

Pilot study to assess the feasibility and patient numbers required prior to a randomised controlled trial of the effect of visual distraction for painful procedures in the burns unit

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 03/11/2015	Condition category 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

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Additional identifiers

Protocol serial number
N0060131616

Study information

Scientific Title

Pilot study to assess the feasibility and patient numbers required prior to a randomised controlled trial of the effect of visual distraction for painful procedures in the burns unit

Study objectives

Does the use of 2D video glasses with DVD film images and stereo sound during dressing changes for patients with burns decrease the intensity of their experience of pain and anxiety?

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

Pain

Interventions

Short visual analogue scale patient rated scores for pain and anxiety, an appropriate questionnaire and short concluding taped interview after intervention. Quantitative methods will be used.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Improved pain control and toleration of regular dressing change and debridement of burns with reduced anxiety and awareness.

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/09/2003

Eligibility**Key inclusion criteria**

Minimum 10 consenting adults suitable to trial criteria and protocols.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Does not meet inclusion criteria

Date of first enrolment

01/04/2003

Date of final enrolment

30/09/2003

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Chelsea & Westminster Hospital

London

United Kingdom

SW10 9NH

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Chelsea and Westminster Healthcare NHS Trust (UK)

Results and Publications

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes