

Racial disparities in low-value surgical care and time to surgery in high-volume hospitals

Destiny K. Jackson¹ | Yaming Li MD, MS² | Mariam F. Eskander MD, MPH² |
Allan Tsung MD² | Bridget A. Oppong MD² | Oindrila Bhattacharyya MSc³ |
Electra D. Paskett PhD⁴ | Samilia Obeng-Gyasi MD, MPH² 

¹Central State University, Wilberforce, Ohio, USA

²Division of Surgical Oncology, Department of Surgery, The Ohio State University, Columbus, Ohio, USA

³Department of Economics, Indiana University Purdue University, Indianapolis, Indiana, USA

⁴Department of Internal Medicine, College of Medicine, The Ohio State University, Columbus, Ohio, USA

Correspondence

Samilia Obeng-Gyasi, The Ohio State University, N924 Doan Hall. 410 West 10th, Columbus, OH 43210, USA.
Email: samilia.obeng-gyasi@osumc.edu

Funding information

Ohio State University Comprehensive Cancer Center using Pelotonia funds

Abstract

Background: The objective of this study is to examine racial differences in receipt of low-value surgical care and time to surgery (TTS) among women receiving treatment at high-volume hospitals.

Methods: Stage I–III non-Hispanic Black (NHB) and Non-Hispanic White (NHW) breast cancer patients were identified in the National Cancer Database. Low-value care included (1) sentinel lymph node biopsy (SLNB) among T1N0 patients age ≥ 70 with hormone receptor–positive cancers, (2) axillary lymph node dissection (ALND) in patients meeting ACOSOG Z0011 criteria, and (3) contralateral prophylactic mastectomy (CPM) with unilateral cancer. TTS was days from biopsy to surgery. Bivariate and logistic regression analyses were used to compare the groups.

Results: Compared to NHWs, NHBs had lower rates of SLNB among women age ≥ 70 with small hormone–positive cancers (NHB 58.5% vs. NHW 62.2% $p < .001$) and CPM (NHB 26.3% vs. NHW 36%; $p < .001$). ALND rates for patients meeting ACOSOG Z0011 criteria were similar between both groups ($p = .13$). The odds of surgery > 60 days were higher among NHBs (odds ratio, 1.77; 95% confidence interval, 1.64–1.91; NHW ref).

Conclusions: NHBs treated at high-volume hospitals have higher rates of surgical delay but are less likely to undergo low-value surgical procedures compared to NHW women.

KEYWORDS

breast cancer, disparities, race, surgery, value

1 | INTRODUCTION

Advancements in the diagnosis and treatment of breast cancer have resulted in earlier stages of diagnosis and increased survival.¹ Current Surveillance Epidemiology and End Result (SEER) Program data show a 90% 5-year relative survival rate for all breast cancer stages.² However, improvements in clinical outcomes such as survival have not extended to non-Hispanic Black (NHB) breast cancer patients. NHB women diagnosed with breast cancer present at younger ages,

with more advanced stages and increased rates of aggressive subtypes.^{3,4} In addition, they face disparities in dosing and frequency of chemotherapy, endocrine therapy adherence, and surgical delays.^{5,6} These differences in treatment are significant as they contribute to worse disease-specific and overall survival.^{7,8}

An avenue to mitigate racial disparities in breast cancer outcomes include evaluations of patterns of care. Studies suggest breast cancer patients treated by oncology specialists or by high-volume providers or centers are more likely to receive guideline-concordant

care and have higher survival rates than patients treated by non-specialists or at low-volume centers.^{9–11} A review of the National Cancer Database (NCDB) suggests that treatment at a high-volume center (defined as annual cases ≥ 298) resulted in an 11% reduction in mortality compared to treatment at low-volume centers.¹² Notably, NHB women in the study did not appear to have any improvement in survival when treated at high-volume centers compared with low-volume centers.¹² In the aforementioned study, hospital volume categories were based on the relationship between annual hospital volume and survival after controlling for potential confounding variables. This approach to defining hospital volume provides a more nuanced understanding of the volume–outcome relationship compared with studies constructing volume categories based on the distribution of cases within a cohort.

To better understand processes of care that contribute to racial disparities in clinical outcomes, the objective of this study was to examine differences in surgical management and time to first definitive surgery between NHB women and Non-Hispanic White (NHW) women receiving treatment in high-volume hospitals in the NCDB. Specifically, we were interested in racial differences in the utilization of low-value surgical care and surgical delays. Low-value surgical procedures are defined as procedures with minimal to no clinical benefit.¹³ The motivation behind focusing on these procedures was to better understand racial differences in the quality of surgical care. Surgical delay was evaluated due to its association with increased disease-specific and overall survival.⁷ We hypothesized that NHB women were more likely to receive low-value surgical care and experience longer wait times from biopsy-proven diagnosis to first definitive surgery at high-volume hospitals compared to their NHW counterparts. To this end, results from this study could provide additional insight into surgically rooted racial differences in outcomes among patients treated at high-volume centers.

2 | METHODS

2.1 | Data source

The NCDB prospectively collects cancer cases from 1500 Commission on Cancer (CoC) accredited facilities and represents 70% of newly diagnosed cancers in the United States.¹⁴ The NCDB is jointly managed by the American Cancer Society and the American College of Surgeons.

2.2 | Study population

The NCDB was queried for NHB and NHW female breast cancer patients ages 18–90, diagnosed with stage I–III breast cancer between 2010 and 2016, who received treatment at high-volume hospitals (Figure 1). Based on the prior work of Greenup et al.¹² high-volume hospitals were defined as facilities that evaluate/treat ≥ 298 patients a year. To define hospital volumes, Greenup et al.¹² used a

multivariable Cox proportional hazards model with restricted cubic splines to account for the nonlinearity of the relationship between hospital volume and survival. Women with inflammatory breast cancer (T4d) and women who were diagnosed on autopsy were excluded.

2.3 | Low value surgical care

This study focused on three established low-value procedures as defined by the Society of Surgical Oncology, the American College of Surgeons, and the American Society of Breast Surgery. The procedures include (1) sentinel lymph node biopsy (SLNB) among patients age ≥ 70 with clinically T1N0 hormone receptor–positive cancers; (2) axillary lymph node dissection (ALND) among patients with clinically T1–2N0 breast cancer with ≤ 2 positive sentinel lymph nodes who undergo breast conservation therapy (BCT). Performance of ALND among patients meeting ACOSOG Z0011 criteria was evaluated from 2012 to 2016 as the ACOSOG Z0011 trial was published in 2011¹⁵; (3) contralateral prophylactic mastectomy (CPM) among women presenting with unilateral breast cancers.^{16–18}

Of note, Choose Wisely guidelines for the omission of SLNB among women age ≥ 70 with small hormone–positive cancers and avoidance of CPM for unilateral cancer were published in 2016.^{19,20} Consequently, our study timeframe includes the period before wide public dissemination of the guidelines. Nonetheless, it should be noted that the data supporting the guidelines were available before 2016.

2.4 | Time to treatment

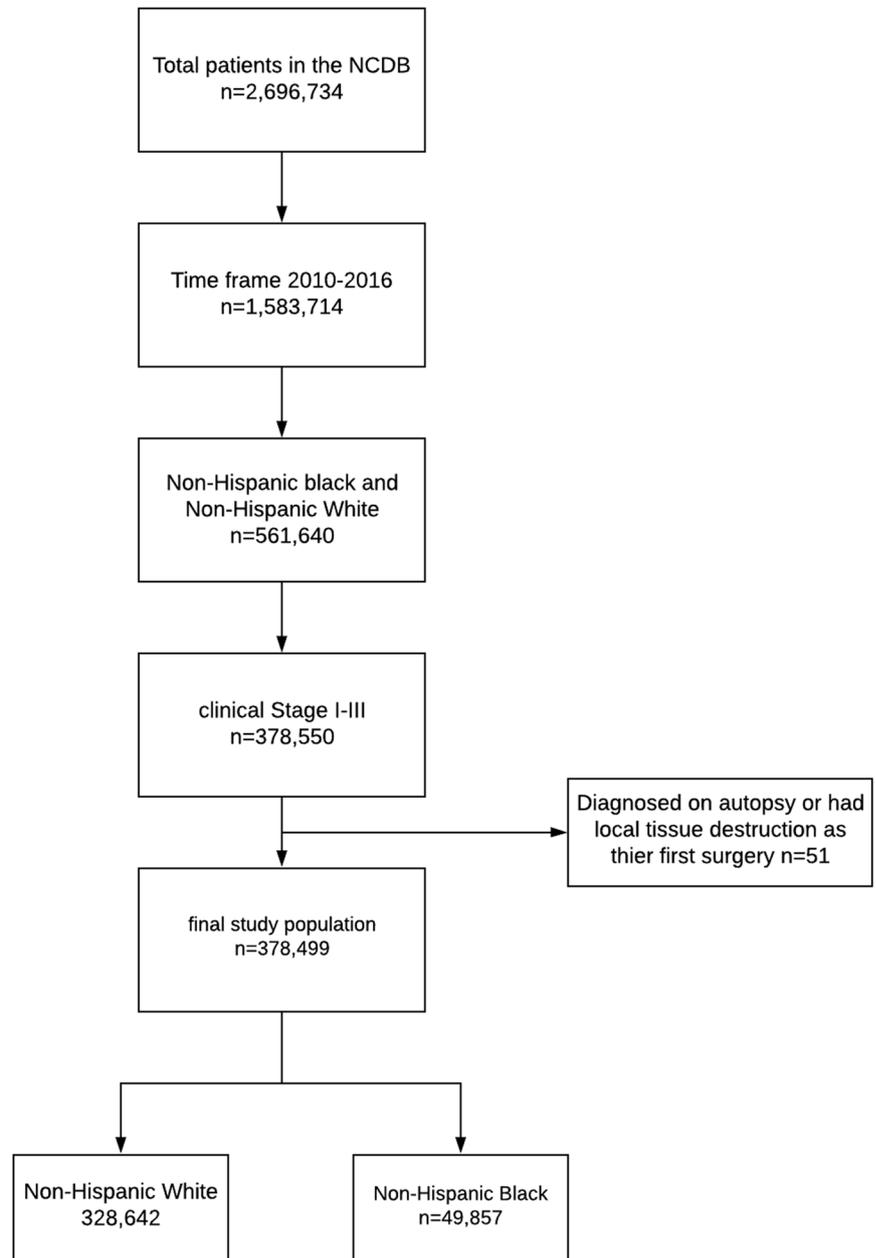
Based on the prior work of Bleicher et al,²¹ surgical delay was defined as the time from biopsy-proven diagnosis to first definitive surgery of >60 days. Patients who received neoadjuvant chemotherapy as their first definitive treatment were omitted from the time to surgery (TTS) analysis as a receipt of neoadjuvant chemotherapy delays time from biopsy to surgery. All patients who had the date of biopsy and surgery on the same date were also excluded due to concerns these cases were also excisional biopsies.

2.5 | Statistical analysis

The study population was dichotomized into NHB women and NHW women. Descriptive analysis of the two groups was conducted with categorical variables calculated as frequencies and continuous variables as medians or means. Sociodemographic, clinical, and treatment characteristics were compared between the two groups using the chi-square test for categorical variables and analysis of variance for continuous variables.

A multivariable logistic regression model was used to evaluate the probability of surgical delay. To account for patients being treated in the same facility we performed clustering at the hospital level.²² Variables in the model included insurance status (not insured, private

FIGURE 1 Study population diagram.
NCDB, National Cancer Database



insurance/managed care, medicaid, medicare, other government, unknown), age (continuous), race (NHB women, NHW women), year of diagnosis (2010–2016), clinical-stage (I–III), comorbidities (0, 1, ≥ 2), education (percentage of people in neighborhood with high school degree, $\geq 21\%$, 13%–20.9%, 7%–12.9%, or $< 7\%$), metropolitan status (metropolitan area of 1M population, metropolitan area of 250K–1M population, metropolitan area of $< 250\text{K}$ population, urban, or rural), surgery type (breast conservation surgery [BCS] vs. mastectomy), reconstruction (yes vs. no), biopsy and surgery facility, and distance traveled to the surgery facility (miles). Patients who received their biopsy and surgery at the same facility were categorized as same facility. Conversely, patients who had their biopsy and surgery in different facilities were categorized as different facility. Distance to the facility, per the NCDB, was based on the distance from the patient's

residence to the hospital in miles.²³ Variables in the model were selected due to their potential relationship to treatment delay.^{6,24–31} All statistical analyses were performed in Stata software version 16.0 with *p* values obtained from a two-tailed test. A *p* value of .05 was considered significant. The Ohio State University Office of Responsible Research Practices deemed this study IRB exempt.

3 | RESULTS

3.1 | Description of study population

There were 378,499 patients who met the study criteria. Overall, NHB women were significantly younger (mean age 58.2 years ± 13.3

vs. 61.4 years \pm 13.3; $p < .001$), more likely to live in a neighborhood with a low median income (neighborhood income $<$ \$38,000, 36.6% 18232/49765 vs. 9.1% 29984/327971; $p < .001$), had lower educational achievement (percent of individuals in neighborhood who did not graduate from high school \geq 21%, 29.0% 14455/49799 vs. 8.8% 28729/328198; $p < .001$) and resided in a large metropolitan area (77.5% 38086/49133 vs. 64.4% 205785/319750; $p < .001$) compared with NHW women (Table 1). In addition, a higher percentage of NHB women (11.4% 5902/49857) were on Medicaid than NHW women (3.5% 11676/328642; $p < .001$).

Significantly more NHB women had ($p < .001$) advanced stages of disease at diagnosis compared NHW women (Table 1). Aggressive cancer subtypes such as Triple Negative Breast Cancer (TNBC) and HER 2+ breast cancer were significantly ($p < .001$) more common among NHB compared to NHW women. Consequently, more NHB women received chemotherapy than NHW women (54.9% 26807/48787 vs 40.3% 129749/322161 $p < .001$). However, despite differences in tumor subtype, there were no differences between the groups on pathologic response after chemotherapy ($p = .23$). NHB women were less likely to have surgery compared to NHW (21.4% 10683/49857 vs 18.1% 59383/328642; $p < .001$). On subset analysis the most common reasons for no surgery included 1) not part of the planned first treatment course and 2) it was recommended by the physician but refused by the patients (not shown). The omission of surgery secondary to patient risk factors was similar between the two groups (not shown). There was no difference in the utilization of BCT (BCS+ radiation) between NHB and NHW women ($p = .55$).

3.2 | TTS and low-value procedures

Median time from biopsy proven diagnosis to first definitive surgery (Table 2) was significantly longer ($p < .001$) for NHB women (43 days [IQR, 28–68]) compared with NHW women (35 days [24–52]). Moreover, 30.6% (9031/29559) of NHB had surgery $>$ 60 days after biopsy compared to 18.0% (39692/219888) of NHW women ($p < .001$).

The groups were similar on the performance of ALND among women meeting criteria for ASOCOG Z0011 ($p = .13$). However, NHB women age \geq 70 with T1 hormone-positive cancers were less likely to have had an SLNB than NHW women (NHB 58.5% 1428/2442 vs. NHW 62.2% 19614/31547; $p < .001$). Moreover, among patients with unilateral breast cancer NHW had higher rates of CPM (36% 38028/105601) than NHB (26.3% 3822/14512) ($p < .001$). A review of CPM over the study period revealed compared to NHB women, NHW women underwent more CPMs during the entire study period (Figure 2).

3.3 | Multivariable analysis

In the multivariable logistic regression analysis, NHB race (odds ratio [OR], 1.77; 95% confidence interval [CI], 1.64–1.91), uninsured status

(OR, 1.61; 95% CI, 1.31–1.96), Medicaid insurance (OR, 1.72; 95% CI, 1.59–1.86), mastectomy (OR, 1.62; 95% CI, 1.53–1.73), later stage at diagnosis (Stage II OR, 1.5; 95% CI, 1.39–1.54, Stage III 3.52 95% CI, 3.17–3.91), biopsy and surgery in different facilities (OR, 1.44; 95% CI, 1.33–1.58) and reconstruction (OR, 1.31; 95% CI, 1.21–1.42) were associated with an increased odds of first definitive surgery of $>$ 60 days after biopsy compared to NHW race, private insurance, BCS, stage I, biopsy and surgery in the same facility and no reconstruction, respectively (Table 3).

4 | DISCUSSION

The results of this study suggest NHB women receiving surgical management at high-volume hospitals are less likely to receive certain low-value breast surgical procedures than NHW women. Specifically, NHB women have lower rates of CPM for unilateral disease and SLNB among women age \geq 70 with small hormone-positive cancers. Unfortunately, NHB breast cancer patients face significant delays in TTS compared to their NHW counterparts and are also more likely to have surgical management omitted. Notably, the omission of ALND among patients meeting ACOSOG Z0011 criteria was similar between both groups of women. We anticipate study findings of lower rates of low-value surgical care among NHB women are most likely secondary to established racial disparities in surgical care rather than disparities in the implementation of guidelines.

When viewed through the lens of receipt of guideline-concordant care, study findings of lower rates of low-value procedures among NHB women are meaningful. Historically, NHB breast cancer patients have been less likely to receive guideline-concordant locoregional and systemic management such as BCT or chemotherapy, respectively, and have higher rates of nonadherence to endocrine therapy.^{32,33} The omission of low-value procedures is consequential as the performance of these procedures does not change the utilization of adjunct treatment modalities, such as chemotherapy or radiation therapy, nor do they improve outcomes including survival.³⁴ In addition, the performance of these procedures adds unnecessary surgical complication risks with no discernable clinical benefit.

The results on low-value care should be interpreted with caution as Choose Wisely guidelines concerning SLNB in women age \geq 70 with small hormone-positive tumors and CPM for unilateral breast cancer were published toward the end of the study period. We anticipate the lower rates of CPM among NHB women in this study is most likely a reflection of higher BCS rates among the population and lower perceived risk of second cancer compared to NHW women.^{35,36} A higher perceived risk of second cancer has been implicated as a factor in decision making for CPM.³⁶ Other factors that may influence the decision for CPM include lack of physician referrals to reconstructive surgeons and the geographic unavailability of reconstructive surgeons.^{37–39} Black race and Medicaid insurance have both been associated with diminished access and utilization of reconstruction.⁴⁰ Consequently, we believe the racial differences in

TABLE 1 Sociodemographic and clinical variables by race*

Variable	Total N = 378,499, N (%)	Non-Hispanic N = 328,642, N (%)	Non-Hispanic N = 49,857, N (%)	P value
Age (continuous) years				<.001
Mean (SD)	61.06 (13.18)	61.44 (13.13)	58.52 (13.27)	
Age				<.001
≤40	22,419 (5.9)	18,125 (5.5)	4294 (8.6)	
41–50	64,142 (16.9)	54,041 (16.4)	10,101 (20.3)	
51–60	94,224 (24.9)	80,619 (24.5)	13,605 (27.3)	
61–70	104,426 (22.6)	92,078 (28.0)	12,348 (24.8)	
71–80	64,234 (17.0)	57,519 (17.5)	6715 (13.5)	
≥81	29,054 (7.7)	26,260 (8.0)	2794 (5.6)	
Income				<.001
<\$38,000	48,216 (12.8)	29,984 (9.1)	18,232 (36.6)	
\$38,000–\$47,999	66,688 (17.7)	56,200 (17.1)	10,488 (21.1)	
\$48,000–\$62,999	96,120 (25.4)	85,838 (26.2)	10,282 (20.7)	
\$63,000+	166,712 (44.1)	155,949 (47.5)	10,763 (21.6)	
Education (% no. high school degree)				<.001
≥21%	43,184 (11.4)	28,729 (8.8)	14,455 (29.0)	
13%–20.9%	80,612 (21.3)	63,658 (19.4)	16,954 (34.1)	
7%–12.9%	122,491 (32.4)	110,382 (33.6)	12,109 (24.3)	
<7%	131,546 (34.8)	125,285 (38.2)	6,261 (12.6)	
Distance to facility (miles)	30.38 (116.19)	32.31 (121.20)	17.67 (73.98)	<.001
Facility type				<.001
Comprehensive community cancer program	114,293 (31.8)	102,876 (32.8)	11,417 (24.7)	
Teaching/research	191,037 (53.1)	163,577 (52.2)	27,460 (59.4)	
Integrated network	54,511 (15.1)	47,171 (15.0)	7340 (15.9)	
Rural/urban				<.001
Metro_1M	243,871 (66.1)	205,785 (64.4)	38,086 (77.5)	
Metro 250K–1M	75,767 (20.5)	68,350 (21.4)	7417 (15.1)	
Metro <250K	15,940 (4.3)	14,348 (4.5)	1592 (3.2)	
Urban	29,931 (8.1)	28,114 (8.8)	1817 (3.7)	
Rural	3354 (0.9)	3153 (1.0)	201 (0.4)	
Region				<.001
New England	23,920 (6.6)	22,712 (7.2)	1208 (2.6)	
Middle Atlantic	67,035 (18.6)	57,539 (18.3)	9496 (20.5)	
South Atlantic	84,497 (23.5)	69,314 (22.1)	15,183 (32.9)	
East North Central	61,822 (17.2)	53,425 (17.0)	8397 (18.2)	
East South Central	18,939 (5.3)	15,570 (5.0)	3369 (7.3)	
West North Central	26,040 (7.2)	24,477 (7.8)	1563 (3.4)	
West South Central	29,828 (8.3)	25,429 (8.1)	4399 (9.5)	
Mountain	14,067 (3.9)	13,647 (4.4)	420 (0.9)	
Pacific	33,693 (9.4)	31,511 (10.0)	2182 (4.7)	
Year of diagnosis				.048
2010	45,976 (12.1)	40,074 (12.2)	5902 (11.8)	
2011	49,550 (13.1)	43,061 (13.1)	6489 (13.0)	
2012	52,116 (13.8)	45,382 (13.8)	6734 (13.5)	
2013	55,053 (14.5)	47,721 (14.5)	7332 (14.7)	

TABLE 1 (Continued)

Variable	Total N = 378,499, N (%)	Non-Hispanic N = 328,642, N (%)	Non-Hispanic N = 49,857, N (%)	P value
2014	57,229 (15.1)	49,665 (15.1)	7564 (15.2)	
2015	59,240 (15.7)	51,369 (15.6)	7871 (15.8)	
2016	59,335 (15.7)	51,370 (15.6)	7965 (16.0)	
Insurance				<.001
Not insured	4235 (1.1)	2934 (0.9)	1301 (2.6)	
Private Insurance/managed care	205,542 (54.3)	180,640 (55.0)	24,902 (49.9)	
Medicaid	17,380 (4.6)	11,676 (3.6)	5704 (11.4)	
Medicare	140,059 (37.0)	123,895 (37.7)	16,164 (32.4)	
Other government	3555 (0.9)	2987 (0.9)	568 (1.1)	
Unknown	7728 (2.0)	6510 (2.0)	1218 (2.4)	
Median follow-up (days)	40.25	40.67	37.62	<.001
IQR	(22.80–60.90)	(23.16–61.21)	(20.63–58.81)	
Clinical stage				<.001
1	234,849 (62.0)	209,704 (63.8)	25,145 (50.4)	
2	119,516 (31.6)	100,111 (30.5)	19,405 (38.9)	
3	24,121 (6.4)	18,817 (5.7)	5304 (10.6)	
Pathologic Stage				<.001
0	13,917 (4.1)	11,247 (3.8)	2670 (6.1)	
1	193,936 (56.8)	173,280 (58.1)	20,656 (47.5)	
2	104,146 (30.5)	88,898 (29.8)	15,248 (35.1)	
3	28,820 (8.4)	24,060 (8.1)	4760 (11.0)	
4	747 (0.2)	636 (0.2)	111 (0.3)	
Pathologic response				.230
None	3,021 (9.8)	2425 (9.6)	596 (10.3)	
Partial	15,646 (50.5)	12,766 (50.6)	2880 (50.0)	
Complete	12,300 (39.7)	10,014 (39.7)	2286 (39.7)	
Tumor size (cm)				<.001
≤2	197,524 (52.2)	175,692 (53.5)	21,832 (43.8)	
2 to ≤5	94,263 (24.9)	79,374 (24.2)	14,889 (29.9)	
≥5	25,602 (6.8)	20,786 (6.3)	4816 (9.7)	
Unknown	61,110 (16.1)	52,790 (16.1)	8320 (16.7)	
Hormone receptor status				<.001
ER+/PR- or +/-HER 2-	261,489 (76.0)	233,649(78.0)	27,840 (62.6)	
ER+/PR+/HER 2+	25,238 (7.3)	21,853 (7.3)	3385 (7.6)	
ER-/PR-/HER 2-	43,343 (12.6)	32,576 (10.9)	10,767 (24.2)	
ER-/PR-/HER 2+	14,064 (4.1)	11,561 (3.9)	2503 (5.6)	
Grade				<.001
Well differentiated	80,529 (21.3)	73,988 (22.5)	6,541 (13.1)	
Moderately differentiated	164,017 (43.3)	145,832 (44.4)	18,185 (36.5)	
Poorly differentiated	111,016 (29.3)	89,132 (27.1)	21,884 (43.9)	
Undifferentiated, anaplastic	595 (0.2)	497 (0.2)	98 (0.2)	
Cell type not determined	22,342 (5.9)	19,193 (5.8)	3149 (6.3)	
Histology				<.001
Ductal	292,982 (77.4)	252,945 (77.0)	40,037 (80.3)	
Lobular	39,820 (10.5)	35,867 (10.9)	3,953 (7.9)	
Mixed (ductal+lobular)	23,503 (6.2)	21,282(6.5)	2,221 (4.5)	
Other	22,194 (5.9)	18,548 (5.6)	3,646 (7.3)	

(Continues)

TABLE 1 (Continued)

Variable	Total N = 378,499, N (%)	Non-Hispanic N = 328,642, N (%)	Non-Hispanic N = 49,857, N (%)	P value
Chemotherapy				<.001
No	214,392 (57.8)	192,412 (59.7)	21,980 (45.1)	
Yes	156,556 (42.2)	129,749 (40.3)	26,807 (54.9)	
Neoadjuvant chemotherapy				<.001
No	309,269 (88.3)	272,543 (89.2)	36,726 (82.5)	
Yes	40,795 (11.7)	33,002 (10.8)	7793 (17.5)	
Immunotherapy				<.001
No	350,558 (92.6)	304,955 (92.8)	45,603 (91.5)	
Yes	27,035 (7.1)	22,940 (7.0)	4095 (8.2)	
Unknown	906 (0.2)	747 (0.2)	159 (0.3)	
Radiation				.025
No	157,572 (41.6)	136,982 (41.7)	20,590 (41.3)	
Yes	219,224 (57.9)	190,213 (57.9)	29,011 (58.2)	
Unknown	1703 (0.4)	1447 (0.4)	256 (0.5)	
Breast conservation therapy				.550
No	30,654 (17.7)	26,830 (17.7)	3824 (17.5)	
Yes	142,828 (82.3)	124,831 (82.3)	17,997 (82.5)	
Surgery				<.001
No surgery	70,066 (18.5)	59,383 (18.1)	10,683 (21.4)	
Partial mastectomy	173,958 (46.0)	152,100 (46.3)	21,858 (43.8)	
Mastectomy	134,134 (35.4)	116,855 (35.6)	17,279 (34.7)	
Unknown	341 (0.1)	304 (0.1)	37 (0.1)	
Time to surgery (days)				<.001
Mean (SD)	40.59 (39.30)	39.30 (37.01)	50.11 (52.30)	
Median (IQR)	33.00 (21.00–48.00)	32.00 (21.00–47.00)	37.00 (22.00–58.00)	
Patient treated at more than one CoC facility				<.001
Yes	83,246 (22.0)	73,540 (22.4)	9,706 (19.5)	
No	295,253 (78.0)	255,102 (77.6)	40,151 (80.5)	
Lymph node surgery				<.001
None (code 0)	26,296 (11.5)	22,248 (11.2)	4048 (13.5)	
Sentinel lymph node biopsy (code 2)	160,733 (70.1)	142,497 (71.5)	18,236 (61.0)	
Axillary lymph node dissection (code 6 and 7)	42,157 (18.4)	34,563 (17.3)	7594 (25.4)	
Comorbidities				<.001
0	318,866 (84.2)	280,038 (85.2)	38,828 (77.9)	
1	46,791 (12.4)	38,582 (11.7)	8209 (16.5)	
≥2	12,842 (3.4)	10,022 (3.0)	2820 (5.7)	

Abbreviations: CoC, Commission on Cancer; IQR, interquartile range.

*Data may not add up to 100% due to rounding.

CPM noted in this study are most likely driven by an interplay between patient preferences, lack of physician referrals, and access to reconstructive surgeons.

Concerning omission of SLNB in women age ≥ 70 with small hormone-positive cancers, the most likely explanation is that these results are a reflection of underlying racial disparities in the

surgical management of the axilla rather than an earlier implementation of guidelines. Notably, the CALGB 9394 trial that served as the justification for omission of SLNB among women age ≥ 70 with small hormone-positive tumors was published in 2013.⁴¹ However, without information in the NCDB about physician knowledge or awareness of CALGB 9394 findings, the implications

TABLE 2 Differences in receipt of low-value surgical care practices by race*

	Total N = 378,499, N (%)	Non-Hispanic White N = 328,642, N (%)	Non-Hispanic Black N = 49,857, N (%)	P value
Time from biopsy to surgery (continuous)	36.00 (24.00–54.00)	35.00 (24.00–52.00)	43.00 (29.00–68.00)	<.001
Time from biopsy to surgery				<.001
≤60 days	200,724 (80.5%)	180,196 (81.9%)	20,528 (69.4%)	
>60 days	48,723 (19.5%)	39,692 (18.1%)	9031 (30.6%)	
ACOSOG Z0011 Criteria				.13
No	8493 (79.9%)	7504 (80.1%)	989 (78.3%)	
Yes	3133 (20.1%)	1859 (19.9%)	274 (21.7%)	
Contralateral prophylactic mastectomy				<.001
No	78,263 (65.2%)	67,573 (64%)	10,690 (73.7%)	
Yes	41,850 (34.8%)	38,028 (36.0%)	3822 (26.3%)	
Sentinel lymph node surgery age ≥70				<.001
No	12,947 (38.1%)	11,933 (37.8%)	1014 (41.5%)	
Yes	21,042 (61.9%)	19,614 (62.2%)	1428 (58.5%)	

*Data may not add up to 100% due to rounding.

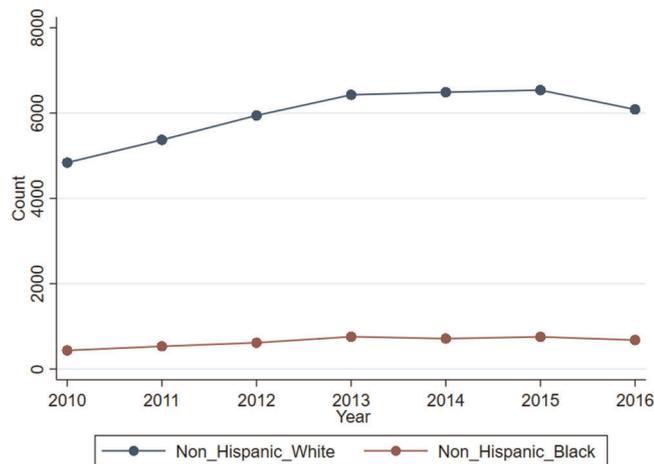


FIGURE 2 Distribution of contralateral prophylactic mastectomy [Color figure can be viewed at wileyonlinelibrary.com]

of the study results on surgical decision making are difficult to surmise.

Higher rates of surgical delay among NHB patients in this study are consistent with previous studies.⁶ Surgical delay is probably attributable to a mixture of institutional and patient-related factors. For example, NHB women receiving treatment at high-volume centers in the NCDB may be experiencing higher rates of transportation issues, financial toxicity, insurance issues, difficulty securing child-care, and lack of time off from work which are all factors that could influence delay.^{42,43} In addition, Black patients tend to be less knowledgeable about their disease and are subjected to physician–patient relationships in which communication styles prohibits cultural and linguistically appropriate information sharing

TABLE 3 Multivariable analysis evaluating time to surgery*

Variable	Odds ratio	Confidence interval	P value
Age, years (continuous)	0.99	(0.98–0.99)	<.001
Insurance			
Not insured	1.61	(1.31–1.96)	<.001
Private	Ref		
Medicaid	1.72	(1.59–1.86)	<.001
Medicare	1.00	(0.95–1.04)	.933
Other government	1.16	(1.00–1.35)	.048
Race			
Non-Hispanic White	Ref		
Non-Hispanic Black	1.77	(1.64–1.91)	<.001
Year of diagnosis			
2010	Ref		
2011	1.09	(1.03–1.16)	.002
2012	1.12	(1.03–1.20)	.004
2013	1.20	(1.11–1.29)	<.001
2014	1.26	(1.16–1.37)	<.001
2015	1.46	(1.33–1.61)	<.001
2016	1.55	(1.41–1.71)	<.001
Clinical stage			
1	Ref		
2	1.46	(1.39–1.54)	<.001
3	3.52	(3.17–3.91)	<.001
Receptor status			
ER+/PR– or+/HER 2–	Ref		
ER+/PR+/HER 2+	1.32	(1.24–1.40)	<.001
ER–/PR–/HER 2–	1.13	(1.07–1.20)	<.001
ER–/PR–/HER 2 +	1.38	(1.27–1.50)	<.001

(Continues)

TABLE 3 (Continued)

Variable	Odds ratio	Confidence interval	P value
Comorbidities			
0	Ref		
1	0.99	(0.94–1.03)	.627
≥2	1.09	(1.01–1.17)	.026
% No. high school degree			
≥21%	Ref		
13%–20.9%	0.92	(0.86–0.99)	.044
7%–12.9%	0.91	(0.83–0.99)	.044
<7%	0.84	(0.76–0.93)	.001
Surgery type			
Mastectomy	1.62	(1.53–1.72)	<.001
Partial mastectomy	Ref		
Reconstruction			
Yes	1.31	(1.21–1.42)	<.001
No	Ref		
Biopsy+surgery facility			
Same	Ref		
Different	1.44	(1.32–1.57)	<.001
Distance traveled (continuous) miles	1.00	(0.99–1.00)	.226

*This table evaluates the probability of surgical delay (probability of surgery >60 days vs probability ≤60 days).

which diminishes patient self-advocacy.^{44,45} This interplay between NHB patient's socioeconomic position and the deficits of the doctor–patient relationship could be adversely affecting the timeliness of care. Nevertheless, these findings are worrisome, as surgical delay >60 days has been implicated in higher overall and disease-specific mortality.⁷ Study findings of an association between surgical delay and Medicaid or uninsured status, mastectomy, later stage of diagnosis, reconstruction and biopsy, and surgery in different facilities are consistent with prior studies in the literature.^{26,46}

The findings from this study highlight some of the ongoing surgical disparities NHB breast cancer patients face compared to their NHW counterparts when receiving care at high-volume hospitals. Although lower rates of low-value surgery and similarities in receipt of BCT show adherence to guideline-concordant locoregional management, higher rates of surgical delay and omission of surgery are worrisome. Furthermore, when placed within the context of a study timeframe before the publication of some of the low-value guidelines these results underscore persistent racial differences in the management of the breast and axilla. The benefits of study findings of having lower rates of low-value surgical procedures on outcomes, such as mortality or recurrence, are unclear and warrants further investigation.

The limitations of this study include a lack of generalizability of study results beyond CoC accredited hospitals. In addition, receipt of care at high-volume hospitals may not be reflective of receipt of care

at all institutions across the United States both within and outside of the NCDB. The NCDB does not collect data on genetic information; therefore, patients who received CPM due to genetic mutations were not identified. As of the writing of this manuscript, the NCDB only has data available until 2016, therefore, part of the study timeframe was before publication and dissemination of some of the low-value surgical procedure guidelines. This data restriction makes it difficult to extrapolate the implications of these results on post-publication practices. The strengths of the study include the large sample size and the fact that the NCDB represents approximately 70% of newly diagnosed cancer cases in the United States.¹⁴

5 | CONCLUSION

Results from this study suggest there are racial disparities in surgical management among patients receiving care at high-volume CoC Centers. These results infer that NHB breast cancer patients receiving care at high-volume facilities face barriers in receiving timely surgical care. Future studies will need to discern if the omission of low-value surgical care is reducing surgical complications rates among NHB women and delineate how this may translate into improved clinical outcomes such as survival in this population. Moreover, more granular studies are needed to better understand the drivers of surgical delay among NHB breast cancer patient receiving care at high-volume hospitals.

