Bisphosphonate-functionalized gold nanoparticles for contrast-enhanced X-ray detection of breast microcalcifications

Lisa E. Cole a, c, Tracy Vargo-Gogola b, c, Ryan K. Roeder a, c, *

* Department of Aerospace and Mechanical Engineering, Bioengineering Graduate Program, University of Notre Dame, Notre Dame, IN 46556, USA
b Department of Biochemistry and Molecular Biology, Indiana University Simon Cancer Center, Indiana University School of Medicine – South Bend, South Bend, IN 46617, USA

A R T I C L E   I N F O

Article history:
Received 1 October 2013
Accepted 26 November 2013
Available online 18 December 2013

Keywords:
Breast cancer
Computed tomography
Contrast agent
Gold nanoparticles
Mammary gland
Microcalcification

A B S T R A C T

Microcalcifications are one of the most common abnormalities detected by mammography for the diagnosis of breast cancer. However, the detection of microcalcifications and correct diagnosis of breast cancer are limited by the sensitivity and specificity of mammography. Therefore, the objective of this study was to investigate the potential of bisphosphonate-functionalized gold nanoparticles (BP-Au NPs) for contrast-enhanced radiographic detection of breast microcalcifications using two models of breast microcalcifications, which allowed for precise control over levels of hydroxyapatite (HA) mineral within a low attenuating matrix. First, an in vitro imaging phantom was prepared with varying concentrations of HA uniformly dispersed in an agarose hydrogel. The X-ray attenuation of HA-agarose compositions labeled by BP-Au NPs was increased by up to 26 HU compared to unlabeled compositions for HA concentrations ranging from 1 to 10 mg/mL. Second, an ex vivo tissue model was developed to more closely mimic the heterogeneity of breast tissue by injecting varying concentrations of HA in a Matrigel carrier into murine mammary glands. The X-ray attenuation of HA-Matrigel compositions labeled by BP-Au NPs was increased by up to 289 HU compared to unlabeled compositions for HA concentrations ranging from 0.5 to 25 mg/mL, which included an HA concentration (0.5 mg/mL) that was otherwise undetectable by micro-computed tomography. Cumulatively, both models demonstrated the ability of BP-Au NPs to enhance contrast for radiographic detection of microcalcifications, including at a clinically-relevant imaging resolution. Therefore, BP-Au NPs may have potential to improve clinical detection of breast microcalcifications by mammography.

© 2013 Elsevier Ltd. All rights reserved.

1. Introduction

Breast cancer is the most common cancer and second leading cause of cancer-related deaths among women in the United States [1]. Early detection by mammography is the current clinical gold standard in breast cancer screening and is considered to be the best hope for controlling the disease due to a demonstrated decrease in breast cancer mortality [2,3]. Microcalcifications are one of the most common abnormalities detected by mammography [4,5] and are coincident with 30–50% of breast cancers detected by mammograms [6–8]. Microcalcifications are deposits of hydroxyapatite (HA) or calcium oxalate (CaOx) mineral within the breast tissue [9–12], which exhibit high X-ray attenuation relative to the low attenuating breast tissue facilitating detection by mammography.

The detection of microcalcifications and correct diagnosis of breast cancer are limited by the sensitivity and specificity of mammography. Mammographically-detected microcalcifications range in size from 0.1 to 1.0 mm, but smaller microcalcifications are detected histologically [6]. Limitations in sensitivity can result in the missed detection of cancerous lesions at an early stage, or false negatives, and therefore lost opportunity for early treatment [13]. Studies have reported that 10–30% of all breast lesions are missed during routine mammographic screening [6,14]. Limitations in specificity can result in false positives [15] and unnecessary biopsies, which increase patient anxiety and healthcare costs [16]. The detection of microcalcifications is limited by their small size and the presence of other structures in the breast, such as fibrous tissue, ducts and blood vessels, which may be mistaken for microcalcifications due to exhibiting greater X-ray attenuation than surrounding tissue [6]. Therefore, improvements are needed for...
detecting microcalcifications during mammographic screening for breast cancer.

X-ray contrast agents [17] could be used to improve the sensitivity and specificity of detecting breast microcalcifications by mammography. Improved sensitivity could be achieved by providing enhanced X-ray contrast for small microcalcifications that are otherwise below the detection limit of mammography, thus enabling earlier detection. Improved specificity could be achieved using a targeted X-ray contrast agent.

