

# The emotional impact of restoring sight

<b>Submission date</b>	<b>Recruitment status</b>	<input checked="" type="checkbox"/> Prospectively registered
14/03/2025	Recruiting	<input type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
27/03/2025	Ongoing	<input type="checkbox"/> Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
26/01/2026	Eye Diseases	<input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

More treatment options than ever before are available for young people with vision impairment, but very little is known about the impact of these sight restoration therapies on wellbeing and mental health. In adults, sight restoration is often associated with depression and a reduction in mental wellbeing.

The aim of this study is to fully understand the wellbeing and mental health of adolescents undergoing sight restoration therapy and will develop clinical guidelines for the psychological support of young people receiving this treatment.

### Who can participate?

Children and adolescents (age 10-20 years) with vision impairment who are receiving treatment expected to improve their vision.

### What does the study involve?

Completing age-appropriate questionnaires five times: before receiving sight restoration treatment, then 1, 3, 6 and 12 months after treatment. Participating in a short interview before treatment and at the final visit. Participants' parent(s)/carer(s) will also complete a questionnaire at each assessment and a short interview at the first and last assessment.

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

**Benefits:** This is an exploratory study and is not likely to directly benefit participants. However, if needed, participants will be referred or signposted to appropriate mental health services.

**Risks:** It is possible that some of the interview questions may raise uncomfortable issues for participants or their parents/carers. The researchers have well-established risk management protocols, written in conjunction with clinical psychologists, which have been used for similar studies to ensure appropriate signposting and referral for participants who express signs of mental ill-health, up to and including emergency care.

### Where is the study run from?

UCL Institute of Ophthalmology, part of University College London, in association with Moorfields Eye Hospital, Great Ormond Street Hospital for Children, and the NIHR Moorfields Biomedical Research Centre (UK)

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

January 2025 to December 2027

Who is funding the study?

1. The Medical Research Foundation (UK)
2. Moorfields Eye Charity (UK)

Who is the main contact?

Michael Crossland, m.crossland@ucl.ac.uk

## Contact information

### Type(s)

Public, Scientific, Principal investigator

### Contact name

Dr Michael Crossland

### ORCID ID

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### Contact details

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## Additional identifiers

### Integrated Research Application System (IRAS)

356506

## Study information

### Scientific Title

The emotional impact of restoring sight (EIRS): well-being and mental health in children and adolescents receiving sight restoring therapy for eye disease

### Acronym

EIRS

### Study objectives

This study will use mixed qualitative and quantitative research methods to understand how sight-restoring treatment (for example, gene therapy for inherited retinal disease) affects the wellbeing and mental health of young people with vision impairment.

### Ethics approval required

Ethics approval required

**Ethics approval(s)**

approved 20/10/2025, London - Brighton & Sussex Research Ethics Committee (2 Redman Place, Stratford, London, E20 1JQ, United Kingdom; -; brightonandsussex.rec@hra.nhs.uk), ref: 25/LO /0718

**Study design**

Longitudinal observational study

**Primary study design**

Observational

**Study type(s)**

Quality of life

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

Any eye disease causing bilateral vision impairment

**Interventions**

Participants will complete age-appropriate questionnaires to assess their mental health and wellbeing five times: before receiving sight restoration treatment, and 1, 3, 6 and 12 months after treatment. Before treatment and at the end of the study, participants will also complete a short qualitative interview.

Participants' parent(s)/carer(s) will also complete a questionnaire at each assessment and a short interview at the first and last assessment.

**Intervention Type**

Not Specified

**Primary outcome(s)**

Mental wellbeing assessed using the Warwick Edinburgh Mental Wellbeing Scale at pre-treatment, months 1, 3, 6 and 12

**Key secondary outcome(s)**

1. Participation assessed using the Child and Adolescent Scale of Participation at pre-treatment, months 1, 3, 6 and 12
2. Depression and anxiety assessed using the Revised Children's Anxiety and Depression Scale at pre-treatment, months 1, 3, 6 and 12
3. Sleep assessed by parental completion of the Children's Sleep Habits Questionnaire at pre-treatment, months 1, 3, 6 and 12
4. Qualitative findings from participant and parent interviews at pre-treatment and month 12

**Completion date**

31/12/2027

**Eligibility**

**Key inclusion criteria**

1. Young people aged 10-20 years
2. Meeting the ICD-11 criterion of having mild, moderate or severe vision impairment, or blindness, in their better eye
3. Receiving treatment intended to improve their visual function
4. With sufficient English comprehension to understand the age-appropriate questionnaires

**Participant type(s)**

Carer, Patient

**Healthy volunteers allowed**

No

**Age group**

Mixed

**Lower age limit**

10 years

**Upper age limit**

20 years

**Sex**

All

**Total final enrolment**

0

**Key exclusion criteria**

Does not meet the inclusion criteria

**Date of first enrolment**

08/01/2026

**Date of final enrolment**

31/12/2026

## Locations

**Countries of recruitment**

United Kingdom

England

**Study participating centre**

**NIHR Moorfields Biomedical Research Centre**  
Moorfields Eye Hospital NHS Foundation Trust  
162 City Road

London  
England  
EC1V 2PD

#### **Study participating centre**

**Dept. of Clinical & Academic Ophthalmology**  
Great Ormond Street Hospital  
Great Ormond Street  
London  
England  
WC1N 3JH

## **Sponsor information**

#### **Organisation**

University College London

#### **ROR**

<https://ror.org/02jx3x895>

## **Funder(s)**

#### **Funder type**

Charity

#### **Funder Name**

Medical Research Foundation

#### **Alternative Name(s)**

MedResFdn, The Medical Research Foundation, MRF

#### **Funding Body Type**

Private sector organisation

#### **Funding Body Subtype**

Trusts, charities, foundations (both public and private)

#### **Location**

United Kingdom

**Funder Name**

Moorfields Eye Charity

**Alternative Name(s)****Funding Body Type**

Private sector organisation

**Funding Body Subtype**

Research institutes and centers

**Location**

United Kingdom

## Results and Publications

**Individual participant data (IPD) sharing plan**

Datasets generated and/or analysed during the current study will be made available upon request from Michael Crossland (m.crossland@ucl.ac.uk). To preserve the confidentiality of the participants, potentially identifiable data will be redacted.

**IPD sharing plan summary**

Available on request

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Participant information sheet</a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes