

Screening for Breast Cancer

Joann G. Elmore, MD, MPH

Katrina Armstrong, MD

Constance D. Lehman, MD, PhD

Suzanne W. Fletcher, MD, MSc

BREAST CANCER SCREENING, ESPECIALLY with mammography, has been recommended for many decades,¹ and the majority of women older than 40 years in the United States participate in screening activities.^{2,3} Meanwhile, new screening modalities have been introduced, and some of these have been increasingly incorporated into community practice. However, none of the new technologies has been evaluated for its effect on breast cancer mortality.

Community practice of screening may differ from the care provided within randomized clinical trials and is less often discussed in review articles. Reviews of breast cancer screening usually emphasize efficacy and results of randomized trials, particularly those involving screen-film mammography.⁴⁻⁷ Efficacy of a screening tool is measured in experimental studies under ideal circumstances.⁸ In contrast, *effectiveness* is defined as the extent to which a specific intervention "when deployed in the field in routine circumstances, does what it is intended to do for a specific population."⁸

We systematically reviewed what is known about the community practice of mammography, clinical breast examination, and breast self-examination, when possible, comparing the results from community studies with

CME available online at
www.jama.com

Context Breast cancer screening in community practices may be different from that in randomized controlled trials. New screening modalities are becoming available.

Objectives To review breast cancer screening, especially in the community and to examine evidence about new screening modalities.

Data Sources and Study Selection English-language articles of randomized controlled trials assessing effectiveness of breast cancer screening were reviewed, as well as meta-analyses, systematic reviews, studies of breast cancer screening in the community, and guidelines. Also, studies of newer screening modalities were assessed.

Data Synthesis All major US medical organizations recommend screening mammography for women aged 40 years and older. Screening mammography reduces breast cancer mortality by about 20% to 35% in women aged 50 to 69 years and slightly less in women aged 40 to 49 years at 14 years of follow-up. Approximately 95% of women with abnormalities on screening mammograms do not have breast cancer with variability based on such factors as age of the woman and assessment category assigned by the radiologist. Studies comparing full-field digital mammography to screen film have not shown statistically significant differences in cancer detection while the impact on recall rates (percentage of screening mammograms considered to have positive results) was unclear. One study suggested that computer-aided detection increases cancer detection rates and recall rates while a second larger study did not find any significant differences. Screening clinical breast examination detects some cancers missed by mammography, but the sensitivity reported in the community is lower (28% to 36%) than in randomized trials (about 54%). Breast self-examination has not been shown to be effective in reducing breast cancer mortality, but it does increase the number of breast biopsies performed because of false-positives. Magnetic resonance imaging and ultrasound are being studied for screening women at high risk for breast cancer but are not recommended for screening the general population. Sensitivity of magnetic resonance imaging in high-risk women has been found to be much higher than that of mammography but specificity is generally lower. Effect of the magnetic resonance imaging on breast cancer mortality is not known. A balanced discussion of possible benefits and harms of screening should be undertaken with each woman.

Conclusions In the community, mammography remains the main screening tool while the effectiveness of clinical breast examination and self-examination are less. New screening modalities are unlikely to replace mammography in the near future for screening the general population.

JAMA. 2005;293:1245-1256

www.jama.com

Author Affiliations: Department of Medicine (Dr Elmore) and Department of Radiology and Seattle Cancer Care Alliance (Dr Lehman), University of Washington School of Medicine, Seattle; Department of Medicine, University of Pennsylvania School of Medicine, Philadelphia (Dr Armstrong); and the Department of Ambulatory Care and Prevention, Harvard Pilgrim Health Care and Harvard Medical School, Boston, Mass (Dr Fletcher).

Corresponding Author: Joann G. Elmore, MD, MPH, Harborview Medical Center, 325 Ninth Ave, Box 359780, Seattle, WA 98104 (jelmore@u.washington.edu).

Clinical Review Section Editor: Michael S. Lauer, MD. We encourage authors to submit papers for consideration as a "Clinical Review." Please contact Michael S. Lauer, MD, at lauer@cfc.org.

those of randomized clinical trials. In addition, we reviewed what is known about newer screening modalities, specifically digital mammography, computer-aided detection programs for mammography, ultrasound, and magnetic resonance imaging (MRI).

METHODS

The evaluation of screening modalities, especially in the community setting, is challenging for methodological, clinical, and ethical reasons. Randomized clinical trials are considered the gold standard for evaluating a new screening test. The long-term breast cancer mortality rate of women randomized to receive a new screening test is compared with that of women randomized to receive standard care. However, such trials are difficult to conduct. They require tens of thousands of women who need to be followed up for more than 15 years. Furthermore, because mammography screening has been shown to be effective in some trials, it would likely be even more difficult to demonstrate any additional efficacy of new tests. Finally, as treatment for breast cancer has improved over time,⁹⁻¹¹ the impact of screening on breast cancer mortality may be increasingly difficult to establish.

Because of these challenges, new screening tests are often first studied by establishing characteristics of the tests themselves, rather than by studying their effect on patient outcome such as breast cancer mortality. Important test characteristics include sensitivity, specificity, safety, cost, simplicity, and patient and clinician acceptability. We review what test characteristics have been studied and the findings for each new modality. We also indicate the study design and the end points studied for each screening test. Although it is important to use resources wisely when considering a screening test for a large segment of the population, cost-effectiveness analyses are not reviewed.

It is important to determine the characteristics of a screening test in a community setting if the test is to be used in that setting. However, test character-

istics of new modalities are usually evaluated among women for whom the rate of breast cancer is higher than average, such as women at increased risk of breast cancer or women in a diagnostic setting with breast symptoms or known breast abnormalities. The reported sensitivity and specificity of a test in these high-risk women may be different from the sensitivity and specificity of the same test used in a general screening population.¹² We therefore indicate if a test has been evaluated as a diagnostic or screening test and if as a screening test, whether it has been evaluated in women thought to be at increased risk or in the general population.

For this review, searches of MEDLINE, The Cochrane Library, the National Guideline Clearinghouse Web site, the US Preventive Services Task Force recommendations and reviews,^{5,13} and the International Agency for Research on Cancer Handbook of Cancer Prevention (IARC)⁴ were performed to identify English-language articles about breast cancer screening. Examples of search terms included *mass screening and breast* and then the specific modality (eg, *mammography, breast examination, digital mammography, computer assisted diagnosis or computer aided detection, magnetic resonance imaging, ultrasound or ultrasonography*). The bibliographies of retrieved articles were also scanned to retrieve additional relevant articles. Details regarding search terms are available from the authors.

RESULTS

Screening Mammography

Eight reported randomized trials have studied mammography's effectiveness in the United States,^{14,15} Sweden,¹⁶⁻²⁰ Canada,^{21,22} and the United Kingdom.²³ Concerns related to flaws of these randomized clinical trials have been raised.^{24,25} In-depth independent reviews of the criticisms of the trials have concluded that these flaws do not negate mammography's efficacy in reducing breast cancer mortality, especially in women aged 50 to 69 years.^{4,5,7,26} Trials comparing mammography with or without clinical breast ex-

amination to usual care (with little or no screening mammography) demonstrated remarkably consistent results for women older than 50 years. Meta-analyses that included all trials demonstrated statistically significant reductions of 20% to 35% in mortality from breast cancer for women aged 50 to 69 years.⁷ The majority of participants in clinical trials of mammography were white and information on *BRCA* mutation status was not known.

In general, breast cancers detected by mammography screening are smaller and have more favorable histological and biological features than tumors detected between mammography screening rounds or tumors found outside of screening.²⁷⁻³² Because the favorable prognoses of women with breast cancer detected by mammography screening may be attributable to selection bias, length bias, lead-time bias, and overdiagnosis,^{33,34} randomized controlled trials with breast cancer mortality as the outcome have been particularly important in excluding such biases.

