

Lectins are sugar-binding proteins or glycoproteins that recognize specific carbohydrate structures and agglutinate various types of animal cells. In marine animals, lectins are believed to contribute as non-self recognition factors to the defense mechanism. Interestingly, it has been theorized that some lectins from marine animals mediate the interaction between symbiont and host. SLL-2 is a D-galactose binding lectin isolated from an octocoral, *Sinularia lochmodes*. It was found that SLL-2 was distributed densely on the surface of symbiotic dinoflagellate *Symbiodinium* sp. cells. Previous report showed that SLL-2 transforms free-swimming stage *Symbiodinium* cells into non-motile stage *Symbiodinium* cells and keep them in their non-motile stage [1,2]. These results show that SLL-2 is a chemical cue in the symbiosis between dinoflagellates and coral. The three-dimensional structure of SLL-2 will provide information about the symbiosis mechanism.

The structure of SLL-2 was determined by the molecular replacement method using atomic coordinates of *Helix pomatia* agglutinin (HPA lectin, PDB code: 2ccv) as a search model. Three SLL-2 monomers form a trimer around a non-crystallographic 3-fold axis, and two trimers form a hexameric assembly using hydrogen bonds of three pairs of elongated strands from each monomer. The crystal structure of SLL-2 monomer shows the beta-sandwich lectin fold with six strands. The sites of N-glycosylation and sugar binding ("site 1") were identified clearly in the monomer structure. In the hexameric molecule, two of the six "site 1"s possess galactopyranoside derivative that might come from the N-glycosylation site, three contain the precipitant molecule, and the remaining one accommodate a water molecule. Crystals from low monosaccharide (GalNAc: N-acetyl-D-Galactosamine) concentration holds three GalNAcs and three precipitant molecules in its "site 1"s. The crystal structure of SLL-2-GalNAc complex from sugar-rich environment indicated that GalNAc molecules bind to all "site 1"s. These observations reveal that SLL-2 can maintain both unsymmetrical and symmetrical hexameric molecule stably across various environments. In addition, a large electron density, which appears for a part of oligosaccharides but was not enough to ensure the bound species and its orientation, was observed at extra sugar binding site, "site 2". From these results, we could propose binding mode of the pentasaccharide, which is the unbranched polysaccharide recognized by HPA lectin, to SLL-2, and the function of the SLL-2 in the symbiosis between dinoflagellates and coral.

[1] M. Jimbo, *et al.*, *Comp. Biochem. Physiol. B Biochem. Mol. Biol.* **125** (2000) 227-236. [2] K. Koike, *et al.*, *Biol. Bull.* **207** (2004) 80-86.

Keywords: lectin, coral, symbiosis

MS64.P13

Acta Cryst. (2011) A67, C636

Interface Plasmon in Pseudo One-dimensional Si/SiO₂ core-shell Nanostructures by Electron Energy Loss Spectroscopy

Quan Li,^a Juan Wang,^a Mingwen Chu,^b Marek Malac,^c ^aDepartment of Physics, The Chinese University of Hong Kong, Shatin, New Territory, Hong Kong. ^bCenter for Condensed Matter Sciences, national Taiwan University, Taipei, Taiwan. ^cNational Institute for Nanotechnology, 11421 Saskatchewan Drive, Edmonton, Alberta, (Canada). E-mail: liquan@phy.cuhk.edu.hk

Surface/interface plasmons (IP) in nanostructured materials has generated much interest for their potential applications in plasmonic devices, in particular, those can be operated beyond the diffraction limit of light. While the plasmonic properties of nanomaterials are commonly studied using optical approaches, the best spatial resolution achieved is a fraction of optical wavelength, i.e., ~50 nm. Consequently, mapping of plasmonic properties at higher spatial resolution and imaging of nanostructures of smaller than about 50 nm remains

difficult. On the other hand, TEM-EELS has been recently employed as alternative technique for such purposes [1]. The excellent spatial resolution of the technique enables the mapping of local variations in the material's dielectric response at nanometer scale. Although caution has to be taken, qualitative agreement between the EELS results and material's photonic response make it a powerful tool in probing the local plasmonic properties of nano-objects. Furthermore, the TEM allows to obtain chemical and structural information from the *same* area as the plasmon maps.

The interface plasmon in pseudo one-dimensional Si/SiO₂ core-shell nanostructures have been investigated using electron energy loss spectroscopy (EELS)-related techniques under parallel illuminations, which results are comparable to those obtained by optical methods.

Elongation from a perfect spheroid Si/SiO₂ core-shell nanoparticle to nanorod, and eventually to a long nanowire, results in splitting of the IP modes to a transverse and a longitudinal branch. The longitudinal IP mode red shifts with the aspect ratio increase of the nanoparticle, due to the larger charge separation distance along the long axis of the nanostructure. Retardation effect comes into play in longer nanostructures, leading to periodical pile up of opposite charges. This is revealed by the uniform intensity along the longitudinal direction of the long Si/SiO₂ nanocable in the energy filtered image. The small diameter of the nanostructure determines the dominance of the longitudinal mode in the IP oscillation. Consequently, the optical absorption of small-diameter nanostructures is significantly different from that of the larger ones.

Assembly of the Si nanoparticles into one-dimensional particle chains leads to interaction between the adjacent nanoparticles, which can also induce the splitting of the interface plasmon into transverse and longitudinal polarizations. In addition, such coupling causes spatial re-distribution of the IP intensity, leading to local field enhancement in-between the two nanoparticles. By controlling the growth of Si nanostructures into different morphologies, we demonstrate that the material's optical properties can be manipulated.

This work is supported by the grants from GRF of HKSAR under project No. 402007,414908, and 414709. Additional support was obtained from the NINT and NSERC, Canada.

[1] J. Nelayah, M. Kociak, O. Stéphan, F.J. Garcia de Abajo, M. Tencé, L. Henrard, D. Taverna, I. Pastoriza-Santos, L.M. Liz-Marzán and C. Colliex, *Nat. Phys.*, 2007, **3**, 348.

Keywords: EELS, plasmon, silicon

MS65.P01

Acta Cryst. (2011) A67, C636-C637

Radiation damage and electron energy loss spectroscopy of Au particles on amorphous Ge substrates

Marek Malac, Ray Egerton, Juan Wang^c National Institute for Nanotechnology, 11421 Saskatchewan Drive, Edmonton, T6G 2M9, (Canada). E-mail: marek.malac@gmail.com

Ideally the morphology, structure, and electronic properties from an individual nanoparticle would be obtained in a (scanning) transmission electron microscope (S)TEM experiment at an irradiation dose that does not lead to appreciable sample damage. Gold nanoparticles are of interest as catalysts and as building blocks of plasmonics devices. Irradiation of Au particles by an electron beam results in both hopping of individual Au atoms, observed in extremely small clusters, and in frequent change of orientation of particle with respect to the incident electron beam, observed even in particles with volume more than (10)³ nm³ [1].

A TEM/STEM equipped with a cold field electron source and an electron energy loss (EEL) spectrometer can be used to obtain all the