Clinical and cost-effectiveness of contingency management for cannabis use in early psychosis: the CIRCLE randomised clinical trial

Luke Sheridan Rains, Louise Marston, Mark Hinton, Steven Marwaha, Thomas Craig, David Fowler, Michael King, Rumana Z. Omar, Paul McCrone, Jonathan Spencer, Joanne Taylor, Sophie Colman, Catherine Harder, Eleanor Gilbert, Amie Randhawa, Kirsty Labuschagne, Charlotte Jones, Theodora Stefanidou, Marina Christoforou, Meghan Craig, John Strang, Tim Weaver and Sonia Johnson

Abstract

Background: Cannabis is the most commonly used illicit substance amongst people with psychosis. Continued cannabis use following the onset of psychosis is associated with poorer functional and clinical outcomes. However, finding effective ways of intervening has been very challenging. We examined the clinical and cost-effectiveness of adjunctive contingency management (CM), which involves incentives for abstinence from cannabis use, in people with a recent diagnosis of psychosis.

Methods: CIRCLE was a pragmatic multi-centre randomised controlled trial. Participants were recruited via Early Intervention in Psychosis (EIP) services across the Midlands and South East of England. They had had at least one episode of clinically diagnosed psychosis (affective or non-affective); were aged 18 to 36; reported cannabis use in at least 12 out of the previous 24 weeks; and were not currently receiving treatment for cannabis misuse, or subject to a legal requirement for cannabis testing. Participants were randomised via a secure web-based service 1:1 to either an experimental arm, involving 12 weeks of CM plus a six-session psychoeducation package, or a control arm receiving the psychoeducation package only. The total potential voucher reward in the CM intervention was £240. The primary outcome was time to acute psychiatric care, operationalised as admission to an acute mental health service (including community alternatives to admission). Primary outcome data were collected from patient records at 18 months post-consent by assessors masked to allocation. The trial was registered with the ISRCTN registry, number ISRCTN33576045.

(Continued on next page)
Background
Cannabis is the most commonly used illicit substance amongst people with psychosis [1]. In longitudinal studies, cannabis use in first episode psychosis is associated with substantially higher acute psychiatric admission rates [2]: an Australian study reported a 51% admission rate over 15 months follow-up amongst substance users (mostly cannabis) compared with 17% amongst non-users [3], accompanied by a threefold difference in inpatient admission rates. Similarly, a Dutch study reported a 42% admission rate amongst persistent cannabis users compared with 17% amongst those who never used or stopped round the time of first onset [4]. A dose-response relationship between severity of cannabis misuse and time to admission was also reported in this study. Problematic cannabis use in psychosis is also associated with marked delays in remission [5], and lower engagement in work or education, as well as poorer outcomes in other clinical and social domains [3, 6, 7]. Despite the clear need for effective treatments in this group, so far none have been identified. A Cochrane review [8] of psychosocial interventions for substance misuse in severe mental illness identified 32 randomised controlled trials but found little evidence that any type of therapy was more effective than treatment as usual, including Cognitive Behavioural Therapy and Motivational Interviewing.

Contingency management (CM) is an operant conditioning-based intervention for substance misuse that typically uses financial rewards to reinforce target behaviours, such as abstinence from substance use or treatment adherence. CM has an increasingly substantial evidence base in a variety of contexts, including cannabis misuse [9–11], smoking cessation [12], heavy drinking [13], and other illicit drug misuse [14, 15]. The National Institute for Health and Care Excellence (NICE) advocates its adoption in England in the management of substance use [16]. In a review conducted by NICE as part of its guidance of psychosocial interventions for substance misuse, 14 trials of CM were identified, all from the USA, of which three involved cannabis use. A consistent finding of a benefit for CM was reported, with most studies using abstinence at 12 weeks as their outcome measure. In the context of severe mental illness, a number of trials have found CM to be effective in reducing cannabis use in psychosis, at least at the level of voucher reward offered.

Conclusions: Overall, these results suggest that CM is not an effective intervention for improving the time to acute psychiatric admission or reducing cannabis use in psychosis, at least at the level of voucher reward offered.

Keywords: Financial incentives, Contingency management, Cannabis, Psychosis, Early intervention, Substance misuse
time to acute psychiatric admission amongst Early Intervention in Psychosis (EIP) service users. The objectives were (1) to conduct a pilot study of a CM intervention for cannabis use in early psychosis, (2) if the pilot was successful, to proceed with a full multi-centre pragmatic randomised controlled trial, (3) to test whether the intervention results in an increase in the time to acute psychiatric admission (the primary outcome), (4) to test whether the intervention results in a decrease in cannabis use, reduced positive psychotic symptoms, and in an increase in participation in work or education (secondary outcomes), (5) to assess the cost-effectiveness of the intervention from an NHS perspective.

Methods

Trial design and participants

CIRCLE was a rater-blind, multi-centre randomised controlled trial with two arms. The experimental (CM) group received a 12-week CM intervention together with a Treatment-As-Usual (TAU) package targeting cannabis use, which was a standardised and optimised psychoeducational package of the type of psychoeducational intervention recommended in EIP practice [28]. The control group received the optimised TAU psychoeducational intervention only. Assessments were performed at the time of consent, at 3 months (treatment end), and at 18 months following trial inclusion. The primary outcome was time to acute psychiatric admission, operationalised as admission to an acute mental health service. Ethical approval for the trial was received on 16 March 2012 from the London – South East National Research Ethics Service committee (REC reference 11/LO/1939). The trial was conducted according to the trial protocol published by Johnson et al. [29], and trial procedures are described here only in brief. Oversight was provided by an external Trial Steering Group that was appointed as agreed by the funding body.

Participants were recruited via EIP services throughout the Midlands and South East of England. Sites were added to the trial until the recruitment target was achieved. EIP services accept individuals who have developed psychotic illness for the first time and aim to treat the initial episode early and effectively, minimising acute psychiatric admissions and optimising clinical and social recovery. They are now standard provision in English mental health Trusts following policy mandates, and typically work with service users for 3 years prior to discharge to primary care or to other mental health teams for continuing care [16]. Inclusion criteria were (1) on an EIP service caseload, (2) having used cannabis at least once in 12 out of the previous 24 weeks, (3) age 18–36, (4) living in stable accommodation, (5) sufficient English to understand fully and answer the assessment instruments, (6) able to give informed consent to participate, (7) not subject to a compulsory community treatment or court order requiring urine testing for cannabis, (8) not in receipt of treatment for cannabis use from another agency, and (9) not compulsorily detained in hospital, or in prison.

Randomisation and masking

Following the baseline assessment interview, participants were randomised with a 1:1 ratio, stratified on severity of cannabis use (1–3 uses per week, > 3 uses per week). An independent randomisation service was used, coordinated by Priment Clinical Trials Unit based at University College London. Primary outcome assessors were masked to allocation throughout the trial. Secondary outcome assessors were masked to allocation at follow-up assessments in the main trial. However, masking was not feasible at the 3-month assessment interview in the pilot as sufficient researchers were not available for different staff to conduct initial and follow-up assessments. The statistical and health economics analysis plans were finalised before database lock and statisticians and health economists undertaking the analysis were masked to trial arm allocation.

Procedures

The intervention under evaluation was 12 weeks of CM delivered by clinical staff, who included graduates without mental health professional qualifications, such as assistant psychologists. Participants attended weekly CM sessions at which they were immediately rewarded with vouchers if urinalysis indicated cannabis abstinence since the previous session. Urinalysis was performed using a small bench-top analyser capable of providing rapid test results of drug of misuse urinary concentration (Kaiwood CHR-110). To perform the analysis, the clinician pipetted a fixed amount of urine into a buffer solution tube to give a 7:1 serial dilution. This allowed a standard 50 ng/ml marijuana test cassette placed in the analyser to provide a urine cannabis concentration reading between 0 and 350 ng/ml. Guidelines were provided to all staff delivering the intervention to allow interpretation of the test results. The guidelines set out upper and lower thresholds for urinary tetrahydrocannabinol (THC) based on recommendations from the suppliers of the urinalysis equipment, Surescreen Diagnostics. Results greater than the upper threshold (350 ng/ml) clearly indicated use within the last week, so were not rewarded. Results below the lower threshold indicated urinary THC concentration below 50 ng/ml, the accepted standard for detecting urinary THC, so were rewarded. Within these thresholds, urinary THC was expected to fall each week until it reached the lower THC threshold, which should be reached within 1 month. A drop in urinary THC would receive the reward. A
similar result to, or higher than, the previous week would fail. A temperature strip on the side of the specimen cup allowed staff to check whether the sample had been tampered with.

The CM schedule and rules were adapted from Budney et al. [10, 11], which both investigated CM for cannabis misuse. It used a variable reward schedule that began at £5 for providing a baseline sample in the first week. In the pilot, voucher values rose by £2 each week, with a £10 bonus voucher every other session. Based on feedback from clinicians and service users, the schedule was simplified in the main trial. The voucher value rose by £5 every two clean samples and the bonuses were removed. In total, participants could receive £240 (US$307, €267 equivalent (11 Jan. 2019)) in vouchers in both versions of the reward schedule. If a participant submitted a urine specimen indicating cannabis use in the previous week, they would receive a £5 voucher. Patients who failed to attend intervention sessions or submit a scheduled specimen did not receive a voucher. If the participant attended the next week and provided a negative sample, they were rewarded with £10. In the subsequent week, if the participant provided a second consecutive negative sample, the voucher values resumed from the highest previous level of reward. Participants could arrange in advance to miss scheduled sessions and still receive the reward for that week if they had a valid commitment that prevented them from attending and they subsequently demonstrated that they had not used since the previous sessions. They could do this on a maximum of two occasions for 1 week only each time. Vouchers were for major supermarkets.

The TAU was a standardised version of a good quality psychoeducation, which was delivered by EIP staff in a digital format. It was designed to be sufficiently highly structured for staff without high-level clinical qualifications, such as support workers or assistant psychologists, to be able to deliver it competently following brief training. The package was comprised of six modules that were delivered via a standard PC or laptop. Full delivery of all six modules was intended to take approximately 3 h, normally offered over six regularly programmed sessions of 30 min duration. It included a PDF package for clinicians to work through with their clients, which presents information regarding the effects of cannabis, motivational materials, and strategies for coping, minimizing potential harms, and abstaining from cannabis. It also included video materials, short quizzes, audio files, and further information and written records of the modules for the service user to keep. The primary aim was to deliver information to meet psychoeducation goals that were tailored to the individual needs of the participant, but not to act as a psychological intervention. It adopted a harm minimisation approach, with an acknowledgement that in a young person with psychosis, cannabis abstinence may be required to ensure that no harm is done. The content was based on motivational interviewing principles, relapse prevention, and harm reduction strategies [link to psychoeducation package - https://www.ucl.ac.uk/psychiatry/research/epidemiology-and-applied-clinical-research-depa/projects/circle-trial/trial-materials]. A full day’s training to provide CM and the TAU psychoeducational package was delivered to EIP clinical staff by members of the CIRCLE research team. Training was tailored to the knowledge and experience of clinicians, some of whom were assistant psychologists with psychology degrees, but not mental health professional training. Clinicians were provided with written documentation on delivering the CM and the psychoeducation and supported by the CIRCLE team throughout the trial. Records of urinalysis results and voucher rewards received for each participant were checked by the CIRCLE research team to ensure fidelity to the intervention protocol.

Outcomes

The primary outcome was time to admission to an acute psychiatric service, including psychiatric hospital, crisis resolution team or crisis house, or other acute mental health service intended as an equivalent to hospital. The primary outcome was assessed over an 18-month period using electronic patient records.

Secondary outcomes were between-group differences at follow-up for the following: (1) severity of positive symptoms of psychosis. (2) Social functioning based on self-reports of engagement in work or study. (3) Self-reported number of days’ cannabis use in the previous 3 months at 3-month follow-up, or the previous 6 months at 18 months. (4) Proportion of cannabis-free urines at assessment. (5) Number of admissions over 18 months follow-up. (6) Quality-adjusted life years (QALYs) (SF-12 and EQ-5D) and service use (CSRI) were used in the cost-effectiveness analyses. Baseline and secondary outcome data were collected primarily during assessment interviews. Some secondary outcome data were checked or collected using electronic patient records at each assessment. The following measures were performed at all assessment interviews:

- Demographics, most recent diagnosis, and social information. Where feasible, data were checked using patient records.
- Cannabis use
  ○ The timeline followback (TLFB) [30] was used to record self-reported substance use, including cannabis, other illicit substances, and alcohol, over the previous 6 months at baseline and 18 months,
and over the previous 3 months at the 3-month follow-up assessment interview.

- Structured Clinical Interview for DSM IV (SCID) part E was used to assess history of alcohol and substance misuse disorders.
- Urinalysis for cannabis, performed using a commercially available immunoassay test for metabolites of the cannabinoid tetrahydrocannabinol (THC) using the standard cut-off of 50 ng/ml [31].

- Psychotic symptoms
  - The positive and negative symptom subscales of the Positive and Negative Syndrome Scale (PANSS) [32].

- Service use and health economic analysis:
  - The Client Service Receipt Inventory (CSRI) [33] was used to record clinical service use, medication use, receipt of state welfare benefits, and use of other state-funded services including the criminal justice system. Data were collected for the previous 6 months at baseline and 18 months, and for the preceding 3 months at 3-month assessment. Data were collected at assessment interview and checked using patient records. To minimise attrition, at 18 months, data were collected from patient records for a subset of the resources deemed most likely to contribute to higher costs and that feasibly could be collected. Patient record data were collected for all patients still in the trial, unless they had withdrawn or their patient records could not be identified.
  - 12-Item Short Form Survey (SF-12) [34] and the EQ-5D [35] were used to derive quality-adjusted life years (QALYs).
  - Details of the participant’s referral to the EIP service, history of admission to acute mental health services, and time spent on a Community Treatment Order (CTO) in the last 6 months at baseline. At 3 months and 18 months, history of admission to acute mental health services and time spent on CTO were recorded since study inclusion. Data were obtained from patient records.

**Statistical methods**

The sample size for the trial was 544. This was based on data suggesting a usual acute psychiatric admission rate of around 50% over the study timeframe in cannabis users [3, 4]. A 15% decrease in this rate due to the intervention was considered to be clinically beneficial. Using a power of 90% and a significance level of 5%, a total sample size of 460 subjects was determined to be required. This sample size was based on an analysis of time to acute psychiatric admission and allowed a 37% decrease in the hazard of admission (hazard ratio of 0.63) in the intervention group to be detected using a Cox proportional hazards model. The sample size was inflated by a factor of 1.06, which assumed that each clinician delivering the intervention would see an average of four trial participants, and an intraclass correlation coefficient of 0.02 for clinician clustering, which gave a total sample size of 488. Finally, the sample size was inflated by 10% to account for attrition for the primary outcome, giving a total sample size of 544. This calculation was performed using Stata version 11 [36].

All analyses were carried out comparing the CM and control groups as randomised using all available data. The continuous variables were summarised using mean (standard deviation (SD)) or median (interquartile range (IQR)) as appropriate. Categorical variables were presented as frequencies and percentages. Logistic regression was used to determine baseline predictors of missingness. Kaplan-Meier survival curves by randomised groups were used to examine the primary outcome (time to acute psychiatric admission) descriptively. A Cox proportional hazards model was used to compare the primary outcome in the intervention and control groups, adjusting for severity of cannabis use (the stratification variable; 1 to 3 times a week versus 4 or more times a week) at baseline and whether the participant was part of the pilot trial. The assumption of proportional hazards was checked using Schoenfeld residuals [37]. The supportive analyses for the primary outcome were as follows:

- Adjusting for significant baseline predictors of missingness.
- Excluding those participants who had no secondary outcome data (for 12 weeks and 18 months separately).
- Including those in the main trial only as some minor changes were made to the protocol at the end of the pilot before the main trial commenced.
- Adjusting additionally for the number of psychoeducation sessions attended (which was offered in both arms of the trial).
- Adjusting additionally for the number of admissions in the 6 months prior to baseline.
- Adjusting for the same factors as the primary analysis but using centre as the clustering variable instead of clinician.

Secondary outcomes were analysed separately at 3 and 18 months. Models were adjusted for severity of cannabis use at baseline and whether the participant was part of the pilot trial. For the dichotomous outcomes (cannabis-positive urine, engaged in work or study), logistic regression was used. The residuals for the positive and
negative symptoms from PANSS outcomes were not normally distributed and were therefore log transformed and analysed using linear regression models. Zero-inflated Poisson regression was used for count outcomes including number of cannabis days and number of admissions over follow-up, as there were excess zeros in these outcomes. The number of cannabis days at 3 months was analysed using Poisson regression. All secondary outcomes were also analysed after adjusting for predictors of missingness. An additional supportive analysis for number of admissions was to include all those who were discharged from psychiatric services within 18 months and assuming they did not have any admissions over 18 months. Robust standard errors were used in all regression models to account for clustering by a clinician delivering the CM in the analyses [38]. Results from all supportive and secondary analyses are presented as estimates and 95% confidence intervals (CI) as specified in the statistical analysis plan. All analyses were carried out using Stata version 14 [39].

Health economic analyses

The cost-effectiveness analysis was conducted from an NHS/social care perspective over 18 months. Costs of the CM intervention and psychoeducation were calculated using the salaries of staff delivering the CM and psychoeducation, supervision time, and overheads. Other service use was measured with the CSRI and from data collected from patient records. Costs were calculated by combining resource use with 2015/2016 unit costs. Average costs were used to cover the gap between 3 and 12 months. This did not apply to inpatient costs which were available throughout the follow-up period. Imputation for missing non-inpatient costs was carried out with predictors being community costs for available periods and inpatient costs. (This was done so that analyses could be conducted on the sample with complete inpatient data.) Costs were compared between arms at 18-month follow-up using a regression model, controlling for baseline, and with bootstrapped confidence intervals based on the percentile method and 10,000 resamples.

Quality-adjusted life years (QALYs) were derived from the EQ-5D-3L and SF-12 (for secondary analyses) using area under the curve methods incorporating scores at baseline and 3- and 18-month follow-ups. Missing utility scores were imputed from all available scores. QALY comparisons were made after adjusting for baseline scores. Costs and QALY differences were obtained from 10,000 bootstrapped resamples. Cost-effectiveness planes and cost-effectiveness acceptability curves (CEACs) were generated.

Results

Recruitment and follow-up

Recruitment to the pilot phase occurred between June 2012 and March 2013. The pilot study successfully achieved its aims by recruiting within 10% of its target (n = 68), demonstrating the feasibility of recruiting, delivering the interventions, and retaining at least 60% of participants at 3 months. Subsequently, a full RCT was performed. Recruitment to the main trial occurred between October 2013 and March 2016. Follow-up data collection was completed by October 2017. Figure 1 shows the flow of service users through the trial. Seventy EIP teams took part in the trial from 23 NHS trusts. Two thousand four hundred two service users were initially approached by EIP clinicians. Of these, 551 service users gave consent and were randomised into the trial. Primary outcome data were obtained for 530 participants (92%). Assessment interviews were performed with 371 participants at 3-month follow-up (67%) and 278 participants at 18 months (50%).

Baseline characteristics

Most baseline characteristics were similar between the two groups. In both groups, over 85% of participants were male, with a mean age of around 25 (SD 4). 72% were using cannabis more than three times a week and reported using on 111 days on average in the previous 6 months. Around a third of participants were diagnosed with schizophrenia or schizo-affective disorder and half had diagnoses of other types of psychosis, perhaps in part because EIP services tend to be reticent about making a diagnosis of such as schizophrenia because of concerns regarding stigma and diagnostic instability. A quarter of participants were currently engaged in some form of work or education, but a large majority (over 80%) had held open market employment at some point. Median PANSS positive symptom score was 12 (IQR 9, 17) in the control group and 13 (IQR 9, 16) in the CM group. Lifetime rates of cannabis dependence were very high (87% of control and 85% of intervention group members), and around three quarters in both groups were dependent at baseline. Lifetime rates of alcohol misuse or dependence, and of reports of using substances other than cannabis, were also fairly high (e.g. 47% of control and 52% of intervention group members reported using cocaine; 36% of control and 32% of intervention group members met the criteria for alcohol dependence) (Table 1). In the 6 months prior to baseline, median days of substance use other than cannabis were low (0) in the CM group (IQR 0, 4) and in the control group (0; IQR 0, 1). Median alcohol using days was 6 in both the CM (IQR 0, 26) and control groups (IQR 0, 26).
Participation in CM and psychoeducation

Participants in the CM group obtained a mean of £64 (out of a maximum of £240) in voucher rewards and attended a median number of 9 (IQR 3, 12) (maximum of 12) CM sessions. Forty-six participants declined the CM intervention or did not attend any sessions. Participants attended a median of 6 and 4 (maximum of 6) psychoeducation sessions in the CM and control groups respectively. However, 86 participants in the control group and 63 in the CM group declined the psychoeducation or did not attend any sessions.

Primary outcome

For the primary outcome, there was no statistically significant difference in time to admission to an acute mental health service between the randomised groups (hazard ratio (HR) 1.03, 95% CI 0.76, 1.40) (Fig. 2 and Tables 2, 3, and 4).

Supportive analyses

Results from the supportive analyses were similar. The odds of at least one admission over 18 months of follow-up were similar across randomised groups (OR 1.02,
Table 1 Baseline demographics

| Characteristic                        | Control n/N or median or utility score | %, SD, or IQR | Contingency Management (CM) n/N or median or utility score | %, SD, or IQR |
|--------------------------------------|----------------------------------------|---------------|-------------------------------------------------------------|---------------|
| Male                                 | 240/273                                | 88            | 238/278                                                     | 86            |
| Age mean (SD)                        | 25                                     | (4)           | 24                                                         | (4)           |
| Ethnicity                            |                                        |               |                                                            |               |
| White                                | 144/273                                | 53            | 148/277                                                     | 53            |
| Black                                | 62/273                                 | 23            | 65/277                                                     | 23            |
| Asian                                | 30/273                                 | 11            | 29/277                                                     | 10            |
| Other                                | 37/273                                 | 14            | 35/277                                                     | 13            |
| Marital status                       |                                        |               |                                                            |               |
| Single                               | 253/273                                | 93            | 259/278                                                     | 93            |
| Married or cohabiting                | 14/273                                 | 5             | 17/278                                                     | 6             |
| Other                                | 6/273                                  | 2             | 2/278                                                      | 1             |
| Educational attainment               |                                        |               |                                                            |               |
| No qualifications                    | 48/273                                 | 18            | 43/277                                                     | 16            |
| GCSE or equivalent                   | 104/273                                | 38            | 133/277                                                    | 48            |
| A level or equivalent                | 67/273                                 | 25            | 58/277                                                     | 21            |
| Post 18 education (including HND, trade, degree) | 54/273 | 20 | 43/277 | 16 |
| Ever had open market employment      | 223/273                                | 82            | 240/278                                                    | 86            |
| Any current work or study            | 67/273                                 | 25            | 73/278                                                     | 26            |
| Diagnosis                            |                                        |               |                                                            |               |
| Schizophrenia or schizo-affective disorder | 80/256 | 31 | 90/268 | 34 |
| Bipolar affective disorder           | 26/256                                 | 10            | 19/268                                                     | 7             |
| Depression with psychotic symptoms   | 11/256                                 | 4             | 5/268                                                      | 2             |
| Other psychosis                      | 139/256                                | 54            | 154/268                                                    | 57            |
| Cannabis use                         |                                        |               |                                                            |               |
| 1–3 times a week                     | 77/273                                 | 28            | 78/278                                                     | 28            |
| More than 3 times a week             | 196/273                                | 72            | 200/278                                                    | 72            |
| Any work or study                    | 58/183                                 | 32            | 58/189                                                     | 31            |
| Cannabis-positive urine              | 210/262                                | 80            | 214/272                                                    | 79            |
| History of cannabis dependence       | 238/273                                | 87            | 236/278                                                    | 85            |
| Current cannabis dependence          | 183/238                                | 77            | 185/236                                                    | 78            |
| PANSS positive symptoms median (IQR) | 12                                    | (9, 17)       | 13                                                         | (9, 16)       |
| PANSS negative symptoms median (IQR) | 14                                    | (11, 19)      | 14                                                         | (10, 19)      |
| EQ-5D-3L utility score               | 0.7619                                 |               | 0.7652                                                     |               |
| SF-6D utility score                  | 0.6789                                 |               | 0.6833                                                     |               |

