ISRCTN90337900 https://doi.org/10.1186/ISRCTN90337900

BIS monitoring and depth anaesthesia for cardioversion

Submission date 28/09/2007	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 28/09/2007	**Overall study status** Completed	
Last Edited 12/10/2017	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Dr M Parsloe

Contact details
Department of Anaesthesia
Conquest Hospital
The Ridge
St Leonards on Sea
United Kingdom
TN37 7RD

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0019186908

Study information

Scientific Title
BIS monitoring and depth anaesthesia for cardioversion

Study objectives
To compare differences in the haemodynamic parameters during cardioversion with sevoflurane and propofol anaesthesia.

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)
Hospital

Study type(s)
Treatment

Participant information sheet
Not available in web format, please use the contact details to request a patient information sheet

Health condition(s) or problem(s) studied
Surgery: Anaesthesia

Interventions
BIS monitoring vs standard treatment

Intervention Type
Drug

Phase
Not Applicable

Drug/device/biological/vaccine name(s)
Sevoflurane, propofol

Primary outcome measure
Optimisation of depth of anaesthesia

Secondary outcome measures
To assess appropriateness of using BIS monitor when anaesthetising patients for cardioversion

Overall study start date
08/11/2006

Completion date
01/06/2007

Eligibility

Key inclusion criteria
Patients undergoing elective day case cardioversions for cardiac arrhythmias

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
60

Key exclusion criteria
Any patient not willing to consent or not able to consent, with history of malignant hyperthermia or hypersensitivity to propofol

Date of first enrolment
08/11/2006

Date of final enrolment
01/06/2007

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Conquest Hospital
St Leonards on Sea
United Kingdom
TN37 7RD

Sponsor information

Organisation
Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Sponsor details
The Department of Health, 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 Sussex Hospitals 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