

A comparison of two online cognitive-behavioural interventions for symptoms of depression in a student population: the role of therapist responsiveness

Submission date 20/08/2009	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 09/11/2011	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 10/11/2011	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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2

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

OTD

Study information

Scientific Title

A randomised parallel group controlled trial of online delivered cognitive behavioural therapy treatment with an adult student population

Acronym

OTD

Study objectives

Online delivered cognitive behavioural therapy (CBT) treatments for depression are effective and the presence of a responsive therapeutic relationship to treatment adds value to the treatment delivered and consequently has the potential to influence a successful outcome.

Please note, as of 10/11/2011 the public title for this trial has been updated. The previous title was as follows:

Online Treatments for Depression: a randomised controlled trial on an adult student population

Ethics approval required

Old ethics approval format

Ethics approval(s)

University of Dublin, Ireland, 27 November 2007

Study design

Randomised parallel group controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Mild to moderate depression

Interventions

The trial had two active treatment conditions, the first being cCBT (Computerised Cognitive Behaviour Therapy), self-administered by the user with no support. The second was eCBT (Email Cognitive Behaviour Therapy), delivered by a clinician. Both used the same CBT protocol. In addition the latter had the added element of the therapists free text writing as a response to the clients free text alongside the specific CBT aspects of treatment. No other condition was present such as waiting list. The duration of the treatment was 8 sessions over 8 weeks. Baseline measures were administered for screening purposes, they were the Beck Depression Inventory (BDI-II), Brief Symptom Inventory (BSI), Clinical Outcomes in Routine Evaluation - Outcome Measure - 10 items (CORE-OM-10), history collected data on previous experience of counselling and medication for depression, whether they had a previous diagnosis of a mental health disorder, alcohol and other drug use. Thereafter the principal measures were administered at Week 8, end of treatment, again at week 16 Follow-up and at week 32 Follow-up.

The online interventions included two CBT-based online treatments for depression. One consisted of therapist-led online counselling, the second was a self-administered software program called Beating the Blues, with minimal therapist contact. Beating the Blues is a cCBT (computerised cognitive behavioural therapy) program utilised to treat depression and anxiety. Each intervention includes 8 sessions of online treatment comprising cognitive-behavioural interventions, including psycho-education, written assignments, self monitoring, cognitive-restructuring, behavioural change, and relapse prevention strategies. In addition to the specific CBT aspects of the treatment, the online counselling condition included counsellors, through text, cultivating non-specific treatment factors that develop the therapeutic relationship including such generic skills as conveying of empathy, normalising, responding to the client's emotions, and validating successes. Treatment was delivered by trained and experienced professional therapist with additional training in online counselling.

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Beck Depression Inventory (BDI)

Measured at baseline, week 8 (end of treatment), week 16 follow-up and week 32 follow-up

Secondary outcome measures

1. Clinical Outcomes in Routine Evaluation - Outcome Measure (CORE-OM-10) measured at baseline, week 8 (end of treatment), week 16 follow-up and week 32 follow-up
2. Working Alliance Inventory (WAI)-was administered at weeks 2, 4 and 6
3. Helpful Aspects of Therapy (HAT)-was administered from session 2 to session 8
4. Satisfaction with treatment measure was administered at week 8, end of treatment

Overall study start date

01/08/2008

Completion date

01/06/2010

Eligibility

Key inclusion criteria

Participant scores between 14 and 29 on the Beck Depression Inventory (BDI)

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

120

Key exclusion criteria

1. Below 14 or above 29 on BDI
2. Suicidal ideation and/or plans
3. Diagnosed organic mental health disorder
4. Recent medical illness diagnosis
5. Drug or alcohol dependence
6. On medication for depression less than 6 months

Date of first enrolment

01/08/2008

Date of final enrolment

01/06/2010

Locations**Countries of recruitment**

Ireland

Study participating centre

199-200 Pearse Street

Dublin

Ireland

2

Sponsor information**Organisation**

University of Dublin (Ireland)

Sponsor details

c/o Dr. Derek Richards
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Trinity College Dublin
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2

Sponsor type

University/education

Website

<http://www.tcd.ie>

ROR

<https://ror.org/05m7pjf47>

Funder(s)**Funder type**

Research organisation

Funder Name

ESB Electric Aid (Ireland)

Funder Name

HSE Innovation Fund (Ireland)

Funder Name

National Office for Suicide Prevention (Ireland)

Results and Publications**Publication and dissemination plan**

Not provided at time of registration

Intention to publish date**Individual participant data (IPD) sharing plan**

IPD sharing plan summary

Not provided at time of registration