

An evaluation of Motivational (MI) plus Cognitive Therapy (CBT) for Schizophrenia and Substance Misuse

Submission date 29/08/2003	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 01/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 03/08/2009	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<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

Protocol serial number

G0200471

Study information

Scientific Title

Acronym

Motivational Interventions for Drug and Alcohol Use in Schizophrenia (MIDAS)

Study objectives

The aim of the study is to evaluate the benefits of the adjunct of a psychological treatment over standard available care for patients with schizophrenia and a co-morbid drug or alcohol problem.

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

Mental and behavioural disorder

Interventions

The experimental treatment (MI plus CBT) will consist of motivational interviewing to increase motivation to reduce substance use; CBT for help with both substance use reduction and symptom management; and relapse prevention strategies. Patients randomised to receive MI plus CBT will be offered up to 26 sessions over 12 months with treatment being located at home or clinic according to the patient's choice. The MI plus CBT treatment will be in addition to standard psychiatric care. The control group will receive standard psychiatric care alone.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Either readmission to hospital for reason related to psychosis or death from any cause/not admitted to hospital in the 12 month post-treatment period.

Key secondary outcome(s))

Added as of 6 February 2007: Measures of symptomatology, relapses, substance misuse and health economic analyses.

Completion date

02/05/2009

Eligibility

Key inclusion criteria

1. Diagnostic and Statistical Manual of Mental Disorders, Fourth edition (DSM IV) diagnostic criteria for schizophrenia, schizophreniform or schizoaffective disorder
2. Recorded contact and treatment from mental health services at the point of recruitment
3. Prescribed anti-psychotic medication
4. Alcohol use exceeding 28 units for males, 21 units for females on at least half the weeks in the previous 3 months
5. DSM IV diagnosis of drug and/or alcohol dependence or abuse
6. No significant history of organic factors implicated in the aetiology of psychotic symptoms
7. English speaking
8. Informed patient consent
9. Having a fixed abode

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

See inclusion criteria

Date of first enrolment

03/10/2004

Date of final enrolment

02/05/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Academic Division of Clinical Psychology
Manchester
United Kingdom
M23 9LT

Sponsor information

Organisation
University of Manchester (UK)

ROR
<https://ror.org/027m9bs27>

Funder(s)

Funder type
Research council

Funder Name
Medical Research Council (MRC) (UK)

Alternative Name(s)
Medical Research Council (United Kingdom), UK Medical Research Council, MRC

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
United Kingdom

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	01/10/2009		Yes	No