

A randomised controlled trial of Internet based cognitive behavioural psychotherapy for depression

Submission date 13/08/2004	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 08/11/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 14/09/2009	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Study information

Scientific Title

Study objectives

To evaluate the clinical and cost effectiveness of Internet based cognitive behavioural therapy (webCBT) for depression compared to a waiting list control in primary care. A qualitative study will be done as part of the trial to assess the acceptability of the intervention to patients and health care providers.

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

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Depression

Interventions

Intervention: Up to eight one hour sessions of cognitive behavioural therapy (CBT) delivered online by a qualified psychologist.

Control: The control group will be on a waiting list for CBT of 8 months and will receive 'usual care' from their GP. This may include pharmacological treatment but will not include psychotherapy.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

The Beck depression inventory score at 2 months adjusted for baseline.

Key secondary outcome(s)

EQ5D and SF-12 at 8 months

Completion date

30/09/2008

Eligibility**Key inclusion criteria**

Patients with GP diagnosed depression in primary care. Diagnosis must be confirmed by standardised psychological measure.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/10/2005

Date of final enrolment

30/09/2008

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre**Division of Primary Care**

Bristol

United Kingdom

BS6 6JL

Sponsor information**Organisation**

University of Bristol (UK)

ROR

<https://ror.org/0524sp257>

Funder(s)

Funder type

Charity

Funder Name

BUPA Foundation (UK) (ref: 683/G14)

Alternative Name(s)

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	22/08/2009		Yes	No