



A novel cognitive behaviour therapy for bipolar disorders (Think Effectively About Mood Swings or TEAMS): study protocol for a randomized controlled trial

DOI:
[10.1186/1745-6215-15-405](https://doi.org/10.1186/1745-6215-15-405)

Document Version
Proof

[Link to publication record in Manchester Research Explorer](#)

Citation for published version (APA):

Mansell, W., Tai, S., Clark, A., Akgonul, S., Dunn, G., Davies, L., Law, H., Morriss, R., Tinning, N., & Morrison, A. P. (2014). A novel cognitive behaviour therapy for bipolar disorders (Think Effectively About Mood Swings or TEAMS): study protocol for a randomized controlled trial. *Trials*, *15*(1), Article 405. <https://doi.org/10.1186/1745-6215-15-405>

Published in:
Trials

Citing this paper

Please note that where the full-text provided on Manchester Research Explorer is the Author Accepted Manuscript or Proof version this may differ from the final Published version. If citing, it is advised that you check and use the publisher's definitive version.

General rights

Copyright and moral rights for the publications made accessible in the Research Explorer are retained by the authors and/or other copyright owners and it is a condition of accessing publications that users recognise and abide by the legal requirements associated with these rights.

Takedown policy

If you believe that this document breaches copyright please refer to the University of Manchester's Takedown Procedures [<http://man.ac.uk/04Y6Bo>] or contact uml.scholarlycommunications@manchester.ac.uk providing relevant details, so we can investigate your claim.



The Protocol for a Randomised Controlled Trial for a Novel Cognitive Behaviour Therapy for Bipolar Disorders: TEAMS (Think Effectively About Mood Swings)

Warren Mansell, Sara Tai

School of Psychological Sciences, University of Manchester¹

Alexandra Clark, Savas Akgonul,

Psychosis Research Unit, Greater Manchester West Mental Health NHS Foundation Trust²

School of Psychological Sciences, University of Manchester¹

Graham Dunn

Centre for Biostatistics, Institute of Population Health, University of Manchester⁴

Linda Davies

Centre for Health Economics, Institute of Population Health, University of Manchester³

Heather Law

Psychosis Research Unit, Greater Manchester West Mental Health NHS Foundation Trust

Richard Morriss

Institute of Mental Health, School of Medicine, University of Nottingham⁵

Neil Tinning

Bipolar UK, London⁶

Anthony P. Morrison

Psychosis Research Unit, Greater Manchester West Mental Health NHS Foundation Trust²

School of Psychological Sciences, University of Manchester¹

Emails: warren.mansell@manchester.ac.uk; sara.tai@manchester.ac.uk; alexandra.clark@gmw.nhs.uk; savas.akgonul@gmw.nhs.uk; graham.dunn@manchester.ac.uk; linda.davies@manchester.ac.uk; heather.law@gmw.nhs.uk; richard.morriss@nottingham.ac.uk; neiltinning@me.com; tony.morrison@gmw.nhs.uk

¹ School of Psychological Sciences, University of Manchester, Oxford Road, Manchester M13 9PL

² The Psychosis Research Unit, GMW Mental Health NHS Foundation Trust, Harrop House, Bury New Road, Prestwich, Manchester, M25 3BL

³ Centre for Biostatistics, Institute of Population Health, Jean McFarlane Building (1st Floor), Oxford Road, Manchester, M13 9PL

⁴ Centre for Health Economics, Institute of Population Health, Jean McFarlane Building (1st Floor), Oxford Road, Manchester, M13 9PL

⁵ Institute of Mental Health, School of Medicine, University of Nottingham, Wollaton Road, NG8 1BB

⁶ Bipolar UK, 11 Belgrave Road, London, SW1V 1RB

Abstract

Background. Existing psychological therapies for bipolar disorders have been found to have mixed findings, with a consensus that they provide a significant, but modest, effect on clinical outcomes. Typically, these approaches have focused on promoting strategies to prevent future relapse. An alternative treatment approach, termed Think Effectively About Mood Swings (TEAMS) addresses current symptoms, including subclinical hypomania, depression and anxiety; and promotes long-term recovery. Following the publication of a theoretical model, and range of research studies testing the model, and a case series have demonstrated positive results. The current study reports the protocol of a feasibility randomised controlled trial to inform a future multi-centre trial.

Methods/Design. A target number of 84 patients with a diagnosis of bipolar I or II disorder, or bipolar disorder not-otherwise-specified are screened, allocated to a baseline assessment and randomised to either 16 sessions of TEAMS therapy plus treatment-as-usual (TAU) or TAU. Patients complete self-report inventories of depression, anxiety, recovery status, and bipolar cognitions targeted by TEAMS. Assessments of diagnosis, bipolar symptoms, medication, access to services, quality of life are conducted by assessors blind to treatment condition at 3, 6, 12, and 18 month post-randomisation. The main aim is to evaluate recruitment and retention of participants into both arms of the study as well as adherence to therapy to determine feasibility and acceptability. It is predicted that TEAMS+TAU will reduce self-reported depression in comparison to TAU at six months post-randomisation. The secondary hypotheses are that TEAMS will reduce the severity of hypomanic symptoms and anxiety; reduce bipolar cognitions, and improve social functioning and promote recovery compared to TAU at post-treatment and follow-up. The study also incorporates semi-structured interviews about the experiences of previous treatment and the experience of TEAMS therapy that will be subject to qualitative analyses to inform future developments in the approach.

298 words

Trial Registration: Current Controlled Trials [ISRCTN83928726](https://www.clinicaltrials.gov/ct2/show/study?term=ISRCTN83928726)

Keywords: cognitive behavioural therapy; controlled trial; bipolar disorder; recovery

Background

Bipolar disorders are characterised by periods of depression and periods of elated or irritable mood (mania or hypomania), with around 2 to 6% of the general population meeting criteria for diagnosis [1, 2]. The cost of bipolar disorders annually in the UK is £5.2 billion, projected to rise to £8.2 billion by 2026. Despite treatment, people with bipolar disorders remain depressed for the majority of time when outside an episode. After apparent recovery from a bipolar episode, subsyndromal symptoms of depression remain in the long term, estimated at 30-50% of subsequent weeks [3, 4]. Depressive symptoms form the largest contribution to impaired functioning in bipolar disorder [5, 6]. At present, antidepressant treatment is relatively ineffective for these individuals [7].

There is a clear need for a more effective, cost-efficient intervention such as CBT (Cognitive Behavioural Therapy). A number of reviews and meta-analyses of existing psychological interventions for bipolar disorders demonstrate mixed outcomes; yet overall, there are significant but modest effects [8-10]. A similar mixed set of outcomes were found in the most recently published trials, with most participants continuing to experience significant sub-clinical symptoms and disruption to life functioning after receiving an intervention [11-13]. Thus, there remains a need for an alternative approach to managing bipolar disorders. Our group has been developing and piloting a case series for a new CBT approach known as TEAMS (Think Effectively About Mood Swings) based on an empirically supported model [14]. It involves a new integrative treatment approach that has the potential to benefit those people whose symptoms of depression, hypomania and anxiety have otherwise remained treatment resistant and with reduced functioning.

Our approach specifically targets current problems as specific to the patient, in contrast to earlier forms of CBT for bipolar disorder that have focused on preventing future relapse. In a range of other psychiatric disorders, CBT is also designed to work on current problems. Treatment is focused on developing a formulation (a shared understanding) of how thinking styles and behaviours maintain and escalate current symptoms - such as unipolar depression [15, 16], anxiety disorders [17, 18], and eating disorders [19]. In

our own research, people diagnosed with bipolar disorders reported that their recovery involved not simply remaining free of relapse but also regaining a sense of purpose in their lives and facing their longstanding problems [20, 21].

Therefore, the cognitive model [22] that guides TEAMS focuses on understanding what is maintaining people's current problems, such as anxiety, depression and irritability; and facilitates people regaining control over their lives. The model proposes that people with bipolar disorders experience mood difficulties because they strive to control their mood in extreme ways, which negatively affects their ability to control and achieve their broader goals in life. As such, their mood rarely reaches a stable state and their attempts to pursue their life goals are disrupted. The goal of TEAMS is to help people to develop awareness of the ways in which they manage their moods and develop alternative ways in which to live a more fulfilling and balanced life.

The current study represents the fourth phase in the development and testing of TEAMS. In the first phase, we conducted a series of literature reviews [23-25] and qualitative analyses of personal accounts [21, 26] to inform our treatment model [22]. In the second phase, we tested the model through a range of self-report, observational and experimental studies that provided strong evidence for the central role of extreme appraisals of internal states as a key mechanism in bipolar disorders and mood swings (e.g.[27-29]. In the third phase we reported a number of case studies and a case series [30] that demonstrated promising findings regarding acceptability and effect sizes; and so established the rationale for a pilot randomised controlled trial at the fourth stage. At this fourth stage it is important to establish whether there is evidence that TEAMS added to usual care has evidence of clinical effectiveness of usual care. Hence the TEAMS CBT will be compared to usual care. The fifth stage is envisaged to be a multi-site, definitive randomised controlled trial to establish a more precise estimate of the clinical and cost effectiveness of the intervention as well as the generalizability of the results.

Method

The study was approved by the London Queens-Square Research Ethics Committee (REC ref: 11/LO/1326).

Objectives

To determine whether the TEAMS approach is feasible and acceptable to individuals with Bipolar Disorders compared with treatment as usual and establish an estimate of treatment effect size to inform a larger definitive randomised controlled trial.

Main research questions:

1. We aim to assess feasibility, acceptability, and information to design a definitive RCT such as uptake, service barriers and attrition,
2. We predict that TEAMS will reduce symptoms of depression, hypomania, and anxiety; improve social functioning, and promote recovery in comparison to treatment as usual at 6 months post-baseline (post-treatment in the TEAMS wing), and at 12 and 18-month post-baseline.
3. We aim to evaluate the effectiveness and cost-effectiveness of an intervention which has the potential for direct benefits for patients

Trial design

A parallel group rater-blind randomised controlled trial comparing i) treatment as usual with ii) up to 16 sessions of TEAMS therapy. Allocation to the two groups will be one to one and has an exploratory framework. The design for this study involves a quantitative component (pilot trial) and a qualitative component (interviews with patients and service providers described later).

Primary outcome measure

The main objective of the pilot randomised control trial (RCT) relates to evaluation of recruitment and retention of participants into both arms of the study as well as adherence to therapy to determine feasibility and acceptability. We predict that TEAMS will reduce depression in comparison to treatment as

usual at 6 and 12 months post-randomisation with the Beck Depression Inventory [31] at 6 months as the primary outcome measure.

Secondary outcome measures

The secondary hypotheses are that TEAMS will reduce the severity of hypomanic symptoms and anxiety; change thinking styles responsible for maintaining symptoms which will be targeted by the therapy, and improve social functioning and promote recovery compared to treatment as usual at post-treatment and follow-up.

