
Seminar

Novel Hybrid Complexes from Proteins and Polymer Surfactants

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Proteins represent an extensive class of soft biological colloids with a multitude of advantages that can be exploited for fabricating functional smart materials and bio-based devices. Protein-based biomaterials with distinct structural and physical characteristics, such as elastic fibronectin-elastin films, ultrathin nano-filtration ferritin membranes, micro-porous bovine serum albumin membranes, silk fibroin patterned films and food-protein films, have been recently described. However, incorporation and retention of protein functions such as catalytic activity remain a considerable challenge due to problems associated with surface-induced denaturation, molecular aggregation, steric hindrance at the active site, and incompatibility with non-biological components. Clearly, there is a pressing requirement to retain both structure and function if high performance biomaterials in the form of films, hydrogels, nanoparticles, and fibers are to be successfully exploited for use in bio-sensing, tissue regeneration, molecular recognition, drug delivery, and opto-electronic devices.

In my talk, I will show my present work related to the development of a new class of protein-based nano-conjugated systems based on the molecular integration of structurally and functionally compatible polymer surfactants. I will show how the accessibility of reactive amino acid side chains present on the surface of globular proteins provides a spatially organized chemical platform for integrating hybrid components that can then be used to tune various biomaterial properties via site-specific surface functionalization. Specifically, I will talk about my recent work on the synthesis and characterization of single-component stoichiometric protein-polymer surfactant conjugates that melt at ambient temperatures to form solvent-free liquids with protein concentrations as high as 20-25 wt.%. These novel protein biofluids offer remarkable benefits over aqueous protein dispersions, including exceptionally high concentrations of protein molecules, remarkable levels of thermal stability (half denaturation temperatures of around 160°C), persistent protein dynamics and sustained biological functions. For example, equilibrium dioxygen binding and reversible redox transitions have been demonstrated in solvent-free myoglobin melts, and enzyme activity reported in solvent-free liquid lipases. Working on solvent-free liquid myoglobin and lipase, I also developed an experimental methodology that resulted in the first known examples of enzymatically active self-standing films obtained from the hierarchical assembly of protein-polymer surfactant bio-conjugates. This represented a novel and generic template-free hierarchical self-assembly methodology to fabricate self-supporting functional biomaterials comprising protein concentrations of ca. 35-40 wt.%.

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11:30 AM (Tea/Coffee at 11:15 AM)

Seminar Hall, TCIS