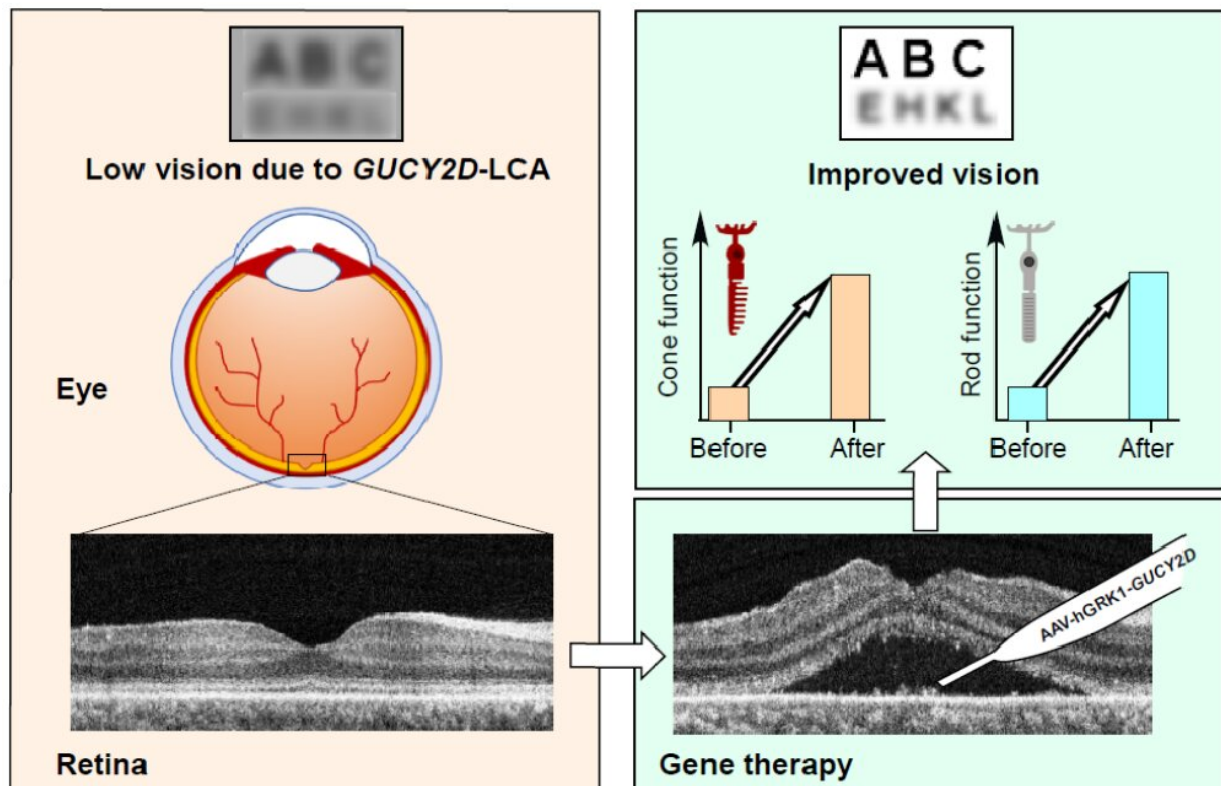


Gene therapy shows promise in initial trial for patients with childhood blindness

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Childhood visual loss in Leber congenital amaurosis due to GUCY2D mutations affects both night vision and clarity, but the eyes' photoreceptors remain relatively intact. An injection of gene therapy led to improved night (rod) and day (cone) vision. Credit: Alexander Sumaroka and Alexandra V. Garafalo

A new gene therapy for one of the most common forms of congenital

blindness was safe and improved patients' vision, according to initial data from a clinical trial led by researchers at the Scheie Eye Institute in the Perelman School of Medicine at the University of Pennsylvania.

The [therapy](#) delivers working copies of GUCY2D to the eyes of patients who have severe [vision](#) impairments caused by mutations in the gene. Each of the first three treated patients experienced improvement in some aspects of vision, without [serious side effects](#), according to the new study, published in the journal *iScience*.

"We found sustained improvements in both day and night vision, even with a relatively low dose of the [gene therapy](#)," said study lead author Samuel G. Jacobson, MD, Ph.D., a professor of ophthalmology in the Perelman School of Medicine.

The GUCY2D gene is one of about 25 different human [genes](#) whose mutations cause problems in the retina, leading to severe vision impairment from birth or early childhood. This family of inherited retinal disorders, collectively known as Leber congenital amaurosis (LCA), accounts for a considerable portion of blindness in children worldwide.

Normal copies of GUCY2D encode an enzyme in the key pathway that light-sensitive rod and cone [cells](#) in the retina use to convert light into electrochemical signals. A lack of this enzyme blocks the recovery of this pathway, preventing the reset needed for further signaling. As a result, the signal from rod and cone cells becomes very weak—which equates to severe vision loss.

Even in adults who have lived for decades with this condition, it is often the case that many light-sensing retinal cells remain alive and intact despite their dysfunction. Thus, adding functional copies of GUCY2D via a gene therapy could get those cells working again and restore some

vision.

In 2019, Jacobson and co-investigator Artur V. Cideciyan, Ph.D., a research professor of ophthalmology in the Perelman School of Medicine, began the first clinical trial of a GUCY2D gene therapy, a solution of a harmless virus that carries the gene and is injected beneath the retina—initially in just one eye per patient. They are following each patient for two years after treatment. In the new report, they described their findings after nine months in the first three patients treated.

The first patient experienced a substantial increase in light-sensitivity in rod cells, which are more light-sensitive than cone cells and are chiefly responsible for low-light or "night vision." This patient also showed improved pupil responses to light.

The second patient showed a smaller but sustained increase in light-sensitivity in rod cells, starting about two months after the gene therapy.

The third patient showed no improvement in rod cell sensitivity, but did show significantly improved [visual acuity](#) over the nine-month follow-up period, an improvement that the researchers tied to better function in the patient's [cone cells](#), the predominant cells for daylight and color vision.

"These initial results from the first-ever trial of a GUCY2D gene therapy are very encouraging and will inform our ongoing and future trials of this therapy," said Cideciyan.

There were no serious adverse side effects, and any side effects that occurred in the patients' retinas resolved.

The gene therapy dose used in these first three patients was the lowest of the doses the researchers plan to use in the study, so they are hoping to see continued safety and greater efficacy in later-enrolled patients who

will receive higher doses.

The ongoing clinical trial is registered at clinicaltrials.gov as trial NCT03920007.

Jacobson and Cideciyan in earlier studies have reported success for gene-based therapies against two other forms of Leber congenital amaurosis involving other genes.

More information: Samuel G. Jacobson et al. Safety and Improved Efficacy Signals following Gene Therapy in Childhood Blindness Caused by GUCY2D Mutations, *iScience* (2021). [DOI: 10.1016/j.isci.2021.102409](https://doi.org/10.1016/j.isci.2021.102409)

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