

# BIS monitoring and depth anaesthesia for cardioversion

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

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## 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