

Vergelijking tussen algehele anesthesie met en zonder toevoeging van een ruggenprik bij hartoperaties

No registrations found.

Ethical review	Not applicable
Status	Suspended
Health condition type	-
Study type	Interventional

Summary

ID

NL-OMON27352

Source

Nationaal Trial Register

Brief title

Easycab

Health condition

CABG, OPCAB, HTEA, coronary artery disease

Sponsors and support

Primary sponsor: Not applicable

Source(s) of monetary or material Support: Not applicable

Intervention

Outcome measures

Primary outcome

Combined endpoint of 30 days all cause mortality, myocardial infarction, stroke or acute renal failure requiring dialysis

Secondary outcome

Separate components of the primary endpoint, e.g.: 30 days all cause mortality, myocardial infarction, stroke or acute renal failure requiring dialysis

Study description

Background summary

N/A

Study objective

Thoracic epidural anesthesia with local anesthetics produces intense sympathicolysis and protects the heart against ischemia during off pump coronary artery surgery

Study design

30 days and 1 year

Intervention

High thoracic epidural anesthesia with local anesthetics

Contacts

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Scientific

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Eligibility criteria

Inclusion criteria

Patients > 18 year, scheduled for elective coronary artery surgery without heart lung machine. (Off pump CABG, OPCAB)

Exclusion criteria

Pre specified coagulation disorders

Study design

Design

Study type:	Interventional
Intervention model:	Parallel
Allocation:	Randomized controlled trial
Masking:	Open (masking not used)
Control:	Active

Recruitment

NL	
Recruitment status:	Suspended
Start date (anticipated):	01-01-2014
Enrollment:	700
Type:	Anticipated

Ethics review

Not applicable	
Application type:	Not applicable

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL3994
NTR-old	NTR4166
Other	: geen
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Study results

Summary results

N/A