

Controlled evaluation of Cognitive Behaviour Therapy (CBT) for Obsessive Compulsive Disorder (OCD) in children and adolescents: comparison of standard, therapist intensive therapy versus brief therapy

Submission date 11/03/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 11/05/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 04/07/2011	Condition category Mental and Behavioural Disorders	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

MREC 0312045

Study information

Scientific Title

Study objectives

The aims of the study are:

1. To determine the effectiveness of Cognitive Behaviour Therapy (CBT) for children and adolescents with Obsessive Compulsive Disorder (OCD) and the extent to which this can be sustained by a treatment with relatively low intensity of therapist contact
2. To produce a treatment practice manual suitable for dissemination
3. To identify patient characteristics that may be related to differential treatment response
4. To determine whether treatment improves quality of life
5. To investigate service use associated with the two treatments

Ethics approval required

Old ethics approval format

Ethics approval(s)

No ethics information 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

Obsessive Compulsive Disorder

Interventions

Participants will be randomly assigned to one of three conditions:

1. Standard CBT: comprising 12 sessions therapist contact and handouts explaining the features

of OCD and its treatment. Both this and the brief CBT will be delivered over a three month period.

2. Brief CBT: comprising five sessions therapist contact and use of a detailed multi-section workbook intended to supplement the therapy sessions.

3. Waiting-list/delayed treatment, for three months. At the end of this period if the child still has diagnosable OCD they will be offered treatment in either of the first two conditions on a randomized basis. This group will act as the control group for the study.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Child Yale-Brown Obsessive Compulsive Scale (CY-BOCS) (Scahill et al. 1997).

Secondary outcome measures

1. Anxiety Disorders Interview Schedule for Children and Parents (ADIS-C/P, Silverman & Nelles, 1988)
2. Obsessive Compulsive Inventory (OCI) adapted for use with children (Foa et al. 1997)
3. Responsibility Attitude Scale (Salkovskis et al. 2000)
4. Child OCD Impact Scale (COIS) child and parent version (Piacentini et al. 2001)
5. Manchester Short Assessment of Quality of Life (MANSA) (Priebe et al. 1999)

Overall study start date

01/10/2003

Completion date

31/03/2007

Eligibility

Key inclusion criteria

1. Aged 10 to 18
2. Diagnosis of OCD according to Diagnostic and Statistical Manual of Mental Disorders, Fourth edition (DSM-IV) criteria
3. If on medication, on a stable dosage for at least six weeks

Participant type(s)

Patient

Age group

Child

Lower age limit

10 Years

Upper age limit

18 Years

Sex

Both

Target number of participants

110

Key exclusion criteria

1. Immediate suicide risk
2. Intelligence Quotient (IQ) less than 70
3. Comorbid diagnosis of autism or marked autistic spectrum features, or psychosis, which commonly raise problems about differential diagnosis
4. Comorbid condition which has greater treatment priority (such as some cases of depression, or school-refusal)

Date of first enrolment

01/10/2003

Date of final enrolment

31/03/2007

Locations**Countries of recruitment**

England

United Kingdom

Study participating centre

Department of Psychology

London

United Kingdom

SE5 8AF

Sponsor information**Organisation**

King's College London (UK)

Sponsor details

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Sponsor type
University/education

ROR
<https://ror.org/0220mzb33>

Funder(s)

Funder type
Charity

Funder Name
The Health Foundation (UK) (reference Number 1206/1782)

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	01/12/2011		Yes	No