Help from home for depression: A randomised controlled trial comparing internet-delivered cognitive behaviour therapy with bibliotherapy for depression

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A R T I C L E   I N F O

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A B S T R A C T

Major Depressive Disorder (MDD) is a leading cause of the Global Burden of Disease. Cognitive Behavioural Therapy (CBT) is an effective treatment for MDD, but access can be impaired due to numerous barriers. Internet-delivered CBT (iCBT) can be utilised to overcome treatment barriers and is an effective treatment for depression, but has never been compared to bibliotherapy. This Randomised Controlled Trial (RCT) included participants meeting diagnostic criteria for MDD (n = 270) being randomised to either: iCBT (n = 61), a CBT self-help book (bCBT) (n = 77), a meditation self-help book (bMED) (n = 64) or wait-list control (WLC) (n = 68). The primary outcome was the Patient Health Questionnaire 9-item scale (PHQ-9) at 12-weeks (post-treatment). All three active interventions led to large within-group reductions in PHQ-9 scores at post-treatment (g = 0.88-1.69), which were maintained at 3-month follow-up, although there was some evidence of relapse in the bMED group (within-group g [post to follow-up] = 0.09-1.04). Self-help based interventions could be beneficial in treating depression, however vigilance needs to be applied when selecting from the range of materials available. Replication of this study with a larger sample is required.

1. Introduction

Depressive Disorders – including Major Depressive Disorder (MDD) – are a global public health concern and a leading cause of disease burden (Ferrari et al., 2013), with burgeoning individual and societal costs (Ustun et al., 2004). Recent NICE (2009) guidelines recommend non-drug interventions such as cognitive behavioural therapy (CBT) as a first line treatment for mild to moderate depression (www.nice.org.uk/CG90), but dissemination of CBT can be limited due to inadequate resources (Shapiro et al., 2003), particularly in certain geographical locations (Cavanagh, 2014). To address these treatment barriers, there has been a recent surge in innovative ways to deliver CBT (Andersson et al., 2012), specifically in the development of more accessible internet and computerised therapies for the treatment of common mental disorders (Marks et al., 2003).

Internet-delivered CBT (iCBT) programs are based on face-to-face CBT treatment manuals, and typically deliver psycho-education and the key CBT skills (e.g., cognitive restructuring, behavioural activation) via lessons or modules, supplemented with homework activities and practical exercises to complete between lessons. The concept of ‘guided iCBT’ typically takes the form of a therapist or coach supporting the patient throughout the course; this is a distinct difference to unguided iCBT which takes place without support - a purely ‘self-help’ format. Although unguided iCBT programs exist, recent findings indicate that guided iCBT is a more effective treatment for depression (Johansson and Andersson, 2012). Indeed, ‘clinician-assisted’ guidance (whereby a technician provides support and encouragement to patients and responds to general questions about the course) has been shown to be equally as effective as ‘clinician-assisted’ guidance provided by a qualified clinical psychologist (Titov et al., 2010).
The main strengths of iCBT include its accessibility to large amounts of people (Andersson, 2006), the privacy and convenience it invokes for the user (Wootton et al., 2011), and the ability to offer immediate feedback (Andersson et al., 2005). This treatment modality is also scalable, with the scope to be utilised within a stepped-care framework (Andrews and Williams, 2015). iCBT is cost-effective (Hedman et al., 2012) with a recent meta-analysis revealing that guided iCBT and face-to-face CBT produced equivalent overall effects in treating depression ($g = 0.05$ (95% CI: $-0.19$ to $0.30$) (Andersson et al., 2014). Although the majority of randomised trials have found positive effects of internet and computerised CBT on depression, there are some exceptions. A recent study found that guided computerised CBT did not achieve better outcomes for people with depression compared to usual (GP) care alone (Gilbody et al., 2015), although there are some concerns with the probity of the REEACT trial (Andrews et al., 2016).

The stages in CBT for depression can be generalised and represented in manuals and ‘self-help’ books. Bibliotherapy - including self-help books - are commonly used to teach self-help tools and strategies (Den Boer et al., 2004), with their usefulness to treat depression - when compared to controls - largely supported (Gregory et al., 2004). However as numerous self-help books are widely available for depression - whereby a specific self-help program is presented and is typically worked through independently - the evidence-base to support them is relatively small (McKendree-Smith et al., 2003), making their impact difficult to establish.

It is not known whether guided iCBT represents an advance in efficacy over the numerous CBT self-help books widely available for treating depression. This question is especially pertinent in an increasingly digital age; however access to the internet is not completely ubiquitous, especially for those on lower incomes (Cline and Haynes, 2001) and those living in rural and remote areas where internet access can be limited. Therefore the first aim of the current study was to compare an evidence-based guided iCBT course (Perini et al., 2009), the ‘Sadness’ program, versus unguided access to a self-help book based on CBT for depression. Participants in the CBT self-help book group were provided with a leading self-help book free of charge. ‘Beating the Blues’ (Tanner and Ball, 2012), which they completed without any form of guidance.

We also compared these two interventions to a third group who received a self-help meditation book, ‘Silence your Mind’ (Manocha, 2013). The self-help meditation book was also unguided and served as an active comparison condition to control for the effects of assessment, monitoring, and active learning and skills practice. We aimed to explore whether there were any significant differences between these active intervention groups at post-treatment. A variety of meditation-based interventions for depression are currently available; given that meditation can result in reduced psychological stress (Goyal et al., 2014), and has shown to be beneficial in treating acute and sub-acute depression (Jain et al., 2015), we expected the meditation book to have some advantageous effects on depression symptoms and to outperform a waiting list control group, but we anticipated that both CBT groups would outperform the meditation book in reducing depression. To our knowledge, this is the first study to compare guided iCBT with two self-help books, one book based on CBT and one based on meditation, for the treatment of MDD.

2. Method

2.1. Design

The current study utilised a randomised controlled trial design to compare an iCBT program, a CBT self-help book, a meditation self-help book, to a wait-list control for participants meeting diagnostic criteria for MDD. Participants were assessed at pre-treatment, mid-treatment, post-treatment and at 3 months post-treatment.

2.2. Inclusion/exclusion criteria

Inclusion criteria were: (i) aged over 18 years (ii) prepared to provide contact details of their General Practitioner (iii) had access to internet and printer, (iv) resident in Australia, (v) fluent in written and spoken English, (vi) a score between 5 and 23 on the Patient Health Questionnaire 9-item scale (PHQ-9), (vii) met criteria on the Mini International Neuropsychiatric Interview Version 5.0.0 (MINI) (Sheehan et al., 1998) for DSM-IV criteria for MDD on structured telephone diagnostic interview, (vii) if taking medications for anxiety or depression, were on a stable dose for at least 8 weeks at time of intake interview, and (viii) had not commenced face to face CBT within 4 weeks at time of intake interview. Exclusion Criteria included psychosis or bipolar disorder, drug or alcohol dependency, benzodiazepine use, severe depression PHQ-9 total scores > 24, or current suicidality.

