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Cognitive Behavioural Case Management (CBCM) in First-Episode Schizophrenia and Related Psychotic Disorders - a pilot study

Submission date	Recruitment status	[ ] Prospectively registered
29/09/2006	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
29/09/2006	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
12/08/2016	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
N0038161724

Study information

Scientific Title
Cognitive Behavioural Case Management (CBCM) in First-Episode Schizophrenia and Related Psychotic Disorders - a pilot study

Study objectives
Will a group of participants with first episode psychosis (FEP) receiving CBCM report lower levels of psychopathology, better psychosocial functioning and quality of life, greater service satisfaction and be more engaged with treatment than a group of participants with FEP receiving treatment-as-usual at the end of a 12 month intervention and at an 18 month follow-up?

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Pilot 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
Mental and Behavioural Disorders: Schizophrenia

Interventions
[A] Cognitive Behavioural Case Management
[B] Treatment-as-usual

Intervention Type
Other

Phase

Not Applicable

Primary outcome measure
Quality of Life Survey

Secondary outcome measures
1. Brief psychiatric rating scale (expanded version)
2. Montgomery-Asberg Depression Rating Scale
3. WHOQOL-BREF
4. Social and Occupational Functioning Assessment Scale
5. Addition Severity Scale
6. Illness Duration Interview
7. Pathways to care item (from CORS)
8. Premorbid Adjustment Scale
9. Verona Service Satisfaction Scale
10. An engagement measure
11. Socio-demographic & Service Receipt Inventory
12. SCID-P
13. OCPRIT

Overall study start date
08/03/2005

Completion date
07/03/2006

Eligibility

Key inclusion criteria
12-15 patients with first-episode psychosis, recruited from Bath & North East Somerset and North Somerset localities in the Avon & Wiltshire Mental Health Partnership NHS Trust.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
15

Key exclusion criteria
1. Presence of organic mental disorder
2. Mental retardation
3. Inadequate command of language

Date of first enrolment

08/03/2005

Date of final enrolment
07/03/2006

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Bristol
Bristol
United Kingdom
BS6 6JL

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Avon and Wiltshire Mental Health Partnership NHS Trust (UK) NHS R&D Support Funding

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