Enduring effects of a 5-week behavioral activation program for subthreshold depression among late adolescents: an exploratory randomized controlled trial

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Background: No significant effect of psychological treatment has been reported from meta-analysis of subthreshold depression patients and control subjects at 1-year follow-up. However, behavioral activation is a simpler and more cost-effective treatment than cognitive behavioral therapy. The primary purpose of this study was to assess by comparison to an assessment-only control group whether the effects of behavioral activation program for depressive symptoms can persist up to 1-year follow-up without the use of antidepressants or other psychotherapy.

Patients and methods: Late adolescent students were the population targeted in this study. Participants were allocated randomly to an intervention group (n=62) or a control group (n=56). Treatment consisted of five-weekly 60-minute sessions. Participants underwent a structured interview and completed self-report scales at 1 year post-assessment.

Results: Late adolescent students receiving treatment had significantly lower mean Beck Depression Inventory, second edition scores at 1-year follow-up than control group students. The effect size (Hedges’ g) for between-group differences at 1-year follow-up was –0.41.

Conclusion: Our behavioral activation program is simple and short. Nevertheless, the results obtained at 1-year follow-up of the control group and late adolescent students receiving treatment indicated a significant difference in their Beck Depression Inventory, second edition scores. Our 5-week behavioral activation program based on behavioral characteristics for subthreshold depression might be promising for subthreshold depression. The sample examined for this study imposed some study limitations.

Keywords: behavioral activation, first-year university students, randomized controlled trial, subthreshold depression

Introduction
Subthreshold depression is defined as the exhibition of clinically significant depressive symptoms that do not meet the diagnostic criteria for major depressive disorder (MDD).1 Subthreshold depression is a major risk factor for subsequent development of more severe depressive symptoms.2–5 The probability of MDD is known to increase steadily during adolescence, that is, 12–20 years.6 Although not meeting the full criteria, subthreshold depression has been associated with severe problems such as negative impact on academic performance and social activities,7 which might be as severe as those associated with full depression.8
An earlier report described that the use of antidepressant medication by adolescent patients is associated with a modestly increased risk of suicidality. It is, therefore, especially important to assess the effectiveness of psychological treatment programs only, without antidepressants, for late adolescents with subthreshold depression. Several earlier reports of studies with longer follow-up (1 year) have described that cognitive behavioral therapy (CBT) can reduce the risk of MDD. Nevertheless, a meta-analysis of psychological treatments for subthreshold depression found no significant effect of psychotherapy for depressive symptoms at 1-year follow-up and found no significant effect of CBT on the incidence of MDD at follow-up.

For adolescent and late adolescent patients with depression, earlier reports have described behavioral activation as an effective treatment. For university students with moderately depressive symptoms, behavioral activation was found to have significantly greater reduction in depressive symptoms from pretreatment to posttreatment. Although the meta analysis of behavioral activation for children and adolescents reported that behavioral activation was effective from pretreatment to posttreatment, the long-term effect has not yet been elucidated for children and adolescent patients. An earlier systematic review of behavioral activation programs for depression revealed an effect size of −0.35 (Hedges’ g) in comparison with control conditions at 6–9 months of follow-up. The effectiveness of behavioral activation in follow-up might not have a sufficient effect. No report of the relevant literature describes a study that examined the long-term efficacy of brief behavioral activation for subthreshold depression. Recently, we conducted a randomized controlled trial (RCT) to examine the efficacy of a brief five-session behavioral activation program for late adolescent patients with subthreshold depression. Results demonstrated that late adolescent students receiving this treatment exhibited significantly greater improvement in self-reported depressive symptoms than the assessment-only control group. The effect size was Hedges’ g of −0.90. It has not been reported whether the effects of this behavioral activation program can be maintained over the long term.

Therefore, the primary purpose of this study was to assess whether a behavioral activation program without antidepressants for depressive symptom can be maintained for depressive symptoms up to 1 year and be compared with the assessment-only control group. The secondary aim was to explore whether the behavioral activation program can decrease the risk of developing MDD or not.

**Patients and methods**

**Research design**

Details of the study design of this RCT were presented in an earlier report (UMIN000011967). The 1-year follow-up study was already planned in the original RCT. Furthermore, the same student sample as that examined in an earlier study was followed up 1 year later in the present study.

**Procedures**

Details of procedures of this RCT were presented in an earlier report. Participants were recruited from Hiroshima University over a 2-year period during 2013–2014 (Figure 1). The Beck Depression Inventory, second edition (BDI-II) is administered annually to all first-year students of the university as a part of their health examination, which is conducted at the Health Service Center of Hiroshima University. We randomly selected the first-year students with a BDI-II score ≥10 using the Microsoft Excel randomization function. Recruitment took place through email at a public information sharing center. Then, we held a meeting to explain the study contents. After all the participants had signed a written informed consent form, participants received the Composite International Diagnostic Interview (CIDI) and completed self-report scales via the Internet. An expert of the Department of Clinical Research, who was independent of the research team that conducted this study, developed a sequential assignment list using computer-generated random numbers to allocate the participants to a treatment or a control group randomly at a 1:1 ratio. The random sequence was stratified by sex and depression severity during screening (BDI-II score ≤13, BDI-II score ≥14). The group allocation was masked in the entry and in the CIDI assessment. A psychiatrist interviewed the participants and assessed the risk of a future acute suicide attempt if a participant had a history of attempted suicide based on CIDI criteria. Furthermore, participants who met the criteria for a possible acute suicide attempt received counseling from a psychiatrist. Participants joined the RCT. We contacted the participants by email and telephone 1 year later. Participants received telephone interviews by CIDI and completed self-report scales via the Internet 1 year after the assessment by blind testers who did not know the allocation. In CIDI assessment of 1-year follow-up, allocation to the treatment group or control group was masked.

**Participants**

We recruited late adolescent students with subthreshold depression during a 2-year period of 2013–2014 at Hiroshima
University in Japan. For this study, the inclusion criteria were the following: 18–19-year-old first-year university student at Hiroshima University, BDI-II score ≥ 10 according to earlier studies, participants who did not have a major depressive episode during the previous year as determined by the structured clinical interview with the CIDI, and not undergoing psychopharmacological or psychological treatment. The exclusion criteria were the following: a lifetime history of

**Figure 1 Study flow chart.**

**Abbreviations:** BDI-II, Beck Depression Inventory, second edition; CIDI, Composite International Diagnostic Interview; ITT, intention-to-treat.
bipolar disorder according to earlier studies, possibility of an acute suicidal attempt, difficulty understanding the study purpose, or having a severe physical disease and “difficulty in completing self-report scales”.

Intervention
Our behavioral activation program consisted of five-weekly 60-minute sessions. The program emphasized increased access to positive reinforcement activities to improve the rate of response-contingent positive reinforcement. Our behavioral activation program was modeled after the behavioral activation component of a CBT program developed based on earlier studies of adult depression in the USA and which has been demonstrated in an RCT to be effective for subthreshold depression in the workplace in Japan. Details of intervention of this RCT can be found in an earlier report.

Control group
The control group participants were examined for outcome measures. They received baseline, posttreatment, and follow-up assessments at approximately the same time as the intervention group did. The control group participants in this study received no medical treatment, usual care, or any educational information.

Measures
Primary outcome
The Japanese version of the BDI-II BDI-II consists of 21 self-report items scored on a four-point scale. It is used to measure depressive symptoms. Good reliability and validity have been reported for the Japanese version of BDI-II. BDI-II was administered at baseline, at the end of treatment, and at 1-year follow-up. The primary outcome of this report is BDI-II scores at 1-year follow-up. The internal consistencies of the BDI-II in pretreatment, posttreatment, and 1-year follow-up were assessed using Cronbach’s . The internal consistencies for all time BDI-II were also acceptable: pretreatment = 0.79, posttreatment = 0.74, and 1-year follow-up = 0.87.

Secondary outcome
Composite International Diagnostic Interview
The CIDI is a structured interview for assessing mental disorders. Good reliability and validity have been reported for the Japanese ver 3.0 of the CIDI. We used the Japanese 3.0 version of the CIDI in screening and 1-year follow-up. We assessed for 12-month history of MDD, generalized anxiety disorder, panic disorder, social phobia, and post-traumatic stress disorder. In addition, each participant was assessed for the lifetime history of bipolar disorder. Because earlier reports have described the comorbidity rate for depression and anxiety as high, we included generalized anxiety disorder, panic disorder, social phobia, and post-traumatic stress disorder in screening for this study.

We administered CIDI to the participants before baseline and at 1-year follow-up. At the follow-up assessment, we limited the time frame to the previous year to ascertain the new incidence of MDD. The secondary outcome of this report is the incidence of MDD during 1 year of follow-up.

Sample size calculation
Sample size calculations were conducted at the study design stage.