The benefit of screening women in their 40s is slower to appear and is somewhat less than that of women older than 50 years. Women in their 40s have a lower incidence of disease, denser breast tissue (which can lower the sensitivity of mammography), and, on average, faster-growing cancers.^{27,35,36} A randomized trial of mammography screening for women in their early 40s is under way in the United Kingdom.³⁷ Clinicians and patients are often surprised at the large number of women who need to be screened to prevent 1 death due to breast cancer. For example, it has been estimated that between 500 and 1800 women who are 40 years of age would need to undergo regular screening mammography to prevent 1 breast cancer death after 14 to 20 years.^{7,13}

Randomized trials have included few or no women older than 70 years. One case-control study found that screening women between 65 and 74 years of age was associated with mortality reduction.³⁸ Pooled data from a community study showed that the sensitivity and specificity of screening mammog-

raphy were highest in older women, especially those older than 80 years.³⁶ Screening mammography may not be beneficial for women with significant comorbidity or with a life expectancy of less than 5 years, and may actually necessitate work-up that does not result in any benefit.³⁹⁻⁴¹ However, if a woman is in reasonably good health and would be eligible for and interested in treatment, continued screening should be supported. Encouraging individualized decisions may be especially appropriate for women older than 70 years.⁴²

The majority of women with abnormalities noted on screening mammograms ($\approx 95\%$) do not have breast cancer with variability based on multiple factors including the radiologist's assessment and the woman's age.⁷ Because the risk of breast cancer increases with age, the likelihood of a woman with an abnormal mammogram result having cancer also increases with age.^{43,44} On the other hand, having a normal mammogram result does not rule out the possibility of having breast cancer, because false-negative mammography examination results do occur. In such cases, either the cancer is not visible on mammography examination or the radiologist fails to notice the lesion prospectively.⁴⁵⁻⁴⁷

Results from 7 population-based community screening programs in the United States on 463 372 screening mammograms revealed an overall sensitivity of 75.0% and specificity of 92.3%.³⁶ A sensitivity of 75% means that 25% of women (or 25 of 100 women) who were diagnosed with breast cancer had normal mammogram results between 12 and 24 months before their cancer diagnosis (eg, false-negative examination result). This sensitivity of 75% in the community was similar to that reported in the randomized trials (68% to 88%) but the specificity was lower than in most of the trials (range, 82%-93% for Canadian National Breast Screening Study 1 to 98.5% for Health Insurance Plan of New York).⁴⁸

Breast density and age are important predictors of accuracy.³⁶ Ad-

Table 1. Breast Imaging Reporting and Data System Assessment Categories Used in the United States for Mammography Examinations and Associated Likelihood Ratio for Breast Cancer Diagnosis*

Assessment Category	Assessment	Definition	Likelihood Ratio for Breast Cancer Diagnosis†
1	Negative	Breasts appear normal	0.1
2	Benign finding	A negative mammogram result, but the interpreter wishes to describe a finding	0.1
3	Probably benign finding; short-interval follow-up suggested	Lesion with a high probability of being benign noted on mammogram	1.2
0	Need additional imaging evaluation	A lesion is noted for which additional imaging evaluation is needed; used almost always in a screening situation	7.0
4	Suspicious abnormality—biopsy should be considered	A lesion is noted for which the radiologist has sufficient concern to recommend a biopsy	125
5	Highly suggestive of malignancy; appropriate action should be taken	A lesion is noted that has a high probability of being cancer	2200

*Categories also include an assessment of 6, used when there is a known diagnosis of breast cancer before the mammogram.⁵³
†Likelihood ratios for risk of breast cancer diagnosis at first screening mammography.⁵⁴

justed sensitivity ranged from 63% in women with extremely dense breasts to 87% in women with almost entirely fatty breasts; adjusted sensitivity increased with age from 69% in women aged 40 through 49 years to 83% in women aged 80 through 89 years.³⁶ Adjusted specificity increased from 89% in women with extremely dense breasts to 97% in women with almost entirely fatty breasts.³⁶

Guidelines for quality assurance have been issued by several bodies, such as the Commission of the European Communities^{49,50} and the US Mammography Quality Standards Act.⁵¹ The Breast Imaging Reporting and Data System, used in the United States to standardize reports, includes categories for assessment ranging from 0 to 5⁵² (TABLE 1). The associated likelihood ratios for a breast cancer diagnosis for first screen are shown in the table.⁵⁴ Use of the Breast Imaging Reporting and Data System has not eliminated the variability among radiologists⁵⁵ that had been noted before.^{36,57}

Large differences have been noted between the recall rates (or percentage of screening mammograms considered as

positive) of community-based mammography programs in the United States and those in other countries. The recall rate in the United States is twice the recall rate in the United Kingdom (eg, 12.5%-14.4% vs 7.6%), with no difference in cancer detection rate.⁵⁸ Elmore et al⁵⁹ noted comparable differences between North American screening programs and those in other locations, which persisted after adjusting for differences such as age of women screened, use of single vs double reading, and use of 1 vs 2 views of each breast for examinations. Other possible reasons for the regional variability noted include differences in the characteristics of the population screened (eg, presence of risk factors or symptoms) and features of the mammography examination (eg, equipment type and year, technician training).^{59,60} The experience of the physician interpreting the mammograms has also been raised as a possible reason for the variability.⁶⁰⁻⁶² Recommendations vary regarding the minimum number of mammograms that the physician should interpret yearly, from 480 in the United States⁵⁰ to 5000 in the United King-

dom.⁵¹ Educational training has been shown to improve sensitivity with no change in specificity.⁶³ Finally, features of the health care system, including malpractice concerns, financial incentives, quality control procedures, and auditing procedures, may be related to some of the variability.⁶⁰

Double-reading of films, where 2 or more radiologists interpret each film, is offered in some US screening programs and in about half of the other countries that use mammography screening.^{4,58} However, double reading is performed in various ways (eg, 2 interpretations may be completely independent, 2 radiologists may perform interpretations together, or a third reader may serve as a tie-breaker). A systematic review of 10 cohort studies found that double reading increased the cancer detection rate by 3 to 11 per 10000 women screened; recall rates increased or decreased depending on the method of double reading used.⁶⁴ The heterogeneity of the definitions of double reading and of screening practices precludes an accurate assessment of benefits and harms.

Full-Field Digital Mammography and Computer-Aided Detection Programs

Both screen-film mammography and full-field digital mammography use x-rays to obtain images. With screen-film mammography the image is captured on film; with full-field digital mammography the image is captured digitally. Digital images can then be printed on film for viewing, or the images can be interpreted directly from a computer monitor. The digital acquisition process improves logistics and work flow by allowing electronic transmission, storage, and retrieval. Radiologists viewing the image on a monitor can alter the contrast and brightness of the image and magnify specific areas without additional x-ray exposure. Digital imaging also allows easier use of computer-aided detection software. One disadvantage, however, is the increased cost associated with full-field digital mammography.⁴

Three community-based studies have compared full-field digital mammography to screen-film mammography (TABLE 2). Two studies found the sensitivity of full-field digital mammography (64% and 74%) to be less than that of screen-film mammography (79%, 90%), but these studies had a small number of women with breast cancer (42 and 31, respectively) and the display systems and experience of radiologists may have improved since these studies.^{65,66,70} A larger randomized study reported similar cancer detection rates (per all screened), with higher recall rates for full-field digital mammography.⁶⁷ A trial now being conducted, which aimed to enroll 49400 women, will compare the diagnostic accuracy of full-field digital mammography from 4 different manufacturers with that of traditional screen-film mammography.⁷¹

Computer-aided detection programs, which recognize patterns in breast images associated with cancer (FIGURE 1), may potentially help radiologists improve their diagnostic accuracy, but presently data are limited (Table 2).^{68,69,72,73} Computer programs that can mark calcifications, masses, or other potential lesions on the mammogram may increase the number of cancers detected compared with unassisted interpretations. In a study of 12860 women,⁶⁸ computer-aided detection increased radiologists' overall screening recall rate from 6.5% to 7.7% while increasing the number of cancers detected from 41 without computer-aided detection to 49 with the technology. However, in the largest clinical series to date, which included 59139 mammograms interpreted with computer-aided detection and 56432 interpreted without, recall rates and cancer detection rates did not differ significantly.⁶⁹ Computer-aided detection may prove helpful in reducing variability among radiologists of differing expertise. Such programs are used by a small but growing number of mammography facilities in the United States.^{69,74} Medicare and Medicaid allow for additional billing for computer-aided interpretations of mammo-

grams. As technologies continue to improve, larger multisite studies will be needed for more definitive evidence.

