

Advances in vestibular rehabilitation: use of videos and visual motion

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 02/02/2017	Condition category Ear, Nose and Throat	<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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0263152080

Study information

Scientific Title

Advances in vestibular rehabilitation: use of videos and visual motion

Study objectives

Will rehabilitation incorporating a home video with visual motion provide similar objective & subjective improvements in balance & symptoms compared to a programme using exposure to full field optokinetic?

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)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied

Ear, Nose and Throat

Interventions

1. Equipment-based visual
2. Equipment-based visual via video

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

At each assessment validated questionnaires and objective tests to assess the effect of treatment on symptom severity, balance, quality of life, activities of daily living, and emotional state will be used. Objective tests include posturography.

Secondary outcome measures

Not provided at time of registration

Overall study start date

23/11/2004

Completion date

31/07/2007

Eligibility

Key inclusion criteria

75 patients with intractable vestibular symptoms from Neuro-otology

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

75

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

23/11/2004

Date of final enrolment

31/07/2007

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

National Hospital for Neurology & Neurosurgery
London
United Kingdom
WC1N 3BG

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House
79 Whitehall
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SW1A 2NL
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dhmail@doh.gsi.org.uk

Sponsor type

Government

Website

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

Funder(s)

Funder type

Government

Funder Name

UCL Hospitals NHS Foundation Trust (UK)

Funder Name

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