

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

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

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## Additional identifiers

## 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

**Study type(s)**

Treatment

**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(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**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

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**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)

**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**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration