

A randomised double-blind comparative trial investigating the level of pain experienced on induction of anaesthesia using either standard Propofol 1% or Propofol-Lipuro 1% with or without the addition of lignocaine 2% (1:10 ratio)

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 05/09/2012	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr A Mallick

Contact details
Anaesthetics
D Floor, Jubilee Building
Leeds General Infirmary
Great George Street
Leeds
United Kingdom
LS1 3EX
+44 (0)113 243 2799
Abhiram.Mallick@leedsth.nhs.uk

Additional identifiers

Protocol serial number

N0436125555

Study information

Scientific Title

Study objectives

To determine which propofol preparation affords the least pain on injection.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised double-blind comparative trial

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Surgery: Anaesthesia

Interventions

1. Standard Therapy
2. New Therapy

Intervention Type

Drug

Phase

Not Specified

Drug/device/biological/vaccine name(s)

Standard Propofol 1% or Propofol-Lipuro 1% with or without the addition of lignocaine 2%

Primary outcome(s)

Injection pain/no pain comparison between groups

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/04/2006

Eligibility

Key inclusion criteria

Patients undergoing uncomplicated elective surgery

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

01/08/2003

Date of final enrolment

01/04/2006

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Anaesthetics

Leeds

United Kingdom

LS1 3EX

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/05/2007		Yes	No