

A feasibility study to test the effect of a set of behavioural change strategies in improving physical activity levels in adults aged 45 and over

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
07/05/2018	No longer recruiting	<input checked="" type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
10/05/2018	Completed	<input checked="" type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
06/01/2026	Other	

Plain English summary of protocol

Background and study aims

It is well known that being physically inactive is good for mental and physical health and that adults who are involved in regular physical activity are protected from heart disease, cancer and other illnesses. On the other hand, adults who are inactive are more likely to get mental and physical health conditions. Inactive adults tend to have poorer balance and mobility and fall more often and often they become socially isolated. Many countries worldwide are trying to implement national strategies to get older adults moving and success has been limited.

Research has shown that behavioural change strategies can result in improved physical activity levels for adults in community settings. Move For Life (MFL) is designed to improve and maintain physical activity levels. MFL is an intervention comprised of a set of behavioural change techniques and will be delivered by an instructor who is trained in these strategies. The instructor will then train a member of the community in the skills necessary to sustain the class in the future. This peer-mentor may then train up more members of the community in time. Before any intervention like this can come about, there must be evidence that it works and this is typically done by a trial. The aim of this study is to establish if it is feasible to conduct this type of trial and to measure the outcomes. The primary outcome is measuring time spent in daily physical activity.

Who can participate?

People aged over 45 years

What does the study involve?

Participants will be invited to attend a recruitment evening where they will be asked questions about their age and ability to participate in the physical activity programmes to ensure that they are suitable for the trial. Any participants deemed unsuitable will be asked to leave the study at this point. After participants have been screened to ensure they are suitable for the trial, they will be enrolled on the trial. All the participants in each local sports partnership hub will be allocated to the same arm of the study.

Group 1: Move For Life Intervention. Participants will be given a choice of four different physical activity programmes: Men on the Move is a 12-week structured physical activity class for men; Women on Wheels is an 8-week structured cycling class for women; Get Ireland Walking is an 8-week walking initiative for inactive men and women; Go for Life is a 8-week structured physical activity class for women and men aged 50 and over. Each programme will be run by a professional instructor who has been trained in behavioural change techniques. The instructor will train a 'peer mentor' who will be chosen from the local community to help to facilitate the group. The participants will have data collected before they commence the physical activity programme and which will run for 8-12 weeks. Repeat data will be collected at 3 and 6 months.

Arm 2: Usual care group. Participants will be enrolled in their preferred option of the four physical activity programmes described above. All classes are run by a professional instructor. The instructor will not have been trained in behavioural change techniques and will not have a 'peer-mentor' in the group. The participants will have data collected before they commence the physical activity programme and which will run for 8-12 weeks. Repeat data will be collected at 3 and 6 months.

Arm 3: True control. Participants will be given information about physical activity and will be offered entry into programmes when the trial has finished. They will have data collected after enrolment and at 3 and 6 months.

What are the possible benefits and risks of participating?

The benefits of physical activity are well documented and it is hoped that the participants in the intervention arm in particular will become more active and will maintain physical activity levels for life. There may be a risk of injury as with any form of exercise. However, all of the physical activity classes are run by a professional instructor and have a proven track record in safety. Participants will be allocated to suitable levels of physical activity at enrolment.

Participants may leave the classes or the trial at any stage.

Where is the study run from?

University of Limerick

When is the study starting and how long is it expected to run for?

March 2018 to November 2019

Who is funding the study?

Health Service Executive

Who is the main contact?

Dr Andrew O'Regan, andrew.oregan@ul.ie

Contact information

Type(s)

Scientific

Contact name

Dr Andrew O'Regan

Contact details

Graduate Entry Medical School

Faculty of Education and Health Sciences

University of Limerick

Limerick
Ireland
V94 YDE9

Additional identifiers

Protocol serial number

2018_02_15_EHS

Study information

Scientific Title

An evaluation of a peer mentoring intervention designed to help inactive 45+ become more active: a cluster randomised feasibility trial of the Move for Life Programme.

Acronym

Move for Life

Study objectives

As this is a feasibility study we do not have a null hypothesis. We are testing the feasibility of measuring an improvement in moderate to vigorous physical activity levels in adults.

Ethics approval required

Old ethics approval format

Ethics approval(s)

University of Limerick Faculty of Education and Health Sciences Research Ethics Committee, 09 /04/2018, 2018_02_15_EHS

Study design

Cluster randomized controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Feasibility of increasing physical activity in people aged over 45 years

Interventions

Move for Life (MFL) is a feasibility cluster randomised control trial (RCT) where the Local Sports Partnership (LSP) hubs are the units of randomisation (the clusters), and individuals within the hubs are the units of analysis (the participants). The advantage of a cluster RCT is that it overcomes contamination problems that arise if simple random allocation is used. When testing a complex intervention, randomisation of participants risks contamination if individuals from both arms attend the one group. This study will enrol a total of eight hubs and randomise physical activity groups to each of three arms. This framework is in line with the recommendations of Eldridge et al. (Eldridge et al. 2016a) and the CONSORT (Eldridge et al.

