

## Patient Views and Correlates of Radiotherapy Omission in a Population-Based Sample of Older Women with Favorable Prognosis Breast Cancer

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**Running head:** Patient views on radiation omission

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**Abstract****Background**

Radiotherapy omission after lumpectomy is a reasonable option for many older women with favorable prognosis breast cancer. We sought to evaluate patient perspectives regarding decision-making about radiotherapy (RT).

**Methods**

Women age 65-79 with stage I and II breast cancer reported to the Georgia and Los Angeles County SEER registries were surveyed (response rate=70%) regarding radiotherapy decisions, the rationale for omitting RT, decision-making values, and understanding of recurrence risk. We also surveyed their corresponding surgeons (response rate=77%). We evaluated patient characteristics associated with omission of RT using multilevel, multivariable logistic regression, accounting for patient clustering within surgeons.

**Results**

Of 999 patients, 135 omitted RT (14%). Older age, lower grade, and estrogen receptor-positive disease were each strongly associated with omission of RT in multivariable analyses, whereas number of comorbidities was not. Non-English speakers were more likely to omit RT (adjusted OR 5.9, 95% CI 1.4-24.5).

The most commonly reported reasons for RT omission were that a physician advised the patient it was not needed (54% of patients who omitted RT) and patient choice (41%). Local recurrence

risk was overestimated by all patients, by about 2-fold among those who omitted radiation and 8-fold among those who received radiotherapy. Distant disease recurrence risk was overestimated 3-fold on average.

### **Conclusions**

To some extent, decisions about radiotherapy omission are appropriately influenced by age, grade, and estrogen receptor status, but do not appear to be optimally tailored according to competing comorbidities. Many women who are candidates for radiotherapy omission overestimate their risk of recurrence.

### **Condensed Abstract:** (2 concise sentences)

Radiotherapy omission among older women with early stage breast cancer was appropriately associated with age, grade, and estrogen receptor status, but did not appear to be tailored according to comorbid disease. Despite their favorable prognosis, many older women with early stage breast cancer markedly overestimate the risk of local and distant recurrence.

## Introduction

There is growing concern about overtreatment in older women with early stage breast cancer because they are often more likely to die of competing comorbidities than breast cancer.<sup>1</sup> Two trials (CALGB 9343 and PRIME II) have shown that among older women with stage I, estrogen receptor (ER)-positive invasive breast cancer that is treated with breast conserving surgery and endocrine therapy, adjuvant radiotherapy significantly reduces the incidence of local recurrence, but without an apparent influence on the rate of metastasis or breast cancer mortality.<sup>2,3</sup> Guidelines now consider omission of RT after BCS as an acceptable treatment option for women age  $\geq 70$  with stage T1, clinically node negative, ER-positive breast cancer who receive endocrine therapy.<sup>4</sup>

Some have argued that all older women with characteristics similar to the eligibility requirements for CALGB 9343 and PRIME II should not receive radiotherapy, whether infirm or fit.<sup>5</sup> Others have advocated for a more individualized approach to decisions about RT that accounts for tumor characteristics, comorbidity, and patient preferences.<sup>3</sup> The lack of consensus regarding the treatment approach in this scenario is evident in several practice patterns studies that revealed only a modest decline in RT utilization after publication of CALGB 9343 results.<sup>6-10</sup> Our previous work evaluating clinician views on radiotherapy omission in this context found that many clinicians overestimate the benefits associated with radiotherapy and continue to consider RT omission to be substandard therapy.<sup>11</sup> The frequency with which older patients are offered treatment without RT is uncertain, and little is known about their understanding of the risks and benefits of this treatment approach.

In this setting of evolving views on the treatment paradigm for older women with favorable prognosis breast cancer, we sought to evaluate patients' perspectives on the decision about radiotherapy omission as part of a survey that included a sizable sample of older women recently diagnosed with early stage breast cancer, as identified by the population-based Georgia and Los Angeles SEER registries. To our knowledge, this is the first study to evaluate patient views on radiotherapy omission. Our objectives were 1) to evaluate patient characteristics associated with radiotherapy omission; 2) explore patients' rationale for not receiving radiotherapy; and 3) assess patients' understanding of recurrence risk.

## **Methods**

### **Patient Sample and Data Collection**

The Individualized Cancer Care (iCanCare) Study is a large survey study of women with early-stage breast cancer between age 20-79 years who were reported to the population-based Surveillance, Epidemiology and End Results (SEER) registries of Los Angeles County, California and Georgia. Racial minorities were oversampled. Patients with tumors larger than 5cm or stage III to IV disease were excluded. From the iCanCare study, we selected women age  $\geq 65$  for the present analysis.

Between July 2013 and August 2015, we identified 7,303 women who were confirmed to be eligible for the study. Surveys were completed a median of 6.8 months (SD 3.2) after diagnosis, with a response rate of 69.6% (n = 5080). The analytic sample (n=999) for the present study

consisted of patients age 65-79 with unilateral invasive breast cancer treated with breast conserving surgery (online supplementary figure 1). Within this sample, 74% of patients had stage I disease and 23% stage II (table 1). Given that we observed a non-negligible rate of radiotherapy omission in stage II patients (12%), we also included stage II patients in multivariable models of radiotherapy omission.

Surveys were mailed with a \$20 cash incentive; a modified Dillman method was used to improve the response rate.<sup>12</sup> Materials were mailed in English; Spanish-translated materials were added for women with surnames that suggested Hispanic ethnicity.<sup>13</sup> Each SEER registry provided SEER data that were stripped of identifiers and merged to survey data. This study was approved by the University of Michigan Institutional Review Board, the University of Southern California, Emory University, and the public health departments of Georgia and California.

### **Measures**

We developed the questionnaire iteratively with input from survey design experts and cognitive interviews with patients and clinicians to assess content validity, as described previously.<sup>14</sup>

The definitive surgical procedure was determined by asking patients to indicate the surgery that was performed after biopsy, and whether additional surgeries were performed. As the primary outcome measure, radiotherapy receipt was determined by asking patients, “Did you or are you planning to have radiation therapy to treat your breast cancer?” as well as whether radiotherapy (RT) was completed, ongoing, or planned. Information on endocrine therapy

receipt was available for 62% of the analytic sample (n=619) who completed the survey module on endocrine therapy.

Patient preferences and values were assessed by asking, "When decisions were being made about your treatments, how important was it to you that your treatments..." followed by several prompts (detailed in figure 4), such as, "kept you from worrying about the cancer coming back," each rated on a 5 point Likert-type scale ranging from "not at all important" to "very important." Women who omitted radiotherapy were asked to indicate the reasons for their decision in a "mark all that apply" format. Responses were aggregated into higher (i.e. a lot, quite a bit, or somewhat) and lower (i.e. not at all or a little bit) categories for analysis.

