Breast Cancer Chemoprevention: A Review of Selective Estrogen Receptor Modulators

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In the United States, breast cancer is the most common cancer diagnosis among women (American Cancer Society, 2005). The risk for breast cancer development increases with age, with the highest incidence occurring from age 75–79 (American Cancer Society, 2003). An estimated 211,240 cases of invasive breast cancer and 58,490 cases of in situ breast cancer will be diagnosed in 2005, and an estimated 40,410 deaths from breast cancer are projected in women in 2005 (American Cancer Society, 2005).

Breast cancer chemoprevention plays an important role for women at high risk for development of the disease. Chemoprevention is the delay or reversal of cancer by the use of pharmacologic agents that inhibit its development (Torrisi, Decensi, Formelli, Camerini, & De Palo, 2001). Selective estrogen receptor modulators (SERMs) have antiestrogen and estrogen-like effects. Tamoxifen and raloxifene are first- and second-generation SERMs, respectively. These SERMs have improved side-effect profiles is needed. Oncology nurses play a pivotal role in helping patients to understand the current status of breast cancer prevention as well as the future direction of research.

Breast Cancer Chemoprevention Clinical Trials

In 2001, the National Cancer Institute (NCI) Prevention Cancer Task Force reviewed the current status of breast cancer prevention as well as the future direction of research. The NCI Prevention Cancer Task Force (2001) reported that the use of selective estrogen receptor modulators (SERMs) tamoxifen and raloxifene has demonstrated mixed results in breast cancer prevention clinical trials. In the United States, tamoxifen is approved for use for breast cancer prevention in women at high risk. The continued development of SERMs with improved side-effect profiles is needed. Oncology nurses play a pivotal role in helping patients to understand the current status of breast cancer prevention as well as the future direction of research.

The Physiology of Estrogen and Effects on Breast Tissue

Estrogens are steroid hormones that are produced in the ovaries, adrenal glands, and placenta during pregnancy. The hypothalamus secretes gonadotropin-releasing hormone (GnRH), which stimulates the anterior pituitary to release follicle-stimulating hormone (FSH) and luteinizing hormone (LH). FSH and LH act on the ovaries to produce estrogen in the form of estradiol and estrone. These estrogens bind with estrogen receptors in target tissues of the breast, uterus, brain, bone, liver, and heart. Estrogen receptors are molecules inside the cell to which only estrogens or closely related molecules can bind; however, not all cells contain receptors specific for estrogen. Once an estrogen has combined with a receptor in a cell, the shape of the receptor changes to combine with the DNA sites inside the target tissue. The DNA stimulates gene activation and production of RNA, which, in turn, stimulates the synthesis of protein. This protein produces changes in the cell according to tissue type and underlying conditions. The cycle is completed when high levels of estrogen in the blood send negative feedback to the hypothalamus to suppress the release of GnRH (Huether & McCance, 2004).

Estrogen released from the ovaries during adolescence stimulates the growth of the ductal system in breast tissue, which is not developed fully until pregnancy in preparation for lactation (Varney, Kriebes, & Gegg, 2004). Prolonged exposure to high levels of estrogen may cause changes in cell proliferation from normal growth to hyperplasia to neoplasia with DNA mutations and uncontrolled cell proliferation (Clemens & Goss, 2001) (see Figure 1). Women may be at increased risk for breast cancer from prolonged exposure to estrogen such as with early menarche before age 12, first full-term pregnancy after age 30, late menopause after age 55, or recent long-term use of postmenopausal hormone replacement therapy (HRT) (American Cancer Society, 2003; Clemens & Goss; Fabian & Kimler, 2002).

Most cases of breast cancer develop from the ductal epithelium and may be classified as invasive or noninvasive. Invasive breast cancers have broken through the basement membrane of the ducts or lobules into the fatty tissue of the breast (Huether & McCance, 2004). About 70%–80% of all breast cancers are termed estrogen receptor positive (ER+) if they express the ER protein α. These ER+ breast cancers are more likely to respond to hormonal therapy such as SERMs. Breast cancers that do not express the ER protein α are termed estrogen receptor negative (ER−). These ER− breast cancers are less likely to respond to hormonal therapy such as SERMs.