# **Dietary Soy Intake and Breast Cancer Risk**

Carol A. Enderlin, PhD, RN, Elizabeth Ann Coleman, PhD, RNP, AOCN®, Carol Beth Stewart, BS, and Reza Hakkak, PhD

bout one in eight women born in the United States today is at risk for receiving a breast cancer diagnosis during his or her lifetime. In 2009, an estimated 192,370 women will be diagnosed with and 40,170 will die from breast cancer (National Cancer Institute [NCI], 2009). The risk for developing invasive breast cancer increases as women age, with about 66% aged 55 years or older at diagnosis (American Cancer Society [ACS], 2009). Age-related breast cancer risks, however, do not reflect an individual woman's risks, which may be greater or smaller depending on a number of factors. In addition to older age and female gender, being of Caucasian race, having inherited certain genetic mutations, having a family or personal history of breast cancer, having more dense breast tissue, having begun menstrual periods before the age of 12 years or menopause after the age of 55 years, radiation treatment to the chest area early in life, and prior treatment (mothers and daughters) with diethylstilbestrol are considered nonmodifiable risk factors. Modifiable risk factors include having not had children or having had the first pregnancy after the age of 30 years, having no history of breastfeeding, being overweight or obese, having a lack of exercise, recent use of birth control pills, postmenopausal hormone therapy, and alcohol use (ACS). The pressing need to discover some means to decrease the risk of breast cancer development has led scientists to examine soy foods as possible prevention strategies.

# Soy Isoflavones, Sources, and Effects Soy Isoflavones

Soybeans, also referred to as soy or soya, are plants of Asian originthat produce beans used in a variety of food products (Biology Online, 2005). Soybeans and soy products are a major source of phytoestrogen, an estrogenlike substance (MedlinePlus, 2008) also referred to as plant estrogen. Isoflavones, one class of phytoestrogens, are structurally similar to mammalian estrogens, have estrogenic properties, and are potential anticarcinogens (Peeters, Keinan-Boker, van der Schouw, & Grobbee, 2003). Isoflavones also are considered a subclass of

**Purpose/Objectives:** To conduct a metasynthesis of the literature on human studies of the relationship between dietary soy intake and breast cancer risk.

**Data Sources:** Publications in English reporting human studies were searched with the terms soy and breast cancer, using Ovid®, PubMed, and EBSCO databases. Only human studies investigating the relationship of soy intake to breast cancer development in women published from January 1997 through June 2008 were included in the review.

**Data Synthesis:** A total of 364 publications were located; 18 of the studies met the inclusion criteria and 18 additional studies were located through other publications identified in the search. Because four articles reported on the same two studies, a total of 34 studies were included in the review.

**Conclusions:** The naturally occurring dietary intake of soy food or its components appears safe for women without breast cancer; however, the safety of high supplements of soy or its components is less certain.

**Implications for Nursing:** Nurses should become more knowledgeable about soy foods and supplements and include soy intake in dietary assessments. Nurses caring for women at high risk for or with a history of breast cancer should confer with dietitians on current practice recommendations. Women with health issues should avoid initiating high intake of soy dietary supplements until the possible effects are better understood.

flavonoids, a large family of compounds synthesized by plants and thought to have potential antioxidant properties (Linus Pauling Institute, 2008). Antioxidants are substances that protect cells from damage caused by free radicals produced by oxidation during normal metabolism, thought to play a role in cancer development (NCI, 2004). The most common dietary isoflavones are genistein, daidzein, and glycitein (Linus Pauling Institute), sometimes also referred to as isoflavonoids (Kelly, Nelson, Waring, Joannou, & Reeder, 1993). The major metabolite of daidzein is equol, a nonsteroidal estrogen produced in the intestines (Medicinenet.com, 2004). Only about 33% of the population from Western cultures are capable of producing equol based on the findings of urinary excretion studies (Setchell, Brown, & Lydeking-Olsen, 2002). Because equol has greater estrogenic activity than daidzein or other metabolites, differences in its production may account for some of the variation in cancer association reported among different racial or ethnic populations (Linus Pauling Institute). Another daidzein metabolite, O-demethylangolensin, also is thought to be formed in the intestines (Adlercreutz, 1995) and has variable urinary excretion rates possibly related to individual variability in the metabolism of dietary isoflavones (Kelly et al.).

#### **Soy Dietary Sources**

Soybeans are the main dietary source of isoflavones. Foods made from soybeans in the traditional Asian diet include tofu, miso, natto, tempeh, and edamame. Foods made from soybeans eaten in Western countries include soy milk, soy cheese, soy yogurt, and soy meat substitutes. Soy isoflavone supplements and extracts are available as dietary supplements, although their actual isoflavone content is not standardized and may vary (Linus Pauling Institute, 2008). Higdon (2007) identified some common foods made from soybeans, their serving sizes, and respective isoflavone content (see Table 1). An estimated upper limit for isoflavone intake is 100 mg per day, or the approximate amount in three servings of traditional soy foods (Messina, 2008).

## Soy Estrogenic and Nonestrogenic Effects

Soy isoflavones and their metabolites exert weak estrogenic activity by binding to estrogen receptor sites on cells, mimicking estrogen in some types of tissue, and blocking estrogen effects in others (Wang, 2002). Soy isoflavones also can act independently of estrogen receptors, including inhibition of enzymes involved in estrogen metabolism and cell proliferation as well as antioxidant activities (Barnes et al., 2000). Consequently, possible relationships between dietary soy intake and the development of hormone-associated cancers, including breast cancer, remain a current focus of research interest.

**Table 1. Isoflavone Content of Selected Soy Foods** 

| Food                   | Serving  | Total<br>Isoflavones<br>(mg) | Daidzein<br>(mg) | Genistein<br>(mg) |
|------------------------|----------|------------------------------|------------------|-------------------|
| Tofu                   | 3 ounces | 20                           | 8                | 12                |
| Miso                   | 1/2 cup  | 59                           | 22               | 34                |
| Tempeh                 | 3 ounces | 37                           | 15               | 21                |
| Edamame                | 1/2 cup  | 12                           | 6                | 6                 |
| Soy milk               | 1 cup    | 30                           | 12               | 17                |
| Soy cheese, mozzarella | 1 ounce  | 2                            | 0.3              | 1                 |
| Tofu yogurt            | 1/2 cup  | 21                           | 7                | 12                |
| Soy meat substitute    |          |                              |                  |                   |
| Hot dog                | 1 dog    | 11                           | 3                | 6                 |
| Sausage                | 3 links  | 3                            | 0.6              | 2                 |

Note. From An Evidence-Based Approach to Dietary Phytochemicals (p. 132), by J. Higdon, 2007, New York: Thieme Medical. Copyright 2007 by Thieme Medical. Adapted with permission.

# **Clinical Evidence of Dietary Soy and Breast Cancer Development**

#### Literature Review

The current review was guided by experts in the process of reviewing and synthesizing literature (Cooper, 1998; Stock, 1994). A search was conducted for publications in English reporting human studies using the terms *soy* and *breast cancer* with Ovid®, PubMed, and EBSCO databases, including CINAHL® and Academic Search Elite.

Only human studies that investigated the relationships between dietary soy intake and breast cancer development in women published from January 1997 through June 2008 were selected as representative of a period of intense investigation of this topic. The intervention studies selected were analyzed, summarized, and synthesized to answer three questions on the relationship between dietary soy intake and breast cancer development in women: what are the characteristics of research studies over the past decade, what do current research findings say about the relationship of soy intake to the development of breast cancer, and what general conclusions can be drawn from these findings?

Using the search terms of *soy* and *breast cancer*, 305 articles were located from OVID, 105 from PubMed, and 114 from EBSCO. After deletion of duplicates, 364 studies remained. Eighteen studies were human studies published from January 1997 through June 2008 that investigated the relationship of soy with breast cancer development in women. Eighteen additional studies were located through publications identified in the search. A total of 36 published studies were included in this review. Because four articles reported on the same two studies, 34 actual studies were reviewed (see Table 2). The 346 excluded publications included 31 literature reviews or meta-analyses related to soy intake and breast cancer risk and 61 general articles (e.g., commentaries, issues, informational, research briefs).

The other excluded articles addressed topics that were related to soy intake, breast, or other cancers but were not the focus of this review. They included mammographic density or benign breast disease (n = 15); menopausal symptom management (n = 16); tamoxifen, breast cancer treatment, or survival (n = 19); cognition (n = 1); diabetes (n = 1); biochemical, genetic, cellular, or endocrine effects (n = 113); nutrition or nutrition interventions (n = 55); prostate cancer (n = 6); general cancer prevention (n = 8); dermatologic uses (n = 2); and mouse, rat, or other animal studies (n = 18).

#### **Selected Study Characteristics**

**Design:** The 34 studies selected included cohort and case-control studies. Six had a

cohort design, including three prospective cohort studies. Twenty-eight had a case-control design, including 1 familial matched, 11 nested, 6 hospital-based, and 17 population-based case-control studies.

**Race or ethnicity:** The populations from which the study samples were derived represented 12 countries and multiple races and ethnicities. Study populations were: Chinese (n = 6); Japanese (n = 6); British (n = 3); Dutch (n = 3); Australian (n = 2); Canadian (n = 2); Filipino (n = 2); German (n = 2); American and predominantly Caucasian (n = 1); Asian American, including Chinese and Japanese, and non-Asian American such as African American, Latino, and Caucasian (n = 1); American and Canadian, predominantly Caucasian (n = 1); South Asian residing in England (n = 1); French (n = 1); Italian (n = 1); Greek (n = 1); and South Korean (n = 1).

Menopausal status: Four of the studies addressed breast cancer risk in premenopausal women only, two in postmenopausal women only, and the remainder addressed breast cancer risk in women from premenopausal through postmenopausal status.

**Dietary soy forms:** The forms of dietary soy intake included soy foods (total, fermented, and unfermented), soybeans (fresh and dried), soy proteins, tofu (soybean curd), and tofu-containing foods (particularly miso soup and natto), phytoestrogens (traditional and nontraditional soy foods, soy-enriched flour, and soy protein-enriched canned foods), isoflavones, and the specific isoflavones daidzen and genistein.

Dietary soy intake assessment: Assessment of dietary soy intake varied among the studies. A food frequency questionnaire was used in 17 studies, administered either by self-report or by an interviewer. Structured or semistructured interviews or a food diary were used in 11 studies. A nutrient database or food composition table, including U.S. Department of Agriculture and Taiwanese databases, was used in nine studies to determine dietary soy intake.

**Dietary soy measures:** Direct measures of dietary soy intake included times eaten per day or week, soy proteins in grams per week, tofu in times per week or year, phytoestrogens in micrograms per day, isoflavones in milligrams per day or milligrams per week, and daidzen and genistein in micrograms or milligrams per day. Indirect measures of dietary soy intake included serum and urine levels of total isoflavonoid (isoflavones) and of a variety of specific isoflavonoids (isoflavones) including daidzen, genistein, glycitein, equol, and O-desmethyangolensin. Serum isoflavonoid (isoflavone) levels were reported in nanograms per milliliter, and urine isoflavonoid (isoflavone) levels in nanomols per milligram of urinary creatinine, or micrograms per millimol of urinary creatinine. Serum isoflavone levels were reported in micrograms per milliliter, and daidzen and genistein in nanograms per milliliter.

## **Findings of Reduced Breast Cancer Risk**

Eighteen studies suggested some reduction in breast cancer risk associated with the dietary intake of soy or its components or the plasma levels or urinary excretion of isoflavone metabolites.

Premenopausal: Higher dietary intake of tofu and isoflavones (Hirose et al., 2005; Hirose, Takezaki, Hamajima, Miura, & Tajima, 2003; Witte et al., 1997), and specifically the isoflavones daidzein and genistein (Linseisen, Piller, Hermann, & Chang-Claude, 2004), were associated with a reduced risk of premenopausal breast cancer.

**Postmenopausal:** Higher urinary excretion of the specific isoflavone diadzein was found in postmenopausal women without breast cancer compared to patients with breast cancer, indirectly suggesting a protective association between higher dietary isoflavone intake and breast cancer (Murkies et al., 2000).

Premenopausal through postmenopausal: Higher dietary soy food intake has been associated with decreased breast cancer risk in pre- and postmenopausal women (Dai et al., 2001; Shu et al., 2001; Wu, Yu, Tseng, Hankin, & Pike, 2003) with a stronger risk reduction for hormone-responsive types of breast cancer (Dai et al.; Shu et al.), although Lee et al. (2005) found a reduced risk only in women aged 40 years or older.

Specific to the type of soy food, a higher dietary intake of miso soup has been associated with reduced breast cancer risk (Yamamoto, Sobue, Kobayashi, Sasaki, & Tsugane, 2003) as has a higher intake of yellow or black soybeans (Do, Lee, Jung, & Lee, 2007; Do, Lee, Kim, Jung & Lee, 2007). Lastly, specific to the age at which soy was consumed, a higher dietary phytoestrogen and soy food intake by adolescents (Thanos, Cotterchio, Boucher, Kreiger, & Thompson, 2006; Wu et al., 2002) and a higher intake by adult women (Wu et al., 2002) was associated with reduced breast cancer risk.

Two studies found that a higher dietary intake of isoflavones and, more specifically, the isoflavones daidzein and genistein, has been associated with reduced breast cancer risk (Dos Santos Silva et al., 2004; Yamamoto et al., 2003). Higher serum plasma levels of the isoflavone genistein have been associated with reduced breast cancer risk in two studies (Iwasaki et al., 2008; Lampe et al., 2007; Verheus et al., 2007), indirectly suggesting a protective association between a higher dietary isoflavone intake and breast cancer. Higher urinary excretion rates of total isoflavonoids (isoflavones), the specific isoflavone glycitein (Zheng et al., 1999), and the daidzein metabolite equol (Ingram, Sanders, Kolybaba, & Lopez, 1997) have all been associated with reduced breast cancer risks, again indirectly suggesting a protective association between higher dietary isoflavone intake and breast cancer. Specific to the age at which isoflavones were consumed, a higher dietary intake of isoflavones during adolescence has been associated with reduced breast cancer risk (Thanos et al., 2006).

**Table 2. Studies Investigating the Relationship Between Dietary Soy Intake and Breast Cancer Risk** in Women

| Study  | Design                                | Sample   | Soy Type   | Findings  |
|--|---------------------------------------|--|--|---|
| Bosetti et al.,<br>2005  | Case control;<br>hospital-based       | Italian: pre- and post-<br>menopausal cases<br>(n = $2,569$ ) and con-<br>trols (n = $2,588$ )           | Dietary isoflavones  | No significant association was found between dietary isoflavone intake and breast cancer risk.  |
| Boucher et al.,<br>2008  | Case control;<br>population-<br>based | Canadian: pre- and postmenopausal cases (n = 372) and controls (n = 356)                                 | Infant soy formula   | Exclusive soy feeding from $0-4$ months of age was associated with a 58% reduction in breast cancer risk (odds ratio [OR] = 0.42, 95% confidence interval [CI] = 0.18–1.9) and, from 5–12 months of age, with a 41% reduction in breast cancer risk (OR = 0.59, 95% CI = 0.18–0.9).   |
| Cui et al.,<br>2007  | Case control;<br>population-<br>based | Chinese: pre- and postmenopausal cases $(n = 1,459)$ and controls $(n = 1,556)$                          | Vegetable-soy<br>or meat-sweet<br>dietary patterns   | No association was found between dietary patterns and breast cancer risk.   |
| Dai et al.,<br>2001; Shu et<br>al., 2001                         | Case control;<br>population-<br>based | Chinese: pre- and postmenopausal cases $(n = 1,459)$ and controls $(n = 1,556)$                          | Dietary soy<br>protein and<br>isoflavones  | Highest compared to lowest soy intake was associated with a 34% reduction in breast cancer risk (OR = 0.66, 95% CI = 0.46–0.95, p for trend = 0.28) and a 56% reduction in estrogen and progesterone receptor site-positive breast cancer risk (OR = 0.44, 95% CI = 0.25–0.78, p for trend = 0.05).                         |
| den Tonkelaar<br>et al., 2001                                    | Case control;<br>population-<br>based | Dutch (Caucasian):<br>postmenopausal cases<br>(n = 88) and controls<br>(n = 268)                         | Urine genistein  | No significant association was found for the highest compared to the lowest urinary genistein excretion and breast cancer risk.   |
| Do, Lee, Kim,<br>et al., 2007;<br>Do, Lee, Jung,<br>et al., 2007 | Case control;<br>hospital-based       | South Korean: pre-<br>and postmenopausal<br>cases (n = 359) and<br>controls (n = 708)                    | Soy food grams   | No association was found between total soy intake and breast cancer risk in pre- or postmenopausal women. Higher compared to lower intake of yellow or black soybeans was associated with an approximate 33% risk reduction for breast cancer (OR = 0.67, 95% CI = 0.45–0.91, p for trend $<$ 0.02).                        |
| Dos Santos<br>Silva et al.,<br>2004                              | Case control;<br>population-<br>based | South Asian women<br>in England: pre- and<br>postmenopausal cases<br>(n = 240) and controls<br>(n = 477) | Dietary isoflavones  | Highest compared to lowest dietary isoflavone intake was associated with a 42% reduction in breast cancer risk (OR = $0.58$ , $95\%$ CI = $0.33-1$ , p for trend = $0.08$ ).  |
| Grace et al.,<br>2004  | Case control;<br>population-<br>based | United Kingdom: pre-<br>and postmenopausal<br>cases (n = 333) and<br>controls (n = 114)                  | Dietary phyto-<br>estrogens and<br>serum isoflavones<br>(daidzein, genistein<br>glycitein, and their<br>metabolites O-des-<br>methylangolensin<br>and equol) | Higher compared to lower serum daidzen, serum equol, and urine equol were associated with a 22%, 45%, and 34% increase in breast cancer risk (adjusted odds ratio [AOR] = $1.22$ , $1.005-1.481$ ; p = $0.044$ ; AOR = $1.455$ , $1.051-2.017$ ; p = $0.024$ ; AOR = $1.344$ , $1.063-1.699$ ; p = $0.013$ , respectively). |
| Hirose et al.,<br>2003   | Case control                          | Japanese: pre- and postmenopausal cases $(n = 2,385)$ and controls $(n = 9,013)$                         | Tofu   | Higher compared to lower tofu intake was associated with a 16% reduction in breast cancer risk in premenopausal women only (OR = $0.84$ , $95\%$ CI = $0.67$ – $1.04$ , p for trend = $0.02$ ).   |
| Hirose et al.,<br>2005   | Case control;<br>hospital-based       | Japanese: pre- and<br>postmenopausal cases<br>(n = 167) and controls<br>(n = 854)                        | Soybean products, including tofu and dietary isoflavones   | Highest compared to lowest tofu and isoflavone intake was associated with a 51% and 56% reduction in premenopausal breast cancer risk (AOR = 0.49, 95% CI = 0.25–0.95, p for trend = 0.03; AOR = 0.44, 95% CI = 0.22–0.89, p for trend = 0.02).  (Continued on next page)   |

Table 2. Studies Investigating the Relationship Between Dietary Soy Intake and Breast Cancer Risk in Women (Continued)

| Study                        | Design                                | Sample   | Soy Type  | Findings   |
|------------------------------|---------------------------------------|--|---|--|
| Horn-Ross et<br>al., 2001    | Case control;<br>population-<br>based | Non-Asian American (African American, Latino, and Caucasian): cases (n = 1,272) and controls (n = 1,610) | Dietary<br>phytoestrogens<br>(traditional and<br>nontraditional soy<br>foods)   | No significant association was found for the highest compared to the lowest phytoestrogen intake and breast cancer risk for pre- or postmenopausal women, ethnic group, or phytoestrogen type.                     |
| Horn-Ross et al., 2002       | Prospective<br>cohort                 | 111,526 Caucasian<br>American women:<br>breast cancer cases<br>(n = 711)                                 | Dietary<br>phytoestrogen<br>and genistein   | No significant association was found for phytoestrogen or genistein intake and breast cancer risk  |
| Ingram &<br>Sanders, 1997    | Case control;<br>hospital-based       | Australian: pre- and postmenopausal cases $(n = 144)$ and controls $(n = 144)$                           | Dietary<br>isoflavones (daid-<br>zein, genistein,<br>and equol)   | Highest equol urinary excretion compared to lowest was associated with a 73% reduction in breast cancer risk (AOR = $0.27$ , 95% CI = $0.1-0.69$ , p for trend = $0.009$ )   |
| lwasaki et al.,<br>2008      | Case control;<br>population-<br>based | Japanese: pre- and postmenopausal cases (n = 144) and controls (n = 288)                                 | Dietary genistein,<br>daidzein, and iso-<br>flavones (genistein<br>plus daidzein); se-<br>rum genistein and<br>daidzein | Highest compared to lowest plasma genistein levels were associated with a 66% reduction in breas cancer risk (AOR = $0.34$ , 95% CI = $0.16$ – $0.74$ , p for trend = $0.02$ ).                                    |
| Keinan-Boker<br>et al., 2004 | Case control;<br>population-<br>based | Dutch: pre- and post-<br>menopausal cases<br>(n = 280) and controls<br>(n = 15,555)                      | Phytoestrogens  | No significant association was found between phy toestrogen intake and breast cancer risk.   |
| Key et al.,<br>1999          | Prospective<br>cohort                 | Japanese: pre- and postmenopausal women (N = 11,067)   | Soy food  | No significant association was found between soy intake and breast cancer risk.  |
| Lampe et al.,<br>2007        | Case control;<br>population-<br>based | Chinese: pre- and postmenopausal cases $(n = 196)$ and controls $(n = 1,002)$                            | Plasma genistein<br>and daidzein  | Highest compared to lowest plasma genistein levels were associated with a 74% reduction in breas cancer risk (OR = 0.26, 95% CI = 0.13–0.5, p for trend $< 0.0001$ ).  |
| Lee et al.,<br>2005          | Case control                          | Chinese: pre- and postmenopausal cases (n = 250) and controls (n = 219)                                  | Soy rich foods  | Higher compared to lower soy intake was associated with a 50% reduction in breast cancer risk in women older than 40 years only (OR = $0.5$ , 95% CI = $0.3$ – $1$ ).  |
| Linseisen et<br>al., 2004    | Case control;<br>population-<br>based | German: premenopausal cases (n = 278) and controls (n = 666)   | Dietary daidzein<br>and genistein   | Higher compared to lower genistein intake was associated with a 53% reduction in estrogen and progesterone receptor-positive breast cancer risk (OR = $0.47$ , 95% CI = $0.29$ – $0.74$ , p for trend = $0.002$ ). |
| Murkies et al.,<br>2000      | Case control;<br>hospital-based       | Australian: postmeno-<br>pausal cases (n = 18)<br>and controls (n = 20)                                  | Urine daidzein<br>and genistein   | Controls had significantly higher urinary excretion levels of daidzein than breast cancer cases (p = $0.03$ ).   |
| Nishio et al.,<br>2007       | Prospective<br>cohort                 | 30,454 Japanese<br>women: breast cancer<br>cases (n = 145)   | Soy foods   | No significant association was found between soy food intake and breast cancer risk.   |
| Peterson et al.,<br>2003     | Case control                          | Greek: pre- and post-<br>menopausal cases<br>(n = 820) and controls<br>(n = 1,548)                       | Dietary flavonoids  | No significant association was found between higher compared to lower dietary isoflavone intak and breast cancer risk.   |
| Piller et al.,<br>2006       | Case control;<br>population-<br>based | German: premeno-<br>pausal cases (n = 220)<br>and controls (n = 237)                                     | Dietary and plas-<br>ma genistein   | No significant association was found between plas<br>ma genistein concentration and premenopausal<br>breast cancer risk.   |
|                              |                                       |  |   | (Continued on next page  |

**Table 2. Studies Investigating the Relationship Between Dietary Soy Intake and Breast Cancer Risk in Women (***Continued***)** 

