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Attentional style in bipolar disorder

Submission date	Recruitment status	[ ] Prospectively registered
12/05/2010	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
12/05/2010	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
11/08/2014	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
5864

Study information

Scientific Title

Study objectives
1. Do patients with bipolar disorder display an attentional bias towards threat?
2. Can this bias be modified using a simple computer-based intervention?
3. Does modifying the bias lead to changes in the emotional response to stressful situations?

Ethics approval required
Old ethics approval format

Ethics approval(s)
MREC approved, ref: 08/H0603/36

Study design
Single-centre interventional treatment trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Topic: Mental Health Research Network; Subtopic: Bipolar affective disorder; Disease: Bipolar affective disorder

Interventions
Intervention - positive attention training: this involves completing a computer-based task on the day of testing. The task requires that particiapnts pay less attention to negative information. Participants complete the task only during the testing session. The task takes approximately 30 minutes to complete.
Control - neutral attention training: this is identical to the intervention task except that the participants are not required to direct their attention away from negative information.

Study entry: registration and one or more randomisations.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure

Attentional bias as measured using a computerised task (the dot-probe task) on the day of testing

Secondary outcome measures
1. Symptoms of anxiety, measured on a 10-point scale and collected via SMS messaging three times a day for the week following testing
2. Symptoms of low mood (measured using the Positive Affect Negative Affect Schedule [PANAS] scale) and anxiety (measured using the State-Trait Anxiety Inventory [STAI] scale) immediately following testing

Overall study start date
13/03/2009

Completion date
01/10/2011

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Planned sample size: 80; UK sample size: 80

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
13/03/2009

Date of final enrolment
01/10/2011

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University Department of Psychiatry
Oxford
United Kingdom
OX3 7JX

Sponsor information

Organisation
University of Oxford (UK)

Sponsor details
Research Services
Clinical Trials and Research Governance
Headley Way
Headington
Oxford
England
United Kingdom
OX3 9DU

Sponsor type
University/education

Website
http://www.ox.ac.uk/

ROR
https://ror.org/052gg0110

Funder(s)

Funder type
Charity

Funder Name
The Wellcome Trust (UK)

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