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Anesthesiological service and treatment quality

Submission date 27/08/2007	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 03/10/2007	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 06/09/2011	**Condition category** Surgery	[_] Individual participant data [_] 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 (Universitaetsmedizin 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
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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