

Randomised controlled trial of excision and grafting with same flap, deffating of skin flap and treatment with steristrips of pre-tibial skin lacerations

Submission date 30/09/2005	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 30/09/2005	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 17/04/2015	Condition category Surgery	☐ Individual participant data
		☐ 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

Protocol serial number

N0224149878

Study information

Scientific Title

Randomised controlled trial of excision and grafting with same flap, deffating of skin flap and treatment with steristrips of pre-tibial skin lacerations

Study objectives

What is the best treatment option in an A and E setting for cases of pre-tibial lacerations? Three different treatments will be compared to try to establish the better option for patients attending A and E. Healing times will be specifically looked at in each category and pain scores to help improve patient care for the future.

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)

Treatment

Health condition(s) or problem(s) studied

Surgery: Skin transplantations

Interventions

Excision and grafting with the same flap versus deffating of the skin flap versus put ting steristrips to hold the skin.

Intervention Type

Procedure/Surgery

Phase

Not Specified

Primary outcome(s)

Is there any reduction in mean healing time with any suggested treatments?
Any improvement in pain scores?

Key secondary outcome(s))

Not provided at time of registration

Completion date

30/04/2007

Eligibility

Key inclusion criteria

Patients attending A and E with pre-tibial lacerations.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

1. Patients who are unable to decide for themselves, patients with dementia, mentally ill
2. Those who cannot understand English/need an interpreter
3. Patients on anticoagulants except aspirin

Date of first enrolment

24/06/2004

Date of final enrolment

30/04/2007

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

South Devon Health Care NHS Trust

Torquay

United Kingdom

TQ2 7AA

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

South Devon Healthcare NHS Trust (UK), NHS R&D Support Funding

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration