

# Falls prevention in the visually impaired

<b>Submission date</b> 20/05/2015	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
<b>Registration date</b> 21/05/2015	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 14/06/2023	<b>Condition category</b> Other	<input type="checkbox"/> Individual participant data

## Plain English summary of protocol

### Background and study aims

Falls and fear of falling have a big impact on older people's lives. Many people stop going out altogether, or reduce their activity so that they are much less active than they used to be. This in turn has a negative impact on their social life and general well-being. Research has shown that exercise programmes can help with this, by building up a person's confidence as well as their physical strength and stamina. This can help make them less likely to fall, more likely to keep active and more likely to have a better quality of life. However, there are a lot of older people who are visually impaired or blind and cannot easily access or participate in these kinds of exercise programmes. The aim of this study is to work with a group of older people with vision problems to adapt an existing exercise programme designed to help prevent falls so that it can also be used by people who are visually impaired. To do this, a group of older, visually impaired people will work alongside health and exercise professionals to adapt the programme and agree on ways of measuring how well it works (e.g. measuring confidence, number of falls, fear of falling and general health) for those with sight loss. The main focus of this small study will be to see if older people with sight loss in Newcastle and Glasgow are prepared to join in and stick with the exercise programme and complete the assessments. The results of this study will be used to see whether a larger study could be carried out.

### Who can participate?

Adults aged 60 and over with visual impairment.

### What does the study involve?

Participants are randomly allocated into one of two groups. Those in group 1 (intervention group) take part in a 12-week exercise programme. Those in group 2 (control group) are given standard primary health care advice. Measurements, including confidence, fear of falling, number of falls etc. are taken at the start of the trial, at the end of the exercise programme, and again 6 months later.

### What are the possible benefits and risks of participating?

Not provided at time of registration.

### Where is the study run from?

University of Newcastle upon Tyne (UK)

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

May 2015 to May 2016

Who is funding the study?

National Institute for Health Research (NIHR) - Public Health Research (UK)

Who is the main contact?

Ms R Lampitt

## Contact information

### Type(s)

Scientific

### Contact name

Ms Rosy Lampitt

### Contact details

University of Newcastle upon Tyne  
Neurosurgical Trials Unit  
3-4 Claremont Terrace  
Newcastle University  
Newcastle upon Tyne  
United Kingdom  
NE2 4AE

## Additional identifiers

### Protocol serial number

18789

## Study information

### Scientific Title

Visually Impaired OLder people's Exercise programme for falls prevention (VIOLET): a feasibility study

### Acronym

VIOLET

### Study objectives

This study will assess the feasibility and acceptability of, and adherence to, an exercise programme aimed to increase fitness in older people with sight loss. The exercise programme aims to decrease physical injury from falls and increase confidence in participants.

### Ethics approval required

Old ethics approval format

### Ethics approval(s)

Ref: 15/NE/0057

**Study design**

Randomised interventional study

**Primary study design**

Interventional

**Study type(s)**

Quality of life

**Health condition(s) or problem(s) studied**

Fitness of visually impaired older people

**Interventions**

1. Treatment group takes part in an adapted exercise programme designed for older, visually impaired people for 12 weeks
2. Control group receives standard care

**Intervention Type**

Behavioural

**Primary outcome(s)**

Fear of falling measured using Short Falls Efficacy Scale – International (FES-I) at baseline, 12 weeks and 24 weeks.

**Key secondary outcome(s)**

Not available at time of registration

**Completion date**

23/05/2016

## Eligibility

**Key inclusion criteria**

1. Age >60 years
2. Attend a low vision clinic and/or are members of organisations for the visually impaired such as NSBP in Newcastle or Visibility in Glasgow
3. Live in their own home
4. Can walk indoors without the help of another person but may use a walking aid such as a stick
5. Can walk outdoors but may need the help of another and/or walking aid
6. Physically able to take part in a group exercise class
7. Participant has given informed consent (as appropriate to each older person with visual impairment) to participate in the study prior to any study-specific procedures

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Sex**

All

**Key exclusion criteria**

1. Unable to comprehend or follow simple movement instructions in English (to an extent of being unable to follow simple movement instructions)
2. Acute or uncontrolled medical problems which the participant's General Practitioner considers would exclude them from undertaking the exercise programme (e.g. uncontrolled heart disease, poorly controlled diabetes, acute systemic illness, neurological problems, severe chronic obstructive pulmonary disease (COPD)) in addition to visual impairment
3. Conditions requiring a specialist exercise programme e.g. uncontrolled epilepsy, severe neurological disease or impairment, unable to maintain a seated upright position or unable to move independently indoors
4. Current involvement in other falls prevention exercise programmes (but not excluding walking programmes), investigational studies or trials

**Date of first enrolment**

01/05/2015

**Date of final enrolment**

30/04/2016

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

**University of Newcastle upon Tyne**

Neurosurgical Trials Unit

3-4 Claremont Terrace

Newcastle University

Newcastle upon Tyne

United Kingdom

NE2 4AE

**Sponsor information****Organisation**

University of Northumbria at Newcastle

**ROR**

<https://ror.org/049e6bc10>

## Funder(s)

**Funder type**

Government

**Funder Name**

Public Health Research Programme

**Alternative Name(s)**

NIHR Public Health Research Programme, The Public Health Research (PHR), PHR

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

United Kingdom

## Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	results	12/12/2018		Yes	No
<a href="#">Results article</a>		01/02/2019	14/06/2023	Yes	No
<a href="#">Protocol article</a>		02/08/2016	14/06/2023	Yes	No
<a href="#">HRA research summary</a>			28/06/2023	No	No
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes