Hormone Therapy for the Prevention of Chronic Conditions in Postmenopausal Women: Recommendations from the U.S. Preventive Services Task Force

U.S. Preventive Services Task Force*

This statement summarizes the U.S. Preventive Services Task Force recommendations on hormone therapy for the prevention of chronic conditions in postmenopausal women and the supporting scientific evidence, and updates the Task Force’s 2002 recommendations on hormone replacement therapy. The updated statement is based on the results of the Women’s Health Initiative randomized, controlled trial, as well as the information in the 2002 summary of the evidence on this topic, which is available on the USPSTF Web site (www.preventiveservices.ahrq.gov).


For author affiliations, see end of text.

SUMMARY OF THE RECOMMENDATIONS

The U.S. Preventive Services Task Force (USPSTF) recommends against the routine use of combined estrogen and progestin for the prevention of chronic conditions in postmenopausal women. This is a grade D recommendation. (See Appendix Table 1 for a description of the USPSTF classification of recommendations.)

The USPSTF found good evidence that the use of combined estrogen and progestin results in both benefits and harms. Benefits include reduced risk for fracture (good evidence) and colorectal cancer (fair evidence). Combined estrogen and progestin has no beneficial effect on coronary heart disease (CHD) and may even pose an increased risk (good evidence). Other harms include increased risk for breast cancer (good evidence), venous thromboembolism (good evidence), stroke (fair evidence), cholecystitis (fair evidence), dementia (fair evidence), and lower global cognitive function (fair evidence). Because of insufficient evidence, the USPSTF could not assess the effects of combined estrogen and progestin on the incidence of ovarian cancer, mortality from breast cancer or CHD, or all-cause mortality. The USPSTF concluded that the harmful effects of unopposed estrogen are likely to exceed the chronic disease prevention benefits in most women.

The U.S. Preventive Services Task Force (USPSTF) recommends against the routine use of unopposed estrogen for the prevention of chronic conditions in postmenopausal women who have had a hysterectomy. This is a grade D recommendation.

The USPSTF found good evidence that the use of unopposed estrogen results in both benefits and harms. The benefits include reduced risk for fracture (good evidence). Harms include increased risk for venous thromboembolism (fair evidence), stroke (fair evidence), dementia (fair evidence), and lower global cognitive function (fair evidence). There is fair evidence that unopposed estrogen has no beneficial effect on CHD. Because of insufficient evidence, the USPSTF could not assess the effects of unopposed estrogen on the incidence of breast cancer, ovarian cancer, or colorectal cancer, as well as breast cancer mortality or all-cause mortality. The USPSTF concluded that the harmful effects of unopposed estrogen are likely to exceed the chronic disease prevention benefits in most women.

CLINICAL CONSIDERATIONS

The balance of benefits and harms for a woman will be influenced by her personal preferences, her risks for specific chronic diseases, and the presence of menopausal symptoms. A shared decision-making approach to preventing chronic diseases in perimenopausal and postmenopausal women involves consideration of individual risk factors and preferences in selecting effective interventions for reducing the risks for fracture, heart disease, and cancer.

Other USPSTF recommendations for prevention of chronic diseases (screening for osteoporosis, high blood pressure, lipid disorders, breast cancer, and colorectal cancer, and counseling to prevent tobacco use) are available at www.preventiveservices.ahrq.gov.

The USPSTF did not consider the use of hormone therapy for the management of menopausal symptoms, which is the subject of recommendations by other expert groups. Women and their clinicians should discuss the balance of risks and benefits before deciding to initiate or continue hormone therapy for menopausal symptoms. For example, for combined estrogen and progestin, some risks (such as the risks for venous thromboembolism, CHD, and stroke) arise within the first 1 to 2 years of therapy, and other risks (such as the risk for breast cancer) appear to...
increase with longer-term hormone therapy. The populations of women using hormone therapy for symptom relief may differ from those who would use hormone therapy for prevention of chronic disease (for example, age differences). Other expert groups have recommended that women who decide to take hormone therapy to relieve menopausal symptoms use the lowest effective dose for the shortest possible time.

