Young Women with Breast Cancer in Resource-Limited Settings: What We Know and What We Need to Do Better

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Abstract: Young women with breast cancer (YWBC) account for a variable proportion of patients diagnosed with breast cancer around the globe, with a higher prevalence in resource-limited settings than in high-income countries. This group represents a unique population that warrants special attention due to specific biological considerations and age-specific supportive care issues. This review aims to explore existing knowledge regarding YWBC’s needs, particularly in resource-restricted settings. To date, scarce information regarding the care of YWBC in resource-constrained countries is available, with most reports describing suboptimal care in terms of survivorship needs. Health care providers should implement actions to improve endocrine treatment adherence, referrals for fertility counseling and preservation, contraceptive use compliance, timely body image and sexual function interventions, comprehensive genetic risk assessments, and early quality of life and psychosocial health interventions. While high costs act as a barrier for optimal care in resource-limited settings, improving patient education represents a promising and cost-effective solution to improve patient care. Future research on developing tailored educational resources for YWBC in resource-limited settings should be considered a priority.

Keywords: breast cancer, young women, unmet needs, disparities, resource-limited settings, low- and middle-income countries

Introduction

Globally, breast cancer (BC) is the most frequently diagnosed malignancy and the leading cause of cancer-related death in women.1 In low- to middle-income countries (LMICs), BC is also considered the leading cause of disability among young women.2-4 Young women with breast cancer (YWBC), defined as women aged 40 years or less at diagnosis, represent an often underserved population with age-specific needs.5 While BC in young women is considered a relatively rare clinical entity that accounts for 5–7% of all BC cases in high-income countries (HICs),6 LMICs such as Iran (23%),7 Nigeria (21%),8 Mexico (15%),9 India (15%),10 and Brazil (11%)11 report a substantially higher proportion of cases diagnosed at a young age. Hence, understanding the particular needs of this population could be vital in optimizing BC care in resource-limited countries.

YWBC are diagnosed at more advanced stages, with more aggressive pathological features, such as a higher proportion of grade 3, triple-negative, and HER2-positive tumors, and higher rates of recurrence and death compared to their older counterparts.6 Additionally, YWBC face not only the threat of a potentially fatal illness and...
burdensome treatment, but also experience an added burden of unique concerns at the peak of their careers, motherhood plans, and family life. Therefore, attention to age-specific cancer and supportive care issues, including endocrine treatment-related toxicities, fertility counseling and preservation, contraceptive use, body image and sexual function/satisfaction, genetic risk, quality of life (QoL), and psychosocial health, is of utmost importance in this unique population.

Although comprehensive care of YWBC has been found to be suboptimal worldwide, efforts to improve awareness and knowledge among patients and providers about the specific needs of YWBC have been undertaken predominantly in HICs, such as the United States and Canada. In LMICs, scant resources, limited access to healthcare, and cultural and socioeconomic barriers may further aggravate YWBC’s prognosis. Furthermore, limited data exist on YWBC’s special needs, especially in LMICs. Therefore, the main aim of this review is to explore existing knowledge regarding YWBC’s unique needs specifically in resource-limited settings.

Methods

The authors conducted a comprehensive search using the PubMed database for publications on YWBC in resource-limited settings. Resource-limited settings were defined as all countries classified as low-income, lower-middle-income, and upper-middle-income economies according to the World Bank Group report, based on its gross national income per capita. Furthermore, young women were defined as those aged 40 years or younger at BC diagnosis. Only publications in the English language were considered eligible. The following keywords were searched: breast cancer, young women, special needs, supportive care, limited resource settings, low- and middle-income countries, developing countries, fertility counseling, fertility preservation, contraceptive use, body image, sexual function, sexual satisfaction, genetic risk, quality of life, and psychosocial health. Studies that described YWBC in resource-limited settings, including epidemiological, clinical-pathological, treatment, and supportive care characteristics, were considered for this narrative synthesis.

Endocrine Treatment-Related Concerns

Although YWBC have a higher proportion of triple-negative and HER2-positive tumors than postmenopausal women, young age by itself is not associated with worse outcomes in these specific subtypes. Nonetheless, most YWBC will present with hormone receptor (HR)-positive tumors, and patients with this molecular subtype have a higher risk of recurrence and death than their older counterparts. Several theories have been proposed for the unfavorable prognosis in HR-positive tumors in YWBC including treatment failure/resistance, a higher proportion of luminal B than luminal A tumors, and relatively low rates of adherence to adjuvant endocrine therapy (ET).

