# **Evaluation of Palpable Breast Masses**

SUSAN KLEIN, M.D., Southern Illinois University School of Medicine, Decatur, Illinois

Palpable breast masses are common and usually benign, but efficient evaluation and prompt diagnosis are necessary to rule out malignancy. A thorough clinical breast examination, imaging, and tissue sampling are needed for a definitive diagnosis. Fine-needle aspiration is fast, inexpensive, and accurate, and it can differentiate solid and cystic masses. However, physicians must have adequate training to perform this procedure. Mammography screens for occult malignancy in the same and contralateral breast and can detect malignant lesions in older women; it is less sensitive in women younger than 40 years. Ultrasonography can detect cystic masses, which are common, and may be used to guide biopsy techniques. Tissue specimens obtained with core-needle biopsy allow histologic diagnosis, hormone-receptor testing, and differentiation between in situ and invasive disease. Core-needle biopsy is more invasive than fine-needle aspiration, requires more training and experience, and frequently requires imaging guidance. After the clinical breast examination is performed, the evaluation depends largely on the patient's age and examination characteristics, and the physician's experience in performing fine-needle aspiration. (Am Fam Physician 2005;71:1731-8. Copyright© 2005 American Academy of Family Physicians.)

See page 1635 for strength-of-recommendation labels.

Nearly one third of women

diagnosed with breast can-

cer between 1996 and 2000

were younger than 50 years.

reast masses have a variety of etiologies, benign and malignant. Fibroadenoma is the most common benign breast mass; invasive ductal carcinoma is the most common malignancy. Most masses are benign, but breast cancer is the most common cancer and the second

leading cause of cancer deaths in women.<sup>2</sup> Although most breast cancers occur in women older than 50 years, 31 percent of women diagnosed with breast cancer between 1996 and 2000 were younger than 50 years.<sup>3</sup> An

efficient and accurate evaluation can maximize cancer detection and minimize unnecessary testing and procedures.

# Initial Evaluation HISTORY

A thorough patient history is necessary for the physician to identify risk factors for breast cancer. Some risk factors are well established, and others indicate probable or possible increased risk (*Tables 1 and 2*).<sup>4-14</sup>

## PHYSICAL EXAMINATION

A complete clinical breast examination (CBE) includes an assessment of both breasts and the chest, axillae, and regional lymphatics. In premenopausal women, the CBE is best done the

week following menses, when breast tissue is least engorged. With the patient in an upright position, the physician visually inspects the breasts, noting asymmetry, nipple discharge, obvious masses, and skin changes, such as dimpling, inflammation, rashes, and unilateral nipple retraction or inversion.<sup>15</sup>

With the patient supine and one arm raised, the physician thoroughly palpates breast tissue on the raised-arm side in the superficial, intermediate, and deep tissue planes (i.e., the "triple touch" technique); axilla; supraclavicular area; neck; and chest wall, assessing the size, texture, and location of any masses (Figure 1).15 The physician should note the size of the masses to document changes over time. Next, the physician should inspect the areola-nipple complex for any discharge. CBE sensitivity can be improved by longer duration (i.e., five to 10 minutes) and increased precision (i.e., using a systematic pattern, varying palpation pressure, and using three finger pads and circular motions). 15,16

Benign masses generally cause no skin change and are smooth, soft to firm, and mobile, with well-defined margins. Diffuse, symmetric thickening, which is common in the upper outer quadrants, may indicate fibrocystic changes. Malignant masses generally are hard, immobile, and fixed to surrounding skin and soft tissue, with poorly defined

Key clinical recommendation	Label	References
Ultrasonography-guided CNB should be considered to diagnose malignancy in women with palpable breast lesions.	С	1
In young women with dense breast tissue, ultrasonography should be used rather than mammography to detect breast lesions.	С	18, 20
Mammography should be used rather than magnetic resonance imaging to detect in situ carcinomas of the breast.	С	27, 28, 33
Diagnostic mammography is indicated in women older than 40 years if FNA reveals a solid mass.	С	34, 48
Excisional biopsy should be performed in women with clinically suspicious lesions, or lesions that are equivocal on imaging, FNA, or CNB.	С	40, 41, 42
Cystic lesions that resolve after FNA do not require further evaluation unless they recur.	С	48
If CBE, FNA, and imaging indicate benign disease, the CBE should be repeated in four to six weeks.	С	48

CNB = core-needle biopsy; FNA = fine-needle aspiration; CBE = clinical breast examination.

