

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

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

Contact name

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Contact details

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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