A randomised controlled trial of treatment of unilateral straight eyed visual acuity deficit

Prospectively registered Submission date Recruitment status 23/01/2004 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 23/01/2004 Completed [X] Results Individual participant data **Last Edited** Condition category 09/11/2022 **Eve Diseases**

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

Protocol serial number

RRCC7R R1903/6624

Study information

Scientific Title

A randomised controlled trial of treatment of unilateral straight eyed visual acuity deficit

Study objectives

Not provided at time of registration

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

Study type(s)

Not Specified

Health condition(s) or problem(s) studied

Eve diseases

Interventions

The effects of correction of refractive error, alone and in combination with patching will be compared to no treatment.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

The main outcome measure at both the preliminary and final analysis will be LogMAR acuity. Anova will be used to test the effect of group and sub-group analyses will test for heterogeneity between and at different initial acuities.

Key secondary outcome(s))

Not provided at time of registration

Completion date

28/02/2002

Eligibility

Key inclusion criteria

Children aged 3-4 years, with lateral visual acuity defects of 6/9 to 6/24.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Child

Lower age limit

3 years

Upper age limit

4 years

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

28/02/1999

Date of final enrolment

28/02/2002

Locations

Countries of recruitment

United Kingdom

England

Study participating centre Royal Victoria Infirmary

Newcastle upon Tyne United Kingdom NE1 4LP

Sponsor information

Organisation

NHS R&D Regional Programme Register - Department of Health (UK)

Funder(s)

Funder type

Government

Funder Name

NHS Executive Northern and Yorkshire (UK)

Results and Publications

Individual participant data (IPD) sharing plan

Not provided at time of registration

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		01/01/2005		Yes	No
Other publications		29/11/2003		Yes	No
Other publications		01/08/2004		Yes	No
Other publications		01/07/2006		Yes	No