A = consistent, good-quality patient-oriented evidence; B = inconsistent or limited-quality patient-oriented evidence; C = consensus, disease-oriented evidence, usual practice, opinion, or case series. See page 1635 for more information.

or irregular margins.<sup>15</sup> However, mobile or nonfixed masses can be cancerous. Infections such as mastitis and cellulitis tend to be erythematous, tender, and warm to the touch; they may be more circumscribed if an abscess has formed. Similar symptoms may occur in patients with inflammatory breast cancer. Therefore, caution should be used in assessing patients with suspected breast infections.

Digital palpation of the breast is effective in detecting masses and can help determine whether a mass is benign or malignant. 15,17 CBE can detect up to 44 percent of cancers, up to 29 percent of which would not have been detected by mammography. 15,17 Despite its accuracy, CBE alone is not adequate for definitive diagnosis of breast cancer. Further evaluation, including follow-up examinations, imaging, and tissue sampling, is required in all patients with breast masses.

## **Imaging ULTRASONOGRAPHY**

Ultrasonography can effectively distinguish solid masses from cysts, which account for approximately 25 percent of breast

TABLE 1	
Relevant History in Women with Palpable Breast Ma	sses

Breast lump characteristics	Medical and surgical history
Changes in size over time	Personal history of breast cancer
Change relative to menstrual cycle	Previous breast masses and biopsies
Duration of mass	Recent breast trauma or surgery
Pain or swelling	Recent radiation therapy or chemotherapy
Redness, fever, or discharge	Other exposure to radiation
Diet and medications	Personal characteristics
Current medications	Age at first childbearing
History of hormone therapy	Age at menarche
Family history	Age at menopause
History of breast disease	Current age
Relationship to patient	Current lactation status
Relative's age at onset	History of breastfeeding
	Number of children
	Social history
	Radiation and chemical exposure
	Smoking

### TABLE 2

## Risk Factors for Breast Cancer

## Well-established risk factors

Age 50 or older

Benign breast disease, especially cystic disease, proliferative types of hyperplasia, and atypical hyperplasia

Exposure to ionizing radiation

First childbirth after age 20

Higher socioeconomic status

History of breast cancer

History of breast cancer in a first-degree relative

Hormone therapy

Nulliparity

Obesity (i.e., BMI  $\geq$  30 kg per m<sup>2</sup>)\*

### Probable risk factors

Alcohol consumption

Did not breastfeed

Elevated endogenous estrogen levels

High BMI\*

Hormonal contraception therapy

Increased mammographic density of breast tissue

Menarche before age 12

Menopause after age 45

Mutations in BRCA 1 and BRCA 2 genes

## Possible risk factors

Chemical exposure

Diet high in fat

Diet low in beta carotene, folate, and vitamins

A and C

Diet low in fruits and vegetables

BMI = body mass index.

\*—Breast cancer risk increases by 3 percent with every 1 kg per m<sup>2</sup> increase in BMI.

Information from references 4 through 14.

lesions.<sup>18,19</sup> When strict criteria for cyst diagnosis are met, ultrasonography has a sensitivity of 89 percent and a specificity of 78 percent in detecting abnormalities in symptomatic women.<sup>18</sup> Recurrent or complex cysts may signal malignancy; therefore, further evaluation of these lesions is required.<sup>19</sup>

Although ultrasonography is not considered a screening test, it is more sensitive than mammography in detecting lesions in women with dense breast tissue. <sup>18,20</sup> It is useful in discriminating between benign and malignant solid masses, <sup>18,21</sup> and it is superior to mammography in diagnosing clinically benign palpable masses (i.e., up to 97 percent accuracy versus 87 percent for mammography). <sup>21</sup>

## DIAGNOSTIC MAMMOGRAPHY

Diagnostic mammography can help physicians determine whether a lesion is potentially malignant, and it also screens for occult disease in surrounding tissue. A radio-opaque ball bearing marks the location of the mass, and spot compression and magnification

views can clarify the breast mass and determine its density. If old films are available, they are compared with the new images. Diagnostic mammography is up to 87 percent sensitive in detecting cancer.<sup>22</sup> Its specificity is 88 percent, and its positive predictive value may be as high as 22 percent.<sup>22</sup>

# DIGITAL MAMMOGRAPHY

Digital mammography allows images to be enhanced and transmitted electronically.

The ability to alter contrast and brightness permits further evaluation of abnormal areas to identify features diagnostic of benign and malignant disease.<sup>23-26</sup> Although the overall cancer-detection rate is similar in screen-field and full-field

reliable in detecting breast abnormalities in symptomatic women when strict diagnostic criteria are met.

Ultrasonography is highly

mammography, screen-field imaging has better image quality and less artifact, and requires fewer patient recalls.<sup>24,25</sup>

In addition to its usefulness in telemammography, digital mammography may be more accurate than traditional mammography; studies comparing the methods are underway. Potential new techniques include three-dimensional imaging, lower-dose radiation, dual energy subtraction, contrast-enhancement imaging, and computer-assisted diagnosis.<sup>23,24,26</sup>

## MAGNETIC RESONANCE IMAGING

Magnetic resonance imaging (MRI) is being studied to determine its usefulness in diagnosing breast masses. Gadolinium contrast is used to enhance the vascularity of malignant lesions. Although MRI is highly sensitive (85 to 100 percent), it lacks specificity

The rightsholder did not grant rights to reproduce this item in electronic media. For the missing item, see the original print version of this publication.

Figure 1.

(47 to 67 percent).<sup>27,28</sup> MRI is inferior to mammography in detecting in situ cancers and cancers smaller than 3 mm, and it provides no cost benefit over excisional biopsy for verifying malignancy.<sup>27,28</sup> Research suggests two potential roles for MRI in breast mass diagnosis: evaluating patients with silicone breast implants<sup>29</sup> and assessing patients in whom evaluation by ultrasonography and mammography is problematic. The latter group includes patients who have had breastconserving surgery; patients with known carcinoma in whom multifocal, ipsilateral, or contralateral disease must be ruled out; patients with axillary metastasis and an unknown primary; patients with extensive postoperative scarring; and patients with extremely dense parenchyma. <sup>28,30-32</sup> A recent study<sup>33</sup> compared the effectiveness of mammography and MRI in women with a family history of breast cancer or a genetic susceptibility to the disease. The sensitivity of MRI was higher than that of mammography in detecting breast cancer, and MRI was better able to discriminate between benign and malignant lesions. Although MRI improves detection of early breast cancers in carriers of BRCA mutations, it has a lower specificity than mammography, which requires additional evaluations. It also has a limited sensitivity in detecting ductal carcinoma in situ.

# Tissue Studies FINE-NEEDLE ASPIRATION

The first step in evaluating patients with palpable breast masses often is fine-needle aspiration (FNA), in which a 22- to 25-gauge needle is used to aspirate cystic fluid or sample solid lesions for cytology. In some patients, the lesion completely resolves after FNA, and no further diagnostic work-up is required. However, when imaging is indicated after FNA, cyst wall disruption caused by the procedure may make imaging more difficult to evaluate. The problem may be avoided by scheduling imaging studies up to two weeks after FNA and notifying the radiologist of the recent procedure.<sup>34</sup>

FNA also is used with ultrasonography or stereotactic imaging to further assess poorly defined palpable masses. When sampling in

The triple test is the

sampling.

combination of results

from clinical breast exami-

nation, imaging, and tissue

patients with solid lesions is adequate, FNA is highly sensitive for malignancy (98 to 99 percent) and has a positive predictive value of 99 percent and a negative predictive value of 86 to 99 percent.<sup>35</sup> Sample adequacy is of some concern; one study<sup>36</sup> rated 28 percent of samples as inadequate and another 22 percent as less than optimal. The physician's training and experience may be a key factor in obtaining adequate samples.<sup>36</sup>

## **CORE-NEEDLE BIOPSY**

Core-needle biopsy (CNB) produces a larger tissue sample than FNA and may be used in conjunction with ultrasonography or stereotactic imaging for small or difficult-to-palpate lesions. Local anesthesia is required. A 14- to 18-gauge cutting needle is used to obtain two to six slender cores of tissue for histology.<sup>37,38</sup> The sensitivity of ultrasonography-guided CNB may be as high as 99 percent in diagnosing malignancy in palpable lesions and 93 percent in nonpalpable lesions. Specimens can be used to differentiate between in situ and invasive carcinoma, and to identify hormone-receptor levels.<sup>39</sup> Results vary depending on radiographic guidance, the size of the needle, and the number of cores sampled. A minimum of four cores is suggested to achieve greater accuracy. 37,38 Insufficient specimens are rare.<sup>1,38</sup> Compared with FNA, CNB takes more time and requires specific training and patient anesthesia, but it has a higher positive predictive value for suspicious and atypical results and may provide an overall cost benefit.38