Current international guidelines recommend adjuvant ET with tamoxifen or an aromatase inhibitor (AI) for at least five and up to 10 years with or without ovarian function suppression (OFS) for premenopausal women with HR-positive BC. Two studies have focused solely on describing ET adherence rates in YWBC in limited-resource settings. Remarkably, both of these studies report high adherence rates to ET in YWBC, contrasting to previous reports from HICs. The first study to evaluate tamoxifen adherence specifically in YWBC in a resource-constrained setting was recently published by the “Joven & Fuerte” program. In this report, a cross-sectional survey among 141 YWBC receiving adjuvant tamoxifen was conducted at the National Cancer Institute in Mexico City. Adherence to tamoxifen was measured subjectively through self-reported surveys and objectively through the medication possession ratio (MPR). Regarding subjective adherence, 95% expressed taking tamoxifen regularly, with 70% reporting not missing any doses in the past 30 days. For objective adherence, 74.8% of patients had an MPR ≥80%.

Subsequently, the same research group evaluated premenopausal BC patients’ adherence to ET and physicians’ prescribing practices in three Mexican referral centers (including two public and one private hospitals). Seventy-two percent and 27% of participants reported complete (100%) and acceptable (>80%) adherence, respectively. Being employed/studying, worrying about long-term ET use, and experiencing ET-related symptoms were associated with incomplete adherence. Treating physicians prescribed guideline-endorsed regimens with tamoxifen or AI plus OFS only in 64% of patients. However, 88% of the patients should have received this treatment modality.

Surprisingly, both studies report high adherence rates to ET in YWBC, contrasting to previous reports from HICs. Causes for the high treatment adherence rates
described in these two reports may be associated with
the fact that patients were being treated in a YWBC
specialized program, with increased rates of ET patient-
provider discussions before treatment initiation, higher
perception of the potential benefits of adjuvant ET, and
free of charge medications dispensed in public hospitals.
The main reasons associated with patients’ low adher-
ence and high discontinuation rates of ET included for-
getfulness, adverse effects, and unwillingness to take the
medication.\textsuperscript{32,33} As for physicians’ suboptimal treatment
recommendations with OFS, a possible explanation may be
the prevailing limited patient access to gonadotropin-
releasing hormone analogs (GnRHa), which are not rou-
tinely covered by Mexican public healthcare insurance
schemes, and are considered expensive.\textsuperscript{33}

Another essential factor that might impact adherence
and persistence in this reproductive-aged group is fertility con-
cerns associated with long-term use ET and possibly reduced
chances of successful conception.\textsuperscript{34–36} The POSITIVE study
is currently investigating the impact of temporary ET inter-
ruption to allow pregnancy in premenopausal women with
HR-positive early BC who had received 18–30 months of
adjuvant ET and wished to interrupt ET for pregnancy, deliv-
ergy, and breastfeeding, followed by ET resumption to com-
plete the planned duration. Patient accrual has closed, and
results on safety of adjuvant ET interruption are currently
awaited. Unfortunately, only 6 (1.2\%) of the 518 women
enrolled in this study, live in a LMIC.\textsuperscript{35}

Although young age has been associated as a risk factor for
low adherence to adjuvant ET in previous studies in HICs,\textsuperscript{28,29}
the scarce available evidence from an upper-middle-income
country, Mexico, suggests ET adherence might not be as low
as expected in YWBC from LMICs.\textsuperscript{32,33} However, further
research concerning this important topic is needed in other
limited-constrained settings as YWBC women potentially
have longer life expectancies than their older counterparts,
and low adherence rates to ET have been associated with
worse long-term survival.\textsuperscript{37} Developing setting-adapted strat-
gies to improve physicians’ adherence to guideline recom-
endations and YWBC’s adherence to adjuvant ET as well as
guaranteeing adequate access to optimal ET are urgently
needed in limited-resource settings.

**Fertility Counseling and Preservation**

Current BC international guidelines recommend oncfo-
tility counseling to be started as early as possible in the
treatment plan of every YWBC, irrespective of subtype
and stage of disease, and even if there is no interest in
future offspring. A comprehensive fertility counseling is
comprised of treatment-related gonadotoxicity assessment,
and a discussion on the different strategies available for
ovarian function and/or fertility preservation.\textsuperscript{38} Even
though oncology counseling is an essential component in
the management of YWBC, particular barriers in
resource-limited settings may be encountered, as attention
is mainly directed to cancer diagnosis and treatment,
whilst survivorship concerns may be neglected. Such bar-
riers might arise from a lack of engagement from health
care providers, limited public coverage for supportive
needs heightened by significant socioeconomic gaps, and
inequitable distribution of cancer centers and specialists.\textsuperscript{39}

In a 2017 Mexican report from a tertiary care center, only 30.6\% of YWBC recalled receiving information from
their treating physicians about the infertility risk associ-
ated with cancer treatment.\textsuperscript{40} When surveying Mexican
physicians on their attitudes and knowledge toward ferti-
ity preservation in YWBC, only 58\% claimed they always
informed patients about infertility risks, and only 38\% always
discussed fertility preservation procedures. Strikingly, 24\% of physicians wrongly considered preg-
nancy and 20\% GnRHa use during chemotherapy to be
detrimental to the prognosis of BC.\textsuperscript{41} Such misconceptions
and neglect towards interventions aimed at fertility pre-
servation are not restricted to the Latin American region;
a report from East Asia also describes a reduced uptake on
fertility preservation strategies due to lack of discussions
and referrals to fertility specialists.\textsuperscript{42}

Not only physician-related barriers have led to scarce
oncology referrals in limited-resource settings. In
fact, high cost is regarded as one of the most significant
barriers for fertility preservation.\textsuperscript{41,43,44} Without insur-
ance coverage, access to fertility preservation strategies
is only through out-of-pocket spending, and for a large
proportion of BC patients in LMICs, these treatments are
unaffordable.\textsuperscript{39,45} This was illustrated in the previously
mentioned 2017 Mexican survey, as only 3\% of the
surveyed patients considered themselves able to afford
extra expenses beyond oncologic treatment.\textsuperscript{46}
Furthermore, the short window of time that cancer
patients have before starting gonadotoxic cancer treat-
ment further limits fertility preservation referrals, parti-
cularly in young Latin American patients with a higher
burden of locally advanced disease.\textsuperscript{46}
Multinational efforts are being made to assess fertility preservation access in resource-limited settings. The Repro-Can-OPEN Study surveyed oncofertility centers from 14 developing countries (Egypt, Tunisia, Brazil, Peru, Panama, Mexico, Colombia, Guatemala, Argentina, Chile, Nigeria, South Africa, Saudi Arabia, and India) on the availability and utilization of different fertility preservation strategies. The most frequent methods for fertility preservation in female patients with BC identified were gonadal shielding in case of irradiation (62.5%), fractionation of chemo- and radiotherapy (62.5%), and oocyte/embryo freezing (58.9% and 55.4%, respectively). The systematic development of international efforts among LMICs, such as the Repro-Can-OPEN, represents one of the first steps to enhance quality fertility counseling and access to preservation, but much effort is still needed as these programs were developed mainly in tertiary health care centers. In Mexico, the “Joven & Fuerte” program has made significant advances to improve access to fertility preservation strategies. In their prospective study on fertility preservation, out of 140 YBCW who desired future offspring before systemic cancer treatment, 38% could preserve fertility. Among fertility preservation techniques, oocyte/embryo freezing was the most frequently used method (59%), followed by temporary ovarian suppression with GnRHα during chemotherapy (26%) or a combination of both strategies (15%).

**Contraceptive Use**

Although BC treatment may have detrimental effects on ovarian function, such as treatment-induced amenorrhea and early menopause, it does not exclude the possibility of pregnancy. Therefore, contraceptive counseling and adherence are extremely important because reproductive-age BC patients may have an unintended pregnancy during treatment, restricting the use of some local and systemic therapeutic options, such as radiation therapy, monoclonal antibodies, and ET, and representing added risks for the fetus. Current studies from HICs show that the use of effective contraceptive methods by YWBC is suboptimal. However, information on contraceptive counseling and use in YWBC in LMICs is limited.

In a Turkish qualitative study, premenopausal women with BC were interviewed to evaluate the information needed and received by those women regarding contraception, early menopause, infertility, fertility preservation, and sexuality. Twenty premenopausal women with BC reported receiving insufficient counseling on these issues, despite their desire to receive information regarding these issues. Only five of the 18 women who did not want to become pregnant again were using an effective contraceptive method, and only eight had received contraceptive counseling after BC diagnosis. This study demonstrated that premenopausal women with BC have unmet information needs about contraception and the other related topics.

A cross-sectional survey was conducted regarding contraceptive use and counseling among YWBC who had completed chemotherapy in the previous five years or were currently receiving adjuvant ET and/or trastuzumab at a large tertiary health care facility in Mexico. Of 104 participating women, only 51.1% and 45.7% reported using contraception during chemotherapy and adjuvant ET and/or trastuzumab, respectively. Of the 51 patients who were sexually active during chemotherapy, only 29.4% used an effective contraceptive method. When asked about contraceptive counseling, only 16.7% recalled being advised by their health care provider about effective strategies. As anticipated, women who received contraception counseling used contraceptives more often than women who were not.

A qualitative study of in-depth interviews among 24 women aged 18–49 years diagnosed with BC at a tertiary hospital in Cape Town, South Africa. Among women using a contraceptive method, the most used strategy after BC diagnosis was the non-hormonal intrauterine device. Remarkably, all women reported receiving limited information from health care providers about contraceptive use, future fertility planning, impact of cancer treatment on fertility and fertility preservation options.

Thus, only three studies have reported on contraceptive use and counseling in LMICs (Turkey, Mexico, and South Africa) from different continents. These reports demonstrate that YWBC contraception use is low and that women need and are willing to receive information on effective contraceptive options. Notably, these three studies took place in referral cancer centers. Hence, patients in less specialized institutions may have lower use and less counseling on effective contraception. Informing all premenopausal patients about effective contraceptive use during all treatment duration should be an essential component of the supportive care of YWBC.

**Body Image and Sexual Function/Satisfaction**

Sexual function is a fundamental aspect of patients’ well-being. BC treatment modalities can lead to various
sexual changes among BC survivors. \(^{58-60}\) Locoregional treatments, including surgery and radiotherapy, can produce intense emotional distress, cause a detrimental change in body image, and reduce sensitivity in the breasts. Furthermore, systemic treatments, including chemotherapy and ET, can cause premature menopause, decreased libido, dyspareunia, and failure to reach orgasm, leading to reduced sexual activity and satisfaction. \(^{59}\)

In a qualitative study conducted in Iran, 21 BC survivors under 51 years of age were included for in-depth semi-structured interviews regarding sexual life. The participants experienced changes in sexual function as a result of their BC diagnosis that led to an unfulfilled sex life. Four main barriers leading to an unfulfilled sexual experience were identified: a) undesirable sexual function, characterized by decreased sexual desire, arousal and satisfaction, less frequent sexual relationships, and perception of diminished sexual attractiveness; b) context-based beliefs, including religious, cultural and family beliefs; c) unmet information and support needs by health care providers; and d) emotional crisis, with feelings of loss of femininity and guilt. \(^{57}\)

In another qualitative methods study from Nigeria, fifteen women ≤45 years old who underwent mastectomy for BC were interviewed over a 6-month period. This study identified several major themes on the impact of mastectomy on YWBC’s lives. Removal of the breast considerably impacted participants’ perception of femininity, making them look physically less like women and more like men. Also, women in this study believed that breasts were foci for sexual arousal. Therefore, they had difficulty becoming sexually aroused and experienced decreased libido after mastectomy, further inflicting them concern about satisfying their husbands. \(^{61}\)

The “Joven & Fuerte” group has also reported from a prospective study assessing sexual health using the Female Sexual Function Index (FSFI) and the Sexual Satisfaction Inventory (SSI) at baseline and 6-month follow-up. Mexican YWBC experienced a high percentage of sexual dysfunction, as well as an elevated proportion of low sexual satisfaction at both assessments, with a worse sexual function at follow-up than at baseline. \(^{62}\) Similarly, in another cross-sectional study from Iran, 144 premenopausal women with BC who underwent surgical therapy were assessed using the questionnaires FSFI and Meston’s Sexual Satisfaction scale for women. Remarkably, all patients had sexual dysfunction in all dimensions and even lower sexual satisfaction scores with total mastectomy. \(^{63}\)

More knowledge about the sexual difficulties and adaptation process of young BC survivors and their partners is needed. The timely identification of women at risk for sexual dysfunction is imperative to provide them with the necessary tools for a fulfilled sexual experience during and after BC diagnosis and treatment. Tailored interventions to address concerns regarding sexual function and satisfaction need to be developed according to the different cultural and religious backgrounds of YWBC in LMICs.

**Genetic Risk**

Among unselected BC patients, the reported prevalence of germline \(BRCA\,\,1/2\) \((gBRCA)\) pathogenic variants (PVs) is approximately 5%. \(^{64}\) When testing only for YWBC the proportion rises to 12.1%, \(^{65}\) and is even higher when considering only YWBC with triple-negative tumors in whom the prevalence of \(gBRCA\) PVs is reported in up to 23%. \(^{66}\)

Current National Comprehensive Cancer Network’s (NCCN) Guidelines on Hereditary Cancer Testing recommend testing for high penetrance BC susceptibility genes \((BRCA, CDH1, PLB2, PTEN, and TP53)\) for every woman with a personal history of BC diagnosed at 45 years or less. Hence, every YWBC should undergo a comprehensive genetic assessment. \(^{67}\) Furthermore, \(BRCA\) status might also impact treatment-related decisions. The EMBRACA, OlympiAD, and OlympiA phase III trials have shown that BC patients that carry a \(BRCA\,\,1/2\) germline mutation might be amenable to treatment with a poly (adenosine diphosphate–ribose) polymerase (PARP) inhibitor in both the advanced and localized settings. \(^{68-70}\) Unfortunately, diagnostic genomic tools and genetic counseling services require complex technology and specialized personnel that are not fully accessible in limited-resource settings. \(^{71}\)

Access to genetic counseling has been reported as an essential area to address among minorities. \(^{71}\) Although it has been described that the rate of mutations among breast and ovarian cancer patients does not differ by either ethnicity or race (except for Jewish Ashkenazi), non-Hispanic white females with a family history of breast/ovarian cancer are statistically more likely to be referred to genetic counseling, exemplifying prevailing disparities among underserved populations. \(^{72}\) Also, among those patients that were identified as \(BRCA\) PVs carriers, non-Hispanic whites were more likely to undergo cancer screening and risk-reducing surgeries compared to patients of other ethnicities. \(^{73}\) Furthermore, a qualitative report from the
US–Mexico border, an underserved area, described that only 3% of Latina women have knowledge about basic genetic concepts, and this was found to be irrespective of educational status. Among this group, only 1–4% of women had BRCA testing performed even though most of them declared they would undergo genetic testing within six months if it was offered free of cost.  

Barriers to genetic counseling reported from an underdeveloped setting in Europe are mainly related to limited access to healthcare and lack of integrated services. An important reason for patients to forego genetic testing could be the associated costs. A 2021 review by Grant et al. described that the proportion of patients who declined BC genetic testing due to cost ranged from 13.6–70.4%. In a survey of East Asian patients at high risk of developing BC (ie, personal history of breast/ovarian cancer or a known family member carrier for a BRCA PVs), 71.3% would not have undergone self-financed genetic screening, emphasizing the importance of sponsored cancer genetic testing services.

One of the main examples of how to address genetic counseling in a limited setting arises from the Genomic Risk Assessment for Cancer Implementation and Sustainment (GRACIAS) intervention in Mexico, which used implementation science methods to develop and deliver low-cost BRCA testing with a 115 BRCA panel (HISPANEL) and comprehensive genetic cancer risk assessment across four sites in Mexico. Patient eligibility for genetic assessment included a diagnosis of BC at a young age and triple-negative BC. Among the four recruiting sites in Mexico, over 1300 women received a comprehensive genetic risk assessment, with 15.6% identified as carriers of BRCA PVs. Furthermore, a model was adapted to evaluate the proportion of new patients meeting NCCN criteria who participated in genetic assessment among the four sites, reaching 76% to 90% of eligible patients. A critical finding from the GRACIAS intervention is the patients’ feedback recommending the development of Spanish resources to improve learning.

