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TITLE:  Detection of Breast Microcalcifications Under Ultrasound Using Power Doppler and Acoustic Resonance Imaging

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**Title andSubtitle**
Detection of Breast Microcalcifications Under Ultrasound Using Power Doppler and Acoustic Resonance Imaging

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**Supplementary Notes**
report contains color

**Abstract (Maximum 200 Words)**
Intracranial breast carcinoma (DCIS) represents approximately one third of mammographically detected breast carcinoma. Currently, DCIS and benign breast microcalcifications can only be reliably be evaluated utilizing x-ray mammography. Our goal with our current project was to utilize breast sonography coupled with the technique of acoustic resonance to image and evaluate the breast microcalcifications in patients prior to biopsy. We have been successful in evaluating the calcifications. However, thus far, our analysis demonstrates no significant difference between the malignant and benign calcifications. Currently, work is ongoing with patient accrual.

**Subject Terms**
breast cancer, Doppler imaging, ultrasound, diagnosis, mammography

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Introduction:

Technical advances in the field of mammography are resulting in higher detection rates of early breast cancers than in the past, of both solid masses and microcalcifications. These technical advances include digital mammography, high frequency ultrasound transducers and breast magnetic resonance imaging (MRI). Mammography is still the only reliable method of evaluating microcalcifications in the breast. Breast MRI, although has a sensitivity rate nearing 100% for invasive carcinoma, the reported sensitivity for DCIS has been reported to be as low as 40% (1-3). It is estimated that DCIS represents 20-30% of breast carcinomas detected on screening mammography (4) and has been steadily been rising in incidence since the 1970’s (5). Our goal is to image breast microcalcifications utilizing sonography, a readily available equipment in breast imaging centers, coupled with acoustic resonance. The concept of acoustic resonance imaging (ARI) is based on the size of the microcalcifications and the binding strength with the surrounding tissues in which it is imbedded. When subjected to a wide frequency range, different sized particles will resonate at different frequencies given the same binding environment. By “tuning” into the appropriate frequency range, it would be possible to selectively visualize microcalcifications of varying sizes
Body:

At the writing of this report, we are currently 21.5 months into our project. Dr. Weinstein coordinated the project with Dr. Seghal. Dr. Weinstein has been involved in patient selection and in performing the imaging evaluation along with the research coordinator. Dr. Seghal has organized the imaging protocol and is overseeing the digitizing of the images and data analysis performed in the Ultrasound Research Laboratory.

Task 1: See report from 7/1/00-6/30-02.

Task 2: Patient recruitment period completed

Task 3: Ongoing patient recruitment. Continuation of Task 2.

Since the writing of our last report, we have been involved in patient accrual for the study. To this date, we have accrued thirty-two patients. In thirty-one of those patients, data was obtained for the study. Due to technical factors, data could not be obtained from one of the patients. We have made significant progress in patient recruitment since the writing of our last annual report. Although we are slightly behind schedule, we are optimistic that we will continue to accrue at the current pace.
The patients present to the Breast Imaging Section of the Hospital of the University of Pennsylvania on the day of their biopsy procedure. The patients recruited for the study have breast microcalcifications that are considered suspicious. The patients may undergo excisional biopsy in the operating suite by a surgeon. Alternatively, they may undergo percutaneous core needle biopsy in the Breast Imaging Section in order to obtain a tissue diagnosis. Prior to either of these conventional biopsy procedures the patients are recruited for the study.

The age of the patients have ranged from 40 to 83 years. The breakdown of the age of the patient population is shown in Figure 1 by the decade. The demographics of the patient population is shown in Figure 2. Thus far, out of the 31 patients, 20 patients had benign calcifications on pathologic evaluation and 11 patients had calcifications that were demonstrated to be malignant (Figure 3). Figure 4 shows the age of the patients relative to the pathology results.

First gray scale imaging was performed at the site of the microcalcifications. Next, the same area was scanned in Power Doppler mode in conjunction with acoustic resonance. The acoustic resonance device emitted low frequency vibrations in 10 Hz steps between 50-600 Hz. In the first nine patients, 100 Hz steps were utilized, and it was decided that the steps too large and a decision was made to utilize smaller incremental steps. The scanning was repeated using 3 different vibration amplitudes to determine variation and the frequency of response of the calcifications to the variation in the amplitude. All the images were stored on a videotape and used for quantitative analysis.

The mammogram from each patient was digitized at 300 DPI and stored in the database for future review. The ultrasound examination was performed using state of the
art ATL 5000 ultrasound scanner. All imaging was performed with 12.5 MHz broadband transducer. The instrument reset determined from the studies conducted in the first year were used for all patients.

The acoustic resonance scans involving low frequency vibrations were tolerated by all the patients. We have had no adverse reactions to date. Average total of scan time was approximately 15 minutes. Imaging was completed successfully in 31/32 patients. In 1 case, the examination was not completed due to problems with the scanning equipment.

We have completed quantitative analysis of images from all of the patients enrolled in the study to date. This involved digitizing the images from the videotape. The images of each patient were compiled in a single file and used for quantitative measurement of mean mean color level (MCL), percent area of color (FA) and color weighted fractional area (CWFA). The analysis was performed by selecting the entire image as proposed in the application. However, the results show that structures other than calcifications, such as connective tissue are also enhanced by the acoustic resonance imaging. In view of these results we believe that the images may have to be re-analyzed by choosing specific area of interest to decrease the background noise.

Of the 31 patients studied, 20 cases (64.5 %) were benign and the remaining 11 cases (35.5 %) were malignant. Figure 5 shows an example of images obtained from a patient with malignant calcifications. The mammogram in 5a shows a cluster of calcifications. The biopsy proven malignant calcifications are circled. Figures 5b-d show 3 representative ultrasound images at 50 Hz, 350 Hz and 480 Hz from a total of 165 images. In the gray scale image Figure 5b (50 Hz), calcifications cannot be seen with any
level of confidence. As the frequency of vibration increases, at 350 Hz, calcifications can be seen in color superimposed on gray scale image. At 350 Hz (Figure 5c), the power Doppler pixels, representing calcifications, are in a distribution pattern that is very similar to that seen on the mammogram Figure 5a. In addition to the calcifications, enhancement of secondary structures associated with ducts and connective tissue are also observed (arrow) in Figure 5c. At higher frequency, this effect becomes significant and can mask the calcifications. This is shown in Figure 5 d imaged at 480 Hz. The inset panel (Figure 5e) showing a graph represents change in image enhancement (CWFA) as a function of vibration frequency (Hz). The error bars on the data point represent +/- image deviation of the three measurements at three different scanning parameters. On all three different scans, the graph demonstrates a clear resonance peak at 480 Hz.

Figure 6 shows the results from a patient with benign calcifications. Figure 6a shows the mammographic image of the calcifications. There is a focal cluster of calcifications that is circled. Figure 6b demonstrates the region of the calcifications in gray scale. Again, the calcifications are difficult to discern. However, there is enhancement seen at 280 Hz. The expected region of calcifications is circled. However, there is significant enhancement of secondary structures present. Conclusions are similar to those obtained from Figure 5.

The peak position (P) and the width of the peak at half height (W) were measured for each patient. These values are summarized in Table 1. In all patients with malignant calcifications, enhancement was seen in the power Doppler mode and a peak in CWFA vs. frequency curve was observed in all the cases. The mean +/- standard deviation for the peak position (P) was 363 +/- 132. The peak width (W) was 132 +/- 50 Hz. In
patients with benign calcifications, the image enhancement was observed in 18 (90%) of 20 cases studied. Two cases did not show any measurable enhancement. The mean +/- standard deviation for the peak position and peak width were 381 +/- 111 Hz and 122 +/- 54 Hz respectively. There appears to be no significant difference in the peak width and peak position between the benign and malignant calcifications.

**Technical Issues:**

In the last annual report review, a technical issue was raised. The size of the particles used in the phantoms was questioned. The particle size used in the phantom work ranged from 400-800\(\mu\). While some malignant breast microcalcifications may be smaller than this size, benign breast calcifications may vary greatly in size and certainly fall within this size range.

Once we are able to recruit a pathologist for the study. The actual size of the particles will be measured from the pathologic specimen.

**Key Research Accomplishments:**

- We have been able to visualize calcifications with power Doppler and acoustic resonance in patient studies in 29/31 patients. No significant enhancement could be seen in two patients who both had benign calcifications.
- The visualized calcifications demonstrated a resonance peak.
• There was no significant difference in the resonance peaks between the malignant and benign calcifications

**Reportable Outcomes:**


**Conclusions:**

The results of this study to date show that while we were able to observe calcifications using power Doppler and acoustic resonance in the majority of the cases (29/31), there is no significant difference could be observed between the malignant and the benign calcifications. At this stage of the study, we speculate, that the lack of differentiation between the two types may be associated with the enhancement from the secondary structure such as ducts and connective tissues of the breast that would be seen with both benign and malignant calcifications.
At present analysis, we have analyzed the entire region present in the image field of view. This includes the enhancement of secondary structures. To overcome this limitation, the image data must be re-analyzed. We propose to review the images and select out the regions that represents connective tissues. These areas and the image will be discarded from the analysis to determine if the enhancement of calcifications did mask in the differentiation between the malignant and benign calcifications.
**References:**


**Appendices:**

See attached
Figure 1

Age of the Patients

Number of Patients

Age of the Patients
Figure 2

Benign Vs. Malignant Calcifications

Number of Patients

Benign

Malignant
Figure 3

Age of the Patients vs. Pathology

Number of Patients

Age of the Patient
Figure 4

Demographics

Number of Patients

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Figure Legends:

Figure 1. Figure 1 demonstrates the age distribution of the patients who participated in the study.

Figure 2. Figure 2 shows the number of benign and malignant calcifications that have thus far been imaged and evaluated.

Figure 3. Figure 3 shows the age distribution of the patients and the number of benign and malignant calcifications in each of the age groups.

Figure 4. Figure 4 shows the demographics of the patients who participated in the study.

Figure 5.
A. A mammographic image of the malignant calcifications is demonstrated. The calcifications are circled.
B. A sonographic image of the region of the calcifications is shown. The calcifications are difficult to see at 50 Hz.
C. At 350 Hz, there are power Doppler pixels demonstrated. The expected region of the calcifications is circled.
D. At 480 Hz, there is increased enhancement of the secondary structures.
E. The graph shows the degree of enhancement as a function of frequency. There is a clear resonance peak at 480 Hz.

Figure 6.
A. A mammographic image from a patient with biopsy proven benign calcifications is shown. The calcifications are circled.
B-C. No significant enhancement is seen at 50 Hz. At 280 Hz, enhancement in the region of the calcifications (circled) is seen but there is also enhancement of the surrounding secondary structures in the breast.
D. The graph of degree enhancement vs. frequency again shows a resonance peak at approximately at 480 Hz, similar to the value seen with malignant calcifications.

Table 1.
Table 1 shows the peak position (P) and the width of the peak at half height (W) for malignant and benign calcifications. There appears to be no significant difference between these two values for malignant and benign calcifications.
years. Patients were excluded if pathologic (lumpectomy or mastectomy) specimens contained invasive carcinoma in addition to DCIS. We used commercially available ultrasound equipment (Acuson, Mountain View, CA) with a high frequency transducer (8-15MHz). Cases were analyzed for (1) histologic grade, (2) mammographic findings, and (3) ultrasound findings.

RESULTS: 13 patients met the above criteria. All patients presented with mammographic clustered pleomorphic calcifications. These calcifications were sonographically identified in 7 (54%). Sonographically, these calcifications were present in focal clusters of 1-8 (18.2%) in size, with a mean size of 3.8 (8.44) mm. Patients had the following DCIS histologic grades: grade 1/61% high, grade 2/22% intermediate, grade 3/11% low. Cases with grade 2 or 3 disease represented 9/69% (grade 2) and 1/63 (grade 3) disease. There was no statistically significant difference (p>0.05). Sonographic analysis of the 3 disease categories had the following characteristics: mean size was 12.1 mm. All cases had had malignant margins. Shape was irregular in all cases. The cases had the following echogenicity: hypoechoic, 2 heterogeneous. Six cases were hypoechoic (2 had edge shadowing, 2 had shadowing adjacent to skin, 2 had lesions obscuring the mass). Four cases were associated with architectural distortion and spiculation.

