

COINS Seminar #5

Block Copolypept(o)ides: Introducing PeptoMicelles, NanoPeptoGels, PeptoSomes and PeptoPlexes

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Time: 5:00PM – 6:00PM

**Venue: Room #56, Engineering building #5,
The University of Tokyo**

—Abstract—

Polypept(o)ides combine the multifunctionality of polypeptides with the shielding properties of the polypeptide polysarcosine (poly(N-methyl glycine)). Unlike other materials under evaluation for drug delivery applications, these systems are not only biocompatible but biodegradable by proteases. Since only endogenous amino acids are used in the synthesis of polypept(o)ides non-toxic metabolites, amino acids, are likely to occur.

Block copolypept(o)ides possess the ability to self-assemble into core-shell structures by hydrophobic interactions (PeptoMicelles) or complex formation (PeptoPlexes), where PSar exposes targeting moieties and shields the cargo, which can be either physically entrapped or covalently attached to the peptidic block. Furthermore a novel class of cysteine derivatives enables core-cross linking of micelles by disulfide bond formation in a highly controlled manner. Thus, PeptoMicelles, NanoPeptoGels, PeptoSomes and PeptoPlexes are able to act as a novel generation of nano drug delivery systems for hydrophobic drugs, proteins, pDNA or siRNA.



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