To this date, one of the main limitations for genetic testing in LMICs has been the lack of readily accessible tools. However, through the recent development of low-cost tests, improved access is expected soon. Nevertheless, the cost of risk-reducing strategies may still represent a barrier to the management of hereditary cancer syndromes. As the cost of genome sequencing is gradually descending, the cost of BC drugs is rapidly rising. To address this issue in a high-volume referral center in Latin America, discussions have taken place to find a solution for patient overload. However, only limited actions had taken place at the moment. Policymakers in Latin America need to carefully balance the benefits and costs of interventions in the rationale of cancer care.

Therefore, efforts should be made to generalize cost-accessible genetic testing and comprehensive counseling for all YWBC, irrespective of their economic background.

### QoL and Psychosocial Health

Quality of life (QoL) has been reported to be worse in YWBC compared to their older counterparts. YWBC experience more significant psychological distress characterized by higher levels of depression and anxiety over the future, particularly regarding motherhood and fear of cancer recurrence. Limited data from LMICs have been published on this topic, and even less information focusing exclusively on YWBC is available.

Results from the pilot phase of the “Joven & Fuerte” cohort showed that QoL improved significantly two years after BC diagnosis as measured by mean QLQ-C30 global QoL, emotional functioning, pain, and financial difficulty scales. However, physical functioning, nausea/vomiting, and constipation were found to be statistically worse.

Another Mexican study performed one-on-one semi-structured interviews at the National Cancer Institute in Mexico City, including 25 YWBC who had survived five or more years after diagnosis. The participating women reported psychological distress, anxiety at the transition to survivorship care, distress when entering the hospital, and explicit requests for psychological care. Notably, the participants expressed that unmet informational needs contributed to their anxiety.

Likewise, in another qualitative Mexican study, 29 YWBC recently diagnosed felt that they were not provided with enough information regarding BC diagnosis, treatment, and specific side effects affecting fertility, menopause, and sexuality. They reported that such lack of information further exacerbated uncertainty, distress, anxiety, and fear. The participants wished that information was delivered in an empathic and personalized manner, with additional take-home materials to help them remember, understand and/or expand information received during medical appointments. Similarly, another report from Mexican BC survivors revealed that 97% YWBC would like to receive more educational information in BC, with a particular interest in the adverse effects of treatment.
The development of tailored strategies for YWBC has yielded positive results among Mexican BC patients. After surveying YBCW who are part of the Mexican “Joven & Fuerte: Program for Young Women with Breast Cancer in Mexico”, 97% reported the program to be useful/very useful, and 82% believed that the information and support delivered by the program aided to better cope with their illness. Noteworthy, psychological support was identified as the most helpful resource according to participants.

YWBC represent an underserved population, not only regarding specific biologic needs such as fertility preservation and treatment-related sexual dysfunction but also suffer higher levels of anxiety and depression, leading to a worse QoL. Early interventions such as offering psychological counseling and further information on BC diagnosis and treatment may ameliorate these symptoms and enhance the patients’ experience.

Conclusions

There is scarce information regarding YWBC in limited-resource countries. Further research about the clinical behavior, as well as the supportive care and information needs of YWBC should be a priority, as this population represents a higher proportion of patients in LMICs. Evidence from HICs has shown that dedicated comprehensive programs for YWBC impact patients’ cancer and survivorship care and enhance patients’ experience. Only a few initiatives have been developed in Latin America, and much effort still needs to be made. Therefore, limited-resource countries could aim to replicate successful specialized programs for YWBC, especially in regions speaking the same language. Furthermore, partnerships between governments and non-profit organizations might also help bridging the resource gap in LMICs.

Many strategies and interventions to enhance clinical care in YWBC are costly and might need resource allocation for their implementation, hindering their adoption in limited economies. However, one high-yield and cost-effective intervention that might be useful in every scenario is education. YWBC have externalized on numerous reports their unmet need for information. Hence, every cancer-care team should develop tailored educational resources to optimize patient care.

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Martinez-Cannon and Barragan-Carrillo are both to be considered first authors as they contributed equally to this manuscript.

Disclosure

The authors report no conflicts of interest in this work.

References


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