

The effect of a mobile safety alarm on going outside, feeling safe, fear of falling and quality of life in community living older persons.

Submission date 20/07/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 02/09/2010	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 02/09/2010	Condition category Other	<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

Protocol serial number
N/A

Study information

Scientific Title

The effect of a mobile safety alarm on going outside, feeling safe, fear of falling and quality of life in community living older persons: A randomised controlled trial

Study objectives

What is the effectiveness of a mobile alarm in changing the frequency of going outside of older people and what are their experiences about safety, fear of falling and quality of life?

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Medical Ethical Committee of University of Amsterdam/Academic Medical Centre Amsterdam reviewed this study and confirmed that as a social study, it did not require ethics approval.

Study design

2 armed randomised controlled trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

New communication technology and services for older persons

Interventions

The intervention consisted of a mobile safety alarm with an inbuilt drop sensor. The mobile alarm was a prototype in development and uses a positioning system patented Europe-wide. The alarm went off by pressing a button. An integrated hand free function allowed the user to speak to personnel of an emergency call centre, even if the user didn't hold the device at their ear. If the user had fallen, the system automatically registered this and independently made a voice call to the emergency call centre. The Butler used a new kind of positioning system. The user of the alarm was located in 3 steps: their rough location was found via the cell phone network: from close-range, their position was pinpointed via an integrated tracking device and a beeping sound emitted from the device. This procedure allowed first-aiders to reliably find someone, even where conventional positioning systems such as GPS fail, such as in buildings or underground garages.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Primary outcome measure was change in the frequency of going outside. The frequency of going outside will be assessed in a telephone interviews at baseline and at 1, 2, 4 and 6 months follow-up.

Key secondary outcome(s)

The following secondary outcomes will be assessed in a telephone interview at baseline, and at 1, 2, 4 and 6 months follow-up:

1. Fear of Falling (Visual Analogue Scale [VAS]-FOF)
2. Unsafe feelings (VAS for Feeling Unsafe)
3. Quality of Life (VAS-EQ-5D).
4. Falls (data on falls were collected prospectively with use of fall calendars)

Completion date

31/03/2010

Eligibility**Key inclusion criteria**

1. Having a home-based alarm
2. Community-dwelling
3. Being able to go outside alone
4. Willingness to participate

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

All

Key exclusion criteria

Confined to bed

Date of first enrolment

16/07/2009

Date of final enrolment

31/03/2010

Locations**Countries of recruitment**

Netherlands

Study participating centre

Academic Medical Center
Amsterdam

Netherlands
1105 AZ

Sponsor information

Organisation
City council Amsterdam (Netherlands)

Funder(s)

Funder type
Government

Funder Name
City council of Amsterdam (Netherlands)

Results and Publications

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?
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes