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

Submission date 16/09/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 05/01/2005	**Overall study status** Completed	
Last Edited 26/08/2009	**Condition category** Surgery	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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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 anesthesia

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