Although estrogen alone or in combination with progestin reduces the risk for fractures in women, other effective medications (for example, bisphosphonates and calcitonin) are available for treating women with low bone density to prevent fractures. The role of chemopreventive agents in preventing fractures in women without low bone density is unclear. The USPSTF addressed screening for osteoporosis in postmenopausal women in 2002 (1).

Unopposed estrogen increases the risk for endometrial cancer in women who have an intact uterus. Clinicians should use a shared decision-making approach when discussing the possibility of using unopposed estrogen in women who have not had a hysterectomy (2).

**DISCUSSION**

The median age of menopause in women in the United States is 51 years (range, 41 to 59 years), but ovarian production of estrogen and progestin begins to decrease years before the cessation of menses. The average woman in the United States who reaches menopause has a life expectancy of nearly 30 more years. The probability that a menopausal woman will develop various chronic diseases during her lifetime has been estimated to be 46% for CHD, 20% for stroke, 15% for hip fracture, 10% for breast cancer, and 2.6% for endometrial cancer (3). In North America, an estimated 7% to 8% of people 75 to 84 years of age have dementia, and more than 90% of cases of colorectal cancer occur after the age of 50 years (4).

**Benefits of Hormone Therapy**

**Osteoporosis and Fractures**

Good evidence from observational studies and randomized clinical trials demonstrates that estrogen therapy increases bone density and reduces the risk for fractures. The combined estrogen–progestin arm of the Women’s Health Initiative (WHI) trial (5), a fair-quality study, found significant reductions in total fracture risk (hazard ratio [HR], 0.76 [adjusted 95% CI, 0.63 to 0.92]) among healthy women taking estrogen and progestin. This arm of the WHI trial also showed reductions for hip and vertebral fracture, although these did not achieve statistical significance (5). (In its analysis, the USPSTF used nominal 95% CIs for the primary outcomes and adjusted 95% CIs for all secondary outcomes.)

The estrogen-only arm of the WHI trial also reported decreased risk for hip and vertebral fracture, which also did not reach statistical significance (6). A meta-analysis of 22 trials of estrogen reported an overall 27% reduction in nonvertebral fractures (relative risk [RR], 0.73 [CI, 0.56 to 0.94]), although the quality of individual studies varied (7). The Heart and Estrogen/progestin Replacement Study (HERS) and its unblinded follow-up study, HERS II (8), a fair-quality trial of combined estrogen–progestin for the secondary prevention of heart disease that reported many other outcomes, found no reduction in hip, wrist, vertebral, or total fractures with hormone therapy (relative hazard for total fractures, 1.04 [CI, 0.87 to 1.25]). Overall, a good-quality body of evidence supports the efficacy of hormone therapy in increasing bone density and decreasing fracture risk.

**Colorectal Cancer**

Results from the WHI study (5) and HERS (8) showed a trend toward reduced incidence of colon cancer (HR, 0.63 [adjusted CI, 0.32 to 1.24], and relative hazard, 0.81 [CI, 0.46 to 1.45], respectively), but the trend did not reach statistical significance. The estrogen-only arm of the WHI trial showed neither benefit nor harm for colorectal cancer risk (HR, 1.08 [adjusted CI, 0.63 to 1.86]) (6). A meta-analysis of 18 observational studies of postmenopausal women reported a 20% reduction in colon cancer (RR, 0.80 [CI, 0.74 to 0.86]) and a 19% reduction in rectal cancer (RR, 0.81 [CI, 0.72 to 0.92]) among women who had ever used combined estrogen–progestin or estrogen alone compared with women who had never used hormone therapy (9). This decrease in risk was more apparent when current users were compared with those who had never used hormone therapy (RR, 0.66 [CI, 0.59 to 0.74]). Overall, the evidence suggesting a trend toward reduction of colorectal cancer risk with combined hormone therapy should be interpreted cautiously until controlled trials clarify whether therapy has either no benefit or modest benefit.

**Harms of Hormone Therapy**

**Breast Cancer**

The estrogen–progestin arm of the WHI study was terminated after an average of 5.2 years of follow-up because “evidence for breast cancer harm, along with evidence for some increase in CHD, stroke, and pulmonary embolism, outweighed the evidence of benefit for fractures and possible benefit for colon cancer” (5). This study showed an increased invasive breast cancer incidence (HR, 1.26 [nominal CI, 1.00 to 1.59]). However, no effect on breast cancer mortality was observed. Comparable increases in breast cancer incidence were observed among women taking estrogen and progestin over 6.8 years of follow-up in HERS (8). The U.K. Million Women Study, a fair-quality study, showed an increased risk for breast cancer in current users of combined estrogen–progestin (RR, 2.00 [CI, 1.91 to 2.09]) compared with those who had never used hormone therapy (10). Results from 2 good-quality cohort studies conflict on the effects of long-term hormone therapy on breast cancer mortality (11, 12). Overall, there is a good-quality body of evidence indicating that com-
Hormone Therapy for the Prevention of Chronic Conditions in Postmenopausal Women

Clinical Guidelines

Volume 142 • Number 10

857

Venous Thromboembolism (Deep Venous Thrombosis and Pulmonary Embolism)

In a meta-analysis of 12 studies (3 randomized, controlled trials; 8 case–control studies; and 1 cohort study), hormone therapy (estrogen alone or in combination with progestin) was associated with an increased risk for venous thromboembolism (RR, 2.14 [CI, 1.64 to 2.81]) (15, 16). Five of 6 studies that examined the effects of hormone therapy over time reported that the risk was highest within the first year of use (RR, 3.49 [CI, 2.33 to 5.59]). These results are consistent with the findings in the estrogen–progestin arm of the WHI (5), which reported a 2-fold increased rate of venous thromboembolic disease, including deep venous thrombosis and pulmonary embolism, in women taking combined estrogen–progestin daily. The estrogen-only arm of the WHI trial showed a trend toward increased risk for venous thromboembolism with unopposed estrogen use (HR, 1.33 [adjusted CI, 0.86 to 2.08]) (6).

Cognition and Dementia

While earlier studies showed a beneficial effect of hormone therapy on cognition, these studies had marked heterogeneity and variation in assessment of outcomes. For example, 9 randomized, controlled trials examining the effect of hormone therapy on cognition in women showed improvement in verbal memory, vigilance, reasoning, and motor speed; however, these trials may have biased results, since they were conducted with women experiencing menopausal symptoms at baseline. A meta-analysis of 12 observational studies (1 of good quality, 3 of fair quality, and 8 of poor quality) showed a reduction in the risk for dementia among postmenopausal women taking hormone therapy (RR, 0.66 [CI, 0.53 to 0.82]) (17). Because of issues of internal and external validity from these previous studies, the more recent, fair-quality WHI memory studies are more likely to represent the effects of hormone therapy use in the healthy postmenopausal population. The WHI memory study showed decreased global cognitive function (measured by the modified Mini-Mental Status Examination) in women taking estrogen alone and in the pooled group of women taking estrogen alone or estrogen–progestin (18). The WHI memory study also showed an increased risk for probable dementia or mild cognitive impairment in both the estrogen-alone (HR, 1.38 [CI, 1.01 to 1.89]) and estrogen–progestin (HR, 1.44 [CI, 1.04 to 1.99]) arms of the trial (19). The overall evidence supports harmful effects of hormone therapy on cognitive function, although the clinical relevance of this difference in cognitive function is unclear.

Endometrial and Ovarian Cancer

Results of a meta-analysis of 29 good-quality observational studies of endometrial cancer reported a relative risk of 2.3 for users of unopposed estrogen compared with non-users (20). Risks increased with increasing duration of use

CHD

In the WHI study, women who took combined estrogen–progestin daily, compared with women taking placebo, had an increased risk for CHD (fatal and nonfatal myocardial infarctions), which became evident shortly after initiation of the study (HR, 1.29 [nominal CI, 1.02 to 1.63]) (5). However, mortality from CHD was not significantly increased among the women taking combined hormone therapy daily. One meta-analysis of observational studies showed a statistically significant reduction in CHD (RR, 0.80 [CI, 0.68 to 0.95]) among current hormone therapy users, but not among those who had used hormone therapy in the past or among those who had never used it (13). This meta-analysis also showed that CHD mortality in observational studies was reduced among current hormone therapy users (RR, 0.62 [CI, 0.40 to 0.90]) but was not reduced among those who had used hormone therapy in the past. However, among studies that controlled for socioeconomic status (social class, education, or income), no CHD benefit was seen among current hormone therapy users, suggesting that the observed difference may be due to confounding by socioeconomic status and other lifestyle factors (for example, exercise or alcohol use) rather than use of hormone therapy. Thus, selection bias (in this case, the tendency of healthier women to use hormone therapy) appears to explain the apparent protective effect of estrogen against CHD seen in observational studies. The estrogen-only arm of the WHI trial showed no decreased risk for CHD (6).

Stroke

A meta-analysis of 9 observational primary prevention studies suggests that hormone therapy is associated with a small increase in stroke incidence (RR, 1.12 [CI, 1.01 to 1.23]), due primarily to an increase in thromboembolic stroke (RR, 1.20 [CI, 1.01 to 1.40]) (13, 14). The risk for subarachnoid bleeding and hemorrhagic stroke was not increased, and the overall stroke mortality was marginally reduced (RR, 0.81 [CI, 0.71 to 0.92]). These results are consistent with findings from the WHI, which reported increased incidence of stroke in women taking combined estrogen–progestin daily (HR, 1.41 [adjusted CI, 0.86 to 2.31]) (5). The estrogen-only arm of the WHI trial, which was terminated after an average of 6.8 years of follow-up, showed a trend toward increased stroke risk with unopposed estrogen use (HR, 1.39 [adjusted CI, 0.97 to 1.99]) (6).

bined estrogen–progestin increases breast cancer risk. It is unclear whether the combination of estrogen and progestin confers a greater breast cancer risk than estrogen alone. In studies of estrogen alone, the results are conflicting: The Million Women Study showed an increased risk for breast cancer in current users of estrogen only (RR, 1.30 [CI, 1.22 to 1.38]) compared with those who had never used it (10), but the estrogen-only arm of the WHI trial showed a trend toward breast cancer prevention (HR, 0.77 [nominal CI, 0.59 to 1.01]) (6).
(RR, 9.5 for 10 years of use), and the risk for endometrial
cancer remained elevated 5 or more years after discontin-
uation of unopposed estrogen therapy. Estrogen and pro-
gestin did not increase the risk for endometrial cancer in
HERS (4) or in the WHI (5).

Data on the association between the use of hormone
therapy and the risk for ovarian cancer is inconsistent.
Two good-quality cohort studies reported increased risks
for ovarian cancer or ovarian cancer mortality among
women who had taken hormone therapy for 10 years or
more (21, 22). However, a third study found no effect
of hormone therapy on ovarian cancer mortality (23). One
study suggested higher risk with unopposed estrogen than
with estrogen–progestin therapy (21), but data are insuffi-
cient to resolve the effects of different formulations or
doses of hormone therapy on ovarian cancer risk. Neither
the WHI nor HERS reported risk for ovarian cancer.

Cholecystitis

Results from the Nurses’ Health Study, a good-quality
cohort study, reported an increased risk for cholecystitis
among current hormone therapy users and long-term users
(>5 years) compared with nonusers (24). Risk for chole-
cystitis remained elevated among past users. An increase in
biliary tract surgery during 6.8 years of follow-up was re-
ported among women taking estrogen plus progestin com-
pared with those taking placebo in HERS (8, 25); the
WHI has not reported on outcomes for biliary tract disease
among women taking hormone therapy.

CONCLUSION

Combined estrogen–progestin therapy may reduce the
risk for fractures and colorectal cancer but has no beneficial
effect on CHD. The use of combined estrogen–progestin
may lead to increased risk for breast cancer, venous throm-
boembolism, stroke, cholecystitis, dementia, and lower
global cognitive function. The excess absolute combined
risks for CHD and breast cancer that can be attributed to
hormone therapy are low; for example, according to WHI
results, there would be 7 more CHD events, 8 more
strokes, 8 more pulmonary embolisms, and 8 more cases of
invasive breast cancer each year for every 10 000 women
taking hormone therapy. The absolute risk reduction for
every 10 000 women would be 6 fewer cases of colorectal
cancer and 5 fewer hip fractures. The evidence is insuffi-
cient to determine the effects of hormone therapy on the
incidence of ovarian cancer, mortality from breast cancer
or CHD, or all-cause mortality. Evidence about the effects
of different dosages, types, and delivery modes of hormone
therapy remains insufficient. Overall, the harmful effects of
combined estrogen and progestin are likely to exceed the
benefits of chronic disease prevention for most women.

Since unopposed estrogen increases a woman’s risk for
endometrial cancer, it has been used in postmenopausal
women without a uterus to prevent chronic disease. While
estrogen alone may decrease a woman’s risk for fractures, it
has no beneficial effect on CHD. The use of estrogen alone
may lead to increased risk for thromboembolism, stroke,
dementia, and lower global cognitive function. The evidence
is insufficient to determine the effects of unopposed
estrogen on the incidence of breast cancer, ovarian cancer,
or colorectal cancer, as well as breast cancer mortality or
all-cause mortality. Overall, the harmful effects of unop-
posed estrogen are likely to exceed the chronic disease pre-
vention benefits in most women.

RECOMMENDATIONS OF OTHER GROUPS

The American College of Obstetricians and Gynecolo-
gists (26), American Heart Association (27), North Amer-
ican Menopause Society (28), and Canadian Task Force on
Preventive Health Care (29, 30) recommend against use of
hormone therapy for the prevention of chronic diseases in
postmenopausal women.

APPENDIX

Members of the U.S. Preventive Services Task Force are
Ned Calonge, MD, MPH, Chair (Colorado Department of Pub-
lic Health and Environment, Denver, Colorado); Janet D. Allan,
PhD, RN, CS, Vice-Chair (University of Maryland, Baltimore,
Maryland); Alfred O. Berg, MD, MPH (University of Wash-
ington, Seattle, Washington); Paul S. Frame, MD (Tri-
County Family Medicine, Cohocton, and University of Roches-
ter, Rochester, New York); Leon Gordis, MD, MPH, DrPH
(Johns Hopkins Bloomberg School of Public Health, Baltimore,
Maryland); Kimberly D. Gregory, MD, MPH (Cedars-Sinai
Medical Center, Los Angeles, California); Russell Harris, MD,
Appendix Table 2. U.S. Preventive Services Task Force Grades for Strength of Overall Evidence*

<table>
<thead>
<tr>
<th>Grade</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>Good</td>
<td>Evidence includes consistent results from well-designed, well-conducted studies in representative populations that directly assess effects on health outcomes</td>
</tr>
<tr>
<td>Fair</td>
<td>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</td>
</tr>
<tr>
<td>Poor</td>
<td>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</td>
</tr>
</tbody>
</table>

* The U.S. Preventive Services Task Force (USPSTF) grades the quality of the overall evidence for a service on a three-point scale (good, fair, poor).

MPH (University of North Carolina School of Medicine, Chapel Hill, North Carolina); Mark S. Johnson, MD, MPH (University of Medicine and Dentistry of New Jersey–New Jersey Medical School, Newark, New Jersey); Jonathan D. Klein, MD, MPH (University of Rochester School of Medicine, Rochester, New York); Carol Loveland-Cherry, PhD, RN (University of Michigan School of Nursing, Ann Arbor, Michigan); Virginia A. Moyer, MD, MPH (University of Texas Health Science Center, Houston, Texas); Judith K. Ockene, PhD (University of Massachusetts Medical School, Worcester, Massachusetts); Diana B. Petitti, MD, MPH (Kaiser Permanente Southern California, Pasadena, California); Albert L. Siu, MD, MSPH (Mount Sinai Medical Center, New York, New York); Steven M. Teutsch, MD, MPH (Merck & Co., Inc., West Point, Pennsylvania); and Barbara P. Yawn, MD, MSc (Olmstead Research Center, Rochester, Minnesota).

This list includes members of the Task Force at the time this recommendation was finalized. For a list of current Task Force members, go to www.ahrq.gov/clinic/uspsfab.htm. Steven M. Teutsch, MD, MPH, recused himself from voting on this topic.

From the U.S. Preventive Services Task Force, Agency for Healthcare Research and Quality, Rockville, Maryland.

Disclaimer: Recommendations made by the USPSTF are independent of the U.S. government. They should not be construed as an official position of the Agency for Healthcare Research and Quality or the U.S. Department of Health and Human Services.

Requests for Single Reprints: Reprints are available from the USPSTF Web site (www.preventiveservices.ahrq.gov) and in print through the Agency for Healthcare Research and Quality Publications Clearinghouse (800-358-9295).

References


