

Eye-movement training improves reading function in children and young people with eye-movement disorders after acquired brain injury

Submission date
17/09/2024

Recruitment status
No longer recruiting

Prospectively registered

Protocol

Registration date
22/09/2024

Overall study status
Completed

Statistical analysis plan

Results

Last Edited
08/10/2024

Condition category
Injury, Occupational Diseases, Poisoning

Individual participant data

Plain English summary of protocol

Background and study aims

Acquired brain injury (ABI) often causes a worsening of reading function. This may decrease motivation for school activities. Eye-movement training has been shown to have positive effects, but its feasibility and effectiveness are unknown, as is its optimal duration, frequency and intensity. The aim was to develop and evaluate eye-movement training for children and young people with ABI.

Who can participate?

Children and young people aged between 10-19 years old with eye-movement disorders after ABI

What does the study involve?

The intervention involves 20 minutes of training each day for three weeks. Each participant and caregiver will be introduced to the home training program and will receive individual support by phone calls and/or text messages during the intervention. Vision assessments, a reading speed test and a survey will record results at baseline, at the end of the training, and three months after the end of the training.

What are the possible benefits and risks of participating?

The participants might benefit regarding their reading function, and possible risks including headaches from the training are expected to be minimal.

Where is the study run from?

The study is run from the Child and Youth Rehabilitation Center in Lund, Sweden.

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

January 2020 and February 2024

Who is funding the study?

Region Skåne (Sweden)

Who is the main contact?

Dr. Katarina Lauruschkus, katarina.lauruschkus@med.lu.se

Contact information

Type(s)

Public, Scientific, Principal Investigator

Contact name

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

Nil known

Study information

Scientific Title

Eye-movement training improves reading function in children and young people with eye-movement disorders after acquired brain injury

Study objectives

Three weeks of eye-movement training is feasible and improves the reading function in children and young people with eye-movement disorders after acquired brain injury.

Ethics approval required

Ethics approval required

Ethics approval(s)

Approved 20/11/2020, Swedish Ethical Review Authority (Box 2110, Uppsala, SE-750 02, Sweden; +46 10-475 08 00; registrator@etikprovning.se), ref: EPM-Dnr 2020-04154

Study design

Single-centre interventional study

Primary study design

Interventional

Secondary study design

Non randomised study

Study setting(s)

Care home, Hospital, Internet/virtual, Telephone

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet.

Health condition(s) or problem(s) studied

Eye-movement disorders after acquired brain injury

Interventions

The materials for the eye-movement training with instructions were provided for each participant by the occupational therapist or the special education teacher. The materials are described and visualized in the manuscript that will be submitted and is available upon request.

The intervention involves 20 minutes of training each day for three weeks. Each participant and caregiver is introduced to the home training program and receives individual support during the intervention. Vision assessments, a reading speed test and a survey records results at baseline, at the end of the training, and three months after the end of the training.

The paediatrician, the occupational therapist and the special education teacher all worked at the Child and Youth Rehabilitation Center and were trained for the intervention. They were all well experienced for working with the target group.

Intervention Type

Behavioural

Primary outcome measure

Reading speed measured using a reading speed test at baseline, at the end of the training, and three months after the end of the training

Secondary outcome measures

Clinical signs of convergence insufficiency measured using vision assessments and the modified Convergence Insufficiency Symptom Survey (CISS) at baseline, at the end of the training, and three months after the end of the training

Overall study start date

25/01/2020

Completion date

20/02/2024

Eligibility

Key inclusion criteria

1. Aged 10-19 years old
2. Eye-movement disorders after ABI
3. Live in the Skåne Region in Southern Sweden
4. Being followed up by the Child and Youth Rehabilitation Center in Lund, Sweden
5. Able to read independently before the injury

Participant type(s)

Patient

Age group

Mixed

Lower age limit

10 Years

Upper age limit

19 Years

Sex

Both

Target number of participants

17

Total final enrolment

16

Key exclusion criteria

Not meeting the participant inclusion criteria

Date of first enrolment

20/11/2020

Date of final enrolment

12/11/2023

Locations

Countries of recruitment

Sweden

Study participating centre
Child and Youth Rehabilitation Center
Lovisastigen 9
Lund
Sweden
22100

Sponsor information

Organisation

Region Skåne

Sponsor details

Department of Habilitation, Committee on Psychiatry, Habilitation and Technical Aids,
Dockplatsen 26
Malmö
Sweden
211 74
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Sponsor type

Government

Website

<https://www.skane.se/>

Funder(s)

Funder type

Government

Funder Name

Region Skåne

Alternative Name(s)

Funding Body Type

Government organisation

Funding Body Subtype

Local government

Location

Sweden

Results and Publications

Publication and dissemination plan

Planned publication in a peer-reviewed journal

Intention to publish date

01/03/2025

Individual participant data (IPD) sharing plan

The datasets generated and analyzed during the current study will be available upon request from Katarina Lauruschkus, katarina.lauruschkus@med.lu.se

IPD sharing plan summary

Available on request

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Poster results		06/09/2022	20/09/2024	No	No