

Design of the Optimal Technique in Cardiac Anaesthesia Recovery: The OPTICARE Trial

Submission date 16/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 05/01/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 26/08/2009	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Acronym

Opticare I

Study objectives

Not provided at time of registration

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**Health condition(s) or problem(s) studied**

Cardiac surgery patients

Interventions

Compare the effect of intravenous anaesthesia combined with thoracic epidural anaesthesia (TEA group), with total intravenous anaesthesia (general anesthetic [GA] group) without epidural anaesthesia

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/04/2003

Completion date

30/04/2004

Eligibility

Key inclusion criteria

Consecutive patients scheduled for elective cardiac surgery (coronary artery bypass graft [CABG], valve, redo CABG)

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/04/2003

Date of final enrolment

30/04/2004

Locations

Countries of recruitment

Netherlands

Study participating centre

PO Box 10500

Zwolle

Netherlands

8000 GM

Sponsor information

Organisation

Isala Clinics, Zwolle (Isala Klinieken) (Netherlands)

Sponsor details

Health Care Improvement Project

PO Box 10500

Zwolle

Netherlands

8000 GM

Sponsor type

Not defined

ROR

<https://ror.org/046a2wj10>

Funder(s)**Funder type**

Hospital/treatment centre

Funder Name

Isala Clinics, Zwolle (Isala Klinieken) (Netherlands)

Funder Name

Het Groene Land-Achmea Insurance (Netherlands)

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