CONCLUSIONS: High resolution ultrasound can detect a high rate (94%) of DCIS calcifications and is frequently (65%) associated with other sonographic abnormalities (located ducts or mass). Furthermore, grade 2 DCIS was much more likely to display these additional abnormalities than non-high grade DCIS (p<0.005). Further investigation is needed to determine whether these sonographic findings may add to mammographic prediction of breast malignancy or provide information to direct sonographic biopsy guidance for patients who cannot tolerate mammographic biopsy positioning (BEH, DMR, and SMB test equipment for Acuson Corp., Mountain View, CA).

1179 · 2:57 PM
The Specificity of Combined Mammographic and Ultrasonographic Evaluation of Palpable Lumps and Palpable Thickening
O.B. Kopans, MD, Boston, MA · R.H. Moore, AB · P.J. Sliomek, MD, MPH · E.D. Yeh, MD · O.A. Hall, MD · K.A. McCamby, MD
PURPOSE: To determine the likelihood of breast cancer when both the mammogram and ultrasound show normal appearing breast tissue in women referred with a clinically palpable lump or thickening.

METHOD AND MATERIALS: The computerized records of the Breast Imaging Division of the Massachusetts General Hospital were reviewed to determine the number of women evaluated between 1/1/95 and 12/31/98; the number of these who were referred for evaluation of a palpable lump or thickening, the number of these who had a normal mammogram and ultrasound, the number of these who had a breast biopsy and the number of these who proved to have cancer.

RESULTS: 37,301 women were evaluated. 3,323 had been referred for a palpable lump or thickening. 320 had normal appearing mammograms and ultrasound studies. 67 of these had breast biopsies. 1 cancer was diagnosed.

CONCLUSIONS: If the mammogram and ultrasound show completely normal breast tissue, the likelihood of a palpable finding being malignant is extremely low.

1180 · 3:06 PM
Targeting and Core Biopsy of Breast Microcalcifications under Ultrasound Using Acoustic Resonance
S.P. Weensten, MD, Philadelphia, PA · E.F. Conant, MD · J. Patton, BS, MS · C.M. Sengal, PhD
PURPOSE: We have developed a novel method by which calcium particles can be brought to acoustic resonance and visualized with the aid of power Doppler ultrasound. The goal of this study is to demonstrate that breast microcalcifications detected with this method can be core biopsied under ultrasound guidance in gelatin phantom and tissue models.

METHOD AND MATERIALS: Our previous studies have demonstrated that breast microcalcifications may be detected using acoustic resonance imaging. Gelatin phantoms were made with calcium carbonate particles (10-800um). The particles were evaluated with power Doppler imaging while resonating the particles at 30-500 Hz. At maximum color visibility the particles were targeted and biopsied using a 14g core needle. The 14g needle was subcutaneously placed to demonstrate that the area of color detection represented the calcium particles. The images were video-recorded and analyzed quantitatively for mean color level (MCL), % area of color (CA) and color weighted fractional area (WFA). The experiments were repeated using turkey phantom models with embedded calcium particles. RESULTS: The plot of MCL vs. WFA was flat. At 500 Hz, the particles were not present. At 30-500 Hz, the most color was present at a MCL of approximately 0.005. The color was not detected at 1000 Hz. At this frequency, color visualization of the particles was observed and enabled localization and biopsy of the microcalcifications under ultrasound guidance.
CONCLUSIONS: Data using phantoms demonstrated that microcalcifications may be targeted under ultrasound with power Doppler and acoustic resonance allowing for dynamic ultrasound guided core biopsy.

