Immunohistochemical and Clinicopathological Characteristics of Invasive Breast Carcinoma in Nigeria

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Abstract

Objectives: We aimed to study the immunohistochemical and clinicopathological characteristics of invasive breast carcinoma among Nigerian women.

Methods: We conducted a retrospective assessment of the female patients diagnosed with breast carcinoma at a tertiary hospital in Nigeria between 2012 and 2019. Archived pathology request forms and processed specimens (tissue blocks and slides) were used as source data in addition to the patients’ demographic and other relevant data.

Results: Reports pertaining N = 113 patients were assessed. Their age range was 30 to 80 years (mean 52.1 ± 12.1 years). Breast carcinoma was most common in patients aged 40 to 49 years (32.7%), closely followed by those aged 50 to 59 years (30.1%). Invasive ductal carcinoma (NOS) was the most common histopathological subtype (94.7%). Nottingham grade III and grade II breast carcinoma accounted for 41.6% and 40.7% of the cases respectively. Mastectomy specimens formed 68.1% of the samples. The most common tumor size (75.9%) was > 5cm (mean 6.8 ± 3.2cm), consistent with the most common staging of T3 (46.0%). The most common lymph node involvement was N1 (56.6%). Immunohistochemical assessment of these tumors with estrogen receptor (ER), progesterone receptor (PR), and human epidermal growth factor receptor-2 (HER-2) biomarkers expressed positivity of 36.3%, 28.3%, and 41.6%, respectively. These tumors were immunohistochemically classified into luminal A (16.8%), luminal B (20.4%), HER-2 enriched (20.4%), and triple-negative (42.5%) subtypes.

Conclusions: The most common immunohistochemical subtype of invasive breast carcinoma among in this sample of Nigerian women was the triple-negative subtype (TNBC), similar to the finding among African Americans.

Keywords: Invasive Breast Carcinoma; Immunohistochemical; Clinicopathological; Estrogen Receptor; Progesterone Receptor; Human Epidermal Growth Factor Receptor Type 2; Nigeria.

Introduction

Breast carcinoma is the most prevalent cancer in women worldwide.1 The global burden of breast cancer, which saw 19.3 million new cases in 2020, is expected to rise to 28.4 million cases by 2040 with a 47% rise from 2020.2,3 Breast
carcinoma in African and African-American women is characterized by hormone receptor triple-negative tumors, late presentation, younger age, advanced stage, and higher grade, leading to worse prognoses compared to Caucasian western women. For optimal management, it is essential to have an accurate diagnosis of the state of the hormone receptors including the estrogen receptor (ER), progesterone receptor (PR), and human epidermal growth factor receptor-2 (HER-2). Breast carcinomas are classified into four major distinct immunohistochemical (IHC) molecular subtypes based on these intrinsic biologic factors using the following IHC markers: luminal A (ER-positive, PR-positive, and HER-2 negative); luminal B (ER-positive, PR-positive, and HER-2 positive); HER-2 overexpression (ER-negative, PR-negative, and HER-2 positive); and triple-negative/basal-like (ER-negative, PR negative and HER2 negative).

Oncologists have also classified breast cancers based on therapy choices using immunohistochemistry as a stand-in: (a) cancers with estrogen receptors (ER) and treated with chemotherapy; (b) cancers having HER-2/neu and treated with trastuzumab or lapatinib; and (c) cancers having negative hormone receptors for ER, PR, and HER-2 (i.e., the ‘triple-negative subgroup’) where chemotherapy is the only treatment option. The triple-negative subtype is reported to disproportionately impact young African-American women, and exhibit aggressive clinical behavior and high histopathological grades.

This study aimed to study the immunohistochemical and clinicopathological characteristics of breast carcinoma in the Nigerian city of Ilorin, including their histopathological types and grades, staging, and immunohistochemical subtypes.

Methods

We made a retrospective assessment of patients with breast carcinoma who received a diagnosis at the Department of Pathology, University of Ilorin Teaching Hospital, between 2012 and 2019. The research was conducted at the Department of pathology of the University of Ilorin Teaching Hospital (UITH), Kwara State, Nigeria.