An earlier report described a mean effect size of in behavioral activation group vs a control group at posttest. The mean effect size found for psychotherapies for subthreshold depression was . Because effect sizes suggest that indicates a medium effect, we set the effect size in our study as a medium effect ( ). We estimated a minimum of 64 participants in each group at the study design stage (power=0.80 and α=0.05, d=0.50 for an unpaired t-test). Therefore, the power might be insufficient to detect differences in incidence of MDD that occurs over 1 year.

Statistical analysis
All statistical analyses were conducted using software (SPSS Statistics 22; IBM Corp.).

Primary outcome analysis
The primary purpose of this study was exploratory examination of whether behavioral activation program effects persisted at 1-year follow-up. The primary point was estimation of the intervention effect on depressive symptoms between the original assigned groups at 1-year follow-up. Analysis of variance with repeated measures is limited to cases with complete data or with imputation of missing data. Some statistical approaches are useful for dealing with missing data that do not necessarily require formal imputation methods. For example, the linear mixed model for repeated measure approach was used for analyzing outcomes measured repeatedly over time. To examine the effects of treatment on primary outcomes at 1-year follow-up, the primary analysis was conducted using a linear mixed model for repeated measures. The random effects of this model were intercepts.
Therefore, because individual differences are explained as a variable effect, the mode of analysis of this research is a model showing change from the individual baseline. The model predictors were categorical time (pretreatment, post-treatment, 1-year follow-up), treatment condition, and time x treatment interaction as fixed effects. We applied intention-to-treat (ITT) analysis between the original assigned groups. We conducted post hoc analysis if an interaction was found. Effect sizes were calculated and reported as Hedges’ g37 for between-group differences in 1-year follow-up, using the observed standard error (SE) at follow-up.

Some participants might not complete the full protocol of a longitudinal study. Sensitivity analyses can be conducted using different approaches to ascertain how robust the primary analysis is for the chosen method for handling missing data.34 For sensitivity analysis, we conducted a complete case analysis to assess the discrepancy between ITT and complete case analysis.

Secondary exploratory outcome analysis
Next, we applied Cox proportional hazard analysis for exploratory assessment of the differences in time for the development of MDD between the intervention group and the control group to adjust for lifetime diagnosis of MDD. In CIDI, participants who met the criteria for MDD reported enduring effects of behavioral activation during the follow-up period. Therefore, missing data of the intervention group were those of ten participants from posttreatment to 1-year follow-up. Missing data of the control group were those of six participants from posttreatment to 1-year follow-up. No significant difference was found in the rates of missing data ($\chi^2(1)=0.73$, $P=0.43$).

Depressive symptoms (primary outcome)
Results obtained from a linear mixed model
The primary point was estimation of the intervention effect on depressive symptoms between the original assigned groups at 1-year follow-up. To assess the effects of treatment on primary outcomes at 1-year follow-up, primary analysis was conducted using a linear mixed model for a repeated measures approach in the original assigned groups (62 participants in the intervention group and 56 participants in the control group). Significant interaction was found between groups and times ($F$ value=9.01, degrees of freedom=216.45, $P<0.01$; Figure 2). Significant difference was found between the mean estimates for the intervention group and the control group at 1-year follow-up (intervention group, mean estimate=11.00, SE=0.89, 95% CI: 9.24 to 12.76; control group, mean estimate=13.83, SE=0.91, 95% CI: 12.00 to 15.61; estimated mean difference=2.83, 95% CI: 0.28 to 5.33, $P=0.03$), as shown in Table 2. The effect size (Hedges’ g) for between-group differences at 1-year follow-up was $-0.41$ (95% CI: $-0.77$ to $-0.04$). Additionally, a significant difference was found in the mean estimates between the intervention group and control group at posttreatment (intervention group, mean estimate=7.03, SE=0.84, 95% CI: 5.38 to 8.68; control group,

**Table 1**
Descriptive data of groups

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>Intervention group</th>
<th>Control group</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>n=62</td>
<td>24/38</td>
<td>21/35</td>
<td>1.00</td>
</tr>
<tr>
<td>Mean age (years)</td>
<td>18.23 (0.42)</td>
<td>18.20 (0.40)</td>
<td>0.70</td>
</tr>
<tr>
<td>Live alone/live with family</td>
<td>48/14</td>
<td>43/13</td>
<td>1.00</td>
</tr>
<tr>
<td>Enter the university of first choice</td>
<td>33/29</td>
<td>33/23</td>
<td>0.66</td>
</tr>
<tr>
<td>Lifetime diagnosis of major depressive disorder (yes/no)</td>
<td>5/57</td>
<td>8/48</td>
<td>0.43</td>
</tr>
<tr>
<td>Comorbidity</td>
<td>Diagnosis of any anxiety disorder in the previous year (yes/no)</td>
<td>7/55</td>
<td>14/42</td>
</tr>
<tr>
<td>BDI-II (pretreatment)</td>
<td>12.76 (6.66)</td>
<td>13.30 (5.95)</td>
<td>0.64</td>
</tr>
</tbody>
</table>

*Note:* Values in parentheses represent SD.

*Abbreviation:* BDI-II, Beck Depression inventory, second edition.

**Ethics approval**
Approval for the study was obtained from the ethics committee of Hiroshima University. All participants signed a written informed consent form.

**Results**
**Participant enrollment**
Figure 1 presents the study flow. Of the 208 students who gave their informed consent to participate, 205 (98.6%) completed screening. The mean score of the sample consisting of 118 students was 13.02. In CIDI, participants who met the criteria for MDD reported $0.77$ to $-0.04$.
mean estimate = 12.77, SE = 0.88, 95% CI: 11.03 to 14.51; estimated mean difference = 5.74, 95% CI: 3.34 to 8.14, P < 0.01. The effect size (Hedges’ g) for between-group differences in 1-year follow-up was −0.87 (95% CI: −1.24 to −0.49).

Next, because this study had missing data, we conducted complete case analysis (intervention group, n = 51; control group, n = 49) and then conducted sensitivity analysis and examined the discrepancy between ITT and complete case analysis. Significant interaction was found between groups and times (F value = 9.42, degrees of freedom = 196, P < 0.01; Figure 2). Significant difference was found between the mean estimates for the intervention group and the control group at 1-year follow-up (intervention group, mean estimate = 10.77, SE = 0.93, 95% CI: 8.91 to 12.60; control group, mean estimate = 13.83, SE = 0.96, 95% CI: 11.90 to 15.65; estimated mean difference = 3.01, 95% CI: 0.39 to 5.63, P = 0.03). The effect size (Hedges’ g) for between-group differences at 1-year follow-up was −0.45 (95% CI: −0.85 to −0.05). Additionally, a significant difference was found in estimated means between the intervention group and the control group at posttreatment (intervention group, mean estimate = 6.45, SE = 0.93, 95% CI: 4.62 to 8.29; control group, mean estimate = 12.77, SE = 0.96, 95% CI: 10.95 to 14.69; estimated mean difference = 6.37, 95% CI: 3.75 to 8.99, P < 0.01). The effect size (Hedges’ g) for between-group differences at 1-year follow-up was −0.90 (95% CI: −1.33 to −0.51). No discrepancy was found between the results of ITT and complete case analysis.

**Prophylactic effects for onset of MDD**

We assessed the significance of differences between the intervention group (n = 51) and the control group (n = 49) in developing MDD during the 1-year follow-up. In CIDI, participants who met the criteria for MDD responded that they had developed depressive symptoms. In addition, participants reported the time as 1) within 1 month, 2) between 2 and 6 months, and 3) between 7 and 12 months after baseline assessment.

Two participants in the intervention group developed MDD during 1-year follow-up (one participant within the previous 1 month, one participant in the previous 2–6 months: 2/51, 3.92%). Four control group participants developed MDD (one participant 2–6 months before, three participants 7–12 months before: 4/49, 8.16%). Three participants of the intervention group and eight participants in the control group had an MDD history longer than 1 year before the screening of this study. We applied Cox proportional hazard analysis to assess differences in survival probabilities between the intervention group and the control group. We used the MDD

**Table 2** Results of linear mixed model

<table>
<thead>
<tr>
<th>Score</th>
<th>Time</th>
<th>Intervention group (n = 62)</th>
<th>Control group (n = 56)</th>
<th>95% CI</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Estimate (SE)</td>
<td>Estimate (SE)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BDI-II</td>
<td>Pretreatment</td>
<td>12.76 (0.83)</td>
<td>13.30 (0.88)</td>
<td>−2.93 to 1.89</td>
<td>0.65</td>
</tr>
<tr>
<td></td>
<td>Posttreatment</td>
<td>7.03 (0.84)</td>
<td>12.77 (0.88)</td>
<td>3.34 to 8.14</td>
<td>0.00</td>
</tr>
<tr>
<td></td>
<td>1-year follow-up</td>
<td>11.00 (0.89)</td>
<td>13.83 (0.91)</td>
<td>0.28 to 5.33</td>
<td>0.03</td>
</tr>
</tbody>
</table>

**Complete case analysis**

<table>
<thead>
<tr>
<th>Score</th>
<th>Time</th>
<th>Intervention group (n = 51)</th>
<th>Control group (n = 49)</th>
<th>95% CI</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Estimate (SE)</td>
<td>Estimate (SE)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>BDI-II</td>
<td>Pretreatment</td>
<td>12.63 (0.93)</td>
<td>13.18 (0.96)</td>
<td>−2.06 to 3.18</td>
<td>0.65</td>
</tr>
<tr>
<td></td>
<td>Posttreatment</td>
<td>6.45 (0.93)</td>
<td>12.77 (0.96)</td>
<td>3.75 to 8.99</td>
<td>0.00</td>
</tr>
<tr>
<td></td>
<td>1-year follow-up</td>
<td>10.77 (0.93)</td>
<td>13.83 (0.96)</td>
<td>0.39 to 5.63</td>
<td>0.03</td>
</tr>
</tbody>
</table>

*Note:* Values in parentheses are SEs.

*Abbreviations:* BDI-II, Beck Depression Inventory, second edition; SE, standard error.
Results of our earlier report indicated that late adolescents with subthreshold depression had a lower frequency of environmental rewards than healthy late adolescents.\(^\text{39}\) Our behavioral activation program emphasized increased access to positive reinforcement activities to improve the rate of response-contingent positive reinforcement. These results demonstrate that late adolescent students receiving treatment had significantly different BDI-II scores from those of the assessment-only control group at 1-year follow-up.

However, the effect size for between-group differences at 1-year follow-up was \(-0.41\). The effect size of intervention can be regarded as small. Our earlier study revealed the effect size of behavioral activation program as \(-0.90\) for between-group differences in acute treatment.\(^\text{19}\) Therefore, the behavioral activation program effect decreased during 1 year. Moreover, we attempted to ascertain whether a 5-week short-term behavioral activation program can decrease the risk of developing MDD compared to a control group or not. Results of Cox proportional hazard analysis revealed no significant difference between the intervention group and the control group in the incidence of MDD. Although a significant difference was found in the depressive symptoms at 1-year follow-up between the intervention group and the control group, one must interpret the results carefully. In terms of the considerable reduction in effects between posttreatment and follow-up, although some earlier studies conducted booster sessions or continuation sessions to maintain the effects of intervention, we conducted no such booster sessions. The knowledge that continued support would be provided via booster sessions might have reinforced participants’ confidence in managing depression or anxiety.\(^\text{42}\) An earlier report described from a meta-analysis that CBT intervention with a booster session is more effective in terms of sustainability than CBT interventions without booster sessions.\(^\text{42}\) To maintain behavioral activation effects, it is necessary to review activities in booster sessions. Moreover, it is very difficult to obtain large effects if baseline depressive symptoms are as low as they were in this study. Furthermore, the behavioral activation intervention was very brief. It can be regarded as a low-intensity intervention. Therefore, long-term effects of such treatment might be less likely to be apparent.

Some study limitations are related to the sample used for this study. The sample size calculations were conducted at the study design stage. However, this sample size was based on the study design of an earlier study.\(^\text{19}\) An earlier report described that 200 participants per condition must be tested for a difference in the incidence rate.\(^\text{40}\) Although we assessed differences in the incidence of MDD between the...
intervention group and the control group in a preliminary study, the sample size of this study was small. A future study using a large sample size must be undertaken to examine the incidence of MDD. Secondly, in this study, we confirmed whether the participants were undergoing psychopharmacological or psychological treatment in CIDI or not. Participants who were not undergoing psychopharmacological or psychological treatment were included in this study. We confirmed the current treatment in this study. Therefore, future studies must be conducted to target people who have never received psychopharmacological or psychological treatment. Despite the limitations described above, the results of this study show a significant difference in BDI-II scores between the intervention group and the control group at 1-year follow-up.

Conclusion
Although our behavioral activation program is simple and brief, the results obtained at 1-year follow-up indicate a significant difference between the BDI-II scores of the late adolescent control group students and the students receiving treatment. Our 5-week behavioral activation program based on behavioral characteristics for subthreshold depression appears to be promising for treating subthreshold depression.

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Author contributions
All authors contributed to the design, acquisition of data, interpretation of data drafting, and critical revision of the paper, and agree to be accountable for all aspects of the work. All authors have approved the final article.

Disclosure
The authors have no conflict of interest, financial or otherwise, related to this study.

References


