Incidentally Detected Enhancing Breast Lesions on Chest Computed Tomography

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Objective: To evaluate the nature and imaging appearance of incidental enhancing breast lesions detected on a routine contrast-enhanced chest CT.

Materials and Methods: Twenty-three patients with incidental enhancing breast lesions on contrast-enhanced chest CT were retrospectively reviewed. The breast lesions were reviewed by unenhanced and enhanced CT, and evaluated by observing the shapes, margins, enhancement patterns and backgrounds of breast lesions. A histopathologic diagnosis or long-term follow-up served as reference standard.

Results: Sixteen (70%) patients had malignant breast lesions and seven (30%) had benign lesions. In 10 patients, the breast lesions were exclusively detected on contrast-enhanced CT. Using unenhanced CT, breast lesions with fibroglandular backgrounds were prone to be obscured \((p < 0.001)\). Incidental primary breast cancer showed a non-significant trend of a higher percentage irregular margin \((p = 0.056)\). All of the four incidental breast lesions with non-mass-like enhancement were proven to be malignant.

Conclusion: Routine contrast-enhanced chest CT can reveal sufficient details to allow for the detection of unsuspected breast lesions, in which some cases may be proven as malignant. An irregular margin of incidental enhancing breast lesion can be considered a suggestive sign of malignancy.

Index terms: Breast; Chest; Computed tomography (CT); Incidental breast lesions

INTRODUCTION

Mammography, breast sonography, and magnetic resonance imaging are currently the preferred examination methods for the detection and characterization of breast diseases. Although dedicated breast computed tomography (CT) had been investigated for potential clinical applications in the evaluation of breast (1), it was not considered the primary method to evaluate specific breast lesions.

Chest CT is routinely used for diseases of the lung, mediastinum, pleura, chest wall, and diaphragm, but not breast tissue. However, the entire breast tissue usually appears on a chest CT. With the increased use of chest CT, either for diagnostic or screening purposes, incidental breast lesions are increasingly being encountered. In some cases, CT may be the first imaging modality to demonstrate a breast lesion. Incidental breast lesions detected on unenhanced or contrast-enhanced CT of chest have been discussed in a few recent English literatures (2–8), which
evaluated the incidence and pathologic outcomes of incidental breast lesions. We undertook a study emphasizing the discovery of incidental breast lesions with enhancement on routine contrast-enhanced chest CT, and evaluated the nature and imaging appearance of the incidental enhancing breast lesions.

**MATERIALS AND METHODS**

**Patients**

Our study has been approved by the institutional review board. All patients provided written informed consent for the diagnostic procedures performed.

From October 2006 to June 2009, a total of 2250 patients underwent routine contrast-enhanced chest CT at our institution, of which, 34 were identified as having enhancing breast lesions. Of the 34 patients, seven cases were excluded from our study due to a known history of breast disease prior to CT (breast cancer, n = 6; breast abscess, n = 1); another four patients were excluded due to the absence of additional diagnostic confirmation. Ultimately, a total of 23 patients, all of which were female with a mean age of 59.2 ± 19.5 years (range, 22–85 years), were included as part of this retrospective study.

Clinical indications for chest CT in the 23 patients included a follow-up study in patients with lung cancer (n = 7), abnormal chest radiograph (n = 6), follow-up study or a staging work-up for non-pulmonary malignancies (n = 5; thyroid cancer, tongue cancer, lymphoma, renal cell carcinoma, cervical cancer - one for each type), abnormal nuclear medicine bone scan (n = 1), chest trauma (n = 2), a suspicious case of aortic aneurysm (n = 1), and hemoptysis (n = 1).

**CT Technique**

The chest CT was performed in the supine position, using a 64-detector row scanner (Brilliance; Philips Medical Systems, Cleveland, OH). Both unenhanced and contrast-enhanced scans were performed and the standard protocol at our institution included an unenhanced helical CT from the lung apices through the adrenal glands, using the following imaging parameters: a section thickness of 0.625 mm, a rotation speed of 0.75 second, a pitch of 1.05–1.25, and 120 kVp; effective tube current × time product ranged between 150 and 200 mAs. Subsequently, a contrast-enhanced helical CT was performed with the intravenous administration of 100 mL iopromide (Ultravist 300; Schering, Berlin, Germany) using a mechanical power injector at a rate of 2.0–3.0-mL/sec. The scanning was performed 50 seconds after the injection of the contrast medium. The imaging parameters were identical to those used for unenhanced CT.

