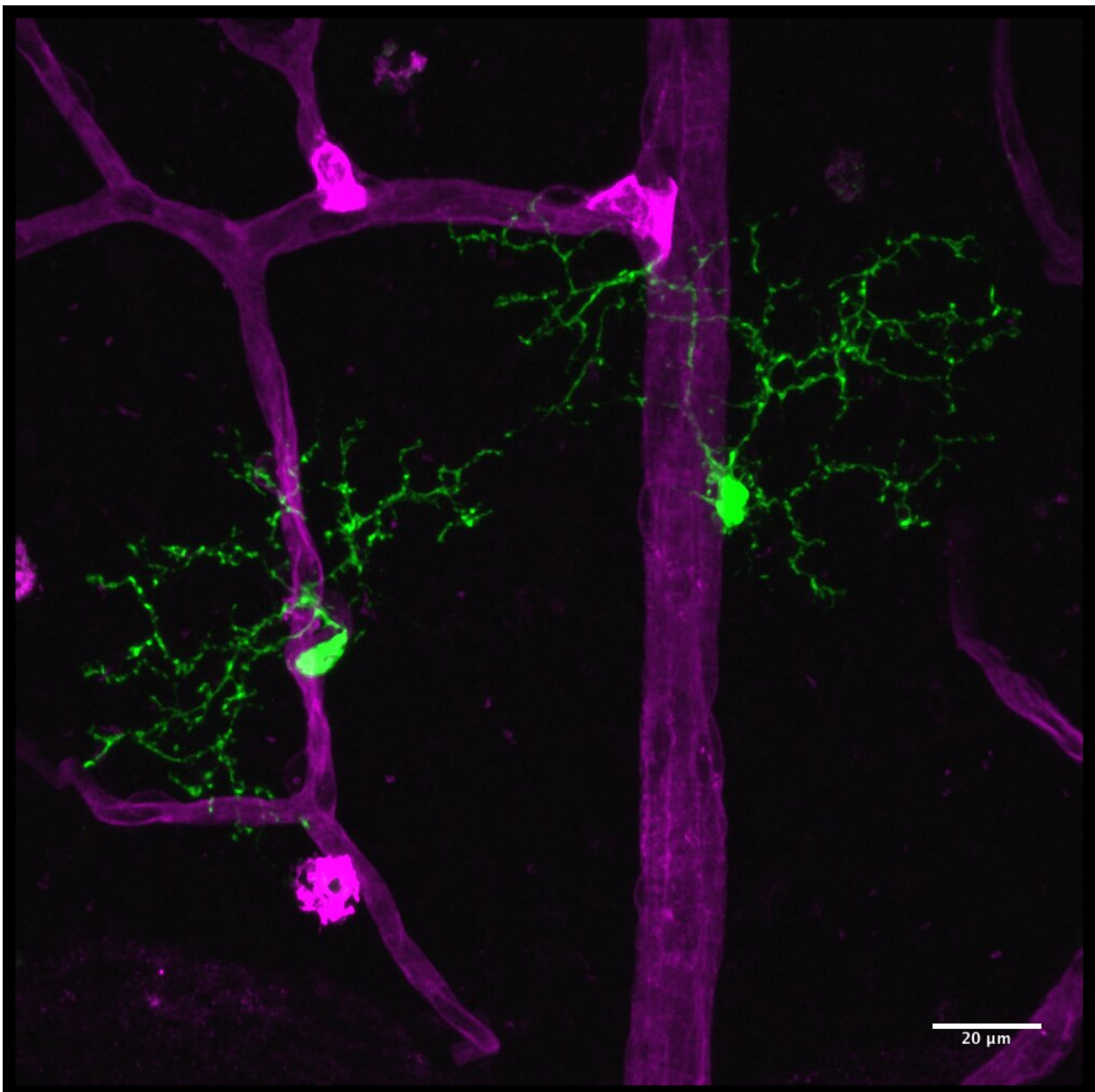


# Microglia, immune cells of the central nervous system, shown to regulate neuroinflammation

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Retinal microglia (green), the resident immune cell of the central nervous system, and the retinal vasculature (magenta), in a retinal flat mount. Credit: Connor Laboratory - Dong Ho Park, M.D.

A research team at Massachusetts Eye and Ear has shown that microglia, the primary immune cells of the central nervous system—including the retina of the eye—serve as "gatekeepers," or biosensors and facilitators, of neuroinflammation in a preclinical model of autoimmune uveitis. Uveitis is one of the leading causes of blindness, accounting for approximately 10% of significant visual impairment worldwide.

In a report published online today in *Proceedings of the National Academy of Sciences (PNAS)*, the researchers describe, for the first time, a role for microglia in directing the initiation of autoimmune uveitis by orchestrating the [inflammatory response](#) within the retina. In reaction to disease induction, microglia closely associate with the retinal vasculature and facilitate inflammatory immune cell entry past the blood brain, or ocular, barrier into the retina. When the researchers depleted microglia in this model, they observed that the disease was completely blocked.

"Normally, the blood brain barrier serves as an impediment and prevents the [immune response](#) from going into tissues of the central nervous system, including the retina. However, our results provide [clear evidence](#), that in the context of uveitis, microglia can facilitate entry of inflammatory immune cells into the retina, and enable the host immune responses to attack cells that are not normally recognized by the immune system," said senior author Kip M. Connor, Ph.D., vision researcher at Mass. Eye and Ear and Associate Professor of Ophthalmology at Harvard Medical School. "Until now, the role of microglia in retinal

disease has not been fully understood, but our research shows—for the first time—that these cells serve as gatekeepers from the immune system to the central nervous system. This gateway not only has implications for treating uveitis, but may provide future avenues for drug delivery across the blood brain barrier for other diseases of the central nervous system."

Despite significant advances in research and therapeutics, the prevalence of uveitis has not been reduced in the past 30 years. Uveitis is characterized as inflammation of the retina as well as the uveal tissues, optic nerve and vitreous, wherein a large influx of immune cells into the eye coincides with elevated inflammatory destruction. Uveitis caused by an autoimmune disease occurs in a variety of diseases including Bechet's disease, sarcoidosis, and Vogt-Koyanagi-Harada disease. Patients with uveitis often suffer serious visual loss after persistent inflammation due to immune mediated damage to the neuronal cells of the retina.

Since microglia have multiple phenotypes and/or different stages of activation that can be associated with either harmful or [beneficial effects](#) in disease pathogenesis, their role and function in disease progression is not well defined. Researchers across all fields of medicine have recently begun to elucidate the function of microglial cells in various conditions. For example, in Alzheimer's, Parkinson's and other neurodegenerative diseases of the brain, microglia are thought to be harmful. In ophthalmology, it is known that microglial [cells](#) are activated in response to a number of developmental and disease indications and their roles in disease are thought to be context dependent, where they can be either beneficial or harmful.

"These findings provide the first insights into how microglia respond and function during a systemic autoimmune [disease](#) targeting the eye," said lead author Yoko Okunuki, MD, Ph.D., an investigator in Dr. Connor's laboratory and Instructor of Ophthalmology at Harvard Medical School.

"This novel work by Dr. Connor and colleagues identifies that microglia regulate entry through the blood-retinal barrier, and it is our hope that these finding can be harnessed for future targeted therapies for uveitis," says Joan W. Miller, MD, the David Glendenning Cogan Professor and Chair of Ophthalmology at Harvard Medical School, Chief of Ophthalmology at Mass. Eye and Ear and Massachusetts General Hospital, and Ophthalmologist-in-Chief at Brigham and Women's Hospital. "It is becoming increasingly clear that [microglia](#) are involved in a number of retinal disorders as well as neuroinflammatory disorders of the central nervous system.

**More information:** Yoko Okunuki et al., "Retinal microglia initiate neuroinflammation in ocular autoimmunity," *PNAS* (2019).  
[www.pnas.org/cgi/doi/10.1073/pnas.1820387116](http://www.pnas.org/cgi/doi/10.1073/pnas.1820387116)

Provided by Massachusetts Eye and Ear Infirmary

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