

BIS monitoring and depth anaesthesia for cardioversion

Submission date 28/09/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 28/09/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 12/10/2017	Condition category Surgery	<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
Dr M Parsloe

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

Additional identifiers

Protocol serial number
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

Study type(s)

Treatment

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(s)

Optimisation of depth of anaesthesia

Key secondary outcome(s)

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

Completion date

01/06/2007

Eligibility

Key inclusion criteria

Patients undergoing elective day case cardioversions for cardiac arrhythmias

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

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**

United Kingdom

England

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

Funder(s)**Funder type**

Government

Funder Name

East Sussex Hospitals NHS Trust (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration