

Water warming garment versus forced air warming system in prevention of intraoperative hypothermia during liver transplantation: a randomised controlled trial

Submission date 29/08/2002	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 29/08/2002	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 05/09/2007	Condition category Digestive System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Study objectives

Not provided at time of registration

Ethics approval required

Old ethics approval format

Ethics approval(s)

The institutional review board at Vanderbilt University approved the study protocol.

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

Liver disease

Interventions

The intraoperative use of either forced-air warmer or water warming garment for maintenance of patient temperature during surgery

Intervention Type

Other

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

01/01/2001

Completion date

01/01/2002

Eligibility

Key inclusion criteria

Patients with liver disease undergoing liver transplantation or liver resection under general anesthesia

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

24

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2001

Date of final enrolment

01/01/2002

Locations

Countries of recruitment

United States of America

Study participating centre

Department of Anesthesiology

Nashville

United States of America

TN 37232-4125

Sponsor information

Organisation

Vanderbilt University Medical Center (USA)

Sponsor details

Department of Anesthesiology

504 Oxford House

1313 21st Ave S

Nashville

United States of America

TN 37232-4125

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/05dq2gs74>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Vanderbilt University Medical Center (USA) - Department of Anesthesiology

Funder Name

MTRE Advanced Technologies Ltd (Israel)

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?
Results article	Results	19/11/2002		Yes	No