

Taurolidine-citrate vs heparin as catheter lock solutions in paediatric patients

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
07/12/2006	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
16/01/2007	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
08/02/2011	Infections and Infestations	<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Uwe Querfeld

Contact details

Department of Pediatric Nephrology
Charité Universitätsmedizin Berlin
Berlin
Germany
13353

Additional identifiers

Protocol serial number

EA2/084/06

Study information

Scientific Title

Acronym

Taurolock study

Study objectives

Use of taurolidine-citrate for catheter locking leads to a significant reduction of bacterial growth and catheter-related infections in paediatric patients with implanted central venous catheters

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approval received by local ethics committee (Ethikkommission 2 am Campus Virchow Klinikum) on 29 September 2006 (ref: EA2/084/06).

Study design

Randomized prospective study in pediatric hospital with stratification for patients treated in renal/oncology/hematology departments

Primary study design

Interventional

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Catheter-related infections

Interventions

Taurolidine-citrate vs heparin as catheter lock solutions

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Bacterial growth/biofilm formation in removed catheters

Key secondary outcome(s)

Catheter-related infections, catheter occlusions

Completion date

31/12/2007

Eligibility

Key inclusion criteria

Paediatric patients with newly implanted central venous catheters

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Sex

Not Specified

Key exclusion criteria

Bacteremia/sepsis ongoing; allergy against heparin or taurolidine

Date of first enrolment

01/01/2007

Date of final enrolment

31/12/2007

Locations

Countries of recruitment

Germany

Study participating centre

Department of Pediatric Nephrology

Berlin

Germany

13353

Sponsor information

Organisation

Tauropharm (Germany)

Funder(s)

Funder type

Industry

Funder Name

Tauropharm GmbH (Germany)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration