

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

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

Contact name

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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