| Study                     | Design   | Sample   | Soy Type   | Findings   |
|---------------------------|--|--|--|--|
| Shannon et<br>al., 2005   | Case control   | Shanghai Chinese: pre-<br>and postmenopausal<br>cases (n = 378) and<br>controls (n = 1,070)                      | Soy food (total<br>unfermented and<br>fermented)   | No significant association was found between soy intake and breast cancer risk.  |
| Thanos et al.,<br>2006    | Case control;<br>population-<br>based                        | Canadian: pre- and postmenopausal cases $(n = 3,024)$ and controls $(n = 3,420)$                                 | Soy foods and dietary isoflavones  | Higher adolescent phytoestrogen and isoflavone intake was associated with a 29% and 19% reduction in breast cancer risk (AOR = 0.71, 95% CI = 0.62–0.82, p for trend < 0.0001; AOR = 0.81, 95% CI = 0.71–0.94, p for trend < 0.01, respectively).  |
| Touillaud et<br>al., 2006 | Prospective<br>cohort  | 26,868 French premenopausal women: breast cancer cases (n = 402)   | Dietary isoflavones  | No significant association was found between iso-<br>flavone intake and premenopausal invasive breast<br>cancer.   |
| Travis et al.,<br>2008    | Prospective cohort   | 37,643 British<br>women: breast cancer<br>cases (n = 585)  | Dietary isoflavones  | No significant association was found between isoflavone intake and breast cancer risk in pre- or postmenopausal women.   |
| Verheus et al.,<br>2007   | Case control;<br>population-<br>based                        | Dutch: pre- and post-<br>menopausal cases<br>(n = 383) and controls<br>(n = 383)                                 | Plasma<br>isoflavones (daid-<br>zein, genistein,<br>glycitein, O-des-<br>methylangolensin,<br>and equol)           | Highest compared to lowest plasma genistein levels were associated with a 32% reduction in breast cancer risk (OR = 0.68, 95% CI = 0.47–0.98, p for trend = 0.07).   |
| Ward et al.,<br>2008      | Case control;<br>population-<br>based                        | 1,189 British pre-<br>and postmenopausal<br>women: breast cancer<br>cases (n = 237)                              | Serum and urine<br>phytoestro-<br>ens (daidzein,<br>genistein glycitein,<br>O-desmethylango-<br>lensin, and equol) | Higher compared to lower total urinary isoflavone levels were associated with an 8% increase in breast cancer risk (OR = 1.08, 95% CI = 1–1.16, p = 0.055); higher compared to lower urinary equol levels were associated with a 7% increase in estrogen receptor site-positive breast cancer risk (OR = 1.07, 95% CI = 1.01–1.12, p = 0.013). |
| Witte et al.,<br>1997     | Case control;<br>familial-<br>matched, pop-<br>ulation-based | American and Canadian Caucasian: premenopausal cases (n = 140) and controls (n = 222)                            | Tofu and soybeans  | Higher compared to lower dietary soy intake was associated with a 50% reduction of premenopausa bilateral breast cancer risk (AOR = $0.5$ , $98\%$ CI = $0.2-1.1$ ).   |
| Wu et al.,<br>2002        | Case control;<br>Population-<br>based                        | Asian American (Chinese, Japanese, and Filipino): pre- and postmenopausal cases (n = 501) and controls (n = 594) | Dietary soy and isoflavones  | High adult and adolescent soy intake compared to low was associated with a 47% reduction in breast cancer risk (OR = 0.53, 95% CI = 0.36–0.78, p for trend = 0.001); high adolescent and low adult soy intake was associated with a 23% reduction in breast cancer risk (OR = 0.77, 95% CI = 0.51–1.16, p for trend = 0.001).                  |
| Wu et al.,<br>2003        | Case control;<br>population-<br>based                        | Asian American: pre-<br>and postmenopausal<br>cases (n = 501) and<br>controls (n = 594)                          | Dietary isoflavones  | High compared to low soy intake during both adolescence and adulthood was associated with a 60% reduction in breast cancer risk (OR = 0.4, 95% CI = 0.24–0.66).  |
| Yamamoto et al., 2003     | Prospective<br>cohort  | 21,852 Japanese<br>women: breast cancer<br>cases (n = 179)   | Soy, miso soup,<br>and dietary<br>isoflavones  | Highest compared to lowest miso soup and dietary isoflavone intake was associated with a 40% and 54% reduction in breast cancer risk (adjusted risk ratio [ARR] = 0.6, 95% CI = 0.34–1.1, p for trend = 0.042; ARR = 0.46, 95% CI = 0.25–0.84 p for trend = 0.043, respectively).  |
| Zheng et al.,<br>1999     | Case control;<br>population-<br>based                        | Chinese: pre- and postmenopausal cases (n = 60) and controls (n = 60)  | Urine daidzen,<br>genistein, glycitein,<br>O-desmethyangol-<br>ensin, and equol                                    | Higher compared to lower urinary excretion of phenols plus total isoflavonoids and glycitein was associated with an 86% reduction in breast cancer risk (AOR = 0.14, 95% CI = 0.02–0.88; AOR = 0.14, 95% CI = 0.03–0.79).  |

## **Findings of No Breast Cancer Risk Reduction**

Twelve studies (13 articles) reported no significant overall reduction of breast cancer risks associated with a higher dietary intake of one or more forms of soy or its components (Bosetti et al., 2005; Cui et al., 2007; Do, Lee, Jung, et al., 2007; Do, Lee, Kim, et al., 2007; Horn-Ross et al., 2001, 2002; Keinan-Boker, van der Schouw, Grobbee, & Peeters, 2004; Key et al., 1999; Nishio et al., 2007; Peterson et al., 2003; Shannon et al., 2005; Touillaud, Thiebaut, Niravong, Boutron-Ruault, & Clavel-Chapelon, 2006; Travis et al., 2008). No significant association was found between plasma genistein levels and breast cancer risks (Piller, Chang-Claude, & Linseisen, 2006), indirectly suggesting no difference in breast cancer risk related to dietary isoflavone intake. Nor was a significant association between urinary genistein levels and breast cancer risks found (den Tonkelaar et al., 2001), again indirectly suggesting no difference in breast cancer risk related to dietary isoflavone intake.

# **Findings of Increased Breast Cancer Risk**

Two studies reported an association between exposure to isoflavones and increased breast cancer risk. Serum levels of the specific isoflavone diadzein and of its metabolite equol in serum (Grace et al., 2004) and urine (Grace et al.; Ward et al., 2008) were significantly associated with an increased risk of developing breast cancer. Additionally, urine equol was significantly associated with estrogen receptor site-positive breast cancer risk in premenopausal and perimenopausal women only (Ward et al.).

# **Findings by Nationality**

North American: Two studies of Caucasian Americans and non-Asian Americans found no significant relationship between dietary phytoestrogens or phytoestrogens and genistein and breast cancer risks (Horn-Ross et al., 2001, 2002). Two studies of Asian Americans found decreased breast cancer risks associated with higher dietary soy or isoflavone intake (Wu et al., 2002, 2003). One study of Caucasian Americans and Canadians found decreased breast cancer risks associated with increased dietary tofu intake (Witte et al., 1997), and two studies of Canadians found decreased breast cancer risks associated with infant soy formula use or higher dietary isoflavone intake (Boucher et al., 2008; Thanos et al., 2006).

**European:** One British study reported no relationship with breast cancer risk (Travis et al., 2008), two reported contradictory increased breast cancer risks for dietary isoflavone intake and urinary equol excretion (Grace et al., 2004; Ward et al., 2008), and one of South Asians living in England reported a decreased breast cancer risk associated with higher dietary isoflavone intake (Dos Santos Silva et al., 2004). One French study found decreased breast cancer risks and higher dietary isoflavone intake (Touillaud et al., 2006). Two German studies found no relationship be-

tween dietary genistein intake or plasma genistein levels and breast cancer risks (Linseisen et al., 2004; Piller et al., 2006). Two Dutch studies found no relationship between dietary phytoestrogen intake or urinary genistein excretion and breast cancer risks (den Tonkelaar et al., 2001; Keinan-Bokar et al., 2004), although one study found decreased breast cancer risks associated with higher plasma genistein levels (Verheus et al., 2007). One Italian (Bosetti et al., 2005) and one Greek study (Peterson et al., 2003) found no relationship between dietary phytoestrogen or isoflavone intake and breast cancer risks.

Asian: One South Korean study (two journal articles) found no relationship between dietary soy intake and breast cancer risks (Do, Lee, Kim, et al., 2007; Do, Lee, Jung, et al., 2007). Two Chinese studies found no relationship between dietary vegetable-soy patterns or soy intake (Cui et al., 2007; Shannon et al., 2005), although four Chinese studies (five journal articles) found decreased breast cancer risks associated with higher dietary soy or isoflavone intake and higher plasma genistein levels (Dai et al., 2001; Shu et al., 2001; Lampe et al., 2007; Lee et al., 2005; Zheng et al., 1999). Two Japanese studies found no relationship between dietary soy, phytoestrogen, or genistein intake (Nishio et al., 2007), although four Japanese studies found decreased breast cancer risks associated with dietary tofu, miso soup or isoflavone intake, and higher plasma genistein levels (Hirose et al., 2003; 2005; Iwasaki et al., 2008; Yamamoto et al., 2003).

Lastly, two Australian studies found decreased breast cancer risks associated with urinary daidzein and equal excretion levels. The ethnicity of participants were not described (Ingram & Sanders, 1997; Murkies et al., 2000).

In summary, the sample nationality most consistently reporting a decrease in the risk for developing breast cancer was Chinese, and it should be noted that Asian Americans and South Asians living in England also reported decreased breast cancer risks.

#### **Limitations**

Limitations to these studies are related to difficulties in accurate quantification of soy intake and inadequate knowledge of the bioavailability, interactions, dose response, and metabolism and excretion of soy and isoflavones. Unmeasured sources of soy intake and unidentified bioactive substances also may have confounded the findings. Because different forms, methods, and measures of soy were used in the studies, only general conclusions can be offered.

### **Conclusions**

This review of studies on the dietary intake of soy or soy products suggests a protective or no relationship to breast cancer risk in women, although two studies suggest an associated increased risk for breast cancer development. Clearly, nurses should become more knowledgeable of

#### What is soy?

- Soy refers to soybeans, an Asian plant used in many foods.
- Soy is a major source of plant estrogens (phytoestrogens).
- Isoflavones are one type of plant estrogen, and include daidzein and genistein.

#### What foods commonly contain soy?

 Soy is most common in Asian diets as tofu, miso, natto, tempeh, and edamame. Some foods gaining in popularity in Western culture include tofu yogurt, soy milk, cheese, and meat substitutes. Foods also may be fortified with soy.

#### How might dietary soy be related to the risk of breast cancer?

- Soy or its components might block estrogen and exert a protective effect.
- Soy or its components might act like estrogen and stimulate the growth of some types of cancer.
- Soy or its components may act as antioxidants, and protect cells from damage which may lead to cancer development.

# What have researchers learned about dietary soy intake and related breast cancer risk?

- About 50% of studies found a protective effect for breast cancer.
- About 50% of studies found no protective effect for breast cancer.
- Two studies have found an increased risk of breast cancer.
- The age at which soy is eaten may influence breast cancer risk.
- Soy intake may actually represent a healthy lifestyle overall.

#### Recommendations for women regarding soy intake and breast cancer development:

- Dietary soy supplements and medications are not considered "drugs" and are not regulated by the U.S. Food and Drug Administration.
- Consuming foods which naturally contain soy is probably safer than consuming foods fortified with concentrated amounts of soy additives.
- The safety of consuming dietary supplements or medications with high doses of soy or soy components (isoflavones, daidzein, or genistein) is unknown and is questionable for women who have had or are at high risk of developing breast cancer.

# Figure 1. What Patients Should Know About Soy and Breast Cancer Risk

Note. Based on information from Linus Pauling Institute, 2008; MedlinePlus, 2008; Messina, 2008.

soy foods and products that are less familiar in Western culture and more familiar with alternative therapies such as soy-based supplements. Questions on the intake of soy foods and over-the-counter soy-based products should be included when obtaining patient nutrition histories. Nurses caring for women who are not at a high

risk for developing breast cancer may include general information about the possible breast cancer-protective properties of routine dietary soy intake in client education (see Figure 1), although no certain recommendations can be made based on the current state of the science. In the meantime, nurses caring for women at high risk for developing breast cancer or those with a previous diagnosis should collaborate closely with dieticians regarding the most current practice recommendations and closely monitor emerging scientific releases regarding dietary soy. Women at high risk for developing breast cancer or who have breast cancer histories should be told to avoid initiating high dietary intake or supplementation with soy or its components until their possible cancer-related interactions are better understood.

Future research recommendations should include the study of dietary soy intake and breast cancer risks in particular age groups as well as racial or ethnic groups, such as African American and Hispanic women. Given the complexity of the possible soy-breast cancer relationship, an interdisciplinary approach and uniformity of methods will strengthen future research findings. Consistent and comprehensive assessment of both soy food and component intake, as well as urinary excretion of these substances, is essential.

Carol A. Enderlin, PhD, RN, is an assistant clinical professor in the Department of Nursing Education in the College of Nursing; Elizabeth Ann Coleman, PhD, RNP, AOCN®, is a professor and the Elizabeth Cooper-Stanley Endowed Oncology Chair in the Department of Nursing Science in the College of Nursing and Department of Internal Medicine in the College of Medicine; Carol Beth Stewart, BS, is a program specialist in the College of Nursing; and Reza Hakkak, PhD, is a professor and chairman of the Department of Dietetics and Nutrition in the College of Health Related Professions and a professor in the Department of Pediatrics in the College of Medicine, all at the University of Arkansas for Medical Sciences in Little Rock. This project was supported, in part, by the John A. Hartford Foundation Building Academic Geriatric Capacity Award Program. The views expressed in this article are those of the authors and do not necessarily represent the views of the John A. Hartford Foundation. Enderlin can be reached at caenderlin@uams.edu, with copy to editor at ONFEditor@ons.org. (Submitted June 2008. Accepted for publication December 8, 2008.)

