Effectiveness of a computerised cognitive behavioural therapy programme (Overcoming Depression) for patients suffering from depression as compared to a waiting list control group.

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	☐ Results
Last Edited	Condition category	☐ Individual participant data
11/04/2014	Mental and Behavioural Disorders	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Robert Dudley

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0573142164

Study information

Scientific Title

Study objectives

Will people receiving Computerised Cognitive Behavioural Therapy (CCBT) show reductions in depressive and anxious symptoms in comparison to those receiving no intervention?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Mental and Behavioural Disorders: Depression

Interventions

Computerised Cognitive Behavioural Therapy (CCBT) vs no intervention

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Beck Depression Inventory (BDI); Beck Anxiety Inventory; Clinical Outcomes in Routine Evaluation: Hospital Anxiety and Depression Scale.

The BDI is the most frequently used measure for considering the effect size. Drawing on similar previous trials we could expect a reduction in BDI scores of approximately 10 points.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2003

Completion date

30/06/2004

Eligibility

Key inclusion criteria

Male and female adult out patients who have been referred for specialist cognitive behavioural therapy for treatment of depression.

Participants will have:

- 1. Primary diagnosis of depression disorder or anxiety.
- 2. No change in medication for the previous 12 weeks.
- 3. Willingness to try a trial of CCBT.
- 4. No evidence of severe depression or suicidal intent.
- 5. No evidence of current substance abuse.
- 6. No evidence of organic impairment.
- 7. Agreement of RMO for participation.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

40

Key exclusion criteria

- 1. Patients with organic disorders
- 2. Patients with personality disorder
- 3. Patients who can not comprehend English

Date of first enrolment

01/06/2003

Date of final enrolment

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Newcastle Cognitive & Behavioural Therapies Centre
Newcastle upon Tyne
United Kingdom
NE1 6UR

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type

Government

Funder Name

Newcastle, North Tyneside and Northumberland Mental Health NHS Trust (UK), NHS R&D Support Funding

Results and Publications

Publication and dissemination planNot provided at time of registration

Intention to publish date

Individual participant data (IPD) sharing plan

IPD sharing plan summaryNot provided at time of registration