2.3. Procedure

Participants were recruited via the Virtual Clinic (www.virtualclinic.org.au), the research arm of the Clinical Research Unit for Anxiety and Depression, based at St Vincent’s Hospital, Sydney and the University of New South Wales. Potential applicants viewed advertisements in the form of leaflets and posters and on social media. Applicants completed online screening questionnaires about demographic information and depressive symptoms as assessed on the PHQ-9 (Kroenke et al., 2001), after reading details about the study. Participants who met online screening criteria then participated in a brief telephone interview. Trained interviewers administered a structured diagnostic interview which consisted of the Mini International Neuropsychiatric Interview Version 5.0.0 (MINI) (Sheehan et al., 1998) MDD and risk assessment modules to confirm whether applicants met for DSM-IV criteria for MDD. Full risk assessment modules (where necessary) were completed by a psychiatry registrar to assess suicidal ideation and determine suitability for the study.

2.4. Participant flow

Details of the participant flow are displayed in Fig. 1. A total of 1143 applicants applied online for the study between May 2013 and February 2015. Of these, 549 applicants were excluded after completing initial online screening questions and received an email with information about alternative services. Five hundred and ninety four applicants passed the online screening phase and were telephoned for a diagnostic interview. A further 324 individuals were excluded at telephone interview stage, leaving 270 applicants who met inclusion criteria and were randomised.

Eligible participants were randomised based on a random number sequence generated at www.random.org by an independent person not involved in the study. Group allocation numbers were concealed from the interviewer with the use of opaque sealed envelopes, which were opened once the applicant was deemed eligible to participate. Participants provided electronic informed consent before being enrolled in the study. This study was approved by the Human Research Ethics Committee (HREC/13/SVH/29) of St Vincent's Hospital Sydney, Australia. The trial was prospectively registered on ANZCTR (ACTRN12613000502730).

2.5. Interventions

2.5.1. Sadness program (iCBT)

The Sadness program was delivered through www.virtualclinic.org.au. The program has been evaluated in four previous trials and its efficacy has been established (Perini et al., 2008, 2009, Titov et al., 2010, Watts et al., 2013) (between-groups effect sizes: $d = 0.8–1.3$, within-group pre-post effect sizes: $d = 0.9–1.0$). The program consisted
1,143 individuals applied to the Help from Home for Depression Program within timeframe (22.05.13 - 27.02.15)

Unsuccessful Application (n=549)
- Incomplete Application (n=249)
- PHQ-9 Score >23 (n=72)
- PHQ-9 Score <5 (n=10)
- Use of illicit drugs/more than 3 drinks/day (n=68)
- Taking Benzodiazepine medication (n=41)
- Suicidal ideation (n=28)
- Bipolar or Psychotic Disorder (n=40)
- No access to internet/computer/printer (n=27)
- Non-resident/under 18 years of age (n=13)
- Unable to speak English (n=1)

594 individuals met inclusion criteria and proceeded to telephone diagnostic interview with MINI 5.0.0

Unsuccessful Diagnostic Interview (n=224)
- Could not contact (n=85)
- Subclinical (n=104)
- Exclusion medications (n=21)
- Incorrect online application (n=3)
- Recent start or change of medication (n=7)
- About to start psychotherapy (n=8)
- Other reasons (e.g., moving house) (n=8)
- Participant in similar e-course (n=22)
- Risk issues (n=15)
- Bipolar Disorder (n=3)
- Decided not to proceed (n=47)
- Substance abuse (n=1)

270 participants met all inclusion criteria and were randomized into Intervention or WLC Groups

iCBT (n=61)
- Did not consent (n=1)
- Did not complete pretreatment questionnaires (n=6)
  Completed pretreatment questionnaires (n=54)
  - Withdrew during trial to start treatment/medication (n=2)
  - Withdrew for other reasons (n=4)
  Completed 6 lessons (n=32)
  Completed 5 lessons (n=33)
  Completed 4 lessons (n=38)
  Completed 3 lessons (n=42)
  Completed 2 lessons (n=44)
  Completed 1 lesson (n=48)
  Completed post-treatment questionnaires (n=33)
  Completed 3-month follow-up questionnaires (n=30)
  Completed 3-month follow-up interview (n=30)

hCBT (n=77)
- Did not consent (n=1)
- Withdrew before prequestionnaires (n=2)
  Completed prequestionnaires (n=74)
  - Did not complete questionnaires (n=20)
  - Withdrew during trial to start treatment/medication (n=5)
  - Withdrew for other reasons (n=2)
  Completed postquestionnaires (n=47)

hMED (n=64)
- Did not consent (n=1)
- Withdrew before prequestionnaires (n=2)
  Completed prequestionnaires (n=61)
  - Did not complete questionnaires (n=16)
  - Withdrew during trial to start treatment/medication (n=4)
  - Withdrew for other reasons (n=2)
  Completed postquestionnaires (n=41)
  Completed 3-month follow-up questionnaires (n=32)
  Completed 3-month follow-up interview (n=40)

WLC (n=68)
- Did not consent (n=3)
- Did not complete prequestionnaires (n=6)
  Completed prequestionnaires (n=59)
  - Withdrew during trial to start treatment/medication (n=2)
  - Withdrew for other reasons (n=2)
  - Did not complete questionnaires (n=7)

Fig. 1. Participant flow diagram.
of 6 online lessons completed over a 12 week period and participants were instructed to complete one lesson every 1–2 weeks of the program. It follows an illustrated story, with a main character who learns how to manage her depressive symptoms using CBT skills. The program content delivers evidence-based CBT including psycho-education, cognitive restructuring, graded exposure, problem solving, effective communication and relapse prevention, with downloadable weekly homework documents (PDF) containing practical tasks to implement between lessons as well as extra resources and patient ‘recovery stories’ to read. The course content is presented in Table 5. All participants in the iCBT program received regular automatic and manual email communication to notify them that a lesson was available and encouragement to complete the lesson. Participants were contacted via email or telephone by research staff (‘The Technician’, JS) after the first two lessons, then as requested, or by a clinician (psychiatry registrar) in response to an increase in distress or suicidal intent.

2.5.2. Beating the Blues (bCBT) (Tanner and Ball, 2012)

This is a self-help book designed to help people overcome depression and was posted to participants in this group, along with a short (one page) welcome letter from the book authors introducing participants to the concept of cognitive therapy and advising them to tackle the issues presented in the book systematically and one at a time. Participants were asked to work through each of the 12 chapters over 12 weeks and advised to read roughly one chapter per week. The book is self-guiding; the content is based on CBT and organised into various parts, including: Understanding Depression, Our Thinking Habits, Looking After Your Needs and Living with Someone Who is Depressed. The course content is presented in Table 6. Questionnaires and reminders to complete questionnaires were emailed to participants in this group with clinical contact occurring only if a participant reported active suicidal intent. In cases of elevated or increased distress response scores, participants were emailed to advise them where to gain additional support - including telephone counselling numbers, visiting their general practitioner or Emergency Services. No guidance or encouragement was offered.

2.5.3. Silence your mind (bMED) (Manocha, 2013)

This is a self-help book aimed at teaching people to meditate and was posted to participants in this group, along with a short DVD, information booklet and a brief (two page) letter from the author introducing participants to the mediation technique described in the book and advising them to practice the technique for at least 10 min per day. Participants were asked to work through each of the 13 chapters over 12 weeks and advised to read roughly one chapter per week. The book is self-guiding; the content is based on a meditative approach called Mental Silence and is organised into various parts, including: What is Meditation, Meditation is Mental Silence, Health, Wellbeing and the non-mind, Helping our Young People, Flow and Optimal Being, and The Brain in Meditation. The course content is presented in Table 7. Questionnaires and reminders to complete questionnaires were emailed to participants in this group with clinical contact only occurring if a participant reported active suicidal intent. In cases of elevated or increased distress response scores, participants were emailed to advise them where to gain additional support - including telephone counselling numbers, visiting their general practitioner or Emergency Services. No guidance or encouragement was offered.

2.5.4. Wait-list control group (WLC)

Participants randomised to the wait-list control (WLC) were able to continue with any course of treatment already specified at intake interview but were withdrawn from the study if they commenced a new treatment during the course of the 12 week waiting period, for example starting psychotherapy or medication for anxiety or depression. Questionnaires and reminders to complete questionnaires were emailed to participants in this group with clinical contact occurring in cases of elevated or increased distress response scores or when a participant reported suicidal intent. Participants had the choice of enrolling in either iCBT or one of the two self-books once their 12 week waiting period had ceased and they had completed all questionnaires.

2.6. Clinician contact

All participants completing the iCBT course received email and/or phone contact with the technician after the first two lessons to answer any questions about the program and to encourage them with the remainder of the course, then as requested. In cases of increased distress, particularly elevated distress and or suicidal intent was reported, the clinician made contact with the participant. Participants in the bCBT and bMED groups with elevated distress were emailed to inform them that their distress was elevated and where to get appropriate external support. The clinician did make contact with participants in these groups who reported suicidal intent.

2.7. Measures

2.7.1. Diagnostic status

The Mini International Neuropsychiatric Interview Version 5.0.0 (MINI (Sheehan et al., 1998)) MDD and risk assessment modules were administered at intake, and then again at 3 month follow-up for the intervention group participants (assessors were not blinded to the intervention condition).

2.7.2. Primary outcome measure

The Patient Health Questionnaire – 9 (PHQ-9, (Kroenke et al., 2001)) is a 9-item self-report scale that was used to assesses DSM-IV criteria for MDD. Participants were asked to rate the frequency of symptoms (including frequency of suicidal ideation with the question “thoughts that you would be better off dead, or of hurting yourself in some way”), over the past two weeks. The scale range is: 0 = not at all, 1 = several days, 2 = more than half of the days and 3 = nearly every day, with total scores ranging from 0 to 27. The PHQ-9 has shown to have excellent reliability and validity (0.86; (Kroenke et al., 2001)).

2.7.3. Secondary outcome measures

The Kessler-10 Psychological Distress Scale (K10; (Kessler et al., 2002)) is a 10-item self-report scale that was used to measure non-specific psychological distress over the past two weeks. Rated on a 5-point scale, higher scores indicate higher distress levels. The K10 has excellent psychometric properties (Furukawa et al., 2003).

The Generalised Anxiety Disorder 7- item scale (GAD-7; (Spitzer et al., 2006)) was used to measure generalised anxiety disorder symptoms over the past two weeks on the same 4-point scale as the PHQ-9. Scores range from 0 to 21, and this scale has good reliability and validity (0.83; (Spitzer et al., 2006)).

The NEO-Five Factor Inventory - Neuroticism Subscale (NEO-FFI, (Costa and McCrae, 1985)) is a 5-point self-report scale which was used to measure the personality dimension of Neuroticism and has good psychometric properties (Cuypers et al., 2005).

2.7.4. Expectancy of benefit and patient satisfaction

Prior to the start of their respective intervention, participants in iCBT, bCBT and bMED were asked to provide a treatment expectancy rating. Participants were asked to provide a rating ranging from 1 to 9 about how logical the therapy offered to them seemed, and how successful they thought the treatment would be in reducing their symptoms of depression (where 1 = not at all, 5 = somewhat, and 9 = very). To examine treatment satisfaction, participants were asked at post-treatment i) how satisfied they were that the program taught them the skills to manage depression and ii) their confidence in recommending the program to a friend with similar problems (range: 1 = not at all, 5 = somewhat, and 9 = very).
2.8. Outcome measurement

At baseline, participants provided basic demographic details (e.g., age, marital status, gender, employment and educational history), depression history and treatment, and current comorbid physical illnesses (6 questions from the 2007 National Survey of Mental Health and Wellbeing (Teesson et al., 2011).

All four groups completed outcome measures at baseline, mid-point (Lesson 4 for iCBT and week 6 for the book groups and WLC), post-treatment (one week after Lesson 6 for iCBT or at the end of the 12 weeks for the book groups and WLC) and at 3-month follow-up (for iCBT and the book groups only). The iCBT group completed the K10 before commencing every lesson.

2.9. Power calculations

Prior to commencement of the trial, a power calculation was conducted to set the minimum sample size needed. At 0.8 power (alpha = 0.05), 100 participants per group were needed to have the power to detect a 0.4 effect size difference in efficacy between groups. However, due to unanticipated difficulties with recruitment during the time-period set out for the study, the final sample (N = 270) fell short of the sample size needed (N = 400).

2.10. Statistical analyses

Groups were compared at baseline using t-tests and chi square analyses where the data consisted of categorical data. To explore the impact of the treatment groups on outcome, linear mixed models were conducted separately for each of the dependent variable measures with time, treatment group, and the time by group interaction entered as fixed factors in the model. For each group, planned contrasts were used to compare changes within and between-groups from baseline to post-treatment and 3-month follow-up. Between-group effect sizes using the pooled standard deviation and adjusted for sample size (Hedges g) were calculated to compare between groups at post-treatment and 3-month follow-up. Within-group effect sizes (Hedges g) were calculated between pre- and post-treatment and between pre- and 3-month follow-up for each of the intervention groups.

3. Results

3.1. Baseline demographics and between-groups differences

Two hundred and forty-eight participants completed baseline questionnaires and were included in the data analysis. For baseline demographics and sample characteristics see Table 1. There was no significant difference between the groups in age, F (50,197) = 1.09, p > 0.05, or pre-treatment PHQ-9 (F (22,225) = 0.452, p > 0.05), GAD-7 (F (19,183) = 0.768, p > 0.05), K-10 (F (28,219) = 1.41, p > 0.05), or NEO (F (25,225) = 1.35, p > 0.05) scores. Chi-square analyses demonstrated that there were no between-group differences in any other demographic characteristics including gender, marital status, educational status or employment status (ps > 0.05).

