Promoting personal recovery from psychosis: A randomised controlled trial in first episode schizophrenia

| Submission date | Recruitment status No longer recruiting | Prospectively registered | | |
|----------------------------------|--|--|--|--|
| 23/01/2004 | | ☐ Protocol | | |
| Registration date 23/01/2004 | Overall study status Completed | Statistical analysis plan | | |
| | | [X] Results | | |
| Last Edited 28/01/2010 | Condition category Mental and Behavioural Disorders | Individual participant data | | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number RGC00447

Study information

Scientific Title

Study objectives

The objective of the study is to evaluate whether a form of cognitive therapy (recovery intervention) is an effective intervention to promote personal adjustment to psychosis and reduce depression, trauma and other characteristic negative consequences of psychosis.

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

Schizophrenia and other psychoses

Interventions

- 1. The intervention group will receive cognitive therapy over a 6 month period
- 2. The control group will receive treatment as usual

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

- 1. Post-psychotic depression, suicidal ideation (Calgary scale)
- 2. Symptoms of Post Traumatic Stress Disorder (Impact of Events scale)
- 3. Adaptation to psychosis
- 4. Particularly perceived control over psychosis

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/05/2003

Eligibility

Key inclusion criteria

Patients conforming to International Statistical Classification of Diseases and Related Health Problems, Tenth revision (ICD-10) schizophrenia or related disorder

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/05/2001

Date of final enrolment

01/05/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre North Birmingham Mental Health NHS Trust

Birmingham United Kingdom B6 5UG

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Executive West Midlands (UK)

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 | 01/06/2009 | | Yes | No |