Breast Cancer in a Male with BRCA2 Mutation: Is Screening Mammography Necessary in Men?

Jessica L. Buicko*, Michael A. Lopez and Beth Ann Lesnikoski
Department of General Surgery, University of Miami-Regional Medical Campus, USA

Abstract
We present the case of a 76 year-old male with BRCA2 and a personal history of left-sided breast cancer who presented 20 years later with right-sided breast cancer found on screening mammography. The role of screening mammography and its relevance to males with breast cancer is poorly understood. There are currently no screening guidelines for male breast cancer. Further research is needed to determine the benefit of screening males at higher risk of developing breast cancer.

Case Presentation
A 76 year-old male presented to our surgical clinic for evaluation. He has a known BRCA mutation and a personal history of left breast cancer, treated with a left sided modified radical mastectomy 20 years prior. He was undergoing intermittent mammography for surveillance. He denied ever feeling a palpable mass or experiencing any nipple discharge. Recent screening mammography—after 4 year interval, showed a nodule in the right retroareolar region, which was 1.2 x 1.3cm in size. He underwent diagnostic workup with spot compression mammography views and ultrasound. Ultrasound showed an irregularly marginated 8 x 5 x 7mm hypoechoic lesion, with a surrounding echogenic region measuring 1.1 x 0.7cm. Ultrasound guided biopsy revealed intermediate grade invasive ductal carcinoma. He subsequently underwent a right total mastectomy with sentinel lymph node biopsy. Final pathology confirmed an estrogen receptor positive, progesterone receptor positive, Her-2-neu non-amplified, 2.0cm invasive ductal carcinoma. Sentinel lymph node was negative for carcinoma.

Discussion
Male breast cancer accounts for less that 1% of cancers diagnosed in the United States. The female-to-male ratio for breast cancer is 100:1 [5]. Approximately 90% of male breast cancers are invasive ductal carcinomas. The male breast lacks terminal lobules, unless exposed to high doses of endogenous or exogenous estrogens. Lobular type accounts for less than 1.5% of male disease, whereas it accounts for greater than 10% in women [6].

It has been found that most male breast cancers express high levels of hormone receptors. The estrogen and progesterone receptors are more likely to be positive in male patients than in female patients. The proportion of hormone-receptor-expressing tumors increases with age, comparable to post-menopausal women. The proto-oncogene HER2/neu is expressed in only 5% of male patients, and in approximately 15% of female patients [6].

There have been many documented risk factors related to male breast cancer. Some of the more common include Klinefelter syndrome, BRCA1 and BRCA2, a family history in a first degree male or female relative, hyperestrogenism, advanced age, and a history of chest radiation [5]. In contrast to women with breast cancer, white race is not a risk factor in males [7].

Germ line mutations in the high penetrance BRCA2, and with a lower frequency BRCA1, have been shown to predispose men to primary breast cancer. Men with a high-risk family history have shown a 60-76% incidence of BRCA2 mutation and a 10-16% BRCA1 mutation [6]. The median age of male breast cancer in BRCA2 positive men is 58.8, and is 69.7 years in those without the mutation. The frequency of mutation of these genes is highly variable. In both men and women, and prevalence has been shown to depend on mainly on ethnicity and country. In Iceland, BRCA2 mutations have been seen in 40% of male breast cancer patients. In Ashkenazi Jews, BRCA1 and BRCA2 mutations are essentially equal between men and women. Family history of BRCA2...
mutations carries the highest likelihood of both male breast cancer and prostate cancer, and warrants genetic testing for male family members [8,9]. Male patients with a BRCA2 mutation have an 8% estimated lifetime risk of developing breast cancer [10]. This is comparable to subgroups of women in the US who are routinely encouraged to undergo mammography screening, such as African American, Latina and Asian women. In total, 4-14% of male patients with breast cancer not selected on basis of family history will have a BRCA2 mutation [10]. In regards to other genetic risk factors, there is mixed data in regards to the association of PALB2, CYP17, and CHEK2 in their risk of male breast cancer [7]. In female breast cancer, it has been established that there is an association of leisure-time physical activity (LTPA) with risk reductions of about 25% among post-menopausal normal weight women compared with no association for obese women [11]. There is some evidence in small studies that exercise reduces the risk of breast cancer in males but this has not been corroborated by larger studies [12]. Arem et al. [11] evaluated the association of physical activity and risk of male breast cancer and found that compared to low-LTPA, medium, and high LTPA were not associated with male breast cancer risk (OR 1.01; 95% CI, 0.79-1.29; 0.90, 0.69-1.18, respectively). There is no current agreement regarding what subgroup of men would benefit from screening mammography. Male breast cancers are often detected at a late stage, commonly when a mass is palpable or nipple discharge is present [1]. Multiple reasons exist for the delay in diagnosis. Factors may include lack of screening in high-risk men, poor education on male breast cancer, and decreased breast tissue in males, which may lead to early lymphatic spread [1,13]. Historically, compared to female breast cancer patients, there is approximately a 22-month delay between onset of symptoms and diagnosis, which has a dramatic impact on treatment and survival [3]. Women, specifically those who are BRCA 1 and BRCA 2 positive, are more likely to be diagnosed at an earlier stage because of guidelines that dictates screening protocols [1]. For women who are BRCA positive, the National Comprehensive Cancer Network (NCCN) recommends yearly breast MRIs between 25 and 29 years old and then adding yearly mammograms starting at age 30. In regards to males at high risk, the NCCN recommends considering mammogram at age 40 [14]. Brenner et al proposed that any male patient with a BRCA mutation should undergo screening mammography [4]. Arguments against intensive screening included false-positive results, unnecessary imaging, and unneeded surgery. In a Dutch multicenter screening study for women with hereditary breast cancer, the false positive rate of mammography and MRI was 5.5 vs. 14.0% (P<.001) [15]. No such data exists for men.

Conclusion
At this time there are no specific screening guidelines for male breast cancer. Although there is convincing evidence for an increased risk of cancer associated with certain gene mutations, specifically BRCA2, no recommendations have been made to detect disease in its early stages. This has unfortunately led to the late diagnosis of cancer in men that may have been managed before its progression to a terminal state. A thorough history can prompt genetic screening, which should encourage early screening of the patient, and both male and female relatives. Further data is needed to determine if men at increased risk, for example those with Klinefelter syndrome or history of chest radiation, would benefit from screening.

References


