Thursday, April 13, 2017 12:15 pm, Room 1315 Chemistry

Analytical Seminar

Department of Chemistry

PROF. YOSHITAKA ISHII School of Life Science & Technology Tokyo Institute of Technology



"Solid-state NMR studies of graphenebased systems and misfolded Amyloid-β Assem-

This work involves two separate topics on solid-state NMR (SSNMR) applications to graphene-related nano-materials and protein nano-assemblies associated with Alzheimer's disease (AD). First, we present SSNMR structural studies on graphene- and graphitebased nano materials. It is demonstrated that SSNMR is a powerful tool to analyze the structures and chemical reactions for graphene/graphite oxide,^{1,2} which are widely used as precursors of mass-production for graphene -based materials.

Second, we discuss structural studies of amyloid misfolding for 42-residue Alzheimer's amyloid b (A β). Misfolded fibrillar assemblies of Ab called amyloid fibrils are a primary component of senile plaque, a hallmark of AD. Increasing evidence suggests that formation and propagation of misfolded assemblies of 42-residue Ab (Ab42), rather than the more abundant 40-residue Ab40, provokes the Alzheimer's cascade. Our group recently presented the first detailed atomic model of A β 42 amyloid fibril based on SSNMR data.³ The result revealed a unique structure that was not previously identified for Ab40 fibril. Based on the results, we discuss our ongoing efforts to analyze a structural conversion in misfolding of Ab42 from oligomeric intermediates to fibrils. The metastable oligomers is associated with brain-derived Ab oligomer called amylo-spheroid (ASPD).^{4,5} Additionally, cross seeding of Ab42 with other Ab isoforms will be discussed. The results provide insight into amyloid misfolding of Ab42 in Alzheimer's disease. Other topics such as ultrahigh-field SSNMR for biological applications will be discussed.^{6,7}