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# References

Adlercreutz, H. (1995). Phytoestrogens: Role in cancer protection. *Environmental Health Perspectives*, 103(Suppl. 7), 103–112.

American Cancer Society. (2009). What causes cancer? Retrieved July 24, 2009, from http://www.cancer.org/docroot/CRI/content/CRI\_2\_2\_2X\_What\_causes\_breast\_cancer\_5.asp?sitearea=

Barnes, S., Boersma, B., Patel, R., Kirk, M., Darley-Usmar, V.M., Kim, H., et al. (2000). Isoflavonoids and chronic disease: Mechanisms of action. *Biofactors*, 12(1–4), 209–215.

Biology Online. (2005). Soy. Retrieved August 26, 2008, from http://www.biology-online.org/dictionary/Soy

Bosetti, C., Spertini, L., Parpinel, M., Gnagnarella, P., Lagiou, P., Negri, E., et al. (2005). Flavonoids and breast cancer risk in Italy. Cancer Epidemiology Biomarkers and Prevention, 14(4), 805–808. Boucher, B.A., Cotterchio, M., Krieger, N., & Thompson, L.U. (2008). Soy formula and breast cancer risk. *Epidemiology*, *19*(1), 165–166

Cooper, H. (1998). Synthesizing research: A guide for literature reviews. Thousand Oaks, CA: Sage.

Cui, X., Dai, Q., Tseng, M., Shu, X.O., Gao, Y.T., & Zheng, W. (2007). Dietary patterns and breast cancer risk in the Shanghai breast cancer study. Cancer Epidemiology Biomarkers and Prevention, 16(7), 1443–1448.

Dai, Q., Shu, X.O., Jin, F., Potter, J.D., Kushi, L.H., Teas, J., et al. (2001). Population-based case-control study of soy food intake and breast cancer risk in Shanghai. *British Journal of Cancer*, 85(3), 372–378.

den Tonkelaar, I., Keinan-Boker, L., Veer, P.V., Arts, C.J., Adlercreutz, H., Thijssen, J.H., et al. (2001). Urinary phytoestrogens and post-

- menopausal breast cancer risk. Cancer Epidemiology Biomarkers and Prevention, 10(3), 223–228.
- Do, M.H., Lee, S.S., Jung, P.J., & Lee, M.H. (2007). Intake of fruits, vegetables, and soy foods in relation to breast cancer risk in Korean women: A case-control study. *Nutrition and Cancer*, 57(1), 20–27.
- Do, M.H., Lee, S.S., Kim, J.Y., Jung, P.J., & Lee, M.H. (2007). Fruits, vegetables, soy foods, and breast cancer in pre- and postmenopausal Korean women: A case-control study. *International Journal of Vitamin* and Nutrition Research, 77(2), 130–141.
- Dos Santos Silva, I., Mangtani, P., McCormack, V., Bhakta, D., Mc-Michael, A.J., & Sevak, L. (2004). Phytoestrogen intake and breast cancer risk in South Asian women in England. *Cancer Causes and Control*, 15(8), 805–818.
- Grace, P.B., Taylor, J.I., Low, Y.L., Luben, R.N., Mulligan, A.A., Botting, N.P., et al. (2004). Phytoestrogen concentrations in serum and spot urine as biomarkers for dietary phytoestrogen intake and their relation to breast cancer risk in a European prospective investigation. *Cancer Epidemiology Biomarkers and Prevention*, 13(5), 698–708.
- Higdon, J. (2007). An evidence-based approach to dietary phytochemicals. New York: Thieme Medical.
- Hirose, K., Imaeda, N., Tokudome, Y., Goto, C., Wakai, K., Matsuo, K., et al. (2005). Soybean products and reduction of breast cancer risk: A case-control study in Japan. *British Journal of Cancer*, 93(1), 15–22.
- Hirose, K., Takezaki, T., Hamajima, N., Miura, S., & Tajima, K. (2003). Dietary factors protective against breast cancer in Japanese premenopausal and postmenopausal women. *International Journal of Cancer*, 107(2), 276–282.
- Horn-Ross, P.L., Hoggatt, K.J., West, D.W., Krone, M.R., Stewart, S.L., Anton, H., et al. (2002). Recent diet and breast cancer risk: The California Teachers Study (USA). Cancer Causes and Control, 13(5), 407–415.
- Horn-Ross, P.L., John, E.M., Lee, M., Stewart, S.L., Koo, J., Sakoda, L.C., et al. (2001). Phytoestrogen consumption and breast cancer risk in a multiethnic population: The Bay Area Breast Cancer Study. *American Journal of Epidemiology*, 154(5), 434–441.
- Ingram, D., Sanders, K., Kolybaba, M., & Lopez, D. (1997). Case-control study of phytoestrogens. *Lancet*, 350(9083), 990–994.
- Iwasaki, M., Inoue, M., Otani, T., Sasazuki, S., Kurahashi, N., Miura, T., et al. (2008). Plasma isoflavone level and subsequent risk of breast cancer among Japanese women: A nested case-control study from the Japan Public Health Center-based prospective study group. *Journal of Clinical Oncology*, 26(10), 1677–1683.
- Keinan-Boker, L., van der Schouw, Y.T., Grobbee, D.E., & Peeters, P.H. (2004). Dietary phytoestrogens and breast cancer risk. American Journal of Clinical Nutrition, 79(2), 282–288.
- Kelly, G.E., Nelson, C., Waring, M.A., Joannou, G.E., & Reeder, A.Y. (1993). Metabolites of dietary (soya) isoflavones in human urine. Clinica Chimoca Acta, 223(1–2), 9–22.
- Key, T.J., Sharp, G.B., Appleby, P.N., Beral, V., Goodman, M.T., Soda, M., et al. (1999). Soya foods and breast cancer risk: A prospective study. *British Journal of Cancer*, 81(7), 1248–1256.
- Lampe, J., Nishino, Y., Ray, R., Wu, C., Li, W., Lin, M.G., et al. (2007). Breast cancer among women in Shanghai, China. Cancer Epidemiology, Biomarkers and Preventions, 16(12), 2579–2586.
- Lee, M.M., Chang, I.Y., Horng, C.F., Chang, J.S., Cheng, S.H., & Huang, A. (2005). Breast cancer and dietary factors in Taiwanese women. Cancer Causes and Control, 16(8), 929–937.
- Linseisen, J., Piller, R., Hermann, S., & Chang-Claude, J. (2004). Dietary phytoestrogen intake and premenopausal breast cancer risk in a case-control study. *International Journal of Cancer*, 110(2), 284–290.
- Linus Pauling Institute. (2008). *Phytochemicals*. Retrieved September 18, 2008, from http://lpi.oregonstate.edu/infocenter/phytochemicals.html
- Medicinenet.com. (2004). Medterms dictionary: Definition of equol. Retrieved October 14, 2008, from http://www.medterms.com/script/main/art.asp?articlekey=32272
- MedlinePlus. (2008). Soy. Retrieved October 23, 2008, from http://www.nlm.nih.gov/medlineplus/druginfo/natural/patient-soy.html
- Messina, M. (2008). Soy and health: Isoflavones. Retrieved September 22, 2008, from http://www.soyconnection.com/health\_nutrition/pdf/ IsofavonesFactSheet.pdf