## **EXCISIONAL BIOPSY**

Excisional biopsy is the gold standard for evaluating breast masses. It is performed in an operating room under local or general anesthesia and results in the removal of the entire lesion. Excisional biopsy is diagnostic and therapeutic: a completely removed mass with good margins of normal tissue may mean that further surgery is not required. An incisional biopsy (i.e., removal of a portion of the lesion) generally is used for tissue diagnosis in large tumors when CNB is nondiagnostic. Excisional biopsy is indicated in patients with clinically suspicious lesions

and lesions in which imaging or tissue studies are equivocal.<sup>40-43</sup> With the increased use of CNB, the need for diagnostic excisional biopsy has declined.<sup>44</sup>

## **Triple Test**

The triple test is the combination of results from CBE, imaging, and tissue sampling. 41,45-47 When the three assessments are performed

adequately and produce concordant results, the triple test diagnostic accuracy approaches 100 percent. 41,45,46 Discordant results or results that cannot be evaluated may indicate the need for excisional biopsy. 45

y. • 5

The Triple Test Score (TTS) was developed to help physicians interpret discordant triple test results. A three-point scale is used to score each component of the triple test (1 = benign, 2 = suspicious, 3 = malignant). A TTS of 3 or 4 is consistent with a benign lesion; a TTS of 6 or more indicates possible malignancy that may require surgical intervention. Excisional biopsy is recommended in patients with a TTS of 5 to obtain a definitive diagnosis.

## **DIAGNOSTIC EVALUATION PROCESS**

After the patient history is obtained and the CBE is performed, the next diagnostic step is determined by the patient's age and the physician's experience with performing office-based FNA (*Figure 2*). Physicians trained in FNA may choose this procedure for two reasons: it is office-based and may be performed during the same clinical visit, and cystic lesions that resolve on aspiration spare the patient further work-up and cost.<sup>45</sup>

If FNA reveals a solid lesion, evaluation with diagnostic mammography should be performed next<sup>34,48</sup>; ultrasonography may be considered in women younger than 40 years.<sup>41</sup> If all three elements (CBE, FNA, and imaging) indicate benign disease (i.e., TTS of 3), the patient may be followed with another examination in four to six weeks.<sup>42,48</sup> If all triple test elements are positive (i.e., suggestive of malignancy), surgical intervention is indicated.<sup>45</sup> Patients with discordant results and a TTS of 4 may be followed with

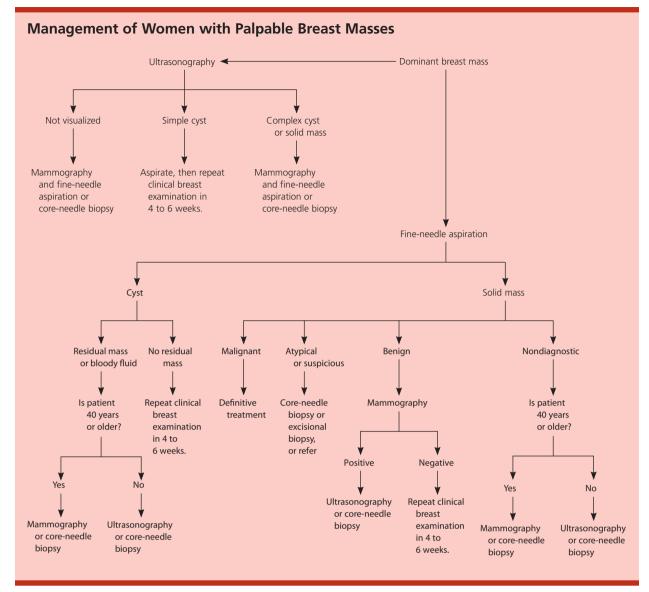


Figure 2. Diagnostic algorithm for patients with palpable breast masses.

repeat examination, but excisional biopsy or referral to a breast specialist is indicated in patients with a TTS of 5 or higher.<sup>42</sup> CNB may be performed to enhance the triple test accuracy if it was not used previously.<sup>38,47</sup>

If FNA is not feasible during the ini-

## The Author

SUSAN KLEIN, M.D., is assistant professor of family and community medicine at Southern Illinois University School of Medicine, Decatur Family Practice Center. She received her medical degree from the University of Illinois at Chicago College of Medicine and completed a family medicine residency at Southern Illinois University School of Medicine, Decatur Family Practice Center.

Address correspondence to Susan Klein, M.D., Southern Illinois University School of Medicine, SIU Family Physicians, 250 W. Kenwood Ave., Decatur, IL 62526-4372 (e-mail: sklein1@siumed.edu). Reprints are not available from the author.

tial presentation, ultrasonography should be considered to rule out cystic disease and delineate lesion margins. <sup>22,48</sup> Cystic lesions may be aspirated. Solid lesions should be evaluated with mammography, which often can be done during the same visit, to further delineate lesion margins and to screen for occult disease in the ipsilateral and contralateral breast, particularly in women older than 40 years. <sup>49</sup> Solid lesions will then require FNA or CNB<sup>48</sup> to complete the triple test. Further management should be implemented as described in *Figure 2*.

The author indicates that she does not have any conflict of interest. Sources of funding: none reported.

Members of various family medicine departments develop articles for "Problem-Oriented Diagnosis." This

is one in a series from the Department of Family and Community Medicine at Southern Illinois University School of Medicine, Springfield. Guest editor of the series is John G. Bradley, M.D., professor and director of the Decatur Family Practice Residency Program.

### REFERENCES

- Schoonjans JM, Brem RF. Fourteen-gauge ultrasonographically guided large-core needle biopsy of breast masses. J Ultrasound Med 2001;20:967-72.
- American Cancer Society. Cancer facts and figures 2003. Accessed online October 14, 2004, at: http://www.cancer.org/downloads/STT/CAFF2003PWSecured.pdf.
- National Cancer Institute. SEER 1973-2001 public-use data. Accessed online October 14, 2004, at: http://seer. cancer.gov/publicdata/.
- Madigan MP, Ziegler RG, Benichou J, Byrne C, Hoover RN. Proportion of breast cancer cases in the United States explained by well-established risk factors. J Natl Cancer Inst 1995;87:1681-5.
- National Cancer Institute. Breast cancer (PDQ): prevention. Accessed online October 14, 2004, at: http://www.cancer.gov/cancertopics/pdq/prevention/breast/healthprofessional.
- Newcomb PA, Titus-Ernstoff L, Egan KM, Trentham-Dietz A, Baron JA, Storer BE, et al. Postmenopausal estrogen and progestin use in relation to breast cancer risk. Cancer Epidemiol Biomarkers Prev 2002;11:593-600.
- Weiss LK, Burkman RT, Cushing-Haugen KL, Voigt LF, Simon MS, Daling JR, et al. Hormone replacement therapy regimens and breast cancer risk (1). Obstet Gynecol 2002;100:1148-58.
- Key TJ, Appleby PN, Reeves GK, Roddam A, Dorgan JF, Longcope C, et al. Body mass index, serum sex hormones, and breast cancer risk in postmenopausal women. J Natl Cancer Inst 2003;95:1218-26.
- Boyd NF, Jensen HM, Cooke G, Han HL, Lockwood GA, Miller AB. Mammographic densities and the prevalence and incidence of histological types of benign breast disease. Reference Pathologists of the Canadian National Breast Screening Study. Eur J Cancer Prev 2000:9:15-24.
- Byrne C, Connolly JL, Colditz GA, Schnitt SJ. Biopsy confirmed benign breast disease, postmenopausal use of exogenous female hormones, and breast carcinoma risk. Cancer 2000;89:2046-52.
- Kumle M, Weiderpass E, Braaten T, Persson I, Adami HO, Lund E. Use of oral contraceptives and breast cancer risk: the Norwegian-Swedish Women's Lifestyle and Health Cohort Study. Cancer Epidemiol Biomarkers Prev 2002;11:1375-81.
- Breast cancer and hormonal contraceptives: collaborative reanalysis of individual data on 53 297 women with breast cancer and 100 239 women without breast cancer from 54 epidemiological studies. Collaborative Group on Hormonal Factors in Breast Cancer. Lancet 1996:347:1713-27.
- Rossouw JE, Anderson GL, Prentice RL, LaCroix AZ, Kooperberg C, Stefanick ML, et al. Risks and benefits of estrogen plus progestin in healthy postmenopausal women: principal results from the Women's Health Initiative randomized controlled trial. JAMA 2002;288:321-33.

