Group rumination-focused cognitive-behavioural therapy (CBT) v. group CBT for depression: phase II trial

Morten Hvenegaard, Stine B. Moeller, Stig Poulsen, Matthias Gondan, Ben Grafton, Stephen F. Austin, Morten Kistrup, Nicole G. K. Rosenberg, Henriette Howard and Edward R. Watkins

Abstract

Background. Although cognitive-behavioural therapy (CBT) is an effective treatment for depression, less than half of patients achieve satisfactory symptom reduction during treatment. Targeting known psychopathological processes such as rumination may increase treatment efficacy. The aim of this study was to test whether adding group rumination-focused CBT (RFCBT) that explicitly targets rumination to routine medical management is superior to adding group CBT to routine medical management in treating major depression.

Methods. A total of 131 outpatients with major depression were randomly allocated to 12 sessions of RFCBT v. group CBT, each in addition to routine medical management. The primary outcome was observer-rated symptoms of depression at the end of treatment measured on the Hamilton Rating Scale for Depression. Secondary outcomes were rumination at post-treatment and depressive symptoms at 6 months follow-up (Trial registered: NCT02278224).

Results. RFCBT significantly improved observer-rated depressive symptoms (Cohen’s d 0.38; 95% CI 0.03–0.73) relative to group CBT at post-treatment on the primary outcome. No post-treatment differences were found in rumination or in depressive symptoms at 6 months follow-up, although these secondary analyses may have been underpowered.

Conclusions. This is the first randomized controlled trial providing evidence of benefits of RFCBT in major depression compared with CBT. Group RFCBT may be a beneficial alternative to group CBT for major depression.

Cognitive-behavioural therapy (CBT) is a recommended psychological treatment for unipolar depression with many randomized controlled trials (RCTs) providing evidence for its efficacy (DeRubeis et al., 2005; Cuijpers et al., 2016). However, it only achieves remission for less than half of treated patients (DeRubeis et al., 2005; Cuijpers et al., 2014). CBT targets key mechanisms in the maintenance of depression such as negative thinking and behavioural avoidance. One potential way to improve the efficacy of CBT is to adapt it to specifically target another key mechanism in depression, namely rumination (Watkins, 2015). Rumination, defined as repetitive negative thinking about the symptoms of depression and their causes and consequences (Nolen-Hoeksema and Morrow, 1991), has been shown to predict the onset, severity and duration of depressive episodes (Nolen-Hoeksema, 2000; Nolen-Hoeksema et al., 2008), and is associated with slower treatment response and poorer rates of recovery when using antidepressant medication and cognitive therapy (Ciesla and Roberts, 2002; Jones et al., 2008). Moreover, because rumination is shown to exacerbate negative affect, impair problem-solving, reduce motivation, and block individuals from connecting with both direct positive experience and evidence disconfirmatory of negative beliefs (Nolen-Hoeksema et al., 2008; Watkins, 2008), tackling rumination is likely to enhance the treatment benefits of cognitive-behavioural approaches. Further, as a transdiagnostic process also contributing to anxiety disorders (Watkins, 2008), targeting rumination may improve treatment for depression with co-morbid anxiety. As a consequence, directly tackling rumination has been recommended to improve interventions for depression (e.g. Topper et al., 2010; Drost et al., 2014; Grierson et al., 2016; Spinhoven et al., 2018). Rumination-focused CBT (RFCBT) was therefore developed as a modification of CBT to explicitly target depressive rumination (Watkins, 2016) and features two key novel adaptations of standard CBT: (1) based on a theoretical conceptualization of rumination-as-a-mental-habit (Watkins and Nolen-Hoeksema, 2014), it uses functional...
analysis to change rumination by identifying its triggers and practicing alternative behaviours to these cues; (2) based on experimental research indicating that the consequences of repetitive thought depend on the information processing style adopted (Watkins et al., 2008), it trains patients to shift into a more adaptive style of processing (Watkins, 2008). It differs from standard CBT by not involving direct thought challenging and by focusing on shifting the process of thinking rather than the content. RFCBT has been shown to improve outcomes in treatment-resistant residual depression (Watkins et al., 2011). Although the reduction in depressive symptoms in that study reported for RFCBT was better than the reduction reported in a RCT of standard CBT for residual depression (Paykel et al., 1999), to date, no RCT has directly compared RFCBT v. standard CBT, nor directly investigated RFCBT for patients with a current major depressive episode. This study therefore reports the first RCT directly comparing RFCBT v. CBT for major depression. A group format for delivering therapy was chosen to improve cost-effectiveness and vicarious learning, and to reduce experiences of loneliness and shame, through sharing and normalization within the group. Even though a group format may reduce experiences of loneliness and shame, through sharing and normalization within the group. Evidence suggests that group therapy has equivalent outcomes compared with individual therapy (Burlingame et al., 2016). The aim of this study was to test the hypothesis that group RFCBT would be superior to group CBT in reducing symptoms of depression post-treatment, when added to standard medical management.

**Method**

The study was approved by the National Committee on Health Research Ethics in Denmark (case no. H-1-2013-049) and the trial was registered at ClinicalTrials.gov (registration no. NCT02278224) on 28 October 2014. The study protocol was published in Trials on 17 August 2015 (Hvenegaard et al., 2015).

**Design**

The study was conducted as a two-arm, assessor-blinded, randomized superiority trial. Participants were randomly allocated in a 1:1 ratio to groups of seven to nine participants providing CBT plus medical management or RFCBT plus medical management. Medical management was defined as clinical management and treatment by a trained and experienced psychiatrist at the outpatient service, including the potential prescription of antidepressant medication. Randomization was performed by an external statistical agency (Statcon, DK) according to an independent pre-study off-site computer-generated schedule with randomly ordered permutable blocks sized 6–10. A researcher (MH) masked and kept blind to treatment allocation assessed all participants at baseline ($T_0$) and 12 weeks later after completing treatment ($T_1$) with all primary and secondary measures and at the 6 months post-treatment follow-up ($T_2$) with the primary measure only. After completing each follow-up $T_2$ assessment or following the point in time in which the $T_2$ assessment was scheduled for those who did not attend, the assessor completed a forced guess of treatment allocation for each participant, and the accuracy of the guesses was at chance level (48.9%), consistent with blindness.

**Participants**

Recruitment occurred from December 2013 to July 2015 from a public health system outpatient clinic north of Copenhagen, Denmark, which treats 200–250 patients with a diagnosis of major depression per year. The clinic is a secondary mental health care facility and offers treatment for patients referred from primary care with affective disorders, post-traumatic stress disorder and personality disorders, including specialized treatment for difficult-to-treat depression. Most patients with depression in the outpatient clinic had received treatment with antidepressant medication and/or psychotherapy in primary care prior to the referral. Consecutive referrals to the outpatient service were approached, and those patients who met inclusion criteria and gave written informed consent to participate were randomly allocated to group RFCBT or to group CBT. When baseline assessment was completed, the off-site randomization administrator informed the relevant therapist to contact the patient and initiate the allocated intervention.

Inclusion criteria were: aged between 18 and 65 years, meeting Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (American Psychiatric Association, 1994) criteria for a current episode of unipolar major depression in a structured M.I.N.I. 5.0 interview (Sheehan and Lecrubier, 1998) and with a score of $\geq 13$ on the 17-item Hamilton Rating Scale for Depression (HRSD; Hamilton, 1960). Exclusion criteria were: a history of bipolar disorder, psychosis, current (past 6 months) drug or alcohol abuse or dependence, a primary diagnosis of any anxiety disorder, anorexia, or bulimia, all determined by the M.I.N.I. 5.0 interview, imminent and substantial suicide risk as assessed by an experienced psychiatrist or clinical psychologist, and concurrent psychotherapy at point of entry to the study. There were no exclusion criteria with respect to co-morbid anxiety disorders or the use of antidepressants.

**Outcome measures**

The primary outcome was severity of depressive symptoms measured with the 17-item interviewer-rated HRSD at post-treatment ($T_1$). All other measures were secondary outcomes and included change between $T_0$ and $T_1$ in self-reported rumination, worry, anxiety and severity of depressive symptoms. Self-report measures of behavioural activation, well-being, a neuropsychological test of task switching and a computer-based test of visual emotional attention bias were also included but will not be reported in this paper. Suicidal behaviour/ideation was monitored during the trial in accordance with the guidelines from the Danish Health Authorities.

**Primary outcome measure**

**Hamilton Rating Scale for Depression**

The HRSD (Hamilton, 1960) is a standardized clinical interview developed to assess severity of depression that includes scoring the test persons answers as well as direct observation of the test person. Higher scores suggest higher levels of symptoms of depression (range 0–52). A Danish version of the 17-item HRSD interview guide was used (Bech and Larsen, 2012). Masked ratings of randomly selected recorded interviews (18%) indicated moderate to strong inter-rater reliability between the interviewer and the masked rater, all $\kappa$ coefficients $>0.76$. The HRSD was conducted as a face-to-face structured interview at $T_0$ and $T_1$. The HRSD at $T_2$ was conducted as a mixture of face-to-face interviews (43%) and telephone interviews (57%); telephone interviews were used for convenience to increase patient retention.
Secondary outcomes measures

Ruminative Response Scale of the Response Style Questionnaire
The Ruminative Response Scale (RRS) (Nolen-Hoeksema and Morrow, 1991) consists of 22 items that assess ruminative responses to sad and depressed mood. Participants rate the frequency that they use unhelpful ruminative strategies, and higher scores suggest higher levels of rumination (range 22–88).

Generalized Anxiety Disorder 7-item Scale
The Generalized Anxiety Disorder 7-item Scale (GAD-7) (Spitzer et al., 2006) consists of seven items that assess the severity of generalized anxiety. Participants rate the frequency that they experience symptoms of anxiety, and higher scores suggest higher frequency of symptoms (range 0–21).

Penn State Worry Questionnaire
The Penn State Worry Questionnaire (PSWQ) (Meyer et al., 1990) consists of 16 items that assess the general disposition to worry. Participants rate statements about worry on a scale of 1 (‘not at all typical of me’) to 5 (‘very typical of me’). Higher scores suggest higher level of worry (range 16–80).