We evaluated patient perceptions of recurrence risk by asking, "After receiving all the planned treatments, what do you think is the chance that your cancer will come back in the breast or the area around it within 10 years?" with instructions to write in a number from 0% to 100%. A similar question was asked about "the chance that your cancer will spread to other parts of your body within 10 years." We asked patients how often they had worried about their cancer coming back within the past month, with responses on a 5-point scale ranging from "almost never" to "almost always," dichotomizing those who reported "sometimes," "often," or "almost always" worrying from those reporting worrying "rarely" or "almost never." We asked how much doctors discussed the chance of cancer recurrence, ranging from "not at all" to "a lot" on a 5-point scale. Patients' decision control preferences were evaluated by asking if they



“preferred to make [their] own decisions,” with responses as “quite a bit of the time” or “all of the time” dichotomized from “some of the time,” “a little of the time,” and “none of the time.”

Additional covariates included patient-reported information on comorbidities, race/ethnicity (white, black, Asian, Latina, other), education (no college vs at least some college), income (<\$20, \$20-40, \$40-60, \$60-90, >\$90, in thousands), insurance (private, Medicare, Medicaid, other), marital status (married, divorced/separated, never married, widowed), and travel time to the nearest radiation oncology facility (<15, 15-30, 31-60, >60 minutes). We asked patients to indicate which language they primarily speak.

### **Surgeon Sample and Data Collection**

Patients were asked to identify their surgeons. From the patient analytic sample, 960 women were linked to 311 treating surgeons, of whom 240 completed a surgeon-specific survey (77%).

A mean of 3 patients (IQR 1-4) were linked to each surgeon. Surgeons were asked about their annual breast cancer patient volume, whether the practice included residents and fellows, and number of years in practice. Surgeons were also asked, “How involved are you in the selection of adjuvant radiation therapy approach in your post-lumpectomy patients?” with answers ranging from “not at all involved” to “very involved” on a 5-point scale.

**Statistical Analysis.** We first calculated the proportion of women omitting radiotherapy overall and by all demographic and treatment factors. Bivariate associations with radiotherapy omission were evaluated using the Rao-Scott  $\chi^2$  test. Multivariable, multilevel logistic

regression was used to explore the adjusted associations with radiotherapy omission, with patients as the primary units of observation and the surgeon identifiers as the secondary units (i.e. patients clustered within surgeon)<sup>15</sup>. Models were constructed beginning with patient-level covariates, incorporating surgeon clustering, and finally adding surgeon-level covariates. Area under the receiver-operator curve (AUC) was reported to measure the model's discriminatory ability. All statistical analyses incorporated weights to account for the differential probability of sample selection and survey nonresponse. Additionally, though survey and SEER item nonresponse was low (<5%) for most covariates, we multiply imputed missing items using sequential multiple imputation techniques<sup>14, 16</sup> to prevent potential bias when using complete-case methods in the presence of missing data. P-values 5% or less were considered significant throughout. All analyses were conducted using the SAS system version 9.4 (Cary, NC, USA).

## Results

### *Radiotherapy Receipt*

Table 1 and online supplementary table 1 show the distribution of patient and surgeon characteristics. Overall, 14.4% of women in this sample of older women omitted radiotherapy after breast conserving surgery, with 15.7% omission in the subset of patients with ER+, Stage I disease. Among those who received radiotherapy, 48.1% received conventionally fractionated whole breast RT, 26.6% received hypofractionated whole breast RT, 11% received accelerated partial breast RT, 2.2% reported "other" or the duration was not specified, and 12.1% were

scheduled to receive RT but had not yet received it at the time of survey. On bivariate analysis, age, grade, ER status, SEER region, and income were associated with RT omission. Figure 1 shows the results of a multilevel logistic regression model that includes patient-level variables, SEER site, and surgeon identity. Older patient age, lower grade, and ER-positive disease were each strongly correlated with RT omission. Age 75-79 had a dominant effect, with an odds ratio of 14.4 (95% CI 5.6-37.1) when compared to age 65-69. The odds of non-English speakers omitting RT were greater when compared to English speakers (OR 5.9, 95% CI 1.4-24.5). It is noteworthy that patient comorbidities were not associated with RT omission on either bivariable or multivariable analysis (OR 1.3 for  $\geq 2$  vs 0 comorbidities, 95% CI: 0.6-2.9). Surgeon variables were not significantly associated with radiotherapy omission and were therefore not retained in the final model.

The multilevel model predicted RT omission well, with an AUC of 0.83 (95% CI 0.79-0.87). When analyzing clustering according to the surgeon identifier, the odds of a patient omitting radiotherapy would be predicted to increase approximately two-fold (OR 1.83, 95% CI 0.87-3.87) if she were to see a surgeon with a practice approach one standard deviation above the RT omission rate of an average surgeon (while adjusting for other model covariates). However, this trend for surgeon influence was not statistically significant. In a model restricted to the subgroup of patients with information on hormonal therapy, use of endocrine therapy was not significantly associated with radiotherapy omission, although there appeared to be a trend for women who omit radiotherapy to also omit endocrine therapy (OR 2.22, 95% CI 0.87-5.65; data not shown). In a model limited to patients with ER+, stage I disease, we again observed that

age, grade, and non-English speakers were significantly correlated with radiotherapy omission, without significant association with comorbidities, similar to the model derived from the larger analytic sample.

### ***Risk Perception and Communication***

In the subset of patients with ER+, Stage I disease, 33.1% of women overestimated their risk of local recurrence after all treatments were received as being >10% at 10 years. Among those who omitted RT, when asked to approximate their risk of local recurrence at 10 years, the mean estimate was 19% at 10 years (compared to the CALGB finding of 10%), and among women who received RT, the mean estimate was 17% (in contrast to CALGB 9343 finding of 2%; figure 2).<sup>2</sup>

The risk of distant disease recurrence risk was similarly overestimated, with 46% of women approximating their risk of distant recurrence as being >5% at 10 years. The mean estimate of 10-year distant recurrence risk was 16.0% (compared to the CALGB finding of 5% both among those who received and omitted RT).<sup>2</sup> In this group of older women with favorable prognosis, approximately a quarter (26.9%) reported that within the last month, they had “sometimes,” “often,” or “almost always” worried about their cancer coming back, without significant differences for women who received or omitted RT.

Regarding communication with providers, 46.2% of patients in the overall sample reported that their physicians used numeric estimates to describe the risk of the cancer coming back.

Approximately one-third (37.9%) reported that their doctors discussed the chance of the cancer coming back “not at all” or “a little bit.” Although the majority of women (70.0%) reported that

they preferred that their doctors tell them what to do for breast cancer treatment, women who omitted radiotherapy were more likely to report that they preferred to make their own decisions about breast cancer treatments than women who received RT (45.8% vs 35.7%,  $p=0.03$ ).

### ***Patient Preferences and Values***

Among women who omitted radiotherapy, the most commonly reported reasons were that a physician told the patient it was not needed (53.8% of patients who omitted RT) and that the decision was left to the patient and she chose to omit RT (40.9%, figure 3). Concerns about placing an excessive burden on family and absence of discussion with a doctor about RT were uncommon reasons for omitting RT ( $\leq 5\%$ ). Although 11.8% of patients reported “quite a bit” or “a lot” of worry about current or future financial problems as a result of breast cancer and treatments,  $<1\%$  of women who omitted radiotherapy reported that cost motivated their treatment decision.