In our study, only transverse images were reviewed, by reconstruction using a 5-mm thickness and 5-mm increments. The images were obtained by using a standard soft-tissue algorithm (window width, 350 Hounsfield unit [HU]; level, 40 HU) and a retrospective lung algorithm (window width, 1000 HU; level, –700 HU).

**Image Interpretation**

Of the 23 patients included in the study, all CT images were retrospectively reviewed by two experienced radiologists (with 11 and 21 years of experience in the interpretation of chest CT, respectively) in consensus. The reviewers had knowledge about the presence of breast lesions on chest CT; however, they were blinded to the locations and diagnosis of the breast lesions.

In each patient, unenhanced CT was reviewed to determine whether the breast lesion could be seen. A contrast-enhanced CT was conducted immediately after interpretation of unenhanced CT. Reviewers were requested to describe the size, location, shape, margin characteristics, and enhancement patterns of the enhancing breast lesions detected on contrast-enhanced CT. An enhancing breast lesion referred to a breast lesion with attenuation higher than that of normal breast glandular tissue on contrast-enhanced CT.

The shape of the incidental enhancing breast lesion was described as a round, ovoid, lobulated, and irregular mass or non-mass. A breast lesion with a non-mass shape, which was also known as a non-mass-like enhancement, meant that the lesion could only be detected on contrast-enhanced CT, which appeared as regional enhancement of the breast. The area of enhancement did not appear as a mass-like configuration, in which the internal features could be described as having a discretely different appearance from that of normal gland parenchyma. The margin of the incidental enhancing breast lesion was described as well-circumscribed, irregular, and spiculated.

Reviewers were also requested to determine the background of each breast lesion; namely, adipose or fibroglandular tissues. An adipose tissue breast lesion meant that more than half of the border was surrounded by adipose tissue. Similarly, a fibroglandular tissue breast
lesion referred to a lesion with more than half its border surrounded by fibro glandular tissue.

Final Diagnosis and Follow-Up Studies

All patients subsequently underwent sonography of the breast or mammography for the incidental breast lesions. Sonographically-guided core-needle biopsy or surgery was performed for definitive diagnosis in 20 patients. The diagnoses in the three remaining patients were determined on the basis of long-term follow-up CT or sonography for at least 12 months.

Statistical Analysis

The statistical analysis was performed with commercially available software (SPSS version 10.0, SPSS, Chicago, IL). Continuous variables were expressed as mean ± standard deviation (SD); and the Student's t-test was used for comparison of the two variables. The two-tailed Fisher's exact test was used for analysis of the nominal variables. A p value of 0.05 was chosen as the threshold for statistical significance.
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RESULTS

Pathologic Outcomes

Sixteen (70%) of the 23 patients had incidental enhancing breast lesions which were subsequently proven to be malignant. Of the 16 patients, nine had invasive ductal carcinoma (IDC), four patients had ductal carcinoma in situ (DCIS), and one patient had two lesions located in bilateral breasts. The remaining three patients had breast metastases from lung cancer (n = 2) or renal cell carcinoma (n = 1) (Table 1). Sonographically-guided core-needle biopsy and/or mastectomy confirmed the diagnoses in 15 patients. The remaining patient with renal cell carcinoma underwent a sonography of the breast, which disclosed multiple breast tumors. However, she did not have further diagnostic work-up for the breast lesions due to her advanced stage of disease including multiple metastases of the lung and mediastinum. With chemotherapy, the follow-up CT performed six and 12 months later, showed complete remission of the breast lesions. The incidental enhancing breast lesions in this patient were assumed as metastases from renal cell carcinoma.

Our records showed a total of 2250 patients that underwent routine contrast-enhanced CT of the chest during the study period. Malignant breast tumors were incidentally detected in 16 (0.7%) patients, whereas primary breast cancer was incidentally detected in 13 (0.6%) patients.

Seven (30%) patients had incidental enhancing benign breast lesions, including four fibroadenomas, one surgical scar, one breast abscess, and one lymphoid hyperplasia.

