

Chemoprevention of Breast Cancer

Recommendations and Rationale

U.S. Preventive Services Task Force

This statement summarizes the current U.S. Preventive Services Task Force (USPSTF) recommendations for the chemoprevention of breast cancer and the supporting scientific evidence. Explanations of the ratings and of the strength of overall evidence are given in Appendix A and Appendix B, respectively. The complete information on which this statement is based, including evidence tables and references, is available in the article Chemoprevention of Breast Cancer: A Summary of the Evidence for the U.S. Preventive Services Task Force¹ (which follows this recommendation) and in the Systematic Evidence Review² on this topic. These documents can be obtained through the USPSTF Web site (www.preventiveservices.ahrq.gov) or the National Guideline Clearinghouse (www.guideline.gov). The summary of the evidence and the recommendation statement are also available in print through the Agency for Healthcare Research and Quality Publications Clearinghouse (call 1-800-358-9295 or e-mail ahrqpubs@ahrq.gov).

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Summary of Recommendations

- The U.S. Preventive Services Task Force (USPSTF) recommends against the routine use of tamoxifen or raloxifene for the primary prevention of breast cancer in women at low or average risk for breast cancer. (See “Clinical Considerations” for a discussion of risk.)
D recommendation

The USPSTF found fair evidence that tamoxifen and raloxifene may prevent some breast cancers in women at low or average risk for breast cancer, based on extrapolation from studies of women at higher risk. The USPSTF concluded, however, that the potential harms of chemoprevention may outweigh the potential benefits in women who are not at high risk for breast cancer.

- The USPSTF recommends that clinicians discuss chemoprevention with women at high risk for breast cancer and at low risk for adverse effects of chemoprevention. (See “Clinical Considerations” for a discussion of risk.) Clinicians should inform patients of the potential benefits and harms of chemoprevention. **B recommendation**

The USPSTF found fair evidence that treatment with tamoxifen can significantly reduce the risk for invasive estrogen-receptor-positive breast cancer in women at high risk for breast cancer and that the likelihood of benefit increases as the risk for breast cancer increases. The USPSTF found consistent but less abundant evidence for the benefit of raloxifene. The USPSTF found good evidence that tamoxifen and raloxifene increase the risk for thromboembolic events (for example, stroke, pulmonary embolism, and deep venous thrombosis) and symptomatic side effects (for example, hot flashes) and that tamoxifen, but not raloxifene, increases the risk for endometrial cancer. The USPSTF concluded that the balance of benefits and harms may be favorable for some high-risk women but will depend on breast cancer risk, risk for potential harms, and individual patient preferences.

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Clinical Considerations

- Clinicians should consider both the risk for breast cancer and the risk for adverse effects when identifying women who may be candidates for chemoprevention.

Risk for breast cancer: Older age; a family history of breast cancer in a mother, sister, or daughter; and a history of atypical hyperplasia on a breast biopsy are the strongest risk factors for breast cancer. Table 1 (p. 30) indicates how the estimated benefits of tamoxifen vary depending on age and family history. Other factors that contribute to risk include race, early age at menarche, pregnancy history (nulliparity or older age at first birth), and number of breast biopsies. The risk for developing breast cancer within the next 5 years can be estimated using risk factor information by completing the National Cancer Institute Breast Cancer Risk Tool (the “Gail model,” available at <http://cancer.gov/bcrisktool/> or 800-4-CANCER). Clinicians can use this information to help individual patients considering tamoxifen therapy estimate the potential benefit. However, the validity, feasibility, and impact of using the Gail model to identify appropriate candidates for chemoprevention has not been tested in a primary care setting. The Gail model does not incorporate estradiol levels or estrogen use, factors that some studies suggest may influence the effectiveness of tamoxifen.

Risk for adverse effects. Women are at lower risk for adverse effects from chemoprevention if they are younger; have no predisposition to thromboembolic events such as stroke, pulmonary embolism, or deep venous thrombosis; or do not have a uterus.

- In general, the balance of benefits and harms of chemoprevention is more favorable for (1) women in their 40s who are at increased risk for breast cancer and have no predisposition to thromboembolic events and (2) women in their

50s who are at increased risk for breast cancer, have no predisposition to thromboembolic events, and do not have a uterus. For example, a woman who is 45 years of age and has a mother, sister, or daughter with breast cancer would have approximately a 1.6% risk for developing breast cancer over the next 5 years (Table 1). On average, treating such women with tamoxifen for 5 years would prevent about 3 times as many invasive cancers (8 per 1,000) as the number of serious thromboembolic complications caused (1 stroke and 1 to 2 pulmonary emboli per 1,000). Among women 55 years of age, benefits exceed harms only for those who are not at risk for endometrial cancer; and the margin of benefit is small unless risk for breast cancer is substantially increased (for example, 4% over 5 years).

- Women younger than 40 years of age have a lower risk for breast cancer, and thus will not experience as large an absolute benefit from breast cancer chemoprevention as older women. Women 60 years of age and older, who have the highest risk for breast cancer, also have the highest risk for complications from chemoprevention with a less favorable balance of benefits and harms.
- The USPSTF found more evidence for the benefits of tamoxifen than for the benefits of raloxifene. Currently, only tamoxifen is approved by the U.S. Food and Drug Administration (FDA) for the specific indication of breast cancer chemoprevention. Although there are biological reasons to suspect that raloxifene should have similar benefits, trial data currently are limited to 1 study in which the primary outcome was fracture prevention. Additional trials to further evaluate this drug’s efficacy for breast cancer chemoprevention are under way, including a trial comparing efficacy and safety of raloxifene and tamoxifen. Raloxifene is approved by the FDA for preventing and treating osteoporosis.

Scientific Evidence

Epidemiology and Clinical Consequences

Breast cancer is the most common non-skin cancer in women. An estimated 203,500 new cases of invasive breast cancer will be diagnosed in 2002, and 39,600 women will die from the disease.³ Although the USPSTF concluded that early detection of breast cancer through mammography has reduced deaths from breast cancer, the effectiveness of mammography is limited. Another approach to reducing breast cancer deaths is chemoprevention for primary prevention of cancer.

Potential Benefits of Chemoprevention

The use of agents to prevent the development of breast cancer was suggested by trials of breast cancer treatment with tamoxifen, a compound with both estrogen-like and anti-estrogen properties (a selective estrogen receptor modulator).⁴ A meta-analysis of 55 studies evaluating tamoxifen for the treatment of women with breast cancer found that the drug was associated with an approximately 50% reduction in the risk for developing new cancers in the opposite breast among women who took the drug for 5 years.⁵

The USPSTF found and evaluated 4 randomized controlled trials (RCTs) of breast cancer chemoprevention in women who had never had breast cancer.⁴ Three of these trials used tamoxifen as the chemopreventive agent⁶⁻⁸; 1 trial used raloxifene, another selective estrogen receptor modulator.⁹

Of the 3 RCTs of tamoxifen, the largest (the Breast Cancer Prevention Trial — BCPT), with 13,388 women enrolled, found a risk reduction of invasive cancer of 49% among women at high risk for breast cancer (estimated 5-year risk of 1.66% or greater).⁷ Over the course of the BCPT, a total of 264 women were diagnosed with invasive breast cancer: 175 in the placebo group and 89 in the tamoxifen group (RR, 0.51; 95% CI, 0.39-0.66).

The absolute risk reduction was 21.4 cases per 1,000 women over 5 years.

The 2 other tamoxifen RCTs did not show a similar benefit. The relative risk reduction for breast cancer was 0.94 (95% CI, 0.59 -1.43) for the Royal Marsden Hospital study⁶ and 0.87 (95% CI, 0.62-2.14) for the Italian Tamoxifen Prevention Study.⁸ Although the reasons for these discrepant results are not definitively established, possible explanations include differences in the duration of therapy and differences between women enrolled in each study.¹ The average duration of therapy was shorter in the European trials and, compared with the women enrolled in BCPT, the women in these trials were younger, had more estrogen-receptor-negative cancers, and were more likely to be taking hormone replacement therapy or to have had an oophorectomy.¹

The study evaluating raloxifene in postmenopausal women with osteoporosis found a 76% risk reduction (RR, 0.24; 95% CI, 0.13-0.44) in the development of invasive breast cancer.⁹ After a median follow-up of 40 months, the absolute risk reduction among women taking raloxifene was 7.9 cases per 1,000 women (number needed to treat, 126).⁹ When effective, both raloxifene and tamoxifen were effective only against estrogen receptor-positive tumors.¹

Potential Harms of Chemoprevention

Both tamoxifen and raloxifene increase the risk for thromboembolic events and hot flashes; tamoxifen increases the risk for endometrial cancer.¹ The number of total thromboembolic events in all 4 trials was small, and differences in specific complication rates between the treatment and placebo arms were statistically significant only for pulmonary embolism.¹ Among women aged 50 and older, for whom the potential harms of tamoxifen and raloxifene are more common than they are for younger women, the BCPT reported that after a median of 55 months of use, tamoxifen increased the rate of stroke from 1.3 cases/1,000 women in

the placebo group to 2.2 cases/1,000 women in the study group (RR, 1.75; 95% CI, 0.98-3.20); increased the rate of pulmonary embolism from 0.3 cases/1,000 women in the placebo group to 1.0 cases/1,000 women in the study group (RR, 3.19%; 95% CI, 1.12-11.15); increased the rate of deep vein thrombosis from 0.9 cases/1,000 women in the placebo group to 1.5 cases/1,000 women in the study group (RR, 1.71; 95% CI, 0.85-3.58).⁷

