $\chi^2_3=5\cdot 3$	32 (p=0·03)				
Lattimore et al (1990) <sup>50</sup> 50/138 50/109 <b>Random effects model</b> Heterogeneity: $l^2=79\%$ , $\chi_1^2=4.73$ (p=0-15) <b>Therapeutic community</b> Prendergast et al (2004) <sup>31</sup> 258/341 196/235 <b>Sacks et al</b> (2012) <sup>30</sup> 27/207 29/163 <b>Random effects model</b> Heterogeneity: $l^2=0\%$ , $\chi_1^2=0.1$ (p=0-76) <b>Random effects model</b> Heterogeneity: $l^2=54\%$ , $\chi^2=28\cdot33$ (p<0.01) <b>Random effects model</b> Heterogeneity: $l^2=54\%$ , $\chi^2=28\cdot33$ (p<0.01)	Psychoeducational					
Random effects model       12-2%       1-11 (0-38         Heterogeneity: $l^2$ =79%, $\chi_1^2$ =4.73 (p=0-15)       11-10-38         Therapeutic community       Prendergast et al (2004) <sup>31</sup> 258/341       196/235         Therapeutic community       12-2%       1-11 (0-38         Sacks et al (2012) <sup>30</sup> 27/207       29/163       27/207         Random effects model       15.5%       0-64 (0-46         Heterogeneity: $l^2$ =0.1 (p=0-76)       100-0%       0-87 (0-68         Heterogeneity: $l^2$ =54%, $\chi^2$ =28-33 (p<0-01)	Lewis (1983) <sup>51</sup>	43/53	37/55	•	4.6%	2.09 (0.86–5.09
Heterogeneity: $l^2 = 79\%$ , $\chi_1^2 = 4.73$ (p=0-15) Therapeutic community Prendergast et al (2004) <sup>31</sup> 258/341 196/235 Sacks et al (2012) <sup>30</sup> 27/207 29/163 Random effects model Heterogeneity: $l^2 = 0\%$ , $\chi_1^2 = 0.1$ (p=0-76) Random effects model Heterogeneity: $l^2 = 54\%$ , $\chi^2 = 28-33$ (p<0-01) Heterogeneity: $l^2 = 54\%$ , $\chi^2 = 28-33$ (p<0-01)	Lattimore et al (1990)50	50/138	50/109		7.6%	0.67 0.40-1.12)
Therapeutic community         Prendergast et al $(2004)^{31}$ 258/341 196/235         Sacks et al $(2012)^{30}$ 27/207 29/163         Random effects model         Heterogeneity: $l^2$ =0%, $\chi^2_1$ =0.1 (p=0.76)         Random effects model         Heterogeneity: $l^2$ =5%, $\chi^2_1$ =0.3 (p<0.01)	Random effects model				12.2%	1.11 (0.38-3.20
Prendergast et al (2004) <sup>31</sup> $258/341$ $196/235$ +       8.4% $0.62$ (0.41         Sacks et al (2012) <sup>30</sup> $27/207$ $29/163$ 7.0% $0.69$ (0.39         Random effects model       15.5% $0.64$ (0.46         Heterogeneity: $l^2=0.9$ , $\chi^2_1=0.1$ (p=0.76)       100.0% $0.87$ (0.68         Heterogeneity: $l^2=54\%$ , $\chi^2=28.33$ (p<0.01)	Heterogeneity: $l^2 = 79\%$ , $\chi_1^2 = 4.7$	73 (p=0·15)				
Sacks et al (2012) <sup>30</sup> 27/207 29/163 Random effects model Heterogeneity: l <sup>2</sup> =0%, $\chi_1^2$ =0.1 (p=0.76) Random effects model Heterogeneity: l <sup>2</sup> =54%, $\chi_1^2$ =28-33 (p<0.01) 100-0% 0-87 (0-68	Therapeutic community					
Random effects model         15.5%         0.64 (0.46           Heterogeneity: l²=0%, χ²=0.1 (p=0.76)         100.0%         0.87 (0.68           Random effects model         100.0%         0.87 (0.68	Prendergast et al (2004)31	258/341	196/235		8.4%	0.62 (0.41–0.9
Heterogeneity: l <sup>2</sup> =0%, \chi <sup>2</sup> =0.1 (p=0.76) Random effects model Heterogeneity: l <sup>2</sup> =54%, y <sup>2</sup> =28-33 (p<0.01) 0.87 (0.68	Sacks et al (2012) <sup>30</sup>	27/207	29/163		7.0%	0.69 (0.39–1.23
Random effects model         100.0%         0.87 (0.68           Heterogeneity: l²=54%, y²=28-33 (p<0.01)	Random effects model				15.5%	0.64 (0.46–0.9
Heterogeneity: /²=54%, y²=28-33 (p<0.01)	Heterogeneity: $l^2=0\%$ , $\chi^2_1=0.1$	(p=0·76)				
Heterogeneity: $l^2=54\%$ , $\chi^2_{13}=28\cdot33$ (p<0.01) 0.2 0.5 1.0 2.0 5.0	Random effects model				100.0%	0.87 (0.68-1.1
$\longleftarrow \longrightarrow$	Heterogeneity: $l^2=54\%$ , $\chi^2_{13}=28$	-33 (p<0·01)	0.2	0.5 1.0 2.0 5.0		
Decreased risk of recidivism Increased risk of recividism				$\longleftarrow \longrightarrow$		

Figure 5: Effectiveness of psychological interventions in prison for reducing recidivism, by intervention type

Data are for randomised controlled trials with an intervention group of at least 50 participants, excluding two outlier studies.<sup>43,6</sup> Error bars show 95% CI. CBT=cognitive behavioural therapy.