Several secondary outcome measures will be used: SCID-LIFE interview for episodes of mania or depression [32]; interview measures of manic symptoms (Bech-Rafaelson; [33]) and depression [34], composite internal state scale score (ISS; [35]); anxiety (GAD-7;[36]); recovery measure (The Process of Recovery Questionnaire QPR,[37]); Cognitive style (Hypomanic Attitudes and Positive Predictions Inventory;[38, 39]); Euroqol Health Status Measure (EQ-5D; [40]) Session-by-session measures. Raters were trained at the beginning and during the study in interview measures of outcome by an experienced rater who is a consultant psychiatrist (RM) to improve the reliability and validity of the ratings. Participants are paid a fixed amount of expenses for their time (£10 per assessment after baseline) and travel for the follow-up assessment sessions. If any participant withdraws from treatment, we will seek permission to collect outcome data. If a participant misses their assessment, they are contacted a maximum of three times by the rater through the means of contact that they have specified (e.g. home and mobile telephone, letter, care coordinator), after which they are coded as missed and recontacted for the next follow-up point. Self-report measures are collected where interview assessments are not possible.

Prior to each session, the clients complete the ISS and a personalised 10-15 item version of the HAPPI (Client-HAPPI) which assesses their most prominent beliefs about internal states [30]. At the end of each treatment session, clients rate their level of satisfaction with therapy on a Likert scale from 0 (not at all) to 10 (extremely). Adherence is evaluated by the widely used revised version of the Cognitive Therapy Rating

Scale (CTS-R; [41]) and a brief checklist of adherence specific to the TEAMS model – The TEAMS Treatment Adherence Checklist (T-TAC).

TEAMS therapy

The TEAMS approach involves an individualised treatment plan of CBT based on the integrative cognitive model of bipolar disorder [22]. Specific treatment strategies and therapy techniques employed are determined by an individual formulation using the model. There are several published descriptions of the intervention as well as a therapy manual in development [14, 22, 30, 42].

Control condition group

The control condition consists of Treatment As Usual (TAU) which typically involves regular meetings with a care-coordinator, access to a psychiatrist, medication, and monitoring of risks that require immediate intervention. Treatments, including other psychological interventions are not withheld in the TAU group, but are monitored using a Treatment Documentation Sheet. Following written consent, participants are randomised within two working days.

Randomisation and blinding

An independent Unit - The Manchester Academic Health Science Centre Clinical Trials Unit conducts anonymised randomisation in a permuted blocks design with block sizes varying randomly between 8 and 12. There is no stratification of the sample, but baseline information may be used in a *post hoc* manner to explore the effects of baseline characteristics. The procedure is as follows. An administrator contacts the CTU with a password and the participant identification number, and receives an allocation sequence to assign participants to either group. The administrator then directly contacts participants and therapists.. Due to the nature of the trial only a single blinding (of raters) is needed. Blindness of raters is ensured using a variety of procedures including separate offices for therapists and research assistants, briefings to

participants prior to assessment, and encryption of randomisation information. Unblindings are regularly monitored and recorded. Deliberate unblinding would only occur if there was a serious adverse incident such as suicide or violence to another person by a participant.

Sample size

The primary aim of this study is to evaluate feasibility; therefore a formal power calculation is not appropriate. However, estimates indicate that with 42 participants per group, using a t-test with a two-tailed significance level of 0.05 we will have over 80% power to detect an effect size of 0.8. If the significance level were altered to 15%, 30 per group will have 80% power to detect an effect size of 0.6. A previous case series of seven patients generated a pre-post effect size at one-month follow-up on the primary outcome measure of 3.0, and between 0.7 and 1.4 for the secondary measures [30]. The above information makes a feasibility trial of 30 participants per group an appropriate target.

In aiming for 60 participants in total, the target recruitment is 84 participants, allowing a 29% attrition rate by the 12-month post-randomisation point. This drop-out rate was based on a conservative estimation just above the highest reported rate of attrition found in other published studies of CBT for bipolar disorder (13% [43]; 25% [44]; 21% [10]).

Recruitment

Eighty-four participants aged 16 or over who meet DSM-IV criteria for a diagnosis of bipolar disorder, will be recruited. Recruitment is taking place across a number of NHS Trusts in and around Greater Manchester. The lead Trust for the research is Greater Manchester West Mental Health NHS Foundation Trust.

Advertisements including posters and flyers to promote the study have been distributed in Community Mental Health Teams (CMHT), outpatient clinics, voluntary services (including Manic Depression Fellowship) and GP practices. Local media and websites are used to maximise potential referrals from individuals who are not currently accessing other services. The trial is also supported by the UK Mental

Health Research Network (MHRN) who provide Clinical Studies Officers (CSO) to facilitate recruitment. The research team and CSOs present verbal and written information outlining the study to clinicians, health professionals and voluntary services. Potential participants are offered a Participant Information Sheet (PIS) containing an overview of the study by a member of their care team or directly by the research team, if preferred.

