Two complementary studies in separate components of the Women’s Health Initiative (WHI) examined relationships among weight loss, diet composition and breast cancer incidence and outcome in postmenopausal women. In the WHI Observational Study, 61,335 postmenopausal women had their weight change determined over a 3-year period with subsequent follow-up. Women with weight loss greater than or equal to 5% had significantly lower breast cancer incidence compared to women with stable weight. In the WHI Dietary Modification randomized clinical trial involving 48,835 postmenopausal women, implementation of a low-fat eating pattern significantly reduced deaths after breast cancer. Thus, moderation regarding dietary composition and body weight maintenance can reduce a postmenopausal woman’s risk of being diagnosed with breast cancer and of dying after breast cancer.

The study population included 61,335 postmenopausal women, entered between 1993 and 1998 at 40 US clinical centers, who were aged 50–79 years and had no prior breast cancer with a normal baseline mammogram. Anthropometrics were measured and BMI calculated at baseline and year 3. Weight change at year 3 was categorized as stable (<5% change), loss (≥5% decrease) or gain (≥5% increase). Participants were followed annually for clinical outcomes. Breast cancer incidence was determined through 11.4 years (mean) follow-up after the year 3 weight status assessment and all breast cancers were verified by central medical record review. Survival findings were augmented by serial National Death Index (NDI) queries, which capture 98% of all deaths [6].

Among 61,335 participants, 3,061 developed breast cancer. Women with weight loss (≥ 5%) had a significantly lower breast cancer incidence compared to women with stable weight (hazard ratio [HR] 0.88, 95% confidence interval [CI] 0.78–0.98, P = 0.02), and the findings did not differ by whether the weight loss was intentional or not. We examined associations between weight change category and breast cancer risk by stratifying variables: baseline BMI,
race/ethnicity, age, and menopausal hormone therapy, with none of the interactions found to be statistically significant. Weight gain was not associated with higher breast cancer risk overall, although a higher risk of triple-negative breast cancer was seen (HR 1.54, 95% CI 1.16–2.05).

In addition and not previously reported, we examined the change in waist circumference (WC) at year 3 from baseline with subsequent breast cancer risk. The study population was 60,552 after excluding those with missing WC, and 3,379 breast cancer cases were identified.

The association between change in WC and subsequent breast cancer risk was examined using the Cox proportional hazards regression model, and the proportional hazards assumption was checked and held. The HRs and 95% CIs for the association between WC change and risk of breast cancer are shown in Table 1:

| WC change category (percentage change) | Cases | Age-adjusted HR (95% CI) | Multivariable-adjusted† HR (95% CI) |
|---------------------------------------|-------|--------------------------|-------------------------------------|
| Stable WC (within ±5% change)         | 2004  | Reference                | Reference                           |
| WC gain (≥5%)                         | 875   | 0.98 (0.91 1.06)         | 1.00 (0.92 1.08)                    |
| WC loss (≥5%)                         | 500   | 0.98 (0.89 1.08)         | 0.94 (0.85 1.04)                    |
| Intentional                           | 346   | 0.96 (0.86 1.08)         | 0.92 (0.82 1.03)                    |
| Unintentional                         | 154   | 1.02 (0.87 1.20)         | 1.00 (0.85 1.18)                    |

* a. In multivariable models, we adjusted for Gail score (based on age, race/ethnicity, age at menarche, age of the mother at the birth of her first live child, number of first-degree relatives with breast cancer, and the number of previous breast biopsy examinations) [8], education, smoking pack-years, recreational physical activity, alcohol, history of hormone therapy use, parity, BMI, and waist circumference.

Table 2: Waist circumference (WC) change category by weight change category

| WC change | Frequency | Percent | Stable weight (±5%) | Weight gain (≥5%) | Weight loss (≥5%) - Unintentional | Weight loss (≥5%) - Intentional | Total |
|-----------|-----------|---------|---------------------|-------------------|-----------------------------------|----------------------------------|-------|
| WC gain (≥5%) | 26948     | 44.50   | 26948               | 4654              | 1516                              | 2193                             | 35311 |
| WC loss (≥5%) - Unintentional | 76.32     | 13.18   | 76.32               | 39.32             | 45.86                             | 45.94                            | 58.32 |
| WC loss (≥5%) - Intentional | 66.31     | 11.19   | 66.31               | 2.50              | 4.29                              | 6.21                             | 26.13 |
| Total     | 40637     | 67.11   | 40637               | 11835             | 3306                              | 4774                             | 60552 |

Table 2: Waist circumference (WC) change category by weight change category

| WC change | Percent | WC change | Frequency | Percent | Stable weight (±5%) | Weight gain (≥5%) | Weight loss (≥5%) - Unintentional | Weight loss (≥5%) - Intentional | Total |
|-----------|---------|-----------|-----------|---------|---------------------|-------------------|-----------------------------------|----------------------------------|-------|
| Stable WC circumference (WC) (±5%) | 26948     | 44.50   | 26948     | 4654    | 1516                | 2193               | 35311                             |                                  |       |
| WC gain (≥5%)                         | 76.32  | 13.18   | 76.32     | 39.32   | 45.86               | 45.94              | 58.32                             |                                  |       |
| WC loss (≥5%) - Unintentional         | 66.31  | 11.19   | 66.31     | 2.50    | 4.29                | 6.21               | 26.13                             |                                  |       |
| WC loss (≥5%) - Intentional           | 66.31  | 11.19   | 66.31     | 2.50    | 4.29                | 6.21               | 26.13                             |                                  |       |
| Total                                 | 40637  | 67.11   | 40637     | 11835   | 3306                | 4774               | 60552                             |                                  |       |
studies are needed to determine the differential influence of intentional versus unintentional weight loss on cancer outcomes in postmenopausal women.

Women’s Health Initiative Dietary Modification randomized trial

In the WHI Dietary Modification (DM) trial, 48,835 postmenopausal women, all with no prior breast cancer and normal baseline mammogram, were randomized to a dietary intervention (a low-fat eating pattern incorporating an increase in fruit, vegetables and grain intake) or to a usual diet comparison group between 1993 and 1998. Breast cancer and colorectal cancer were separate primary study endpoints [11]. The dietary modification group participated in 18 group sessions led by centrally trained nutritionists in year 1 and quarterly sessions throughout the dietary intervention period. Caloric restriction or weight loss were not intervention targets. After 1 year, all dietary targets were significantly and favorably changed in the dietary group, and there was a statistically significant 3% weight loss that remained significant at years 3 and 6 (all \( P < 0.001 \)) but attenuated over time [11, 12].

After 8.5 years (median) dietary intervention, among the 1764 incident breast cancer cases, the incidence rate was 8% lower in the dietary intervention group, a finding which was not statistically significant (\( P = 0.09 \)) [11]. However, during that dietary intervention period, a statistically significant reduction in deaths after breast cancer (breast cancer followed by death from any cause) was seen (\( P = 0.01 \)) [13]. Specifically, after 16.1 years cumulative follow-up, with 3,030 incident breast cancers, deaths after breast cancer continued to be significantly reduced (HR 0.82, 95% CI 0.70–0.96) in the dietary group [13]. In further analysis of this trial, breast cancer overall survival (breast cancer diagnosis followed by death from any cause, measured from the time of breast cancer diagnosis) was found to be significantly increased for intervention versus comparison group participants for breast cancers diagnosed during the dietary period (10-year survival of 82% versus 78% for intervention versus comparison, respectively, \( P = 0.01 \)) [14]. The dietary intervention had a greater benefit among obese participants for deaths after breast cancer [13] and breast cancer overall survival [14].

While the dietary intervention goal was to reduce fat intake to 20% of total energy, at one year, mean energy from fat was actually reduced to 24.3% (standard deviation [SD] 7.5%) [11] and was 29.8% at the end of the dietary intervention [12]. The magnitude of this intervention has been previously described as representing “a modest reduction in fat intake with minimal weight loss” which “could be easily achievable by many” [13].

The important question of whether dietary change or weight loss will alter outcome for women following a diagnosis of early-stage breast cancer [15] was not directly addressed in the WHI DM trial. As all participants were free of cancer at entry, depending on their subsequent date

ASSOCIATED CLINICAL STUDIES

WHI OS and endometrial cancer risk

Weight loss has also been associated with significantly lower risk endometrial cancer risk when similarly evaluated in the WHI OS. In this analysis, women with hysterectomy before enrollment (\( n = 33,317 \)) and women with prior endometrial cancer were excluded, leaving 36,793 women in the final analysis. Women with weight loss \( \geq 5\% \) had significantly lower endometrial cancer risk compared to women with stable weight (HR 0.71, 95% CI 0.54–0.95), with the association strongest among obese women with intentional, rather than unintentional, weight loss (HR 0.44, 95% CI 0.25–0.78) [10], a finding not seen in analyses examining breast cancer associations with weight loss [11]. Further studies are needed to determine the differential influence
of diagnosis, women could have either more time on the dietary intervention before or following diagnosis. As it appeared that exposure to the dietary intervention after breast cancer diagnosis had greater effect on breast cancer mortality [14], the study findings could have relevance to the adjuvant breast cancer setting as well.

The association between implementation of a low-fat eating pattern and reduction in deaths after breast cancer was unchanged by the addition of baseline weight and weight change to the analysis [13, 14]. As weight loss was not an intervention target in the trial, the WHI DM trial did not evaluate the hypothesis raised by the WHI Observational Study finding that associated ≥5% weight loss with lower breast cancer incidence [5]. However, the modest 3% weight loss seen in WHI DM dietary participants suggests nutritional composition change rather than weight loss was the driver of the favorable breast cancer outcome. Ongoing studies are exploring this question.

SUMMARY

In summary, complementary findings from the WHI OS and the WHI DM full-scale randomized cancer prevention clinical trials indicate two distinct routes to improve breast cancer outcomes. A low-fat diet rich in vegetables and fruits can improve breast cancer outcome. In addition, for overweight or obese postmenopausal women, modest weight loss can improve breast cancer outcomes, and may enhance the benefits of a healthy low-fat diet. The current message should be encouraging for postmenopausal women, namely, moderation regarding dietary composition and weight can significantly reduce a woman’s chance of being diagnosed with breast cancer and of dying after a breast cancer diagnosis.

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KP and RTC wrote the initial draft of the report. Data analysis was conducted by JL. All authors provided critical revision of the manuscript for intellectual content and gave final approval of the manuscript.

CONFLICTS OF INTEREST

Dr. Chlebowski reported being a consultant for AstraZeneca, Novartis, Amgen, Immunomedics and Pfizer. No other authors reported conflicts.

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