Clinical Breast Examination

Although two thirds of US women older than 40 years receive regular screening clinical breast examinations,² few data about the efficacy of clinical breast examinations alone are available from randomized clinical trials. Four randomized trials of mammography included the clinical breast examination in the screened group.^{15,21-23} One of these trials, the Canadian National Breast Screening Study 2 of women aged 50 through 59 years at entry, compared the results of an annual standardized 10- to 15-minute clinical breast examination and breast self-examination with the results of an annual standardized clinical examination and breast self-examination plus mammography (in other words, there was no typical control group that received no screening).²¹ The trial found that breast cancer mortality was similar in the 2 groups of women although yearly mammography in addition to physical examination and breast self-examination detected more small and lymph node-negative breast cancers than did screening with physical examination alone.²¹ Although the sensitivity of screening clinical breast examination was highest at 63% in the National Breast Screening Study 2,⁷⁵ an overall estimate based on all randomized trials calculated sensitivity at 54% (95% confidence interval [CI], 48%-60%) and specificity at 94% (95% CI, 90%-97%).⁷⁶

We suspect that few clinicians in the community setting perform examinations as carefully as those in the Canadian trial, so accuracy may be lower in the community setting. Results reported from community practices showed sensitivity ranging from 28% to 36%⁷⁷⁻⁷⁹ (TABLE 3). In one study, two fifths of physicians (34 out of 80) who performed a screening breast examination on manufactured breast models used no discernible systematic search pattern at all.⁸⁰ Sensitivity of examinations improved by spending more time^{80,81} and by using a thorough, sys-

tematic technique.^{76,82} However, the number of false-positive examinations may increase with training.⁸¹

In randomized controlled trials, noting an abnormality on a screening clinical breast examination in an asymptomatic average-risk woman increased the likelihood of breast cancer (likelihood ratio [LR], 10.6; 95% CI, 5.8-19.2).⁷⁶ However, in community practice, an abnormal screening breast examination result was associated with an LR of 2.1,⁷⁶ substantially lower than that of women in the same practice presenting with a breast abnormality (LR, 24).⁸³ Noting a suspicious abnormality on a screening mammogram was as-

sociated with LRs ranging from 7 to 2200 (Table 1).⁸⁴

Breast Self-examination

Breast self-examination is appealing as a patient-centered, noninvasive procedure that allows women to become comfortable with their own bodies. However, the extent of current practice is thought to be low.⁴ About one third of US women regularly perform breast self-examination, and the estimated sensitivity is low (20% to 30%).⁸⁵ Among respondents to the Women Physicians' Health Study (N=4501), only 21% reported performing monthly breast self-examination.⁸⁶

Training in breast self-examination, while associated with increased accuracy of detection of lumps in breast tissue, has been associated with increased rates of false-positive findings and thus diminished specificity.⁸⁷⁻⁹⁰ In addition, there is evidence casting doubt on the benefits. A large randomized controlled trial in Shanghai, China, of 266 064 women working in textile factories provided half of the women with intensive initial instruction, including practice with breast models, as well as regular reminders and practice examinations under supervision once every 6 months for 5 years. This study found no positive effect of breast self-examination on breast can-

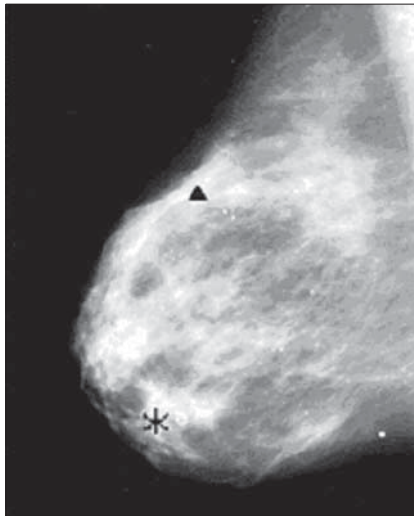
Table 2. Studies of Full-Field Digital Mammography and Computer-Aided Detection Programs in Community Screening Settings

Source/ Location	Population/Design	No./Total (%)				Comment
		Breast Cancer Detection Among Women With Cancer (Sensitivity)		Recall Rate		
		Screen Film	Digital	Screen Film	Digital	
Lewin et al, ⁶⁵ 2002 United States	6736 Examinations for 4489 women Women received both examinations	33/42 (78.6)	27/42 (64.3)	1007/6736 (14.9)	799/6736 (11.9)	2 Institutions
Skaane et al, ⁶⁶ 2003 Norway	3683 Women Women received both examinations	28/31 (90.3)	23/31 (74.2)	128/3683 (3.5)	168/3683 (4.6)	8 Radiologists Initial recall rate before consensus meeting and comparison with previous mammograms available was 7% for screen film and 12% for digital
Skaane and Skjennald, ⁶⁷ 2004 Norway	25 263 Women Randomized to 1 examination	3.0 (45-49 y) 2.5 (50-69 y)	3.7 (45-49 y) 3.8 (50-69 y)	8 Radiologists 4 independent radiologists interpreted all FFDM images for first 4 mo Initial recall rate before consensus meeting and comparison with prior examinations available was 9.8%-10.2% for screen film and 13.7%-14.0% for digital
		Breast Cancer Detection Rate per 1000 Women Screened		Recall Rate, %		
		No CAD	CAD	No CAD	CAD	
Freer and Ulissey, ⁶⁸ 2001 United States	12 860 Women Each mammogram was initially interpreted without CAD, followed by reevaluation of areas marked by CAD	3.2/1000	3.8/1000	6.5	7.7	2 Radiologists
Gur et al, ⁶⁹ 2004 United States	56 432 Examinations without CAD 59 139 Examinations with CAD Pre-post study design after introduction of CAD into a clinical practice setting	3.49/1000	3.55/1000	11.39	11.40	24 Radiologists No adjustment for patient characteristics

Abbreviations: CAD, Computer-aided Detection; FFDM, full-field digital mammography; ellipses indicate that data are not provided on cancer detection among the subgroup of women with cancer (sensitivity). Data presented on cancer detection rates among all women screened were similar for digital and screen film.

cer mortality after 10 years of follow-up but almost double the rate of biopsies due to false-positive findings (1.8% of women

Figure 1. Computer-Aided Detection Markers of Breast Imaging Screening Mammography



Examination with subtle focal asymmetric density on mammography examination of the right breast, mediolateral oblique view. The density is marked by the computer-aided detection program, with a star to call attention to a possible cancer. Breast biopsy confirmed infiltrating ductal carcinoma. The triangle notes calcification in the same breast.

in the instruction group vs 1.0% of women in the control group).^{88,89} The study results should be interpreted with caution because approximately 40% of women in the trial were in their 30s. Also, it is possible that a 10-year follow-up was not long enough to see an effect on breast cancer mortality.

A meta-analysis of the effect of regular breast self-examination on breast cancer mortality or rates of advanced breast cancer (a marker of death) was performed on 20 observational studies and 3 clinical trials.⁹⁰ Bias and confounding may affect the results of studies of women with breast cancer who reported practicing self-examination before diagnosis. No difference in death rate was noted in studies of women who detected their cancers during self-examination (pooled relative risk [RR], 0.90; 95% CI, 0.72-1.12), and no mortality differences were noted in trials of training (pooled RR, 1.01; 95% CI, 0.92-1.12).

Magnetic Resonance Imaging

Although screen-film mammography and full-field digital mammography are the only imaging tools explicitly approved or grandfathered in for breast cancer screening by the US Food and

Drug Administration (FDA), other modalities are under study.⁷³ Those approved by the FDA for diagnostic purposes (not screening) include MRI, ultrasound, scintimammography, thermography, and electrical impedance imaging. Although mammography uses x-ray and sonography uses sound waves to create images, MRI produces images from the combination of a strong magnetic field, radio waves, and computer processing (FIGURE 2).

Screening MRI may be helpful for women for whom mammography is not optimal, such as young women at substantially increased risk for breast cancer because of known *BRCA1* or *BRCA2* mutations. Available data are limited to studies of test characteristics in women at high risk (TABLE 4),⁹¹⁻⁹⁸ and the impact on breast cancer mortality has not been determined. Both retrospective and prospective cohorts have been described. The small number of cases with breast cancer in these studies (the range among studies was 3 to 45 women) means that estimates of sensitivity were not precise. Nevertheless, every study reported higher sensitivity for breast MRI than for mammography, ultrasound, or both.⁹¹⁻⁹⁸ The largest study re-

Table 3. Studies of Screening Clinical Breast Examination Performance in Clinical Trials and Community Settings in the United States

Population (Reference)	No. of Women Screened	Participants and Setting	Years	% (95% Confidence Interval)		Comments
				Sensitivity	Specificity	
Pooled analysis of clinical trials ⁷⁶	6419 With cancer (429 248 with no cancer)	Participants in randomized clinical trial (Health Insurance Plan of NY ⁷⁴), Canadian Breast Screening Study, and nonrandomized controlled trial	1963-1988	54.1 (48.3-59.8)	94.0 (90.2-96.9)	
US National Breast and Cervical Cancer Early Detection Program ⁷⁷	589 048	Low-income women (age ≥ 40 y) enrolled in National Breast and Cervical Cancer Early Detection Program	1995-1998	36.1	96.2	Data shown are for asymptomatic patients only
US Health Plan in Pacific Northwest ⁷⁸	468	Health plan enrollees (age ≥ 40 y) in western Washington attending breast cancer screening program	1988-1994	35.3	Not available	Nurse examiners in dedicated screening program at single health plan
Columbia Medical Center, Department of Radiology ⁷⁹	11 130 (27 825 Screens)	Urban women attending single academic screening program	1995-2000	27.6	99.4	Single radiologist examiner aware of mammography results during examination 78% Had a normal examination 1 mo before the study

ported on 1909 women at increased risk in the Netherlands, with 45 women diagnosed with cancer who had all screening examinations.⁹⁶ The sensitivity of clinical breast examination, mammography, and MRI in this study was 17.9%, 40%, and 71%, respectively. The overall discriminating capacity of MRI was significantly better compared with mammography as assessed by receiver operating characteristic curves (area under the curve 0.827 for MRI vs 0.686 for mammography).

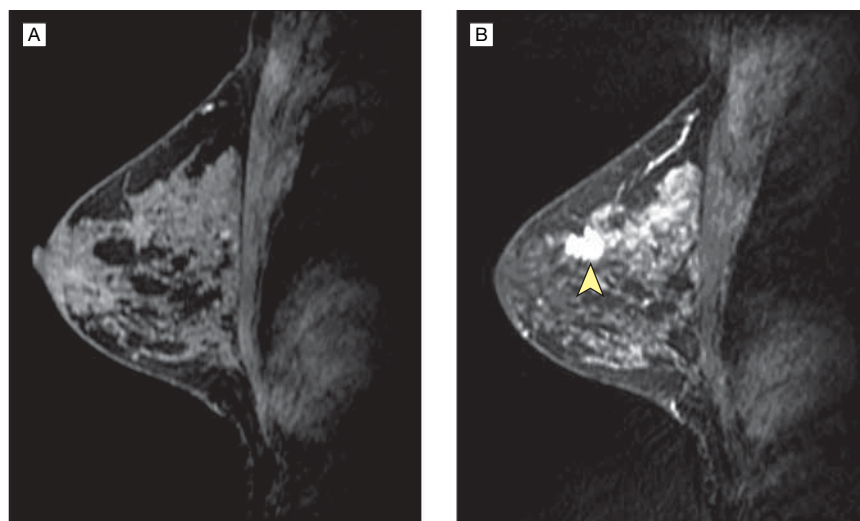
Specificity of MRI tends to be lower than that of mammography; however, data are not consistently presented and specificity is not always easy to calculate. In the study of 1909 women in the Netherlands, the specificity of clinical breast examination, mammography, and MRI was 98.1%, 95.0%, and 89.8%, respectively,⁹⁶ and the authors noted that screening with MRI led to twice as many unneeded additional examinations (420 vs 207) and 3 times as many unneeded biopsies (24 vs 7) as did screening with mammography.⁹⁶ Warner et al⁹⁸ reported a substantial recall rate in the first round of MRI screening (26%), which decreased to 10% in the third round of

MRI screening. Additional studies of MRI screening among high-risk women are under way.⁹⁹

Magnetic resonance imaging has not been studied in the general population as a screening tool, and the results from

MRI screening of high-risk women may not apply to women at average risk. The high cost of MRI (approximately 10 times the cost of mammography) and its relatively low specificity (compared with mammography) probably prohibit its

Figure 2. Magnetic Resonance Imaging Examination of the Breast



A 35-year-old woman at increased risk for breast cancer based on a strong family history of breast cancer obtained screening magnetic resonance imagery examination, before, A, and after contrast injection, B. Examination revealed a 1.3 cm mass in her left breast (arrowhead), occult on screening mammography examinations. Biopsy confirmed infiltrating ductal carcinoma. (See also <http://depts.washington.edu/gim/faculty/elmore.htm> for example of 3-dimensional rotation of magnetic resonance imagery views.)

Table 4. Comparative Results of Nonrandomized Studies of Screening Mammography, Magnetic Resonance Imaging, and Ultrasound in Screening Women at Increased Risk for Breast Cancer

Source	Site	Study Design	Patient Characteristics			No./Total (%)			Cancer Yield From MRI Alone	Biopsies Performed as a Result of MRI	PPV of Biopsies Performed Based on MRI, %
			Total No. of Women	Age, y Mean (Range)	No. (%) Known Mutation Carriers*	Sensitivity					
						Mammography	MRI	Ultrasound			
Kuhl et al, ⁹¹ 2000	Germany	Prospective	192	39 (18-65)	35 (18)	3/9 (33)	9/9 (100)	3/9 (33)	6/192 (3)	14/192 (7.3)	64†
Tilanus-Linthorst et al, ⁹² 2000‡	The Netherlands	Prospective	109	42 (22-68)	12 (11)	0	3/3 (100)	...	3/109 (2.8)	5/109 (4.6)	60
Podo et al, ⁹⁵ 2002	Italy	Prospective	105	46 (25-77)	...	1/8 (13)	8/8 (100)	1/8 (13)	7/105 (6.7)	9/105 (8.6)	89
Morris et al, ⁹⁴ 2003	United States	Retrospective	367	50 (23-82)	19 (5)	NA	14/14 (100)	...	14/367 (3.8)	59/367 (16)	24
Warner et al, ⁹⁸ 2004	Canada	Prospective	236	47 (26-65)	236 (100)	8/22 (36)	17/22 (77)	7/21 (33)§	7/236 (3)	...	46
Kriege et al, ⁹⁶ 2004¶	The Netherlands	Prospective	1909	40 (19-72)	358 (19)	18/45 (40)	32/45 (71)	...	22/1909 (2.2)	56/1909 (2.9)	57

Abbreviations: MRI, magnetic resonance imaging; NA, not applicable; PPV, positive predictive value. Ellipses indicate that data were not available.

*Data are shown for women with known positive results.

†Reported PPV based on 105 women with valid 1-year of follow-up.

‡To detect 3 breast cancers in 109 women, the investigators performed 193 MRI examinations, 51 ultrasounds, 29 fine-needle aspiratory cytology, and 2 benign excision biopsies.

§One patient who had cancer detected only by MRI did not receive ultrasound.

||All women in this study had a normal mammogram result for inclusion.

¶The total number of tumors was 51, but the results in the article were calculated based on 45.

Box. Breast Cancer–Related Information

Calculation of the risk of a breast cancer diagnosis and death at the level of individual women:

<http://bcra.nci.nih.gov/brc/start.htm> (Gail model)

<http://astor.som.jhmi.edu/brcapro> (Gail Model, Claus Model, and a model that predicts the probability of carrying a *BRCA1* or *BRCA2* mutation)

<http://yourcancerrisk.harvard.edu/>

Breast Self-Examination Tutorials

<http://www.komen.org/bse>

<http://www.breastselfexam.ca>

National Guidelines for Breast Cancer Screening

<http://www.guidelines.gov>

Randomized Clinical Trials of New Modalities in Breast Cancer Screening

<http://www.clinicaltrials.gov>

http://www.acrin.org/current_protocols.html

routine use for screening general populations. Also, MRI is time-consuming, requires intravenous contrast administration, and may be problematic for claustrophobic patients.

Ultrasound

Ultrasound, frequently used as a targeted diagnostic examination focusing on a specific area of concern,¹⁰⁰ may help distinguish between cyst and solid masses and also between benign and malignant masses.¹⁰¹ Breast ultrasound data are available from diagnostic populations, with screening studies limited to women with dense breasts on mammography or at increased risk for breast cancer.¹⁰² Although ultrasound may detect 3 to 4 additional breast cancers per 1000 women in these increased-risk populations,^{93,100,101,103-108} there are no data on the use of screening ultrasound in the general population. Breast ultrasound has limitations as a potential screening tool because it requires a well-trained skilled

operator. Examination techniques are not standardized, interpretation criteria are variable, and breast ultrasound does not consistently detect microcalcifications. Preliminary data suggest a higher rate of false-positive examination results with ultrasound than with mammography alone.^{100,104-106} For example, the false-positive rate (based on solid lesion for ultrasound) ranged from 2.4% to 12.9% for ultrasound and 0.7% to 6% for mammography.¹⁰²

National Screening Guidelines

All groups recommend screening mammography for women aged 50 through 69 years.⁷ Within the United States, all recommend it for women in their 40s, but vary in the screening intervals recommended and encourage “informed decision making” with all women about the choice. Being older than 70 years should not preclude women from continuing to undergo screening; however, decisions regarding continued screening should include life expectancy and health status.^{7,41} International policies differ with respect to the target age group to be screened, the intervals between screening, the number of mammographic views taken per breast, and the screening modalities recommended.⁴ Clinical breast examination is recommended by some,¹⁰⁹ but not all, groups. Most national groups no longer recommend breast self-examination, but some encourage women to become familiar with the contour of their own breasts.^{42,109} Other imaging modalities, such as MRI and ultrasound, are not recommended for screening the general population. (See <http://www.Guidelines.gov>.)