Inclusion and exclusion criteria

For the purpose of this study, participants are selected on the basis of following criteria:

Inclusion criteria:

- 1) Meet DSM-IV criteria for bipolar I or II disorder, or bipolar disorder not otherwise specified, characterised by a past major depressive episode and DSM-IV hypomania of two days or more.
- 2) Complete a baseline assessment sessions in an outpatient setting
- 3) A baseline score of at least 15 on the Beck Depression Inventory prior to the study (to ensure presence of significant current distress as targeted in the trial)
- 4) Aged 16 or over

Exclusion Criteria:

- 1) A diagnosis of a non-affective psychotic disorder according to DSM-IV
- 2) Current mania or mixed episode according to DSM-IV
- 3) A primary substance use disorder according to DSM-IV
- 4) Moderate to severe learning disability
- 5) Organic impairment that accounts for mental health problem
- 6) Non-English speaking (owing to the standardised assessment measures)

Procedure

Figure 1 (consort diagram) provides a summary of the study procedure. Potential participants are given at least 24 hours to consider the information provided before being contacted by a researcher to discuss the

study and ensure that they understand the information provided. For willing participants, the research team asks for written permission to contact their GP and/ or other relevant clinician to obtain information that might affect their ability to take part (i.e. risk) and to confirm what clinical support will be ongoing during the trial. The research team then conducts a screening assessment to ensure the participant meets inclusion criteria for the study. This screening process can take place over the telephone or face-to-face (dependent on participant preference), and is based on the Brief Screening Interview and two non-consecutive weeks of exceeding the criteria for depressive symptoms (BDI > 15). If inclusion criteria are met at this stage and the potential participant chooses to enter the trial, full written consent is sought before the initial baseline assessment.

Full assessments will subsequently be conducted on five occasions: initial baseline, 3 months post-randomisation (mid-treatment for the TEAMS group), 6 months post randomisation (post-treatment for the TEAMS group), and 12 months (6 months post-treatment for the TEAMS group) and 18 months. Those participants who have met the inclusion criteria following baseline assessment are randomised (to treatment as usual or TEAMS therapy) and informed of their group allocation by letter. TEAMS treatment usually commences within two weeks of randomisation. See Figure 1 (consort diagram) and Table 1.

---- INSERT TABLE 1 AND FIGURE 1 AROUND HERE ----

Semi-structured interviews and Qualitative Analyses

Qualitative data forms a significant part of the study and will be utilised to guide the subsequent development of the therapist and client treatment manuals. In this study, we conduct two semi-structured interviews with a subgroup of 12-15 randomly selected participants. The interviews were informed on the basis of our previous work on providing TEAMS therapy and through discussions with an experienced service user who is a coinvestigator for the trial. The first interview is completed at the baseline and explores participants' subjective experiences of how they have considered, sought, and experienced

various forms of treatment in the past; and what they feel an appropriate psychological treatment should involve.

The second interview occurs within six months of treatment ending and examines participants' subjective experience of receiving the TEAMS intervention. It focuses on what might have been beneficial and what was considered less effective or unhelpful, and eliciting recommendations for future modifications.

Thematic Analysis will be used to analyse qualitative data. All interviews are audio recorded and transcribed verbatim. Each transcript will be read and subjected to content analysis to identify themes. Emerging themes from each report will be discussed with the research team to ensure agreement. Data saturation will be regarded as achieved when no new themes emerge (it is likely that this will be achieved with approximately 12-15 interviews). Also, a selection of 5 – 10 service providers will be interviewed during the study on their perceptions on their desired outcomes, including referral and retention.

Data monitoring and management

Hard copies of anonymised data are stored at the NHS site in locked filing cabinets separate from identifiable information (names, addresses and dates of birth). Raters enter data into an electronic database that is stored anonymously with the participant ID only, in a password-protected hard drive. All data entry is checked by a second person and any errors corrected. Only researchers, therapists study administrators and lead investigators (ST & WM) have access to personal information. Only the lead investigators, trial statistician (GD) and researchers have access to the final dataset.

Adverse and serious adverse events are collected by the research team and reported to the Chief Investigator or their nominated deputy. There is no data monitoring committee because the nature of the intervention is unlikely to result in any adverse event and there are no interim analyses or stopping rules for the trial. Trial conduct may be audited independently of the research team by the sponsor or independently of both the sponsor and the investigators by the funders. In the unlikely event of harm,

participants would have access to compensation through the NHS indemnity scheme and through the sponsor of the study.

Statistical Analyses

All primary analyses will be based on an intention-to-treat principle, focussing more on descriptive statistics and confidence interval estimation than statistical significance. Primary and secondary outcomes, will be analysed employing analysis of repeated measures with a mixed effects model, accounting for the discrete timing of follow-up assessments, and the random censoring introduced by shorter follow-up periods for participants' recruited towards the end of the trial. The sensitivities of all treatment effect estimates to missing outcome data arising from patient drop-out (noncompliance) will be examined and secondary analyses carried out to estimate the Complier-Average Causal Effect (CACE), allowing for missing outcome data, as described in detail [45,46], again concentrating on confidence interval estimation rather than statistical significance. Basic service data for those refusing randomisation will be collected to assess generalisability of the estimated treatment and economic effects. Other than the CACE analyses, there will be no subgroup analysis, interim analysis nor stopping guidelines because there are no circumstances under which the trial would be stopped.

Cost effectiveness analysis

The incremental cost effectiveness ratio of TEAMS therapy will be estimated and cost effectiveness acceptability analysis conducted, from the perspectives of health and social care providers and patients; the key stakeholders in treatment decisions. Case note review and patient service use questionnaires are being used to collect data about formal and informal service use for each participant to estimate costs for the 3 months prior to entry to the study and from study entry to end of follow up. The main measure of health benefit is the quality adjusted life year (QALY), estimated from survival and the health status of each patient measured by the Euroqol. Secondary analyses will be used to explore uncertainty due to design decisions and inform the design of a definitive trial.

