

Optimising blood-circulation and oxygen delivery in planned abdominal aortic surgery

Submission date 03/08/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 03/09/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 13/09/2017	Condition category Surgery	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Palle Toft

Contact details
Department of Anaesthesiology and Intensive Care
Odense University Hospital
Sdr. Boulevard 29
Odense
Denmark
5000
-
palle.toft@ouh.regionssyddanmark.dk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Optimising stroke volume and oxygen delivery in elective abdominal aortic surgery: a randomised controlled trial

Study objectives

Elective abdominal aortic surgery is performed in patients with aneurism or occlusive atherosclerotic disease. These patients often have severe co-morbidity and are at high risk of postoperative complications. Maintaining optimal circulation during aortic surgery is difficult due to aortic cross clamping and often profound haemorrhage in combination with anaesthetising a patient with general atherosclerotic disease.

Precise and individual circulatory therapy can be performed by continuously monitoring and optimising the patient's stroke volume and oxygen delivery during and after surgery. Optimisation is performed by giving colloid boluses to achieve the individual optimal stroke volume intraoperatively, supplemented by infusion of Dobutamine postoperatively to maintain delivery of oxygen above 600 ml min⁻¹ m⁻².

This protocol may reduce postoperative complications and death, as well as length of stay in the Intensive Care Unit and hospital.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Local Medical Ethics Committee (Den Videnskabetiske Komite for Region Syddanmark) approved in June 2008 (ref: S-20080055)

Study design

Prospective randomised partly blinded controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Prevention

Participant information sheet

Not available in web format, please use contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Elective abdominal aortic surgery; Atherosclerotic abdominal aortic occlusive disease;
Abdominal aortic aneurism

Interventions

Patients were assigned to Individual Goal Directed Therapy (IGDT) or control groups by computer-generated random sequence.

The intervention period started preoperatively, when monitoring with the Lithium Dilution Cardiac Output (LiDCO)-plus-system was established and calibrated at arrival to the operating theatre. The intervention period ended 6 hours postoperatively. Patients were followed for 30 days postoperatively.

Establishment and calibration of the LiDCO-plus-system were carried out by a member of the research team who had no involvement in the peri- and postoperative care and decision making. This allowed complete blinding of both surgical, anaesthetic and Post Anaesthetic Care Unit (PACU) clinical teams to LiDCO-plus-system readings in the control group.

All anaesthetic interventions were at the discretion of the anaesthetist responsible for the perioperative management of the patient. All patients received general anaesthesia with fentanyl, thiopental, rocuronium and sevoflurane in oxygen/air. Before induction of anaesthesia an epidural catheter was inserted at the low thoracic level and an epidural infusion of bupivacain with fentanyl was started and continued until postoperative day 2 or 3.

Standard monitoring for both groups included continuous pulse oxymetry, electrocardiography, invasive arterial and central venous blood pressure monitoring, and spirometry with inspiratory and expiratory oxygen, carbondioxide and anaesthetic gas monitoring. Arterial blood gases were analysed at predefined points in both groups.

Stroke volume index (SVI), cardiac index (CI) and oxygen delivery index (DO₂I) were continuously monitored, by lithium indicator dilution and pulse power analysis using the LiDCO-plus-system in all patients, but data was blinded in the control group.

All patients were treated to achieve a heart rate < 100 bpm or <20% above baseline, a mean arterial pressure (MAP) between 60-100 mmHg, a central venous pressure (CVP) between 4-16, body temperature > 36,5°C, an arterial oxygen saturation (SaO₂) > 94%, a haemoglobin concentration > 6 mmol l⁻¹, and an urine output > 0.5-1.0 ml min⁻¹ kg⁻¹ in the postoperative period.

In all patients crystalloid, colloid, blood products and vasopressors were administered in the peri- and postoperative periods by the anaesthetist based on intra- and postoperative losses, standard haemodynamic parameters and blood-gases.

Intervention: Patients in the IGDT group in the peri- and postoperative period received 250 ml boluses of intravenous colloid solution (Voluven®, Fresenius Kabi AB, Upsala, Sweden) to achieve a sustained rise in SVI of at least 10% for 20 min. Fluid boluses of Voluven® were repeated if SV subsequently decreased or if there was clinical suspicion of hypovolaemia. Furthermore, in the postoperative period, the IGDT group received dobutamine up to a maximum of 10 µg kg⁻¹ min⁻¹ if DO₂I did not reach 600 ml min⁻¹ m⁻² with intravenous fluid alone. During infusion of dobutamine, monitoring was supplemented with 5-lead-electrocardiography, and at signs of myocardial ischemia or heart rate > 100 min⁻¹ or > 20% above baseline, infusion was reduced or discontinued.

Intervention Type

Procedure/Surgery

Phase

Not Applicable

Primary outcome measure

One or more severe postoperative complications:

1. Septic shock
2. Pneumonia
3. Superficial wound infection
4. Deep wound infection
5. Abdominal infection
6. Urinary tract infection
7. Pulmonary embolus
8. Acute Respiratory Distress Syndrome (ARDS)
9. Cardiac arrest
10. Acute coronary syndrome
11. Cardiac arrhythmia (acute treatment needed)
12. Pulmonary oedema
13. Deep venous thrombosis
14. Cerebral thrombosis
15. Cerebral haemorrhage
16. Lower limb paresis
17. Acute kidney insufficiency
18. Intraabdominal hypertension
19. Severe upper gastrointestinal bleeding
20. Gastrointestinal paralysis
21. Creatine Kinase (CK) > 5000
22. Reoperation
23. Readmission to ICU
24. Need of respirator
25. Need of hemodialysis
26. Dead

Secondary outcome measures

1. Flow-related haemodynamic parameters (SVI and Do2I) measured by the LiDCO-plus-system (LiDCO Ltd., Cambridge, UK)
2. Length of stay in Intensive Care Unit
3. Length of hospital stay

Overall study start date

01/06/2008

Completion date

01/01/2010

Eligibility**Key inclusion criteria**

Consecutive patients admitted for elective abdominal aortic surgery

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

85

Key exclusion criteria

1. Chronic renal end-failure
2. Preoperative Lithium therapy
3. Body weight < 40 Kg (88,18 lbs)

Date of first enrolment

01/06/2008

Date of final enrolment

01/01/2010

Locations

Countries of recruitment

Denmark

Study participating centre

Department of Anaesthesiology and Intensive Care

Odense

Denmark

5000

Sponsor information

Organisation

Department of Anaesthesiology Kolding (Denmark)

Sponsor details

Lillebaelt Hospital Kolding

Skovvangen 2-8

Kolding

Denmark
6000
+45 7636 2000
jannie.bisgaard@slb.regionsyddanmark.dk

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/037y5zq83>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Lillebaelt Hospital Kolding (Denmark) - Local research fund

Funder Name

The Toyota Fund (Denmark)

Funder Name

Research Initiative of The Danish Society of Anaesthesiology and Intensive Care Medicine (Denmark)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
-------------	---------	--------------	------------	----------------	-----------------

[Other publications](#)

01/01/2010

Yes

No

[Results article](#)

01/02/2013

Yes

No