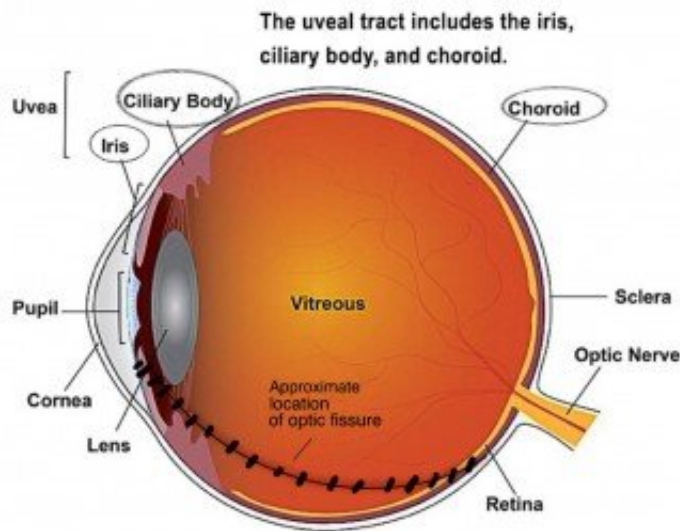


Zebrafish model helps explain eye development

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Uveal coloboma refers to missing parts of the uveal tract—the middle layer of eye tissue that includes the iris, the ciliary body, and the blood vessel-rich choroid. Marked is the approximate location of the optic fissure, which normally closes during embryonic development. Failure of the fissure to close anywhere along this line results in a coloboma. A coloboma's impact on vision varies according to its location and severity. It can cause anything from slight blurriness to blindness. Credit: National Eye Institute

Scientists at the National Eye Institute (NEI) have developed a zebrafish

model of NEDBEH—a rare genetic disorder that can cause coloboma, where parts of the eye are missing due to developmental defects. The model provides a new tool for understanding the eye's embryonic development. The NEI is part of the National Institutes of Health.

"Despite advances in genetics, the underlying cause of coloboma in most families remains unclear," said Brian Brooks, M.D., Ph.D., chief of the NEI Ophthalmic Genetics and Visual Function Branch. The ongoing mystery is partly due to the myriad genes that must be turned on and off at pivotal moments for the eye's normal development.

Coloboma results from a failure of a transient structure called the optic fissure to close as the eye forms. To better understand coloboma, Brooks and NEI Scientists Aman George, Ph.D., explored NEDBEH-associated gene RERE. NEDBEH is short for [neurodevelopmental disorder](#) with or without anomalies of the brain, eye, and heart. The RERE gene produces a scaffold-like protein (*rere*) that enables assembly of transcriptional regulators—proteins that activate or deactivate [gene expression](#).

Brooks and George characterized early eye development in zebrafish with a mutation in the gene *rerea*—the zebrafish equivalent of the human RERE gene. Compared with normal zebrafish, mutants showed eye defects including enlarged optic stalks, eye tissue in the brain, and coloboma. The optic stalk is a transient eye structure that connects the eye to the brain.

The *rerea* mutation altered the expression of key developmental genes. Of note, the *rerea* mutation interfered with signaling of sonic hedgehog (*shh*), an important protein that directs the development of a variety of organs and tissue, including the optic stalk and retina. By inhibiting the [shh pathway](#) at a different point, the scientists rescued coloboma, leading to proper closure of the optic fissure.

"Our [zebrafish model](#) helps clarify the role of the RERE gene in eye development," said George. "Identifying appropriate model systems for studying coloboma is essential to understanding its underlying mechanisms and finding targets for rescuing defects."

The research is published in the journal *Developmental Dynamics*.

More information: Aman George et al, Zebrafish model of RERE syndrome recapitulates key ophthalmic defects that are rescued by small molecule inhibitor of shh signaling, *Developmental Dynamics* (2022). [DOI: 10.1002/dvdy.561](https://doi.org/10.1002/dvdy.561)

Provided by National Eye Institute

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