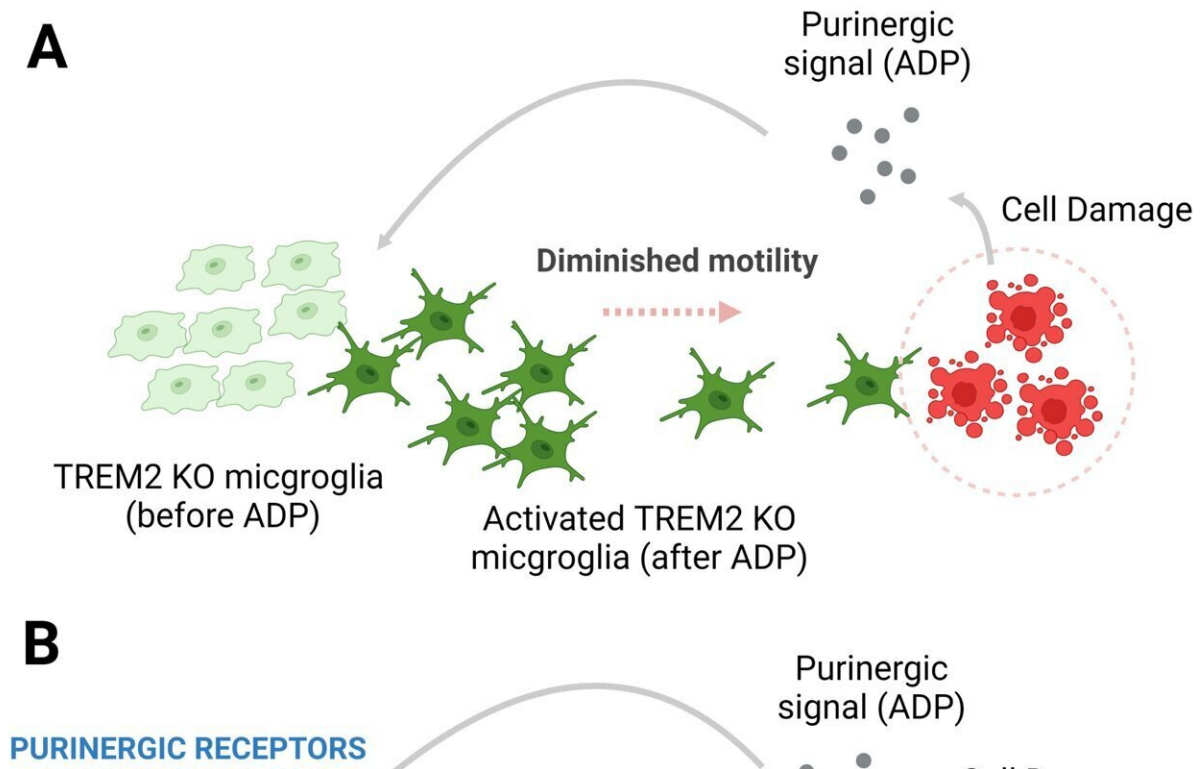


New study reveals how TREM2 gene mutation in brain microglia immune cells can increase Alzheimer's risk

March 23 2022



Induced pluripotent stem cell-derived microglia (iPSC-microglia) deficient in TREM2 and expressing the genetically encoded calcium indicator, Salsa6f. This red-green tandem indicator stably labels the microglial cell in red (tdTomato) and increases its green intensity (GCaMP6f) with rise in cytosolic calcium. The image shows enhanced calcium signaling and process extension in response to purinergic agonist stimulation (ADP). Credit: School of Medicine / UCI

The molecular processes caused by a TREM2 (Triggering Receptor Expressed on Myeloid Cells 2) gene mutation in the brain's microglia immune cells can increase the risk of Alzheimer's disease, according to a recent study led by researchers at the University of California, Irvine. While many immune cell genes have been associated with Alzheimer's, the odds are increased two to three times by mutations in TREM2. However, the processes by which these mutations change the function of microglia cells have not been identified until now.

The study, titled "TREM2 regulates purinergic receptor-mediated calcium signaling and motility in human iPSC-derived microglia" was recently published online in *eLife*.

A critical function of microglia is restoring brain health. Building on their earlier work showing microglia that lack TREM2 have problems moving toward sites of damage in the brain, the research team discovered that this lack of motility is associated with increased levels of P2RY12 and P2RY13 receptors on the microglia. When these receptors detect purinergic molecular danger signals released from nearby damaged cells, they rapidly increase internal messenger calcium within the microglia. The microglia containing higher calcium levels don't turn as frequently and so can't direct themselves to the sites of injury.

"Our study shows that problems arise from having too much signaling between P2RY12 and P2RY13 receptors. We were able to return the microglia [immune cells](#) back to their normal function by blocking these receptors. This suggests that Alzheimer's patients who have mutations in the TREM2 protein would benefit from inhibition of these receptors or related signaling molecules. Manipulating calcium using drugs could be another way to modulate microglia behavior to fight disease," said Michael D. Cahalan, Ph.D., Physiology & Biophysics distinguished

professor and chair at the UCI School of Medicine, and co-corresponding author.

The team used a culture model of microglia derived from human induced [pluripotent stem cells](#) from blood to study the lack of TREM2. A special calcium probe called Salsa6f was engineered into those cells, which allowed calcium to be measured while recording motility characteristics in real time.

"Alzheimer's disease is the most common cause of dementia and currently has no cure," Cahalan said. "Studies of DNA from people with Alzheimer's surprisingly revealed that changes to genes in microglia can have a great impact on whether or not someone gets the disease. Understanding the processes of how these mutations change immune cell function will allow us to develop targeted therapies."

UCI post doctoral scholar Amit Jairaman, and graduate student Amanda McQuade, who is now a postdoctoral fellow at UCSF, are co-first authors of the study.

More information: Murali Prakriya, Editor's evaluation: TREM2 regulates purinergic receptor-mediated calcium signaling and motility in human iPSC-derived microglia, *eLife* (2022). [DOI: 10.7554/eLife.73021.sa0](#)

Provided by University of California, Irvine

Citation: New study reveals how TREM2 gene mutation in brain microglia immune cells can increase Alzheimer's risk (2022, March 23) retrieved 25 April 2024 from <https://medicalxpress.com/news/2022-03-reveals-trem2-gene-mutation-brain.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.