Adherence to Endocrine Therapy in Breast Cancer Adjuvant and Prevention Settings

Rowan T. Chlebowski¹,2, Jisang Kim², and Reina Haque³

Abstract

Adherence to oral endocrine therapy in adjuvant breast cancer settings is a substantial clinical problem. To provide current perspective on adherence to oral endocrine therapies, a comprehensive literature review was conducted. In adjuvant trials, endocrine therapy adherence is relatively high with greater adherence for aromatase inhibitors compared with tamoxifen. In contrast, adherence to adjuvant therapy in clinical practice is relatively poor, with only about 50% of women successfully completing 5-year therapy. Importantly, good adherence (>80% use) has been associated with lower recurrence risk. Endocrine therapy adherence in primary breast cancer prevention trials parallels that seen in adjuvant trials. Factors associated with nonadherence include low recurrence risk perception, side effects, age extremes, medication cost, suboptimal patient–physician communication, and lack of social support. Few prospective studies have evaluated interventions designed to improve adherence. Interventions currently proposed reflect inferences from clinical trial procedures in which clinical contacts are commonly greater than in usual practice settings. In conclusion, for optimal breast cancer outcome, adherence to endocrine therapy must improve. Although general recommendations likely to improve adherence can be made based on clinical trial results and preliminary prospective trial findings, research specifically targeting this issue is needed to establish effective intervention strategies.

Adjuvant endocrine therapy adherence in clinical trials and clinical practice

Clinical adjuvant endocrine therapy trials, in which adherence is commonly closely monitored, did not suggest a major adherence problem. In the National Surgical Adjuvant Breast and Bowel Project (NSABP) B-14 trial in patients with breast cancer receiving adjuvant tamoxifen or placebo, discontinuation rates were 23% in both groups at 60 months median follow-up (11). In the NSABP B-24 intraductal breast cancer trial, 60-month discontinuation rates for placebo were 30% compared with 33% for tamoxifen (12). Adherence rates based on self-report were 89% and 88%, respectively (13). Other studies found overestimated adherence of patients with breast cancer to tamoxifen based on prescription checks (8) or microelectronic monitoring (9). In this regard, in a report comparing nonadherence with adjuvant anastrozole using three separate databases in the same population, estimates of nonadherence varied from 32% to 50% (10); however, subjects in these databases had variable medical insurance coverage, which may partially explain adherence differences. Despite these concerns, consistent general conclusions have emerged from studies using various methods of adherence assessment.

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to 1996, adherence to adjuvant tamoxifen was 83% after 1 year, 68% after 2 years, 61% after 3 years, and only 50% after 4 years. Other studies also found that more than half of patients with breast cancer discontinue endocrine therapy before completion of a recommended 5-year treatment (13, 14). For aromatase inhibitors, the commonly experienced arthralgias raise particular adherence concerns (15–17). However, in several adjuvant clinical trials, adherence to aromatase inhibitors was closely comparable, or even superior, with tamoxifen. In the Arimidex, Tamoxifen, Alone and Combined (ATAC) trial after 5 years, 2.1% of the anastrozole-treated patients and 14.3% of the tamoxifen-treated patients had discontinued use due to adverse events (18). Similar adherence was seen in both treatment groups in adjuvant trials comparing the aromatase inhibitor, exemestane, with tamoxifen (14% discontinued therapy in both arms; ref. 19) and the aromatase inhibitor, letrozole, with placebo (only 10% discontinued therapy in both arms; ref. 20).

Recent systematic reviews on adherence and/or persistence to adjuvant endocrine therapy in clinical practice settings identified 29 reports. Adherence in tamoxifen users ranged from 41% to 88%, whereas adherence in aromatase inhibitor users ranged from 50% to 91% (21). These findings were extended by Huiart et al. (22) who conducted meta-regression analyses to provide summary estimates of nonpersistence in 17 trials. For tamoxifen, 5-year nonpersistence was 47.2% (95% confidence interval, 41.1%–53.5%) compared with 31.0% (95% CI, 25.9%–37.5%) for aromatase inhibitors (Table 1).

The findings are somewhat mixed considering aromatase inhibitor adherence in clinical practices (23, 24); however, in the United Kingdom general practice database, the 1-year discontinuation rate for adjuvant aromatase inhibitor use was 5% compared with about 10% for tamoxifen in women >49 years old and 20% for tamoxifen in women <40 years old (25). Similarly, in the Disease Analyses database (IMS Health), among 16,865 patients with breast cancer, 3-year discontinuation rates were 52% for tamoxifen, 47% for anastrozole, and 44% for letrozole (26). A randomized adjuvant adherence trial found shorter time to treatment discontinuation for exemestane, compared with letrozole (HR, 1.5; 95% CI, 1.1–2.1; ref. 27).

In summary, a substantial problem about adherence and persistence to adjuvant endocrine therapy remains in clinical practice. Somewhat surprisingly, adherence to aromatase inhibitors has been similar or superior to adherence to tamoxifen in several settings.

### Adjuvant endocrine therapy adherence and clinical outcome

Evidence that adherence to adjuvant endocrine therapy could influence clinical outcomes came from a series of randomized adjuvant breast cancer trials evaluating duration of tamoxifen use. As summarized in Early Breast Cancer Trialist Cooperative Group (EBCTCG) analyses, compared with no therapy/placebo, with tamoxifen for 1 year recurrence, reduction was 27%; for 2 years, reduction was 33%; and for 5 years, reduction was 47%; \( P \text{ trend} < 0.00001 \) (28).

Adherence to adjuvant tamoxifen therapy and breast cancer outcome has been examined in several cohort studies. In a U.S. cohort of 1,837 older women with early-stage breast cancer, those who used tamoxifen less than 1 year had substantially higher breast cancer mortality than those who used the drug for 5 or more years (HR, 6.26; 95% CI, 3.10–12.64; ref. 29). Similar findings were reported from a Scottish cohort of 2,080 patients with early-stage breast cancer. In that study, tamoxifen adherence <80% was associated with increased mortality (HR, 1.100; 95% CI, 1.001–1.21; ref. 30).

In the managed care Kaiser Permanente Northern California population, among 8,769 patients with breast cancer, 2,761 (31%) discontinued therapy within 6 months of diagnosis (based on automated pharmacy records); of those who continued, 1,684 (28%) were nonadherent (possession ratios <80%; defined as days with index prescription supplies/total days of follow-up). The survival at 10 years was 80.7% and 73.6% for those who continued therapy compared with those who discontinued therapy, respectively (\( P < 0.001 \); ref. 31). Of those who continued therapy, survival was 81.7% in those adherent to therapy, compared with 73.6% in those nonadherent. In a similar study in Kaiser Permanente Southern California, although breast cancer recurrence was lowest in women with greater adherence (possession ratios >80%), the rates were not markedly different from women with less regular use (32). In a retrospective cohort study of 3,361 Scottish patients with breast cancer, low adherence of <80% to adjuvant tamoxifen in aromatase inhibitor was associated with poor survival (HR, 1.20; 95% CI, 1.03–1.40; \( P = 0.019 \); ref. 33).

| Table 1. Systematic reviews of adherence to adjuvant endocrine therapy |
|--------------------------|---------------------------|
| **Adherence (range)**b | **Tamoxifen**  | **Aromatase inhibitor** |
| Therapy discontinuation (range)a | 41%–88% | 52%–91% |
| Five-year therapy discontinuation from meta-regression analysisb | 15%–20% within year 1 | 5%–25% within 2 years |
| | 47.2% (95% CI, 41.1%–53.5%) | 31.0% (95% CI, 25.9%–37.5%) |

*Murphy et al. (21).

Huiart et al. (22).*
In a prospective cohort of 417 localized patients with breast cancer in Sweden, nonadherence at 1 year was associated with increased early breast cancer events (HR, 2.97; 95% CI, 1.08–8.15; ref. 34). In a study with 857 low-income women with early breast cancer, more recurrences and cancer deaths were observed in women nonadherent to endocrine therapy, but the results were not statistically significant (35). Similarly, in 690 women participating in International Breast Cancer Study Group Trials 13–39 and 14–93, those with ≥4-year selective estrogen receptor modulator (SERM) use had longer disease-free survival compared with those with <4-year use (71% vs. 64%; HR, 1.31; 95% CI, 0.86–1.98; P, 0.20; ref. 36; Table 2). In a small study of 116 men with breast cancer, overall survival was greater in those adherent to tamoxifen adjuvant therapy (37).

