Breast Cancer Racial Disparities: Unanswered Questions

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Running Head: Breast Cancer Disparities

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Abstract

Breast cancer is the most common non-cutaneous cancer diagnosed in women in the United States and is second only to lung cancer as the leading cause of cancer-related mortality. Although mortality rates have been dropping steadily due to a variety of factors including improved treatment modalities and screening, substantial racial differences in outcome between blacks and whites persist. Although differences in healthcare utilization and access, tumor biology, and cancer management have been elucidated as possible reasons for disparities seen it is likely that other interactions exist. The purpose of this review is therefore to present a comprehensive overview of the literature on racial disparities in breast cancer outcome and highlight potential causative factors that may contribute to disparities seen among blacks and whites with breast cancer. In addition, we make research recommendations by discussing some of the remaining gaps in knowledge that may lead to further understanding of disparities and consequently improved outcomes for all women with breast cancer.
Introduction

Projections indicate that approximately 207,000 new cases of breast cancer in the United States and 40,000 deaths will be reported in 2010 alone.(1) Although mortality rates have been dropping since 1990 due to earlier detection and improved treatment modalities, substantial racial disparities in outcome continue to persist. While the incidence rates for breast cancer in blacks are lower than whites (113 vs. 123.5 per 100,000), mortality rates are disproportionately higher (33 deaths vs. 23.9 per 100,000 cases).(1, 2) Although reasons for the racial disparities in breast cancer have not been fully elucidated, it has been suggested that factors such as differences in tumor biology, healthcare access, and disease management may contribute to the poorer outcomes observed. The purpose of this article is to present a comprehensive overview of the literature on racial disparities in breast cancer outcome and highlight potential causative factors that may contribute to disparities seen among blacks and whites with breast cancer. Specifically, we review the role of tumor biology, genetic variants in relation to survival disparities, as well as the impact of factors such as screening and detection and receipt of optimal care on poorer survival among black women with breast cancer. In addition, we discuss some of the remaining gaps in knowledge that may lead to further understanding of disparities and improved outcomes for all women with breast cancer.

Methods

A PubMed search was used to retrieve articles published in English between 1980 and 2010. MeSH terms used included “breast cancer,” “disparities,” “racial,” “mortality,” “incidence,” “treatment,” “screening,” “genetics,” “epidemiology,” and “pathology.”
also utilized editions of the SEER Cancer Statistics Reviews published by the National Cancer Institute in 2001 and 2007. All articles were peer-reviewed with the exception of a conference proceeding. Articles that focused primarily on other disadvantaged groups such as Hispanics were excluded.

Outcome disparities

A disproportionate number of deaths from breast cancer occur in blacks. While the overall incidence rates are higher in whites, blacks with breast cancer have a higher breast cancer specific mortality (33 vs. 23.9 per 100,000). In addition, mortality rates also differ by age-group. In 1980, blacks younger than 45 years had higher mortality rates as compared to whites in the same age group.(3) Between 1993 and 1996, breast cancer mortality rates among those aged 35 years and younger in blacks were 2 fold higher than the rates in whites. For those between 40 and 50 years the rates in blacks were 1.5 fold whites, while the two races had more or less similar outcomes for those 55 years and older.(4) The 5 year survival rates for all stages also differ by race with blacks being disadvantaged. For example, for loco-regional disease 5 year survival rates for blacks are only 72% vs. 85% for whites.(1) Table 1 gives a summary of differences in the epidemiology of breast cancer in blacks and whites.

Due to improved diagnostic and treatment modalities breast cancer mortality has declined over time, but decreases generally have been smaller and less consistent among blacks than whites.(5) In 1975, the age-adjusted death rate from breast cancer for all races was 31.4 per 100,000. By 1990, the rates had increased slightly to 33.1 deaths per 100,000, but subsequently decreased to 22.8 per 100,000 in 2007 reflecting a 27% decrease.
decline from 1975. Comparing death rates from 1975 to rates from 2007, whites have enjoyed a 30% (31.8 to 22.2 per 100,000) reduction in mortality while rates have actually increased by 6.4% (29.5 to 31.4 per 100,000) in blacks in the same period. If the most contemporary period is considered comparing rates in 2000 and 2007, whites have had a 15.3% reduction in mortality as opposed to 8.7% in blacks.

Factors that contribute to these trends are not fully understood, but thought to reflect population differences in tumor biology, genetic underpinnings of aggressive disease, detection, and potential racial differences in management.

**Racial differences in tumor biology**

Although socioeconomic factors and access to care are likely to play a role in poorer survival for black women, they are also more likely to be diagnosed at an earlier age than whites and to have tumors with more aggressive characteristics, which may also factor into poorer outcomes. Black women are frequently diagnosed with tumors with high histologic grade and negative estrogen receptor (ER), progesterone receptor (PR) and HER-2 receptor status, as well as high nuclear atypia, mitotic index, S-phase fraction, and tumor necrosis. (6, 7) They are also more likely to be diagnosed with basal-like breast cancers, which are negative for ER, PR and HER2, and overexpress cytokeratins 5/6 and HER1/EGFR, and have high proliferation rates.(8) This subtype has a less favorable prognosis than the others, particularly Luminal A subtype, which has the highest survival rates.(8) In the Carolina Breast Cancer Study (CBCS), the prevalence of basal-like breast cancer was highest (39%) among black premenopausal women compared to other groups of patients (14-16%).(8) This has been observed in other
studies in the US.(7, 9) It could be suggested that the preponderance of basal-like or hormone receptor negative tumors among black women could be due to their younger age at diagnosis, but even in older age groups, blacks are more likely than whites to present with triple negative disease.(10)

Until recently, the reasons for these differences in subtypes have been unknown. Historically, the majority of risk factors for breast cancer are related to lifetime exposures to estrogens and overall, parity has been associated with reduced risk of breast cancer.(11) In the CBCS, however, parity was only associated with reduced risk of luminal A subtype and, in fact, increased risk of basal-like breast cancer by almost twofold.(12) Of importance, the increased risk of basal-like breast cancer with increasing number of live births was totally ameliorated with breastfeeding. These associations have also been observed in subsequent studies in other populations.(13, 14) In the Black Women’s Health Study, similar associations with parity and breast feeding were observed in relation to age at onset, with higher parity noted to increase risk of breast cancer before age 45 among black women.(12, 15) Because black women are generally more likely to have a greater number of children at a younger age in the US, and not to breastfeed, these reproductive factors could account, in part, for the higher proportion of basal-like breast cancers in black women. However, because basal-like breast cancers are also more prevalent among women in Africa, a biological or genetic component cannot be ruled out.

It is well-established that African-Americans have lower circulating levels of serum vitamin D, perhaps due, in part, to higher skin pigmentation, reducing absorption of ultraviolet rays.(16) In cell culture and rodent models, vitamin D has shown anti-cancer effects.
cancer properties, but the epidemiologic support for a role in breast cancer risk reduction has been inconsistent. It is possible that vitamin D’s effects may be more important in modulating breast cancer aggressiveness and prognosis rather than overall cancer susceptibility. Indeed, mice lacking the vitamin D receptor are more likely to develop aggressive ER/PR negative mammary tumors than wild type mice;(17) and in human breast cancer cells, vitamin D inhibits the expression of a variety of basal/myoepithelial markers, reverting their trans-differentiation, related to the basal-like subtype.(18) Lower serum vitamin D has been associated with higher grade tumors(19) and, in a series of more than 500 women with breast cancer, we observed that serum 25OHD levels were lowest among women with basal-like breast cancers (Ambrosone et al.,unpublished). Thus, it is plausible that lower vitamin D could contribute to the higher proportion of basal-like tumors among black women. Clearly, genetic, biological and epidemiological risk factors for breast cancer subtypes, particularly among black women, needs to be further investigated. However, the observation in several studies that increased risk of basal-like breast cancer is associated with higher parity, and that breastfeeding reduces this risk, warrants further public health recommendations that breastfeeding be encouraged for all women and especially for black women to reduce risk of basal-like breast cancer. Additionally, although associations between vitamin D and breast cancer in black women need to be further investigated, maintenance of adequate vitamin D levels could possibly contribute to lowering risk of aggressive breast cancer in this population.

**Racial differences in germline genetic factors**
While certain founder mutations in *BRCA1* or *BRCA2* genes have been seen more commonly in particular groups such as Ashkenazi Jewish individuals, the spectrum of deleterious mutations appears to be different from that observed in black women. Although the frequency for both *BRCA1* and *BRCA2* mutations may be lower than that observed in white or Ashkenazi Jewish individuals with breast cancer,(20-23) a higher frequency of sequence variation of undetermined significance is seen in black women.(21) It is unclear whether these variants act to modify risk or disease aggressiveness. Breast cancer characteristics seen in blacks are similar to *BRCA1*-associated tumors. As promoter methylation may be a mechanism for inactivating *BRCA1* in sporadic tumors,(24) it is unclear what proportion of blacks with breast cancers have methylation of the *BRCA1* gene that may explain similarities between breast cancer in blacks and *BRCA1*-associated cancers.

Admixture mapping is a useful strategy for discovering disease causing genetic variants that differ in frequency across populations and contribute to complex traits such as oncogenesis. A large genome-wide admixture study in blacks with breast cancer from six population-based studies did not find an association between breast cancer risk and African or European heritage at any loci, suggesting that genetic ancestry may not contribute significantly to risk.(25) This is controversial however, given the striking similarities between breast cancer in blacks living in Africa and black Americans in the United States.(26) We have shown that blacks with breast cancer in Nigeria and the United States have differences in common breast cancer risk factors such as age at menarche, body mass index, number of pregnancies, age at first birth, and lactation duration, despite similar clinicopathologic features.(27) It is therefore plausible that...
genetic factors (and not reproductive and hormonal risk factors) common to the two
groups but different from those observed in whites may explain the biologically
aggressive disease observed in black women irrespective of location. This underscores
the need for pooling of epidemiological studies in black women to have adequate
statistical power to investigate complex interactions between genetic and non-genetic
factors in the etiology of aggressive breast cancer, and evaluation of these and factors
related to access and optimal care to disentangle predictors of poorer survival among
black women with breast cancer. Although polymorphisms have been extensively studied
in relation to risk of overall breast cancer in white women, there are very few studies in
black women, particularly for genetic polymorphisms that infer susceptibility to early
onset aggressive breast cancer.

Racial differences in detection

Although early detection with screening mammography has contributed to
decreases in breast cancer mortality; not all women have benefited equally. Even though
blacks are less likely to undergo regular mammograms than whites,(28) it is unlikely that
differences in detection account for the entirety of outcome disparities. It is difficult to
estimate the attributable risk to overall outcome due to multiple other factors that
influence outcome.

Factors responsible for differences in mammography rates are multifactorial and
include a lower likelihood of receiving a recommendation for mammography from a
health care provider, lack of insurance, or access to regular healthcare.(29, 30) Although
adapting for socioeconomic and insurance status has often resulted in reports that blacks
have greater odds of having a recent mammogram than whites, (31) multivariate models suggest that fewer blacks receive regular mammograms than whites, consequently continued effort is needed to reduce access barriers for blacks not regularly screened. Given guidelines that recommend women begin mammography at age 40 or 50, black women may be at increased risk for delayed diagnosis even if they are compliant with current screening guidelines as greater than 10% of breast cancers in blacks are diagnosed under the age of 40 years, compared to 5% of whites.(32) Therefore, the optimal age for breast cancer screening in blacks still needs to be defined. This is also particularly important given the disproportionate mortality in younger black women with breast cancer. If one considers the increased breast density in this group, mammography may not be the best choice. Perhaps race and/or age-directed screening with ultrasonography or even more sophisticated techniques with novel biomarkers ought to be developed in the future.

**Racial differences in management**

An important factor that also contributes to the racial gap in mortality is the observed difference in clinical management of the disease. It is plausible that black women have more comorbidities that preclude them from receiving established standards; or perhaps they are not receiving standard therapy due to cultural beliefs/distrust; or alternatively they have more problems tolerating equivalent doses of chemotherapy than their white counterparts. These are some potential reasons why black women with breast cancer frequently receive suboptimal care; however more dedicated research to fully elucidate these factors is needed.
For example, a large SEER-Medicare study involving over 55,000 women showed that blacks were less likely than whites to be treated at high-quality hospitals (defined as hospitals with the top quartile rates of radiation following breast conserving surgery) and consequently less likely to receive definitive primary therapy, a finding partially explained by having surgery at a high-quality hospital. When blacks do undergo breast conserving surgery, they are less likely to receive radiation than whites. In this study, the odds of radiation omission were more pronounced in blacks who lived in high-poverty areas or in areas further from cancer centers. Interestingly, these factors did not affect radiation use among whites.

