Protecting the brain from dementia-inducing abnormal protein aggregates

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Researchers from the National Institutes for Quantum Science and Technology prove that the protein p62 eliminates and prevents the formation of toxic tau protein aggregates and inflammation and degeneration of neurons. Credit: Maiko Ono from National Institutes for Quantum Science and Technology, Japan

In order to maintain cellular homeostasis (i.e., a state of equilibrium), cells undergo selective autophagy or self-degradation of unwanted proteins. Autophagy receptors control this process, by mediating the selection of a target protein that is then "cleared."

Tau proteins—which otherwise play an important role in stabilizing and maintaining the internal organization of neurons in the brain—abnormally accumulate inside neurons in conditions like dementia and Alzheimer's disease. This build-up of hyper-phosphorylated tau proteins (or tau oligomers) causes the formation of neurofibrillary tangles (NFTs) and eventual cell death of neurons in the brains of people with dementia, contributing to the disease's progressive neurodegenerative symptoms. Now, while tau proteins can be degraded by selective autophagy, the exact mechanism of how this occurs remains a mystery.

In a recent breakthrough, however, a study done by scientists at the National Institutes for Quantum Science and Technology in Japan proved the critical role played by a certain gene—the p62 gene—in the selective autophagy of tau oligomers. The team included researcher Maiko Ono, and group leader Naruhiko Sahara—both from the Department of Functional Brain Imaging at the National Institutes for Quantum Science and Technology in Japan. Their paper, published in Aging Cell, was made available online on 5 June 2022.

Previous studies have reported that the abnormal accumulation of the tau proteins may be selectively suppressed by autophagy pathways, through the p62 receptor protein (which is a selective autophagy receptor protein). Says Maiko Ono, "This protein's ubiquitin-binding ability helps in the identification of toxic protein aggregates (like tau oligomers), which can then be degraded by cellular processes and organelles."

This study's novelty, however, lay in the demonstration of p62's "neuroprotective" role in a living model, which had never been done before. So, how did the researchers achieve this? They used mouse models of dementia. The p62 gene had been deleted (or knocked out) in one group of these mice, so they did not express p62 receptor proteins.

On studying the brains of these mice using immunostaining and comparative biochemical analyses, an interesting picture was revealed. Neurotoxic tau protein aggregates were found in the hippocampus—the area of the brain associated with memory—and brainstem—the center that
coordinates the body’s breathing, heartbeat, blood pressure, and other voluntary processes—of p62 knockout (KO) mice. When we consider this along with the symptoms of dementia, which include memory loss, confusion, and mood changes, these findings make a lot of sense.

MRI scans revealed that the hippocampus of p62 KO mice was degenerated (atrophied) and inflamed. A postmortem assessment of their brains revealed a greater loss of neurons in their hippocampus. Further immunofluorescent studies showed that the abnormal tau species aggregates can cause cytotoxicity leading to inflammation and cell death of neurons in p62 KO mice. Oligomeric tau, specifically, accumulated more in the brains of p62 KO mice.

Overall, the findings of this study prove that by eliminating and, hence, preventing the aggregation of oligomeric tau species in the brain, p62 played a neuroprotective role in models of dementia.

At a time when researchers across the word are trying to develop drugs for dementia and other related neurodegenerative disorders, the findings of this study will be of great importance in providing evidence for the accurate targeting of tau oligomers. The global population of aging humans is increasing each year; hence, the need to develop methods to slow down the onset and progression of various neurodegenerative diseases is also expanding. This study provides a positive step towards addressing that need.


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