Table 1. Pathologic Outcomes and Imaging Appearance of 16 Patients with Malignant Incidental Enhancing Breast Lesions on Chest CT

| Patient | Age | Number of Lesion | Size (cm) | Background of Lesion | Detected on Unenhanced CT | Shape | Margin | Enhancement | Diagnosis |
|---------|-----|------------------|-----------|----------------------|--------------------------|-------|--------|-------------|-----------|
| 1       | 85  | 1                | 1.5       | Adipose              | Yes                      | Lobulated | Well-circumscribed | Heterogeneous | IDC       |
| 2       | 80  | 1                | 1.4       | Fibroglandular       | No                       | Lobulated | Irregular | Heterogeneous | IDC       |
| 3       | 46  | 1                | 0.9       | Fibroglandular       | No                       | Lobulated | Irregular | Heterogeneous | IDC       |
| 4       | 26  | Multiple         | 1.5       | Adipose              | Yes                      | Round    | Well-circumscribed | Heterogeneous | Metastases |
| 5       | 76  | 1                | 1.7       | Adipose              | Yes                      | Lobulated | Irregular | Heterogeneous | IDC       |
| 6       | 74  | 1                | 1.1       | Adipose              | Yes                      | Lobulated | Irregular | Homogeneous  | IDC       |
| 7       | 72  | 1                | 1         | Fibroglandular       | No                       | Non-mass | Irregular | Homogeneous  | DCIS      |
| 8       | 69  | 1                | 0.6       | Fibroglandular       | No                       | Lobulated | Irregular | Homogeneous  | IDC       |
| 9       | 77  | 1                | 1.2       | Adipose              | Yes                      | Lobulated | Irregular | Homogeneous  | IDC       |
| 10      | 81  | 1                | 1.5       | Adipose              | Yes                      | Lobulated | Well-circumscribed | Heterogeneous | IDC       |
| 11      | 49  | 2                | 2.7       | Fibroglandular       | No                       | Non-mass | Irregular | Heterogeneous | DCIS      |
| 12      | 22  | Multiple         | 1         | Fibroglandular       | No                       | Non-mass | Irregular | Heterogeneous | DCIS      |
| 13      | 53  | Multiple         | 0.8       | Fibroglandular       | No                       | Round    | Well-circumscribed | Homogeneous  | Metastases |
| 14      | 78  | 1                | 1.9       | Fibroglandular       | No                       | Lobulated | Well-circumscribed | Heterogeneous | IDC       |
| 15      | 77  | 1                | 2.3       | Adipose              | Yes                      | Lobulated | Irregular | Homogeneous  | DCIS      |
| 16      | 53  | 1                | 1.3       | Fibroglandular       | No                       | Non-mass | Irregular | Homogeneous  | DCIS      |

Note.— DCIS = ductal carcinoma in situ, IDC = invasive ductal carcinoma.

In case of multiple lesions, only size or background of largest lesion was recorded.
Of the four patients with fibroadenomas, two had multiple lesions in bilateral breasts. Four of the seven patients underwent a sonographically-guided core-needle biopsy to confirm the diagnosis. In the remaining three patients, a surgical incision and drainage was performed for one breast abscess. A comparison with prior chest CT at an interval of more than 12 months showed stability of the breast lesions in two patients with fibroadenoma and surgical scar, respectively.

The mean size of the malignant enhancing breast tumors was 1.4 ± 0.54 cm (range, 0.6–2.7 cm), while the mean size of the benign enhancing breast lesions was 1.5 ± 1.0 cm (range 0.5–3.4 cm). There was no significant difference between the mean size of malignant and benign breast lesions (p = 0.68). All of the nine incidental IDCs were less than 2.0 cm (mean size, 1.3 ± 0.13 cm; range, 0.6–1.9 cm), and at stage T1, which is based on the TNM staging system for breast cancers.

The mean age of patients with incidental malignant enhancing breast tumors was 63.6 ± 19.7 years, whereas the mean age of patients with incidental benign enhancing breast lesions was 49.1 ± 15.9 years. There was no significant difference between the mean age of patients with malignant breast tumors and those with benign breast lesions (p = 0.06). However, the mean age of patients with incidental primary breast cancer (70.5 ± 12.8 years) was significantly higher than that of patients with incidental benign breast lesions (p = 0.002).