Fewer thromboembolic events occurred among women younger than 50, and the trial found no significant difference in incidence between the tamoxifen and placebo groups in this age group.⁷ The relative risk increase in venous thromboembolism from tamoxifen or raloxifene appears similar to the risk for venous thromboembolism from oral contraceptives or hormone replacement therapy.¹

Among women aged 50 and older in the BCPT, participants who received tamoxifen, compared with those who took placebo, had a 4.0 times greater risk (95% CI, 1.70-10.90) of developing Stage 1 endometrial cancer (0.8 cancers/1,000 women taking placebo vs 3.1 cancers/1,000 women taking tamoxifen for a median of 55 months).⁷ Among women younger than 50, the BCPT found no significant difference in endometrial cancer rates between the 2 groups. No deaths attributed to endometrial cancer occurred in the trial.⁷ Raloxifene has not been associated with an increase in endometrial cancer.⁹

The BCPT reported that women in the tamoxifen group were at increased risk for developing cataracts and having cataract surgery compared with placebo (RR, 1.14 [95% CI, 1.01-1.29] and 1.57 [95% CI, 1.16-2.14], respectively).⁷

Quality of life issues have also been of concern and were addressed in the BCPT. Women in the BCPT reported increased rates of bothersome hot flashes (45.7% in the tamoxifen group vs 28.7% in the placebo group) and bothersome vaginal discharge (12.4% in the tamoxifen group vs 4.5% in the placebo group).⁷ Women given raloxifene also noted higher rates of hot flashes than women given placebo (10.7% in the raloxifene group vs 6.4% in the placebo group).⁹

Although long-term adherence for highly motivated women was about 80% in the BCPT trial and about 90% in the raloxifene trial, adherence rates in the general population are unknown.²

Recommendations of Others

The American College of Obstetricians and Gynecologists emphasizes the importance of clinician judgment and recommends that any decision to use tamoxifen be made on an individual basis after consideration of the patient's medical history, risk assessment, and preferences, and with attention to the ability to manage complications of therapy.¹⁰ The American Society of Clinical Oncology suggests that women with a 5-year projected risk for breast cancer greater than or equal to 1.66% may be offered tamoxifen to reduce their risk. They also recommend that raloxifene use should be reserved for treatment of osteoporosis in postmenopausal women.¹¹ The Canadian Task Force on Preventive Health Care recommends that clinicians counsel women at high risk for breast cancer (Gail index \geq 1.66% for 5 years) about the potential benefits and harms of breast cancer prevention with tamoxifen.¹²

References

1. Kinsinger LA, Harris R, Lewis C, Woddell M. Chemoprevention of breast cancer: a summary of the evidence for the U.S. Preventive Services Task Force. *Ann Intern Med.* 2002;137:56-67.
2. Kinsinger LA, Harris R, Lewis C, Woddell M. *Chemoprevention of Breast Cancer.* Systematic Evidence Review No. 8 (Prepared by the Research Triangle Institute – University of North Carolina Evidence-based Practice Center under Contract No. 290-97-0011). Rockville, MD: Agency for Healthcare Research and Quality. July 2002. (Available on the AHRQ Web site at: www.ahrq.gov/clinic/serfiles.htm).
3. Jemal A, Thomas A, Murray T, Thun M. Cancer statistics, 2002. *CA Cancer J Clin.* 2002;52:23-47.
4. Early Breast Cancer Trialists' Collaborative Group. Systemic treatment of early breast cancer by hormonal, cytotoxic, or immune therapy. 133 randomised trials involving 31,000 recurrences and 24,000 deaths among 75,000 women. *Lancet.* 1992;339:1-15.

5. Early Breast Cancer Trialists' Collaborative Group. Tamoxifen for early breast cancer: an overview of the randomised trials. *Lancet*. 1998;351:1451-1467.
6. Powles T, Eeles R, Ashley S, et al. Interim analysis of the incidence of breast cancer in the Royal Marsden Hospital tamoxifen randomised chemoprevention trial. *Lancet*. 1998;352:98-101.
7. Fisher B, Costantino JP, Wickerham DL, et al. Tamoxifen for prevention of breast cancer: report of the National Surgical Adjuvant Breast and Bowel Project P-1 Study. *J Natl Cancer Inst*. 1998;90:1371-1388.
8. Veronesi U, Maisonneuve P, Costa A, et al. Prevention of breast cancer with tamoxifen: preliminary findings from the Italian randomised trial among hysterectomised women. Italian Tamoxifen Prevention Study. *Lancet*. 1998;352:93-97.
9. Cummings SR, Eckert S, Krueger KA, et al. The effect of raloxifene on risk of breast cancer in postmenopausal women: results from the MORE randomized trial. Multiple Outcomes of Raloxifene Evaluation. *JAMA*. 1999;281:2189-2197.
10. American College of Obstetricians and Gynecologists. *Tamoxifen and the Prevention of Breast Cancer in High-risk Women*. ACOG Committee Opinion 224. Washington, DC: ACOG; 1999.
11. Chlebowski RT, Collyar DE, Somerfield MR, Pfister DG. American Society of Clinical Oncology Technology Assessment on breast cancer risk reduction strategies: tamoxifen and raloxifene. *J Clin Oncol*. 1999;17:1939-1954.
12. Levine M, Moutquin JM, Walton R, Feightner J. Chemoprevention of breast cancer. A joint guideline from the Canadian Task Force on Preventive Health Care and the Canadian Breast Cancer Initiative's Steering Committee on Clinical Practice Guidelines for the Care and Treatment of Breast Cancer. *CMAJ*. 2001;164:1681-1690.
13. Gail MH, Costantino JH, Bryant J, et al. Weighing the risks and benefits of tamoxifen treatment for preventing breast cancer. *J Natl Cancer Inst*. 1999;91:1829-1846.

Table 1. Predicted benefits and harms of 5 years of tamoxifen therapy according to age and family history*

Variable†	Women 45 years of age	Women 55 years of age	Women 65 years of age	Women 75 years of age
Predicted 5-year risk of breast cancer, %‡				
No family history	0.7	1.1	1.5	1.6
Family history	1.6	2.3	3.2	3.4
Benefits per 1000 women over 5 y of tamoxifen therapy				
Cases of invasive breast cancer avoided, <i>n</i>				
No family history	3-4	5-6	7-8	8
Family history	8	11-12	16	17
Cases of noninvasive breast cancer avoided, <i>n</i>				
No family history	1-2	2	2-3	2-3
Family history	2-3	3-4	4-5	5-6
Hip fractures avoided, <i>n</i> §	<1	3	5	15
Harms per 1000 women over 5 y of tamoxifen therapy				
Cases of endometrial cancer caused, <i>n</i> §	1-2	12	21	22
Strokes caused, <i>n</i> §	1	3	9	20
Pulmonary emboli caused, <i>n</i> §	1-2	4-5	9	18
Cases of deep venous thrombosis caused, <i>n</i> §	1-2	1-2	3	4

*These estimates are based on the Gail model, outcomes from the Breast Cancer Prevention Trial, and baseline rates of harms from Gail et al.¹³

†No family history = no first-degree relatives with breast cancer; family history = 1 first-degree relative with breast cancer.

‡Based on menarche at 12 years of age, first birth at 22 years of age, and no history of breast biopsy, as calculated from the Gail model.

§Modified from Gail et al.¹³

Appendix A U.S. Preventive Services Task Force - Recommendations and Ratings

The Task Force grades its recommendations according to one of 5 classifications (A, B, C, D, I) reflecting the strength of evidence and magnitude of net benefit (benefits minus harms):

- A. The USPSTF strongly recommends that clinicians routinely provide [the service] to eligible patients. *The USPSTF found good evidence that [the service] improves important health outcomes and concludes that benefits substantially outweigh harms.*
- B. The USPSTF recommends that clinicians routinely provide [the service] to eligible patients. *The USPSTF found at least fair evidence that [the service] improves important health outcomes and concludes that benefits outweigh harms.*
- C. The USPSTF makes no recommendation for or against routine provision of [the service]. *The USPSTF found at least fair evidence that [the service] can improve health outcomes but concludes that the balance of benefits and harms is too close to justify a general recommendation.*
- D. The USPSTF recommends against routinely providing [the service] to asymptomatic patients. *The USPSTF found at least fair evidence that [the service] is ineffective or that harms outweigh benefits.*
- I. The USPSTF concludes that the evidence is insufficient to recommend for or against routinely providing [the service]. *Evidence that [the service] is effective is lacking, of poor quality, or conflicting and the balance of benefits and harms cannot be determined.*

Appendix B U.S. Preventive Services Task Force - Strength of Overall Evidence

The USPSTF grades the quality of the overall evidence for a service on a 3-point scale (good, fair, poor):

- Good:** Evidence includes consistent results from well-designed, well-conducted studies in representative populations that directly assess effects on health outcomes.
- Fair:** Evidence is sufficient to determine effects on health outcomes, but the strength of the evidence is limited by the number, quality, or consistency of the individual studies, generalizability to routine practice, or indirect nature of the evidence on health outcomes.
- Poor:** Evidence is insufficient to assess the effects on health outcomes because of limited number or power of studies, important flaws in their design or conduct, gaps in the chain of evidence, or lack of information on important health outcomes.

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*These current members were not on the Task Force at the time this recommendation was voted.