Abstract ID: 16355 Title: Experimental demonstration of cone-beam polychromatic x-ray fluorescence computed tomography (XFCT) imaging of gold nanoparticle-loaded regions within small animal-sized phantoms

Purpose: To demonstrate the feasibility of determining the location and gold concentration of gold nanoparticle (GNP)-loaded regions within polymethyl methacrylate (PMMA) phantoms using cone-beam polychromatic x-ray fluorescence computed tomography (XFCT) technique.

Methods: Tubes containing water and GNPs at 2%, 1%, and 0.5% by weight were inserted into two 5 cm-long cylindrical PMMA phantoms with 3 cm and 5 cm in diameter, respectively. The phantoms were irradiated by a cone-beam of polychromatic 110 kVp x-rays filtered by 1.0 mm of lead. Energy-sensitive cadmium telluride detectors behind a 2.5 mm diameter lead pinhole collimator collected the spectrum of emitted gold K-shell fluorescence and Compton scattered photons at an angle of 90° relative to the beam central axis as the phantom was rotated to a series of 60 projection angles in 6° steps. Sinograms of gold fluorescence photon signals were constructed by extracting the gold fluorescence peak height from the Compton background, and the image of GNP location and concentration was reconstructed using a maximum likelihood iterative reconstruction algorithm.

Results: Using the measured sinograms, images of GNP location were successfully reconstructed for both the 3 cm and 5 cm phantoms. Additionally, the signal intensity was linearly related to gold concentration. The x-ray dose delivered during the XFCT scanning was measured using the TG-61 protocol, and it was determined that, using an array detector, it would be possible to acquire these images with roughly 30 cGy of x-ray dose delivered.

Conclusions: This study demonstrates, for the first time to our knowledge, the feasibility of imaging GNP location and concentration within small animal-sized objects using an ordinary cone-beam polychromatic x-ray source under realistic constraints on dose and scan time. This investigation strongly suggests a bench-top XFCT system will ultimately become a viable option for in-vivo molecular imaging with conjugated or unconjugated GNPs.