

Evaluation of an external device reflecting infra-red body heat loss to the neck area for chronic mechanical neck pain: a randomised, double-blind controlled trial

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 28/02/2014	Condition category Musculoskeletal Diseases	<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

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0263115136

Study information

Scientific Title

Study objectives

Can the infra-red reflective device reduce neck pain and disability advantageously in patients suffering from chronic mechanical neck pain as compared with a placebo device?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised, double-blind 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

Health condition(s) or problem(s) studied

Musculoskeletal Diseases: Neck pain

Interventions

1. Real neck device
2. Dummy device

Intervention Type

Device

Phase

Not Specified

Primary outcome measure

Pain visual analogue scale, neck disability index and general health status (SF36).

Secondary outcome measures

Not provided at time of registration

Overall study start date

14/05/2002

Completion date

31/07/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

60 patients from Neurosurgery/NSITU.

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

14/05/2002

Date of final enrolment

31/07/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Neurosurgery

London

United Kingdom

WC1N 3BG

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type

Government

Website

<http://www.doh.gov.uk>

Funder(s)

Funder type

Government

Funder Name

University College London Hospitals 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