

# 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

☐ Prospectively registered

☐ Protocol

**Registration date**  
30/09/2005

**Overall study status**  
Completed

☐ Statistical analysis plan

☐ Results

**Last Edited**  
25/04/2014

**Condition category**  
Mental and Behavioural Disorders

☐ Individual participant data

☐ Record updated in last year

## Plain English summary of protocol

Not provided at time of registration

## Contact information

### Type(s)

Scientific

### Contact name

Mr Albert Ngan

### Contact details

Homerton Hospital NHS Trust  
Homerton Row  
London  
United Kingdom  
E9 6SR

## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

### Study design

Randomised controlled trial

### Primary study design

Interventional

### Secondary study design

Randomised controlled trial

### Study setting(s)

Not specified

### Study type(s)

Not Specified

## Participant information sheet

### Health condition(s) or problem(s) studied

Mental and Behavioural Disorders: Psychosis

### Interventions

Two assessment scales

### Intervention Type

Other

### Phase

Not Specified

### Primary outcome measure

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.

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/10/2003

**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

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

30 (study) and 30 (control) patients

**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**

England

United Kingdom

**Study participating centre**

Homerton Hospital NHS Trust

London

United Kingdom

E9 6SR

# Sponsor information

## Organisation

Department of Health

## Sponsor details

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

East London and The City Mental Health NHS Trust (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