

Radiation exposure from diagnostic procedures in patients with newly diagnosed breast cancer

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Radiation exposure is associated with an increased risk of secondary cancers. Knowing the approximate radiation exposure from diagnostic procedures in the first year after a breast cancer diagnosis could help educate patients and allow physicians to monitor them more closely for potential risks.

The use of diagnostic imaging and radiation-based therapies has increased rapidly over the past decade, particularly in oncology.¹ Increased exposure to medical radiation has raised concerns about health risks and in particular, an increased risk of secondary cancers.² The probability of occurrence of the chronic effects of ionizing radiation is a function of the total radiation dose.^{3,4} There are limited data on radiation exposure from diagnostic procedures (DPs) in breast cancer patients. Although patients who have been diagnosed with breast cancer undergo radiation-based diagnostic and staging work-up that is deemed medically necessary, there remains a growing concern about the possible long-term side effects of radiation exposure. Breast cancer is the most common cancer among women worldwide, and the number of breast cancer survivors has increased notably in the past decade. In the United States, as of 2012, nearly 3 million women have a history of breast cancer and constitute more than 40% of female cancer survivors.⁴ The objective of our study was to quantify the radiation dose experienced by breast cancer patients during the first year of their diagnosis. Patients with early-stage breast cancer have a good overall prognosis and survival, hence it is important to be vigilant about possible long-term side effects associated with radiation exposure in such patients. There are currently no additional screening guidelines for cancer survi-

vors to detect secondary cancers, however knowing the approximate radiation exposure and associated potential risks could help educate patients and allow physicians to monitor them more closely.

Methods

We performed a retrospective chart review study to analyze the cumulative radiation doses (millisievert [mSv]) of routine DPs performed during the first year after breast cancer diagnosis. In all, 305 patients with newly diagnosed breast cancer who were treated at West Virginia University during January 2008–October 2010 were included in our study. Data regarding the frequency of DP including mammograms, sentinel lymph node biopsies, X-rays, computed-tomographic (CT) and/or positron emission tomographic (PET) scans, MUGA scans and bone scans were collected. Mean radiation doses of the DPs were obtained from the departments of nuclear medicine and radiation at our institution (Table 1). We used Kruskal-Wallis test and post hoc pairwise comparisons to assess the influence of age, histology, estrogen/progesterone receptor (ER/PR) status, and disease stage on the amount of radiation exposure. We also performed a subgroup analysis to evaluate the effect of those factors on the number of positron emission tomography-computed tomography (PET-CT) scans. All statistical tests with 2-sided $P < .05$ were considered statistically significant.

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Results

Most of our patients were aged 40–60 years ($n = 162$) and most had infiltrating ductal carcinoma (IDC; $n = 198$; Table 2). Patients who were younger than 40 years received

TABLE 1 Radiation doses from various diagnostic procedures

Procedure	Mean dose, mSV
Mammogram (bilateral)	0.4
Stereotactic biopsy	0.2
Sentinel LN biopsy	0.4
Chest X-ray	0.1
PET-CT	24
MUGA scan	9.4
CT	
Brain	2.0
Neck	6.0
Chest	7.0
Chest/abd/pelvis	18.0
Abd/pelvis	10.0
Pelvis	6.0
Bone scan	6.3

Abd, abdomen; CT, computed tomography; LN, lymph node; MUGA, multigated acquisition scan; PET-CT, positron emission tomography-computed tomography

a significantly higher radiation dose compared with those who were older than 60 years (35.9 mSv and 19.2 mSv, respectively; $P = .009$). Patients with ductal carcinoma in situ (DCIS) had significantly less exposure to radiation than did patients with IDC and invasive lobular carcinoma (ILC): 8.5 mSv, 26.7 mSv, and 22.4 mSv, respectively; $P < .0001$. Disease stage IIB or higher was associated with a significantly higher radiation exposure ($P < .0001$). Disease stage IIIA or higher was the only factor associated with a higher radiation dose from PET-CTs ($P < .0001$). The mean radiation exposure relative to various factors is shown in Table 2.

Conclusion

Most of the evidence on radiation-induced health risks is based on data from survivors of the Hiroshima and Nagasaki atomic bomb blasts.⁵ These data provide compelling evidence of radiation-induced cancer risk at doses higher than 100 mSv.³ Radiation-induced risk is more controversial at doses between 10 and 100 mSv because estimating the risk of developing cancer from an individual radiologic study is not clear, given the stochastic nature of harm from ionizing radiation and the limited epidemiologic data. However, there is mounting consensus in the medical and scientific communities that the risk to the patient is real, however small.

The 2006 Biological Effects of Ionizing Radiation (BEIR VII-Phase 2) risk model on lifetime attributable cancer predicts that 1 in 1,000 persons exposed to 10 mSv will develop cancer because of that single exposure.³ The International Commission on Radiological Protection has confirmed that doses for diagnostic exams such as CT scans approach or exceed levels associated with an increase in lifetime cancer risk.⁶

Our study has attempted to quantify the radiation doses associated with diagnostic procedures in breast cancer patients. Although our patients underwent diagnostic procedures and imaging for necessary diagnosis and management of breast cancer, given the high overall survivorship in breast cancer patients, it is imperative to educate both patients and physicians about the possible long-term health risks associated with radiation exposure, including the development of secondary cancers. There are no current recommendations to offer additional screenings to cancer survivors except the general age-appropriate screenings, however knowing the approximate radiation dose experienced by these patients and potential risks associated with it will help patients and physicians monitor closely with details history and physical examinations.

The findings from our study show that radiation exposure from DPs is significant in the first year after a diagnosis of breast cancer, especially for younger patients and those with advanced stage disease, even though the potential risk

TABLE 2 Mean and median of the total radiation doses by groups

Group	No. of patients	Radiation dose, mSV		P
		Mean (SE)	Median (IQR)	
Overall	305	21.94 (1.51)	7.80 (32.10)	—
Age, y				.0187
< 40	19	35.94 (8.65)	25.10 (25.50)	—
40-60	162	22.53 (2.00)	10.30 (33.60)	—
> 60	123	19.18 (2.27)	6.80 (28.90)	—
Histology				< .0001
DCIS	60	8.54 (2.19)	2.10 (5.15)	—
IDC	198	26.69 (2.04)	24.90 (35.60)	—
ILC	26	22.38 (3.43)	25.40 (24.00)	—
Others	13	20.28 (6.52)	19.00 (23.80)	—
ER/PR				.0052
Negative	54	31.79 (4.08)	26.35 (32.90)	—
Positive	224	21.10 (1.69)	7.30 (32.25)	—
Stage				< .0001
0	18	7.62 (4.60)	2.25 (4.30)	—
I	105	13.28 (1.79)	5.80 (22.90)	—
II/IIA	48	24.73 (3.57)	25.25 (29.75)	—
IIB	25	36.03 (3.09)	34.40 (14.30)	—
III/A-C	35	42.04 (4.88)	35.60 (24.90)	—
IV	12	76.68 (7.92)	72.80 (27.30)	—

DCIS, ductal carcinoma in situ; ER/PR, estrogen/progesterone receptor; IDC, infiltrating ductal carcinoma; ILC, invasive lobular carcinoma; IQR, interquartile range

associated with it is small. As far as we know, this is the first attempt to quantify the radiation exposure with DPs. Although we did not find any adverse event directly related to radiation exposure in our group of patients, given the short follow-up and retrospective nature of our study, we need to continue to scrutinize the effects of radiation exposure from DPs. Prospective studies with a longer follow-up are warranted to answer this important health concern.

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