- Budai B, Szamel I, Sulyok Z, Nemet M, Bak M, Otto S, et al. Characteristics of cystic breast disease with special regard to breast cancer development. Anticancer Res 2001:21:749-52.
- Barton MB, Harris R, Fletcher SW. The rational clinical examination. Does this patient have breast cancer? The screening clinical breast examination: should it be done? How? JAMA 1999;282:1270-80.
- Campbell HS, Fletcher SW, Pilgrim CA, Morgan TM, Lin S. Improving physicians' and nurses' clinical breast examination: a randomized controlled trial. Am J Prev Med 1991;7:1-8.
- Baines CJ, Miller AB. Mammography versus clinical examination of the breasts. J Natl Cancer Inst Monogr 1997:(22):125-9.
- Moss HA, Britton PD, Flower CD, Freeman AH, Lomas DJ, Warren RM. How reliable is modern breast imaging in differentiating benign from malignant breast lesions in the symptomatic population? Clin Radiol 1999;54:676-82.
- Berg WA, Campassi CI, Ioffe OB. Cystic lesions of the breast: sonographic-pathologic correlation. Radiology 2003;227:183-91.
- Kolb TM, Lichy J, Newhouse JH. Comparison of the performance of screening mammography, physical examination, and breast US and evaluation of factors that influence them: an analysis of 27,825 patient evaluations. Radiology 2002;225:165-75.
- Lister D, Evans AJ, Burrell HC, Blamey RW, Wilson AR, Pinder SE, et al. The accuracy of breast ultrasound in the evaluation of clinically benign discrete, symptomatic breast lumps. Clin Radiol 1998;53:490-2.
- 22. Barlow WE, Lehman CD, Zheng Y, Ballard-Barbash R, Yankaskas BC, Cutter GR, et al. Performance of diagnostic mammography for women with signs or symptoms of breast cancer. J Natl Cancer Inst 2002;94:1151-9.
- Leichter I, Buchbinder S, Bamberger P, Novak B, Fields S, Lederman R. Quantitative characterization of mass lesions on digitized mammograms for computerassisted diagnosis. Invest Radiol 2000;35:366-72.
- Obenauer S, Luftner-Nagel S, von Heyden D, Munzel U, Baum F, Grabbe E. Screen film vs full-field digital mammography: image quality, detectability and characterization of lesions [published correction appears in Eur Radiol 2002;12:2388]. Eur Radiol 2002;12:1697-702.
- Lewin JM, Hendrick RE, D'Orsi CJ, Isaacs PK, Moss LJ, Karellas A, et al. Comparison of full-field digital mammography with screen-film mammography for cancer detection: results of 4,945 paired examinations. Radiology 2001;218:873-80.
- Fischer U, Baum F, Obenauer S, Luftner-Nagel S, von Heyden D, Vosshenrich R, et al. Comparative study in patients with microcalcifications: full-field digital mammography vs screen-film mammography. Eur Radiol 2002:12:2679-83.
- Hrung JM, Sonnad SS, Schwartz JS, Langlotz CP. Accuracy of MR imaging in the work-up of suspicious breast lesions: a diagnostic meta-analysis. Acad Radiol 1999;6:387-97.
- 28. Kristoffersen Wiberg M, Aspelin P, Perbeck L, Bone B. Value of MR imaging in clinical evaluation of breast lesions. Acta Radiol 2002;43:275-81.
- Herborn CU, Marincek B, Erfmann D, Meuli-Simmen C, Wedler V, Bode-Lesniewska B, et al. Breast augmentation and reconstructive surgery: MR imaging of implant rupture and malignancy. Eur Radiol 2002;12:2198-206.

