Randomised controlled trial (RCT) to evaluate a patient-held shared record card for patients with long term mental illness

Submission date 23/01/2004	Recruitment status No longer recruiting	Prospectively registered		
		[] Protocol		
Registration date 23/01/2004	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 29/01/2010	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers 844

Study information

Scientific Title

Study objectives

From the viewpoint of both primary and secondary care, what is the effect of a patient held shared record card on the pattern of service use and satisfaction with medical care for people with long term mental illness, compared to current practice?

Ethics approval required Old ethics approval format

Ethics approval(s) Not provided at time of registration

Study design Cluster randomised controlled parallel group trial

Primary study design Interventional

Secondary study design Randomised controlled trial

Study setting(s) GP practice

Study type(s) Other

Participant information sheet

Health condition(s) or problem(s) studied Schizophrenia and other psychoses

Interventions

Randomly selected patients entered on the North Birmingham Community District Mental Health Team's "community team case load census" database will be randomised: 1. Patient-held records 2. Current practice 100 patients will be required for each arm of the study. To avoid contamination bias, patients in the study and control arm will be drawn from different general practices.

Other Phase

Intervention Type

Not Applicable

Primary outcome measure

1. Adherence with treatment

- 2. Loss to follow up
- 3. Satisfaction with medical care
- 4. Number of psychiatric admissions/year
- 5. Length of inpatient stay

Secondary outcome measures

Not provided at time of registration

Overall study start date 01/11/1997

Completion date 31/07/2001

Eligibility

Key inclusion criteria

All patients with more than 2 years history of schizophrenia or other psychoses, severe personality disorder, or severe neurosis who require long-term supervision.

Participant type(s) Patient

Age group Adult

Sex Both

Target number of participants 200

Key exclusion criteria

- 1. Dementia, or other brain disorder
- 2. Alcohol or drug misuse
- 3. Learning disability
- 4.15 years or under
- 5.65 year or older.

Date of first enrolment

01/11/1997

Date of final enrolment 31/07/2001

Locations

Countries of recruitment

England

United Kingdom

Study participating centre Dept of General Practice Birmingham United Kingdom B15 2TT

Sponsor information

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

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.doh.gov.uk

Funder(s)

Funder type Government

Funder Name NHS Executive West Midlands (UK)

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?
<u>Results article</u>	results	01/03/2003		Yes	No