

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

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

United Kingdom

WC2E 9RA

+44 (0)20 7257 8000

info@health.org.uk

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?
Results article	results	01/08/2005		Yes	No