3.2. Adherence

In the iCBT group, 32 of the 54 participants completed all 6 lessons resulting in a 59% adherence rate. Of the 54 participants, 33 participants completed post-treatment assessments (61%) and 30 participants completed 3-month follow-up assessments (56%). In the bCBT group, 47 of the 74 participants provided post-treatment assessments (64%) and 43 completed 3-month follow-up assessments (58%). In the bMED group, 41 of the 61 participants provided post-treatment assessments (67%) and 32 completed 3-month follow-up assessments (52%). In the WLC group, 48 of the 59 participants provided complete data at the post-treatment time point (81%). (See Fig. 1 for participant flow). A total of 30/54 participants in the iCBT group (56%), 43/74 participants in the bCBT group (58%), and 40/61 participants in the bMED group (66%), completed a diagnostic interview to assess for MDD at 3-month follow-up.

3.3. Expectancy of benefit

There were no significant between-group differences in treatment expectancy ratings, either logical (F (2186) = 1.57, p > 0.05); iCBT: M = 5.81, SD = 2.10; bCBT: M = 6.41, SD = 2.11; bMED: M = 6.39, SD = 5.69) or successful ((F (2186) = 1.84, p > 0.05); iCBT: M = 5.28, SD = 1.65; bCBT: M = 5.85, SD = 1.77; bMED: M = 1.94, SD = 1.63). Results are reported in Table 2.

3.4. Primary and secondary outcome measures and effect sizes at post-treatment

Table 3 presents the results of all groups on primary and secondary outcome measures, and see Fig. 2 for PHQ-9 graph. There were significant group by time interactions for all of the primary and secondary outcome measures (PHQ-9: F (6401.84) = 6.26, p < 0.001, K-10:F (6, 399.82) = 6.39, p < 0.001, GAD-7: F (6, 372.87) = 7.38, p < 0.001, NEO: F (6, 397.28) = 5.30, p < 0.001).

There were large within-group effect sizes between baseline and post-treatment on all outcome measures in the iCBT, bCBT and bMED groups (g = 0.88–1.69). The reductions in the WLC group were moderate and significant on the PHQ-9 and K-10 (g = 0.56–0.55), but were small and not significant on the GAD-7 (g = 0.38, 95% CI: −0.01–0.76) and NEO (g = 0.13, 95% CI: −0.25–0.51).

Planned pairwise comparisons showed that each of the intervention groups had significantly lower PHQ-9, K-10, GAD-7 and NEO scores relative to WLC at post-treatment (ps < 0.01). There were large between-groups effect sizes on the PHQ-9 (g(5) = 0.86–0.98), and on the K-10 (g(5) = 0.83–0.91), except for a moderate effect size between bMED and WLC on K-10 scores at post-treatment (g = 0.60). We found moderate effect sizes between the intervention groups and WLC on the GAD-7 (g = 0.55–0.77), and moderate (iCBT v WLC; g = 0.64; bMED v WLC; g = 0.77) to large (bCBT v WLC; g = 0.85) effect size differences on the NEO.

There were no significant differences between iCBT, bCBT or bMED groups on any of the outcome measures at post-treatment (ps > 0.05). The between-groups effect sizes revealed small (but not significant) effect sizes favouring the bMED group over the bMED group on the GAD-7 and K-10 (g = 0.23), small but not significant effect sizes favouring iCBT over bMED on the K-10 (g = 0.31), and small but not significant effect sizes favouring bCBT over iCBT on the NEO (g = 0.21).

Of note, post-hoc power calculations indicated that at 0.8 power (α = 0.05), the actual sample size (minimum of 54 per group) had power to detect a between-groups effect size difference of d = 0.54.

3.5. Primary and secondary outcome measures and effect sizes at 3-month follow-up

To explore whether there were any between-groups differences between the iCBT, bCBT and bMED groups between post-treatment and 3-month follow-up, linear mixed-models were conducted separately for each of the dependent variable measures, with time (with post and follow-up scores entered in the model), treatment group, and the group by time interaction entered as fixed factors in the model. Results are reported in Table 4, and see Fig. 2 for PHQ-9 graph.

There was a significant group by time interaction between post-treatment and 3-month follow-up on the PHQ-9 (F (2, 99.60) = 4.19, p < 0.05). Between-group comparisons revealed that 3-month follow-up scores on the PHQ-9 were significantly higher, with moderate between-groups differences in the bMED relative to the bCBT group.
The group by time interactions were not significant for the remaining variables (K-10: F(2, 100.41) = 1.24, p > 0.05, GAD-7: F(2, 99.05) = 1.84, p > 0.05, NEO: F(2, 101.60) = 1.32, p > 0.05). Within-group comparisons between post-treatment and follow-up revealed that the bMED experienced moderate and significant deterioration in PHQ-9 scores between post-treatment and follow-up (within-group g = 0.56, 95% CI: 0.09–1.04). The changes in PHQ-9 scores in the iCBT and bCBT groups were not significant.

The group by time interactions were not significant for the remaining variables (K-10: F(2, 100.41) = 1.24, p > 0.05, GAD-7: F(2, 99.05) = 1.84, p > 0.05, NEO: F(2, 101.60) = 1.32, p > 0.05). Within-group comparisons between post-treatment and follow-up revealed that the bMED experienced moderate and significant deterioration in PHQ-9 scores between post-treatment and follow-up (within-group g = 0.56, 95% CI: 0.09–1.04). The changes in PHQ-9 scores in the iCBT and bCBT groups were not significant.

3.6. Diagnostic status at follow-up

Of the participants that completed a 3-month follow-up telephone interview, 24/30 (80%) in the iCBT group, 34/43 (79%) in the bCBT and 30/40 (75%) in the bMED group no longer met criteria for MDD. There were no statistically significant differences in the proportion of recovered patients in the intervention groups (χ²(4, n = 189) = 1.66,
Table 3
Estimated marginal means and within and between-groups effect sizes at post treatment for internet CBT, book CBT, meditation book and waiting list control groups.

|               | Baseline (T1) Mean (SD) | Mid (T2) Mean (SD) | Post (T3) Mean (SD) | 3 Month FU (T4) Mean (SD) | Within ES (95%CI) T1, T3 | Between ES (95%CI) T3 (post-treatment) |
|---------------|-------------------------|-------------------|---------------------|--------------------------|--------------------------|----------------------------------------|
| **PHQ-9**     |                         |                   |                     |                          |                          |                                        |
| iCBT          | 16.39 (5.03)            | 10.42 (4.88)      | 8.95 (4.77)         | 9.41 (4.71)              | 1.51 (1.00–2.00)         | 0.86 (0.39–1.32)**                    |
| bCBT          | 15.76 (5.03)            | 9.96 (4.93)       | 8.33 (4.78)         | 7.47 (4.72)              | 1.09 (0.70–1.48)         | 0.98 (0.56–1.41)**                    |
| bMED          | 17.26 (5.03)            | 10.95 (4.93)      | 9.05 (4.81)         | 11.73 (4.67)             | 1.55 (1.10–2.00)         | 0.83 (0.40–1.27)**                    |
| WLC           | 16.07 (5.03)            | 13.60 (4.97)      | 13.14 (4.91)        | –                        | 0.50 (0.11–0.89)         | –                                      |
| **K-10**      |                         |                   |                     |                          |                          |                                        |
| iCBT          | 33.67 (6.74)            | 25.63 (6.51)      | 21.98 (6.36)        | 22.92 (6.28)             | 1.69 (1.19–2.19)         | 0.91 (0.44–1.37)**                    |
| bCBT          | 32.26 (6.74)            | 24.99 (6.66)      | 22.49 (6.57)        | 21.83 (6.25)             | 1.47 (1.06–1.88)         | 0.83 (0.41–1.25)**                    |
| bMED          | 34.56 (6.74)            | 26.96 (6.59)      | 23.99 (6.42)        | 26.21 (6.22)             | 1.35 (0.91–1.78)         | 0.60 (0.17–1.03)**                    |
| WLC           | 32.69 (6.74)            | 29.68 (6.66)      | 27.93 (6.57)        | –                        | 0.55 (0.16–0.94)         | –                                      |
| **GAD-7**     |                         |                   |                     |                          |                          |                                        |
| iCBT          | 13.07 (4.58)            | 7.14 (4.43)       | 6.19 (4.32)         | 6.62 (4.26)              | 1.43 (0.95–1.92)         | 0.73 (0.28–1.19)**                    |
| bCBT          | 12.36 (5.39)            | 8.16 (4.54)       | 5.98 (4.38)         | 5.56 (4.33)              | 1.06 (0.67–1.45)         | 0.77 (0.36–1.20)**                    |
| bMED          | 13.27 (5.31)            | 8.03 (4.49)       | 7.01 (4.38)         | 8.11 (4.19)              | 1.00 (0.58–1.42)         | 0.55 (0.12–0.97)**                    |
| WLC           | 11.86 (4.58)            | 10.77 (4.53)      | 9.44 (4.43)         | –                        | 0.38 (–0.01–0.76), ns    | –                                      |
| **NEO**       |                         |                   |                     |                          |                          |                                        |
| iCBT          | 37.59 (6.49)            | 33.57 (6.26)      | 30.72 (6.10)        | 29.90 (6.09)             | 0.94 (0.48–1.39)         | 0.64 (0.18–1.10)**                    |
| bCBT          | 36.16 (6.49)            | 31.90 (6.38)      | 29.39 (6.11)        | 27.54 (6.03)             | 1.02 (0.63–1.41)         | 0.85 (0.43–1.27)**                    |
| bMED          | 36.21 (6.49)            | 32.29 (6.38)      | 29.83 (6.16)        | 30.40 (5.95)             | 0.88 (0.47–1.30)         | 0.77 (0.35–1.21)**                    |
| WLC           | 35.76 (6.49)            | 34.80 (6.40)      | 34.74 (6.31)        | –                        | 0.13 (–0.25–0.51), ns    | –                                      |

Note: **p < 0.001, ns = p > 0.05. GAD-7 = Generalised Anxiety Disorder 7-item scale; iCBT = internet-based cognitive behaviour therapy group; K-10 = Kessler 10-item Psychological Distress scale; NEO = NEO-Five Factor Inventory – Neuroticism Subscale; PHQ-9 = The Patient Health Questionnaire-9, WLC = waitlist control group. Within-group ES = Hedges g between-group ES = Hedges g with Hedges pooled SD. T1 = baseline, T2 = mid-treatment, T3 = post-treatment, T4 = 3-month follow-up. N's T1: iCBT = 54, bCBT = 74; bMED = 61, WLC = 59; T2: iCBT = 40, bCBT = 61; bMED = 50, WLC = 53; T3: iCBT = 33, bCBT = 47; bMED = 41, WLC = 48; T4: iCBT = 30, bCBT = 43; bMED = 32.
### 3.7. Clinical significance

Following Jacobson and Truax (1991) reliable change index (RCI) values were calculated for the PHQ-9 scores to determine the proportion of each group who evidenced reliable improvements (or deterioration) in PHQ-9 scores between baseline and follow-up. RCI values were calculated using test-retest reliability values of 0.84 from Kroenke et al. (2001). In order to calculate standard error of measurement values, standard deviations were derived from current sample (PHQ-9 pre-treatment pooled SD = 4.37). A change of 5 points on the PHQ-9 was considered to be statistically reliable change. Of the iCBT group, 21/54 (61.1%) reliably improved compared to 36/74 (48.6%) in the bCBT group, 31/61 (50.8%) in the bMED group, and 21/59 (35.6%) in the WLC group. There was no evidence of deterioration in the iCBT group, but there was evidence of deterioration on the PHQ-9 for one in the bCBT group (1.4%), one in the bMED group (1.6%), and 6 individuals in the WLC group (10.2%).

### 3.8. Missing data analysis

Because there was a significant proportion of missing data on the primary outcome measure (PHQ-9) at post-treatment, we analysed missing data patterns. First, Little’s MCAR test was not significant for post-treatment scores ($\chi^2 = 0.57, p = 0.45$), or follow-up scores ($\chi^2 = 1.29, p = 0.26$). Second, we conducted independent samples t-tests to explore whether there were any significant differences between those with missing data versus those with complete data at post-treatment. Compared to participants who provided post-treatment data ($n = 169$: M = 32.73, SD = 6.18), those who had missing data were significantly more distressed at baseline on the K-10 ($n = 79$: M = 34.32, SD = 5.13) ($t(246) = 1.99, p = 0.048$), however there were no significant differences between those with and without data at post-treatment on age or other demographics (e.g., marital and education status), baseline PHQ-9 scores, GAD-7 scores, or NEO scores ($p > 0.05$). There was also no statistically significant relationship between treatment group and the proportion of participants with missing data ($\chi^2 (3) = 6.73, p = 0.08$).

In addition to these analyses, a multiple imputation procedure was used to account for the data lost at post-treatment and 3-month follow-up (for the intervention groups only). We assume the value of the data lost relates to observed variables in the dataset. The multiple imputation procedure was implemented in SPSS, using a chained equations multiple imputation procedure, with five imputations. ANOVAs were conducted on imputed post-treatment and follow-up PHQ-9 data to explore whether the significant between-group differences in the main mixed models analysis remained after imputation. The pooled estimates for the imputed data for the four groups at post-treatment were: iCBT: $M = 8.21, SE = 1.03$, bCBT: $M = 8.15, SE = 0.86$, bMED: $M = 8.73, SE = 0.78$, WLC: $M = 13.17, SE = 0.83$. The between-group differences between the three intervention groups and WLC remained significant for post-treatment PHQ-9 scores ($p < 0.001$), and there were no significant differences between the three intervention groups (iCBT, bCBT, bMED). In addition, mirroring the findings from the mixed models in the main analysis, there was a significant group difference on PHQ-9 scores at 3-month follow-up, with the bCBT reporting significantly lower PHQ-9 scores than the bMED group. The pooled estimates for the imputed data for the three treatment groups at follow-up were: iCBT: $M = 8.92$, bCBT: $M = 7.78$, SE = 1.21, SE = 1.39, bMED: $M = 11.64, SE = 1.04$.

### 3.9. Clinician contact

There was a significant difference across groups in the total time

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**Table 4**

| PHQ-9 | Post (T3) | 3 Month FU (T4) | Within iCBT T3, T4 | Within ES (95%CI) T3, T4 | Between ES (95%CI) T4 |
|-------|----------|----------------|-------------------|-------------------------|------------------------|
|       | Mean (SD) | Mean (SD)      |                   |                         |                        |
| iCBT  | 8.70 (5.28) | 9.04 (5.20) | 0.61 (1, 100.10) | 0.06 (−0.43-0.55) | −0.27 (−73.20), ns |
| bCBT  | 8.34 (5.28) | 7.62 (5.25) | 0.92 (1, 97.24)  | 0.13 (−0.29-0.54) | 0.46 (−0.04-0.97), ns |
| bMED  | 8.81 (5.31) | 11.47 (5.15) | 3.02 (1, 100.21)* | 0.56 (0.09-1.04)* | 0.73 (0.26-1.20)* |
| K-10  |          |               |                   |                         |                        |
| iCBT  | 21.89 (7.35) | 22.50 (7.23) | 0.49 (1, 101.46) | 0.07 (−0.43-0.56) | −0.10 (−0.56-0.37), ns |
| bCBT  | 22.16 (7.34) | 21.79 (7.21) | 0.37 (1, 98.87)  | 0.05 (−0.36-0.47) | 0.42 (−0.08-0.93), ns |
| bMED  | 23.53 (7.43) | 25.60 (7.13) | 1.77 (1, 100.66) | 0.30 (−0.16-0.77) | 0.52 (−0.06-0.99), ns |
| GAD-7 |          |               |                   |                         |                        |
| iCBT  | 6.12 (4.60) | 6.32 (4.55) | 0.77 (1, 100.39) | 0.13 (−0.36-0.63) | −0.18 (−0.64-0.29), ns |
| bCBT  | 5.90 (4.66) | 5.50 (4.59) | 0.60 (1, 96.85)  | 0.10 (−0.31-0.52) | 0.49 (−0.01-1.00), ns |
| bMED  | 7.06 (4.67) | 8.56 (4.47) | 2.01 (1, 99.59)* | 0.28 (−0.19-0.74) | −0.68 (0.21-1.16)* |
| NEO   |          |               |                   |                         |                        |
| iCBT  | 30.26 (7.64) | 29.58 (7.61) | 0.54 (1, 102.05) | 0.07 (−0.43-0.56) | −0.28 (−0.75-0.18), ns |
| bCBT  | 29.19 (7.68) | 27.41 (7.54) | 1.71 (1, 2.94)  | 0.25 (−0.16-0.67) | 0.16 (−0.33-0.66), ns |
| bMED  | 30.07 (7.75) | 30.83 (7.41) | 0.63 (1, 102.50) | 0.10 (−0.36-0.56) | 0.45 (0.001-0.91), ns |

Note. **p < 0.001, *p < 0.01, ns = p > 0.05. GAD-7 = Generalised Anxiety Disorder 7-item scale; iCBT = internet-based cognitive behaviour therapy group; K-10 = Kessler 10-item Psychological Distress scale; NEO = NEO-Five Factor Inventory – Neuroticism Subscale; PHQ-9 = The Patient Health Questionnaire-9, WLC = waitlist control group. Within-group ES = Hedges g; between-group ES = Hedges g with Hedges pooled SD. T3 = post-treatment, T4 = 3-month follow-up. T3: iCBT = 33, bCBT = 47; bMED = 41, WLC = 48; T4 iCBT = 30, bCBT = 43; bMED = 32. 32
(minutes) spent contacting the participants (total time spent: \(F(3, 244) = 15.35, p < 0.001\)); given the iCBT group was guided, planned comparisons indicated significantly more time was spent by the technician contacting iCBT participants compared to each of the other three groups (\(p's < 0.001\)). The technician spent an average of 9.7 min (SD = 5.02, range = 3–21) emailing and calling each participant in the iCBT group during the treatment course (including follow-up period). There were no overall between-group differences in the time spent contacting the participants by the clinician (\(F(3, 244) = 1.84, p > 0.05\)). The majority of participants across all groups did not require contact with a clinician (iCBT: 81%; bCBT: 95%; bMED: 92%; WLC: 90%).

### 3.10. Patient satisfaction

There were no overall between-group differences in either treatment satisfaction (\(F(2, 118) = 1.98, p > 0.05\)); iCBT: M = 6.45, SD = 1.84; bCBT: M = 6.02, SD = 2.14; bMED: M = 5.49, SD = 2.25), or confidence in recommending the program to a friend (\(F(2, 118) = 1.57, p > 0.05\)); iCBT: M = 6.73, SD = 2.32; bCBT: M = 6.15, SD = 2.41; bMED: M = 5.71, SD = 2.62).

### 4. Discussion

The current study had a unique aim, to explore whether there were any post-treatment differences between unguided self-help books and guided iCBT in treating MDD. At post-treatment, all three interventions - when compared with the wait-list control group - were associated with significantly lower depressive symptoms, distress, anxiety and neuroticism. We found large between-group differences (\(d = 0.83\) to 0.98) between the active intervention groups compared to WLC on the PHQ-9, and medium to large differences on the secondary outcome measures (K10: \(d = 0.60\) to 0.91; GAD7: \(d = 0.55\) to 0.77; NEO: \(d = 0.64\) to 0.85). These findings provide support for the efficacy of all three intervention groups compared to waiting list.

Although there were no significant differences between the intervention groups at post-treatment, there were significant differences at 3-month follow-up on one outcome (the PHQ-9). Results revealed that although the meditation book was effective in reducing depressive symptoms immediately after treatment, these gains were not sustained at follow-up. In the group that received the meditation self-help book, there was evidence of worsening of symptoms on the PHQ-9 between post-treatment and 3-month follow-up, corresponding to a medium effect size (\(g = 0.56\)). In contrast, the changes in PHQ-9 scores within the iCBT and bCBT groups between post-treatment and follow-up were not significant, suggesting the improvements in these groups were maintained. We note that the relapse in symptoms in the bMED group appeared only on the self-reported depressive symptoms on the PHQ-9, and not the other outcome measures assessing anxiety, distress and disability. In addition there were no group differences in the diagnostic status of participants according to diagnostic interviews at 3-month follow up, with a high proportion of participants in all three intervention groups no longer meeting criteria for MDD at follow-up; 75% in bMED, 79% in iCBT and 80% in iCBT. We acknowledge that the group difference at follow-up was only found on the depression outcome measure (the PHQ-9), and not the other outcome measures (e.g., anxiety, distress, neuroticism) so this finding needs to be replicated before definitive conclusions can be made. However, the study outcomes provide evidence that the benefits of CBT appear to last beyond the completion of treatment, and that the longer-term benefits of self-help meditation-based interventions for treatment of depression warrant further investigation (Manocha et al., 2011).