When asked about considerations that were important in their decision-making, the most commonly reported priorities were that the treatment kept them from worrying about the cancer coming back (74.0%), had a low possibility of complications (73.9%), and allowed them to continue caring for their home and family (73.6%, figure 4). Women who omitted radiotherapy more commonly endorsed the importance of avoiding exposure to radiation (69% vs 37%,  $p<0.001$ ) and the need for fewer trips for treatment visits (49.2% vs 37.2%,  $p=0.008$ ).

## Discussion

In this large, contemporary survey of older women with early stage breast cancer, we observed that to a substantial extent, decisions about radiotherapy omission appear to be appropriately tailored based on older age, ER-positive disease, and lower grade tumor. However, the higher rate of RT omission in non-English speakers and the lack of association with comorbidity observed here are concerning. Incorporation of age and pathology findings into the decision to omit RT may be viewed as a starting point, but there remains a great need to further consider comorbidity status and remaining life expectancy to ensure that decisions about radiotherapy are appropriately individualized.<sup>17</sup>

We found that despite having an excellent prognosis, a sizeable proportion of older women with Stage I, ER+ breast cancer overestimate their risk of local recurrence, with an average two-fold overestimation in women who omitted RT and 8-fold overestimation in women who received RT. Nearly half of women perceived that their risk of distant recurrence was higher than has been reported in clinical trials, with an average 3-fold overestimation. This unrealistically pessimistic view of recurrence risk is reflected in the report from a quarter of women that they had frequently worried about cancer recurrence within the preceding month. Our observations are consistent with prior reports that notwithstanding a favorable prognosis, a large proportion of the most favorable subgroup in our sample of older women felt they were likely to develop a local or distant recurrence and die from breast cancer, resulting in psychological distress, frequent worry, and lower quality of life.<sup>18, 19</sup> Overestimation of recurrence risk may lead to a multiplicative overestimation of risk reduction from

interventions,<sup>20</sup> resulting in overly generous attribution of benefits obtained from treatment. This might underlie the correlation that has been observed between worry about recurrence and receipt of radiotherapy in prior research.<sup>21, 22</sup> It is notable that one of the highest priorities that influenced decision-making for women in our study was that the treatment minimize the worry about cancer recurrence. Although radiotherapy undoubtedly does reduce the risk of local recurrence,<sup>2, 3</sup> our results indicate a need to more clearly communicate the favorable prognosis in this group, and to more directly address worry about recurrence and ensure that decisions are optimally informed.

The effects of patient overestimation of recurrence risk may well relate to physician overestimation that has been demonstrated in other work. In a nationwide survey, 19% of radiation oncologists and 32% of surgeons overestimated the 10-year risk of local recurrence.<sup>11</sup> The reluctance of many surgeons and radiation oncologists to consider radiotherapy omission to be a reasonable option in select older women is particularly problematic given the findings of the current study demonstrating that the most common reason given by women for the decision to omit radiotherapy was advice from a doctor that radiotherapy was not needed. Furthermore, most women (70%) preferred that their doctors tell them what to do regarding their breast cancer treatment. Therefore, physicians' attitudes and approaches to communication may be particularly important to ensure that patients do indeed consider all options and their risks and benefits. Many potential mechanisms have been considered as possible drivers of physicians' poor communication practices in this setting, including lack of up-to-date knowledge of clinical trial results,<sup>23</sup> heuristics like risk aversion and anticipatory regret

that may motivate physicians and patients alike to embrace overly aggressive treatments,<sup>24</sup> lack of training in effective risk communication skills,<sup>25</sup> and financial incentives that reimburse the delivery of care rather than its omission.<sup>7</sup> Our data suggest a compelling need to evaluate the relative roles of these underlying drivers further in order to develop appropriately targeted interventions that encourage clinicians to improve communication in this regard. We find it striking that one-third of women reported minimal discussions with their providers about the risk of recurrence, which is absolutely essential for a patient to understand the relative impact of interventions like radiotherapy.

Our data suggest that surgeons, who are the first breast cancer clinicians to outline a plan of care, may play an important role in decisions regarding radiotherapy omission. Although we did not detect a statistically significant impact of the surgeon on the likelihood of radiotherapy omission, this may be due to the sample size and distribution of patients across surgeons within our sample. We did, however, observe a notable trend, and others have shown that among women with short life expectancy, the probability of receiving RT varies substantially across primary surgeons.<sup>26</sup>

We also found that non-English speakers were significantly more likely to omit radiotherapy, even after adjustment for race, income, employment and education. This finding may reflect barriers to high quality decision-making in a vulnerable population. Numerous reports describe non-English speakers as a vulnerable population, particularly Latinas with low acculturation.<sup>27-30</sup> Latina women who are less acculturated have previously been found to have greater desire for



information, lower satisfaction with breast cancer decision-making,<sup>30</sup> and perceived powerlessness in medical encounters,<sup>31</sup> highlighting a need for greater attention to support these patients and identify potential barriers.<sup>32</sup> Although disparities are often viewed in the context of undertreatment of an aggressive cancer, with vulnerability related to known risk factors such as lower education or minority race, it is interesting to note that vulnerability to overtreatment may be an entirely distinct concept that primarily affects classically privileged populations, as has been observed with trends in contralateral prophylactic mastectomy.<sup>33, 34</sup> The association between language barrier and RT omission is worthy of further exploration in future studies.

We observed a trend for patients who omit radiotherapy to also omit endocrine therapy. This is a particularly concerning finding that merits further evaluation, as receipt of endocrine therapy is known to have a substantial impact in reducing local recurrence<sup>35</sup> and is believed to be a key factor in leading to the acceptable rates of local failure observed in trials like CALGB 9343 as compared to historical studies where endocrine therapy was not required.<sup>36, 37</sup> Endocrine therapy non-adherence and discontinuation are known to be an issue for nearly half of women with breast cancer in general,<sup>38</sup> and if women who omit radiotherapy are even more likely to omit endocrine therapy, rates of recurrence may be higher than expected.

Aspects of the study merit comment. Strengths include a contemporary, diverse, population-based sample with a high response rate and specific measures of patients' clinical decision making. Multiple imputation and weighting were used to account for potential bias related to

missing data, and to ensure that the results were representative of the overall population. Limitations include data obtained from 2 large SEER regions (Georgia and Los Angeles County), which might not reflect the entire US population. The number of patients who omitted radiotherapy is small. Patient responses are necessarily retrospective and may be subject to recall bias. Finally, our study evaluated patients' views of radiotherapy omission, but did not explore views about omitting endocrine therapy rather than radiotherapy.<sup>39</sup> Nonetheless, our study offers a novel and clinically relevant view of decision-making in older women with favorable prognosis breast cancer.

