

Buried absorbable sutures are an acceptable alternative to removed non-absorbable sutures for closure of wounds of the face

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| Submission date 30/09/2004 | Recruitment status No longer recruiting | <input type="checkbox"/> Prospectively registered |
| | | <input type="checkbox"/> Protocol |
| Registration date 30/09/2004 | Overall study status Completed | <input type="checkbox"/> Statistical analysis plan |
| | | <input type="checkbox"/> Results |
| Last Edited 28/11/2014 | Condition category Injury, Occupational Diseases, Poisoning | <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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Contact details
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RH19 3DZ

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0190085892

Study information

Scientific Title

Buried absorbable sutures are an acceptable alternative to removed non-absorbable sutures for closure of wounds of the face

Study objectives

Are buried absorbable sutures an acceptable alternative to non-absorbable sutures in closure of facial 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)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Facial wounds

Interventions

Absorbable sutures vs non-absorbable sutures

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Assessment of end results of scarring using the Vancouver system for measuring height, width & thickness of scar
2. Pain, irritation & redness and patient satisfaction

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/09/2000

Completion date

01/12/2005

Eligibility

Key inclusion criteria

Any facial lesion in an adult for direct single layer closure under local anaesthetic as an out-patient

Participant type(s)

Patient

Age group

Adult

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

01/09/2000

Date of final enrolment

01/12/2005

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

The Queen Victoria Hospital NHS Trust

East Grinstead

United Kingdom

RH19 3DZ

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

Queen Victoria 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