

Comparison between sheet grafts and 1:1 mesh grafts in burnt patients

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
11/04/2012

Condition category
Injury, Occupational Diseases, Poisoning

☐ Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0190155442

Study information

Scientific Title

Study objectives

To compare outcome in terms of intervention rate, complication rate and cosmesis for sheet graft vs. 1:1 meshed graft

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

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Injury, Occupational Diseases, Poisoning: Burns

Interventions

Sheet grafts vs 1:1 mesh grafts.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Complication rates
2. Scar control
3. Patient and PT visual assessment using Vancouver scale.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/06/2005

Completion date

01/06/2006

Eligibility

Key inclusion criteria

All patients aged above 16 years of age undergoing a split skin grafting procedure that would be suitable for either sheet or 1:1 mesh (currently used interchangeably)

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

50

Key exclusion criteria

Those who cannot understand the principles of the study or are incapable of making judgments; those on steroids, those who have conditions which impair healing, those with burn areas greater than 20% and those whom it will not be possible to follow up.

Date of first enrolment

01/06/2005

Date of final enrolment

01/06/2006

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
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Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

The Queen Victoria Hospital NHS Trust (UK), NHS R&D Support Funding

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	18 - month follow up results	01/02/1987		Yes	No