

Hibiscrub as cleansing solution for pin site care

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| Submission date 30/09/2005 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol |
| Registration date 30/09/2005 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results |
| Last Edited 07/06/2017 | Condition category Surgery | <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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0205157956

Study information

Scientific Title

Hibiscrub as cleansing solution for pin site care

Study objectives

To confirm the null hypothesis that there is no difference in the clinical effectiveness of water and hibiscrub versus the current standard of aseptic technique and Na CL 0.9% as cleansing agent for pin sites in patients with external fixators.

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)

Hospital

Study type(s)

Prevention

Participant information sheet

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

Health condition(s) or problem(s) studied

Surgery: Orthopaedic

Interventions

Water and hibiscrub versus the current standard of aseptic technique and Na CL 0.9%

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome measure

1. Patient infection rate: patients with more than 2 pin site infection in any of the five visits
2. The length of the hospital stay
3. Length of time spent by nursing staff performing pin site care and equipment used

Secondary outcome measures

Not provided at time of registration

Overall study start date

20/12/2004

Completion date

19/12/2008

Eligibility

Key inclusion criteria

Patients aged over 16 with an external fixator in any of the Royal London Hospital wards - outpatients department

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

20/12/2004

Date of final enrolment

19/12/2008

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Royal London Hospital

London

United Kingdom

E1 1BB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House

79 Whitehall

London

United Kingdom

SW1A 2NL

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Sponsor type

Government

Website

<http://www.dh.gov.uk/Home/fs/en>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Barts and The London NHS Trust (UK)

Funder Name

NHS R&D Support Funding

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