Dissemination Plans

The findings will be disseminated within peer-reviewed journals authored by the investigators, in addition to presentations at national and international practitioner conferences, and workshops delivered to health professionals as part of training in the TEAMS approach.

Discussion

This study has been designed to assess the feasibility and acceptability of TEAMS therapy for bipolar disorder and to estimate its effect size ahead of a multi-site randomised controlled trial. TEAMS is based on an integrative cognitive model of bipolar disorder that converges existing theoretical approaches and service-user experiences. Whilst we have established a strong evidence-base for several components of the model, the impact of the intervention has been based on case studies and a case series to date.

A key strength of the study is its focus on specific needs and problems as identified by participants. This psychological intervention aims to address the huge impact of current symptoms (including depression, anxiety, irritability and hypomania) and functioning (including life goals around work and social life) that are less emphasised within relapse prevention approaches. The importance of this focus is reflected in the evidence for pervasive subsyndromal symptoms and through the ambivalence reported by many 'recovered' people with bipolar disorder as to whether their current well-being is satisfactory or complete, given the symptoms and difficulties with functioning they still experience. The importance of focusing treatment on current experiences converges with our model, which suggests that internal states during remission are on a continuum with those in episodes. Therapist and patient develop a shared understanding of the processes that may contribute to the maintenance and escalation of these states and are encouraged to 'broaden their bandwidth' of tolerance for such states. This involves understanding, facing, tolerating, enduring and even utilising internal states (e.g. fear, anger, excitability, happiness, sadness, tiredness) rather than struggling to suppress or manipulate them. While this therapeutic process often involves accessing past memories of these states and preparing for future situations within which they may arise, the focus is on their experience in the present and how this is managed.

The design of the study incorporates a number of additional strengths, including multiple longitudinal assessment points, which at the developmental stage of a treatment provides important information regarding therapeutic change. Also, researcher-blinded assessments, session-by-session process and adherence measures, and a qualitative analysis of participants' experiences of treatment as a whole, and TEAMS specifically. These components should ensure that the quality of the therapy is considered carefully and that it is modified and enhanced where necessary for future trials and implementations of TEAMS. The model would predict that TEAMS + TAU would lead to reductions in extreme appraisals of internal states compared to TAU and that, in turn, this cognitive style post-treatment (six month post-baseline), will be correlated with bipolar symptoms at 12 and 18 months when controlling for other demographic and clinical factors.

There are a number of limitations of the study. The clearest of these is that there is no active psychological intervention of equal intensity in the control group. At this stage of treatment development, it is first important to establish superiority to treatment-as-usual, especially as this patient group is rarely offered such an intervention in routine clinical practice. However, especially given recent developments for bipolar disorder within the Improving Access to Psychological Therapies system, it will be increasingly important to compare TEAMS with an alternative psychological approach. There are many potential active ingredients within TEAMS, including a variety of strategies and techniques, and so this study will not be able to identify the necessary, or most efficient, components. Nevertheless, we will draw upon the TEAMS interview for feedback from participants. Furthermore, participants have consented to recording their sessions for later analysis and so a potential focus of future research is to analyse sessions to identify the components that precede sudden gains in symptoms and functioning (cf. Tang & DeRubeis).

The study follows up participants no longer than 18 months post-randomisation. Given the scope of the study, this was felt to be appropriate, but longer follow-ups are essential to confirm the efficacy of an intervention for a mental health condition that can involve recurrent relapse. The study is also limited in

that it is focused on one regional area, and utilises only five clinically qualified clinical psychologists and/or CBT therapists who engage in weekly supervision with the developers of the therapy model. Thus, the potential for disseminating TEAMS to other groups, regions and health professionals requires further exploration. TEAMS also has the capacity to be applied to problematic mood swings outside the context of bipolar disorder (e.g. personality disorder; schizoaffective disorder), yet this was not explored in the current study.

In summary, this protocol is appropriately designed to establish the feasibility, acceptability and initial effect sizes ahead of a multi-site trial with a larger number of participants, therapists and a longer follow-up period. The provision of adherence and process measures and participant interviews will provide additional information and allow convergence with the burgeoning empirical literature regarding the TEAMS model of bipolar disorder.

Trial status

Participants are being recruited into the trial

Competing Interests

None.

Authors Contributions

WM and ST conceived of the trial, led the design and contributed to the writing of the report. SA AC and HL led the writing of the report. LD led the economic design and GD led the statistical design for the trial. NT provide service user input into trial design. RM provided led the diagnostic components of the trial and AM provided guidance for the design of the trial from its inception.

Role of funders and sponsors.

The trial sponsor is the University of Manchester and the contact is Lynn MacRae, [Faculty Research Practice Coordinator](#), [FMHS Research Office, 3.53 Simon Building, University of Manchester, Manchester, M15 6FH](#),

United Kingdom email: Lynne.k.macrae@manchester.ac.uk; telephone: [0161 275 5436](tel:01612755436). Neither the study sponsor or funders had any role in the study design; the collection, management, analysis and interpretation of the data; writing of the report, and the decision to submit the report for publication, nor did they have any ultimate authority over these decisions.

Acknowledgements

This trial has been supported by an NIHR Research for Patient Benefit Grant ID 11119 and additional support costs and infrastructure were provided by Greater Manchester West Mental Health NHS Foundation Trust. Philip Brawn, Ruth Searson, and Robert Griffiths provided therapy and clinical supervision for the trial. Professor Morriss's research time is funded by the National Institute of Health Research (NIHR) Collaboration for Leadership in Applied Health Research and Care (CLAHRC) East Midlands.

