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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 [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 12/09/2003	**Overall study status** Stopped	
Last Edited 19/10/2011	**Condition category** Signs and Symptoms	

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
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
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Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
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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 summary
Not provided at time of registration