Gold nanoparticles (Au NPs) have received recent attention as a potential X-ray contrast agent due to exhibiting greater X-ray attenuation and improved vascular retention compared to iodinated molecular agents [18–20]. Additionally, Au NPs are widely reported to be non-cytotoxic, readily synthesized, and readily surface functionalized for coiicolidal stability and targeted delivery [21]. Au NPs surface functionalized with ligands such as antibodies [22–25], peptides [26] or other tumor-specific biomarkers [27] have demonstrated potential as an actively targeted X-ray contrast agent for cancer cells in vitro or tumors in vivo.

Bisphosphonate-functionalized Au NPs (BP-Au NPs) were recently investigated as a targeted X-ray contrast agent for microdamage in bone tissue [28,29]. Bisphosphonates, such as alendronate, are used as a pharmaceutical in the treatment of osteoporosis [30] due to exhibiting a high binding affinity for HA, the mineral component of bone [31]. BP-Au NPs exhibited colloidal stability in physiological media, a high binding affinity for HA crystals in vitro, and targeted labeling of damaged bone tissue in vitro [28,29]. Interestingly, HA is also the mineral component of the type of breast microcalcifications known to be associated with malignant lesions [9–12,32].

Therefore, the objective of this study was to investigate the potential of BP-Au NPs for contrast-enhanced radiographic detection of breast microcalcifications. Two models of breast microcalcifications were developed for precisely controlling levels of HA mineral within a low attenuating matrix. First, an in vitro imaging phantom was prepared with varying concentrations of HA uniminerals within a low attenuating matrix. The hydrodynamic diameter was measured using DLS after diluting as-prepared BP-Au NPs in DI water to a final gold concentration of ~0.5 µM and reported as the mean of three samples. Colloidal stability was verified by measuring the surface plasmon resonance (SPR) peak using UV–Vis spectroscopy after diluting as-prepared BP-Au NPs in DI water to a final gold concentration of 0.4 mM. Gold concentrations were measured using inductively coupled plasma-optical emission spectroscopy (ICP-OES, Optima 3000, Perkin Elmer, Inc. Waltham, MA) after acidifying with 3% aqua regia (3 HCl:1 HNO3). Calibration curves were created by diluting certified standard gold solutions (Assurance Grade, SPEX CertiPrep, Metuchen, NJ).

2.2. In vitro imaging phantom

An in vitro imaging phantom of breast microcalcifications was created by mixing varying concentrations of HA crystals in an agarose hydrogel. Whisker-shaped HA crystals, exhibiting a mean length and width of ~18 x ~2 µm, were synthesized using the chelate decomposition method, as described elsewhere [34]. The specific surface area of the HA crystals was 5.63 m2/g as measured using Brunauer–Emmett–Teller (BET) N2 adsorption (Autosorb-1, Quantachrome Instruments, Boynton Beach, FL) [28]. HA crystals were added to a 2 wt% agarose hydrogel (Sigma–Aldrich, St. Louis, MO) at varying HA concentrations of 0.5, 1.0, 2.5, 5.0, 10, and 25 mg/mL and gelled into 10 µL pellets at 4°C. The HA concentration in each pellet was verified by measuring the calcium concentration with ICP-OES using the methods described above. Calibration curves were created by diluting certified standard calcium solutions (Assurance Grade, SPEX CertiPrep).

HA-agarose pellets labeled by BP-Au NPs were prepared by labeling HA crystals with BP-Au NPs prior to forming the pellets. HA concentrations of 0.5, 1.0, 2.5, 5.0, 10, and 25 mg/mL were incubated in DI water containing 40 mg/mL BP-Au NPs for 24 h to allow for maximum binding of the BP-Au NPs to the HA crystals [29]. Thus, each HA concentration was exposed to the same BP-Au NP concentration for the same length of time. Unbound BP-Au NPs were separated from HA crystals by centrifugation at ~700 g for 2 min. HA crystals labeled by BP-Au NPs were collected using 0.45 µm filter paper (Nylaflo, Pall Corporation, Ann Arbor, MI), rinsed with 10 mL DI water, dried overnight in an oven at 37°C, and stored under vacuum. Labeled HA crystals were mixed with 2 wt% agarose solution and gelled into 10 µL pellets at 4°C.

Labeled and unlabeled HA-agarose pellets were imaged in Eppendorf tubes by micro-computed tomography (micro-CT, pCT-80, Scanco Medical AG, Bruttisellen, Switzerland) at 45 kVp, 177 µA, 600 ms integration time, and 10 µm resolution for 250 slices. The standard 0.5 mm aluminum filter was removed to increase the number of low energy photons to approximate mammographic imaging. The mean ± standard deviation) sampled volume for each pellet was 5.2 ± 1.3 mm3. The measured mean linear attenuation coefficient of the HA-agarose pellets was converted to Hounsfield units (HU) using an internal sample calibration (air (−1000 HU) and water (0 HU) controls measured from 10 slices in the same sample tube. The X-ray attenuation measurement for each labeled and unlabeled HA concentration was reported in HU as the mean ± standard deviation) of five samples. The percent of BP-Au NPs bound to HA crystals was measured as the concentration of unbound BP-Au NPs subtracted from the initial concentration of BP-Au NPs, divided by the initial BP-Au NP concentration. The initial gold concentration and the concentration of unbound BP-Au NPs were measured using ICP-OES, as described above. The mass of BP-Au NPs bound per HA surface area (mg Au/m2 HA), or surface density, was calculated using the measured specific surface area of the HA crystals. The mean ± standard deviation) percent of BP-Au NPs bound and mass of BP-Au NPs bound per HA surface area were measured from three samples for each HA concentration. The presence and surface density of BP-Au NPs on HA crystal surfaces were also verified using field emission scanning electron microscopy (FE-SEM, 400 XHR, FEI, Hillsboro, OR). Samples were prepared by dropping a solution of HA crystals labeled with BP-Au NPs in 90% ethanol on a heated SEM stub and coating with 2.5 nm iridium by sputter deposition. Samples were imaged at an accelerating voltage of 5 kV and current of 6.3 pA.

2.3. Ex vivo tissue model

Mature mammary glands (MGs) from 8 to 12-week-old nulliparous FVB female mice (Charles River Laboratories International, Inc., Wilmington, MA) were used to develop an ex vivo tissue model of microcalcifications. All studies were approved by the Institutional Animal Care and Use Committee at the University of Notre Dame and were conducted in accordance with the guidelines of the U.S. Public Health Service Policy for Humane Care and Use of Laboratory Animals. All efforts were made to minimize suffering of the mice. Mice were euthanized and dissected to expose the left and right number 4 MGs. HA crystals were mixed with Matrigel (BD Matrigel, BD Sciences, Bedford, MA), a hydrogel comprised of extracellular matrix proteins, at concentrations of 0.5, 0.5, 5.0, and 25 mg/mL, and HA-Matrigel compositions were immediately injected into the fat pad of MGs at a dose of 50 µL. HA-Matrigel compositions labeled with BP-Au NPs were prepared by labeling HA crystals with BP-Au NPs prior to mixing with Matrigel. Three concentrations of
HA crystals, 0.5, 5 and 25 mg/mL, were labeled with two different concentrations of BP-Au NPs (10 mg/L and 74 mg/L) for 24 h using the same methods described above for the in vitro imaging phantom. After labeling, unbound BP-Au NPs were separated from HA crystals by centrifugation at ~700 g for 2 min. HA crystals labeled by BP-Au NP were collected using 0.45 μm filter paper (Nylaflo), rinsed with 10 mL DI water, dried overnight in an oven at 37°C, and stored under vacuum. Labeled HA crystals were mixed with Matrigel and injected into the MGs, as described above.

MGs containing HA-Matrigel compositions were imaged ex vivo by micro-CT within 24 h after injection. Labeled and unlabeled HA-Matrigel compositions were allowed to set for 1 min, before MGs were dissected and stored in phosphate buffered saline at 4°C allowed to set for 1 min, before MGs were dissected and stored in phosphate buffered saline at 4°C. In order to eliminate movement of the MGs and maintain tissue hydration during micro-CT imaging, MGs were embedded in 2% agarose and placed in a polyethylene specimen tube (14 mm diameter). MGs were imaged by micro-CT (μCT-80, Scanco Medical AG) at 45 kVp, 177 μA, and 600 ms integration time at a high-resolution (10 μm) and a more clinically-relevant lower resolution (100 μm). The standard 0.5 mm aluminum filter was removed to increase the number of low energy photons to approximate mammographic imaging. The volume of interest (VOI) for analysis was 3.01 mm³ at 10 μm resolution or 5.67 mm³ at 100 μm resolution for each MG, which included both the HA-Matrigel composition and MG tissue. The measured mean linear attenuation coefficient of the VOI was converted to Hounsfield units (HU) using an internal calibration for air (~1000 HU) and water (0 HU) controls that were included in the first and last micro-CT scan performed each day. The X-ray attenuation measured for each labeled and unlabeled HA concentration was reported in HU as the mean (±standard deviation) physical particle diameter and hydrodynamic diameter were 12.8 (1.6) nm and 45.8 (1.5) nm, respectively. (d) UV-vis spectra of BP-Au NPs showing the surface plasmon resonance peak at ~527 nm.

3. Results and discussion

3.1. Characterization of BP-Au NPs

BP-Au NPs were synthesized using the citrate reduction method and surface functionalized with alendronate, a bisphosphonate with a primary amine for binding to gold surfaces opposite phosphonate groups for targeting calcium ions on HA crystal surfaces. (Fig. 1a). Resulting BP-Au NPs were spherical and monodispersed with a mean (±standard deviation) particle diameter of 12.8 (1.6) nm (Fig. 1b), as measured by TEM [28]. The colloidal stability of BP-Au NPs was characterized using DLS and UV–Vis spectroscopy. The mean (±standard deviation) hydrodynamic diameter was 45.8 (1.5) nm, as measured by DLS (Fig. 1c), which was consistent with previously published results for 13 nm BP-Au NPs [29]. The characteristic SPR peak was observed at ~527 nm in UV–Vis spectroscopy (Fig. 1d), which was also consistent with previously published results for 13 nm BP-Au NPs [28,29]. The hydrodynamic diameter and characteristic SPR peak were characterized for each batch of BP-Au NPs synthesized and over time to ensure that BP-Au NPs remained well-dispersed and stable.

Colloidal stability is critically important for a targeted nanoparticle contrast agent as nanoparticles must remain dispersed in relevant media for targeted delivery. As-prepared Au NPs are stable in water, but are typically unstable in physiological media, requiring the addition of appropriate functional ligands [29,35]. BP-Au NPs...
were previously shown to exhibit colloidal stability in phosphate buffered saline (PBS) and fetal bovine serum (FBS) for at least 7 d due to a combination of the steric and electrostatic stabilizing effects of bisphosphonate ligands [29].

3.2. In vitro imaging phantom

An in vitro imaging phantom was developed to investigate the potential of BP-Au NPs for contrast-enhanced X-ray detection of microcalcifications. Microcalcifications in breast tissue were modeled by mixing varying concentrations of HA crystals in an agarose solution. Agarose provided a matrix with low X-ray attenuation for uniformly dispersing the HA crystals. The range of HA concentrations was used to model different stages of microcalcification burden, including the earliest stages when microcalcifications may be too small to be detected by mammography [6].

The accuracy and precision of the model was validated by correlating the expected HA concentration within HA-agarose pellets against the measured HA concentration from ICP-OES (Fig. 2). Measured HA concentrations were less than expected HA concentrations (p < 0.05, paired t-test), most likely due to losses of HA during mixing and subsequent analysis, but the mean error was only −8.3%, indicating reasonable accuracy. More importantly, linear least squares regression indicated that HA concentrations were highly reproducible (R² = 0.992). Therefore, the in vitro imaging phantom was determined to exhibit sufficient accuracy and precision for investigating the effects of the HA concentration and BP-Au NP labeling on X-ray attenuation.

The X-ray attenuation of unlabeled and BP-Au NP labeled HA concentrations was measured using micro-CT. X-ray attenuation increased with increasing HA concentration for both unlabeled and labeled HA groups (p < 0.001, ANOVA) (Fig. 3). The X-ray attenuation of unlabeled HA exhibited a linear relationship with HA concentration (R² = 0.997), as expected. All unlabeled HA concentrations exhibited significantly greater X-ray attenuation compared to the agarose-only (0 mg/mL HA) control (p < 0.005, t-test). Labeling by BP-Au NPs enhanced the X-ray attenuation of HA concentrations ranging from 1 to 10 mg/mL compared to unlabeled HA at the same concentration (*p < 0.005, t-test).

The X-ray attenuation of the lowest (0.5 mg/mL) and highest (25 mg/mL) HA concentrations were not increased when labeled by BP-Au NPs. Therefore, the percent of the BP-Au NP dose bound to HA and the mass of BP-Au NPs bound per HA surface area (mg Au/m² HA) were measured to investigate differences with HA concentration. Complete binding of the BP-Au NP dose occurred at HA concentrations ≥2.5 mg/mL and the percent binding decreased with decreasing HA concentration (p < 0.0001, ANOVA) (Fig. 4a). On the other hand, HA crystals surfaces were saturated with BP-Au NPs at HA concentrations ≤2.5 mg/mL and the surface density decreased with increasing HA concentration (p < 0.0001, ANOVA) (Fig. 4b). The decreased surface density of BP-Au NPs with increasing HA concentration was also directly confirmed by electron microscopy (Fig. 5). Thus, BP-Au NPs were unable to provide enhanced X-ray contrast at 0.5 mg/mL and 25 mg/mL due to a limited number of total available binding sites at low HA concentrations and a limited number of BP-Au NPs available for binding at high HA concentrations, respectively.

These results suggest a limit to contrast enhancement at HA concentrations ≤0.5 mg/mL using this model, since HA crystal surfaces were saturated with BP-Au NPs (Fig. 4b). Contrast enhancement at the lowest HA concentration (0.5 mg/mL) would seem to be advantageous for detecting HA concentrations otherwise virtually undetectable over the background signal. Therefore,

![](image_url)
contrast enhancement at HA concentrations ≤0.5 mg/mL would require either more sensitive imaging methods or the delivery of a greater mass concentration of Au NPs. On the other hand, a lower limit in HA concentration for contrast enhancement might also be advantageous to prevent false positives. Thus, the level of such a limit or the minimum microcalcification burden for clinical breast cancer diagnosis is critically important but currently unknown.

These results also suggest that contrast enhancement could be achieved at higher HA concentrations (e.g., 25 mg/mL) by simply increasing the BP-Au NP dose. However, contrast enhancement is not necessary at high HA concentrations (e.g., 25 mg/mL) that are readily detected without contrast enhancement. Moreover, an upper limit in HA concentration for contrast enhancement may be advantageous for determining microcalcification burden. Therefore, an optimal BP-Au NP dose must be determined for providing sufficient contrast enhancement over the most appropriate range of HA concentrations, preferably below the detection limits in the absence of contrast enhancement. For this reason, a higher and lower BP-Au NP concentration was investigated in the ex vivo tissue model described below.

Interestingly, the greatest X-ray contrast enhancement in this model occurred at an HA concentration of 5 mg/mL. Unlabeled HA at 5 mg/mL exhibited ~21 HU, which is similar to the X-ray attenuation of the glandular tissue component of human breast tissue at 45 kVp [36], and labeling with BP-Au NPs increased the X-ray attenuation to ~47 HU. This suggests that BP-Au NPs may be particularly useful for providing enhanced contrast to detect microcalcifications in dense breast tissue, which is composed of relatively greater amounts of glandular tissue. This finding is clinically significant because sensitivity for the detection microcalcifications is dramatically reduced in women with radiographically dense breast tissue [37–39], while increased breast density is associated with an increased risk of cancer [40].