References

1. Akiskal HS, Bourgeois ML, Angst J, Post R, Möller H-J, R. H: **Re-evaluating the prevalence of and diagnostic composition within the broad clinical spectrum of bipolar disorders.** *J Affect Disord* 2000, **59**(supplement):S5-S30.
2. Bebbington P, Ramana R: **The epidemiology of bipolar affective disorder.** *Social Psychiatry and Psychiatric Epidemiology* 1995, **30**(6):279-292.
3. Judd LL, Akiskal HS, Schettler PJ, Coryell W, Endicott J, Maser JD, Solomon DA, Leon AC, Keller MB: **A prospective investigation of the natural history of the long-term weekly symptomatic status of bipolar II disorder.** *Archives of General Psychiatry* 2003, **60**(3):261-269.
4. Judd LL, Akiskal HS, Schettler PJ, Endicott J, Maser J, Solomon DA, Leon AC, Rice JA, Keller MB: **The long-term natural history of the weekly symptomatic status of bipolar I disorder.** *Archives of General Psychiatry* 2002, **59**(6):530-537.
5. Gitlin MJ, Swendsen J, Heller TL, Hammen C: **Relapse and impairment in bipolar disorder.** *American Journal of Psychiatry* 1995, **152**(11):1635-1640.
6. Morriss RK, Faizal MA, Jones AP, Williamson PR, Bolton C, McCarthy JP: **Interventions for helping people recognise early signs of recurrence in bipolar disorder.** *Cochrane database of systematic reviews (Online)* 2007(1):CD004854.
7. Sachs GS, Nierenberg AA, Calabrese JR, Marangell LB, Wisniewski SR, Gyulai L, Friedman ES, Bowden CL, Fossey MD, Ostacher MJ *et al*: **Effectiveness of adjunctive antidepressant treatment for bipolar depression.** *New England Journal of Medicine* 2007, **356**(17):1711-1722.
8. Beynon S, Soares-Weiser K, Woolacott N, Duffy S, Geddes JR: **Psychosocial interventions for the prevention of relapse in bipolar disorder: systematic review of controlled trials.** *The British journal of psychiatry : the journal of mental science* 2008, **192**(1):5-11.
9. Richardson TH: **Psychosocial interventions for bipolar disorder: A review of recent research.** *Journal of Medical Sciences* 2010, **10**(6):143-152.
10. Scott J, Colom F, Vieta E: **A meta-analysis of relapse rates with adjunctive psychological therapies compared to usual psychiatric treatment for bipolar disorders.** *The international journal of*

neuropsychopharmacology / official scientific journal of the Collegium Internationale

Neuropsychopharmacologicum 2007, **10**(1):123-129.

11. Gomes BC, Abreu LN, Brietzke E, Caetano SC, Kleinman A, Nery FG, Lafer B: **A randomized controlled trial of cognitive behavioral group therapy for bipolar disorder.** *Psychotherapy and psychosomatics* 2011, **80**(3):144-150.
12. Parikh SV, Zaretsky A, Beaulieu S, Yatham LN, Young LT, Patelis-Siotis I, Macqueen GM, Levitt A, Arenovich T, Cervantes P *et al*: **A randomized controlled trial of psychoeducation or cognitive-behavioral therapy in bipolar disorder: a Canadian Network for Mood and Anxiety treatments (CANMAT) study [CME].** *The Journal of clinical psychiatry* 2012, **73**(6):803-810.
13. Perich T, Manicavasagar V, Mitchell PB, Ball JR, Hadzi-Pavlovic D: **A randomized controlled trial of mindfulness-based cognitive therapy for bipolar disorder.** *Acta psychiatrica Scandinavica* 2013, **127**(5):333-343.
14. Mansell W: **An integrative formulation-based cognitive treatment of bipolar disorders: application and illustration.** *Journal of clinical psychology* 2007, **63**(5):447-461.
15. Beck AT: **Depression : clinical, experimental, and theoretical aspects.** London:: Staples Press; 1969.
16. Watkins E, Scott J, Wingrove J, Rimes K, Bathurst N, Steiner H, Kennell-Webb S, Moulds M, Malliaris Y: **Rumination-focused cognitive behaviour therapy for residual depression: A case series.** *Behaviour research and therapy* 2007, **45**(9):2144-2154.
17. Clark DM: **A cognitive approach to panic.** *Behaviour research and therapy* 1986, **24**(4):461-470.
18. Ehlers A, Clark DM: **A cognitive model of posttraumatic stress disorder.** *Behaviour research and therapy* 2000, **38**(4):319-345.
19. Fairburn CG, Harrison PJ: **Eating disorders.** *The Lancet* 2003, **361**(9355):407-416.
20. Mansell W, Paszek G, Seal K, Pedley R, Jones S, Thomas N, Mannion H, Saatsi S, Dodd A: **Extreme Appraisals of Internal States in Bipolar I Disorder: A Multiple Control Group Study.** *Cognitive Therapy and Research* 2010, **35**(1):87-97.