The favorable impact of adjuvant chemotherapy on outcomes is minimized when full doses of therapy are not given as planned. Griggs et al. revealed that despite controlling for clinical and sociodemographic characteristics, blacks with breast cancer had systematic differences in chemotherapy administration as opposed to whites. They had a higher tendency to receive less relative dose intensity and more 1st cycle dose-reductions. Underlying reasons for these discrepancies remain unclear as biologic reasons such as lower neutrophil counts or chemotherapy tolerance were not associated with these disparities. In addition to receiving less than full doses of chemotherapy, significant delays in initiating chemotherapy are associated with lower survival. Although, Hershman et al. did not find significant racial differences in chemotherapy initiation delays in women ≥65 years with early stage breast cancer, a larger study involving almost 108,000 women found a higher risk of delays in blacks and Hispanics compared with whites. Blacks treated on chemotherapy protocols appear to benefit from chemotherapy with racial differences in stage-specific outcome versus whites.
analysis of patients treated on Southwest Oncology Group protocols between 1974 and 2001 demonstrated that blacks with breast cancer had increased mortality than whites despite being on the same chemotherapy protocols. This finding does not support the assumption that standardized treatments result in consistent outcome but supports the argument that black race has an independent detrimental effect on outcome.

Underuse of endocrine therapy in early-stage hormone receptor positive disease is also associated with inferior outcomes. A study between 1999 and 2000 found that blacks were less likely to receive adjuvant hormonal therapy than whites (71% vs. 80%) even after controlling for stage, comorbidity, age, insurance, and medical oncology referral.

Given the worse health outcome for blacks compared with whites, ensuring access to clinical trials to improve therapies for this group has been a focus of the National Institutes of Health. Unfortunately, the proportion of non-white clinical trial participants declined significantly between 1996 and 2002, particularly for breast, lung, and colorectal cancers. Lack of trust in the healthcare system, not being offered participation by physicians and lack of access to healthcare in general and thus, clinical trials are possible explanations for these differences. In addition, ethnic-related variability in stage of diagnosis and trial exclusion criteria may also contribute to disparities in clinical trial participation.

In summary, although the literature has provided important insights to the magnitude of disparities in clinical management; a major unanswered question is at which point in the process of care disparities arise, particularly for women within similar socioeconomic classes. Several programs to evaluate the quality of breast cancer care...
have been developed by professional organizations. These processes are potentially modificable and are therefore useful tools in addressing racial disparities. In fact, studies have shown that vulnerable groups such as older blacks with early-stage breast cancer are less likely to receive adequate care compared with whites even after adjusting for tumor characteristics and comorbidities. Data specific to younger women with breast cancer are not readily available yet fundamental to fully understanding differences in outcome in this population of patients. Omission of requisite care may be related to failed referral and this may be more common in minorities. Programs to address these disparities must therefore include systems to track or "navigate" vulnerable patients in the medical system.

**Conclusion**

Although several intervention programs to reduce breast cancer disparities have been instituted recently, the persistent magnitude suggests that current strategies may be inadequate. We have demonstrated that while a substantial amount of information is known about racial differences in outcome and associated factors, several gaps in knowledge exist. Addressing racial disparities in breast cancer therefore requires a multidisciplinary approach with innovative research to answer questions that are generally considered “outside the box” of traditional healthcare research. Disparities in access and quality of care may be eliminated by understanding and addressing cultural and economic barriers. Biologic differences in tumors in all racial groups need careful study to allow personalization of care and optimization of outcomes. Closing these gaps
in knowledge will in the future hopefully translate to interventions that lead to improved outcomes for all women with breast cancer.
References:

Table 1
Breast cancer epidemiology by race

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<tr>
<th>Characteristic</th>
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<th>White</th>
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<tr>
<td>Mortality per 100,000</td>
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<td>Stage distribution (%)</td>
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<td>Age of presentation under 50 (%)</td>
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<td>20.9</td>
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Modified from references 1 & 5
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