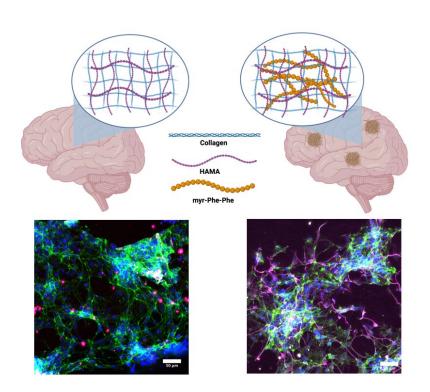


Study reveals insights into Alzheimer's disease mechanisms through novel hydrogel matrix

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By culturing healthy neuronal progenitor cells (NPCs) within this amyloidmimicking environment and comparing results to those in a natural-mimicking matrix, the researchers observed elevated levels of neuroinflammation and apoptosis markers. This suggests a significant impact of amyloid-like structures on NPC phenotypes and behaviors. Credit: Terasaki Institute



Researchers at the Terasaki Institute for Biomedical Innovation (TIBI) have unveiled a pioneering study shedding light on the intricate mechanisms underlying Alzheimer's disease (AD). The study, titled "Effects of amyloid- β -mimicking peptide hydrogel matrix on neuronal progenitor cell phenotype," represents a significant leap forward in understanding the interplay between amyloid-like structures and neuronal cells.

Led by Natashya Falcone and co-first authors Tess Grett Mathes and Mahsa Monirizad, the research team delved into the realm of selfassembling peptide-based hydrogels, renowned for their versatility in mimicking extracellular matrices (ECMs) of diverse microenvironments.

AD presents an intricate challenge in neurodegenerative research. Traditional two-dimensional (2D) models have limitations in capturing the complexity of the disease. Through their innovative approach, the team developed a multi-component <u>hydrogel</u> scaffold, named Col-HAMA-FF, designed to mimic the amyloid-beta (β) containing microenvironment associated with AD.

The study's findings, published in *Acta Biomaterialia*, illuminate the formation of β -sheet structures within the hydrogel matrix, mimicking the nanostructures of amyloid- β proteins.

By culturing healthy neuronal progenitor cells (NPCs) within this amyloid-mimicking environment and comparing results to those in a natural-mimicking matrix, the researchers observed elevated levels of neuroinflammation and apoptosis markers. This suggests a significant impact of amyloid-like structures on NPC phenotypes and behaviors.

Dr. Ali Khademhosseini, the study's corresponding author, said, "This foundational work provides a promising scaffold for future investigations into AD mechanisms and drug testing. By bridging the gap



between 3D hydrogel models and the complex reality of AD pathological nanostructures, we aim to understand this interaction on healthy <u>neuronal</u> <u>cells</u> so that we can accelerate the development of effective therapeutic strategies."

The study represents a crucial step towards unraveling the mysteries of the b-amyloid-like environment which can be found in AD and marks a milestone in the quest for innovative solutions to combat neurodegenerative disorders.

More information: Tess Grett Mathes et al, Effects of amyloid-βmimicking peptide hydrogel matrix on neuronal progenitor cell phenotype, *Acta Biomaterialia* (2024). DOI: <u>10.1016/j.actbio.2024.05.020</u>

Provided by Terasaki Institute for Biomedical Innovation

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