

Investigation in rates of bacterial contamination of tunnelled lines with and without elimination of presumed static electrical charge

Submission date
12/09/2003

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
12/09/2003

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
16/12/2008

Condition category
Infections and Infestations

☐ 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

N0012120116

Study information

Scientific Title

Study objectives

Does pre-injection of the line packet with sterile saline prior to opening reduce the bacterial contamination rate?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Infections and Infestations: Bacterial contamination

Interventions

Random allocation to (A) Treatment for bacterial contamination (B) Standard treatment.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Rates of bacterial contamination of the lines before and after tunnelling

Secondary outcome measures

Long term infection rates of the line

Overall study start date

01/11/2002

Completion date

30/04/2003

Eligibility

Key inclusion criteria

All children undergoing placement of a percutaneous, tunnelled silastic venous access line in the interventional radiology suite will be eligible for inclusion.

Participant type(s)

Patient

Age group

Child

Sex

Not Specified

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2002

Date of final enrolment

30/04/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre**Surgery Unit**

London

United Kingdom

WC1N 1EH

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

Great Ormond Street Hospital for Children NHS Trust / Institute of Child Health (UK)

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	01/07/2005		Yes	No