

# A randomised controlled trial of the effectiveness of prism spectacles in reducing the disability of patients with age-related macular degeneration (ARMD)

<b>Submission date</b> 12/09/2002	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 12/09/2002	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 08/06/2011	<b>Condition category</b> Eye Diseases	<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

## Secondary identifying numbers

1071/530

# Study information

## Scientific Title

### Acronym

Prism Spectacles Study

### Study objectives

To study prism spectacles in reducing the disability of patients

### 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)

Treatment

## Participant information sheet

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

Age-related macular degeneration

### Interventions

1. Prism spectacles prescribed to established protocol
2. Prism spectacles with prism of standard value
3. Placebo spectacles made to appear like prism spectacles

### Intervention Type

Other

### Phase

Not Specified

**Primary outcome measure**

Not provided at time of registration

**Secondary outcome measures**

Not provided at time of registration

**Overall study start date**

01/02/2001

**Completion date**

01/02/2004

## **Eligibility**

**Key inclusion criteria**

1. Bilateral ARMD
2. Visual Acuity (VA) 6/18 to 1/60 binocularly
3. Clinically stable (by visual acuity) with bilateral central scotoma
4. Reporting difficulty with everyday tasks

**Participant type(s)**

Patient

**Age group**

Adult

**Lower age limit**

18 Years

**Sex**

Both

**Target number of participants**

225

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

01/02/2001

**Date of final enrolment**

01/02/2004

## **Locations**

**Countries of recruitment**

England

United Kingdom

**Study participating centre**

**Department of Optometry and Neuroscience**

Manchester

United Kingdom

M60 1QD

## **Sponsor information**

**Organisation**

The Health Foundation (UK)

**Sponsor details**

90 Long Acre

London

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**Sponsor type**

Charity

**Website**

<http://www.pppfoundation.org.uk/>

**ROR**

<https://ror.org/02bj4420>

## **Funder(s)**

**Funder type**

Charity

**Funder Name**

The Health Foundation (UK)

# 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?
<a href="#">Results article</a>	results	01/08/2005		Yes	No