95% CI 0.70, 1.48). Approximately, a third of participants experienced at least one acute admission to a mental health service, including hospital alternatives such as crisis teams and crisis houses, by 18 months in both the CM (90/272) and control groups (85/259). Amongst those who experienced an acute psychiatric admission, the median number of days until admission was 245 (IQR 99, 382) in the control group and 196 (IQR 82, 364) in the CM group.

Secondary outcomes
For participants who had a full 18 months’ follow-up, those in the CM group had a slightly higher rate ratio for number of admissions than those in the control group (incidence rate ratio (IRR) 1.08, 95% CI 0.75, 1.54); this changed little when assuming those who were discharged from services had no admissions during follow-up or when including predictors of missingness. Those in the CM group had lower odds of paid work or
study at both 3 and 18 months (OR 0.95, 95% CI 0.62, 1.46; OR 0.82, 95% CI 0.50, 1.35 respectively). However, those in the CM group also had slightly lower odds of cannabis-positive urine at 3 and 18 months (OR 0.86, 95% CI 0.56, 1.34; OR 0.84, 95% CI 0.49, 1.41 respectively). None of these results approached statistical significance at the 5% level. Illicit substance use other than cannabis was very low at both follow-ups (3-month median days of use: 0; IQR 0, 1 in both groups; 18-month: 0 in both groups, IQR 0, 2 in control; IQR 0, 1 in CM). The median number of alcohol-using days was 4 (control IQR 0, 12; CM IQR 0, 15) in both groups at 3 months and 6 (IQR 0, 24) in both groups in 18 months. For the log-transformed PANSS positive outcome at 3 months, the CM group score was on average 7% lower (better) than the control group (95% CI −14%, −0%). There were 58 reported serious adverse events (SAE). In the CM group, these were 24 psychiatric hospital admissions and 3 deaths. In the control group, there were 28 psychiatric hospital episodes, 2 deaths, and 1 arrest.1

CM in the context of psychoeducation
A post hoc analysis was performed to help understand the results in the context of whether people had received psychoeducation as planned. Attending at least four of the six psychoeducation sessions planned was selected as the measure of compliance (137/263 (52%) in the control group and 168/273 (62%) in the CM group). There were no marked differences in demographic, social, or clinical baseline characteristics between compliers and non-compliers between randomised groups. A dichotomous variable indicating compliance was created. A Cox model with robust standard errors was conducted. The primary outcome of the trial (time to acute psychiatric admission) was used as the dependent variable in the analysis. Randomisation group, compliance, severity of cannabis use at baseline, and whether the participant was part of the pilot trial were included as covariates, and an interaction term between compliance and randomisation groups was used. The interaction term was statistically significant (p value for interaction 0.016), which suggests that CM might be effective in participants who receive sufficient psychoeducation. This may be due to the psychoeducation being effective amongst those who engaged with treatment, and engagement was higher in the CM group.

Health economics
Intervention costs were on average £298 for the CM group and £140 for controls. Inpatient use (based on 531 participants) over the follow-up period was virtually the same between trial arms, with around one quarter receiving this and for about 3 months across the period (Table 4). Other service use (based on the smaller sample of 231 who were followed up with the CSRI) was relatively similar between the arms over the follow-up period. Costs for drug and alcohol services were however greater for the CM group. After imputation, the mean (SD) health and social care costs over the 18-month period were £15,614 (£29,360) for the CM group and £16,620 (£33,283) for controls. After adjusting for baseline costs, the CM group had costs that were on average £1625 lower than for controls (bootstrapped 95% CI, −£3355 to £6869).

EQ-5D-3L and SF-6D utility scores increased gradually over time for both groups (Table 2). The total mean QALYs over the follow-up based on the EQ-5D-3L were 1.2218 for the CM group and 1.1855 for controls. Adjusting for baseline utility, CM resulted in 0.034 more QALYs than controls. QALYs based on the SF-6D scores were 1.0682 for CM and 1.0585 for controls. With adjustment, CM resulted in 0.0063 more QALYs than controls. Therefore, CM had lower costs and produced more QALYs and so ‘dominated’ the control. There is uncertainty around the results shown by the cost-effectiveness planes. With the EQ-5D-3L (Fig. 3), the most likely result is that CM has lower costs and better outcomes (72% of replications), followed by higher costs and better outcomes (26%), lower costs and worse outcomes (2%), and higher costs and worse outcomes (1%). With the SF-6D, there is more uncertainty (Fig. 4), but the most likely outcome is still CM having lower costs and better outcomes (52%), followed by lower costs and
worse outcomes (22%), higher costs and better outcomes (18%), and higher costs and worse outcomes (8%). The CEACs (Fig. 5) show that even with a zero value attached to a QALY there is still a high probability of CM being the most cost-effective option. At a threshold value of £20,000, the probabilities are 0.81 for the EQ-5D-3L and 0.75 for the SF-6D.

**Discussion**

The results of this trial indicate that CM confers no clinical advantage over TAU for patients with psychosis who use cannabis. Neither was any effect seen in our secondary outcomes, including cannabis use, engagement in work or education, and positive psychotic symptoms. However, a post hoc analysis found that compliance with psychoeducation in the CM arm resulted in a statistically significant improvement in time to acute psychiatric admission, while the same was not true for the control group. This suggests that CM had a clinical benefit amongst those who also engaged with psychoeducation.

The economic analyses show that the costs associated with CM were less than for the TAU, although the difference was not statistically significant and this analysis

---

**Table 2** Outcomes at 3 months (treatment end) and 18 months

| Outcome                                                                 | Control n/N, mean, or median | CM n/N, mean, or median |
|------------------------------------------------------------------------|------------------------------|-------------------------|
| Admitted to an acute mental health service 18 months                   | 85/259 33 (IQR)              | 90/272 33 (IQR)         |
| Number of admissions 18 months                                         | 0 (0, 1)                     | 0 (0, 1)                |
| Any work or study 3 months                                            | 58/183 32 (IQR)              | 58/189 31 (IQR)         |
| Any work or study 18 months                                           | 45/135 33 (IQR)              | 42/145 29 (IQR)         |
| Cannabis-positive urine 3 months                                      | 122/170 72 (IQR)             | 128/184 70 (IQR)        |
| Cannabis-positive urine 18 months                                      | 76/124 61 (IQR)              | 77/136 57 (IQR)         |
| Number of days using cannabis* (median) 3 months                      | 30 (3, 84)                   | 26 (1, 67)              |
| Number of days using cannabis* (median) 18 months                     | 26 (1, 142)                  | 26 (0, 118)             |
| PANSS positive symptoms (median) 3 months                             | 11 (8, 16)                   | 10 (8, 14)              |
| PANSS positive symptoms (median) 18 months                            | 10 (8, 15)                   | 11 (8, 13)              |
| PANSS negative symptoms (median) 3 months                             | 14 (10, 18)                  | 12 (9, 17)              |
| PANSS negative symptoms (median) 18 months                            | 12 (8, 17)                   | 12 (9, 17)              |
| Number of days of using illicit substances other than cannabis (median)| 0 (0, 1)                     | 0 (0, 1)                |
| Number of days of using illicit substances other than cannabis (median)| 0 (0, 2)                     | 0 (0, 1)                |
| Number of days using alcohol (median) 3 months                        | 4 (0, 12)                    | 4 (0, 15)               |
| Number of days using alcohol (median) 18 months                       | 6 (0, 24)                    | 6 (0, 24)               |
| EQ-5D-3L utility score (mean) 3 months                                 | 0.7729                      | 0.8175                  |
| EQ-5D-3L utility score (mean) 18 months                               | 0.8032                      | 0.8336                  |
| SF-6D utility score (mean) 3 months                                   | 0.7018                      | 0.7114                  |
| SF-6D utility score (mean) 18 months                                  | 0.7110                      | 0.7205                  |
| Number of psychoeducation sessions attended (median)                  | 4 (0, 6)                    | 6 (1, 6)                |
| Number of contingency management sessions attended (median)           | 9 (3, 12)                   |                         |
| Number who attended 4 or more PE sessions (used in the post hoc analysis) | 137/263 52 (IQR)            | 168/273 62 (IQR)        |

*At 3 months, this was for the previous 12 weeks/84 days. At 18 months, this was for the previous 168 days.*

---
### Table 3: Analysis of primary and secondary outcomes in terms of contingency management

| Outcome                                                                 | Estimate* | 95% CI      | N   | Estimate† | 95% CI | N   |
|------------------------------------------------------------------------|-----------|-------------|-----|-----------|--------|-----|
| Time to acute psychiatric admission (HR)                              | 1.03      | (0.76, 1.40)| 531 | 1.02      | (0.75, 1.40)| 531 |
| Cannabis-positive urine sample 3 months (OR)                         | 0.86      | (0.56, 1.34)| 354 | 0.85      | (0.55, 1.32)| 352 |
| Cannabis-positive urine sample 18 months (OR)                        | 0.84      | (0.49, 1.41)| 260 | 0.85      | (0.50, 1.43)| 260 |
| log PANSS positive symptoms 3 months                                | −0.07     | (−0.14, −0.00) | 366 | −0.07    | (−0.14, −0.00) | 364 |
| log PANSS positive symptoms 18 months                                | −0.04     | (−0.13, 0.05) | 276 | −0.04    | (−0.13, 0.04) | 276 |
| log PANSS negative symptoms 3 months                                 | −0.08     | (−0.16, 0.00) | 362 | −0.08    | (−0.16, 0.00) | 360 |
| log PANSS negative symptoms 18 months                                | 0.01      | (−0.08, 0.11) | 276 | 0.02    | (−0.08, 0.11) | 276 |
| Paid work or study at 3 months (OR)                                  | 0.95      | (0.62, 1.46)| 372 | 0.94      | (0.60, 1.47)| 370 |
| Paid work or study at 18 months (OR)                                 | 0.82      | (0.50, 1.35)| 280 | 0.82      | (0.50, 1.35)| 280 |
| Number of days cannabis used in the previous 12 weeks (3 months follow-up) (IRR) | 0.89 | (0.75, 1.04) | 371 | 0.89 | (0.75, 1.04) | 369 |
| Number of days cannabis used in the previous 6 months (18 months follow-up) (IRR) | 1.09 | (0.88, 1.36) | 274 | 1.09 | (0.87, 1.33) | 274 |
| Number of admissions over 18 months follow-up                         | 1.08      | (0.75, 1.54)| 374 | 1.09      | (0.76, 1.55)| 374 |
| At least one admission over 18 months follow-up (OR)                 | 1.02      | (0.70, 1.48)| 531 | 1.01      | (0.69, 1.48)| 531 |

CI confidence interval, HR hazard ratio, OR odds ratio, IRR incident rate ratio

* Adjusting for level of cannabis use at baseline and whether in the pilot study
† Additionally adjusting for baseline predictors of missingness. These are:
Time to admission, number of admissions, at least one admission: any work or study
Urine positive 3 months, PANSS positive 3 months, PANSS negative 3 months, any work or study 3 months, number of days cannabis use 3 months: exempt from work due to disability
Urine positive 18 months: any work or study
PANSS positive 18 months, PANSS negative 18 months, any work or study 18 months, number of days cannabis use 18 months: voluntary work

### Table 4: Service use for the health economics

| Service                                      | Control % users | Mean (SD contacts) | Mean (SD) cost | CM % users | Mean (SD contacts) | Mean (SD) Cost |
|----------------------------------------------|-----------------|--------------------|----------------|------------|--------------------|----------------|
| Inpatient stays                              | 25.5            | 90.9               | 11,931         | 24.6       | 89.4               | 11,339         |
| Early intervention team                      | 68.9            | 11.1               | 303            | 68.7       | 11.7               | 302            |
| GP                                           | 44.7            | 2.9                | 134            | 51.1       | 3.1                | 127            |
| Psychiatrist                                 | 53.1            | 3.2                | 479            | 54.3       | 3.2                | 404            |
| Psychologist                                 | 20.9            | 5.6                | 445            | 22.7       | 5.0                | 416            |
| Home treatment/crisis team                   | 10.3            | 12.8               | 21             | 9.0        | 10.7               | 302            |
| Mental health nurse                          | 13.9            | 7.0                | 129            | 15.1       | 9.1                | 127            |
| Adult education class                        | 3.7             | 9.0                | 18             | 2.5        | 5.4                | 404            |
| Assertive outreach team                      | 1.1             | 9.0                | 5              | 1.8        | 8.4                | 4              |
| Class/group at a leisure centre              | 4.4             | 25.2               | 33             | 4.0        | 14.6               | 10             |
| Community mental health centre               | 2.6             | 7.6                | 40             | 4.3        | 8.8                | 79             |
| Day care centre/day hospital                 | 0.7             | 27.0               | 3              | 1.4        | 5.2                | 4              |
| Drop-in centre                               | 4.8             | 10.8               | 33             | 1.4        | 16.5               | 12             |
| Drug/alcohol service                         | 4.8             | 6                  | 23             | 4.0        | 16.0               | 163            |
| Drug and alcohol advisor                     | 10.6            | 7.9                | 116            | 9.4        | 5.5                | 291            |
| Occupational therapist                       | 7.0             | 3.7                | 62             | 6.1        | 2.1                | 37             |
| Other counsellor/therapist                   | 5.1             | 7.6                | 156            | 4.0        | 7.6                | 53             |
| Other doctor                                 | 8.1             | 3.7                | 75             | 7.6        | 2.1                | 26             |
| Self-help/support group                      | 5.1             | 6.5                | 9              | 7.2        | 6.8                | 16             |
| Social worker                                | 10.3            | 7.5                | 58             | 7.9        | 10.4               | 44             |
| Medication                                   | 288             |                    |                |            |                    |                |
was carried out after imputation for a large proportion of cases, while we had primary outcome data for most who entered the trial. QALY differences based on the EQ-5D-3L were relatively large compared to other studies, but less so with the SF-6D. The lower costs and greater number of QALYs mean that CM is more cost-effective than TAU even though the differences for both costs and QALYs were not statistically significant. While interpretation of this is complex as there was little difference between CM groups in the intention-to-treat analyses, one possible explanation is that time to acute psychiatric admission was better amongst those who complied with psychoeducation in the CM group, but not the control group. The admission rate was lower than anticipated in the trial as a whole. Only around a third in each group required admission to acute care, while it was expected [3, 4] for the purposes of our sample size calculation and based on previous literature that half would experience an acute psychiatric admission. This suggests that a comparatively stable group of

![Fig. 3 Cost-effectiveness plane based on EQ-5D-3L](image)

![Fig. 4 Cost-effectiveness plane based on SF-6D](image)
patients was recruited to trial and/or that the psychoeducation and/or the extra attention of being in a research study had a beneficial impact. Although the majority still had cannabis-positive urine, the number of reported cannabis-using days in the previous 6 months fell from over 100 to 26 in each group by 18-month follow-up, indicating that cannabis use declined in both groups over this period. This could be due to the psychoeducation intervention, the normal progression of use in this group, regression to the mean, or a combination of these factors. Low levels of self-reported illicit substance and alcohol use at follow-up indicate that people were not substituting them for cannabis.

Overall, the results of the trial are mixed, and stand in contrast to the frequently positive benefits reported for CM interventions [15, 16], including for substance misuse in psychosis [17–19, 21]. Reasons for this are unclear; however, some possible explanations include the following: Firstly, offering more frequent CM sessions or offering a higher reward might have been more clinically effective. The CM schedule was adapted from two trials by Budney et al. [10, 11], which both found a positive effect in cannabis misuse. However, in the two trials by Budney et al. [10, 11] and Bellack et al. [21], all participants were dependent, and so it is unlikely that the CM intervention failed to provide a benefit because of high rates of dependence. An alternative explanation is that participants may have been using more highly potent forms of the cannabis and consequently may have found it more difficult to abstain than those using less-potent forms. There is good evidence that this cohort typically uses more potent forms of cannabis than non-psychiatric groups [40] and that high potency cannabis use is relatively prevalent in London, where much of our sample was recruited [41]. It is therefore plausible that use of highly potent types of cannabis may have been relatively widespread in our sample. However, we did not systematically record the type of cannabis participants were using, making further exploration of this possibility difficult.

Fig. 5 Cost-effectiveness acceptability curves
Thirdly, while engagement was good with people attending a median of 4 psychoeducation sessions in the control group and 6 in the CM group, a substantial proportion of people (around one third of people in the control group and one fifth in the CM group) declined to take part in either the CM and/or the PE or did not attend any sessions. This suggests there may have been a substantial minority who were not particularly motivated to quit cannabis, but perhaps participated due to the enthusiastic approach to recruitment by staff and researchers and/or the small payment received to acknowledge baseline interview assessment participation. Recruitment was slower than anticipated, partly because fewer service users than expected wanted to enter treatment for cannabis use. The post hoc analysis results suggested that people who were motivated to adhere to the psychoeducation had better outcomes. It may be that a more engaged cohort would have benefited more from the CM.

Fourthly, patients in this cohort are often multiply disadvantaged [2, 5, 42], which is likely to make behavioural change much more challenging. This includes being less likely to be in work or education, having poorer social networks, and thereby being more socially isolated but also the nature of psychotic illness itself, which can include greater disorganisation, poorer social skills, and lower motivation [43, 44]. It may be that interventions are needed in this cohort that have a broader focus than just reducing cannabis use. It may be that a better approach would support patients in becoming less socially isolated or spending less time with cannabis-using peers, as well as helping them back into work or engaging in other meaningful activities.

Finally, the trial focused only on cannabis, but baseline data indicated considerable history of alcohol and substance use disorders. It may be that the CM would have been more effective if it had addressed these additional substances as well.

There were several limitations to the study. Firstly, CIRCLE was a pragmatic trial of a CM intervention delivered in EIP services in the UK. The CM and psychoeducation treatments were designed to be feasible for EIP clinicians to deliver in routine practice, which required limiting the additional workload to clinicians. As such, during the intervention period, urinalysis samples were only collected from the CM group and not controls. This makes it more challenging to analyse whether the CM group participants reduced their use more than the controls during the intervention period. However, self-reported days of cannabis use at treatment end did not differ between groups, suggesting no impact from the CM. However, CIRCLE should be viewed as a trial of a pragmatic CM intervention rather as a definitive trial of whether CM reduces cannabis use in psychosis. Studies that more rigorously assess drug use in both treatment groups are needed to investigate whether CM could be effective at reducing use in psychosis.

Secondly, the inclusion of an active control makes it more difficult to interpret results. The fall in cannabis use across the trial population suggests there may have been benefit from the enhanced TAU. While it was intended to be a standardised form of treatment as usual, it became apparent that it was a much better developed and more ambitious psychoeducation than was otherwise available in many of the participating EIP teams.

Thirdly, originally, the CM (and psychoeducation) was originally intended to be delivered by care coordinators (the nurses, occupational therapist or social workers with primary responsibility for keeping in touch with patients and organising their care). However, few care coordinators were prepared to deliver the interventions, largely due to concerns regarding time pressures and potential disruption to therapeutic relationships. Instead, other clinical staff, such as support workers or assistant psychologists, were trained to deliver it. Greater integration into routine care may have improved its effectiveness. However, data on delivery of the CM intervention suggested that those delivering it did adhere well to the intended protocols, and it may be that care coordinators would have been less successful in this.

Fourthly, while the follow-up rate for the primary outcome was very high, attrition was greater than anticipated on the interview measures, potentially introducing response bias.

Fifthly, while a standard threshold was used for the urinalysis at assessment interview (50 ng/ml), a lower threshold would have given a slightly more accurate measure of abstinence rates. A lower threshold may have identified a small difference between groups. However, as there was no difference in either the proportion of cannabis-free urine using a threshold of 50 ng/ml or self-reported days of use at either follow-up, it seems unlikely that a clinically significant difference would be identified.

Sixthly, while we approached all EIP patients who were identified to trial researchers as potentially eligible by their clinicians and who agreed to be contacted by a trial researcher, it may be that gate-keeping by EIP clinicians or self-selection by patients could have introduced bias. However, this was a pragmatic RCT and it is likely that the trial sample is a good reflection of the range of characteristics of the EIP service users who would enter treatment if the intervention was offered through EIP or specialist drug treatment services. Similar issues seem likely if the intervention was introduced as part of routine care.

Seventhly, we did not systematically record the type of cannabis participants were using. It may be that those using more potent types of cannabis would respond differently to...
the intervention compared to those using milder forms, either by finding it harder to change behaviour, or potentially by showing greater benefit from the CM treatment in terms of acute psychiatric service use.

Eighthly, although not a limitation in the analyses, the finding that CM has over 80% likelihood of being cost-effective needs to be treated with caution. In the sample that were followed up at both time points, the costs were lower and QALY outcomes better for the CM group compared to controls, and this led to a favourable cost-effectiveness ratio for CM which applied to most bootstrapped resamples. The cost and QALY differences were though limited and not statistically significant.

Finally, using acute psychiatric admission as the outcome is a pragmatic choice because it is routinely available data that are relatively accessible via patient records. But it is limited in that some acute psychiatric episodes are likely to be contained without acute service use, for example, it may be handled by the EIP team instead, and also that thresholds for admission to acute care may vary considerably, for example by clinician and by area.

Conclusions
Overall, the lack of effectiveness of CM in these intention-to-treat analyses means that it cannot be recommended as an intervention to reduce cannabis use in patients with recent onset psychosis. However, the adherence and health economic analyses suggest that effectiveness of CM might be better amongst those who engage well with psychoeducation, or if a different reward value or schedule was offered. Further investigation of stakeholder perspectives would be useful to explore the latter possibility. However, overall, this is another trial that demonstrates how challenging it is to address the problem of cannabis use in psychosis. It may be that a substantially different approach is required to address this significant clinical problem. It has been noted that young people who have psychosis and problematic cannabis use are often multiply disadvantaged. Despite this, trials of interventions in this area are often narrowly focused on changing cannabis use. A more inclusive management that takes in patients' social contexts, including engagement in work or education, might prove more fruitful.