	β	SE	p value
Year of publication: ≥1990 vs <1990	-0.195	0.335	0.560
Study location: USA vs elsewhere	0.097	0.274	0.722
Sample size (continuous)	0.000	0.000	0.671
Sex of participants: single sex vs both sexes	-0.404	0.371	0.276
Mean age (continuous)	-0.016	0.018	0.372
Age group: adolescents vs adults	-0.161	0.284	0.570
Intervention type: cognitive behavioural therapy-based vs all other types	-0.217	0.270	0.422
Comparator type: usual care vs waitlist or other	0.396	0.301	0.189
Follow-up time period (continuous)	0.074	0.063	0.239
Intervention format: individual vs group or combination	-0.055	0.348	0.875
Intervention aimed at people in prison with a substance use disorder (dichotomous)	-0.283	0.256	0.269
Risk of bias: high vs low or unclear	-0.146	0.266	0.583
Table 2: Meta-regression analyses assessing links between study character	istics and re	ecidivism	risk

earlier meta-analyses because of inclusion of non-experimental designs. $^{\rm s7}$ 

Our review highlights several evidence gaps. Further research is needed to determine whether generic psychological interventions are effective in specific groups of incarcerated populations, such as people living with mental disorders other than substance misuse. Research suggests that tailored individualised interventions are associated with better treatment outcomes.<sup>89</sup> Furthermore, to improve transition to the community, future research should develop and evaluate the effects of follow-up treatments in the community after release. Greater consideration should be given to understanding the influence of environmental factors within prisons on treatment effects. Potential effects could be limited by the setting, because prisons are not primarily therapeutic environments and they prioritise security over health and rehabilitation needs.<sup>90</sup> To better understand this possibility, research comparing the effectiveness of the same treatment modality in prison versus in a community setting could provide information on whether the prison environment sustains behavioural change and what adaptations could improve treatment effectiveness in prisons.

To our best knowledge, we report the first metaanalysis of RCTs on the effectiveness of psychological interventions delivered in prisons for recidivism outcomes. Some limitations should be noted. The study selection process leading up to the full-text screening stage was done by a single reviewer. The included trials were delivered in high-income countries. In addition, the number of included studies was not large (n=29), which underlines the legal, practical, and ethical challenges of doing high-quality research in prisons.<sup>58,90,91</sup> One specific problem encountered in doing clinical research in these settings is high dropout rates, which often result in small and selective samples. Prisons have high turnover rates and participants are likely to be released or transferred unexpectedly.92 Furthermore, despite limiting inclusion to the most robust study design of RCT, only two (7%) of 29 of the included studies had low risk of bias. The most affected domains were randomisation and deviations from the intended interventions. Difficulties associated with masking staff and participants to the assigned intervention are likely to have contributed to an increased risk of bias in these two domains. There was also evidence of selective publication of small studies on the basis of their effect size (ie, some studies with small effect sizes were missing), which indicated that our initial pooled estimate of all studies (OR 0.72) was overestimated because of publication bias.93 Sex-specific analyses comparing estimates in females and males could not be done, because of insufficient numbers of female-only samples.

In conclusion, we have provided a synthesis of current research on the effectiveness of psychological interventions delivered in prisons aimed at reducing post-release recidivism. We report modest effects, at best, for psychological interventions delivered in prison. Trials of therapeutic community interventions and related approaches that facilitate continuity of treatment after prison release should be prioritised. Considering high rates of recidivism<sup>3,4</sup> and the consequences for public health and safety,<sup>5,6</sup> simple, large RCTs on the effectiveness of psychological interventions in prison are necessary.

# Contributors

GB, RY, and SF designed the study. GB did the data search, extraction, analyses, and drafted the tables under the supervision of RY and SF. GB, RY, SF, and AEP interpreted the findings. GB, RY, and SF drafted the article and all authors critically revised it. GB and RY accessed and verified the data. All authors had full access to all the data in the study and had final responsibility for the decision to submit for publication.

## **Declaration of interests**

We declare no competing interests.

# Data sharing

Data are based on the results of published studies listed in the appendix and available online. The study protocol and statistical analysis plan are available online.

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# For the study protocol and statistical analysis plan see https://www.crd.york.ac.uk/ prospero/display\_record. php?RecordID=167228

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