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

Submission date	Recruitment status	[ ] Prospectively registered
30/09/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2005	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
08/04/2014	Injury, Occupational Diseases, Poisoning	[ ] Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr Michael Bater

Contact details
Maxillofacial Department
Poole Hospital NHS Trust
Longfleet Road
Poole
United Kingdom
BH15 2JB

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