Acknowledgements
CIRCLE was funded by the National Institute for Health Research (NIHR) Health Technology Assessment (HTA) Programme (09/144/50). The views expressed in this publication are those of the authors and not necessarily those of the NIHR, HTA programme, NHS, or the Department of Health. We would like to thank all the participants who took part in the trial, as well as all the Early Intervention in Psychosis (EIP) clinical staff who assisted with recruitment or delivered the interventions. We are also grateful to the members of the Trial Steering Committee (Professor Thomas Barnes, Dr. Jonathan Haynes, Dr. Sara Brookes, Maurice Arbuthnott, Katherine Barrett, Nick Barber, and Beverley Chipp) and the Data Monitoring and Ethics Committee (Professor David Kingdon, Dr. Chris Metcalfe, Dr. Rob McPherson, and Dr. David Li) for providing oversight throughout, and the principal investigators (Dr. Charlot Johnston-Webber, Mr. Simon Clark, Dr. Nikola Raman, Dr. Manoj Sukumaran, Dr. Peter Carter, Dr. Rick Fraser, Dr. Nilamadhab Kar, Dr. Sahaj Yousef, Dr. Jonathan West, Mr. Nicholas Chamberlain-Kent, Dr. Abu Abraham, Prof. Hana Soliman, Dr. John Joseph Scanlon, Dr. Nandini Chakraborty, Dr. Saggar Hamad, Dr. Eileen O'Regan, Dr. Chris Harrop, Dr. Sanjoo Chengappa, Prof. Belinda Lennon, Dr. Jesus Perez) for supporting the trial. Finally, we would like to thank Dr. Barnaby Major, Kelly Wilson, Leila Hughes, Melanie Whyman, Simon Spencer, Alyssa Milton, Danielle Lamb, Kate Fullarton, Johanna Freirichs, Kirsty Shepherd, Danielle Pettman, Ruth Mintah, Sherene Olusesi, Tayla McCloud, and the clinical studies officers who contributed to the trial for all their efforts.

Authors’ contributions
SJ was the chief investigator who also managed the primary research centre, UCL, and designed the study with contributions from SM, DF, TC, LM, RO, MK, TW, JS, and PM. MH had a major role in the conception and design of the trial. LSR managed the main trial and drafted the final manuscript. LM and RO authored and performed the statistical analyses. PM authored the cost-effectiveness analyses and with JSJ performed the analysis. MK through the PRIMENT clinical trial unit contributed to the study design, statistical analyses, and the interpretation of the findings. SM, DF, and TC managed the other research centres. MCR managed the pilot phase of the trial and with MH developed the psychoeducation package. MH was largely responsible for the development of the contingency management intervention. SC, KL, CH, JT, CI, EG, AR, MCh, and TS were researchers on the trial, recruited and assessed participants, and supported clinicians in delivering the interventions. JSt and TW provided expertise to the study design and delivery. All authors read and approved the final manuscript.

Funding
This study was funded by the National Institute for Health Research, Health Technology Assessment programme (09/144/50). The funder of the study had no role in study design, data collection, data analysis, data interpretation, or writing of the report.

Availability of data and materials
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

Ethics approval and consent to participate
Ethical approval for the trial was received on 16 March 2012 from the London – South East National Research Ethics Service committee (REC reference 11/LO/1939). All participants provided written informed consent.

Consent for publication
Not applicable.

Competing interests
This trial was funded by the National Institute for Health Research, Health Technology Assessment programme (09/144/50), and MK is a member of CTUs funded by NIHR and the Rapid Trials and Add-on Studies Board. RM is a member of the HTA General Board. JS is supported by the National Institute for Health Research (NIHR) Biomedical Research Centre for Mental Health at South London and Maudsley NHS Foundation Trust and King’s College London. JS reports grants from Camurus (Lund, Sweden), Martindale Pharma (now Ethypharm UK, Woodburn Green, UK), Mundipharma (Mundi-pharma International Limited, Cambridge, UK) and Braeburn Pharma (Braeburn Inc., Plymouth Meeting, PA, USA), and other from Martindale Pharma,
Braeburn Pharma outside the submitted work. In addition, John Strang has patents unrelated to the work described in this article: a patent EuroCeltique issued and a patent King’s College London pending.

Author details
1Division of Psychiatry, University College London, London, UK. 2Mental Health and Wellbeing, Warwick Medical School, University of Warwick, Coventry, UK. 3Addictions Department, Institute of Psychiatry, Psychology, and Neuroscience, King’s College London, London, UK. 4Health Service and Population Research Department, Institute of Psychiatry, Psychology and Neuroscience, King’s College London, London, UK. 5Mental Health Social Work & Inter-professional Learning, Middlesex University London, London, UK. 6Department of Health Services and Population Research, King’s Health Economics, Institute of Psychiatry, Psychology & Neuroscience, King’s College London, London, UK. 7Department of Psychology, University of Sussex, Brighton, UK. 8Department of Primary Care and Population Health and Priment Clinical Trials Unit, University College London, London, UK. 9Department of Statistical Science, University College London, London, UK. 10Institute for Mental Health, School of Psychology, University of Birmingham, Birmingham B15 2TT, UK. 11Camden and Islington NHS Foundation Trust, 4 St Pancras Way, London NW1 0PE, UK. 12Centre for Posttraumatic Mental Health, Department of Psychiatry, University of Melbourne, Melbourne, Australia.

Received: 23 January 2019 Accepted: 22 July 2019
Published online: 15 August 2019