Benefits and Harms

The primary goal of breast cancer screening is to reduce subsequent breast cancer mortality through early detection. Theoretically this should translate into reduced morbidity from the disease.¹¹⁰ In addition, many women report feeling reassured by screening,¹¹¹ especially after having a so-called normal screen result.¹¹²

Possible harms include pain and discomfort, especially noted during com-

pression of breast tissue during mammography. Compression of breast tissue reduces motion artifact and improves image quality. Reports of the level of discomfort, however, vary widely.¹¹³⁻¹¹⁵ Anxiety about screening is another concern.^{116,117} Breast cancer screening yields both false-positive and false-negative results. False-positive results have been associated with anxiety, additional costs, and morbidity.^{117,118} After 10 years of annual screening in the United States, it is estimated that 1 in 2 women will have at least 1 false-positive mammogram result, and 1 in 5 women will have at least 1 false-positive clinical breast examination result.¹¹⁸ False-negative mammography examinations occur in approximately 20% to 40% of women with breast cancer.³⁶

Overdiagnosis and overtreatment of clinically insignificant disease is possible, especially ductal carcinoma in situ noted by mammography.^{119,120} Theoretical concerns about radiation-induced breast cancer from exposure to repeated mammography have been raised, but the potential benefits are thought to outweigh the risks. For example, the benefit-to-harm ratio is estimated to be 48.5 lives saved per 1 life lost due to radiation exposure.¹²¹ A mortality paradox has been noted in women aged 40 through 49 years, whereby increased mortality is noted among women screened for the first 3 to 10 years after the initiation of screening.¹²²⁻¹²⁴ Tumor dissemination after needle biopsy has also been suggested although the clinical significance is unclear.¹²⁵

Observer variability among radiologists who interpret mammography examinations has been noted both in a test situation and in community practices.^{55-57,126} A decision to perform a breast biopsy may depend heavily on the radiologist's interpretation; therefore, interpretive variability can directly affect patient management.¹²⁶

The benefit-to-harm ratio of screening increases as women age because screening accuracy improves and prevalence of breast cancer increases. Younger women, however, have more potential years of life to be gained from screening.

Communicating With Patients

Effective communication of information on benefits and harms is challenging. Multiple studies document inaccurate or incomplete comprehension of risk information, cognitive biases that affect how patients process risk information, and poor communication skills on the part of physicians.¹²⁷⁻¹³⁰ Use of frequencies with specific reference groups (“23 out of 1000 women your age”) instead of percentages (“0.23%”) may facilitate the comparison of small risks.¹³¹⁻¹³³ Presentation of both positive (“23 in 1000 women your age will develop breast cancer”) and negative framing (“977 in 1000 women your age will not develop breast

cancer”) can reduce biases in decision making.^{130,132} Visual aids, such as bar graphs or pie charts, can increase the comprehension and saliency of information.¹³⁴ Risk-prediction models can be used to calculate the probability of being diagnosed with breast cancer (BOX). Effective physician-patient partnership, for which gaps in comprehension are frequently assessed and resolved may have the greatest impact on patients’ understanding of information.¹³⁵

CONCLUSION

Reviews of breast cancer screening usually concentrate on the results of randomized trials of mammography to re-

duce breast cancer mortality. We have emphasized data on effectiveness in the community setting among the general population of women, which can often be different from the ideal setting of a randomized trial or the setting of a study among high-risk women. We have also emphasized the challenge of evaluating new screening modalities. Newer screening tests such as MRI and ultrasound have been studied in women at increased risk of breast cancer (eg, carriers of *BRCA1* or *BRCA2* mutations). None of the newer tests has been evaluated for its effect on breast cancer mortality in the general population and no data support screening to

Table 5. Summary of Breast Cancer Screening Modalities for General Average-Risk Populations

Screening Modality	Test Characteristics Investigated in the General Population by Type of Study Design		Effect on Breast Cancer Studied	FDA Approval	Medicare Reimbursement, \$*	HCPCS Codes	Comments
	RCT	Cohort or Other Design in Community Setting					
Screen-film mammography	Yes	Yes	Yes	Yes†	90	76 092	Most extensively studied screening modality
Clinical breast examination	Yes	Yes	Yes	NA	39	G0101	Higher sensitivity associated with longer duration of examinations Sensitivity noted in community setting appears lower than trials
Breast self-examination	Yes	Yes	Yes	NA	NA	NA	No positive effect on mortality noted and increased biopsies because of false-positive findings
Digital mammography	Yes	Yes	No	Yes	144	G0202	Improves logistical practice, facilitates use of computer-aided detection programs Comparable cancer detection rate to screen-film mammography The impact on recall rates is unclear
Computer-aided detection	No	Yes	No	Yes	21	76 083	Cost in addition to that of mammography May help radiologists improve accuracy but no data to date to definitively support
Ultrasound	No	No	No	No	74	76 645	Requires skilled operator, examination technique not standardized, interpretation criteria variable, does not detect microcalcifications
Magnetic resonance imaging	No	No	No	No	1108	76 094	More sensitive and slightly less specific than mammography in women at increased risk No radiation Requires intravenous contrast, time-consuming, may not be feasible for some women, such as those with pacemaker, aneurysm clips, or claustrophobia

Abbreviation: FDA, US Food and Drug Administration; HCPCS, Healthcare Common Procedure Code System; NA, not applicable; RCT, randomized clinical trial.

*Represents the 2004 Medicare reimbursement rate for Seattle, Wash.¹³⁶ The cost for computer-aided detection is in addition to the cost for the mammography examination.

†Screen-film mammography was grandfathered in by the FDA.

general population with these technologies. Careful evaluation of newer modalities in the populations for which they will be used is critical, especially since these modalities are usually more expensive than current approaches and the risk of increased false-positives is present. An overview of breast cancer screening modalities is shown in TABLE 5.

Most national groups recommend screening with mammography, with or without clinical breast examination, beginning at 40 years of age. Data on clinical breast examination as performed in the community suggest a lower level of cancer detection than would be anticipated from trials. Breast self-examination is no longer recommended by most expert groups. Limitations and potential harms have been identified for all existing screening tools. Quality control needs to be emphasized for established screening methods.

Author Contributions: Dr Elmore had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

Study concept and design: Elmore, Lehman, Armstrong, Fletcher.

Analysis and interpretation of data: Elmore, Lehman, Armstrong, Fletcher.

Drafting of the manuscript: Elmore, Lehman.

Critical revision of the manuscript for important intellectual content: Elmore, Lehman, Armstrong, Fletcher.

Administrative, technical, or material support: Elmore, Lehman.

Study supervision: Elmore, Lehman, Armstrong.

Financial Disclosure: Dr Armstrong has served as an expert witness about breast cancer risk. No other authors reported any disclosures.

Funding/Support: Dr Elmore was supported by grant K05-CA10469 from the National Cancer Institute. Dr Armstrong was supported by the Robert Wood Johnson Generalist Physician Award.

Role of the Sponsor: The Robert Wood Johnson Foundation did not participate in the design and conduct of the study; in the collection, analysis, and interpretation of the data; or in the preparation, review, or approval of the manuscript.

Disclaimer: The views expressed in this article are those of the authors and do not necessarily reflect the opinion of the Robert Wood Johnson Foundation.

Acknowledgment: We thank Carol Munch and Sue Peacock for their administrative assistance.

