

# The Chinese University of Hong Kong Department of Chemistry

Research Seminar Series

**Speaker:** Prof. Yue Zhao

Department of Chemistry University of Sherbrooke

Canada

**Title:** Control of Stimult-Responsive Polymers by

New Methods

**Date:** December 1, 2014 (Monday)

**Time:** 2:30 p.m.

**Venue:** Room G35

Lady Shaw Building





**Speaker:** Prof. Petr Štěpánek

Institute of Macromolecular Chemistry

The Czech Academy of Sciences

**Title:** Interaction of polymer nanoparticles with biological media

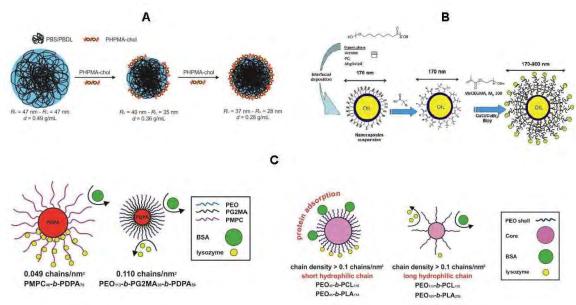
**Date:** December 5, 2014 (Friday)

**Time:** 4:30 p.m.

**Venue:** L1, Science Centre

#### << Abstract >>

Non-specific protein adsorption from complex biological media, especially from blood plasma, is an urgent challenge for the application of nanoparticles as delivery systems, diagnostics and other biomedical applications. The surface modification of nanoparticles by physically anchoring hydrophilic biocompatible polymers (A), by coating the NPs core with polymer brush layers with well-controlled thickness at the nanometer scale through SI-ATRP techniques in water or phosphate buffered saline (B) or by the synthesis of defined hydrophobic-hydrophilic block copolymers (C), are simple and commercially attractive approaches to produce drug-delivery systems resistant to nonspecific protein adsorption. Herein, we report the preparation, characterization and the evaluation of the interaction by placing the aforementioned systems in contact with complex biological media. Several techniques were employed, such as dynamic (DLS), static (SLS) and electrophoretic (ELS) light-scattering techniques, as well as isothermal titration calorimetry (ITC), circular dichroism (CD) spectroscopy and surface plasmon resonance (SPR).



- 1. E. Jäger, A, Jäger, T. Etrych, F.C. Giacomelli, P. Chytil, A. Jigounov, J-L. Putaux, B. Říhová, K. Ulbrich, P. Štěpánek, *Soft Matter* **2012**, 8, 9563-9575.
- C. Rodriguez-Emmeneger, A. Jäger, E. Jäger, P. Štěpánek, A.B. Alles, S.S. Guterres, A.R. Pohlmann, Colloids Surf., B 2011, 83, 376-381.
- 3. F.C. Giacomelli, P. Štěpánek, V. Schmidt, E. Jäger, A. Jäger, C. Giacomelli, Nanoscale 2012, 4, 4504-4514.
- 4. C.A. de Castro, B. Mattei, K.A. Riske, E. Jäger, A. Jäger, P. Štěpánek, F. Giacomelli, Langmuir 2014, 30, 9770-9779.





Speaker: Prof. Michinori Suginome

Department of Synthetic Chemistry and

**Biological Chemistry** 

**Kyoto University** 

**Title:** New Functions of Chirality-Switchable

Macromolecules

**Date:** December 12, 2014 (Friday)

**Time:** 4:30 p.m.

Venue: L1

Science Centre





**Speaker:** 

Dr. Yaowen Wu

Chemical Genomics Centre of the

Max Planck Society

Germany

Title: Protein labeling

Date: December 18, 2014 (Thursday)

Time: 2:30 p.m.

Venue: 1.1

Science Centre





**Speaker:** Dr. Yaowen Wu

Chemical Genomics Centre of the Max Planck Society

Germany

**Title:** Elucidation of membrane trafficking using novel chemical probes

**Date:** December 19, 2014 (Friday)

**Time:** 10:00 a.m.

**Venue:** L5, Science Centre

#### < Abstract >

Protein chemical modification provides large possibilities for modulation of the structure of a polypeptide chain in order to understand protein function. In this talk I will show you the development for a set of site-specific protein modification methods by use of chemoselective reactions, including native chemical ligation, oxime ligation and click chemistry. These approaches enabled us to make single-, dual- or triple-labeled proteins, facilitating investigation of protein-protein interactions, autophagosome formation, protein unfolding and refolding. Recently, we have developed a rapid and fluorogenic affinity conjugation method for labeling of protein inside the cell and a small-molecule switch system for controlling protein function in live cells. I will also show you the biological applications of using these methods to investigate membrane trafficking. Using oxime ligation, we have prepared a set of PEGylated (polyethylene glycol modified) Rab GTPases. Studies of the localization and function of such semisynthetic Rab proteins in cells reveal a comprehensive model for Rab membrane targeting.

- 1. Yi, L.; et al. Angew. Chem. Int. Ed. 2011, 50 (36): 8287-90.
- 2. Yi, L.; et al. Chembiochem, 2011, 12 (16): 2413-7.
- 3. Liu, W.; et al. J. Am. Chem. Soc. 2014, 136 (12): 4468-71.
- 4. Liu, P.; et al. Angew. Chem. Int. Ed. 2014, 53(38):10049-55.
- 5. Li, F.; et al. Proc. Natl. Acad. Sci. U. S. A. 2014, 111 (7), 2572-7.



Yaowen Wu received his BS in Chemistry from Sun Yat-sen University in 2001 and his MS in Organic Chemistry from Tsinghua University in 2004 in China. After graduating as Dr rer. nat. (2008) at the Technische Universität Dortmund working at the Max Planck Institute of Molecular Physiology in Dortmund and a postdoctoral study in cell biology at King's College London, he has been leader of an Otto Hahn group at the Max Planck Institute in Dortmund since 2010. Since 2012 he has been group leader of Chemical Genomics Centre of the Max Planck Society. His research interests are in chemical biology and biochemistry of membrane trafficking.

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## The Chinese University of Hong Kong Department of Chemistry

Research Seminar Series

Speaker: Dr. Xiulan Xie

NMR-Abteilung Fachbereich Chemie

Philipps-Universitat Marburg

Germany

**Title:** NMR Structure Determination of Lasso

Peptides

**Date:** December 19, 2014 (Friday)

**Time:** 11:15 a.m.

Venue: L5

Science Centre

