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

| Submission date | Recruitment status | [ ] Prospectively registered |
|---|---|---|
| 30/09/2005 | No longer recruiting | [ ] Protocol |
| **Registration date** | **Overall study status** | [ ] Statistical analysis plan |
| 30/09/2005 | Completed | [ ] Results |
| **Last Edited** | **Condition category** | [ ] Individual participant data |
| 25/04/2014 | 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

**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

### Study design
Randomised controlled trial

### Primary study design
Interventional

### 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