In conclusion, our findings indicate several targets for interventions to improve the quality of older women's decision-making in the context of radiotherapy after lumpectomy. First, although decisions about radiotherapy omission are influenced by some clinical factors, interventions are necessary to ensure that decisions reflect not only considerations of age and tumor characteristics but also patients' health status and remaining life expectancy. Second, communication must improve, as patients cannot share in the making of preference-concordant decisions when they are not optimally informed about key facts such as recurrence risk and report that their providers had little discussion with them regarding this. Ultimately, we believe that a combination of physician-facing and patient-facing interventions are necessary, as it appears that both patients and physicians play important roles in the predominant intervention bias that has resulted in overtreatment of many older women with early stage breast cancer.<sup>7</sup> Our results demonstrate that most older women with breast cancer care about avoiding complications and about avoiding worry about recurrence. Therefore, decision aids

that present risk information in understandable formats, such as pictographs,<sup>40</sup> constitute particularly promising avenues for the improvement of decision quality in practice. Clear information and communication is critical, both for the subset of older patients with favorable risk disease in whom radiotherapy omission is a reasonable and guideline-concordant option if it accords with individual preferences, and also for the subset with more aggressive or advanced tumors, for whom radiotherapy remains a fundamental component of cure.

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### **Figure Legends**

**Figure 1:** Odds ratios from a model estimating radiotherapy omission. Odds ratios from a multilevel logistic regression model estimating radiotherapy omission. The model was adjusted for race, income, education, insurance, marital status, BMI, and SEER site. The odds ratio for the surgeon effect represents the amount by which a patient's odds of radiotherapy omission are multiplied if they see a surgeon associated with a rate of radiotherapy omission that is one standard deviation above the average surgeon.

**Figure 2:** Patient-reported estimates of 10-year risk of local recurrence in patients aged 65-79 with stage I, estrogen receptor positive invasive breast cancer compared to CALGB 9343 results.

**Figure 3.** Patient reported reasons for omission of radiotherapy. Responses are not mutually exclusive.

**Figure 4.** Patient reported considerations that influenced decisions about breast cancer treatment.

**Online supplementary figure 1:** Flow diagram of development of analytic sample.

**Table 1:** Patient Characteristics

	No. (N=999)	Weighted %	% Omitting RT*	<i>P</i> <sup>†</sup>
<b>Age</b>				
65-69	410	40.2	5.4	<0.0001
70-74	337	34.1	16	
75-79	252	25.7	26	
<b>SEER Stage</b>				0.5404
1	749	73.5	14.3	
2	218	23.2	12.2	
Not reported	32	3.2	28.2	
<b>ER status</b>				0.0164
Positive	857	85.5	15.3	
Negative	111	11.5	5.2	
Not reported	31	3.0	20.2	
<b>SEER Grade</b>				0.0029
1	346	34.8	18.1	
2	430	43.4	14.4	
3	187	18.2	6.1	
Not reported	36	3.7	17.8	
<b>Comorbidities</b>				0.2995
0	482	49.5	13.6	
1	315	30.3	12.8	
2+	190	18.9	17.8	
Not reported	12	1.3	21.8	
<b>Receipt of endocrine therapy<sup>‡</sup></b>				0.2222
Yes	449	71.1	13.6	
No	170	26.8	17.7	
Not reported	15	2.1	18.5	
<b>Site</b>				0.0113
Georgia	522	49.3	11.2	
Los Angeles County	477	50.7	17.2	
<b>Primarily speak language other than English</b>				0.1279
Yes	896	90.9	13.5	
No	89	7.8	20.4	
Not reported	14	1.3	30.4	
<b>Race</b>				0.3019
White	631	67.5	14.9	
Black	160	13.9	10.3	
Latina	136	11.3	14.9	
Asian	52	5.2	10.9	

Other, unknown, or missing	20	2.1	26.6	
<b>Income</b>				0.0265
<20K	163	15.3	15.3	
20K - <40K	183	17.6	9.6	
40K - <60K	136	13.4	15.5	
60K - <90K	134	14.4	10.4	
90K+	147	16.8	11.4	
Don't know/not reported	236	22.6	21	
<b>Education</b>				0.5094
At least some college	614	63.4	13.8	
No college	365	34.8	15.4	
Not reported	20	1.8	8.6	
<b>Type of insurance</b>				0.1035
Medicaid	104	9.9	15.3	
Medicare	641	66.1	14.5	
Private	121	12.2	6	
Other	7	0.6	14.7	
Not reported	126	11.2	20.9	
<b>Marital status</b>				0.5347
Married/partnered	523	53.3	13.5	
Not partnered	457	44.8	15	
Not reported	19	1.9	20.5	
<b>BMI</b>				0.1281
Underweight (<18.5)	8	0.7	33.5	
Normal weight (18.5-25)	262	27.8	17.4	
Overweight (>25-30)	323	32.7	12.5	
Obese >30	371	35.7	12.7	
Not reported	35	3.2	18.4	
<b>Bra cup size</b>				0.5723
A/B	289	29.0	15.9	
C	322	32.1	14.7	
D	197	19.7	13.3	
DD+	156	15.8	11	
Not reported	35	3.4	17.1	
Distance to nearest radiation oncology clinic <sup>‡</sup>				0.5430
≤30 minutes	350		11.3	

>30 minutes	126		11.3	
Not reported	34		54.8	

\*Percent omitting RT calculated within the weighted sample

† *P* values for differences in the proportion of RT omission; the 'not reported' category (if present) was excluded from the calculation.

‡ Not all patients were asked to provide this information due to differences in survey versions

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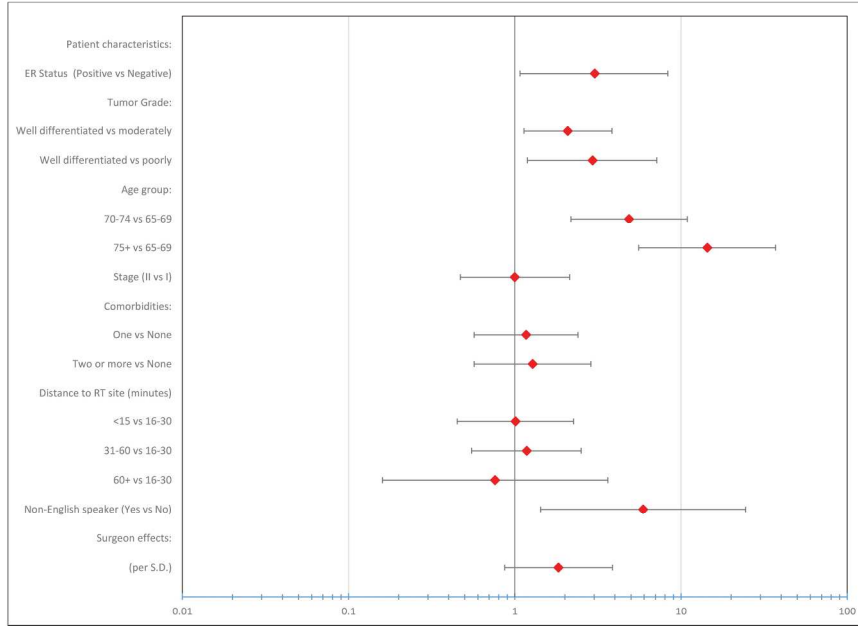


Figure 1: Odds ratios from a model estimating radiotherapy omission. Odds ratios from a multilevel logistic regression model estimating radiotherapy omission. The model was adjusted for race, income, education, insurance, marital status, BMI, and SEER site. The odds ratio for the surgeon effect represents the amount by which a patient's odds of radiotherapy omission are multiplied if they see a surgeon associated with a rate of radiotherapy omission that is one standard deviation above the average surgeon.

215x166mm (300 x 300 DPI)

Autho

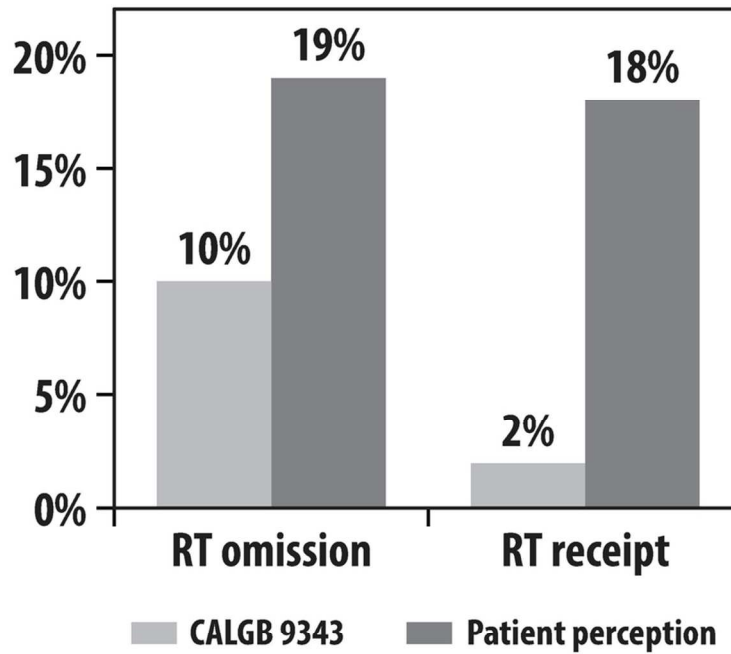


Figure 2: Patient-reported estimates of 10-year risk of local recurrence in patients aged 65-79 with stage I, estrogen receptor positive invasive breast cancer compared to CALGB 9343 results.

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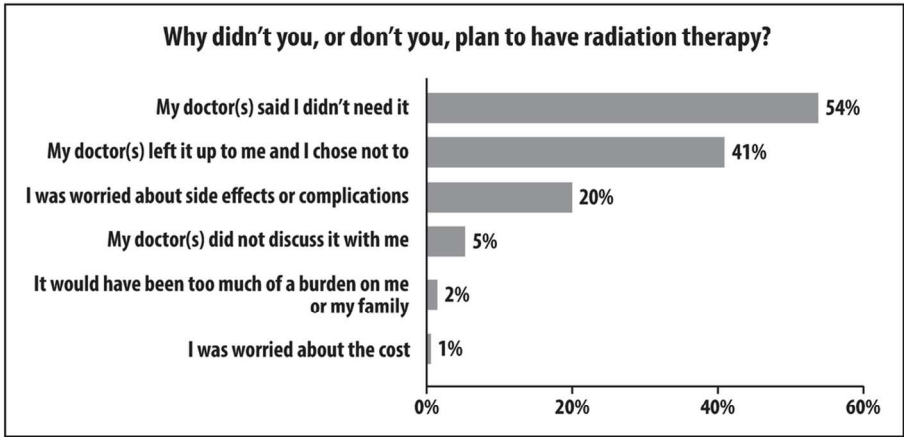


Figure 3. Patient reported reasons for omission of radiotherapy. Responses are not mutually exclusive.

106x59mm (300 x 300 DPI)

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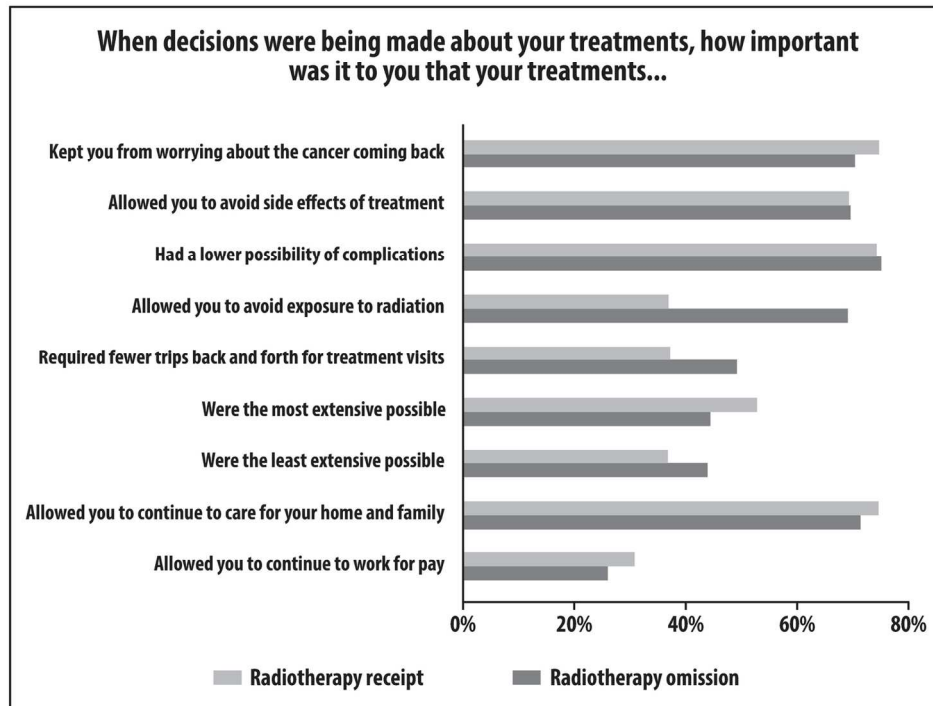


Figure 4. Patient reported considerations that influenced decisions about breast cancer treatment.

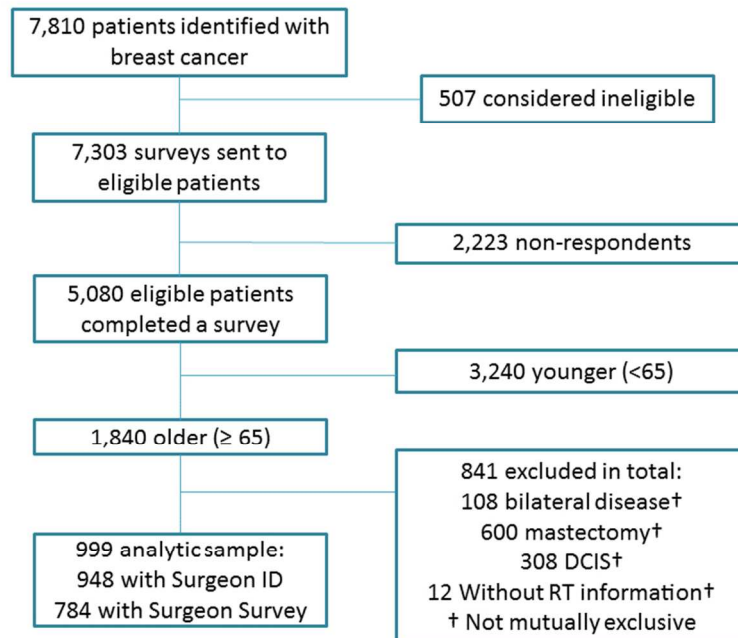
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**Online supplementary table 1:** Physician Characteristics

	<b>No.* (N = 240)</b>	<b>Weighted %</b>
<b>Years in practice</b>		
0-10 yrs	46	19.2
11-20	71	29.6
≥21 yrs	118	49.2
Missing	5	2.1
<b>Site</b>		
Georgia	120	50.0
LA	120	50.0
<b>Work with residents and/or fellows</b>		
Yes	67	27.9
No	169	70.4
Missing	4	1.7
<b>Breast cancer volume within last 12 months</b>		
≤50	142	59.2
>50	88	36.7
Missing	10	4.1
<b>Surgeon involvement in RT decisions</b>		
Not at all or slightly	93	38.8
Somewhat	46	19.2
Moderately or very	94	39.2
Missing	7	2.9
*Number of surgeons who were linked to a patient in table 1 and completed a survey.		



254x190mm (96 x 96 DPI)

Author

A12. Please tell us whether you have ever been told by a doctor that you had any of the following health conditions:

Yes	No	
<input type="radio"/>	<input type="radio"/>	a. Chronic bronchitis or emphysema
<input type="radio"/>	<input type="radio"/>	b. Heart disease, such as coronary artery disease or congestive heart failure
<input type="radio"/>	<input type="radio"/>	c. Diabetes
<input type="radio"/>	<input type="radio"/>	d. Blood clots in the legs or the lung
<input type="radio"/>	<input type="radio"/>	e. Connective tissue disease, such as lupus or scleroderma
<input type="radio"/>	<input type="radio"/>	f. Stroke
<input type="radio"/>	<input type="radio"/>	g. Depression
<input type="radio"/>	<input type="radio"/>	h) High cholesterol
<input type="radio"/>	<input type="radio"/>	i) High blood pressure or hypertension

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C1. **Before your surgery**, did you consult with any of these providers about your treatment decisions?

Yes No

- |                       |                       |  |
|-----------------------|-----------------------|--|
| <input type="radio"/> | <input type="radio"/> | a) Radiation oncologist (a doctor who specializes in radiation treatment)  |
| <input type="radio"/> | <input type="radio"/> | b) Medical oncologist (a doctor who specializes in chemotherapy)   |
| <input type="radio"/> | <input type="radio"/> | c) Plastic surgeon (a doctor who performs breast reconstruction)   |
| <input type="radio"/> | <input type="radio"/> | d) Primary care provider (a doctor, physician assistant, or nurse practitioner who sees you for common problems or standard checkups)                            |
| <input type="radio"/> | <input type="radio"/> | e) Genetic counselor (someone who reviews your family history and explains the purpose and results of genetic tests)   |
| <input type="radio"/> | <input type="radio"/> | f) Navigator (often a nurse, someone who helps you with your treatment schedule, test results, or who gives you and your family information about breast cancer) |

### Surgery

C2. What was the first surgery that you had to remove your breast cancer after the biopsy test?

- I did not have any surgery after the biopsy →
- I had a mastectomy (removal of the entire breast) →
- I had a lumpectomy (removal of the cancer and some surrounding tissue)

**Please go to C3 at the top of the next page**

- a. Did you have a second lumpectomy to remove more breast tissue from the same breast?
- Yes – I had another lumpectomy to remove more breast tissue from the same breast
- No – I only had one lumpectomy
- b. Did you have a mastectomy later, on the same breast?
- Yes – I had a mastectomy after my lumpectomy
- No – I did not have a mastectomy
- c. How strongly did your doctor recommend that you have a mastectomy after your initial lumpectomy?
- |                       |                       |                       |                       |                       |
|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|
| <input type="radio"/> | <input type="radio"/> | <input type="radio"/> | <input type="radio"/> | <input type="radio"/> |
| Very strongly         | Strongly              | Moderately            | Weakly                | Not at all            |
- d. How strongly did you request to have a mastectomy after your initial lumpectomy?
- |                       |                       |                       |                       |                       |
|-----------------------|-----------------------|-----------------------|-----------------------|-----------------------|
| <input type="radio"/> | <input type="radio"/> | <input type="radio"/> | <input type="radio"/> | <input type="radio"/> |
| Very strongly         | Strongly              | Moderately            | Weakly                | Not at all            |

**Please continue to C3 on the next page**

C3. What kind of mastectomy did you have?

- I did not have a mastectomy
- Mastectomy only – no reconstruction
- Mastectomy with reconstruction and I kept my own nipple, called a nipple sparing or nipple saving mastectomy
- Mastectomy with reconstruction and my original nipple was removed

C4. What type of breast reconstruction did you have?

- I have not had any breast reconstruction surgery
- A DIEP flap, TRAM flap, or latissimus dorsi flap (uses your own tissue from the abdomen or back)
- An implant (silicone or saline)
- Other (please explain): \_\_\_\_\_

C5. Did you have a mastectomy on **both breasts**?

<input type="radio"/>	<input type="radio"/>
Yes	No

Go to C6  
below

Please go to C7 at the  
top of the next page

C6. How important were the following factors in your decision to have a mastectomy on **both breasts**?

	Not at all important	A little important	Somewhat important	Quite important	Very important
a. My age	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. Having a family history of breast cancer	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
c. Wanting both breasts to match after reconstruction	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

Please continue to C7 on the next page

**Radiation Therapy**

Please answer the following questions even if you did not have radiation therapy

C8. **At the time of your breast cancer diagnosis**, how much time did it or would it have taken you to get from your home to the nearest radiation treatment facility?

<input type="radio"/> Less than 15 minutes	<input type="radio"/> 31 to 60 minutes	<input type="radio"/> I don't know
<input type="radio"/> 15 to 30 minutes	<input type="radio"/> More than 60 minutes	

C9. Did any doctor tell you that radiation treatment...

	Yes	No	Don't remember
a. Usually involves daily trips to an outpatient facility	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. Can be completed in 3-4 weeks or less	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

C14. Did you or are you going to have radiation therapy to treat your breast cancer?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Yes – I am finished with radiation therapy	Yes – I am still having radiation therapy	Yes – I plan to have radiation therapy but haven't started yet	No

Please go to C15 below

Please go to C14a

C14a. Why didn't you, or don't you, plan to have radiation therapy?

**Please mark ALL that apply.**

- My doctor(s) did not discuss it with me
- My doctor(s) said I didn't need it
- My doctor(s) left it up to me and I chose not to
- I was worried about side effects or complications
- I was worried about the cost
- It would have been too much of a burden on me or my family
- Other (please explain): \_\_\_\_\_  
\_\_\_\_\_

**Please go to C18 on the next page**

C15. How many total weeks of radiation treatments are planned?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
1 week	2-4 weeks	5 weeks	6 or more weeks

C16. How was, or will, your radiation treatment be given? **Please mark ALL that apply.**

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Using beams from outside my body	Using a balloon placed inside my breast	Using needles placed into my breast	I don't know how my radiation treatment was, or will be, given

C17. Which of the following areas were, or will be, treated with radiation? **Please mark ALL that apply.**

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
My whole breast	Only the part of the breast where my tumor was	The chest wall, after my mastectomy	The lymph node areas

### **Hormonal Therapy**

C20. Hormonal therapy helps block estrogen from getting to cancer cells that may remain in the body. Hormonal therapy is sometimes called “anti-estrogen therapy” or “endocrine therapy.” Examples include **tamoxifen**, **anastrozole** or Arimidex, **letrozole** or Femara, and **exemestane** or Aromasin.

Have you or are you going to be taking any of these medications? **Please mark ONE.**

- Yes, I currently take one of these medications
- Yes, I plan to take one of these medications in the future but haven't started yet
- Yes, I took one of these medications before but no longer take it
- No, I am not taking any of these medications right now and I am unsure whether or not I should start
- No, I have no plans to take any of these medications

## Section D: Decision Making

D1. In **general**, please tell us how often you have these thoughts and feelings when you make decisions.

	Never	Rarely	Sometimes	Often	Always
a. I worry about making a bad decision	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. I struggle to decide what the right decision is	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
c. I get angry at myself when I have made a bad decision	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
d. I worry a lot about the outcomes of my decisions	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

D2. When making decisions about **how to treat my breast cancer**...

	Not at all	A little	Somewhat	Quite a bit	A lot
a. I weighed the pros and cons of all the treatment options	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. I feel like I really thought through all the issues important to the treatment decisions	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
c. I talked with others – family or friends – before making treatment decisions	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
d. I talked with other breast cancer patients before making treatment decisions	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
e. I spent time thinking about all of the treatment options	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

D3. When making decisions about **how to treat my breast cancer**...

	Not at all	A little bit	Somewhat	Quite a bit	Very much
a. I would like to have had more information	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. I would like to have participated more	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
c. I am satisfied with the amount of time I had	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
d. I am satisfied with the amount of involvement I had from family and friends	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>



D4. **When decisions were being made about your treatments**, how important was it to you that your treatments...

	Not at all important	A little important	Somewhat important	Quite important	Very important
a. Reduced the need for more surgery	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
b. Allowed you to avoid side effects of treatment	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
c. Allowed you to avoid exposure to radiation	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
d. Required fewer trips back and forth for treatment visits	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
e. Did not make you feel bad about your body	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
f. Kept you from worrying about the cancer coming back	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
g. Allowed you to feel feminine	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
h. Were the most extensive possible	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
i. Were the least extensive possible	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
j. Allowed you to keep your original breast	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
k. Were what your partner/family wanted you to do	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
l. Were what your doctor wanted you to do	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
m. Were the same treatments that other women you know have received	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
n. Were the newest, most advanced treatments available	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
o. Had the shortest recovery time	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
p. Gave you peace of mind	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
q. Allowed you to avoid having follow-up mammograms	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
r. Did not require you to spend a lot of your own money	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
s. Had a lower possibility of complications	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
t. Allowed you to continue to care for your home and family	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
u. Allowed you to continue to work for pay	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

D5. At the time that decisions were being made about your treatments, how much do you feel that your preferences were considered?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Not at all	Slightly	Moderately	Very much	Completely

## Section E: How You Feel About Your Decisions

Please tell us how you feel about the decisions that were made for your breast cancer treatment. If your doctor did not offer you the test or treatment that is listed, please mark N/A for "Not applicable."

E1. Please rate the **amount of information** you had when the following decisions were made:

	Not enough		Just right		Too much	N/A
e. Whether or not to have radiation therapy	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

E2. I received enough information about the following topics:

	Strongly Agree	Agree	Neither agree nor disagree	Disagree	Strongly Disagree
d. The risk of my breast cancer coming back	<input type="radio"/>	<input type="radio"/>			

E3. How **satisfied** are you with the decision about...

	Not at all satisfied	A little satisfied	Somewhat satisfied	Quite satisfied	Totally satisfied	N/A
e. Whether or not to have radiation therapy	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

## Section H: Communicating with Your Doctors

	None of the time	A little of the time	Some of the time	Quite a bit of the time	All of the time
H1. When it came to getting treatment for breast cancer, I preferred to be told what to do	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
H2. When it came to getting treatment for breast cancer, I wanted <u>my doctor</u> to tell me what to do	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
H3. I preferred to make my <u>own</u> decisions about my treatments for breast cancer	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

## Section I: Your Thoughts and Feelings

The questions below are important to help us better understand how women like you feel towards the end of treatment and the beginning of the recovery period. Please answer these questions the best way you can.

I1. For the question below, please write in a number from 0% to 100% where...

0% = you think there is absolutely no chance that your breast cancer will come back in the breast or the area around it in the next 10 years, and

100% = you think it is absolutely certain that your breast cancer will come back in the breast or the area around it in the next 10 years

After receiving all the planned treatments, what do you think is the chance that your cancer will come back in the breast or the area around it within 10 years?

\_\_\_\_\_ % (0 to 100)

I2. After receiving all of the planned treatments, do you consider the chance of your cancer coming back in the breast or the area around it to be:

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Very low	Low	Moderate	High	Very high

I3. Next, we would like to ask you for your opinion on the chances of your cancer spreading to other parts of your body. For the question below, please write in a number from 0% to 100% where...

0% = you think there is absolutely no chance that your breast cancer will spread to other parts of your body in the next 10 years, and

100% = you think it is absolutely certain that your breast cancer will spread to other parts of your body in the next 10 years

After receiving all the planned treatments, what do you think is the chance that your cancer will spread to other parts of your body within 10 years?

\_\_\_\_\_ % (0 to 100)

13a. Why did you pick this number?

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14. After receiving all the planned treatments, do you consider the chance of your cancer spreading to other parts of your body to be:

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Very low	Low	Moderate	High	Very high

15. How much did your doctors discuss with you the chance of your cancer coming back?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Not at all	A little bit	Somewhat	Quite a bit	A lot

16. When your doctors discussed the chance of your cancer coming back, did they use...

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Only words (For example, "small chance")	Only numbers (For example, "8% chance")	Both words and numbers	My doctors did not discuss the risk with me

17. Compared to other women with similar breast cancer and treatment, how likely do you think it is that your breast cancer will come back?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Much less likely	Less likely	About the same	More likely	Much more likely

18. **In the past month**, how often have you worried about your cancer coming back?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Almost never	Rarely	Sometimes	Often	Almost always

19. **During the past month**, how often has worrying about your cancer coming back...

	Almost never	Rarely	Sometimes	Often	Almost always
a. Made you feel upset?	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>

b. Made it difficult for you to carry out your usual daily activities at home or at work?

c. Made you feel distant from family and friends?

110. When you see your cancer doctors for follow-up care, how often do they ask if you are worried about your breast cancer coming back?

Almost never  Rarely  Sometimes  Often  Almost always

### Section K: Home and Work

K15. How much do you worry about current or future financial problems as a result of your breast cancer and treatments?

Not at all  A little  Somewhat  Quite a bit  A lot

### Section L: Language and Other Preferences

L1. What language do you primarily speak? **Please mark ONE.**

English  Mandarin  Cantonese  Korean  
 Spanish  Vietnamese  Japanese  Other (please explain): \_\_\_\_\_

L2. In general, what language(s) do you read and speak?

Only English  English better than any other language  Both equally  Another language better than English  Only another language

L3. What language do you usually speak at home?

Only English  More English than any other language  Both equally  Another language more than English  Only another language

L4. In what language do you usually think?

Only English  More English than any other language  Both equally  Another language more than English  Only another language

L5. What language do you usually speak with your friends?

Only English  More English than  Both equally  Another language  Only another

any other language

more than English

language

## Section M: A Few More Questions

M1. Today's date is: \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_  
 month      day      year

M2. About how tall are you? \_\_\_\_\_ feet \_\_\_\_\_ inches or \_\_\_\_\_ meters

M3. **At the time of your breast cancer diagnosis**, about how much did you weigh?

\_\_\_\_\_ pounds or \_\_\_\_\_ kilograms

M4. **Before your breast surgery**, what was your bra cup size?

- A       D  
 B       DD  
 C       Other (please explain): \_\_\_\_\_

M5. **In the 12 months before your diagnosis with breast cancer**, what was your experience with your menstrual periods?

- I had no menstrual periods in the 12 months before my breast cancer diagnosis  
 I had regular (or the usual timing of) menstrual periods in the 12 months before my breast cancer diagnosis  
 I had a change in the timing of menstrual periods in the 12 months before my breast cancer diagnosis

M6. **In the 12 months before your breast cancer diagnosis**, did you experience hot flashes or night sweats at any time – even once?

- |                       |                       |
|-----------------------|-----------------------|
| <input type="radio"/> | <input type="radio"/> |
| Yes                   | No                    |

M7. What is your birth date? \_\_\_\_\_ / \_\_\_\_\_ / \_\_\_\_\_  
 month      day      year

M8. When you were diagnosed with breast cancer, what was your marital status?

- Married                       Living with partner  
 Divorced                       Widowed  
 Separated                       Never married

M9. What is the highest level of education you have completed?

- No high school                       Some college or technical school  
 Some high school                       College graduate (Bachelor's degree)  
 High school graduate or G.E.D.                       Graduate degree or higher

M10. Are you of Hispanic, Latino, or Spanish origin?

- Yes, Mexican, Mexican-American, or Chicano  
 Yes, Puerto Rican  
 Yes, Cuban  
 Yes, another Hispanic, Latino, or Spanish origin (please explain): \_\_\_\_\_  
 No

M11. Are you of Jewish descent?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Yes	No	Don't know

M12. For how many years have you lived in the United States? \_\_\_\_\_ years

M13. In what country were you born? \_\_\_\_\_

- Don't know

M14. In what country was your mother born? \_\_\_\_\_

- Don't know

M15. In what country was your father born? \_\_\_\_\_

Don't know

M16. Which of the following best describes your race? **Please mark ALL that apply.**

White

Chinese

Black or African-American

Filipino

American Indian or Alaska Native

Japanese

Native Hawaiian or other Pacific Islander

Korean

Asian Indian

Vietnamese

Other Asian (please explain): \_\_\_\_\_

Other Race (please explain): \_\_\_\_\_

### Section N: Your Doctors

N1. We want to learn from doctors about better ways to communicate with patients and their families about treatment decisions. The information you provide below will help us contact the doctors who treat patients with breast cancer. The doctors may be surveyed about their treatment practices. Importantly, your answers will never be shared with any doctors and your personal information including your name will never be used in any communication.

a. **Surgeon** who performed your first lumpectomy or mastectomy:

Doctor's last name: \_\_\_\_\_ First name: \_\_\_\_\_

Name of hospital or clinic: \_\_\_\_\_

City: \_\_\_\_\_



## Section A: About You and Your Practice

A1. At how many hospitals do you see patients?

- 1       2       3 or more

A2. Does your primary practice have...

	Yes	No
a. Residents and/or fellows	<input type="radio"/>	<input type="radio"/>
b. Cancer care navigators for newly-diagnosed breast cancer patients to help them with treatment decision making	<input type="radio"/>	<input type="radio"/>

A3. **In the past 12 months**, approximately what percentage (%) of your new patients were...

a. \_\_\_\_\_ % Breast cancer (including DCIS) patients

b. \_\_\_\_\_ % Other (non-breast) oncology patients

c. \_\_\_\_\_ % Patients without cancer

**100 % Total for the past 12 months**

A4. **In the past 12 months**, how many new patients who were diagnosed with breast cancer have you seen? (Please include patients who came to you for a second opinion.)

- None       11 – 20       51 – 100  
 1 – 10       21 – 50       More than 100

A5. **In the past 12 months**, approximately what percent of your newly diagnosed breast cancer patients did you discuss in a multidisciplinary meeting (e.g., a tumor board) to get input on the treatment plan?

- 0%       10 – 25%       More than 50%  
 1 – 9%       26 – 50%

## Section I: Radiation Therapy

I3. How involved are you in the selection of adjuvant radiation therapy approach in your post-lumpectomy patients?

<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>	<input type="radio"/>
Not at all involved	Slightly involved	Somewhat involved	Moderately involved	Very involved

**Section L: A Few More Questions**

L1. How many years have you been in practice since completing your residency or fellowship training?

\_\_\_\_\_ years

L2. Please PRINT the name of the location where you practice the most: \_\_\_\_\_

L3. How old are you? \_\_\_\_\_ years

L4. What is your gender?

Male

Female

L5. Which of the following best describes your race? **Please mark ALL that apply.**

White

Chinese

Black or African-American

Filipino

American Indian or Alaska Native

Japanese

Native Hawaiian or other Pacific Islander

Korean

Asian Indian

Vietnamese

Other Asian (please explain): \_\_\_\_\_

Other Race (please explain): \_\_\_\_\_

M3. Which **radiation oncologists** do you interact with the most when you are uncertain about the **management of your newly diagnosed patients with breast cancer**? Please **PRINT** the first and last names of up to 2 radiation oncologist whom you would approach to get advice.

<u>Name</u>	<u>Hospital / Institution / Practice</u>	<u>Do you share patients via a tumor board?</u>	
		Yes	No
_____	_____	<input type="radio"/>	<input type="radio"/>
_____	_____	<input type="radio"/>	<input type="radio"/>

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