Non-surgical treatment of benign paroxysmal positional vertigo (BPPV) - a randomised controlled trial of procedures and practices

Submission date 12/09/2003	Recruitment status Stopped	Prospectively registered
		☐ Protocol
Registration date	Overall study status	Statistical analysis plan
12/09/2003	Stopped	Results
Last Edited	Condition category	Individual participant data
19/10/2011	Signs and Symptoms	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

N0256111276

Study information

Scientific Title

Study objectives

How efficient are the different non-surgical treatments that are in current clinical practice? Is any one method better than the other?

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)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Signs and Symptoms: Benign paroxysmal positional vertigo (BPPV)

Interventions

- 1. Particle repositioning manoeuvre alone
- 2. Particle repositioning manoeuvre plus soft collar & sleeping instructions
- 3. Particle repositioning manoeuvre with bone vibrator
- 4. Particle repositioning manoeuvre with bone vibrator plus soft collar & sleeping instructions
- 5. Control group treated with rehabilitation exercises

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

All the above described methods are recognized standard treatment procedures for BPPV in current clinical practice. Patients will be reviewed in the clinic 8 weeks post-treatment, when they will be required to answer the Dizziness Handicap Inventory (DHI). Questionnaire. A negative Hallpike's test during this visit would be considered as a positive outcome.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/02/2002

Completion date

31/07/2003

Reason abandoned (if study stopped)

Poor recruitment

Eligibility

Key inclusion criteria

Outpatients who have a positive Dix-Hallpike's positional test diagnostic of BPPV Five groups, approx 20 patients per group. 5 Patients recruited 2001- 2002.

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

100

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/02/2002

Date of final enrolment

31/07/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Audiological Medicine Department, RNTNE
London
United Kingdom
WC1X 8DA

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

The Royal Free Hampstead 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 summaryNot provided at time of registration