

Anesthesiological service and treatment quality

Submission date 27/08/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 03/10/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 06/09/2011	Condition category 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N/A

Study information

Scientific Title

Study objectives

The treatment quality of anesthesiological services is being analyzed for the purpose to find independent predictors for outcome quality.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics Committee of Charité - Berlin Medical University (Universitätsmedizin Berlin), Geschaeftsstelle, Schumannstr. 20/21, 10117 Berlin, Germany. Approved on 21 August 2007. (ref: EA1/143/07).

Study design

Retrospective observational study.

Primary study design

Observational

Secondary study design

Cross-section survey

Study setting(s)

Not specified

Study type(s)

Other

Participant information sheet**Health condition(s) or problem(s) studied**

Anesthesia

Interventions

Anesthesia records during the period of study will be checked, pertinent data entered into statistical database, and analyzed afterwards with a statistical software.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

Outcome quality. This will be analyzed in the statistical database. It will be measured in terms of length of hospital stay and length of stay in recovery room.

Secondary outcome measures

Independent predictors: Use of various anesthesia techniques, ventilation techniques, anesthesiological approaches, and utilization of different medications.

Overall study start date

01/07/2006

Completion date

30/06/2009

Eligibility

Key inclusion criteria

All patients receiving anesthesia in our department.

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

150,000 patients

Key exclusion criteria

None

Date of first enrolment

01/07/2006

Date of final enrolment

30/06/2009

Locations

Countries of recruitment

Germany

Study participating centre

Charité - Berlin Medical University

Berlin

Germany

D-13353

Sponsor information

Organisation

Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

Sponsor details

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Sponsor type

University/education

Website

<http://www.charite.de/ch/anaest/>

ROR

<https://ror.org/001w7jn25>

Funder(s)

Funder type

University/education

Funder Name

Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

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

