

Quetiapine augmentation of a serotonin reuptake inhibitor in treatment resistant obsessive-compulsive disorder: a multi-site, placebo-controlled study

Submission date 13/01/2005	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 13/01/2005	Overall study status Completed	☐ Protocol
Last Edited 11/08/2008	Condition category Mental and Behavioural Disorders	☐ Statistical analysis plan
		☒ Results
		☐ 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

Protocol serial number

5077IL/9009

Study information

Scientific Title

Study objectives

Although serotonin reuptake inhibitors are effective in the treatment of obsessive-compulsive disorder (OCD), many patients fail to respond to these agents. Growing evidence from open-label and placebo-controlled trials suggests a role for augmentation of serotonin reuptake inhibitors (SRIs) with atypical antipsychotics in OCD. Quetiapine is generally well tolerated and previous open-label data has produced mixed results in OCD, and additional controlled data is needed.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Obsessive-compulsive disorder

Interventions

Placebo-controlled, double-blind, flexible-dose augmentation with quetiapine of a serotonin reuptake inhibitor maintained at the stable maximum tolerated dose.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration.

Key secondary outcome(s)

Not provided at time of registration.

Completion date

01/11/2003

Eligibility

Key inclusion criteria

Subjects with obsessive-compulsive disorder who failed at least one trial (12 weeks duration, of which six weeks at maximum tolerated dose) of a serotonin reuptake inhibitor.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/05/2002

Date of final enrolment

01/11/2003

Locations**Countries of recruitment**

Canada

South Africa

Study participating centre

MRC Research Unit on Anxiety Disorders

Cape Town

South Africa

7505

Sponsor information**Organisation**

AstraZeneca (South Africa)

ROR

<https://ror.org/04r9x1a08>

Funder(s)

Funder type

Industry

Funder Name

AstraZeneca Pharmaceuticals (South Africa)

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	24/01/2005		Yes	No