The study findings support the notion that iCBT is a reputable treatment for depression (Hedman et al., 2012); yet, we also found strong evidence supporting the CBT self-help book, ‘Beating the Blues’. Our study results seem to replicate recent findings comparing iCBT with

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**Table 5**

| Lesson title          | Lesson content                                                                 | Homework activities                      |
|----------------------|--------------------------------------------------------------------------------|------------------------------------------|
| Lesson 1: The diagnosis | • Depression explained  
• The vicious cycle of depression: physical, cognitive  
+ behavioural symptoms  
• How is depression treated  
• Recognise own symptoms of depression  | • What are your symptoms of depression?  |
| Lesson 2: Monitoring your thoughts and activities | • Role of activity in depression + monitoring activity  
• Recognising unhelpful thinking  
• Reducing ruminating  
• Shifting attention  
• Restoring sleep patterns  | • Activity monitoring form  
• Thought monitoring form  |
| Lesson 3: Learning to improve your activities and thoughts | • Activity planning  
• Challenging unhelpful thinking  
• Positive and negative beliefs about Ruminating + worry  
• Recording positives  
• Learning about the role of avoidance  
• Facing fears gradually with exposure  | • Challenging negative beliefs about ruminating and worry sheet  
• Your activity planning Worksheet  
• Your thought challenging worksheet  
• Your positives hunt worksheet  
• Structured problem solving worksheet  
• Graded exposure: identify feared situation + develop graded hierarchy  
• Facing fears worksheet  
• Your thought challenging worksheet  
• Your positives hunt worksheet  |
| Lesson 4: Facing Fears | • Structured problem solving  
• Learn about the role of avoidance  
• How to get more help with depression  
• Your evaluation  | • Your relapse prevention plan  
• Self-evaluation  |
| Lesson 5: Being assertive | • Being assertive and challenging unhelpful thoughts about assertiveness  
• Effective communication  
• Recording positives  
• Thought challenging  | Extra resources:  
Frequently asked questions (FAQs), 100 Things To Do (pleasant activity planning ideas), In Case of Emergency, Good Sleep Guide, Structured Problem Solving, About Assertiveness  |
| Lesson 6: Preventing relapse and getting even better | • Lapses and relapses explained  
• How to prevent lapses + what to do if a lapse happens  
• How to get more help with depression  
• Your evaluation  | Worksheets (Extra Resources):  
Activity Planning & Monitoring Worksheet, Activity Planning Worksheet, Daily Positives Hunt Worksheet, Thought Challenging Worksheet, Structured Problem Solving Worksheet, Facing Fears Worksheet, Conversation Skills  |
Table 6
Chapter content and activities to complete in 'Beating the Blues' (bCBT).

| Chapter title                                      | Chapter content                                                                 | Tasks to complete                                                                                   |
|---------------------------------------------------|-------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------|
| Chapter 1: Understanding depression               | • Types of depression • Recognising symptoms of depression • How CBT can help • Monitoring your mood | • Exercise: Monitor your moods                                                                       |
| Chapter 2: What makes people vulnerable to depression | • Coping strategies • Factors triggering depression • Protective Factors against depression | • Exercise: Personality Characteristics and Risk of Depression • Exercise: Life Events Questionnaire |
| Chapter 3: How you think your way into feeling down | • The ABC of thinking, feeling + behaving • Automatic Thoughts                | • Exercise: Daily Thought Monitoring Form                                                               |
| Chapter 4: Breaking the lethargy circuit           | • Common negative thoughts • What causes lack of motivation • Good Reasons for Staying Active When Feeling Down • Common negative thoughts and testing them • Setting Weekly Goals | • Exercise: Focus on Doing • Exercise: Activity Plan • List Pleasant Events • Exercise: Treating Yourself |
| Chapter 5: Our thinking habits: The good, the bad and the ugly | • Thinking habits and distortions • Perfectionism and setting unrealistic expectations • Converting positives into negatives • Overgeneralising and exaggeration • Catastrophising • Ruminating | • Exercise: Listing positives • Summary of Unhelpful thinking habits • Exercise: Identifying unhelpful thinking habits |
| Chapter 6: Changing old thinking habits: the secret to being a happier person | • Five steps to becoming a happier person • Re-programming thoughts • Common negative thoughts and alternatives | • Exercise: The Daily Diary • Exercise: Collecting evidence to test out beliefs • Exercise: Gaining perspective • Exercise: Building resilience • Summary: Learning to question + change thinking |
| Chapter 7: Are you fully awake? Cultivating mindful awareness | • What is mindful awareness? • Steps to cultivating mindful awareness • Integrating mindful awareness into everyday life • Breaking old habits through mindful awareness • Calming the nervous system | • Exercise: Awareness of the five senses • Exercise: Awareness of the body • Exercise: Awareness in eating • Exercise: Meditation on the breath |
| Chapter 8: Boosting your self-esteem               | • Belief systems and self-esteem • How to improve your self-esteem             | • Exercise: Challenging perfectionism • Exercise: Enjoy your own company • Exercise: Recording your strengths • Exercise: What did I do well today? • Exercise: Keeping a gratitude book |
| Chapter 9: Overcoming loneliness and jealousy      | • Thinking your way into loneliness • Developing resilience: how you can feel good about being alone • Overcoming jealousy • How partners can help | • Exercise: Managing your time alone                                                                 |
| Chapter 10: Dealing with self-harm, hopelessness and suicidal urges | • Understanding, managing and reducing self-harm • Understanding and managing hopelessness • Suicidal impulses | • Exercise: When you were happy • Exercise: Positives Hunt • Exercise: Challenging suicidal thoughts |}

bibliotherapy; in spite of the scant literature available on self-help treatment books, comparisons between iCBT and bibliotherapy for OCD (Wootton et al., 2013) and for social phobia (Furmark et al., 2009) exist and showed both modes of presentation to be equivalent. However it is unknown whether the same outcomes for the self-help books would be replicated if they were delivered outside a controlled environment, or even in routine clinical practice. Participants in this trial were rigorously assessed at intake, and were instructed to read one chapter per week over a strict 12-week period. They also completed regular routine assessments throughout the study period and were closely monitored for suicidality. It is also unclear whether these results generalise to other self-help treatments for depression – specifically other self-help books – that are widely available within the public domain.

It is not known what the role – if any - individual treatment preferences (for the internet or book mode) prior to study randomisation, and any individual depression treatment history, had on final outcomes. Over a third of the sample reported frequent and episodic depression (8+ episodes). It has been suggested that people who have experienced repeated episodes of depression may gain less from internet-delivered self-help treatments (Andersson et al., 2004), although a recent meta-analysis of psychotherapies for depression in adults purported that relapse can be prevented by using CBT, particularly in those with numerous previous episodes (Cuijpers, 2017).
| Chapter title | Chapter content | Tasks to complete |
|---------------|-----------------|-------------------|
| Chapter 1: The ants | • Introduction to meditation  
• Finding meditation |  |
| Chapter 2: The path to research | • A path less trodden  
• Mental silence – the new frontier  
• Words from meditators |  |
| Chapter 3: Defining Meditation | • Defining meditation  
• The placebo effect |  |
| Chapter 4: Specific effects – understanding the evidence | • Randomised controlled trials and meditation  
• An overview of the research  
• The right RCT | • How this applies to you |
| Chapter 5: Mental silence – a unique discovery | • The architecture of our thinking  
• Contemplation, relaxation, mindfulness and mind-emptiness – what’s the difference?  
• A Buddhist perspective on the mind, mindfulness and mind-emptiness  
• Can beginners experience mental silence easily?  
• What about long term meditators? |  |
| Chapter 6: The ancient paradigm | • Thinking and being  
• The history of mental silence  
• Present day  
• Words from meditators |  |
| Chapter 7: A thought experiment that ends in silence | • Thought experiment  
• Advice about posture  
• Standard affirmations | • How to begin – what am I aiming to experience?  
• Guided meditation sequence  
• Thought-stopping sequence  
• The Affirmations |
| Chapter 8: Stress – the noise in the mind | • The stress epidemic  
• Signs of stress and What is stress?  
• Can meditation modify the way that our mind reacts?  
• Mental silence and work stress  
• Is meditation uniquely effective in reducing stress?  
• A physiological paradox  
• Implications: cognitive reappraisal and stress management  
• Words from meditators |  |
| Chapter 9: Beyond the mind-body connection | • Case studies  
• Menopause  
• Asthma  
• The long term effects of meditation on health  
• Non-thought and wellbeing  
• Mind-body medicine  
• Words from meditators | • Enhancing your experience  
• Short affirmation sequence  
• Tying up your attention and protecting your experience  
• Conscious breathing and Mindful breathing  
• Bringing back your focus  
• Attention-based strategies  
• Other centering strategies  
• Footsoaking and Earthing  
• Ways to assess your progress |
| Chapter 10: Young minds | • Taming the wild things  
• Rebuilding resilience  
• The ADHD clinic  
• Case studies and The Results |  |
| Chapter 11: Meditation in the class room | • Evaluating meditation in schools  
• Emotional and behavioural wellbeing  
• Engagement  
• Teacher stress  
• Relationships  
• Preventing depression  
• Implications  
• Words from meditators | • Meditation for children  
• The magic bubble  
• The golden string  
• Tying up your attention  
• Rainbow protection |
| Chapter 12: Meditation and flow | • About flow  
• Sport, mental clarity, flow and meditation  
• Flow and mental silence in novice meditators  
• Case study  
• Flow and mental silence in novice meditators  
• In conclusion  
• Implications  
• Words from meditators |  |
| Chapter 13: Brain, mind and non-mind | • Introducing the brain  
• Neuroplasticity  
• Changing negative mind habits  
• This is your brain in meditation  
• Mental silence and the brain  
• The molecules of silence  
• The final frontier | • Frequently Asked Questions  
• Appendix and Acknowledgements |
Interestingly, only a small number of the sample were undertaking other psychotherapeutic treatments (e.g. face to face CBT) for their depression at intake assessment (10%), but a significantly larger proportion of the total sample were on stable doses of medications for anxiety or depression (44%). For many therefore, the study interventions were adjuncts to existing treatments highlighting the potential usefulness of self-help treatments for depression with concurrent (pharmacological) treatments.

The issue of clinician time spent is also of note as the majority of participants in all three intervention groups did not require any contact with a clinician throughout the duration of the trial (including follow-up stage), indicating the cost-effectiveness of the study interventions. The lack of post-treatment differences between the guided and unguided interventions is also of note but seemingly not uncommon; in a comparison between guided and unguided internet therapy for depression, no significant effects were found highlighting the ability of internet treatments to be delivered successfully without any form of guidance (Berger et al., 2011). Furthermore, a recent study found that unguided bibliotherapy achieved similar results to an unguided iCBT and guided iCBT group for the treatment of health anxiety, and that all three treatments were significantly more effective than a control group at post-treatment (Hedman et al., 2016), mirroring the current study findings. An interesting finding of this study was that the therapist-guided iCBT group completed significantly more modules than the unguided iCBT group, supporting the association that clinician contact during the iCBT program is associated with improved adherence (Hilvert-Brace et al., 2012).

The current results must be considered in light of limitations. First, the study was underpowered to detect significant differences between the three intervention groups, and as we conducted multiple group comparisons there is risk of Type II error. Future replications of this study with a larger sample are needed to detect these effects. A significant limitation of the study is that no post-treatment data existed for around 30% of the sample within the intervention groups; although high drop-out rates can be a common drawback of self-help trials (Eysenbach, 2005), the adherence for this research study was generally lower than expected. However this was true for iCBT (61% completed post-treatment questionnaires), as it was for the self-help books (64% completed post-treatment questionnaires in iCBT and 58% completed post-treatment questionnaires in BMED). Some participants within the iCBT group did remark that they had issues accessing the internet as regularly as the program required.

Thirdly, the study design excluded follow-up comparison between WLC and the three intervention groups; the follow-up period was also brief (only 3 months) - the enduring benefits of iCBT, and the long-term superiority of iCBT over WLC warrant further examination (Mahoney et al., 2014). It is of note that not all participants in the intervention groups were able to be contacted to complete the three-month follow-up diagnostic assessment, which may lead to an over-estimate of the effects of treatment on remission rates from MDD. A further drawback is that the researchers are not clear, due to the limited contact with these participants, on how much a participant randomised to the self-help books adhered to the content and materials presented within the treatment program of the books themselves. It has been asserted that bibliotherapy can be challenging when the reader is not able to understand the materials, which can lead to a loss of motivation and the individual giving up (Team, H. o. N. D. I. P. and Usher, 2013).

In summary, to overcome the burden of depressive disorders, accessible and effective, low-cost interventions are required, and self-help treatments (including bibliotherapy) could be utilised to address these treatment barriers. These study findings are promising; participants who met criteria for mild, moderate and severe (but not very severe) depression engaging in a CBT or meditation-based intervention had significantly reduced depressive symptoms in comparison to participants waiting for treatment at post-treatment. We do include a cautionary note however, that vigilance does need to be applied when selecting from the vast array of self-help treatments publicly available, and would strongly recommend that patients are formally assessed and properly monitored by professionals when embarking on self-help treatments for depression. Replication of this study with a larger sample is required.

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