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A comparison of two psychological treatments for severe eating disorders

Submission date	Recruitment status	[ ] Prospectively registered
22/07/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
22/07/2005	Completed	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
12/06/2015	Mental and Behavioural Disorders	

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
046386

Study information

Scientific Title
A comparison of two psychological treatments for severe eating disorders

Study objectives
1. To test theories regarding what processes cause clinical eating disorders to persist.
2. To evaluate the relative effectiveness of two different psychological treatments for severe eating disorders.

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)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Eating disorders

Interventions
Randomised controlled trial with embedded psychological tests.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
1. Standardised measures of eating disorder features
2. General psychiatric symptoms
3. Psychosocial impairment

Secondary outcome measures
No secondary outcome measures

Overall study start date
01/10/2001

Completion date
30/09/2007

Eligibility

Key inclusion criteria
1. Patients with any type of severe eating disorder
2. 18 to 60 years
3. Living in central Oxfordshire or Leicestershire
4. Available for treatment

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Both

Target number of participants
Not provided at time of registration

Key exclusion criteria
1. Patients who have been previously exposed to either of the two treatments
2. Pregnancy

Date of first enrolment
01/10/2001

Date of final enrolment
30/09/2005

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Oxford
Oxford
United Kingdom
OX33 1JT

Sponsor information

Organisation
University of Oxford (UK)

Sponsor details
University Offices
Wellington Square
Oxford
England
United Kingdom
OX1 2JD
+44 (0)1865 270143
research.services@admin.ox.ac.uk

Sponsor type
University/education

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

ROR
https://ror.org/052gg0110

Funder(s)

Funder type
Charity

Funder Name
Wellcome Trust (UK) (grant ref: 046386)

Alternative Name(s)

Funding Body Type
Private sector organisation

Funding Body Subtype
International organizations

Location
United Kingdom

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/03/2009		Yes	No
Results article	results	01/07/2015		Yes	No