- Murkies, A., Dalais, F.S., Briganti, E.M., Burger, H.G., Healy, D.L., Wahl-qvist, M.L., et al. (2000). Phytoestrogens and breast cancer in postmenopausal women: A case control study. *Menopause*, 7(5), 289–296.
- National Cancer Institute. (2004). *Antioxidants and cancer prevention:* Fact sheet. Retrieved November 18, 2008, from http://www.nci.nih.gov/cancertopics/factsheet/antioxidantsprevention
- National Cancer Institute. (2009). SEER stat fact sheets. Retrieved July 24, 2009, from http://seer.cancer.gov/statfacts/html/breast.html
- Nishio, K., Niwa, Y., Toyoshima, H., Tamakoshi, K., Kondo, T., Yatsuya, H., et al. (2007). Consumption of soy foods and the risk of breast cancer: Findings from the Japan Collaborative Cohort (JACC) Study. Cancer Causes and Control, 18(8), 801–808.
- Peeters, P.H., Keinan-Boker, L., van der Schouw, Y.T., & Grobbee, D.E. (2003). Phytoestrogens and breast cancer risk. *Breast Cancer Research and Treatment*, 77(2), 171–183.
- Peterson, J., Lagiou, P., Samoli, E., Lagiou, A., Katsouyanni, K., La Vecchia, C., et al. (2003). Flavanoid intake and breast cancer risk: A case-control study in Greece. *British Journal of Cancer*, 89(7), 1255–1259.
- Piller, R., Chang-Claude, J., & Linseisen, J. (2006). Plasma enterolactone and genistein and the risk of premenopausal breast cancer. *European Journal of Cancer Prevention*, 15(3), 225–232.
- Setchell, K.D., Brown, N.M., & Lydeking-Olsen, E. (2002). The clinical importance of the metabolite equol—A clue to the effectiveness of soy and its isoflavones. *Journal of Nutrition*, 132(12), 3577–3584.
- Shannon, J., Ray, R., Wu, C., Nelson, Z., Gao, D.L., Li, W., et al. (2005). Food and botanical groupings and risk of breast cancer. *Cancer Epidemiology Biomarkers and Prevention*, 14(1), 81–90.
- Shu, X.O., Jin, F., Dai, Q., Wen, W., Potter, J.D., Kushi, L.H., et al. (2001). Soy food intake during adolescence and subsequent risk of breast cancer among Chinese women. *Cancer Epidemiology Biomarkers and Prevention*, 10(5), 483–488.
- Stock, W.A. (1994). Systematic coding for research synthesis. In C.H. & L.V. Hedges (Eds.), The handbook of research synthesis (pp. 125–138). New York: Russell Sage Foundation.
- Thanos, J., Cotterchio, M., Boucher, B.A., Kreiger, N., & Thompson, L.U. (2006). Adolescent dietary phytoestrogen intake and breast cancer risk (Canada). *Cancer Causes and Control*, 17(10), 1253–1261.
- Touillaud, M.S., Thiebaut, A.C., Niravong, M., Boutron-Ruault, M.C., & Clavel-Chapelon, F. (2006). No association between dietary phytoestrogens and risk of premenopausal breast cancer in a French cohort study. Cancer Epidemiology Biomarkers and Prevention, 15(12), 2574–2576.
- Travis, R.C., Allen, N.E., Appleby, P.N., Spencer, E.A., Roddam, A.W., & Key, T.J. (2008). A prospective study of vegetarianism and iso-flavone intake in relation to breast cancer risk in British women. *International Journal of Cancer*, 122(3), 705–710.
- Verheus, M., van Gils, C.H., Keinan-Boker, L., Grace, P.B., Bingham, S.A., & Peeters, P.H. (2007). Plasma phytoestrogens and subsequent breast cancer risk. *Journal of Clinical Oncology*, 25(6), 648–655.
- Wang, L.Q. (2002). Mammalian phytoestrogens: Enterodiol and enterolactone. Journal of the Chromatography B: Analytical Technologies in the Biomedical Life Sciences, 777(1–2), 289–309.
- Ward, H., Chapelais, G., Kuhnle, G.G., Luben, R., Khaw, K.T., & Bingham, S. (2008). Breast cancer risk in relation to urinary and serum biomarkers of phytoestrogen exposure in the European Prospective into Cancer-Norfolk cohort study. *Breast Cancer Research*, 10(2), R32.
- Witte, J.S., Ursin, G., Siemiatycki, J., Thompson, W.D., Paganini-Hill, A., & Haile, R.W. (1997). Diet and premenopausal bilateral breast cancer. Breast Cancer Research and Treatment, 42(3), 243–251.
- Wu, A.H., Wan, P., Hankin, J., Tseng, C.C., Yu, M.C., & Pike, M.C. (2002). Adolescent and adult soy intake and risk of breast cancer in Asian Americans. *Carcinogenesis*, 23(9), 1491–1496.
- Wu, A.H., Yu, M.C., Tseng, C.C., Hankin, J., & Pike, M.C. (2003). Green tea and risk of breast cancer in Asian Americans. *International Jour*nal of Cancer, 106(4), 574–579.
- Yamamoto, S., Sobue, T., Kobayashi, M., Sasaki, S., & Tsugane, S. (2003). Soy, isoflavones, and breast cancer risk in Japan. *Journal of the National Cancer Institute*, 95(12), 906–913.
- Zheng, W., Dai, Q., Custer, L.J., Shu, X.O., Wen, W.Q., Jin, F., et al. (1999). Urinary excretion of isoflavonoids and the risk of breast cancer. *Cancer Epidemiology Biomarkers and Prevention*, 8(1), 35–40.