![Fig. 4. BP-Au NP binding to varying concentrations of HA crystals in HA-agarose pellets was characterized by measuring the concentration of unbound BP-Au NPs using ICP-OES. (a) The percent of BP-Au NPs bound to HA crystals showed complete binding of the BP-Au NP dose at HA concentrations ≥2.5 mg/mL and decreased percent binding with decreasing HA concentration (p < 0.0001, ANOVA). (b) The mass of BP-Au NPs bound per HA surface area (mg Au/m² HA) showed that HA crystals surfaces were saturated with BP-Au NPs at HA concentrations ≥2.5 mg/mL and the surface density decreased with increasing HA concentration (p < 0.0001, ANOVA). Error bars show one standard deviation of the mean (n = 3) and error bars not shown were negligibly small.](image)

![Fig. 5. Representative FE-SEM micrographs showing the surface density of BP-Au NPs (arrows) on HA crystal surfaces. The surface density appeared to increase with decreasing HA concentration and became saturated at HA concentrations ≤2.5 mg/mL, in agreement with quantitative measurements of the mass of BP-Au NPs bound per HA surface area (Fig. 4b).](image)
The in vitro imaging phantom demonstrated the ability of BP-Au NPs to enhance X-ray contrast for the detection of microcalcifications (Fig. 3), but was not without limitations. The measured X-ray attenuation of agarose was approximately −14 HU or 1.189 cm\(^{-1}\) at 45 kVp, which provided a good average approximation, but was higher than adipose tissue (−260 HU at 45 kVp) and lower than glandular tissue (20 HU at 45 kVp) [36]. Additionally, HA crystals were uniformly dispersed within the agarose matrix. However, breast microcalcifications may appear as single crystals or clustered aggregates of many crystals and the degree of clustering may be used to distinguish between malignant and benign lesions [41,42]. Therefore, an ex vivo tissue model was developed to more closely recapitulate the heterogeneity of microcalcifications in breast tissue.

3.3. Ex vivo murine tissue model

An ex vivo murine tissue model was developed to more closely mimic the heterogeneity of breast tissue and also lay the groundwork for an in vivo model. Murine MGs are important models for studying human breast cancer [43] and were used in this study to mimic the low X-ray attenuation of adipose tissue in the human breast [36]. Varying concentrations of unlabeled and labeled HA were mixed with Matrigel and injected into the fat pad of excised murine MGs to model the heterogeneity of microcalcifications in breast tissue. Varying HA concentrations were used to model different levels of microcalcification burden, including the earliest stages when microcalcifications may be too small to be detected by mammography [6]. The injection volume and HA concentration were readily tailored providing a straightforward and reproducible model for creating microcalcifications of known HA concentration for investigating X-ray contrast agents. Breast microcalcifications have also been biologically-induced in mice and rats via implanted cancer cells [44] and delivery of exogenous growth factors [45–47], respectively, but these models require separate measurements to quantify the HA concentration or microcalcification burden rather than offering a priori control.

X-ray attenuation increased with increasing HA concentration for both unlabeled and labeled HA-Matrigel compositions imaged by micro-CT at both high (10 \(\mu\)m) and low (100 \(\mu\)m) resolution (\(p < 0.001\), ANOVA) (Fig. 6). The X-ray attenuation of unlabeled HA exhibited a linear relationship with HA concentration at both high and low imaging resolution (\(R^2 > 0.993\)), as expected. Unlabeled HA concentrations of 5 and 25 mg/mL exhibited significantly greater X-ray attenuation compared to the Matrigel control (0 mg/mL HA) at both high (\(p < 0.005\), t-test) and low (\(p < 0.05\), t-test) resolution, but differences between 0.5 mg/mL HA and the Matrigel control were not statistically significant (\(p > 0.28\)) (Fig. 6). Therefore, HA-Matrigel compositions containing 0.5 mg/mL HA were undetectable relative to the Matrigel-only control. Interestingly, the microcalcifications created at 0.5 mg/mL HA appeared to be less than 0.1 mm in size in micro-CT images, and increased in size with HA concentration to greater than 1.0 mm at 25 mg/mL HA (Fig. 7). Therefore, the microcalcifications produced by the model included sizes below the detection limit of mammography (<0.1 mm) and within the detection limits of mammography (0.1–1.0 mm) [6].