1151 • 3:15 PM
Ectogenic Breast Carcinomas
L. A. McCarthy, MD, Reppish, PA, K. M. Harris, MD
PURPOSE: Prior studies, including Stones et al.'s landmark article (Radiology 1986), have suggested that dense and uniform hyperechogenicity allow a breast mass to be classified as benign and avoid biopsy. Contrary to this, we describe the features of a series of ectogenic breast carcinomas.
METHOD AND MATERIALS: Between 1/91 - 1/98, one of the authors (KM) collected a series of 10 ectogenic breast carcinomas from her daily practice. The following features of these were retrospectively reviewed: margin (ill-defined/circumscribed), palpability, shape (oval, round, irregular, lobulated), echotexture (homogeneous, heterogeneous), sound attenuation, reflectance, shadow, vascularity, and whether the lesion was “saw-toothed” or “wider-than-tall.” All lesions underwent surgical excision and were proven to be carcinomas.
RESULTS: There were nine cases of infiltrating ductal carcinoma and one case of ductal carcinoma in situ. All lesions had ill-defined margins. 8/10 lesions were palpable. 8/10 had an irregular shape, and 9/10 were homogeneously hyperechogenic. 4/10 lesions caused posterior acoustic shadowing while 6/10 lesions had neither posterior acoustic shadowing nor enhancement. 1/10 lesions was “saw-toothed” while 5/10 lesions were “wider-than-tall” and 4/10 lesions were of approximately equal height and width.
CONCLUSIONS: Although the great majority of uniformly hyperechogenic breast masses are benign, breast images must remain aware that there are exceptions, and that hyperechogenic breast cancers, though rare, exist. Features associated with hyperechogenic cancers were ill-defined margins, irregular shape and being palpable.

Wednesday Afternoon • Room S403B

Radiation Oncology (Sarcoma, Biology)

1152 • 2:30 PM
Image Fusion: A New Approach for Using MRI Imaging in Brachytherapy
R. C. Kleinpen, MD, Heidelberg, Germany, W. Harms, MD, H. A. Grabowski, PhD, R. W. Hensley, PhD, M. Treiber, MD, M. Wannenmacher, MD, DDS
PURPOSE: Accurate definition of tumor volume is essential for brachytherapy treatment planning. CT accurately localizes sources, but does not visualize the tumor. MRI has several imaging benefits such as improved tissue definition and un restricted multiplanar imaging but does not visualize the source dwell positions. The aim of this study was to assess the possibilities of a new developed system generating synthetic image modalities from matched and fused CT and MRI data in brachytherapy planning.
METHOD AND MATERIALS: The system presented consists of five modules serving for viewing, matching, segmentation, fusion and validation. The automatic registration takes about one minute for two data sets of about 50 slices. Depending on slice thickness the registration algorithm is able to register MRI and CT data in over 95% of the cases without any interaction with an average accuracy of the calculated transformation below 1 mm. The segmentation module consists of two modules, the volume module for three dimensional region growing to segment connected structures and the slice module for interactive slice by slice segmentation of the superimposed previously registered images. Tumor structures or source dwell positions can be segmented in any plane in one imaging modality and then copied into the other. Reconstruction of the fused images in different planes offer the possibility to validate the matching accuracy.
RESULTS: A patient with liposarcoma in the thigh showed a positive tumor margin to the sigmoidal nerve. Intraperitoneal the decision was made to implant brachytherapy guides for postoperative boost radiation on the tumor bed rather than resecting the nerve to preserve the function of the leg. CT accurately localizes source positions but the nerve as the primary target volume was hardly detectable. The MRI visualized the nerve but the source positions could not be detected. The sigmoidal nerve was segmented in MRI and the gray values or the segmented structure were copied in the CT image. The source positions are segmented via seed points in the CT. The position of the nerve can now be visualized in comparison to the source dwell positions. The synthetic images are added into the brachytherapy planning system. The CT-based brachytherapy planning would have resulted in a miscalculation of the target volume while the synthetic images lead to an optimization in therapy planning.
CONCLUSIONS: These preliminary data show that image registration and fusion is feasible for brachytherapy planning. In the future we will fuse the synthetic images in brachytherapy planning lead to an optimized treatment planning.

1184 • 2:48 PM
MRI May Be Used to Evaluate the Effects of Tumor Oxygenating Agents on Radiosensitivity
H. A. Al-Hattab, AB, Chicago, IL, M. A. Zamora, BS, B. L. Fish, BS, J. E. Moulder, PhD, G. S. Karczewski, PhD
PURPOSE: Previous work suggested that MRI could provide noninvasive, high-resolution measurements of the effects of tumor oxygenating treatments (TOP) on tumor blood and tissue oxygen levels. Here we evaluate whether MRI can be used to target TOP according to their effects on increasing tumor response to radiation.
METHOD AND MATERIALS: Studies of BALB/c mouse mammary carcinomas (WAC) have shown that the administration of a tumor oxygenation mimetic, CO (6.5% O2/5% CO2) breathing, significantly increased radiosensitivity relative to either treatment alone. We investigated whether MRI correctly predicts the relative efficacy of treatment with carbogen or PFC alone compared to treatment with PFC - carbogen in a tumor model. We used MRI signal linewidth, i.e., blood-oxygen-level-dependent (BOLD) contrast, as an index of change in tissue oxygen tension. Decreases in linewidth indicate increases in tumor oxygenation, to assess the spatial distribution of response, maps of pixels that responded to treatment were generated for each tumor.
RESULTS: Average MRI signal linewidth decreased by 25% during carbogen breathing, by 23% during PFC - air, and by 45% during PFC - carbogen (n = 7). The change caused by PFC - carbogen was significantly greater than the change caused by either treatment alone (p < 0.01) in paired t-tests demonstrating that the combined treatment increases tumor oxygenation more than PFC or carbogen alone. Percentages of responding pixels in each category were averaged over all 7 tumors. 21% of the total pixels responded to treatment with carbogen while 32% of pixels
Microcalcifications in Breast Tissue Phantoms Visualized with Acoustic Resonance Coupled with Power Doppler US: Initial Observations

Calcium carbonate particles embedded in gelatin and turkey breast tissues were visualized with acoustic resonance imaging and power Doppler ultrasonography. Sonography revealed that the region of color level detection corresponded to the location of the calcium carbonate particles. Correlation between color level detection and the location of the particles was confirmed on radiographs of the specimens obtained at core needle biopsy performed through the region of color level detection.

It is estimated that ductal carcinoma in situ represents 20%–30% of breast carcinomas detected at screening mammography (1). Since the 1970s, the detection rate of ductal carcinoma in situ has steadily increased with the wider use of screening mammography. The detection rate for women younger than 50 years was 2.3 cases per 100,000 women in the 1970s and increased to 6.2 cases per 100,000 women by the 1990s (2). More dramatically, in the group of women older than 50 years, the rate has increased from 14.3 to 54.6 per 100,000 (2) in the same period.

Frequently, ductal carcinoma in situ manifests only as calcifications without an associated mass (3). Because ductal carcinoma in situ represents as much as half of mammographically depicted cancers (3), it would be efficacious to image microcalcifications with as many modalities as possible to allow flexibility in imaging-guided biopsy. Currently, even with all the technical advances in breast imaging, including magnetic resonance imaging, scintigraphy, and ultrasonography (US), mammography is the only reliable method to help detect, characterize, and localize microcalcifications for biopsy.

We have developed a technique in which acoustic resonance is used to visualize microcalcifications. Acoustic resonance is used on the basis of the size of the microcalcifications and the binding strength with the surrounding tissues in which they are embedded. When subjected to a wide frequency range, particles of different sizes will resonate at different frequencies given the same binding environment. "Tuning" to the appropriate frequency range would make it possible to selectively visualize microcalcifications of varying sizes. The purpose of our study was to evaluate US coupled with acoustic resonance to demonstrate small calcified particles for targeting and biopsy.

I Materials and Methods

Phantoms

Gelatin phantoms were constructed by using a mixture of water, gelatin, and glycerol with uniformly dispersed oil-lecithin emulsion (particle diameter, 0.2–2.0 μm). The suspension was placed in a mold and refrigerated until a solid state was reached. Calcium carbonate particles 400–800 μm in diameter were suspended in the gelatin phantom in an area 1.0–1.5 cm in diameter. The phantom was kept refrigerated until use, at which time it was removed from the mold. Ten phantoms were made.

Twelve tissue phantoms were made from boneless turkey breast. The turkey breast was dissected along the tissue planes, and 400–800-μm-diameter calcium carbonate particles mixed in acoustic cou-
Imaging: US Coupled with Acoustic Resonance

Imaging was performed (S.P.W. or C.S., with the assistance of J.A.P.) by using gray-scale and power Doppler US coupled with acoustic resonance (Logic 700; GE Medical Systems, Milwaukee, Wis.). A variable 10–13-MHz transducer was used. All the gelatin and tissue phantoms were scanned. The phantoms were imaged with B-mode US and power Doppler US, while the particles were resonated from 50 to 500 Hz. The frequency that corresponded to the maximum color level detection was noted.

The frequency used to excite the particles into resonance was emitted from a thin (5-mm) lightweight (13-g) piezoelectric speaker element with the capability to transmit low-frequency vibrations from 50 Hz to 2 kHz. The device was housed in a fiberglass case. The piezoelectric vibrator was designed for 8 Ω of impedance to be driven by an audio amplifier. The vibrator was specially designed by the biomedical medical instrumentation group at our university; they specialize in building custom instruments, and this vibrator was approved for human use by the university institutional review board. The disk was coupled with US gel (E-Z-Gel; E-Z-Em, Westbury, NY) and placed adjacent to the tissue to be scanned. The images were videotaped for analysis.

Image Analysis

The videotaped images were subsequently analyzed. Imaging time for each scan was approximately 2 minutes. At the video frame rate of 30 images per second, approximately 120 × 30, or 3,600, images were generated. An algorithm was developed to reduce the data by sorting the images on the basis of the frequency of the vibration. These images were subsequently used in a computer program developed in our laboratory to analyze the vibrational response of the microcalcifications.

Color was analyzed for each image: Mean color level, percentage of fractional area of color, and color-weighted fractional area were computed. To determine the mean color level, the color palate on the image was read by the computer and divided equally on a scale of 0–100. With this scaling system, the computer constructed a lookup table for hue, saturation, and brightness values for the colors in the palate bar. Next, the computer identified colored pixels on the image and used the lookup table to assign a color value to each pixel in the region of interest. The color level of the pixels in the region of interest was summed and divided by the number of color pixels to calculate the mean color level. The percentage of fractional area of color was defined as the area covered by colored pixels divided by the area of the region of interest, multiplied by 100%. Color-weighted fractional area was defined as the mean color level multiplied by the percentage of fractional area of color, all divided by 100, and indicated the presence of net motion in the region of interest. Each parameter was plotted with respect to the frequency range.

Biopsy

All the gelatin and tissue phantoms were scanned from 50 to 500 Hz. The location where the maximum color level was detected and the corresponding frequency were noted. To demonstrate that the area of color level detection in the phantoms corresponded to the calcium carbonate particles, biopsy was performed through the region of color detection level. A 14-gauge disposable core biopsy needle (Monopty Biopsy Instrument; Bard, Covington, Ga) was used to perform biopsy in the region of interest during in real-time scanning by using power Doppler US coupled with acoustic resonance imaging. Approximately seven to 10 samples were obtained for each phantom. The core biopsy samples were placed in a core needle biopsy specimen container (Beekley, Bristol, Conn.). Radiography of the specimens (alpha RT; Instrumentarium, Milwaukee, Wis) was performed at 22 kVp and 6 mAs, with magnification of 1.8, to confirm that the biopsy target, the region of depiction at
power Doppler US, corresponded to the calcium carbonate particles.

I Results

Gelatin Phantom

The calcium carbonate particles were readily visible as white particles in all the gelatin phantoms at B-mode US. Figure 1a shows one of the phantoms. The calcium carbonate particles were readily visible in all 10 specimens. With power Doppler US evaluation, a gradual increase in color level detection was seen in the region of the calcium carbonate particles (Fig 1b). The maximum detection level was reached and then plateaued at frequencies of 200–380 Hz (Fig 1c). This plateau was followed by a gradual decrease in the color level detection (Fig 1d). This trend in color level detection was seen in all the gelatin phantoms evaluated. As seen in Figure 1, there was a minimal color level detected outside the region of the calcium carbonate particles. Graphs of mean color level, percentage of fractional area of color, and color-weighted fractional area versus frequency demonstrate the results seen on the static images in Figure 1, that is, a gradual increase in depiction level, followed by a decrease (Fig 2).

Turkey Phantom

The turkey phantom was also imaged with B-mode US and power Doppler US. In all 12 turkey phantoms, unlike in the gelatin phantoms, the calcium carbonate particles were not as readily visible at B-mode US before use of acoustic resonance imaging coupled with power Doppler US (Fig 3a). When the turkey phantom was imaged with power Doppler US coupled with acoustic resonance imaging, a gradual increase in color level detection was seen, a peak was reached, and a decrease followed (Fig 3b–3d). Similar findings in power Doppler detection were seen in all the turkey phantoms evaluated.