2016b) guidance on the minimum number of clusters required to obtain accurate estimates of rates and proportions in pilot and definitive cluster RCTs respectively. The first arm is the intervention (three hubs); the second is usual exercise classes (three hubs) and the third is a no-physical activity control (two hubs). It is a complex intervention focussed on behavioural change that will use the cascade training model to augment existing programmes and will involve the professional instructor training up one (or more) of the group participants to be a peer-mentor, using an evidence-based educational toolkit. Assessments will be completed at baseline, 3 and 6 months follow up and include a physical health battery, completion of a MFL questionnaire and the monitoring of physical activity and sedentary behaviour over a 7-day period using an ActivPAL accelerometer.

After participants have been screened to ensure they are suitable for the trial, they will be enrolled on the trial. All the participants in each local sports partnership hub will be allocated to the same arm of the study. The study has three arms as described below:

Arm 1: Move For Life (MFL) Intervention.

Participants will be given a choice of four physical activity programmes: Men on the Move is a 12-week structured physical activity class for men; Women on Wheels is an 8-week structured cycling class for women; Get Ireland Walking is an 8-week walking initiative for inactive men and women; Go for Life is a 8-week structured physical activity class for women and men aged 50 years and over. Each programme will be run by a professional instructor who has been trained in behavioural change techniques. The instructor will train a 'peer mentor' who will be chosen from the local community to help to facilitate the group. The participants will have data collected before they commence the physical activity programme and which will run for 8 to 12 weeks. Repeat data will be collected at 3 and 6 months.

Arm 2: Usual care group.

Participants will be enrolled in their preferred option of the four physical activity programmes described above. All classes are run by a professional instructor. The instructor will not have been trained in behavioural change techniques and will not have a 'peer-mentor' in the group. The participants will have data collected before they commence the physical activity programme and which will run for 8 to 12 weeks. Repeat data will be collected at 3 and 6 months.

Arm 3: True control.

Participants will be given information about physical activity and will be offered entry into programmes when the trial has finished. They will have data collected after enrolment and at 3 and 6 months.

Intervention Type

Behavioural

Primary outcome(s)

Time spent in daily moderate to vigorous physical activity. This will be measured by a sophisticated accelerometer that will be worn by the patients for 7 days at baseline, 3 and 6 months. The accelerometer data will then be uploaded by the research team for analysis.

Key secondary outcome(s)

1. Participant recruitment outcomes include demographic profile and the success of each of the recruitment strategies, based on the number, profile and reasons of participants who drop out, which will be recorded by the data collection team.
2. Measures relating to attendance at classes, retention of participants, characteristics of drop outs and refusals. Data will be collected on attendance, and safety and adverse events will be recorded by the LSP tutor.
3. Physical health assessments, including body mass index (BMI) and waist circumference.

Validated tests of functional ability including balance and strength testing will be carried as well as height, weight and grip strength.

4. Participant experience. Participants will be asked to complete a MFL questionnaire comprising of four modules: behavioural change techniques, well-being, cost effectiveness, and process evaluation. After the trial a subset of participants will be asked to undertake interviews to investigate their experience. NVIVO version 11, a qualitative research software package, will be used to assist analysis of the data with thematic analysis (Braun & Clarke, 2006) used to analyse the findings.

5. Description of how concealment of allocation from participants, data collectors and data analysis team was achieved

6. Feasibility of measuring number of minutes of moderate to vigorous physical activity

Completion date

30/11/2019

Eligibility

Key inclusion criteria

Adults aged 45 years and over

Participant type(s)

All

Healthy volunteers allowed

No

Age group

Mixed

Lower age limit

45 years

Upper age limit

100 years

Sex

All

Total final enrolment

733

Key exclusion criteria

Aged under 45 years

Date of first enrolment

01/04/2018

Date of final enrolment

30/11/2018

Locations

Countries of recruitment

Ireland

Study participating centre

University of Limerick

Castletroy

Limerick

Ireland

V94 YDE9

Sponsor information

Organisation

University of Limerick

ROR

<https://ror.org/00a0n9e72>

Funder(s)

Funder type

Government

Funder Name

Health Service Executive - Healthy and Positive Ageing for All (HaPAL)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article		15/05/2024	30/05/2024	Yes	No
Protocol article		09/07/2019	25/07/2019	Yes	No

<u>Other publications</u>		18/03/2025	09/06/2025	Yes	No
<u>Other publications</u>	Process evaluation with cost analysis	05/12/2025	06/01/2026	Yes	No
<u>Study website</u>		11/11/2025	11/11/2025	No	Yes