The patients’ request cards were used to retrieve personal information, tumor size, histopathological diagnosis, Nottingham grading system, lymph node status, and immunohistochemical diagnosis (hormone receptor status and human epidermal growth factor findings). Medical records were used to extract the history of co-morbidities, history of neoadjuvant chemotherapy, local symptoms like breast lumps, deformed breast shapes, tumor sizes, puckering skin changes, peau d’orange, ulcers, fixation to chest wall structures, and treatment received.

Our inclusion criteria restricted the participants to women with invasive breast carcinoma diagnosis whose full pathological data was available (including tumor grade, tumor size, number of harvested positive lymph nodes, clinical staging, and type of surgery performed). Patients with incomplete data were excluded from the study.

Our hospital procedure required all harvested surgical biopsies and mastectomies to be immediately subjected to fixation in 10% neutral buffered formalin in a labelled container and sent to the pathology department for study. There, the specimens were grossed by determining the size, color, consistency, weight, and dimension of the tumor. Following tissue processing, embedding, and routine hematoxylin and eosin (H&E) staining, representative tumor tissue section slides were prepared and used to make histopathological diagnoses. The Nottingham composite histopathologic grading scheme was used to access the histopathological grades (Elston-Ellis modification of Scarff-Bloom-Richardson-Elson grading system). The WHO classification system from 2012 was used to assess the various histopathological types. The TNM system, endorsed by UICC and the American Joint Committee on Cancer and End Results Reporting, was used to assess tumor staging. The immunohistochemistry procedure with avidin-biotin complex (ABC), also known as the Avidin biotin Immunoperoxidase technique, was employed on the formalin-fixed paraffin-embedded (FFPE) tissue blocks used for this study. Therefore, the slides on ER and PR were assessed using nuclear reactivity and the slides on HER-2 using membrane reactivity. Cells with complete membrane staining that appear brownish for HER-2 are regarded positive. Cells with distinct brown colors in the nuclei for ER and PR are also considered positive. Immunostained negative tumor cells, however, show as bluish. Non-specific binding/brown artifacts on cells and connective tissue are ignored.

The staining intensity of immunohistochemical reactions was graded and scored as per the Allred scoring guidelines. The percentage of tumor nuclei stained, and the intensity of staining are the two components that make up the Allred grading system (for ER and PR), where a score ≥ 1 indicates that both proportion and intensity are positive. HER-2/neu was assessed using the Dako Herceptest scoring guideline, yielding results of 0, +1, +2, and +3.
Sections with scores of +3 were viewed as positive, whereas those with scores of 0 and +1 were viewed as negative, and scores of +2 as equivocal (requiring in-situ hybridization studies for confirmation).

IBM SPSS Statistics for Windows, version 20 (IBM Corp., Armonk, N.Y., USA) was used to analyze the data (SPSS, Inc., Chicago, IL, USA). The data was displayed in tables, charts, and photomicrographs. For quantitative variables, data was described with the mean standard deviation (SD), and for qualitative variables, as p-values is significant when less than 0.05 and percentages.

This study followed the ethical principles of the Helsinki Declaration on human subjects in biomedical research. Information about the patients’ identities and personal health was kept private. The ethical approval for the study was obtained from the University of Ilorin Teaching Hospital Ethical Committee with registration number NHREC/02/05/2010 dated December 28, 2016.

Results

This study examined specimens from N = 113 breast carcinoma patients, which included 36 surgery biopsies (31.9%) and 77 mastectomies (68.9%).

The average patient age was 52.1 ± 12.1 (range: 30–80) years. There were more patients in their forties and fifties than in the other age groups. [Table 1]. Invasive ductal carcinoma not otherwise specified (NOS) accounted for 107 of 113 (94.7%) patients. Grade I breast cancers were detected in 20 (17.7%) cases in the Nottingham grading scheme, while grade III and grade II cancers numbered 47 (41.6%) and 46 (40.7%) respectively [Table 2].

Table 1: Age at presentation (in years) of patients with breast cancer (N = 113).

<table>
<thead>
<tr>
<th>Age groups</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 40</td>
<td>20</td>
<td>17.7</td>
</tr>
<tr>
<td>40 – 49</td>
<td>37</td>
<td>32.7</td>
</tr>
<tr>
<td>50 – 59</td>
<td>34</td>
<td>30.1</td>
</tr>
<tr>
<td>60 – 69</td>
<td>10</td>
<td>8.8</td>
</tr>
<tr>
<td>≥ 70</td>
<td>12</td>
<td>10.6</td>
</tr>
<tr>
<td>Mean ± SD</td>
<td>52.1 ± 12.1</td>
<td>30 – 80</td>
</tr>
</tbody>
</table>

Table 2: Histopathological type and grade of breast carcinoma (N = 113).

<table>
<thead>
<tr>
<th>Histopathological type</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>Invasive ductal carcinoma (NOS)</td>
<td>107</td>
<td>94.7</td>
</tr>
<tr>
<td>Metaplastic carcinoma</td>
<td>1</td>
<td>0.9</td>
</tr>
<tr>
<td>Mucinous carcinoma</td>
<td>3</td>
<td>2.7</td>
</tr>
<tr>
<td>Papillary carcinoma</td>
<td>1</td>
<td>0.9</td>
</tr>
<tr>
<td>Plasmacytoma</td>
<td>1</td>
<td>0.9</td>
</tr>
<tr>
<td>Histopathological grade</td>
<td></td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>20</td>
<td>17.7</td>
</tr>
<tr>
<td>2</td>
<td>46</td>
<td>40.7</td>
</tr>
<tr>
<td>3</td>
<td>47</td>
<td>41.6</td>
</tr>
</tbody>
</table>

Lymph nodes in the n = 83 mastectomy specimens were investigated and divided into three groups, (a) N0, where the cancer has not spread to the lymph nodes, (b) N1 (metastatic carcinoma has affected 1–3 lymph nodes), and (c) N3 (significant metastatic carcinoma in lymph nodes). N0 was found in 35 of 83 (42.2%) lymph node samples, N1 in 47 (56.6%) , and N3 in one (1.2%) [Table 3]. The tumor diameters ranged 3.2–16.0 (6.8 ± 3.2 cm). Most patients (75.9%) had tumors larger than 5.0 cm [Table 3].

All cases were subjected to immunohistochemical evaluation using monoclonal ER, PR, and HER-2 antibodies. Most tumor cells (72 of 113; 63.7%) were ER-negative, and the remaining 41 (36.3%) were ER-positive. In contrast 81 (71.7 %) were PR-negative and the remaining 32 (28.3%) PR-positive. HER-2 positivity was found in (47 of 113 (41.6 %) tumor cells [Table 4 and Figure 1]. Additionally, triple-negative (ER-/PR-/HER-2-) accounted for 48 (42.5%), luminal A (ER+/PR+/HER-2-), for 19 (16.8%), luminal B (ER+/PR+/HER-2+) for 23 (20.4%), and HER-
2 enriched (ER-/PR-/HER-2+) for 23 (20.4%) cases, respectively [Figures 1 and 2]. Notably, 52 (46.0%) patients were in stage T3, 29 (25.7%) were in stage T4, and one (0.9%) patient was in stage T1 [Table 5]. We found that the predominant locoregional symptom of breast carcinoma was lumps as found in 105 (92.9%) patients, followed by deformity of the breast (78; 69.0 %) and ulceration (21;18.6 %) [Table 6].

Table 3: Specimen type, tumor size, and lymph node of women with breast cancer.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Frequency</th>
<th>Percentage (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Type of specimen</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Biopsy</td>
<td>36</td>
<td>31.9</td>
</tr>
<tr>
<td>Mastectomy</td>
<td>77</td>
<td>68.1</td>
</tr>
<tr>
<td><strong>Tumor size (cm)</strong></td>
<td>n = 83</td>
<td></td>
</tr>
<tr>
<td>&lt; 2</td>
<td>0</td>
<td>0.0</td>
</tr>
<tr>
<td>2–5</td>
<td>20</td>
<td>24.1</td>
</tr>
<tr>
<td>&gt; 5</td>
<td>63</td>
<td>75.9</td>
</tr>
<tr>
<td><strong>Mean ± SD</strong></td>
<td>6.8 ± 3.2</td>
<td></td>
</tr>
<tr>
<td><strong>Range</strong></td>
<td>0 – 16</td>
<td></td>
</tr>
<tr>
<td><strong>Lymph node</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>N0</td>
<td>35</td>
<td>42.2</td>
</tr>
<tr>
<td>N1</td>
<td>47</td>
<td>56.6</td>
</tr>
<tr>
<td>N3</td>
<td>1</td>
<td>1.2</td>
</tr>
</tbody>
</table>

Table 4: Immunopositivity of estrogen, progesterone receptor, and HER-2 biomarkers.

<table>
<thead>
<tr>
<th>Variables</th>
<th>Positive (%)</th>
<th>Negative (%)</th>
<th>Total (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>ER</td>
<td>41 (36.3)</td>
<td>72 (63.7)</td>
<td>113 (100)</td>
</tr>
<tr>
<td>PR</td>
<td>32 (28.3)</td>
<td>81 (71.7)</td>
<td>113 (100)</td>
</tr>
<tr>
<td>HER-2</td>
<td>47 (41.6)</td>
<td>66 (58.4)</td>
<td>113 (100)</td>
</tr>
</tbody>
</table>

Note: HER-2: epidermal growth factor receptor 2. ER: estrogen receptor, PR: progesterone receptor
**Figure 1:** Immunohistochemical expression of ER, PR, and HER-2 biomarkers.

**Note.** HER-2: epidermal growth factor receptor 2. ER: estrogen receptor; PR: progesterone receptor

**Figure 2:** Immunohistochemical subtypes of breast carcinoma.

**Note:** HER-2: epidermal growth factor receptor 2.

**Table 5:** Staging of breast carcinoma (where T = tumor size) [N = 103].

<table>
<thead>
<tr>
<th>Staging</th>
<th>Frequency</th>
<th>Percentage</th>
</tr>
</thead>
<tbody>
<tr>
<td>T1</td>
<td>1</td>
<td>0.9</td>
</tr>
<tr>
<td>T2</td>
<td>21</td>
<td>18.6</td>
</tr>
<tr>
<td>T3</td>
<td>52</td>
<td>46.0</td>
</tr>
<tr>
<td>T4</td>
<td>29</td>
<td>25.7</td>
</tr>
</tbody>
</table>

**Table 6:** Presenting Symptoms of breast cancer patients* (N= 113).

<table>
<thead>
<tr>
<th>Variable</th>
<th>Frequency (%)</th>
<th>Frequency (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lump</td>
<td>105 (92.9)</td>
<td>8 (7.1)</td>
</tr>
<tr>
<td>Deformity of the breast</td>
<td>78 (69.0)</td>
<td>35 (31.0)</td>
</tr>
<tr>
<td>Ulceration</td>
<td>21 (18.6)</td>
<td>92 (81.4)</td>
</tr>
<tr>
<td>Lump and deformity</td>
<td>57 (50.4)</td>
<td>56 (49.6)</td>
</tr>
<tr>
<td>Lump, deformity, and ulceration</td>
<td>21 (18.6)</td>
<td>92 (81.4)</td>
</tr>
</tbody>
</table>

*Multiple symptoms may have presented in the same patient.

**Discussion**

This study sought to determine the age of breast carcinoma patients, their histopathological types and grades, staging, and immunohistochemistry subtypes. It also aimed to analyze the clinicopathological and immunohistochemical characteristics of breast carcinoma in Ilorin, Nigeria.

The age range of our patients was 30–80 years and the most represented age group (32.7%) was 40–49 years. The mean age at presentation was 52.1 ± 12.1 years. These are comparable to the previous findings at the same center. Our results are also consistent with those from other parts of Nigeria. Research from Ghana, Egypt, Sudan, Saudi Arabia, Malaysia and Eastern Europe also endorse a similar fifth decade peak.
At 94.7%, invasive ductal carcinoma was the most prevalent histological type of breast cancer NOS among our patients, similar to the 82.6% prevalence reported by an earlier study at the same center.11 Comparable findings have also emerged from Ibadan, Calabar, Lagos, and Sokoto.16,23,24 The notable exception was a study from Uyo (Nigeria) which reported much lower prevalence (53.0%) of invasive ductal carcinoma NOS than most other studies including ours.25 Studies in other African countries have also yielded similar prevalence as ours: Sudan (90.0%), Cameroon (90.0%), Ghana (75.8%), Tanzania (78.0%), and Tunisia (87.0%).26-30 Comparable prevalence of invasive ductal carcinoma have been reported from Iraq (95.5%) and India (88.0%) as well.31,32

Mucinous carcinoma represented 2.7% of the cases in this study, compared to lower prevalence reported from Oshogbo and Lagos (1.2%–2.5%).23,31 Results from other parts of Nigeria have been diverse.9,20,30 Kano in northwestern Nigeria reported significantly higher (6.9%) prevalence in mucinous carcinoma than most other Nigerian studies including ours.34 Data from other African nations revealed varied prevalence of mucinous carcinoma ranging from 2% to 9.8%.17,31,35

The prevalence of metaplastic carcinoma in our study was 0.9%, and similar findings elsewhere ranged from 0.19% to 2.4%.12,33,36

In our study, Nottingham grade I breast carcinoma accounted for 17.7% of cases, compared to 12.7% in Lagos, 9.4% in Ibadan, and 5.9% in Jos.23,37,38 Dramatically high prevalence of 27.3% of grade I breast carcinoma was reported from Kano in northwest Nigeria.39 The rates of grade I breast carcinoma in other African nations have been reported at 1.0%, 9.0%, 14.8%, and 25.0%, respectively, in Sudan, Tunisia, Ghana, and Tanzania.20,26,29,30 Outside Africa, a prevalence of 4.2% has been reported in Pakistan, 17.4% in Iraq, and 29.0% in the United States (USA).31,40,41 Geoclimatic, ethnic, and lifestyle differences might be hypothesized to be responsible for much of the observed variations in prevalence of different types of breast cancer.

In this study, Nottingham grade II breast cancer represented 40.7% of all cases, vis-à-vis Ibadan (44.3%) and Lagos (48.0%), Kano (33.4%), Jos (23.5%), and Calabar (23.5%).17,27,34,37,38 The prevalence of grade II breast carcinoma in other countries were widely variable: Ghana (31.5%–39.7%),17,28 Iran (36.2%), US (41.4%), and Pakistan (75.0%).20,34,37,38 The discrepancy in histopathologic grades noted in the various study populations may be due to the Nottingham histopathologic grade system’s interobserver variability, and variations in the disease’s demographic distribution.

Nottingham grade III breast carcinoma had a prevalence of 41.6% among the participants in this study. Prevalence rates reported from different parts of Nigeria include 29.7% of cases in Ilorin12 and 15.6% in Ibadan,37 7.1% in Kano, 66.7% in Calabar, and 70.6% in Jos.16,34,38 Nottingham grade III prevalence in other African nations were 30.3% in Cameroon, 35.1% in Tunisia, 53.7% in Ghana, and 68.0% in Sudan.17,26,27,30 Across the world, the prevalence was 20.3% in Pakistan, 30.3% in US, and 46.4% in Iran.31,40,41 Different types of biopsies (tru-cut, incisional, and excisional) used for assessment and interobserver variability in the assessment of breast carcinoma using the Nottingham histopathologic grade system may partly have may led to the discrepancies of findings in the surveys in Nigeria, other African nations, and developed countries. Due to the minimal amount of breast tissue compared to incisional or excisional breast biopsies, tru-cut biopsies tend to downgrade breast carcinoma. The accuracy of diagnoses might also be affected by geopolitical and socioeconomic factors.

The size of the tumor, clinical stage, and lymph node metastases are significant independent prognostic variables for invasive breast carcinoma. A sizable proportion of women with breast carcinoma in this study presented with lymph node involvement, matching stages 3 and 4, and had tumor sizes greater than 5 cm. Our results are comparable to those from regional and international studies.34,38,42,44