### Imaging Appearance of Incidental Enhancing Breast Lesions

With the unenhanced CT of chest, breast lesions could be detected in 13 (57%) of the 23 patients, which included seven (44%) of 16 malignant cases and six (86%) of seven benign cases. All of the lesions detected on unenhanced CT were enhanced on contrast-enhanced CT.

A total of 10 patients had incidental breast lesions exclusively detected on contrast-enhanced CT, and nine (90%) of the 10 patients had malignant breast lesions.

The backgrounds of incidental enhancing breast lesions were adipose tissues in 11 patients, and fibroglandular tissues in the remaining 12 patients (Figs. 1, 2). For the 13 patients with incidental breast lesions detected on unenhanced CT, only two (15%) had breast lesions with fibroglandular backgrounds; however, all (100%) of the 10 patients with incidental breast lesions that were exclusively detected on contrast-enhanced CT had breast lesions of fibroglandular backgrounds. In those patients with incidental breast lesions exclusively detected on contrast-enhanced CT, a significantly high percentage of patients had incidental breast lesions of fibroglandular backgrounds (p < 0.001).

Of the 16 patients with malignant breast tumors, heterogeneous enhancement of tumors were observed in 10 (63%) patients, and homogeneous enhancement was noted in the remaining six (38%). Of the seven patients with benign breast lesions, three (43%) showed heterogeneous enhancement and four (57%) showed homogeneous enhancement.

### Table 2. Pathologic Outcomes and Imaging Appearance of Seven Patients with Benign Incidental Enhancing Breast Lesions on Chest CT

| Patient | Age | Number of Lesion | Size (cm) | Background of Lesion | Detected on Unenhanced CT | Shape | Margin | Enhancement | Diagnosis              |
|---------|-----|-----------------|-----------|----------------------|--------------------------|-------|--------|-------------|------------------------|
| 1       | 66  | 1               | 0.5       | Fibroglandular       | No                       | Lobulated | Irregular | Homogeneous | Fibroadenoma             |
| 2       | 23  | Multiple        | 2.4       | Adipose              | Yes                      | Lobulated | Well-circumscribed | Heterogeneous | Fibroadenoma             |
| 3       | 62  | Multiple        | 1.2       | Fibroglandular       | Yes                      | Lobulated | Well-circumscribed | Homogeneous | Fibroadenoma             |
| 4       | 60  | 1               | 1         | Adipose              | Yes                      | Irregular | Spiculated     | Homogeneous | Surgical scar           |
| 5       | 33  | 1               | 3.4       | Fibroglandular       | Yes                      | Lobulated | Well-circumscribed | Heterogeneous | Abscess of breast       |
| 6       | 50  | 1               | 0.8       | Adipose              | Yes                      | Lobulated | Well-circumscribed | Heterogeneous | Lymphoid hyperplasia    |
| 7       | 50  | 1               | 1.5       | Adipose              | Yes                      | Ovoid   | Well-circumscribed | Homogeneous | Fibroadenoma             |

Note. — In case of multiple lesions, only size or background of largest lesion was recorded.
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enhancement. There was no significant difference in the percentage of heterogeneous or homogeneous enhancement between the malignant and benign incidental enhancing breast lesions (p = 0.65).

Incidental enhancing breast lesions with a lobulated shape were noted in 15 patients, in whom 10 (67%) had malignant breast lesions, and five (33%) had benign breast lesions (Tables 1, 2). A total of four incidental enhancing breast lesions were categorized as non-mass breast lesions, and all of them were subsequently confirmed to be malignant breast tumors (Fig. 1) (Table 1).

Irregular or spiculated margins were observed in 11 (79%) of the 14 incidental enhancing primary malignant breast tumors, and two (29%) of seven benign breast lesions (Tables 1, 2). Incidental primary breast cancer showed an insignificant trend of a higher percentage of irregular or spiculated margin (p = 0.056) (Fig. 2). Three patients of breast metastases showed multiple breast lesions that were round in shape and had a well-circumscribed margin.

Well circumscribed enhancing breast lesions were noted in five (71%) of seven patients with benign breast lesions. The incidental enhancing benign breast lesion with a spiculated margin was proven to be a surgical scar from a past history of resection for fibroadenoma of the breast, which was obtained and the long-term stability of this lesion was confirmed at follow-up CT. The benign breast lesion with an irregular margin was proven to be a 0.5-cm fibroadenoma by a sonographically-guided core-needle biopsy.