References
1. Barnes TE, Muttsata SH, Hutton SB, Watt HC, Joyce EM. Comorbid substance use and age at onset of schizophrenia. Br J Psychiatry. 2006;188:237–42.
2. Schoeler T, Monk A, Sami MB, Klamerus E, Foglia E, Brown R, et al. Continued versus discontinued cannabis use in patients with psychosis: a systematic review and meta-analysis. Lancet Psychiat. 2016;3:215–25.
3. Wade D, Hannigan S, Edwards J, Burgess PM, Whelan G, McCoy PD. Substance misuse in first-episode psychosis: 15-month prospective followup study. Br J Psychiatry. 2006;189:229–34.
4. Linszen D, Dingemans P, Lenier M. Cannabis abuse and the course of recent-onset schizophrenic disorders. Arch Gen Psychiatry. 1994;51:273–4.
5. Lambert M, Conus P, Lubman DI, Wade D, Yuen H, Moritz S, et al. The impact of substance use disorders on clinical outcome in 643 patients with first-episode psychosis. Acta Psychiatr Scand. 2005;112:141–8.
6. Walsh E, Buchanan A, Fahy T. Violence and schizophrenia: examining the evidence. Br J Psychiatry. 2002;180:490–5.
7. Marwaha S, Johnson S, Bebbington P, Stafford M, Angermeyer MC, Brugha T, et al. Rates and correlates of employment in people with schizophrenia living in the UK, France and Germany: findings from the EuroSC study. Br J Psychiatry. 2007;191:30–7.
8. Hunt GE, Sefried N, Morley K, Sirtharan T, Cleare AJ. Psychosocial interventions for people with both severe mental illness and substance misuse. Cochrane Database Syst Rev. 2013;(10):Art. No.:CD01088. https://doi.org/10.1002/14651858.CD01088.pub3 http://www.cochrane.org/CD001088.
9. Secades-Villa R, García-Rodríguez O, Higgins S, Fernández-Hermida JR, Caballo JL. Community reinforcement approach plus vouchers for cocaine dependence in a community setting in Spain: six-month outcomes. J Subst Abus Treat. 2008;34:202–7.
10. Weaver T, Metrebian N, Hellier J, Pilling S, Charles V, Little N, et al. Use of contingency management incentives to improve completion of hepatitis B vaccination in people undergoing treatment for heroin dependence: a cluster randomised trial. Lancet. 2014;384:153–63.
11. Preibe S, Yeates K, Bremner S, Lauber C, Eldridge S, Ashby D, et al. Effectiveness of financial incentives to improve adherence to maintenance treatment with antipsychotics: cluster randomised controlled trial. BMJ. 2015;347:s5847.
12. Vercarle-Willa R, García-Rodríguez O, Higgins S, Fernández-Hermida JR, Caballo JL. Community reinforcement approach plus vouchers for cocaine dependence in a community setting in Spain: six-month outcomes. J Subst Abus Treat. 2008;34:202–7.
13. Treizier J, Froud J, Huggett K, Pilling S, Edgar A, Roberts S, et al. A randomised controlled trial of the clinical and cost-effectiveness of a contingency management intervention compared to treatment as usual for reduction of cannabis use and of relapse in early psychosis (CIRCLE): a study protocol for a randomised controlled trial. Trials. 2016;17:515.
14. Sobell L, Sobell M. Timeline follow-back. In: Allen J, Litten R, editors. Measuring alcohol consumption: psychosocial and biological methods. New York: Humana Press; 1992.
15. Cany PL. The marijuana detection window: determining the length of time cannabinoids will remain detectable in urine following smoking: a critical review of relevant research and cannabinoid detection guideline for courts. Drug Court Rev. 2005;5:23–56.
16. Kay SR, Fiszbein A, Opler LA. The positive and negative syndrome scale (PANSS) for schizophrenia. Schizophr Bull. 1987;13(2):261–76.
17. Beecham J, Knapp M. Costing psychiatric interventions. In: Thomsricto G, editor. Measuring mental health needs. London: Gaskell; 2001.
18. Amorim P, Gomes de Sa P, Duarte P, Pimentel I, Carvalho F, Correia V, et al. A randomized controlled trial of contingency management for stimulant use in community mental health patients with serious mental illness. Am J Psychiat. 2013;170:94–101.
19. Tidey J, Rohsenow D, Kaplan G, Swift R, Reid N. Effects of contingency management and bupropion on cigarette smoking in smokers with schizophrenia. Psychopharmacol. 2011;217(2):279–87.
20. Murphy SM, McDonell MG, McPherson S, Srebrik D, Angelo F, Roll JM, et al. An economic evaluation of a contingency-management intervention for stimulant use among community mental health patients with serious mental illness. Drug Alcohol Depen. 2015;153:293–9.
21. Bellack AS, Bennett ME, Gearon JS, Brown CH, Ye Yang MS. A randomized clinical trial of a new behavioral treatment for drug abuse in people with severe and persistant mental illness. Arch Gen Psychiatry. 2006;63:426–32.
22. Sigmon SC, Steingard S, Badger GJ, Anthony SL, Higgins ST. Contingent reinforcement of marijuana abstinence among individuals with serious mental illness: a feasibility study. Exp Clin Psychopharmacol. 2000;8:509–17.
23. Sigmon SC, Higgins ST. Voucher-based contingent reinforcement of marijuana abstinence among individuals with serious mental illness. J Subst Abus Treat. 2006;30:291–5.
24. Rabin R, Kozak K, Zakanzis K, Remington G, Stefan C, Budney A, et al. A method to achieve extended cannabis abstinence in cannabis dependent patients with schizophrenia and non-psychiatric controls. Schizophren Res. 2018;194:47–54.
25. Secades-Villa R, García-Rodríguez O, Higgins S, Fernández-Hermida JR, Caballo JL. Community reinforcement approach plus vouchers for cocaine dependence in a community setting in Spain: six-month outcomes. J Subst Abus Treat. 2008;34:202–7.
26. Weaver T, Metrebian N, Hellier J, Pilling S, Charles V, Little N, et al. Use of contingency management incentives to improve completion of hepatitis B vaccination in people undergoing treatment for heroin dependence: a cluster randomised trial. Lancet. 2014;384:153–63.
27. Preibe S, Yeates K, Bremner S, Lauber C, Eldridge S, Ashby D, et al. Effectiveness of financial incentives to improve adherence to maintenance treatment with antipsychotics: cluster randomised controlled trial. BMJ. 2013;347:s5847.
28. NICE, NHS England, and the National Collaborating Centre for Mental Health, Implementing the Early Intervention in Psychosis Access and Waiting Time Standard: Guidance, 2016.
29. Johnson S, Sheridan Rains L, Marwaha S, et al. A randomised controlled trial of the clinical and cost-effectiveness of a contingency management intervention compared to treatment as usual for reduction of cannabis use and of relapse in early psychosis (CIRCLE): a study protocol for a randomised controlled trial. Trials. 2016;17:515.
30. Sobell L, Sobell M. Timeline follow-back. In: Allen J, Litten R, editors. Measuring alcohol consumption: psychosocial and biological methods. New York: Humana Press; 1992.
31. Cany PL. The marijuana detection window: determining the length of time cannabinoids will remain detectable in urine following smoking: a critical review of relevant research and cannabinoid detection guideline for courts. Drug Court Rev. 2005;5:23–56.
32. Kay SR, Fiszbein A, Opler LA. The positive and negative syndrome scale (PANSS) for schizophrenia. Schizophr Bull. 1987;13(2):261–76.
33. Beecham J, Knapp M. Costing psychiatric interventions. In: Thomsricto G, editor. Measuring mental health needs. London: Gaskell; 2001.
34. Brazeur JE, Roberts JR. The estimation of a preference-based index from the SF-12. Med Care. 2004;42:851–9.
35. McCrone P, Patel A, Knapp M, Schene A, Koeter M, Ammadeo F, et al. A comparison of SF-6D and EQ-5D utility scores in a study of patients with schizophrenia. J Ment Health Policy Econ. 2009;12:27–31.
36. StataCorp. Stata Statistical Software: Release 11. College Station: StataCorp LP; 2009.
37. Collett D. Modelling survival data in medical research. 3rd ed. Hoboken: Chapman and Hall; 2014.
38. White H. A Heteroskedasticity-consistent covariance matrix estimator and a direct test for Heteroskedasticity. Econometrica. 1980;48(4):817.
39. StataCorp. Stata statistical software release 14. College Station: StataCorp LP; 2015.
40. Di Forti M, Marconi A, Cana E, Farietta S, Trotta A, Bonomo M, et al. Proportion of patients in South London with first-episode psychosis attributable to use of high potency cannabis: a case-control study. Lancet Psychiatry. 2015;2(3):233–8.
41. Di Forti M, Quattrone D, Freeman T, Tripoli G, Gayer-Anderson C, Quigley H, et al. The contribution of cannabis use to variation in the incidence of psychotic disorder across Europe (EU-GEI): a multicentre case-control study. Lancet Psychiatry. 2019;6(5):427–36.
42. Schimmelmann BG, Corus P, Cotton S, Kupferschmid S, McGorry PD, Lambert M. Prevalence and impact of cannabis use disorders in adolescents with early onset first episode psychosis. Eur Psychiat. 2012;27:463–9.
43. DiClemente C, Nidecker M, Bellack A. Motivation and the stages of change among individuals with severe mental illness and substance abuse disorders. J Subst Abuse Treat. 2008;34(1):25–35.
44. Priester M, Brovne T, lachini a c S, DeHart D, Seay K. Treatment access barriers and disparities among individuals with co-occurring mental health and substance use disorders: an integrative literature review. J Subst Abus Treat. 2016;61:47–59.

Publisher’s Note
Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.
Minerva Access is the Institutional Repository of The University of Melbourne

Author/s:
Rains, LS; Marston, L; Hinton, M; Marwaha, S; Craig, T; Fowler, D; King, M; Omar, RZ; McCrone, P; Spencer, J; Taylor, J; Colman, S; Harder, C; Gilbert, E; Randhawa, A; Labuschagne, K; Jones, C; Stefanidou, T; Christoforou, M; Craig, M; Strang, J; Weaver, T; Johnson, S

Title:
Clinical and cost-effectiveness of contingency management for cannabis use in early psychosis: the CIRCLE randomised clinical trial

Date:
2019-08-15

Citation:
Rains, LS; Marston, L; Hinton, M; Marwaha, S; Craig, T; Fowler, D; King, M; Omar, RZ; McCrone, P; Spencer, J; Taylor, J; Colman, S; Harder, C; Gilbert, E; Randhawa, A; Labuschagne, K; Jones, C; Stefanidou, T; Christoforou, M; Craig, M; Strang, J; Weaver, T; Johnson, S, Clinical and cost-effectiveness of contingency management for cannabis use in early psychosis: the CIRCLE randomised clinical trial, BMC MEDICINE, 2019, 17 (1)

Persistent Link:
http://hdl.handle.net/11343/227623

File Description:
Published version