

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

<b>Submission date</b> 30/09/2005	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 30/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 08/04/2014	<b>Condition category</b> 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

Mr Michael Bater

### Contact details

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## Additional identifiers

### Protocol serial number

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

**Study type(s)**

Not Specified

**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(s)**

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.

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

30/04/2005

**Eligibility****Key inclusion criteria**

Patients with facial lacerations requiring suture

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Not Specified

**Sex**

Not Specified

**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**

United Kingdom

England

**Study participating centre**

**Maxillofacial Department**

Poole

United Kingdom

BH15 2JB

**Sponsor information****Organisation**

Department of Health

**Funder(s)****Funder type**

Government

**Funder Name**

Poole Hospital NHS Trust (UK)

## **Results and Publications**

**Individual participant data (IPD) sharing plan**

**IPD sharing plan summary**

Not provided at time of registration