Thus, the lack of adherence and persistence to prescribed endocrine adjuvant therapy represents a barrier to achieving favorable outcomes for patients with breast cancer. The magnitude of the benefit of being adherent to adjuvant endocrine therapy is comparable with that seen with the addition of adjuvant chemotherapy.

### Table 2. Studies relating duration of and/or adherence to adjuvant endocrine therapy to breast cancer outcome

<table>
<thead>
<tr>
<th>Lead author Study</th>
<th>Findings</th>
</tr>
</thead>
<tbody>
<tr>
<td>EBCTCG—Early Breast Cancer Trialist Collaborative Group 2001 Overview analyses of randomized clinical trials evaluating duration of tamoxifen use</td>
<td>Tamoxifen duration 1 year, recurrence reduced 27%; tamoxifen duration 2 years, recurrence reduced 33%; tamoxifen duration 5 years, recurrence reduced 47%; P trend &lt; 0.00001</td>
</tr>
<tr>
<td>Yood et al. (29) Cohort of 1,837 U.S. patients (≥65 years old) with early-stage breast cancer</td>
<td>Adjuvant tamoxifen &lt;1 year vs. ≥ 5 years with higher breast cancer mortality (HR, 6.26; 95% CI, 3.10–12.64)</td>
</tr>
<tr>
<td>McCowan et al. (30) Retrospective cohort of 2,080 Scottish patients with early-stage breast cancer</td>
<td>Adherence to tamoxifen &lt;80% associated with poorer survival (HR, 1.10; 95% CI, 1.001–1.21)</td>
</tr>
<tr>
<td>Hershman et al. (23) Northern California Kaiser Permanente cohort of 8,769 women with early-stage, hormone-sensitive breast cancer and endocrine therapy adherence (drug availability)</td>
<td>31% discontinued therapy, 10-year survival was 73.6% 69% continued therapy, 10-year survival was 80.7%; P &lt; 0.001</td>
</tr>
<tr>
<td>Xu et al. (37) Cohort of 116 men with early-stage, hormone-sensitive breast cancer and hormone therapy adherence</td>
<td>For those adherent, 10-year survival was 79.6%; for those nonadherent, 10-year survival was 50.5%; P = 0.008</td>
</tr>
<tr>
<td>Markkula et al. (34) Prospective cohort of 417 Swedish patients with early-stage breast cancer and adherence (self-report) to adjuvant endocrine therapy</td>
<td>Nonadherence at the 1-year visit associated with increased early breast cancer events (HR, 2.97; 95% CI, 1.08–8.15)</td>
</tr>
<tr>
<td>Haque et al. (32) Southern California Kaiser Permanente cohort of 22,850 women with early-stage breast cancer and endocrine therapy adherence (drug availability)</td>
<td>Women with high adherence had greater recurrence risk reduction (e.g., HR, 0.42; 95% CI, 0.36–0.47 for tamoxifen) compared with those with less adherence (HR, 0.46; 95% CI, 0.41–0.52 for tamoxifen) but the difference was not statistically significant.</td>
</tr>
<tr>
<td>Pagani et al. (36) International Breast Cancer Study Group trials 12–93 and 14–92 with 690 women with early-stage breast cancer or SERMs</td>
<td>Women with ≥4 years of SERM had longer 10-year disease-free survival (71%) compared with &lt;4-year use (64%), P = 0.20</td>
</tr>
</tbody>
</table>
endocrine therapy (or had discontinued treatment within 6 months) despite having pharmacy coverage (32). This finding stimulated that organization to implement a medication adherence tool in the electronic medical records to potentially improve adherence. In the Women’s Health Initiative cohort, in 3,588 patients with hormone receptor–positive, early-stage invasive breast cancer evaluated within 5 years of diagnosis by survey questionnaire, whereas adjuvant endocrine therapy use was reported by 83%, 17% reported no use. In their response, women cited “lack of physician recommendation” as the most common reason for nonuse (39). Finally, 743 patients, identified from Surveillance, Epidemiology, and End Results (SEER) registries, eligible for adjuvant endocrine therapy were surveyed 4 years after diagnosis, surprisingly 10.8% never initiated therapy, and 15.1% started therapy but discontinued before 4 years (40; Table 3). Although detailed information on the characteristics of those not initiating adjuvant endocrine therapy is not currently available, further exploration of this issue is warranted.

Data are sparse about the communication between oncologists and patients with breast cancer on their therapeutic plan, as it is difficult to conduct linguistic communication studies. However, one study videotaped the initial breast cancer adjuvant therapy discussion in a series of 28 early-stage patients and found the issue of adherence to be poorly addressed. Much of the discussions on endocrine therapy focused on side effects and trial findings rather than on the importance of adherence (41).

Finally, a recent study found substantial discordance in adherence to adjuvant endocrine therapy when comparing results among prescription refill information, patient self-report, and oncologists’ estimates. The oncologists estimated their patients’ adherence at over 94%, which was less than estimated by telephone questionnaire self-report ($P = 0.003$), or by the pharmacy database in which only 67% of women ≥65 years old were identified as having drug available ($P = 0.0001$; ref. 42).

### Table 3. Adjuvant hormone therapy use for hormone receptor–positive postmenopausal women with early-stage breast cancer

<table>
<thead>
<tr>
<th>In Women’s Health Initiative Cohort</th>
<th>Kaiser Permanente Southern California</th>
<th>In SEER Population by Survey</th>
<th>Kaiser Permanente Northern California</th>
</tr>
</thead>
<tbody>
<tr>
<td>Use: AI 33%, SERM 31%, mix 36%</td>
<td>Use: SERM 38%, AI 19%, mix 16%</td>
<td>Use: endocrine 75%</td>
<td>Not examined</td>
</tr>
<tr>
<td>17% none</td>
<td>24% none</td>
<td>10.8% none</td>
<td>30% none</td>
</tr>
<tr>
<td>33% of users became nonadherent</td>
<td>21% users became nonadherent</td>
<td>15.1% uses became nonadherent by year 4</td>
<td>Not examined</td>
</tr>
</tbody>
</table>

Abbreviation: AI, aromatase inhibitor.

*Livaudais et al. (39).
*Haque et al. (32).
*Frease et al. (40).
*Livaudais et al. (38).

### Adjuvant endocrine therapy adherence in long-duration clinical trials

Interest in adherence to long-term adjuvant endocrine therapy regimen was enhanced by the recent report from the worldwide Adjuvant Tamoxifen Longer Against Shorter (ATLAS) adjuvant trial in which continued tamoxifen use for longer than 5 years reduced breast cancer recurrence ($P = 0.002$) and overall mortality ($P = 0.01$; ref. 4). Based on self-report, 5-year adherence was an excellent 84% for continued tamoxifen users. In contrast, the Investigation on the Duration of Extended Adjuvant Letrozole treatment (IDEAL) trial entered 1,250 patients with early breast cancer comparing 2.5 years with 5 years of extended letrozole use after 5 years of adjuvant endocrine therapy found overall nonadherence was 18.4% at 2.5 years (43). It is not clear whether this apparent difference between long-term continued tamoxifen and continued aromatase inhibitor use represents real differences in tolerability, or is the result of the limited data on the aromatase inhibitors currently available. In any event, more information is needed about persistence to long-term aromatase inhibitor adjuvant use.

### Endocrine therapy adherence in breast cancer prevention trials

Available evidence suggests that adherence to endocrine therapy in primary breast cancer prevention trial participants is similar to that seen in the adjuvant setting. In the NSABP P-1 prevention trial, discontinuation rates after 54.6 months mean follow-up were 23.7% on tamoxifen, and 19.7% on placebo (11). Discontinuation rates were somewhat higher in the International Breast Intervention Study-1 (IBIS-1) in which, in a primary prevention setting, the 50-month median follow-up discontinuation rate for tamoxifen was 36% compared with 26% for placebo (44). In the longer intervention duration Royal Marsden Hospital trial comparing tamoxifen with placebo, therapy was prematurely discontinued at a median of 70 months in 46% of tamoxifen and 36% of placebo participants, respectively.
In the NSABP Study of Tamoxifen and Raloxifene (STAR) prevention trial, 5-year adherence was 70.8% for tamoxifen and 73.9% for raloxifene \((P < 0.001; \text{ref. 46})\). 

The aromatase inhibitor, exemestane, has been compared with placebo for primary breast cancer prevention in the Mammary Prevention (MAP)3 trial. After median 35 months follow-up, a 65%, statistically significant, relative reduction in invasive breast cancer incidence was seen for exemestane \((P = 0.04; \text{ref. 46})\). In the MAP trial, exemestane was discontinued because of “intolerable side effects” by 15.4% of participants but, surprisingly, 10.8% of placebo participants discontinued study pills for the same reason. With only a net 5.3% difference, a major influence of factors other than drug side effects likely influenced the adherence results seen. A similar result was seen in the MA.17 adjuvant trial, in which about 20% of patients with breast cancer in the placebo group reported climactic symptoms \((48)\). These results point to the importance of placebo controls to generate the most reliable tolerability information.

In a prevention study, adherence was related to outcome in the Women’s Health Initiative (WHI) trial of estrogen alone. In this study, when 10,739 postmenopausal women with prior hysterectomy were randomized to conjugated equine estrogen alone or placebo, surprisingly, a statistically significant, lower breast cancer incidence was seen in the estrogen-alone group in intent-to-treat analyses \((HR, 0.77; 95\% \text{ CI}, 0.62–0.95; \text{ref. 49})\). However, in sensitivity analyses, censoring participants with less than 80% adherence to the pill-taking regimen, an even stronger association between estrogen-alone use and lower breast cancer incidence was seen \((HR, 0.68; 95\% \text{ CI}, 0.49–0.95)\).

**Factors associated with nonadherence to endocrine therapy in breast cancer prevention trials**

Factors predictive of tamoxifen chemoprevention nonadherence were examined in the P-1 breast cancer prevention trial. Current smokers and heavy alcohol users had lower tamoxifen adherence, whereas obesity and lower physical activity were unrelated to adherence \((50)\). Similar findings were seen in 100 participants in the IBIS-1 study where women with smoking history also were less likely to persist with their randomized drug \((51)\). In addition, in the IBIS-1 trial, use of additional prescribed medication was an important factor in predicting successful completion of therapy \((P = 0.04; \text{ref. 51})\). The latter findings suggest that women already using other prescription medications may represent a potentially favorable population, and thus, be more likely to accept and adhere to endocrine chemoprevention regimens. Lack of influence of obesity and low physical activity on adherence suggests that factors other than an unhealthy lifestyle are related to medication discontinuation.

**Endocrine therapy for prevention in clinical practice**

Currently, use of the two drugs approved for chemoprevention in the United States (tamoxifen and raloxifene) continues to be low \((52)\), and, for this reason, information on adherence in clinical practice settings is not available. However, a review of a clinical experience from the Partners HealthCare System identified 2,938 women with breast lesions with atypia. Women who received no chemoprevention had 10-year breast cancer incidence of 21.3% compared with 7.5% \((P < 0.001)\) in women who did receive chemoprevention \((53)\).

**Factors associated with nonadherence to adjuvant endocrine therapy**

Factors associated with nonadherence to adjuvant hormonal therapy include lack of physician recommendation \((32)\), patient perception of low risk for recurrence \((54)\), adverse effects of therapy \((55, 56, 57)\), age extremes: older age \((23, 58)\) and younger age \((23, 59)\), medication costs \((60, 61, 62)\), low social economic status \((63)\), suboptimal patient–physician communication \((64)\), higher comorbidity \((23, 59, 62)\), cigarette smoking \((50, 51)\), and lack of social support \((65)\); Table 4). Similar factors were associated with adherence in a low-income population in California \((66)\). Findings about adherence by race/ethnicity have produced mixed results \((23, 38, 67)\).

Many oncologists likely consider endocrine therapy side effects to be a major factor influencing therapy adherence. However, the available evidence identifies a less-straightforward relationship. In patients with breast cancer in the Commonly used Medications and Breast Cancer outcomes (COMBO) study, among 538 participants, 18.2% discontinued use before completing 5 years of therapy, whereas 25% discontinued after <1-year use \((68)\). As in several prior reports, women who discontinued therapy were more likely to have been tamoxifen \((43.9\%)\) compared with aromatase inhibitor users \((22.4\%)\). Of interest, the only adverse effect significantly associated with discontinuation of both aromatase inhibitor and tamoxifen was headaches, an adverse event not commonly associated with these therapies. Such findings suggest that, although control of adverse effects is an important clinical consideration, adverse effects of endocrine therapy use may not play a major role in determining adherence and persistence to adjuvant endocrine therapy.

Factors adversely influencing adherence, perhaps in unexpected ways, are anxiety and depression. Following a breast cancer diagnosis, anxiety and depression decrease from about 50% in year one to about 15% in year five \((69)\), a reciprocal to endocrine therapy adherence over the same period \((1, 3, 70; \text{Fig. 1})\). Supporting the concept that greater patient anxiety correlates with better adjuvant hormone therapy adherence are findings from the prospective COM-PAS study in which patients with breast cancer with higher anxiety levels had better adherence to adjuvant endocrine therapy \((P = 0.028; \text{ref. 71})\). In an extremely large breast cancer population from IMS Health with 17,512 patients, depression \((P < 0.002)\) was also associated with decreased risk of treatment discontinuation \((26)\). As anxiety and depression can be linked in a cancer population, unraveling the relative contribution of these two factors on adjuvant endocrine therapy adherence requires further study. In this...
regard, despite early concerns, evidence from the NSABP placebo-controlled clinical prevention trial found depression was not increased by tamoxifen use (72, 73).

Clinical trials to improve endocrine therapy adherence

Few prospective studies have evaluated interventions designed to improve adherence to endocrine adjuvant therapy. However, to guide future study designs, theoretical models of factors influencing adherence and persistence have been proposed (74).

Although adherence to endocrine therapy in breast cancer adjuvant and prevention settings remains problematic, there are limitations to the currently available information. As reviewed (59, 75), only modest information about factors associated with continued hormone therapy use is known and, importantly, few of the factors identified are easily modifiable. In addition, current medical claims databases, commonly used in adherence analyses, contain limited information on healthcare practice patterns or patient characteristics needed to identify new potentially modifiable factors.

Despite the important influence of adjuvant endocrine therapy adherence on clinical outcome, there has only been one full-scale, randomized intervention trial designed to improve adherence completed to date. The Patient’s Anastrozole Compliance to Therapy (PACT) program was a randomized, prospective, multicenter study designed to improve persistence and compliance to adjuvant endocrine therapy (76). In this trial, 4,844 patients were randomly assigned to standard therapy or standard therapy plus mailed educational materials including monthly reminders on persistence and additional letters and brochures. Questionnaires were completed before therapy was initiated, at 12 and 24 months and at treatment discontinuation. At 1 year, there was no difference in the primary endpoint of compliance (88.5% vs. 88.8 %; $P = 0.81$). Thus, provision of education materials did not increase adherence to adjuvant endocrine therapy (76).

A more promising result was seen in a smaller COM-PAS study of 181 patients receiving adjuvant aromatase inhibitor therapy. The randomization was either to a control condition, a letter group in which participants received five mailings in the first year and three in the second, and a telephone group in which participants were contacted by a study nurse using a semistructured interview technique at the same intervals as in the letter group (77). Adherence was determined as a composite of self-report using a standardized questionnaire plus medication possession ratios calculated from pharmacy prescription refill information. At 12 months, 48% in the control group, 63% in the telephone group, and 65% in the letter group were judged adherent. Although the differences between the groups were not statistically significant, a post hoc analysis pooling both interventions versus control indicated a significant difference favoring intervention ($P = 0.039$). These encouraging results

Table 4. Correlates associated with discontinuing endocrine therapy or nonadherence

<table>
<thead>
<tr>
<th>Reason</th>
<th>Study</th>
</tr>
</thead>
<tbody>
<tr>
<td>Side effects</td>
<td>Demissie et al. (55), Kahn et al. (56), Lash et al. (57), Cluze et al. (65)</td>
</tr>
<tr>
<td>Higher comorbidity</td>
<td>Hershman et al. (23), Haddj (26), Sedj and Devine (62)</td>
</tr>
<tr>
<td>Financial considerations or low socioeconomic status</td>
<td>Kimmek et al. (85), Riley et al. (60), Neugut et al. (61), Liu et al. (66)</td>
</tr>
<tr>
<td>Very young or older age</td>
<td>Hershman et al. (23), Land et al. (50), Owusu et al. (58)</td>
</tr>
<tr>
<td>Lack of physician recommendation</td>
<td>Davidson et al. (41)</td>
</tr>
<tr>
<td>Perception of low risk of recurrence</td>
<td>Fink et al. (54)</td>
</tr>
<tr>
<td>Lack of social support</td>
<td>Cluze et al. (86), Land et al. (50)</td>
</tr>
<tr>
<td>Follow-up care with general practitioner vs. oncologist</td>
<td>Murphy et al. (21)</td>
</tr>
<tr>
<td>African American race/ethnicity</td>
<td>Hershman et al. (23)</td>
</tr>
<tr>
<td>Cigarette smoking</td>
<td>Land et al. (50)</td>
</tr>
<tr>
<td>Presence of anxiety/depression linked to better adherence</td>
<td>Kyvenetakis et al. (71), Haddj (26)</td>
</tr>
<tr>
<td>Alcohol use</td>
<td>Land et al. (50)</td>
</tr>
</tbody>
</table>

Figure 1. Nonadherence rates for adjuvant tamoxifen therapy in clinical practice and incidence of depression and/or anxiety in breast cancer patients with early stage disease.
provide a foundation for a future confirmation trial as either intervention would be feasible for implementation in clinical practice settings.

Focus on endocrine therapy patient education: "optimization of expectations"

There is emerging evidence that a patient’s expectation about the benefits and drawbacks of a therapy can influence adverse effects and persistence with therapy. A meta-analysis identified significant associations between cancer patients’ expectation of developing adverse effects and the actual adverse effect experience (78). When 597 patients with early-stage breast cancer who were prescribed tamoxifen were followed for 2 years, 17% discontinued tamoxifen use. Of these, women with neutral or negative beliefs about tamoxifen efficacy were significantly more likely to discontinue than those with more positive beliefs. Based on these and similar findings, several strategies to enhance endocrine therapy adherence are now focused on the development and testing of structured educational sessions implementing at the beginning of therapy with the goal of optimization of expectations. In another study, the balance between efficacy and side effects was assessed in women receiving adjuvant endocrine therapy with an Adaptive Conjoint Analysis customized to each patient. Using such information, a benefit/drawback ratio was calculated and the 16% of women who valued the efficacy less than the adverse effects had substantially lower adherence (79). Based on such findings, an ongoing randomized, controlled trial is evaluating a three-session program of cognitive, behavioral training designed to provide a realistic and balanced view of endocrine therapy (80).

Recommendations for improving adherence and clinical practice

Despite the paucity of full-scale clinical trial evidence, there are strategies for implementation in current clinical practice which would likely have a favorable effect on endocrine therapy adherence resulting in more favorable clinical outcome.

Adherence to endocrine adjuvant therapy has been higher in clinical trials where patient contacts are commonly greater than in clinical practice settings, and where concerned attention is directed at encouraging the maintenance of adherence. Strategies to increase patient contacts, which incorporate emerging technologies such as email reminder programs and use of cell phone apps (81–84) shown to improve adherence in other disease settings, seem promising to evaluate in breast cancer trials. Strategies to increase contacts with patients in practice settings include use of automated telephone refill reminders and implementing medication adherence tools in electronic medical records. The concept that increased contacts with patients would increase endocrine therapy adherence is strengthened by the findings from the COMPAS trial in which both additional mailings and telephone contacts seem to influence favorable adherence (77).

While awaiting results of ongoing clinical studies, one could reasonably conclude that attention to endocrine therapy patient education to optimize realistic patient expectations for adjuvant endocrine therapy, besides being good medical practice, could also improve therapy adherence. For infusional chemotherapy, in many practices, the benefits and risks of therapy, originally discussed by the oncologist, are reinforced in formal chemotherapy education sessions by a mid-level provider. A similar approach to improve adjuvant endocrine therapy patient education could be considered. Educational interventions should focus on increasing patients’ understanding of the benefits and risks of therapy including the relationship between therapy adherence and persistence and higher efficacy of the therapy in reducing cancer recurrence. Implementation of these recommendations is likely to favorably affect adherence in clinical practice at this time. More definitive evidence must come from future activity in the research arena.

Disclosure of Potential Conflicts of Interest

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