REFERENCES

- Adair FE. Clinical manifestations of early cancer of the breast. *N Engl J Med.* 1933;208:1250-1255.
- Blackman DK, Bennett EM, Miller DS. Trends in self-reported use of mammograms (1989-1997) and papanicolaou tests (1991-1997): Behavioral Risk Factor Surveillance System. *MMWR CDC Surveill Summ.* 1999;48:1-22.
- Weir HK, Thun MJ, Hankey BF, et al. Annual report to the nation on the status of cancer, 1975-2000, featuring the uses of surveillance data for cancer prevention and control. *J Natl Cancer Inst.* 2003;95:1276-1299.
- Vainio H, Bianchini F. *Breast Cancer Screening: International Agency for Research on Cancer (IARC) Handbooks of Cancer Prevention.* Vol 7. Lyon, France: IARC Press; 2002.
- US Preventive Services Task Force. Screening for breast cancer: recommendations and rationale. *Ann Intern Med.* 2002;137:344-346.
- US Preventive Services Task Force. *Guide to Clinical Preventive Services.* 2nd ed. Alexandria, Va: International Medical Publishing Inc; 1996.
- Fletcher SW, Elmore JG. Clinical practice: mammographic screening for breast cancer. *N Engl J Med.* 2003;348:1672-1680.
- Last JM. *A Dictionary of Epidemiology.* New York, NY: Oxford University Press; 1995:52.
- Peto R, Boreham J, Clarke M, Davies C, Beral V. UK and USA breast cancer deaths down 25% in year 2000 at ages 20-69 years. *Lancet.* 2000;355:1822.
- Early Breast Cancer Trialists' Collaborative Group. Tamoxifen for early breast cancer: an overview of the randomised trials. *Lancet.* 1998;351:1451-1467.
- Early Breast Cancer Trialists' Collaborative Group. Polychemotherapy for early breast cancer: an overview of the randomised trials. *Lancet.* 1998;352:930-942.
- Barlow WE, Lehman CD, Zheng Y, et al. Performance of diagnostic mammography in women with signs or symptoms of breast cancer. *J Natl Cancer Inst.* 2002;94:1151-1159.
- Humphrey LL, Helfand M, Chan BK, Woolf SH. Breast cancer screening: a summary of the evidence for the U.S. Preventive Services Task Force. *Ann Intern Med.* 2002;137:347-360.
- Shapiro S, Venet W, Strax P, Venet L. Current results of the breast cancer screening randomized trial: the health insurance plan (HIP) of greater New York study. In: Day NE, Miller AB, eds. *Screening for Breast Cancer.* Toronto, Ontario: Hans Harbor; 1988:3-15.
- Shapiro S, Venet W, Strax P, Venet L. *Periodic Screening for Breast Cancer: The Health Insurance Plan Project and Its Sequelae, 1963-1986.* Baltimore, Md: Johns Hopkins University Press; 1988.
- Nystrom L, Andersson I, Bjurstam N, Frisell J, Nordenskjold B, Rutqvist LE. Long-term effects of mammography screening: updated overview of the Swedish randomised trials. *Lancet.* 2002;359:909-919.
- Andersson I, Janzon L. Reduced breast cancer mortality in women under age 50: updated results from the Malmo Mammographic Screening Program. *J Natl Cancer Inst Monogr.* 1997;63-67.
- Tabar L, Fagerberg G, Chen HH, et al. Efficacy of breast cancer screening by age: new results from the Swedish Two-County Trial. *Cancer.* 1995;75:2507-2517.
- Frisell J, Lidbrink E. The Stockholm Mammographic Screening Trial: risks and benefits in age group 40-49 years. *J Natl Cancer Inst Monogr.* 1997:49-51.
- Bjurstam N, Bjorneld L, Duffy SW, et al. The Gothenburg Breast Cancer Screening Trial: preliminary results on breast cancer mortality for women aged 39-49. *J Natl Cancer Inst Monogr.* 1997:53-55.
- Miller AB, To T, Baines CJ, Wall C. Canadian National Breast Screening Study-2: 13-year results of a randomized trial in women aged 50-59 years. *J Natl Cancer Inst.* 2000;92:1490-1499.
- Miller AB, To T, Baines CJ, Wall C. The Canadian National Breast Screening Study-1: breast cancer mortality after 11 to 16 years of follow-up: a randomized screening trial of mammography in women age 40 to 49 years. *Ann Intern Med.* 2002;137:305-315.
- Alexander FE, Anderson TJ, Brown HK, et al. 14 years of follow-up from the Edinburgh randomised trial of breast-cancer screening. *Lancet.* 1999;353:1903-1908.
- Olsen O, Gøtzsche PC. Cochrane review on screening for breast cancer with mammography. *Lancet.* 2001;358:1340-1342.
- Olsen O, Gøtzsche PC. Screening for breast cancer with mammography. Oxford, England: Cochrane Library, Update Software; Issue 3: 2003.
- The Benefit of Population Screening for Breast Cancer with Mammography.* The Hague, the Netherlands: Health Council of the Netherlands; 2002. Publication 2002/3E.
- Porter PL, El-Bastawissi AY, Mandelson MT, et al. Breast tumor characteristics as predictors of mammographic detection: comparison of interval- and screen-detected cancers. *J Natl Cancer Inst.* 1999;91:2020-2028.
- Crosier M, Scott D, Wilson RG, Griffiths CD, May FE, Westley BR. Differences in Ki67 and c-erbB2 expression between screen-detected and true interval breast cancers. *Clin Cancer Res.* 1999;5:2682-2688.
- Klemi PJ, Joensuu H, Toikkanen S, et al. Aggressiveness of breast cancers found with and without screening. *BMJ.* 1992;304:467-469.
- Gilliland FD, Joste N, Stauber PM, et al. Biologic characteristics of interval and screen-detected breast cancers. *J Natl Cancer Inst.* 2000;92:743-749.
- Groenendijk RP, Bult P, Tewarie L, et al. Screen-detected breast cancers have a lower mitotic activity index. *Br J Cancer.* 2000;82:381-384.
- Joensuu H, Lehtimäki T, Holli K, et al. Risk for distant recurrence of breast cancer detected by mammography screening or other methods. *JAMA.* 2004;292:1064-1073.
- Zahl PH, Strand BH, Maehlen J. Incidence of breast cancer in Norway and Sweden during introduction of nationwide screening: prospective cohort study. *BMJ.* 2004;328:921-924.
- Welch HG. *Should I Be Tested for Cancer? Maybe Not and Here's Why.* Berkeley: University of California Press; 2004.
- Ries LAG, Eisner MP, Kosary CL, et al. *SEER Cancer Statistics Review, 1975-2001.* Bethesda, Md: National Cancer Institute; 2004.
- Carney PA, Miglioretti DL, Yankaskas BC, et al. Individual and combined effects of age, breast density, and hormone replacement therapy use on the accuracy of screening mammography. *Ann Intern Med.* 2003;138:168-175.
- Moss S; Trial Steering Group. A trial to study the effect on breast cancer mortality of annual mammographic screening in women starting at age 40. *J Med Screen.* 1999;6:144-148.
- Van Dijk JA, Verbeek AL, Beex LV, et al. Mammographic screening after the age of 65 years: evidence for a reduction in breast cancer mortality. *Int J Cancer.* 1996;66:727-731.
- Satariano WA, Ragland DR. The effect of comorbidity on 3-year survival of women with primary breast cancer. *Ann Intern Med.* 1994;120:104-110.
- Walter LC, Eng C, Covinsky KE. Screening mammography for frail older women: what are the burdens? *J Gen Intern Med.* 2001;16:779-784.
- Walter LC, Covinsky KE. Cancer screening in elderly patients: a framework for individualized decision making. *JAMA.* 2001;285:2750-2756.
- Smith R, Saslow D, Sawyer KA, et al. American Cancer Society guidelines for breast cancer screening: update 2003. *CA Cancer J Clin.* 2003;53:141-169.
- Kerlikowske K, Grady D, Barclay J, Sickles EA, Eaton A, Ernster V. Positive predictive value of screening mammography by age and family history of breast cancer. *JAMA.* 1993;270:2444-2450.
- Brown ML, Houn F, Sickles EA, Kessler LG. Screening mammography in community practice: positive predictive value of abnormal findings and yield of follow-up diagnostic procedures. *AJR Am J Roentgenol.* 1995;165:1373-1377.
- Ma L, Fishell E, Wright B, Hanna W, Allan S, Boyd NF. Case-control study of factors associated with fail-