When filling out the pathology request form, many patients do not have the proper documentation of their clinical staging, which could affect the accuracy of the database and subsequent follow-up and management. To properly prognosticate patients, the pathologist should take careful note of the tumor’s size and thoroughly dissect the axillary tail fat. Due to false positives and false negatives—such as reactive follicular hyperplasia and sinus histiocytosis of the nodes—the clinical assessment of lymph node status is inaccurate (e.g., lymph nodes with small metastatic deposits). Consequently, a biopsy is required for a precise evaluation and diagnosis. The 10-year disease-free survival rate is close to 70% to 80% with no nodal involvement; the rate drops to 35% to 40% with one to three positive nodes, and 10% to 15% with more than ten nodes positive.45 In addition, prior studies only examined the lymph nodes for presence of malignancy without stratifying and specifying the number of positive lymph nodes for staging.45 The majority of patients in this study had advanced tumors that were larger than 5 cm, consistent with past studies in Nigeria and other Sub-Saharan African nations.33,36,40,41 Lack of coordinated breast cancer screening, political apathy,
Breast carcinoma tumor cells in this research had immunohistochemistry reactivity for ER, PR, and HER-2 at rates of 36.3%, 28.3%, and 41.6%, respectively. A retrospective study conducted in Nigeria and Senegal that included 507 patients diagnosed with breast carcinoma among African Americans of Nigerian and Senegalese origin between 1996 and 2007 showed ER, PR, and HER-2 at rates of 24.0%, 20.0%, and 17.0% respectively, however, this contrasts with a study conducted in the University of Ibadan and University of Calabar a few years ago that showed 27.0%, 16.0%, and 30.0% for ER, PR, and HER-2 respectively.4,44 While Agbo et al. in Sokoto discovered that ER, PR, and HER-2 were 47.8%, 41.3%, and 43.5% respectively for immunostained breast carcinoma tumor cells, the corresponding values from Uyo were 18.0%, 14.8%, and 32.8% respectively.45,46 In Ibadan, the ER, PR, and HER-2 expression was 65.1%, 54.7 %, and 79.7 %, respectively. A US study among Caucasians found that ER, PR, and HER-2 expression were 45.0%, 37.0%, and 32.0%, respectively.37,40,42 Such wide variations in immunoreactivity values in studies above might be due to limitations in some studies. These may include the use of archival paraffin-fixed embedded tissue blocks with antigen degradation due to poor storage or suboptimal adherence to prescribed guidelines for handling and storing breast specimens in some studies.48

The immunohistochemical subtypes of breast carcinoma recorded in this study were 16.8%, 20.4%, 20.4%, and 42.5% for luminal A, luminal B, HER-2 enriched, and triple-negative, respectively. These results differed from those from a previous study in our center which were 20.0%, 11.0%, 30.0%, and 25.0%, respectively.44 The results were equivalent to those of a study carried out in Sokoto, Senegal, and Nigeria’s six geopolitical zones.4,46 But a study among Nigerians who were African Americans and Senegalese at six geopolitical zones where University College Hospital in Ibadan was used as coordinating center revealed that the hormonal expression in patients from African-American patients was equivalent to that of Caucasians and people from other parts of the world.37,40,49,50 The differences observed among racial groups and geographical areas could be explained by tumor biology, various antibody clones, antigen retrieval SOPs, and bias in the use of the Allred & Herceptest scoring system.

The chief limitation of this study is its retrospective nature; the principal investigator lacked control over the historical patient data, preanalytical/analytical tissue handling variables including those related to standardized cold ischaemic time in the operating room, fixative composition, and fixation time. Additionally, due to a lack of resources, the breast carcinoma cases with HER-2 membrane staining scores of 2+ could not be subjected to in-situ hybridization tests such as fluorescent in-situ hybridization (FISH) for further characterization.

The next step will be to carry out a prospective study where the lead researcher will oversee these variables. Along with performing molecular biological studies like FISH, polymerase chain reaction (PCR), and next-generation sequencing (NGS) to identify a novel therapeutic target for the prevalent triple-negative subtype of breast carcinoma. Furthermore, a large-scale multicenter study is essential for a better understanding of the nature and prevalence of various types of breast carcinoma in Nigeria.

Conclusion

This study found that breast carcinoma was most prevalent among Nigerian woman in their fifties and that 42.5% of the invasive breast carcinomas had triple-negative immunohistochemical phenotypes similar to the findings among African Americans. Breast cancer is now categorized as a disease of public health importance. Early diagnosis through coordinated breast screening and government support will help mitigate the death toll of breast cancer patients in Nigeria and the other African countries.

Disclosure

The authors’ contributions to this paper were as follows: RMW: conception and design of the work, data acquisition, data analysis, manuscript preparation, critical appraisal, final approval of the version to be published, and agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. UBE: manuscript reviewing and editing, literature review,
editing, critical appraisal. AOS: literature review and critical reviewing, editing. AEA: design, interpretation of data, drafting and revising, final approval. NAI: literature review and critical reviewing, editing. AAA: literature review and critical reviewing, editing. AAT: statistical analysis.

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