

The use of topical antibacterial skin ointment (TASO) in the management of soft tissue wounds: A randomised controlled trial

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 08/04/2014	Condition category Injury, Occupational Diseases, Poisoning	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0186150989

Study information

Scientific Title

Study objectives

Is there any improvement in wound healing or cosmesis by prescription of topical ointments following suture of soft tissue wounds?

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

Injury, Occupational Diseases, Poisoning: Wound healing

Interventions

Antibacterial ointment (TASO) vs placebo vs no ointment - A randomised controlled trial

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

At review for suture removal the wounds would be assessed for healing. At a subsequent review, the wound would be assessed for cosmesis. The groups would be compared and the best treatment modality ascertained.

Secondary outcome measures

Not provided at time of registration

Overall study start date

30/04/2004

Completion date

30/04/2005

Eligibility

Key inclusion criteria

Patients with facial lacerations requiring suture

Participant type(s)

Patient

Age group

Not Specified

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

30/04/2004

Date of final enrolment

30/04/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Maxillofacial Department

Poole

United Kingdom

BH15 2JB

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Poole Hospital NHS Trust (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