- ure to detect breast cancer by mammography. *J Natl Cancer Inst.* 1992;84:781-785.
46. Harvey JA, Fajardo LL, Innis CA. Previous mammograms in patients with palpable breast carcinoma: retrospective vs blinded interpretation. *AJR Am J Roentgenol.* 1993;161:1167-1172.
 47. Ikeda DM, Andersson I, Wattsgård C, Janzon L, Linell F. Interval carcinomas in the Malmö Mammographic Screening Trial: radiographic appearance and prognostic considerations. *AJR Am J Roentgenol.* 1992;159:287-294.
 48. Fletcher SW, Black W, Harris R, Rimer BK, Shapiro S. Report of the International Workshop on Screening for Breast Cancer. *J Natl Cancer Inst.* 1993;85:1644-1656.
 49. Commission of the European Communities. *European Guidelines for Quality Assurance in Mammography Screening.* 3rd ed. Luxembourg: Office for Official Publications of the European Communities; 2001.
 50. National Health Service Breast Screening Programme. *Guidelines on Quality Assurance Visits.* Sheffield, England; 1998. NHSBSP Publication 40. Available at: <http://www.cancerscreening.nhs.uk/breastscreen/publications/qa-12.html>. Accessibility verified February 2, 2005.
 51. Quality mammography standards—FDA; final rule. *Federal Register.* 1997;62:55852-55994.
 52. American College of Radiology. *Illustrated Breast Imaging Reporting and Data System (BI-RADS).* 3rd ed. Reston, Va: American College of Radiology; 1998.
 53. D'Orsi CJ, Bassett LW, Feig SA, et al. *Breast Imaging Reporting and Data System.* 3rd ed. Reston, Va: American College of Radiology; 1998.
 54. Kerlikowske K, Smith-Bindman R, Ljung BM, Grady D. Evaluation of abnormal mammography results and palpable breast abnormalities. *Ann Intern Med.* 2003;139:274-284.
 55. Kerlikowske K, Grady D, Barclay J, et al. Variability and accuracy in mammographic interpretation using the American College of Radiology Breast Imaging Reporting and Data System. *J Natl Cancer Inst.* 1998;90:1801-1809.
 56. Elmore JG, Wells CK, Lee CH, Howard DH, Feinstein AR. Variability in radiologists' interpretations of mammograms. *N Engl J Med.* 1994;331:1493-1499.
 57. Beam CA, Layde PM, Sullivan DC. Variability in the interpretation of screening mammograms by US radiologists: findings from a national sample. *Arch Intern Med.* 1996;156:209-213.
 58. Smith-Bindman R, Chu PW, Miglioretti DL, et al. Comparison of screening mammography in the United States and the United Kingdom. *JAMA.* 2003;290:2129-2137.
 59. Elmore JG, Nakano CY, Koepsell TD, Desnick LM, D'Orsi CJ, Ransohoff DF. International variation in screening mammography interpretations in community-based programs. *J Natl Cancer Inst.* 2003;95:1384-1393.
 60. Elmore JG, Miglioretti DL, Carney PA. Does practice make perfect when interpreting mammography? II. *J Natl Cancer Inst.* 2003;95:250-252.
 61. Esserman L, Cowley H, Eberle C, et al. Improving the accuracy of mammography: volume and outcome relationships. *J Natl Cancer Inst.* 2002;94:369-375.
 62. Beam CA, Conant EF, Sickles EA. Association of volume and volume-independent factors with accuracy in screening mammogram interpretation. *J Natl Cancer Inst.* 2003;95:282-290.
 63. Linver MN, Paster SB, Rosenberg RD, Key CR, Stidley CA, King WV. Improvement in mammography interpretation skills in a community radiology practice after dedicated teaching courses: 2-year medical audit of 38,633 cases. *Radiology.* 1992;184:39-43.
 64. Dinnes J, Moss S, Melia J, Blanks R, Song F, Kleijnen J. Effectiveness and cost-effectiveness of double reading of mammograms in breast cancer screening: findings of a systematic review. *Breast.* 2001;10:455-463.
 65. Lewin JM, D'Orsi CJ, Hendrick RE, et al. Clinical comparison of full-field digital mammography and screen-film mammography for detection of breast cancer. *AJR Am J Roentgenol.* 2002;179:671-677.
 66. Skaane P, Young K, Skjennald A. Population-based mammography screening: comparison of screen-film and full-field digital mammography with soft-copy reading—Oslo I Study. *Radiology.* 2003;229:877-884.
 67. Skaane P, Skjennald A. Screen-film mammography versus full-field digital mammography with soft-copy reading: randomized trial in a population-based screening program—the Oslo II Study. *Radiology.* 2004;232:197-204.
 68. Freer TW, Ulissey MJ. Screening mammography with computer-aided detection: prospective study of 12,860 patients in a community breast center. *Radiology.* 2001;220:781-786.
 69. Gur D, Sumkin JH, Rockette HE, et al. Changes in breast cancer detection and mammography recall rates after the introduction of a computer-aided detection system. *J Natl Cancer Inst.* 2004;96:185-190.
 70. Lewin JM, Hendrick RE, D'Orsi CJ, et al. Comparison of full-field digital mammography with screen-film mammography for cancer detection: results of 4,945 paired examinations. *Radiology.* 2001;218:873-880.
 71. Digital versus film-screen mammography [ACRIN protocol 6652]. Philadelphia, Pa: American College of Radiology Imaging Network (ACRIN). October 27, 2003. Available at: http://www.acrin.org/current_protocols.html. Accessibility verified February 2, 2005.
 72. Huo Z, Giger ML, Vyborny CJ, Metz CE. Breast cancer: effectiveness of computer-aided diagnosis observer study with independent database of mammograms. *Radiology.* 2002;224:560-568.
 73. Nass S, Henderson C. *Mammography and Beyond: Developing Technologies for the Early Detection of Breast Cancer.* Washington, DC: National Academy Press; 2001.
 74. Hendrick RE, Cutter GR, Berns EA, et al. Community-based mammography practice: services, charges, and interpretation methods. *AJR Am J Roentgenol.* 2005;184:433-438.
 75. Fletcher SW, Black W, Harris R, Rimer BK, Shapiro S. Report of the International Workshop on Screening for Breast Cancer. *J Natl Cancer Inst.* 1993;85:1644-1656.
 76. Barton MB, Harris R, Fletcher SW. Does this patient have breast cancer? the screening clinical breast examination: should it be done? how? *JAMA.* 1999;282:1270-1280.
 77. Bobo JK, Lee NC, Thames SF. Findings from 752,081 clinical breast examinations reported to a national screening program from 1995 through 1998. *J Natl Cancer Inst.* 2000;92:971-976.
 78. Oestreicher N, White E, Lehman CD, Mandelson MT, Porter PL, Taplin SH. Predictors of sensitivity of clinical breast examination (CBE). *Breast Cancer Res Treat.* 2002;76:73-81.
 79. 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;225:165-175.
 80. Fletcher SW, O'Malley MS, Bunce LA. Physicians' abilities to detect lumps in silicone breast models. *JAMA.* 1985;253:2224-2228.
 81. 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.
 82. Saunders KJ, Pilgrim CA, Pennypacker HS. Increased proficiency of search in breast self-examination. *Cancer.* 1986;58:2531-2537.
 83. Barton MB, Elmore JG, Fletcher SW. Breast symptoms among women enrolled in a health maintenance organization: frequency, evaluation, and outcome. *Ann Intern Med.* 1999;130:651-657.
 84. Kerlikowske K, Grady D, Barclay J, Sickles EA, Ernster V. Likelihood ratios for modern screening mammography: risk of breast cancer based on age and mammographic interpretation. *JAMA.* 1996;276:39-43.
 85. O'Malley MS, Fletcher SW; US Preventive Services Task Force. Screening for breast cancer with breast self-examination: a critical review. *JAMA.* 1987;257:2196-2203.
 86. Frank E, Rimer BK, Brogan D, Elon L. US women physicians' personal and clinical breast cancer screening practices. *J Womens Health Gend Based Med.* 2000;9:791-801.
 87. Hall DC, Adams CK, Stein GH, Stephenson HS, Goldstein MK, Pennypacker HS. Improved detection of human breast lesions following experimental training. *Cancer.* 1980;46:408-414.
 88. Thomas DB, Gao DL, Self SG, et al. Randomized trial of breast self-examination in Shanghai: methodology and preliminary results. *J Natl Cancer Inst.* 1997;89:355-365.
 89. Thomas DB, Gao DL, Ray RM, et al. Randomized trial of breast self-examination in Shanghai: final results. *J Natl Cancer Inst.* 2002;94:1445-1457.
 90. Hackshaw AK, Paul EA. Breast self-examination and death from breast cancer: a meta-analysis. *Br J Cancer.* 2003;88:1047-1053.
 91. Kuhl CK, Schmutzler RK, Leutner CC, et al. Breast MR imaging screening in 192 women proved or suspected to be carriers of a breast cancer susceptibility gene: preliminary results. *Radiology.* 2000;215:267-279.
 92. Tilanus-Linthorst MM, Obdeijn IM, Bartels KC, de Koning HJ, Oudkerk M. First experiences in screening women at high risk for breast cancer with MR imaging. *Breast Cancer Res Treat.* 2000;63:53-60.
 93. Warner E, Plewes DB, Shumak RS, et al. Comparison of breast magnetic resonance imaging, mammography, and ultrasound for surveillance of women at high risk for hereditary breast cancer. *J Clin Oncol.* 2001;19:3524-3531.
 94. Morris EA, Liberman L, Ballon DJ, et al. MRI of occult breast carcinoma in a high-risk population. *AJR Am J Roentgenol.* 2003;181:619-626.
 95. Podo F, Sardanelli F, Canese R, et al. The Italian multi-centre project on evaluation of MRI and other imaging modalities in early detection of breast cancer in subjects at high genetic risk. *J Exp Clin Cancer Res.* 2002;21(3 suppl):115-124.
 96. Kriege M, Brekelmans CT, Boetes C, et al. Efficacy of MRI and mammography for breast-cancer screening in women with a familial or genetic predisposition. *N Engl J Med.* 2004;351:427-437.
 97. Stoutjesdijk MJ, Boetes C, Jager GJ, et al. Magnetic resonance imaging and mammography in women with a hereditary risk of breast cancer. *J Natl Cancer Inst.* 2001;93:1095-1102.
 98. Warner E, Plewes DB, Hill KA, et al. Surveillance of BRCA1 and BRCA2 mutation carriers with magnetic resonance imaging, ultrasound, mammography, and clinical breast examination. *JAMA.* 2004;292:1317-1325.
 99. Leach MO, Eeles RA, Turnbull LW, et al. The UK national study of magnetic resonance imaging as a method of screening for breast cancer (MARIBS). *J Exp Clin Cancer Res.* 2002;21(3 suppl):107-114.
 100. Gordon PB. Ultrasound for breast cancer screening and staging. *Radiol Clin N Am.* 2002;40:431-441.
 101. Stavros AT, Thickman D, Rapp CL, Dennis MA, Parker SH, Sisney GA. Solid breast nodules: use of sonography to distinguish between benign and malignant lesions. *Radiology.* 1995;196:123-134.
 102. Irwig L, Houssami N, van Vliet C. New technologies in screening for breast cancer: a systematic review of their accuracy. *Br J Cancer.* 2004;90:2118-2122.