21. Mansell W, Powell S, Pedley R, Thomas N, Jones SA: **The process of recovery from bipolar I disorder: A qualitative analysis of personal accounts in relation to an integrative cognitive model.** *British Journal of Clinical Psychology* 2010, **49**(2):193-215.
22. Mansell W, Morrison AP, Reid G, Lowens I, Tai S: **The Interpretation of, and Responses to, Changes in Internal States: An Integrative Cognitive Model of Mood Swings and Bipolar Disorders.** *Behavioural and cognitive psychotherapy* 2007, **35**(05).
23. Mansell W, Colom F, Scott J: **The nature and treatment of depression in bipolar disorder: a review and implications for future psychological investigation.** *Clinical psychology review* 2005, **25**(8):1076-1100.
24. Mansell W, Pedley R: **The ascent into mania: a review of psychological processes associated with the development of manic symptoms.** *Clinical psychology review* 2008, **28**(3):494-520.
25. Mansell W, Scott J: **Dysfunctional beliefs in individuals with bipolar disorders.** In: *The psychology of bipolar disorder: new developments and research strategies.* edn. Edited by Jones SH, Bentall RP. Oxford: Oxford University Press; 2006: 73-90.
26. Seal K, Mansell W, Mannion H: **What lies between hypomania and bipolar disorder? A qualitative analysis of 12 non-treatment-seeking people with a history of hypomanic experiences and no history of major depression.** *Psychology and Psychotherapy: Theory, Research and Practice* 2008, **81**(1):33-53.
27. Dodd AL, Mansell W, Morrison AP, Tai S: **Extreme appraisals of internal states and bipolar symptoms: the Hypomanic Attitudes and Positive Predictions Inventory.** *Psychological assessment* 2011, **23**(3):635-645.
28. Kelly RE, Mansell W, Wood AM, Alatiq Y, Dodd A, Searson R: **Extreme positive and negative appraisals of activated states interact to discriminate bipolar disorder from unipolar depression and non-clinical controls.** *Journal of affective disorders* 2011, **134**(1-3):438-443.
29. Mansell W, Paszek G, Seal K, Pedley R, Jones S, Thomas N, Mannion H, Saatsi S, Dodd A: **Extreme appraisals of internal states in bipolar I disorder: A multiple control group study.** *Cognitive Therapy and Research* 2011, **35**(1):87-97.

30. Searson R, Mansell W, Lowens I, Tai S: **Think Effectively About Mood Swings (TEAMS): a case series of cognitive-behavioural therapy for bipolar disorders.** *Journal of behavior therapy and experimental psychiatry* 2012, **43**(2):770-779.
31. Beck AT, Ward CH, Mendelson M, Mock J, Erbaugh J: **An inventory for measuring depression.** *Archives of general psychiatry* 1961, **4**:561-571.
32. Paykel ES, Abbott R, Morriss R, Hayhurst H, Scott J: **Sub-syndromal and syndromal symptoms in the longitudinal course of bipolar disorder.** *The British Journal of Psychiatry* 2006, **189**(2):118-123.
33. Bech P, Rafaelsen OJ, Kramp P, Bolwig TG: **The mania rating scale: scale construction and inter-observer agreement.** *Neuropharmacology* 1978, **17**(6):430-431.
34. Hamilton M: **A rating scale for depression.** *Journal of neurology, neurosurgery, and psychiatry* 1960, **23**:56-62.
35. Bauer MS, Crits-Christoph P, Ball WA, Dewees E, McAllister T, Alahi P, Cacciola J, Whybrow PC: **Independent assessment of manic and depressive symptoms by self-rating: Scale characteristics and implications for the study of mania.** *Archives of General Psychiatry* 1991, **48**(9):807-812.
36. Spitzer RL KK, Williams JBW, Löwe B. : **A brief measure for assessing generalized anxiety disorder: the GAD-7. .** *Arch Intern Med* 2006, **166**(10):1092-1097.
37. Neil ST, Kilbride M, Pittb L, Nothard S, Welford M, Sellwood W, Morrison AP: **"The questionnaire about the process of recovery (QPR): A measurement tool developed in collaboration with service users": Corrigendum.** *Psychosis: Psychological, Social and Integrative Approaches* 2010, **2**(1):88-91.
38. Dodd AL, Mansell W, Sadhnani V, Morrison AP, Tai S: **Principal components analysis of the hypomanic attitudes and positive predictions inventory and associations with measures of personality, cognitive style and analogue symptoms in a student sample.** *Behavioural and cognitive psychotherapy* 2010, **38**(1):15-33.
39. Mansell W: **The Hypomanic Attitudes and Positive Predictions Inventory (HAPPI): A Pilot Study to Select Cognitions that are Elevated in Individuals with Bipolar Disorder Compared to Non-Clinical Controls.** *Behavioural and cognitive psychotherapy* 2006, **34**(04):467.

40. Kind P: **The EuroQoL instrument: an index of health-related quality of life.** *Quality of life and pharmacoeconomics in clinical trials* 1996, **2**: 191-201.
41. Blackburn I-M, James IA, Milne DL, Baker C, Standart S, Garland A, Reichelt F: **The revised cognitive therapy scale (CTS-R): Psychometric properties.** *Behavioural and Cognitive Psychotherapy* 2001, **29**(4):431-446.
42. Mansell W, Hodson S: **Imagery and memories of the social self in people with bipolar disorders.** In: *Imagery and the threatened self: Perspectives on mental imagery and the self in cognitive therapy.* edn. Edited by Lusia Stopa. Sussex: Routledge; 2009: 232.
43. Lam DH, Watkins ER, Hayward P, Bright J, Wright K, Kerr N, Parr-Davis G, Sham P: **A randomized controlled study of cognitive therapy for relapse prevention for bipolar affective disorder: Outcome of the first year.** *Archives of General Psychiatry* 2003, **60**(2):145-152.
44. Ball JR, Mitchell PB, Corry JC, Skillecorn A, Smith M, Malhi GS: **A randomized controlled trial of cognitive therapy for bipolar disorder: Focus on long-term change.** *Journal of Clinical Psychiatry* 2006, **67**(2):277-286.
45. Dunn G, Maracy M, Dowrick C, Ayuso-Mateos JL, Dalgard OS, Page H, Lehtinen V, Casey P, Wilkinson C, Vázquez-Barquero JL, Wilkinson G and The Outcomes of Depression International (ODIN) Group. **Estimating psychological treatment effects from an RCT with both non-compliance and loss to follow-up.** *British Journal of Psychiatry* 2013, **183**: 323-331.
46. Dunn G, Maracy M, Tomenson B. **Estimating treatment effects from randomized clinical trials with non-compliance and loss to follow-up: the role of instrumental variable methods.** *Statistical Methods in Medical Research* 2005, **14**: 369-395.

Table 1. *Summary of the measures taken at each assessment session and during therapy*

† indicates a self-report measure. All other measures are administered by a researcher as a clinical interview. ¹Pre-session client measures ²Post-session client measures³Adherence measures

Figure 1. *Consort diagram showing design of study*

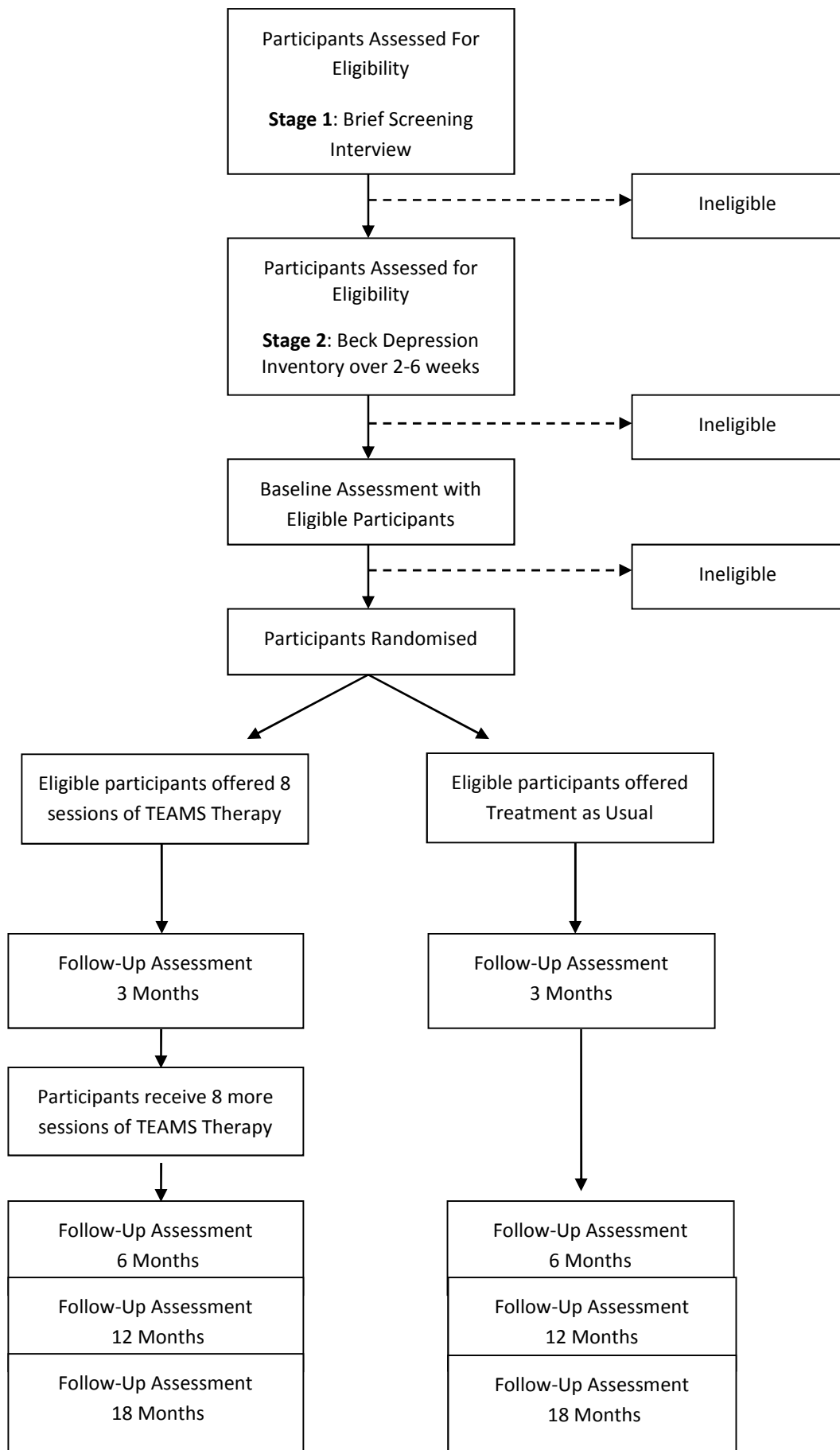


Figure 1

Additional files provided with this submission:

Additional file 1: TEAMSPROTOCOLTable 1.docx, 16K

<http://www.trialsjournal.com/imedia/5063792891396653/supp1.docx>

Additional file 2: 2014_08_15_13_58.pdf, 1667K

<http://www.trialsjournal.com/imedia/5210872731442825/supp2.pdf>