

Seminar

Investigation of the Mechanism of Amyloid Aggregation Using Single-Molecule Fluorescence Techniques

Sourav Das Adhikari

TIFR, Hyderabad

The mechanism of amyloid- β ($A\beta$) deposition in Alzheimer's disease (AD) remains a central yet unresolved question in neurodegenerative research. Understanding how $A\beta$ aggregates, and how lipids and apolipoprotein E (ApoE)—a major genetic risk factor for AD—influence $A\beta$ 42 aggregation pathways, is essential for elucidating AD pathology. In this seminar, I will discuss how distinct phospholipids promote the formation of lipid-peptide condensates with $A\beta$ 42, which act as nucleation sites for aggregation. I will also demonstrate that different ApoE isoforms affect the elongation of $A\beta$ fibrils in different ways. ApoE4, for example, binds to $A\beta$ fibril ends less strongly, making it a less efficient inhibitor of elongation. ApoE4's distinctive function in the promotion of $A\beta$ pathology is further underscored by its ability to enhance condensate-mediated nucleation. Together, these findings reveal the molecular interplay between $A\beta$, lipids, and ApoE, offering new mechanistic insights and potential therapeutic targets for AD.

Wednesday, Oct 22nd 2025

11:30 Hrs (Tea / Coffee 11:15 Hrs)

Auditorium, TIFRH