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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	Recruitment status	[ ] Prospectively registered
30/09/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
30/09/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
03/11/2015	Injury, Occupational Diseases, Poisoning	[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Hospital

Study type(s)
Treatment

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

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/04/2003

Completion date
30/09/2003

Eligibility

Key inclusion criteria
Minimum 10 consenting adults suitable to trial criteria and protocols.

Participant type(s)
Patient

Age group
Adult

Sex
Both

Target number of participants
10

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
England

United Kingdom

Study participating centre
Chelsea & Westminster Hospital
London
United Kingdom
SW10 9NH

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
Hospital/treatment centre

Funder Name
Chelsea and Westminster Healthcare 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