Resonant soft x-ray scattering from structured polymer nanoparticles

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The application of resonant soft x-ray scattering to chemically heterogeneous soft condensed matter materials is presented. Two structured styrene-acrylic polymer composite latex particles ∼230 nm in diameter were utilized to delineate the potential utility of this technique. Angular scans at photon energies corresponding to strong scattering contrast between specific chemical moieties made it possible to infer the effective radii that correspond to the two polymer phases in the nanoparticles. The results show that resonant soft x-ray scattering should be a powerful complementary tool to neutron and hard x-ray scattering for the characterization of structured soft condensed matter nanomaterials. © 2006 American Institute of Physics. [DOI: 10.1063/1.2356306]

Structured macromolecular materials play an important role in many nanotechnology and biotechnology developments (see, e.g., Sundberg and Durant and references therein). Owing to their small size scale and complexity a wide range of powerful tools, e.g., x-ray and neutron scattering, electron microscopy and scanning probe microscopy, and complementary methods are necessary to infer and delineate morphology and composition. Synthetic composite latex particles, which are composed of two or more homo- or copolymers, have an extraordinary range of applications and are good examples of nanomaterials with internal structures. Complex dynamics during synthesis and competing interfacial forces lead to complex structures, including many states of incomplete phase separation. Competent characterization of these materials is often very challenging and sometimes seemingly impossible. Since the various components of these composite particles are often chemically similar to each other, complete characterization requires both high spatial resolution and high chemical sensitivity. Transmission electron microscopy (TEM) is a technique with high spatial resolution and has made great contributions to date. Due to potential preparation artifacts, “round robin” characterization has been used to arrive at sound morphological models. A complementary technique that can provide excellent chemical sensitivity, but is presently limited in spatial resolution to ∼40 nm, is scanning transmission x-ray microscopy (STXM).

Complementary to microscopy, scattering methods can provide spatial information of ensemble averages approaching the $\lambda/2$ limit ($\lambda = $ wavelength) in a backscattering geometry. Neutron scattering typically requires deuteration. For hard x rays, scattering intensity is determined by electron density differences, which are often quite low. Resonant (anomalous) x-ray scattering (RXS) can achieve improved chemical differentiation. RXS with soft x rays has previously focused on the study of magnetic materials, but also offers important advantages for the differentiation of small phases in soft condensed matter materials or for thin film analysis. Tunable RXS contrast was demonstrated for spherical, single phase polymeric structures embedded in a polymeric matrix and for block copolymer samples. In this letter, the utility of RXS to characterize more complex objects such as structured polymeric nanoparticles will be demonstrated, and the results will be discussed in the context of conventional x-ray scattering, STXM, and TEM capabilities.

Two types of styrene-acrylic polymer latex particles synthesized with a two step polymerization process were characterized. Sample A was composed of a poly(methyl methacrylate) (PMMA) first stage polymerization and a second stage of a random copolymer of 30/70 w/w butyl acrylate (BA) and styrene (S) [P(BA-co-S)]. The conditions utilized lead to nonspherical particles. Sample B was composed of a random copolymer first stage of 40/60 w/w methyl acrylate (MA) and MMA [P(MA-co-MMA)] and a second stage polymerization of polystyrene (PS). Although the synthesis conditions were similar to system A, a more complicated structure resulted, in which phase separation is incomplete and small domains of PS might be dispersed within the seed particle core.

Nanoparticles were cast from a diluted aqueous suspension onto 100 nm thick silicon nitride (Si$_3$N$_4$) membranes. STXM measurements on such samples were performed with the STXM at beamline BL5.3.2 of the Advanced Light Source (ALS) with a lateral resolution of ∼40 nm. Scattering measurements on identically prepared or the very same samples were performed at beamline BL6.3.2 of the ALS...
With NEXAFS microscopy, a core-shell-type structure can for selective PS contrast enhancement. The compositional sensitivity of RXS can be appreciated from Fig. 2. Figures 2(a) and 2(b) show the $\beta$ (absorption) and $\delta$ (dispersion) of PS and PMMA in the vicinity of the carbon 1s absorption edge, respectively. Two sharp characteristic peaks at 285.2 eV (C 1s $\rightarrow \pi^*_{C=C}$ transitions in the phenyl ring of PS) and 288.4 eV (C 1s $\rightarrow \pi^*_{C=O}$ transitions in the carbonyl moiety in PMMA) can be observed for $\beta$. Poly(butyl acrylate) and poly(methyl acrylate) have NEXAFS spectra similar to PMMA, but with slightly different C 1s $\rightarrow \pi^*_{C=O}$ intensities. For the present discussion, we will use the PMMA NEXAFS spectrum as the prototypical acrylate NEXAFS spectrum. The x-ray scattered intensity is proportional to $|\delta(E)+i\beta(E)||\delta(E)-i\beta(E)|$ and is at high photon energies dominated by electron density differences. Near resonance, the intensity is modulated strongly as shown in Figs. 2(c) and 2(d), providing energy dependent, “moiety-specific” scattering. These RXS features should have a characteristic signature for a wide range of materials, similar to and related to the well documented sensitivity of carbon NEXAFS to composition in carbonaceous materials.

Resonant scattering experiments of samples A and B were carried out at three photon energies that were selected to provide resonant enhancement for styrene (285.2 eV) and acrylics (288.4 eV), and for scattering of polymer particles “as a whole” (320.0 eV). Figures 3 show scattered intensity at these three photon energies versus $q$. Well defined oscillations can be observed for both samples due to the relatively monodisperse size distribution of these particles. Irrespective of the shape of particles, the low $q$ maxima and minima in these oscillations are related to the effective radius. Interestingly, the maxima and minima depend strongly on the photon energy, reflecting the effective compositional radial distribution of the nanoparticles. The scattering of sample A indicates that the effective radius corresponding to the PS phases is larger than the one corresponding to the PMMA phase. The corresponding minima and maxima for the 285.2 eV data, dominated by the PS signal, are at significantly lower $q$ values than the minima and maxima for photon energies of
288.4 or 320 eV. This is consistent with prior knowledge of sample A with a well defined PMMA core and a styrene-rich “lobed shell.” In contrast to this, the scattering minima and maxima for sample B do not exhibit a strong dependence on photon energy. Furthermore, the minima in the 288.4 eV data are at smaller q values than in the 285.2 or the 320 eV data. The styrenic and acrylic phases have about the same effective radii with maybe a slight acrylic excess on the surface. Thus, sample B does not have a clearly defined average core-shell structure. A more detailed analysis would require use of more sophisticated models. The semiquantitative discussion presented is sufficient to demonstrate the complementary analysis capabilities of RXS.

The application of RXS will not be limited to the materials employed here. The extension of the technique to structured materials for which there are no TEM staining methods (e.g., structured acrylic based nanoparticles) and to multiphasic materials in general is possible. In addition, RXS scattering is a “photon-in and photon-out” technique, and control of the sample condition including in situ liquid environments and temperature control will be similar to that already achieved in STXM.27 The availability of several scattering profiles will also much better constrain sophisticated models of the samples than if only a single profile, as in neutron or hard x-ray scattering, is available.

In summary, we demonstrated the compositional sensitivity of RXS and its utility for the characterization of nanoscale structured polymeric materials. Conventional XS would not have been able to assess effective radii, and neither STXM nor TEM was able to do so. Neutron scattering would have required the use of deuterated species. RXS thus has the potential to become a powerful complement to neutron and conventional x-ray scattering, particularly for polymeric and biological composite particles, vesicles, micelles, multiblock copolymers, various colloids and microgels, miscible polymer blends, polymer solutions, microcapsules, and lipid membranes, all of which are primarily composed of low Z elements with high, tunable contrast near their K-shell absorption edges.

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10λ = 123.9 nm/46 eV. For E = 300 eV, λ/2 = 2.07 nm; for E = 540 eV, λ/2 = 1.15 nm.
26Although the radius of gyration of a particle is well defined, we have chosen here to use the term “effective radius” for the particle instead in order to avoid any confusion with the radius of gyration of an individual polymer chain.