Biopsy Results

The radiograph of a specimen is shown in Figure 4. All radiographs of the specimens depicted the calcium carbonate particles in the specimens.

I Discussion

The concept of acoustic resonance imaging is relatively new. There is a large body of literature that reports various modes of US imaging to characterize motion and elasticity of soft tissues induced by external vibrations or due to natural motion (4–8). None of these authors deal with the issue of detection of microcalcifications, nor do they use the concept of acoustic resonance as a means to identify the properties of inhomogeneities in the tissue. Our goal was to be able to visualize the resonance motion of inhomogeneities relative to that of the surrounding tissue.

The 400–800-μm calcium carbonate particles were excited into resonance when exposed to low-frequency vibrations from 50 to 500 Hz. The vibrations were of low amplitude, with no known harmful side effects. When the particles were in resonance and combined with power Doppler US, the calcium carbonate particles were expected to demonstrate a color level relative to the degree of resonance. At maximal resonance, there would be maximal color level detection. The frequency at which maximal resonance is demonstrated depends on the size of the microcalcifications and the binding properties between the tissue
Figure 3. (a) US scan depicts a turkey breast phantom. Calcium carbonate particles are not as obvious as they are in the gelatin phantom. Color is represented in gray scale. (b–d) Power Doppler US scans coupled with acoustic resonance depict some color pixels in the region of the calcium carbonate particles (arrows) at 155 Hz (b); the number of color pixels continues to increase at 270 Hz (c); followed by a decrease at 355 Hz (d).

and the microcalcifications, which is demonstrated with the phantom models as shown in Figures 1 and 3. Graphs of the mean color level, percentage of fractional area of color, and color-weighted fractional area versus frequency demonstrate an increase in power Doppler US depiction level, followed by a plateau, then a decrease. The plateau, seen in the curves, may represent the particles of different sizes reaching resonance over a different range of frequencies.

In the simple gelatin phantom, the microcalcifications were visible without power Doppler US, which allowed direct correlation between the location of the particles and the area of color-level depiction. Unlike the homogeneous background of the gelatin phantom, the turkey phantom more closely approximates human breast tissue, with muscle striations and specular interfaces similar to the appearance of Cooper ligaments and other specular reflectors in human glandular tissue. Therefore, the location of the calcium carbonate particles is not obvious until power Doppler US coupled with acoustic resonance imaging is used. Core biopsy was performed in the area where color level was detected in both types of phantoms, and radiography of the specimens was performed. Findings on the radiographs of the specimens confirmed that the region of color level detection corresponded to the calcium carbonate particles.

There are multiple potential applications for this method. Once the microcalcifications are detected at US, biopsy could be performed with this dynamic imaging guidance. Results of various studies have demonstrated the cost-effectiveness of core needle biopsy compared with excisional biopsy (9–12) or US-guided core needle biopsy versus stereotactic core needle biopsy (13). With the ability to perform biopsy with vacuum assistance and US guidance, this method would allow performance of biopsy in calcifications by using US and possibly dynamic visualization of the extraction of calcifications, which would allow con-
firmation of targeting and, conceivably, a procedure time shorter than that for stereotactic core needle biopsy. Currently, sonographically guided biopsy is not always feasible because calcifications cannot always be definitively visualized with US alone.

We have shown that power Doppler US coupled with acoustic resonance imaging has the ability to demonstrate microcalcifications in phantom models. Future studies are needed to develop this technique for use in patients. US evaluation may be performed in women prior to conventional stereotactic or excisional biopsy. Correlation between particle size, peak resonance, binding properties, and pathologic results may provide beneficial information. Analysis of acoustic resonance properties may help determine the strength of adhesion between the calcium deposits and the surrounding tissues. Such forces are likely to be related to the molecular characteristics of the deposits and the mechanisms that cause calcifications to develop at specific sites in the tissues. Measurement of adhesive forces with acoustic resonance, in conjunction with morphologic parameters detected at mammography, may help characterize calcifications.

References