DISCUSSION

Although a routine chest CT was usually performed for indications other than breast disease, and the technology was not optimized for breast imaging, the incidental breast lesions could be occasionally detected. In some cases, incidental breast cancer could be detected on chest CT. Swensen et al. (9) reported three (0.4%) cases of breast cancer from 735 women who underwent unenhanced low-dose chest CT for lung cancer screening. Shojaku et al. (3) reported four (0.4%) cases of incidental breast cancer or metastatic breast carcinoma from 1008 patients who underwent unenhanced chest CT. In a recent study, eight (2%) incidental primary breast cancers were detected in 432 chest CT scans (6). We included only incidental enhancing breast lesions in our study, which amounted to 16 (0.7%) patients of 2250 who underwent routine contrast-enhanced chest CT.

A few researches have evaluated the pathologic outcomes of incidental breast lesions and determined the CT features of incidental breast lesions suggestive of malignancy (6-8). In these studies, 24–32% of incidental breast lesions were subsequently proven to be malignant (6–8). Our study showed a much higher rate (70%) of malignancy. This higher result may be partly due to a selection bias from the retrospective study design, and the small number of cases in our study. However, it should be noted that we included only enhancing breast lesions in our study, and therefore some incidental breast lesions which were not enhanced on chest CT may not be included. Porter et al. (8) reported that 17 (50%) normal breast glandular “pseudo-masses” and simple cysts, which may not be enhanced on contrast-enhanced CT, in a total of 34 incidental breast lesions.

By excluding the non-enhancing benign incidental breast lesions, the rate of malignant incidental breast lesions likely causes an increase in the rate of malignancy.

The imaging protocol of routine chest CT may be a reason for the relatively small number of benign incidental breast lesions in our series. The routine contrast-enhanced chest CT was performed at 50 seconds after intravenous administration of contrast medium, and the entire scanning would be completed within two minutes after administration of the contrast medium. This period corresponded to the early arterial phase of a dynamic breast CT. Most malignant breast lesions had strong enhancement in the early arterial phase, and benign breast lesions were well-enhanced in late phase of the dynamic CT study (10, 11). As a result, breast malignancy may be incidentally detected at routine chest CT by the early enhancement of tumor, and some benign breast lesions may not be detected due to lack of sufficient enhancement in the early phase.

By using dynamic CT of breast, the reported signs of malignancy included early enhancement of breast tumors and washout or plateau patterns of the time-density curve for three to eight minutes after injection of the contrast medium (10, 11). In our study, the incidental breast lesions were detected by routine contrast-enhanced chest CT. The contrast-enhanced study was performed at 50 seconds after the injection of contrast medium, and not a dynamic study. Therefore, the analysis of the enhancement patterns at dynamic breast CT study could not apply to a routine chest CT.

The reported CT features suggestive of malignant incidental breast lesions on routine chest CT included rim enhancement of breast tumors, an irregular or spiculated
margin, and an axillary lymphadenopathy (5, 7, 8). In our study, the incidental primary breast cancer showed a higher probability of an irregular margin, with an insignificant trend. However, we have observed that incidental IDC may also demonstrate well-circumscribed margins. Therefore, it would seem reasonable to suggest that well-circumscribed incidental breast lesions should be referred for dedicated assessment, as recommended by previous authors (2–5, 8). We also noted that the age of patients with incidental primary breast cancer was higher than that of patients with incidental benign breast lesions. This may be consistent with the fact that risk of breast cancer rises with increased age.

Incidental malignant and benign breast lesions may share overlapping enhancement characteristics in our study. Heterogeneous or homogeneous enhancement was not a reliable sign to distinguish malignant from benign incidental breast lesions on chest CT. The size of incidental breast lesions was also not a reliable feature to distinguish malignancy from benignity. There was no case demonstrating axillary lymphadenopathy in our study; however, this may be due to small number of cases or early stage of disease. All of the nine incidentally detected IDC in our study were at the T1 stage of breast cancer.

Some incidental breast lesions were exclusively detected on contrast-enhanced CT in our study. The adipose or fibroglandular backgrounds of breast lesions may be one reason for neglected breast lesions on unenhanced CT. Also, all breast lesions not detected on unenhanced CT were of fibroglandular background. In these cases, there was not enough of a difference in the attenuations between a breast lesion and adjacent fibroglandular tissue; therefore, the breast lesions could not be detected on unenhanced CT.