103. Gordon PB, Goldenberg SL. Malignant breast masses detected only by ultrasound: a retrospective review. *Cancer*. 1995;76:626-630.
104. Kolb TM, Lichy J, Newhouse JH. Occult cancer in women with dense breasts: detection with screening US—diagnostic yield and tumor characteristics. *Radiology*. 1998;207:191-199.
105. Buchberger W, DeKoekkoek-Doll P, Springer P, Obrist P, Dunser M. Incidental findings on sonography of the breast: clinical significance and diagnostic workup. *AJR Am J Roentgenol*. 1999;173:921-927.
106. Kaplan SS. The utility of bilateral whole breast ultrasound in the evaluation of women with dense breast tissue [abstract]. *Radiology*. 2000;217:318.
107. Hou MF, Chuang HY, Ou-Yang F, et al. Comparison of breast mammography, sonography and physical examination for screening women at high risk of breast cancer in Taiwan. *Ultrasound Med Biol*. 2002;28:415-420.
108. O'Driscoll D, Warren R, MacKay J, Britton P, Day NE. Screening with breast ultrasound in a population at moderate risk due to family history. *J Med Screen*. 2001;8:106-109.
109. National Guidelines Clearinghouse Web site. Breast cancer screening. Available at: <http://www.guidelines.gov>. Accessibility verified February 2, 2005.
110. Freedman GM, Anderson PR, Goldstein LJ, et al. Routine mammography is associated with earlier stage disease and greater eligibility for breast conservation in breast carcinoma patients age 40 years and older. *Cancer*. 2003;98:918-925.
111. Baines C, To T, Wall C. Women's attitudes to screening after participation in the National Breast Screening Study: a questionnaire survey. *Cancer*. 1990;65:1663-1669.
112. Bartolucci G, Savron G, Fava GA, Grandi S, Trombini G, Orlandi C. Psychological reactions to thermography and mammography. *Stress Med*. 1989;5:195-199.
113. Rutter DR, Calnan M, Vaile MS, Field S, Wade KA. Discomfort and pain during mammography: description, prediction, and prevention. *BMJ*. 1992;305:443-445.
114. Kornguth PJ, Keefe FJ, Conaway MR. Pain during mammography: characteristics and relationship to demographic and medical variables. *Pain*. 1996;66:187-194.
115. Dullum JR, Lewis EC, Mayer JA. Rates and correlates of discomfort associated with mammography. *Radiology*. 2000;214:547-552.
116. Gram IT, Slenker SE. Cancer anxiety and attitudes toward mammography among screening attenders, nonattenders, and women never invited. *Am J Public Health*. 1992;82:249-251.
117. Lerman C, Trock B, Rimer BK, Boyce A, Jepson C, Engstrom PF. Psychological and behavioral implications of abnormal mammograms. *Ann Intern Med*. 1991;114:657-661.
118. Elmore JG, Barton MB, Mocerri VM, Polk S, Arena PJ, Fletcher SW. Ten-year risk of false-positive screening mammograms and clinical breast examinations. *N Engl J Med*. 1998;338:1089-1096.
119. Ernster VL, Barclay J. Increases in ductal carcinoma in situ (DCIS) of the breast in relation to mammography: a dilemma. *J Natl Cancer Inst Monogr*. 1997;151-156.
120. Ernster VL, Barclay J, Kerlikowske J, Grady D, Henderson IC. Incidence of and treatment for ductal carcinoma in situ of the breast. *JAMA*. 1996;275:913-918.
121. Feig SA, Hendrick RE. Radiation risk from screening mammography of women aged 40-49 years. *J Natl Cancer Inst Monogr*. 1997:119-124.
122. Baines CJ. Mammography screening: are women really giving informed consent? *J Natl Cancer Inst*. 2003;95:1508-1511.
123. Baines CJ. Mammography screening: are women really giving informed consent? (countering the counterpoint). *J Natl Cancer Inst*. 2003;95:1512-1513.
124. Berg AO. Mammography screening: are women really giving informed consent? (counterpoint). *J Natl Cancer Inst*. 2003;95:1511-1512.
125. Hansen NM, Ye X, Grube BJ, Giuliano AE. Manipulation of the primary breast tumor and the incidence of sentinel node metastases from invasive breast cancer. *Arch Surg*. 2004;139:634-639.
126. Boyd NF, Wolfson C, Moskowitz M, et al. Observer variation in the interpretation of xeromammograms. *J Natl Cancer Inst*. 1982;68:357-363.
127. Thornton H. Patients' understanding of risk: enabling understanding must not lead to manipulation. *BMJ*. 2003;327:693-694.
128. Schwartz LM, Woloshin S, Black WC, Welch HG. The role of numeracy in understanding the benefit of screening mammography. *Ann Intern Med*. 1997;127:966-972.
129. Braddock CH III, Edwards KA, Hasenberg NM, Laidley TL, Levinson W. Informed decision making in outpatient practice: time to get back to basics. *JAMA*. 1999;282:2313-2320.
130. Edwards A, Elwyn G, Covey J, Matthews E, Pill R. Presenting risk information—a review of the effects of “framing” and other manipulations on patient outcomes. *J Health Commun*. 2001;6:61-82.
131. Paling J. Strategies to help patients understand risks. *BMJ*. 2003;327:745-748.
132. Gigerenzer G, Edwards A. Simple tools for understanding risks: from innumeracy to insight. *BMJ*. 2003;327:741-744.
133. Hoffrage U, Lindsey S, Hertwig R, Gigerenzer G. Communicating statistical information. *Science*. 2000;290:2261-2262.
134. Edwards A, Elwyn G, Mulley A. Explaining risks: turning numerical data into meaningful pictures. *BMJ*. 2002;324:827-830.
135. Alaszewski A, Horlick-Jones T. How can doctors communicate information about risk more effectively? *BMJ*. 2003;327:728-731.
136. National Physician Fee Schedule Payment Amount File [Centers for Medicare & Medicaid Web site]. Available at: <http://www.cms.hhs.gov/providers/pufdownload/carrpuf.asp>. Accessed September 2, 2004. Accessibility verified February 2, 2005.