

# To determine the efficacy of administering the Liverpool University neuroleptic side-effect rating scale (LUNSERS) for identifying side effects of psychotropic medications.

<b>Submission date</b> 30/09/2004	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 30/09/2004	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 19/09/2013	<b>Condition category</b> 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

### Contact name

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### Contact details

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Nottingham  
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## Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

## Secondary identifying numbers

N0172139155

# Study information

## Scientific Title

### Study objectives

Does the use of the LUNSERs increase the reporting of side-effects of psychotropic medication?

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Not provided at time of registration

### Study design

Randomised Controlled Trial (Prospective)

### 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

### Interventions

The trial arms are self-reporting and physician questioning of side-effects versus usual care and administration of the LUNSERs scale to identify side-effects.

### Intervention Type

Other

### Phase

Not Specified

### Primary outcome measure

Use of LUNSERs, side-effects reported.

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

22/02/2004

**Completion date**

31/12/2005

## Eligibility

**Key inclusion criteria**

30 clients in each arm of the trial.

**Participant type(s)**

Patient

**Age group**

Not Specified

**Sex**

Not Specified

**Target number of participants**

60

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

22/02/2004

**Date of final enrolment**

31/12/2005

## Locations

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

Division of Psychiatry

Nottingham

United Kingdom

NG3 6AA

# Sponsor information

## Organisation

Department of Health

## Sponsor details

Richmond House  
79 Whitehall  
London  
United Kingdom  
SW1A 2NL

## Sponsor type

Government

## Website

<http://www.dh.gov.uk/Home/fs/en>

# Funder(s)

## Funder type

Government

## Funder Name

Nottinghamshire Healthcare 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