Central venous catheter use in paediatric oncology patients in the UK: mechanical problems

Submission date	Recruitment status	Prospectively registered
01/07/2001	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
01/07/2001	Completed	Results
Last Edited	Condition category	Individual participant data
03/03/2015	Cancer	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

- - -

Contact details

UKCCCR Register Co-ordinator MRC Clinical Trials Unit 222 Euston Road London United Kingdom NW1 2DA

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers SC 9606

Study information

Scientific Title

Central venous catheter use in paediatric oncology patients in the UK: mechanical problems

Study objectives

Added 07/08/2009:

The aim of this trial is to determine the optimum technique for ensuring central venous catheters are not subject to mechanical failure (eg. falling out).

As of 07/08/2009 this record was extensively updated. All updates can be found under the relevant field with the above update date.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Multicentre randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Miscellaneous paediatric cancer

Interventions

- 1. Group A: Skin only exit site suture on central venous catheter
- 2. Group B: Skin and line exit site suture on central venous catheter

Mechanical problems include dislodgment and migration documented in the post operative period.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Added 07/08/09:

- 1. Mechanical failure rate.
- 2. Correlation with exit suture type and whether additional procedures (eg. bone marrow, lumbar puncture) are undertaken at the same anaesthetic as the central venous line insertion.

Secondary outcome measures

Not provided at time of registration

Overall study start date

15/12/1996

Completion date

31/12/2000

Eligibility

Key inclusion criteria

- 1. Patient under the care of a UKCCSG centre
- 2. Patients must have cancer

Participant type(s)

Patient

Age group

Child

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Patients without cancer who require a central venous catheter for treatment of immunodeficiency, chronic antibiotic administration, dialysis or total parenteral nutrition are excluded.

Date of first enrolment

15/12/1996

Date of final enrolment

31/12/2000

Locations

Countries of recruitment

England

United Kingdom

Study participating centre MRC Clinical Trials Unit London United Kingdom NW1 2DA

Sponsor information

Organisation

Cancer Research UK (CRUK) (UK)

Sponsor details

PO Box 123 Lincoln's Inn Fields London United Kingdom WC2A 3PX +44 (0)207 317 5186 kate.law@cancer.org.uk

Sponsor type

Charity

Website

http://www.cancer.org.uk

ROR

https://ror.org/054225q67

Funder(s)

Funder type

Charity

Funder Name

Cancer Research UK (CRUK) (UK)

Alternative Name(s)

CR_UK, Cancer Research UK - London, CRUK

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

United Kingdom

Funder Name

United Kingdom Children's Cancer Study Group (UKCCSG)

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