Differences in the X-ray attenuation between unlabeled and BP-Au NP labeled HA concentrations within murine MGs were also measured by micro-CT at both high (10 \(\mu\)m) and low (100 \(\mu\)m) resolution. Using high-resolution imaging (10 \(\mu\)m), labeling by either dose of BP-Au NPs enhanced the X-ray attenuation of all HA concentrations compared to unlabeled HA at the same HA concentration (\(p < 0.05\), t-test), including at an HA concentration (0.5 mg/mL) that was otherwise undetectable by micro-CT (Fig. 6a). Using low resolution imaging (100 \(\mu\)m), labeling by either dose of BP-Au NPs enhanced the X-ray attenuation compared to unlabeled HA at an HA concentration of 5 mg/mL (\(p < 0.05\), t-test) (Fig. 6b). Note, however, that non-significant differences between unlabeled and labeled HA-Matrigel compositions at 0.5 and 25 mg/mL HA were nearly significant (\(p < 0.13\)), but underpowered. Post hoc power analysis indicated that these differences would become statistically significant if the sample size were increased from 5 to 9 specimens. The differential contrast measured for BP-Au NP labeled HA concentrations was visually apparent in grayscale micro-CT images (Fig. 7). A maximum differential contrast enhancement of 289 or 242 HU was achieved at 25 mg/mL HA with the higher gold dose when imaging at high or low resolution, respectively (Fig. 6).

Results from the ex vivo murine tissue model showed that BP-Au NPs provided enhanced contrast at all HA concentrations (Fig. 6a), including the lowest (0.5 mg/mL) and highest (25 mg/mL) HA concentrations which were not able to be detected in the in vitro phantom (Fig. 3). Moreover, the magnitude of differential contrast...
obtained with the ex vivo tissue model was significantly greater than that obtained with the in vitro phantom. The greater sensitivity and contrast enhancement in the tissue model was most likely due to the lower mean background signal in the tissue model \((w/C0 \approx 300)\) compared to the imaging phantom \((w/C0 \approx 14 \text{ HU})\).

Imaging resolution was important to investigate for clinical translation as current mammographic imaging systems have a microcalcification detection limit of \(\leq 100 \, \mu \text{m}^{2}\) [48]. Therefore, tissue samples were imaged at both high-resolution (10 \(\mu\text{m}\)) and a more clinically-relevant lower resolution (100 \(\mu\text{m}\)). High-resolution imaging provided excellent visualization of the structure of the MGs, including the ducts which attenuated more than the adipose tissue (Fig. 7a), and detected enhanced contrast for BP-Au NP labeled HA-Matrigel compositions at each HA concentration (Fig. 6a). However, this level of resolution is not yet available clinically. At the more clinically-relevant lower resolution (100 \(\mu\text{m}\)), the ducts were not readily visualized (Fig. 7b), but enhanced contrast was still able to be detected for BP-Au NP labeled HA-Matrigel compositions (Fig. 6b), though some differences were underpowered at the lowest and the highest HA concentration, as discussed above. This is a very promising result, as future studies will continue to investigate BP-Au NPs in in vivo models using pre-clinical imaging systems with resolution lower than 10 \(\mu\text{m}\).

All differences in X-ray attenuation between the low (10 mg/L) and high (74 mg/L) BP-Au NP dose at the same HA concentration were not statistically significant \((p > 0.23, t\text{-test})\). However, the low and high dose were chosen to be below and above, respectively, the equilibrium saturation of Au NPs bound to HA surfaces [28], which was expected to lead to differences in the surface density of BP-Au NPs labeling HA crystals and the measured X-ray attenuation. Therefore, the percent of the BP-Au NP dose bound to HA and the mass of BP-Au NPs bound per HA surface area (mg Au/m\(^2\) HA) were measured to investigate differences with HA concentration and the BP-Au NP dose. Complete binding occurred at HA concentrations \(\geq 5.0 \, \text{mg/mL}\) for either BP-Au NP dose, but the percent binding was significantly lower \((p < 0.001, t\text{-test})\) for the higher BP-Au NP dose at a lower HA concentration \((0.5 \, \text{mg/mL})\) where binding was incomplete (Fig. 8a). On the other hand, the surface density of BP-Au NPs was significantly greater for the higher BP-Au NP dose at each HA concentration \((p < 0.0001, t\text{-test})\) and decreased with increasing HA concentration for either BP-Au NP dose \((p < 0.0001, \text{ANOVA})\) (Fig. 8b). These differences in the surface density of BP-Au NPs were also directly confirmed by electron microscopy (not shown). Thus, a greater concentration of BP-Au NPs was able to drive equilibrium surface adsorption toward a greater surface density of BP-Au NPs labeling HA crystals surfaces, as expected [28]. However, the greater mass concentration of gold labeling HA crystals using the higher dose of BP-Au NPs (Fig. 8b) did not translate into a statistically significant increase in X-ray attenuation (Fig. 6). This suggests that while the higher BP-Au NP dose provided additional Au NPs labeling HA crystal surfaces, the additional dose was unnecessary for contrast-enhanced X-ray detection as the low BP-Au NP dose was alone sufficient to provide enhanced contrast at all HA concentrations (Fig. 6). In other words, the additional BP-Au NP dose provided negligible additional increase in contrast compared to the contrast between labeled and unlabeled HA concentrations. Note, however,
mammography [52], computed tomography [53–55], high-resolution ultrasound [56], near-infrared fluorescence imaging with targeted molecular agents [57,58], and Raman spectroscopy [12,59,60]. However, a targeted X-ray contrast agent, such as BP-Au NPs, could provide a more sensitive and more readily translatable diagnostic tool that could be used in conjunction with current mammography systems.

This study also demonstrated the utility of two models of breast microcalcifications, including an in vitro imaging phantom and ex vivo murine tissue model, for evaluating targeted contrast agents. In vivo models of breast microcalcifications have been developed in mice using implanted tumor cells [44] and in rats using exogenous growth factors [45–47]. These models are more biologically-relevant and will be useful for future investigations for the targeted delivery of BP-Au NPs and radiographic imaging in vivo. However, the in vitro imaging phantom and ex vivo murine tissue model afforded more precise and a priori control over the HA concentration for an initial pre-clinical evaluation of a targeted contrast agent. Moreover, the methods developed to create microcalcifications in the ex vivo tissue model could be applied to an in vivo murine model.

The main limitation of this study was that HA crystals were labeled with BP-Au NPs in DI water prior to being mixed with the agarose solution or Matrigel carrier. Therefore, the mass concentration of BP-Au NPs labeling HA crystal surfaces in these models may represent the best-case scenario compared to delivery in vivo. As discussed above, only a fraction of the initial BP-Au NP dose will reach the targeted site upon in vivo delivery [26,27]. Moreover, the binding of BP-Au NPs to HA crystals was previously shown to be decreased in the presence of serum proteins [28]. Therefore, the contrast enhancement achieved in this study is encouraging since the Au NP concentration was relatively low at 10–74 mg/L. Previous studies on X-ray imaging have utilized Au NP concentrations ranging 0.6–2500 mg/L in vitro [22,27] and 3000–270,000 mg/L in vivo [18–20,23–27].

Another important limitation was the small size and low background intensity of the specimens in this study relative to human breast mammography. The level of contrast enhancement, or signal-to-noise ratio, is expected to decrease with increased specimen thickness or density, due to increased absorption and scattering of photons by the background material or tissue. Nonetheless, this study was an important first step to demonstrate feasibility prior to ongoing studies investigating the targeted delivery of BP-Au NPs and radiographic imaging in mice in vivo, including models of radiographically dense breast tissue.

4. Conclusions

Contrast-enhanced detection of breast microcalcifications using BP-Au NPs was investigated using an in vitro imaging phantom and ex vivo murine tissue model comprising varying and reproducible levels of HA crystals. X-ray attenuation increased linearly with increasing HA concentration, as expected. The X-ray attenuation of HA compositions labeled by BP-Au NPs was significantly increased compared to unlabeled compositions at the same HA concentration, including an HA concentration that was otherwise undetectable by micro-CT. Contrast enhancement was shown to be dependent on achieving a sufficient mass concentration of BP-Au NPs labeling HA crystals surfaces. Thus, BP-Au NPs enabled contrast-enhanced detection of breast microcalcifications by X-ray imaging, including at a clinically-relevant imaging resolution. BP-Au NPs may therefore have potential to improve clinical detection of breast microcalcifications by mammography.
Funding sources

Walther Cancer Foundation, Indiana CTSI (NIH RR025761).

Acknowledgments

This research was supported by the Walther Cancer Foundation and the Indiana Clinical and Translation Sciences Institute (CTSI) (NIH RR025761). The authors acknowledge the Freimann Life Science Center at the University of Notre Dame for the use of BET and ICP-SEMS, and the Center for Environmental Science and Technology at the University of Notre Dame for the use of BET and ICP-OES.

References


