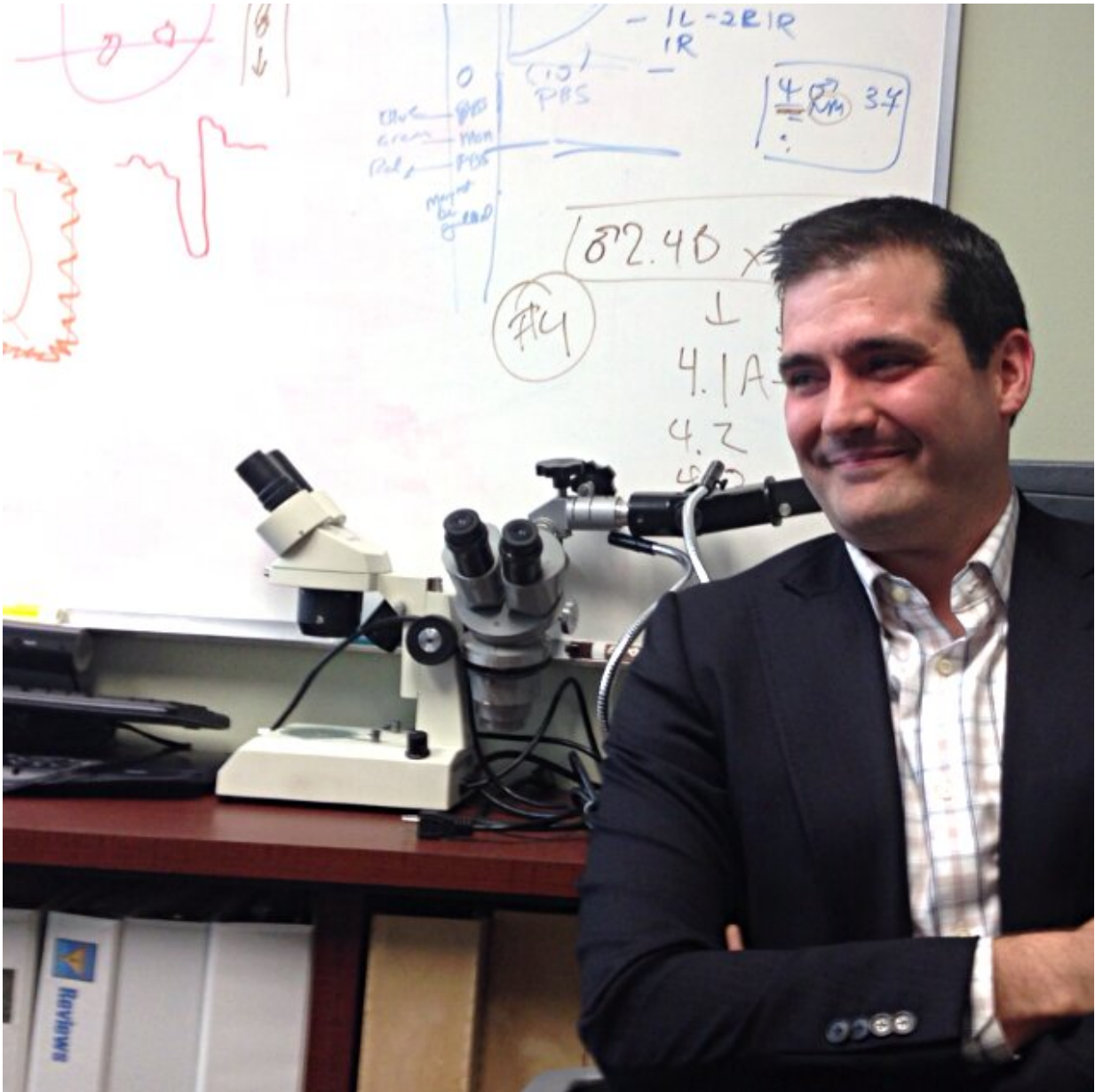


Researchers uncover mechanism blocking retina regeneration

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Dr. Ross A. Poche. Credit: Baylor College of Medicine

A discovery opens the possibility of one day restoring loss of vision by activating the retina's ability to regenerate. Researchers at Baylor College of Medicine, the Cardiovascular Research Institute and the Texas Heart Institute reveal in the journal *Cell Reports* that although the mammalian retina—a layer of specialized nerve cells that mediates vision and is located on the back of the eye- does not spontaneously regenerate, it has a regenerative capacity that is kept dormant by a cellular mechanism called the Hippo pathway. The discovery opens the possibility of activating the retina's ability to restore lost vision by manipulating this pathway.

"Damage to the [retina](#) can lead to irreparable loss of vision in humans and other mammals because their retinas do not regenerate," said lead author Dr. Ross A. Poché, assistant professor of molecular physiology and biophysics and member of the Dan L Duncan Comprehensive Cancer Center at Baylor. "However, other animals such as zebrafish can reverse blindness thanks to specialized cells in the retina called Müller [glial cells](#). When the retina is damaged, Müller glial cells proliferate and differentiate into the lost [retinal neurons](#), effectively replacing injured cells with fully functional ones."

Although Müller glial cells in injured mammalian retina do not restore vision as their counterpart in zebrafish do, other researchers have shown that, when the mammalian retina is injured, a small subset of Müller glial cells takes the first steps needed to enter the proliferation cycle, such as acquiring molecular markers scientists expect to see in a proliferating cell.

"But this attempt to proliferate is transient; after acquiring some of the

cell markers the cells shut off," said Poché, who also is affiliated with the Intellectual and Developmental Disabilities Research Center and the Cullen Eye Institute, both at Baylor. "These observations suggested that the mechanism that drives cell repair in zebrafish also might be present in mammals, but it is actively suppressed. For years, the suppressing mechanism was unknown."

Searching for the proposed suppressing mechanism, the Poché lab combined forces with the lab of co-corresponding author Dr. James Martin, professor of molecular physiology and biophysics, Vivian L. Smith Chair in Regenerative Medicine at Baylor College of Medicine and director of the Cardiomyocyte Renewal Lab at the Texas Heart Institute. The researchers focused their attention on the Hippo [pathway](#), a network of molecular events that contributes to organ growth during development and to the regulation of heart tissue regeneration in response to myocardial infarction. The Martin lab previously showed that the Hippo pathway acts like a 'break' on cardiomyocyte proliferation by inhibiting the activity of another pathway called YAP.

In this study, the researchers first determined that the Hippo pathway is expressed in mammalian Müller glial cells. Then, they investigated whether altering the Hippo pathway in these cells would affect their ability to proliferate. Creating a malfunctioning Hippo pathway by eliminating two of its molecular steps resulted in modest cell proliferation. And when the researchers genetically engineered Müller glial cells to carry a version of YAP called YAP5SA that is impervious to the inhibitory influence of Hippo, the cells showed major proliferation and acquired a progenitor cell identity. Importantly, a small subset of these Müller glia-derived progenitor cells showed signs of spontaneous differentiation into new retinal neurons.

"Up to this point researchers did not know what endogenous blocking mechanism prevented Müller cells from entering a regenerative state.

The Hippo pathway is a new molecular entry point to that mechanism," said Poché. "Our next step is to develop a strategy to guide proliferating Müller glial cells into differentiation pathways leading to retinal [cells](#) capable of restoring vision."

More information: *Cell Reports* (2019). [DOI: 10.1016/j.celrep.2019.04.047](#)

Provided by Baylor College of Medicine

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