Three patients with four incidental breast lesions showed an interesting imaging finding of non mass-like enhancement. All of the four lesions were subsequently proven to be malignant. The description “non mass-like enhancement” had been used in the interpretation of breast magnetic resonance (MR) imaging, which meant that enhancement occurred in an area of fibroglandular tissue that otherwise appeared as normal on unenhanced images (12). Non mass-like enhancement on breast MR imaging may represent DCIS, lobular cancer, mastopathic changes (focal adenosis), hormonal stimulation, or inflammatory changes (12).

Three patients with a known history of non-mammary malignancy showed multiple, well-circumscribed breast tumors. These tumors were confirmed as breast metastases. Although not common, the possibility of breast metastases should be considered in patients with multiple well-circumscribed breast nodules and non-mammary malignancy.

There were limitations in our study; first, the small number of patients making up the study population and retrospective nature of the study design may reduce the power of this study. A routine chest CT is less sensitive than dedicated breast CT or MR imaging in detecting breast lesions (10, 12). Second, it would have been preferable if all the patients had histopathologic diagnoses. However, the 12-month follow-up studies in two patients with benign breast lesions were probably sufficient for diagnosis of benignity. Widespread metastases occurred in one patient with breast metastases; therefore, only the breast sonography was performed, while the further histopathologic examination was reasonably not performed.

In conclusion, it is important for general radiologists to pay attention to the breasts on CT performed for other thoracic indications. Routine contrast-enhanced chest CT can reveal sufficient details to allow the detection of unsuspected breast lesions, in which some cases may be subsequently proven to be malignant. A low threshold for referral of patients with routine chest CT-detected incidental enhancing breast lesions to a breast clinic for assessment is advocated. Irregular margins of incidental enhancing breast lesion can be considered a suggestive sign of a malignant lesion.

REFERENCES

1. Lindfors KK, Boone JM, Nelson TR, Yang K, Kwan AL, Miller DF. Dedicated breast CT: initial clinical experience. Radiology 2008;246:725-733
2. Kim SM, Park JM. Computed tomography of the breast. Abnormal findings with mammographic and sonographic correlation. J Comput Assist Tomogr 2003;27:761-770
3. Shojaku H, Seto H, Iwai H, Kitazawa S, Fukushima W, Saito K. Detection of incidental breast tumors by noncontrast spiral computed tomography of the chest. Radiat Med 2008;26:362-367
4. Yi JG, Kim SJ, Marom EM, Park JH, Jung SI, Lee MW. Chest CT of incidental breast lesions. J Thorac Imaging 2008;23:148-155
5. Harish MG, Konda SD, MacMahon H, Newstead GM. Breast lesions incidentally detected with CT: what the general radiologist needs to know. Radiographics 2007;27:S37-S51
6. Hussain A, Gordon-Dixon A, Almusawy H, Sinha P, Desai A. The incidence and outcome of incidental breast lesions detected by computed tomography. Ann R Coll Surg Engl
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2010;92:124-126

7. Moyle P, Sonoda L, Britton P, Sinnatamby R. Incidental breast lesions detected on CT: what is their significance? Br J Radiol 2010;83:233-240

8. Porter G, Steel J, Paisley K, Watkins R, Holgate C. Incidental breast masses detected by computed tomography: are any imaging features predictive of malignancy? Clin Radiol 2009;64:529-533

9. Swensen SJ, Jett JR, Hartman TE, Midthun DE, Sloan JA, Sykes AM, et al. Lung cancer screening with CT: Mayo Clinic experience. Radiology 2003;226:756-761

10. Miyake K, Hayakawa K, Nishino M, Nakamura Y, Morimoto T, Urata Y, et al. Benign or malignant?: differentiating breast lesions with computed tomography attenuation values on dynamic computed tomography mammography. J Comput Assist Tomogr 2005;29:772-779

11. Inoue M, Sano T, Watai R, Ashikaga R, Ueda K, Watatani M, et al. Dynamic multidetector CT of breast tumors: diagnostic features and comparison with conventional techniques. AJR Am J Roentgenol 2003;181:679-686

12. Kuhl C. The current status of breast MR imaging. Part I. Choice of technique, image interpretation, diagnostic accuracy, and transfer to clinical practice. Radiology 2007;244:356-378