

# Genetic basis for glaucoma uncovered

January 30 2018, by Will Doss

---



Acute angle closure glaucoma of the right eye (intraocular pressure was 42 in the right eye). Credit: James Heilman, MD/Wikipedia

In two recent publications, Northwestern Medicine scientists and international collaborators discovered mutations that cause improper drainage and a buildup of ocular pressure leading to one form of congenital glaucoma, and identified a path towards future treatments for the disease.

Susan Quaggin, MD, chief of Nephrology and Hypertension in the Department of Medicine and director of the Feinberg Cardiovascular Research Institute, was senior author on the studies, the first published in the *Journal of Clinical Investigation* (JCI) and the second published in the *Proceedings of the National Academy of Sciences* of the United States of America (PNAS).

Glaucoma is a leading cause of blindness around the globe, and elevated intraocular pressure (IOP) is an important risk factor for the disease. Developmental defects in the anterior chamber of the eye, including a drainage vessel called Schlemm's canal, can lead to a particularly severe form of glaucoma in children known as primary [congenital glaucoma](#) (PCG).

"This special vessel acts like the drainpipes in a bathtub—to prevent the eye from 'overflowing,' " said Quaggin, also the Charles H. Mayo, MD, Professor and a professor of Medicine in the Division of Nephrology and Hypertension.

Previous studies from Quaggin and her collaborators have shown loss-of-function mutations in the angiopoietin (ANGPT) receptor TIE2/TEK in families with PCG, and that ANGPT/TIE2 pathway activity is critical for Schlemm's canal development.

## **Uncovering the genetic roots of congenital glaucoma**

In the JCI study, Quaggin and her colleagues used mice models to explore the importance of individual components of the ANGPT/TEK pathway, finding mice without the growth factor ANGPT1 had severely deformed and small Schelmm's canals. In addition, loss of TIE2/TEK, the angiopoietin receptor, had a similar effect.

"Both ANGPT1 and TIE2/TEK are essential to form the drainage system of the eye to regulate [intraocular pressure](#) and prevent Glaucoma," Quaggin said. "We believe it's highly likely that this pathway is also important in common forms of elevated IOP glaucoma, with over 60 million individuals affected by this disease worldwide."

Investigators found two human subjects with loss-of-function mutations in ANGPT1 within an international group of PCG patients, further

supporting a causative role for ANGPT1 in the disease, according to Quaggin.

"These discoveries highlight the central role of this molecular pathway in glaucoma and provide sign posts guiding us to a new therapy," Quaggin said.

## Flipping the Switch

In the *PNAS* study, Quaggin and her colleagues searched for ways to influence the molecular pathway they'd identified. In particular, inhibition of TIE2/TEK has been linked to vessel leakage and inflammation, so finding a way to activate TIE2/TEK was a priority.

"To develop a new and much needed therapy to 'fix' these vessels and lower the pressure inside the eye to prevent or treat [glaucoma](#), we need to understand how to turn the molecules 'on' and 'off,' " Quaggin said.

Again using mouse models, investigators found inhibition of a protein called VEPTP allowed ANGPT2 to be used as a TIE2/TEK activator, providing a blueprint for a pharmacological solution, according to Quaggin.

"We are currently well underway to developing new small molecule drugs to improve drainage through the Schlemm's canal and associated outflow vessels," Quaggin said. "We are also screening other patients and looking for other mutations in the pathway."

**More information:** Benjamin R. Thomson et al. Angiopoietin-1 is required for Schlemm's canal development in mice and humans, *Journal of Clinical Investigation* (2017). [DOI: 10.1172/JCI95545](https://doi.org/10.1172/JCI95545)

Tomokazu Souma et al. Context-dependent functions of angiopoietin 2

are determined by the endothelial phosphatase VEPTP, *Proceedings of the National Academy of Sciences* (2018). [DOI: 10.1073/pnas.1714446115](https://doi.org/10.1073/pnas.1714446115)

Provided by Northwestern University

Citation: Genetic basis for glaucoma uncovered (2018, January 30) retrieved 21 November 2023 from <https://medicalxpress.com/news/2018-01-genetic-basis-glaucoma-uncovered.html>

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.