

The effects of neural taping on pain and shoulder function in patients with neck and referred arm pain. A single blind pilot study

Submission date 30/09/2005	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 30/09/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
Last Edited 30/09/2013	Condition category Musculoskeletal Diseases	<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

Miss Elizabeth Anne Foley

Contact details

Colton

Leeds

United Kingdom

LS

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0436154190

Study information

Scientific Title

Study objectives

To investigate the effectiveness of the taping procedure on pain and function in these types of patients

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Single blind pilot study

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 and arm pain

Interventions

Randomised controlled trial

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Outcome measures to be used are a visual analogue scale and the neck disability index

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/07/2004

Completion date

01/09/2004

Eligibility

Key inclusion criteria

Convenience sample of patients with neck and arm pain attending the Neurosurgery clinic of Mr Nick Phillips (Neurosurgeon).

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Target Recruitment: 26

Key exclusion criteria

Known skin conditions or tape allergies, diabetes mellitus, pregnancy, known congenital abnormality of the nervous system, rheumatoid arthritis

Date of first enrolment

01/07/2004

Date of final enrolment

01/09/2004

Locations

Countries of recruitment

United Kingdom

Study participating centre

Colton

Leeds

United Kingdom

LS

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)**Funder type**

Government

Funder Name

Leeds Teaching Hospitals 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