ACKNOWLEDGMENTS

The authors would like to acknowledge Brittany Bernardo, PhD and Toyin Adeyanju, MPH for their support in the preparation of this manuscript. The Ohio State University Comprehensive Cancer Center Undergraduate Summer Research Internship Program in Disparities, funded by Pelotonia supported Ms. Jackson.

FUNDING INFORMATION

Support for this study was provided by the Ohio State University Comprehensive Cancer Center using Pelotonia funds.

DISCLOSURES

Destiny Jackson declares that she has no conflict of interest. Yaming Li declares that she has no conflict of interest. Miriam Eskander declares that she has no conflict of interest. Bridget Oppong declares that she has no conflict of interest. Allan Tsung declares that he has no conflict of interest. Oindrila Bhattacharyya declares that she has no conflict of interest. Electra Paskett declares that she has no conflict of interest. Samilia Obeng-Gyasi declares that she has no conflict of interest.

DATA AVAILABILITY

The National Cancer Database (NCDB) is a joint project of the Commission on Cancer (CoC) of the American College of Surgeons and the American Cancer Society. The CoC's NCDB and the hospitals participating in the CoC's NCDB are the sources of the deidentified

data used herein; they have not verified and are not responsible for the statistical validity of the data analysis or the conclusions derived by the authors.

ORCID

Samilia Obeng-Gyasi  <http://orcid.org/0000-0002-5330-7247>

REFERENCES

- DeSantis CE, Ma J, Gaudet MM, et al. Breast cancer statistics, 2019. *CA Cancer J Clin*. 2019;69(6):438-451.
- American Cancer Society. Survival Rates for Breast Cancer. 2019; <https://www.cancer.org/cancer/breast-cancer/understanding-a-breast-cancer-diagnosis/breast-cancer-survival-rates.html>. Accessed 05/07/2018, 2019.
- Newman LA, Mason J, Cote D, et al. African-American ethnicity, socioeconomic status, and breast cancer survival: a meta-analysis of 14 studies involving over 10,000 African-American and 40,000 White American patients with carcinoma of the breast. *Cancer*. 2002;94(11):2844-2854.
- Newman LA, Kaljee LM. Health disparities and triple-negative breast cancer in African American women: a review. *JAMA Surg*. 2017;152(5):485-493.
- Daly B, Olopade OI. A perfect storm: how tumor biology, genomics, and health care delivery patterns collide to create a racial survival disparity in breast cancer and proposed interventions for change. *CA Cancer J Clin*. 2015;65(3):221-238.
- Fedewa SA, Edge SB, Stewart AK, Halpern MT, Marlow NM, Ward EM. Race and ethnicity are associated with delays in breast cancer treatment (2003-2006). *J Health Care Poor Underserved*. 2011;22(1):128-141.
- Bleicher RJ, Ruth K, Sigurdson ER, et al. Time to surgery and breast cancer survival in the United States. *JAMA Oncol*. 2016;2(3):330-339.
- Roberts MC, Wheeler SB, Reeder-Hayes K. Racial/Ethnic and socioeconomic disparities in endocrine therapy adherence in breast cancer: a systematic review. *Am J Public Health*. 2015;105(Suppl 3):e4-e15.
- Chang CM, Huang KY, Hsu TW, et al. Multivariate analyses to assess the effects of surgeon and hospital volume on cancer survival rates: a nationwide population-based study in Taiwan. *PLoS One*. 2012;7(7):e40590.
- Chang CM, Yin WY, Wei CK, et al. The association of socioeconomic status and access to low-volume service providers in breast cancer. *PLoS One*. 2013;8(12):e81801.
- Kingsmore D, Hole D, Gillis C. Why does specialist treatment of breast cancer improve survival? The role of surgical management. *Br J Cancer*. 2004;90(10):1920-1925.
- Greenup RA, Obeng-Gyasi S, Thomas S, et al. The Effect of hospital volume on breast cancer mortality. *Ann Surg*. 2018;267(2):375-381.
- Wang T, Baskin AS, Dossett LA. Deimplementation of the choosing wisely recommendations for low-value breast cancer surgery: a systematic review. *JAMA Surg*. 2020;155:759.
- American College of surgeons. National Cancer Database. 2020; <https://www.facs.org/quality-programs/cancer/ncdb>. Accessed 8/17/2020, 2020.
- Giuliano AE. Axillary dissection vs no axillary dissection in women with invasive breast cancer and sentinel node metastasis: a randomized clinical trial. *JAMA*. 2011;305(6):569-575.
- Chagpar AB, Hatzis C, Pusztai L, et al. Association of LN Evaluation with survival in women aged 70 years or older with clinically node-negative hormone receptor positive breast cancer. *Ann Surg Oncol*. 2017;24(10):3073-3081.
- Albornoz CR, Matros E, Lee CN, et al. Bilateral mastectomy versus breast-conserving surgery for early-stage breast cancer: the role of breast reconstruction. *Plast Reconstr Surg*. 2015;135(6):1518-1526.
- Mann JM, Wu X, Christos P, Nagar H. The state of surgical axillary management and adjuvant radiotherapy for early-stage invasive breast cancer in the modern era. *Clin Breast Cancer*. 2018;18(4):e477-e493.
- Choosing wisely: An initiative of the ABIM Foundation ASoBS. Don't routinely perform a double mastectomy in patients who have a single breast with cancer. 2016; <https://www.choosingwisely.org/clinician-lists/breast-surgeons-mastectomies-for-single-breast-cancer-patients/>. Accessed 08/09/2020, 2020.
- Choosing wisely: An initiative of the ABIM Foundation. Don't routinely use sentinel node biopsy in clinically node negative women ≥70 years of age with hormone receptor positive invasive breast cancer. 2016; <http://www.choosingwisely.org/clinician-lists/sso-sentinel-node-biopsy-in-node-negative-women-70-and-over/>. Accessed 08/13/2018, 2018.
- Bleicher RJ, Ruth K, Sigurdson ER, et al. Time to Surgery and Breast Cancer Survival in the United States. *JAMA Oncol*. 2015;1:10.
- Abadie A, Athey S, Imbens G, Wooldridge J. When Should You Adjust Standard Errors for Clustering? *NBER Work Pap Ser*. 2017; <https://economics.mit.edu/files/13927>. Accessed 09/10/19.
- The American College of Surgeons. NCDB Data Dictionary. *Medicaid Expansion Status State Group*. 2019. <http://ncdbpuf.facs.org/node/433>. Accessed 5/10/2019.
- Bilimoria KY, Ko CY, Tomlinson JS, et al. Wait times for cancer surgery in the United States: trends and predictors of delays. *Ann Surg*. 2011;253(4):779-785.
- Bleicher RJ, Ciocca RM, Egleston BL, et al. Association of routine pretreatment magnetic resonance imaging with time to surgery, mastectomy rate, and margin status. *J Am Coll Surg*. 2009;209(2):180-187.
- Bleicher RJ, Ruth K, Sigurdson ER, et al. Preoperative delays in the US Medicare population with breast cancer. *Journal of clinical oncology: official journal of the American Society of Clinical Oncology*. 2012;30(36):4485-4492.
- Chaudhry R, Goel V, Sawka C. Breast cancer survival by teaching status of the initial treating hospital. *CMAJ: Canadian Medical Association journal=journal de l'Association Medicale Canadienne*. 2001;164(2):183-188.
- George P, Chandwani S, Gabel M, et al. Diagnosis and surgical delays in African American and white women with early-stage breast cancer. *J Womens Health (Larchmt)*. 2015;24(3):209-217.
- Hulvat M, Sandalow N, Rademaker A, Helenowski I, Hansen NM. Time from diagnosis to definitive operative treatment of operable breast cancer in the era of multimodal imaging. *Surgery*. 2010;148(4):746-750.
- Jones AP, Haynes R, Sauerzapf V, Crawford SM, Zhao H, Forman D. Travel time to hospital and treatment for breast, colon, rectum, lung, ovary, and prostate cancer. *European journal of cancer (Oxford, England: 1990)*. 2008;44(7):992-999.
- Liederbach E, Sisco M, Wang C, et al. Wait times for breast surgical operations, 2003-2011: a report from the National Cancer Data Base. *Ann Surg Oncol*. 2015;22(3):899-907.
- Roberts MC, Wheeler SB, Reeder-Hayes K. Racial/ethnic and socioeconomic disparities in endocrine therapy adherence in breast cancer: a systematic review. *Am J Public Health*. 2015;105(S3):e4-e15.
- Bickell NA, Wang JJ, Oluwole S, et al. Missed opportunities: racial disparities in adjuvant breast cancer treatment. *J Clin Oncol*. 2006;24(9):1357-1362.
- Greenup RA, Prakash I, Sorenson C. "Choosing wisely" in breast cancer surgery: drivers of low value care. *Ann Surg Oncol*. 2020;27(8):2577-2579.
- Thomas P, Killelea BK, Horowitz N, Chagpar AB, Lannin DR. Racial differences in utilization of breast conservation surgery: results

- from the National Cancer Data Base (NCDB). *Ann Surg Oncol*. 2016; 23(10):3272-3283.
36. Kim Y, McCarthy AM, Bristol M, Armstrong K. Disparities in contralateral prophylactic mastectomy use among women with early-stage breast cancer. *NPJ Breast Cancer*. 2017;3:2.
 37. Soran A, Kamali Polat A, Johnson R, McGuire KP. Increasing trend of contralateral prophylactic mastectomy: what are the factors behind this phenomenon? *Surgeon*. 2014;12(6):316-322.
 38. Tseng JF, Kronowitz SJ, Sun CC, et al. The effect of ethnicity on immediate reconstruction rates after mastectomy for breast cancer. *Cancer*. 2004;101(7):1514-1523.
 39. Bauder AR, Gross CP, Killelea BK, Butler PD, Kovach SJ, Fox JP. The relationship between geographic access to plastic surgeons and breast reconstruction rates among women undergoing mastectomy for cancer. *Ann Plast Surg*. 2017;78(3):324-329.
 40. Kruper L, Holt A, Xu XX, et al. Disparities in reconstruction rates after mastectomy: patterns of care and factors associated with the use of breast reconstruction in Southern California. *Ann Surg Oncol*. 2011;18(8):2158-2165.
 41. Hughes KS, Schnaper LA, Bellon JR, et al. Lumpectomy plus tamoxifen with or without irradiation in women age 70 years or older with early breast cancer: long-term follow-up of CALGB 9343. *J Clin Oncol*. 2013;31(19):2382-2387.
 42. Artiga Samantha HE. Beyond Health Care: The Role of Social Determinants in Promoting Health and Health Equity. 2018. <https://www.kff.org/disparities-policy/issue-brief/beyond-health-care-the-role-of-social-determinants-in-promoting-health-and-health-equity/>. Accessed 10/15/2019, 2019.
 43. Carrera PM, Kantarjian HM, Blinder VS. The financial burden and distress of patients with cancer: Understanding and stepping-up action on the financial toxicity of cancer treatment. *CA Cancer J Clin*. 2018;68(2):153-165.
 44. Freedman RA, Kouri EM, West DW, Keating NL. Racial/ethnic disparities in knowledge about one's breast cancer characteristics. *Cancer*. 2015;121(5):724-732.
 45. Cooper LA, Beach MC, Johnson RL, Inui TS. Delving below the surface. Understanding how race and ethnicity influence relationships in health care. *J Gen Intern Med*. 2006;21 Suppl 1(Suppl 1): S21-S27.
 46. Bleicher RJ, Chang C, Wang CE, et al. Treatment delays from transfers of care and their impact on breast cancer quality measures. *Breast Cancer Res Treat*. 2018.

How to cite this article: Jackson DK, Li Y, Eskander MF, et al. Racial disparities in low-value surgical care and time to surgery in high-volume hospitals. *J Surg Oncol*. 2021;123: 676–686. <https://doi.org/10.1002/jso.26320>