## **Breast Masses**

- Liberman L, Morris EA, Dershaw DD, Abramson AF, Tan LK. MR imaging of the ipsilateral breast in women with percutaneously proven breast cancer. AJR Am J Roentgenol 2003:180:901-10.
- Obdeijn IM, Brouwers-Kuyper EM, Tilanus-Linthorst MM, Wiggers T, Oudkerk M. MR imaging-guided sonography followed by fine-needle aspiration cytology in occult carcinoma of the breast. AJR Am J Roentgenol 2000;174:1079-84.
- 32. Liberman L, Morris EA, Kim CM, Kaplan JB, Abramson AF, Menell JH, et al. MR imaging findings in the contralateral breast of women with recently diagnosed breast cancer. AJR Am J Roentgenol 2003;180:333-41.
- Kriege M, Brekelmans CT, Boetes C, Besnard PE, Zonderland HM, Obdeijn IM, et al. Efficacy of MRI and mammography for breast-cancer screening in women with a familial or genetic predisposition. N Engl J Med 2004;351:427-37.
- Hindle WH, Chen EC. Accuracy of mammographic appearances after breast fine-needle aspiration. Am J Obstet Gynecol 1997;176:1286-90.
- Ariga R, Bloom K, Reddy VB, Kluskens L, Francescatti D, Dowlat K, et al. Fine-needle aspiration of clinically suspicious palpable breast masses with histopathologic correlation. Am J Surg 2002;184:410-3.
- 36. Saxe A, Phillips E, Orfanou P, Husain M. Role of sample adequacy in fine needle aspiration biopsy of palpable breast lesions. Am J Surg 2001;182:369-71.
- Fishman JE, Milikowski C, Ramsinghani R, Velasquez MV, Aviram G. US-guided core-needle biopsy of the breast: how many specimens are necessary? Radiology 2003;226:779-82.
- 38. Westenend PJ, Sever AR, Beekman-De Volder HJ, Liem SJ. A comparison of aspiration cytology and core needle biopsy in the evaluation of breast lesions. Cancer 2001;93:146-50.
- Chuo CB, Corder AP. Core biopsy vs fine needle aspiration cytology in a symptomatic breast clinic. Eur J Surg Oncol 2003;29:374-8.

- Smith BL. The breast. In: Ryan KJ, Kistner RW. Kistner's Gynecology and women's health. 7th ed. St. Louis: Mosby, 1999:197-202.
- Morris KT, Vetto JT, Petty JK, Lum SS, Schmidt WA, Toth-Fejel S, et al. A new score for the evaluation of palpable breast masses in women under age 40. Am J Surg 2002;184:346-7.
- 42. Morris KT, Pommier RF, Morris A, Schmidt WA, Beagle G, Alexander PW, et al. Usefulness of the triple test score for palpable breast masses. Arch Surg 2001;136:1008-12.
- Osuch JR, Reeves MJ, Pathak DR, Kinchelow T. BREAST-AID: clinical results from early development of a clinical decision rule for palpable solid breast masses. Ann Surg 2003:238:728-37
- 44. Crowe JP Jr, Rim A, Patrick R, Rybicki L, Grundfest S, Kim J, et al. A prospective review of the decline of excisional breast biopsy. Am J Surg 2002;184:353-5.
- 45. Steinberg JL, Trudeau ME, Ryder DE, Fishell E, Chapman JA, McCready DR, et al. Combined fine-needle aspiration, physical examination and mammography in the diagnosis of palpable breast masses: their relation to outcome for women with primary breast cancer [published correction appears in Can J Surg 1997;40:9]. Can J Surg 1996;39:302-11.
- Kamphausen BH, Toellner T, Ruschenburg I. The value of ultrasound-guided fine-needle aspiration cytology of the breast: 354 cases with cytohistological correlation. Anticancer Res 2003;23:3009-13.
- 47. Clarke D, Sudhakaran N, Gateley CA. Replace fine needle aspiration cytology with automated core biopsy in the triple assessment of breast cancer. Ann R Coll Surg Engl 2001;83:110-2.
- 48. Kerlikowske K, Smith-Bindman R, Ljung BM, Grady D. Evaluation of abnormal mammography results and palpable breast abnormalities. Ann Intern Med 2003;139:274-84.
- Shetty MK, Shah YP, Sharman RS. Prospective evaluation of the value of combined mammographic and sonographic assessment in patients with palpable abnormalities of the breast. J Ultrasound Med 2003;22:263-8.