Hamilton Self-report Questionnaire
The Hamilton Self-report Questionnaire (HAM-D6) (Beck, 1975) consists of six items that assess the severity of symptoms of depression. Participants rate intensity of symptoms, and higher scores suggest higher levels of symptoms of depression (range 0–22).

Interventions

RFCBT is a principle-driven manualized CBT treatment for depression, adopting a behavioural activation perspective (Martell et al., 2001), in which rumination is conceived as a learnt habitual behaviour developed through negative reinforcement (Watkins and Nolen-Hoeksema, 2014). Based on this conceptualization, rather than challenging individual negative thoughts, RFCBT uses functional analysis to change rumination by helping patients to learn to identify antecedent cues and triggers to rumination, control exposure to these cues and repeatedly practice alternative behaviours to these cues. Further, based on experimental research indicating that the consequences of repetitive thought depend on information processing style (Watkins et al., 2008), it trains patients to shift into a more adaptive style of processing. Alternative responses include activity scheduling, imagery, recreating experiences of being absorbed (‘flow’) or of increased compassion to self or others, and/or shifting into a more concrete and specific thinking style (Watkins, 2008). A group version consisting of a one-to-one individual preparatory session of 1 h and 11 group sessions of 3 h with two breaks, scheduled weekly, was developed in collaboration with Edward Watkins (EW) – the original developer of RFCBT (Møller et al., 2017). Trial recruitment, data collection and analysis of data were conducted in Copenhagen independently of EW.

CBT was based on Beck’s CBT manual for depression (Beck, 2011) adapted to a group format, which was the routine treatment already being used in the outpatient service. It consisted of a one-to-one individual preparatory session of 1 h followed by 11 group sessions of 3 h with two breaks, scheduled weekly. Both treatment manuals are described in online Supplement 1.

The therapists in both treatment conditions were employees in the psychiatric clinic in which the patients were recruited. Therapists were not chosen or allocated on the basis of therapeutic allegiance or experience: the therapists in the CBT arm were already delivering CBT groups for depression in the clinic; the RFCBT therapists were chosen on the basis of their availability for training and to deliver new treatment groups. All therapists had prior CBT training and had completed at least one year or more of formal education in CBT. The therapists in both treatment conditions had equivalent levels of training and experience as CBT therapists (9 years on average), and received equivalent levels of video supervision during the trial (1 h a month). In addition, the therapists conducting RFCBT received a 3-day training workshop on RFCBT conducted by the developer of the therapy (EW). Prior to the trial, a pilot group in both conditions was conducted with video supervision provided.

All therapy sessions in the trial were videotaped. For both treatment conditions, a random sample of 16 (18%) videotapes, stratified by therapy group and therapy session, were rated for therapist’s competence and adherence to treatment manual by four independent raters. For each treatment, based on the detailed and structured therapy manual (Watkins, 2016; Møller et al., 2017), there was a checklist of the required and prohibited therapy components. To assess adherence to treatment manuals, the raters used each checklist to record the presence or absence of these key therapy components in the rated sessions for each treatment. For both treatment conditions, no prohibited components were reported and the presence of required therapy key components was high (CBT 98%; RFCBT 98%).

Therapists’ competence was rated using the 11-item Cognitive Therapy Rating Scale (CTRS) for the CBT condition (Young and Beck, 1980). For the RFCBT condition, an adapted version of the CTRS was used. The first six items reflecting general skills common to both therapies (e.g. agenda setting, asking for feedback, therapist empathy, interpersonal effectiveness, collaboration and efficient use of time) were the same. To capture the novel components of RFCBT, other item scales were adapted as required to reflect specific RFCBT competence, e.g. item 8 ‘Focusing on key cognitions or behaviours’ was changed to ‘Focusing on key cognitions or behaviours relevant to functional analysis’, item 9 ‘Strategy for Change’ was adapted to ‘Focus on changing thinking style’ and item 10 ‘Application of Cognitive-Behavioural Techniques’ was adapted to ‘Application of RFCBT techniques’. A total score of 40 or greater on the CTRS represents the standard threshold of acceptable competence in CBT delivery (Dobson et al., 1985). CTRS scores for all the rated sessions for both CBT [M = 43.6, standard deviation (s.d.) = 2.1] and RFCBT [M = 46.3, s.d. = 2.2] were 40 or above for all raters, evidencing good quality of treatments delivered by the CBT and RFCBT therapists. The inter-rater reliabilities in both conditions were moderate-to-good (RFCBT: \( \kappa = 0.65 \); CBT: \( \kappa = 0.66 \)).

Statistical analysis

The primary and secondary outcomes were analysed using a multilevel regression model with treatment condition (RFCBT v. CBT) as main effect, therapy group as random intercept, baseline \( (T_0) \) scores as covariate and \( T_1 \) scores as the dependent variable. The analysis was performed according to the intention-to-treat principle (ITT, i.e. all participants according to randomization), with multiple imputations of missing data. For post-treatment, 12.2% of HRSD scores were missing. Multiple
imputations conducted with MICE package in R-studio (van Buuren and Groothuis-Oudshoorn, 2011) were used to account for missing data for all primary and secondary outcomes. No difference was found on HRSD baseline scores for participants with missing HRSD T1 scores (M = 20.1, s.d. = 6.8) and complete cases (M = 19.9, s.d. = 4.9). See online Supplement 2 for a full description of the missing data and the multiple imputations method.

We calculated the sample size required based on the relative changes in HRSD scores pre- to post-treatment for RFCBCT (Watkins et al., 2011) and CBT (Paykel et al., 1999) for patients with residual depression in prior RCTS. Assuming similar mean changes in HRSD scores from pre- to post-intervention as found by Watkins et al. (2011) for RFCBCT (M = 7.8) and by Paykel et al. (1999) for CBT (M = 3.5) and a conservative estimate of pooled s.d. for change in HRSD of 6.0 (when s.d. = 3.6 for change in HRSD in RFCBCT from Watkins et al., 2011), we estimated a between-treatment effect size of Cohen’s d = 0.7. To detect a difference in effect size of 0.7 between RFCBCT and CBT at a two-tailed significance level of 5%, each treatment arm requires 44 patients to obtain 90% statistical power. Assuming a lost to follow-up rate of 20%, we would recruit 55 patients into each treatment arm. With an average size of the therapy group of m = 8 in both treatment arms and an intraclass correlation of about ρ = 0.05, a design effect of 1 + (m – 1)ρ = 1.35 followed, so that we planned to recruit eight groups in each treatment arm (128 patients in total). Initial sample size (N = 112) was adjusted upwards based on recommendations to control for design effects in group studies – this occurred after recruitment commenced, but before it completed, and was published in the study protocol (Hvenegaard et al., 2015). The analysis plan was decided prior to the data collection and was described in the published study protocol (Hvenegaard et al., 2015).

Results

Patient flow

A total of 140 patients from a public Danish psychiatric outpatient service were screened and 131 patients who agreed to participate and met the inclusion criteria were randomized to either group RFCBCT (n = 66) or group CBT (n = 65). Figure 1 shows the participant flow from screening to follow-up. The main reasons for potentially eligible individuals not participating were that they declined to participate (6.4%) or they did not meet study criteria (3.2%). Main reasons for not meeting the inclusion criteria were: not meeting criteria for an episode of major depression, not meeting criteria (3.2%). Main reasons for not meeting the inclusion criteria were: not meeting criteria for an episode of major depression, or meeting criteria for bipolar depression.

All participants across both treatment conditions were offered clinical management and treatment with antidepressant medication by a trained and experienced psychiatrist at the outpatient service. The number of participants receiving antidepressant medication did not differ between CBT and RFCBCT [59 of 65 (91%) v. 60 of 66 (91%); $\chi^2 = 0.001; p = 0.978$]. See online Table S3 in Supplement 3 for full details on number of participants receiving antidepressant medication, types of antidepressant medications, dosage of antidepressant medications and for statistics showing no significant differences between the uses of medications in the two treatment conditions. All participants were offered at least consultation by a psychiatrist in the outpatient clinic on their use of medication during the treatment.

Participants’ verbal reports of side effects of the medication and non-compliance with the medical treatment were reported in the participants’ medical files. The number of participants reporting no side effects of medications (CBT: n = 44, 67.7% v. RFCBCT: n = 50, 75.8%; $\chi^2 = 1.051, p = 0.305$) and the number of participants reporting non-compliance with medical treatment (CBT: n = 4, 6.1% v. RFCBCT: n = 2, 3.0%; $\chi^2 = 0.731, p = 0.39$) did not differ between the two treatment conditions. See online Table S4 in Supplement 4 for full details on side effects of medical treatment. The number of consultations with a psychiatrist during the trial did not differ between CBT and RFCBCT (M = 1.1, s.d. = 1.4 v. M = 1.1, s.d. = 1.6; t = −0.174, p = 0.862).

One participant was hospitalized for prevention of suicide during the trial. To assess deterioration, we calculated a Reliable Change index (RC; Jacobson and Truax, 1991) for the HRSD of 6.5 points. The RC was calculated using the α coefficient (α = 0.789) from a meta-analysis on the reliability of the HRSD scale (Trajković et al., 2011) and by dividing the HRSD change score with the standard error of difference. No participant showed deterioration exceeding the RC and only two participants (one in CBT; one in RFCBCT conditions) reported a deterioration of more than 3 points on the HRSD.

For both conditions, overall treatment compliance was good: there was no difference in the number of group sessions attended between CBT and RFCBCT (M = 8.3, s.d. = 3.2 v. M = 8.8, s.d. = 2.8; t = −1.2, p = 0.226), nor in the number of participants who dropped out of treatment [11 of 65 (17%) v. 9 of 66 (14%); $\chi^2 = 0.273; p = 0.601$]. A total of 114 (87%) completed the post-treatment assessment (T2). Despite repeated attempts to contact all participants, only half of the participants could be contacted and then participated in the T2 follow-up assessment 6 months post-treatment (70, 53%), reducing our statistical power for T2 analyses. The last patient was randomized on 26 May 2015. Follow-up data were obtained between 4 March 2014 and 15 January 2016. No harms or side effects of psychological interventions, or adverse events were reported during the trial.

Participant characteristics

Table 1 shows participant characteristics of the ITT sample for both the RFCBCT and CBT groups. Twenty-six per cent had chronic depression lasting 2 years or more, 57% had recurrent depression with a history of two or more depressive episodes, and 65% had a comorbid anxiety disorder.†

Primary outcome

As shown in Table 2, as hypothesized, group RFCBCT patients reported a significantly greater reduction in depressive symptoms at post-treatment (T1) than group CBT patients, after adjusting for difference in baseline HRSD scores (M ΔHRSD = 2.8; 95% CI 0.0–5.6, p = 0.049). A complete case analysis (n = 114; 87% of sample) found similar results: RFCBCT resulted in significantly lower-between-treatments HRSD scores at T1 than CBT (M ΔHRSD = 2.7; t = 2.26, 95% CI 0.3–5.1, p = 0.026).

Secondary outcomes

In both treatments, the levels of self-reported depression, rumination, worry and anxiety were reduced, but no statistical difference

†The notes appear after the main text.
was found between RFCBT and CBT for any of these variables at post-treatment (T1), although we note varying levels of missing data on the questionnaires. Missing secondary outcomes included: RRS (41, 31%), PSWQ (42, 32%), HAM-D6 (41, 31%). In a complete case analysis (n = 87; 66% of sample), RFCBT reduced symptoms of anxiety significantly more than CBT (M ΔGAD-7 = 2.4, 95% CI 0.4–4.4). Complete case analyses on other secondary outcomes were not significant. Change scores from baseline to post-treatment for both primary and secondary outcomes are shown in Table 2. No significant between-treatment difference in average depressive symptoms (i.e. average HDRS at T2) was found between RFCBT (M = 9.7, s.d. = 7.5) and CBT (M = 8.7, s.d. = 6.8) in the ITT sample at the 6 months follow-up (M ΔHRSD = −1.1, 95% CI −4.1 to 1.9, p = 0.56, E.S. = 0.15).

Discussion

The primary aim of this study was to compare the efficacy of group RFCBT with the efficacy of group CBT for treating major depression.
Table 1. Demographic and psychiatric characteristics of group rumination-focused cognitive-behavioural therapy (RFCBT) and group cognitive-behavioural therapy (CBT)

|                      | CBT (n = 65) | RFCBT (n = 66) |
|----------------------|-------------|--------------|
| Female, n (%)        | 53 (82)     | 47 (71)      |
| Age (years), mean (s.d.) | 39.4 (12.5)  | 39.8 (13.7)  |
| Marital status, n (%)* |            |              |
| Single               | 31 (50)     | 28 (42)      |
| Married or cohabiting| 31 (50)     | 37 (58)      |
| Levels of education, n (%)* |       |
| No educational qualifications | 0 (0)     | 1 (2)        |
| Some educational qualifications | 13 (20)    | 17 (26)      |
| High school/vocational education | 35 (55)   | 36 (55)      |
| University degree/professional qualifications | 16 (25)  | 11 (17)      |
| Job status, n (%)* |            |              |
| Student              | 9 (15)      | 9 (14)       |
| Unemployed           | 9 (15)      | 11 (17)      |
| Full-time work       | 10 (17)     | 10 (15)      |
| Part-time work       | 5 (8)       | 4 (6)        |
| Sickness allowance   | 21 (36)     | 26 (40)      |
| Other                | 5 (8)       | 5 (8)        |
| Psychiatric characteristics |          |              |
| Baseline depression HRSD, mean (s.d.) | 20.1 (4.4) | 19.8 (5.8) |
| Length of current depression (months), mean (s.d.) | 16.6 (18.1) | 15.2 (22.6) |
| Recurrent depression (two or more episodes), n (%) | 33 (51) | 42 (64) |
| Receiving antidepressant medication, n (%) | 59 (91) | 60 (91) |
| Comorbid generalised anxiety disorder (M.I.N.I.), n (%) | 43 (66) | 42 (64) |
| Baseline GAD-7 scores, mean (s.d.) | 11.0 (4.8) | 11.5 (4.5) |
| Baseline rumination scores, mean (s.d.) | 58.2 (9.2) | 57.8 (11.6) |

HRSD, Hamilton Rating Scale for Depression (17-item version); GAD-7, Generalized Anxiety Disorders scale; RRS, Ruminative Response Scale (22 items).

*Five patients did not give information on marital status, two did not give information on level of education and seven did not give information on job status. Percentages calculated for valid cases.

Treatment effects on depressive symptoms

Consistent with our primary hypothesis, participants in the group RFCBT treatment improved significantly more than those in the group CBT treatment in reducing symptoms of depression at the end of treatment (after 12 weeks). This finding is consistent with the positive results of RFCBT already found for residual depression (Watkins et al., 2011; Teismann et al., 2014) and for adolescents at risk for depressive relapse because of a prior history of depression (Jacobs et al., 2016). Furthermore, the within-group effect of group CBT in this study was similar to that found in other trials (Oei and Dingle, 2008). Because it is difficult to find benefits of an intervention compared with another effective intervention, these findings are encouraging. In the absence of a
definitive RCT of RFCBT v. CBT with a larger sample and a longer follow-up with less missing data, we tentatively suggest that these modifications made to CBT for RFCBT may engender better treatment outcomes.

