

The impact of routine utilization of two assessment scales on management of antipsychotic-induced side effects

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 25/04/2014	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

Protocol serial number
N0146155662

Study information

Scientific Title

Study objectives

To identify the effects of introduction of two assessment scales to regularly appraise antipsychotic-induced side effects in forensic psychiatric patients on management of these side effects and patients' clinical outcome.

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)

Not Specified

Health condition(s) or problem(s) studied

Mental and Behavioural Disorders: Psychosis

Interventions

Two assessment scales

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Changes/differences in the scales ratings at the baseline and the end-assessment within the control and study group and between groups. Differences in the general mental health rating at the beginning and the end-assessment between the control and the study groups. Associations between side effects assessment and the general mental health status.

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/04/2004

Eligibility

Key inclusion criteria

30 (study) and 30 (control) patients in the Forensic Mental Health Unit, Homerton

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

01/10/2003

Date of final enrolment

30/04/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Homerton Hospital NHS Trust

London

United Kingdom

E9 6SR

Sponsor information**Organisation**

Department of Health

Funder(s)**Funder type**

Government

Funder Name

East London and The City Mental Health NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration