The feasibility of Whole Body Vibration in institutionalised elderly persons and its influence on muscle performance, balance and mobility: a randomised controlled trial

Submission date	Recruitment status No longer recruiting	Prospectively registered		
30/06/2005		☐ Protocol		
Registration date 06/07/2005	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 15/04/2008	Condition category Other	[] Individual participant data		

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

Study information

Scientific Title

Study objectives

Whole Body Vibration is feasible in institutionalised elderly persons and improves the mobility and muscle performance compared to control.

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

Institutionalised elderly persons

Interventions

Intervention: A 6-week exercise program on a vertical vibration platform (Power-Plate, Badhoevedorp, The Netherlands), three times per week (with a minimum of 1-day rest in between) and consisting in 6 static exercises targeting lower limb muscles. The exercise volume and intensity being progressively increased according to the overload-principle.

Control: Exactly the same exercise program on the vibration platform as the intervention group, but without vertical vibration (the sound of the motor of the vibration platform being reproduced by a tape recorder during each bout of exercise).

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Feasibility: taking into account continuation of the exercise program and/or occurence of complications related to the Whole Body Vibration exercises.

Improvement due to the intervention: taking into account balance and gait (using the timed upand-go test and Tinetti-test), upper limb and lower body flexibility (using the back scratch and chair sit-and-reach test), maximal grip strength (using a Martin vigorimeter device, Elmed, Addison, USA), closed chain bilateral leg extension (using a linear isokinetic multi-joint dynamometer, Aristokin®, Lode, Groningen, The Netherlands).

Secondary outcome measures

No secondary outcome measures

Overall study start date

01/12/2003

Completion date

15/12/2003

Eligibility

Key inclusion criteria

All residents of a nursing home (Van Zanden, Brussels, Belgium; capacity of 102 beds) within dependence categories O, A and B according to the scale of Katz et al. (1963) for basic activities of daily living.

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

24

Key exclusion criteria

Mainly based on contra-indications for Whole Body Vibration: presence of infectious disease, insulin-dependent diabetes mellitus, endogenous osteosynthethical material, knee or hip prosthesis, pacemaker, epilepsy, musculo-skeletal disorders and cognitive or physical dysfunction interfering with test and training procedures.

Date of first enrolment

01/12/2003

Date of final enrolment

15/12/2003

Locations

Countries of recruitment

Belgium

Study participating centre Geriatric & Gerontology Department

Brussels Belgium B-1090

Sponsor information

Organisation

Free University of Brussels (VUB) - Gerontology (Belgium)

Sponsor details

Laarbeeklaan 103 Brussels Belgium B-1090

Sponsor type

University/education

ROR

https://ror.org/006e5kg04

Funder(s)

Funder type

University/education

Funder Name

Free University of Brussels (VUB) - Gerontology

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results:	22/12/2005		Yes	No