The data available for T2 also indicate that initial treatment effects are stable over the 6 months follow-up. However, the difference in depressive symptoms at 6 months follow-up (T2) numerically disappeared. However, a large proportion (47%) of patients were lost to follow-up at T2 and the most parsimonious explanation is that the study was underpowered at follow-up (T2) to detect a difference on HRSD between the conditions, even if there was a genuine difference in the effect of the treatments. Because of the high attrition at T2, these secondary analyses need to be treated with caution. Alternatively, it may be that both CBT and RFCBT are similarly effective treatments for depression in the long run, but that the benefits of RFCBT manifest earlier. We are unable to discriminate between these different interpretations in the current study.

Mechanisms of the treatment effect

Surprisingly, group RFCBT did not reduce self-reported rumin-ation significantly more than group CBT. In both conditions, the level of rumination was significantly lower at post-treatment compared with baseline. We note several possible accounts for this observation. First, because of missing data on this secondary measure and follow-up attrition (only 66% completion), the study was underpowered to detect a genuine difference in rumination, unless there was a large effect size between RFCBT and CBT. As such, we need to be cautious about making any strong inter-pretation of these findings. Second, it may be that group CBT is also effective at reducing rumination, perhaps because challenging negative thoughts, increased problem solving and activity sched-uling all act to break the vicious circle of rumination, as suggested in a recent meta-analysis (Spinnewyn et al., 2018), although this meta-analysis also found that treatments targeting rumination tended to produce stronger reductions in rumination.

The lack of a differential effect of the treatments on rumin-ation raises the possibility that shifting rumination was not the active mechanism underpinning the effect of RFCBT. RFCBT differs from standard CBT in a number of ways. Elements unique to RFCBT include engendering the ability to recognize pathological rumination and coaching an ability to adopt more functional styles of processing as an alternative through practise in experien-tial/imagery exercises, such as concreteness training, absorption training and self-compassion training. Any or none of these elements might be responsible for the apparent differential efficacy between treatments. It has been posited that a behavioural activation approach may be simpler and more straightforward for peo-ple with depression, with one study finding that behavioural activation outperformed CBT for patients with more severe levels of depression (Dimidjian et al., 2006), but others finding no dif-fERENCE (Richards et al., 2016). The emphasis on habit change in RFCBT may provide a simple and convincing rationale for patients, and may encourage repeated practice of new strategies in daily life engendering more robust change. Because the trial was designed to test the effects of the complete intervention packages, we cannot determine which of the treatment compo-nents within RFCBT are responsible for the observed differential treatment effect. The current RCT was designed to mitigate threats to internal validity when evaluating RFCBT relative to CBT and was successful in this intention. However, it was not designed to investigate construct validity (i.e. to determine what aspect of RFCBT contributes to treatment outcome). Nonetheless, the relative outperformance of RFCBT to CBT post-treatment raises the possibility that some elements found in RFCBT but not in CBT may underpin either improved treatment outcomes or faster recovery. Rigorous trial designs that can decompose the active ingredients of treatment (e.g. dismantling studies or factorial designs) are needed to resolve the question of which elements actively underpin outcome.

It is hypothesized that patients with depression would benefit more from RFCBT than classical CBT when they have severe, chronic and treatment-resistant depression, because rumination is found to exacerbate and prolong depression and interfere with treatment, or when they have co-morbid anxiety disorders, because rumination is identified as a transdiagnostic mechanism. However, these hypotheses were not formally tested in this trial.

Limitations of the study

This study has several limitations. First, the principal limitation is the missing data on secondary outcomes and the high follow-up attrition rate at 6 months, which limit conclusions for these out-comes. Ideally, more participants would have been retained at 6-month follow-up and follow-up would have continued for at least 2 years post-treatment to examine rates of relapse and recur-rence longer term. Resource constraints meant that this was not feasible. Nonetheless, the trial was well-powered to answer the primary aim and there was little missing data on the primary out-come. Second, because we did not evaluate non-specific therapy factors such as patient expectations, therapy allegiance and treat-ment credibility, we cannot rule out the possibility that differences in non-specific factors may account for the observed difference in treatment outcomes. Third, there was no active monitoring of changes in antidepressant medication over the course of the trial making it impossible to assess the impact of any such changes to the primary and secondary outcomes. However, no difference was found in the use of antidepressant medication throughout the trial between the treatment conditions as assessed from medical records (see online Supplement 3). Fourth, the lack of consecutive repeated HRSD assessments at post-treatment limits our ability to assess the proportion of participants who achieved remission (lasting >3 weeks). Fifth, no systematic assess-ment of potential harm effects of psychotherapy was conducted, as now recommended (e.g. Schniebel et al., 2017). Sixth, registra-tion of the trial happened almost 1 year after the trial commenced and sample size was increased during recruitment into the trial to include the recommended design effect to account for the reduced variability of participants treated in the same therapy group (Roberts and Roberts, 2005), although this amendment was included in the published trial protocol. Seventh, we were unable to examine to what extent participants may have received CBT or not (CBT-naïve) prior to the trial, as no record of prior psycho-therapy before entering the secondary outpatient service was rou-tinely collected. However, it is unlikely that participants received CBT prior to the referral, as CBT is not routine treatment in Danish primary care services and patients are typically referred to the secondary service in order to receive CBT for depression: the secondary out-patient service is the principal route to access CBT for depression in the Danish healthcare system.

In conclusion, this study is the first RCT to conduct a head-to-head comparison of group RFCBT and group CBT for patients with major depression. The finding that a novel adaptation of
traditional CBT (Rumination-focused CBT) performs significantly better in reducing observer-rated depressive symptomatology at 12 weeks than an established empirically validated intervention (CBT) in a reasonably well-powered study is noteworthy, as it is rare for new treatments to outperform current treatments. As a minimum, these results suggest the potential benefits of rumination-focused CBT as an alternative to standard CBT for depression in this population. Nonetheless, as a single study, we need to be cautious about this finding and there is a need for larger, multicentre RCTs to replicate these findings in other settings and to examine cost-effectiveness in a definitive Phase III trial.

Note

A between-treatment sensitivity analysis including only the first 112 randomised participants (i.e. the original sample size) did not differ from the primary analysis (M ΔHRSD = 2.8; p = 0.023, 95% CI 0.4–5.2).

Supplementary material.

The supplementary material for this article can be found at https://doi.org/10.1017/S0033291718003835.

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Conflict of interest.

None.

References

American Psychiatric Association (1994) Diagnostic and Statistical Manual of Mental Disorders, 4th Edn. (DSM-IV). Washington, DC: APA.

Bech P (1975) Quantitative rating of depressive states. Acta Psychiatrica Scandinavica 51, 161–170.

Bech P and Larsen ER (2012) Interview-guide til Hamiltons Depressionsskala. Available at https://wwwpsykiatrienrm.dk/siteassets/forsking/afdeling-q---forsking/interview_guide_hamid17_rev_pb_21_12_12_1358151150.pdf.

Beck JS (2011) Cognitive Behavior Therapy: Basics and Beyond. New York, NY: Guilford press.

Burlingame GM, Gleave R, Erekson D, Nelson PL, Olsen J, Thayer S and Beecher M (2016) Differential effectiveness of group, individual, and conjoint treatments: an archival analysis of OQ-45 change trajectories. Psychotherapy Research 26, 556–572.

Ciesla JA and Roberts JE (2002) Self-directed thought and response to treatment for depression: a preliminary investigation. Journal of Cognitive Psychotherapy 16, 435–453.

Cuipiers P, Karyotaki E, Weitz E, Andersson G, Hollon SD and van Straten A (2014) The effects of psychotherapies for major depression in adults on remission, recovery and improvement: a meta-analysis. Journal of Affective Disorders 159, 118–126.

Cuipiers P, Cristea IA, Karyotaki E, Reijnders M and Huibers MJH (2016) How effective are cognitive behavior therapies for major depression and anxiety disorders? A meta-analytic update of the evidence. World Psychiatry 15, 245–258.

DeRubeis RJ, Hollon SD, Amsterdam JD, Shelton RC, Young PR, Salomon RM, O’Reardon JP, Lovett ML, Gladis MMBL, Gallop R, Brown LL and Gallop R (2005) Cognitive therapy vs medications in the treatment of moderate to severe depression. Archives of General Psychiatry 62, 409–416.

Dimidjian S, Hollon SD, Dobson KS, Schmaling KB, Kohlenberg RJ, Addis ME, Gallop R, McGlinchey JB, Markley DK, Gollan JK, Atkins DC, Dunner DL, Dobson KS, Shaw BF and Vallis TM (2006) Reliability of a measure of the quality of cognitive therapy. British Journal of Clinical Psychology 44, 295–300.

Dobson KS, Shaw BF and Vallis TM (1985) Reliability of a measure of the quality of cognitive therapy. British Journal of Clinical Psychology 24, 295–300.

Drost J, Van der Does W, van Hemert AM, Penninx BW and Spinholven P (2014) Repetitive negative thinking as a transdiagnostic factor in depression and anxiety: a conceptual replication. Behaviour Research and Therapy 63, 177–183.

Grierson AB, Hickie JB, Naismith SL and Scott J (2016) The role of rumination in illness trajectories in youth: linking trans-diagnostic processes with clinical staging models. Psychological Medicine 46, 2467–2484.

Hamilton M (1960) A rating scale for depression. Journal of Neurology, Neurosurgery & Psychiatry 23, 56–62.

Hvenegaard M, Watkins ER, Poulsen S, Rosenberg NK, Gondan M, Grafton B, Austin SF, Howard H and Moeller SB (2015) Rumination-focused cognitive behaviour therapy vs. cognitive behaviour therapy for depression: study protocol for a randomised controlled superiority trial. Trials 16, 344.

Jacobs RH, Watkins ER, Peters AT, Feldhaus CG, Barba A, Carbray J and Langenecker SA (2015) Targeting ruminative thinking in adolescents at risk for depressive relapse: rumination-focused cognitive behaviour therapy in a pilot randomized controlled trial with resting state fMRI. PLoS ONE 11, e0163952.

Jacobson NS and Tuax P (1991) Clinical significance: a statistical approach to defining meaningful change in psychotherapy research. Journal of Consulting and Clinical Psychology 59, 12–19.

Jones NP, Siegle GJ and Thase ME (2008) Effects of rumination and initial severity on remission to cognitive therapy for depression. Cognitive Therapy and Research 32, 591–604.

Martell CR, Addis ME and Jacobson NS (2001) Depression in Context: Strategies for Guided Action. New York: W. W. Norton & Company, Inc.

Meyer TJ, Miller ML, Metzger RL and Borkovec TD (1990) Development and validation of the Penn State Worry Questionnaire. Behaviour Research and Therapy 28, 487–495.

Moller SB, Hvenegaard M and Kistrup M (2017) Ruminationsfokusert Kognitiv Adførdetrap for Depression – Manual Til Gruppeterapi. København: Hans Reitzels Forlag.

Nolen-Hoeksema S (2000) The role of rumination in depressive disorders and mixed anxiety/depressive symptoms. Journal of Abnormal Psychology 109, 504–511.

Nolen-Hoeksema S and Morrow J (1991) A prospective study of depression and posttraumatic stress symptoms after a natural disaster: the 1989 Loma Prieta Earthquake. Journal of Personality and Social Psychology 61, 115–121.

Nolen-Hoeksema S, Wisco BE and Lyubomirsky S (2008) Rethinking Rumination. Perspectives on Psychological Science 3, 400–424.

Oei TPS and Dingle G (2008) The effectiveness of group cognitive behaviour therapy for unipolar depressive disorders. Journal of Affective Disorders 107, 5–21.

Paykel ES, Scott J, Teasdale JD, Johnson AL, Garland A, Hons BA, Moore R, Jenaway A, Cornwall PL, Hayhurst H, Abbott R and Pope M (1999) Prevention of relapse in residual depression by cognitive therapy. Archives of General Psychiatry 56, 829–835.

Richards DA, Ekers D, McMillan D, Taylor RS, Byford S, Warren FC, Barrett B, Farrant PA, Glibbode S, Kuyken W, O’Mahan H, Watkins ER, Wright KA, Hollon SD, Reed N, Rhodes S, Fletcher E and Finning K (2016) Cost and Outcome of Behavioural Activation versus Cognitive Behavioural Therapy for Depression (COBRA): a randomised, controlled, non-inferiority trial. The Lancet 388, 871–880.

Roberts C and Roberts SA (2005) Design and analysis of clinical trials with clustering effects due to treatment. Clinical Trials 2, 152–162.

Schneiberg R, Willbartz G, Scholz C, Becker M, Brakemeier EL, Bschor T, Zobel I and Schnoll D (2017) Adverse events of group psychotherapy in the in-patient setting – results of a naturalistic trial. Acta Psychiatrica Scandinavica 136, 247–258.
Sheehan D and Lecrubier Y (1998) The Mini International Neuropsychiatric Interview (MINI): the development and validation of a structured diagnostic psychiatric interview. *Journal of Clinical Psychiatry* 59, 22.

Spinhoven P, Klein N, Kennis M, Cramer AO, Siegle G, Cuijpers P, Ormel J, Hollon S and Bockting CL (2018) The effects of cognitive-behavior therapy for depression on repetitive negative thinking: a meta-analysis. *Behaviour Research and Therapy* 106, 71–85.

Spitzer RL, Kroenke K, Williams JW and Löwe B (2006) A brief measure for assessing generalized anxiety disorder. *Archives of Internal Medicine* 166, 1092–1097.

Teismann T, Von Brachel R, Hanning S, Grillenberger M, Hebermehl L, Hornstein I and Willutzki U (2014) A randomized controlled trial on the effectiveness of a rumination-focused group treatment for residual depression. *Psychotherapy Research* 24, 80–90.

Topper M, Emmelkamp PM and Ehring T (2010) Improving prevention of depression and anxiety disorders: repetitive negative thinking as a promising target. *Applied and Preventive Psychology* 14, 57–71.

Trajković G, Starčević V, Latas M, Leštarević M, Ille T, Bukumirić Z and Marinčović J (2011) Reliability of the Hamilton Rating Scale for Depression: a meta-analysis over a period of 49 years. *Psychiatry Research* 189, 1–9.

van Buuren S and Groothuis-Oudshoorn K (2011) Mice: multivariate imputation by chained equations in R. *Journal of Statistical Software* 45, 1–67.

Watkins ER (2008) Constructive and unconstructive repetitive thought. *Psychological Bulletin* 134, 163–206.

Watkins ER (2015) Psychological treatment of depressive rumination. *Current Opinion in Psychology* 4, 32–36.

Watkins ER (2016) *Rumination-Focused Cognitive-Behavioral Therapy for Depression*. New York, NY: Guildford Press.

Watkins ER and Nolen-Hoeksema S (2014) A habit-goal framework of depressive rumination. *Journal of Abnormal Psychology* 123, 24–34.

Watkins ER, Moberly NJ and Moulds ML (2008) Processing mode causally influences emotional reactivity: distinct effects of abstract versus concrete construal on emotional response. *Emotion* 8, 364–378.

Watkins ER, Mullan E, Wingrove J, Rimes K, Steiner H, Bathurst N, Eastman R and Scott J (2011) Rumination-focused cognitive-behavioural therapy for residual depression: phase II randomised controlled trial. *The British Journal of Psychiatry* 199, 317–322.

Young J and Beck AT (1980) *Cognitive Therapy Scale: Rating Manual. Unpublished manuscript*, Philadelphia: University of Pennsylvania.