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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

Protocol serial number
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

Study type(s)
Not Specified

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(s)
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.

Key secondary outcome(s))
Not provided at time of registration

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

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

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
United Kingdom

England

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

Sponsor information

Organisation
Department of Health (UK)

Funder(s)

Funder type
Government

Funder Name
The Royal Free Hampstead NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration