The impact of socioeconomic status on the psychological distress of breast cancer patients in Kolkata

Arunima Datta¹, Rimi Sharma²

¹Dept. of Psychology, NetajiSubhas Chandra Bose Cancer Research Institute, 16A, Park Lane, Kolkata-700016, Kolkata, West Bengal, India.
²Dept. of Psychology, Neotia University, Kolkata, West Bengal, India.

ABSTRACT

Background: Breast cancer patients experience a variety of psychological symptoms such as anxiety, depression that affect their quality of life. The present study probed whether the socioeconomic status had any significance on psychological distress and quality of life of breast cancer patients in India.

Method: This was a cross-sectional and observational study, conducted over 111 breast cancer patients in a tertiary cancer hospital, Kolkata. Subjects were eligible to participate and they were of over 23rd years of age and had a histologically confirmed diagnosis of breast cancer had no history of recurrence or metastasis, and had no previous psychological problems. Age, marital status, residence, education, occupation and family income were assessed for patients’ socio economic status. Psychological symptoms and quality of life were assessed by using validated tools. Means and standard deviations of each outcome were compared by socioeconomic status and multivariate linear regression models for evaluating the association between socioeconomic status, psychological distress and quality of life.

Results: The patient group was categorized into three groups based on their income level as follows: ≤500=27.9%, 500-1000=51.4% and high income = ≥1000=20.7%. A total of 111 breast cancer patients participated in the study; the mean age of participants was 49.7 (SE=1.31). After using multivariate logistic regression with fully adjusted models, the patients who were living alone had significantly higher level of depression and poor quality of life compared to those who are employed. Higher education and family income both positively associated with quality of life after adjusting for age, marital status and occupation.

Conclusion: The findings revealed that the family income have a significant impact on the psychological distress of the cancer patients.

Keywords: Breast cancer; psychological factors; socioeconomic

1. Introduction

Generally in India, there is limited number of investigations done about the impact of socioeconomic status and its impact on the breast cancer patients. Although many studies have been conducted on cancer patients in general, the number of studies done on breast cancer patients, specifically, is limited. Following are some of the studies conducted in order to understand the link between SES and psychological distress among cancer patients: This study is to understand the impact of socioeconomic status on the Psychological distress of cancer and cardiovascular disease patients in Kerala region of India. This study was conducted on a total of 225 women patients belonging to Kerala, in the year 2016 by Alex. The findings of this study were that individuals belonging to a low socioeconomic background were more concerned about the expenses that were to be born for their treatment. They were also more worried about managing their day-to-day expenses. Thus, the study revealed that the socioeconomic status does play an important role in the Psychological distress of an individual with cancer and cardiovascular disease experiences.
Oliver Chang et al., in 2014 had worked over association between socioeconomic status and altered appearance distress, body image and quality of life among breast cancer patients. Through their study they had said that socioeconomic status is significantly associated with altered appearance distress, body image, and quality of life in Korean women with breast cancer. Patients who suffer from altered appearance distress or lower body image are much more likely to experience psychosocial, physical, and functional problems than women who do not, therefore health care providers should be aware of the changes and distresses that these breast cancer patients go through and provide specific information and psychosocial support to socioeconomically more vulnerable patients.

Socioeconomic determinants of health have been discussed in the literature, and the need to tailor patient care has been widely expressed (Preston, 2010). Studies regarding socioeconomic status and its differing effects on the health of breast cancer survivors have focused primarily on assessing clinical and pathologic characteristics, quality of life, depression, and anxiety (King et al., 2000; Karakoyun-Celik et al., 2010; SchmidBuchi et al., 2013; Wang et al., 2013).

In the year 2009, Limin et al conducted a study on the impact of socioeconomic status on cancer incidence and stage at diagnosis: selected findings from the surveillance, epidemiology and end results. Through their study they concluded that individuals (men and women) whose educational levels were lower than high school degrees, and also those individuals who had an annual income lower than $12500, had higher rates of cancer incidence as compared to the individuals who had acquired college degrees, and also from the group of people having annual income of more than $50,000. They also concluded that lower income was significantly a factor for the increased chances of distant-stage breast cancer in people.

Psychological well-being and the quality of life he or she has, influenced to a large extent by the impact of socioeconomic status. It is known that low socioeconomic status is linked with more occurrences of stressful events and the lack of coping methods (Eron et al 1982). It is seen that poor coping skills and the life events that the individual faces are correlated with psychological well-being and their quality of life. People belonging to a higher socioeconomic status have been seen to have lower chances of acquiring and suffering from most psychiatric disorders and medical conditions. They have been known to have lower mortality rates from these conditions, which also includes cancer (Adler et al., 1994).

We have followed their study to investigate the impact of SES on breast cancer patients of a different population- the eastern region population of West Bengal. How this psychological states as well as socioeconomic status interfere in a patient’s quality of life was also discussed.

2. Materials and Methods

Design:

This was observational study was designed using validated tools and structured face to face interview schedule.

Patient Selection:

For the present study participants were recruited between July 1 and October 30, 2017, from NetajiSubhas Chandra Bose Cancer Research Institute, Kolkata.

The researchers had approached 168 breast cancer survivors; 131 (72.02%) agreed to participate in the study. Among them, women who had previous history of psychological problem (n=3), recurrence of the disease (n=6), or who were stage 4 (n=11) were excluded from the study, resulting in a total of 111 study participants.

To evaluate the impact of socioeconomic status on psychological problems, we used the data of the only 111 diagnosed with breast cancer within less than 12 months. Their consent was also taken. Demographic information was taken from the patients including, their age, marital status, residence, and income. On the basis of their per capital income, patients were divided into three groups: below 500, 500 to 1000 and above 1000. All patients were
made to go through the assessment of depression, anxiety and quality of life. Inclusion and exclusion criteria are
given below.

Inclusion Criteria
- Patients diagnosed with breast cancer;
- Permanent resident of India
- People belonging to initial stage, or adverse stage of cancer;

Exclusion
- People with any other forms of cancer, patients with previous history of mental disorders or illness; newly
diagnosed breast cancer patients.

Data collection
The structured interview was conducted using proper performa for taking demographical and
clinoco-pathological details. After that depression, anxiety and quality of life were assessed by using validated
tools.

STAI-II: State-Trait Anxiety Inventory
The Bengali version of State-Trait Anxiety Inventory was used on the patients. It is a 40-item self-report
questionnaire. Every item is rated by the patient on a 4-point rating scale. The state version of the inventory
measures a temporary anxiety state of the patients, whereas the trait version of the inventory measures a long-term
and more stable anxiety experience of the patients. (Eskelinen M et al., 2011)

Beck Depression Inventory (BDI): This is a 21 item questionnaire consisting of six sub-scales, Mild mood
disturbance, Borderline clinical depression, Moderate depression, severe depression and Extreme depression.
Each item is rated in a 3 point scale. (Eskelinen M et al., 2011)

EORTC : European Organization for Research and Treatment of Cancer breast cancer-specific quality-of-life
questionnaire module

EORTC QLQ-BR23 is a questionnaire used to measure the quality of life in cancer patients. It is a
self-administered questionnaire administered on breast cancer patients who are at various stages of treatment.
There are 5 domains that the questionnaire administers: sexuality, body image, breast symptoms, arm symptoms,
and systemic therapy side effects. (May Leng Tanet al. 2014)

Procedure
After maintaining exclusion and inclusion criteria, patients were selected for the present study. Ethical
concern was approved following ICMR rule. Demographic information were taken from both groups (including,
age, marital status, residence, occupation, family type and family income). After taking data, use proper statistics
for making conclusion.

3. Statistical Analysis
We scored anxiety, depression and quality of life according to the manual of the Beck Depression Inventory,
State trait Anxiety Inventory and EORTC QLQ-BR23 respectively. For assessing the impact of socioeconomic
status on depression , anxiety and quality of life, we calculated the mean and standard error of the each response
and compared them with socioeconomic status. To analyze the relation between socioeconomic status and anxiety,
depression and quality of life we used multivariate linear regression model. In the multivariate analysis, we
adjusted for variables that could be potential variables: age, residence, relationship status, education, occupation
and family income. All statistical analyses were performed using SPSS program version 21. Statistical significance
was defined as p<0.05.
Table 1. Socio demographic characteristics of Participants (N = 111)

<table>
<thead>
<tr>
<th>Socio demographic factor</th>
<th>No of cases</th>
<th>BDI (Mean±SE)</th>
<th>STAI (Mean±SE)</th>
<th>FORTC QLQ-BR23 (Mean±SE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (&lt;30 years)</td>
<td>10 (9.0%)</td>
<td>35.2±9.2</td>
<td>50.1±6.6</td>
<td>17.4±1.2</td>
</tr>
<tr>
<td>30-40</td>
<td>35 (31.5%)</td>
<td>28.4±2.4</td>
<td>49.4±3.1</td>
<td>17.7±1.2</td>
</tr>
<tr>
<td>40-50</td>
<td>21 (18.9%)</td>
<td>30.1±2.1</td>
<td>48.4±3.0</td>
<td>15.0±0.9</td>
</tr>
<tr>
<td>50-60</td>
<td>23 (20.7%)</td>
<td>28.3±1.6</td>
<td>49.1±3.6</td>
<td>16.5±1.0</td>
</tr>
<tr>
<td>&gt;60 years</td>
<td>22 (19.8%)</td>
<td>35.0±1.2</td>
<td>44.5±2.1</td>
<td>17.0±1.3</td>
</tr>
<tr>
<td>Marital Status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Living with spouse</td>
<td>88 (77.5%)</td>
<td>28.6±2.1</td>
<td>48±2.2</td>
<td>16.9±0.6</td>
</tr>
<tr>
<td>Living without spouse</td>
<td>23 (22.5%)</td>
<td>30.6±1.6</td>
<td>49.7±2.6</td>
<td>18.7±1.3</td>
</tr>
<tr>
<td>Residence</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Rural</td>
<td>55 (49.5%)</td>
<td>27.8±1.6</td>
<td>49.0±2.8</td>
<td>17.3±0.8</td>
</tr>
<tr>
<td>urban</td>
<td>56 (50.5%)</td>
<td>31.3±2.7</td>
<td>49.7±2.6</td>
<td>17.3±0.7</td>
</tr>
<tr>
<td>Education</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Primary School</td>
<td>49 (44.1%)</td>
<td>36.3±1.2</td>
<td>39.0±0.9</td>
<td>17.3±0.8</td>
</tr>
<tr>
<td>Higher Secondary</td>
<td>37 (33.3%)</td>
<td>35.6±1.3</td>
<td>41.3±2.1</td>
<td>16.7±1.0</td>
</tr>
<tr>
<td>Graduate</td>
<td>25 (22.5%)</td>
<td>34.2±2.4</td>
<td>41.2±5.3</td>
<td>18.3±1.2</td>
</tr>
<tr>
<td>Precipitate Family Income</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;500</td>
<td>31 (27.9%)</td>
<td>41.7±4.7</td>
<td>53.8±2.7</td>
<td>13.7±1.1</td>
</tr>
<tr>
<td>500-1000</td>
<td>57 (51.4%)</td>
<td>39.5±1.4</td>
<td>54.3±2.8</td>
<td>20.8±0.8</td>
</tr>
<tr>
<td>&gt;1000</td>
<td>23 (20.7%)</td>
<td>37.3±2.9</td>
<td>45.3±1.8</td>
<td>21.2±1.3</td>
</tr>
</tbody>
</table>

4. Results

Table 1 showed the distribution of sociodemographic factors among breast cancer patients. According to age group, it showed that 9% patients belonged to age group <30 years; 31.5% belonged to 30-40 years; 21% belonged to age-group of 40-50 years; 23% belonged to age-group of 50-60 years and 22% patients were above the age of 60.

In the marital status bar graph, we can see that 77.5% people were living with their spouses, and 22.5% people were living alone. The chart of education level indicates that 44.1% individuals had primary level of education, 33.3% people had completed high school degrees and 22.5% people were graduate degree holders.

Out of the total, 49.5% lived in rural areas, whereas 50.5% lived in urban areas. Through this, we found that out of the total breast cancer patients, 20.7% people belonged to income group 3 that is whose monthly income was more than 10000 INR. 51.4% people belonged to income group 2, whose monthly income was between 5000-10000 and 27.9% people belonged to income group 1, with income below 5000 INR.

Breast cancer patients with less education suffered from higher level of psychological distress than those with higher education: depression: (36.3 vs 34.22); anxiety: (39.02 vs 41.25); quality of life: (17.31 vs 18.36) . In terms of monthly family income, patients who had higher income (>1000) reported better psychologically stable than patients with lower income (<500) depression: (41.71 vs 37.36); anxiety: (53.8 vs 45.33); quality of life: (13.71 vs 21.21), and it was statistically significant. Although major psychological symptoms as well as quality of life seemed to be affected by a patient’s age, relationship status and her residence.

Table 2 shows the distribution of clinicopathological factors of breast cancer patients. The menopausal status of the breast cancer patients shows that in pre-menopausal state, 44.1% women were present in total, and 55.9% patients were there in post-menopausal state.

In the histology bar graph, it can be seen that 81.9% patients had ductal and 18.9% patients had lobular problems. The estrogen receptor bar graph displays that 69.4% patients had positive and 30.6% patients had negative feedbacks. In progesterone bar graph we see that a total 65.8% patients had positive and 34.2% patients had negative feedback. The Her2/neu status bar graph showed that over expression occurred in 64.9% patients and
non-over expression was seen in 35.1% patients.

<table>
<thead>
<tr>
<th>Clinical Characteristic</th>
<th>No. of Cases</th>
<th>BDI (Mean±SE)</th>
<th>STAI (Mean±SE)</th>
<th>EORTC QLQ-BR23 (Mean±SE)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Menopausal Status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pre</td>
<td>49(44.1%)</td>
<td>33.6±5.85</td>
<td>52.09±1.93</td>
<td>19.32±0.96</td>
</tr>
<tr>
<td>Post</td>
<td>62(55.9%)</td>
<td>32.1±2.2</td>
<td>53.55±0.41</td>
<td>20.63±0.77</td>
</tr>
<tr>
<td>Histology</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Ductal</td>
<td>90(81.5%)</td>
<td>31.51±2.1</td>
<td>46.14±1.89</td>
<td>20.41±0.71</td>
</tr>
<tr>
<td>Lobular</td>
<td>21(18.5%)</td>
<td>36.4±5.05</td>
<td>48.11±1.33</td>
<td>19.1±1.18</td>
</tr>
<tr>
<td>Estrogen Receptor</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>77(69.4%)</td>
<td>34.2±2.11</td>
<td>49.12±1.79</td>
<td>20.23±0.72</td>
</tr>
<tr>
<td>Negative</td>
<td>34(30.6%)</td>
<td>37.4±2.74</td>
<td>48.71±1.12</td>
<td>19.04±1.12</td>
</tr>
<tr>
<td>Progesterone Receptor</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Positive</td>
<td>73(65.4%)</td>
<td>32.2±1.17</td>
<td>50.67±2.98</td>
<td>20.23±1.10</td>
</tr>
<tr>
<td>Negative</td>
<td>38(34.6%)</td>
<td>33.6±1.89</td>
<td>48.24±1.23</td>
<td>19.7±1.01</td>
</tr>
<tr>
<td>Her2/neu status</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Over express</td>
<td>72(64.9%)</td>
<td>30.8±1.11</td>
<td>47.64±2.72</td>
<td>20.3±0.74</td>
</tr>
<tr>
<td>Not over express</td>
<td>39(35.1%)</td>
<td>36±2.52</td>
<td>50.8±2.46</td>
<td>19.5±1.07</td>
</tr>
<tr>
<td>Number of chemotherapy</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>≤6th cycle</td>
<td>74(71.2%)</td>
<td>31.8±1.59</td>
<td>47.53±2.1</td>
<td>19.9±0.75</td>
</tr>
<tr>
<td>More than 6th</td>
<td>32(28.8%)</td>
<td>28±1.5</td>
<td>43.78±1.95</td>
<td>20.2±0.98</td>
</tr>
<tr>
<td>Treatment Plan</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Neo-adjuvant chemotherapy</td>
<td>68(61.3%)</td>
<td>38.0±1.02</td>
<td>48.1±1.89</td>
<td>21.1±1.11</td>
</tr>
<tr>
<td>Adjuvant chemotherapy</td>
<td>43(38.7%)</td>
<td>32.1±1.96</td>
<td>47.7±2.05</td>
<td>20.2±0.78</td>
</tr>
</tbody>
</table>

Table 2. Clinicopathological Factors of Participants (N = 111)

When considering the number of chemotherapy, it was seen that 71.2% patients had 6 or less numbers of chemotherapy and 28.8% patients had taken more than 6 chemotherapy. In the treatment plan graph, neo-adjuvant chemotherapy was done for 61.3% patients and 38.7% patients had got adjuvant chemotherapy.

When considering number of chemotherapy, it can be seen that patients with 6 or less chemotherapy had higher depression score than those with more than 6 chemotherapy (31.82 vs 28). Patients with 6 or below chemotherapy also displayed higher anxiety score than those with more than 6 chemotherapy (47.52 vs 43.78). The score on quality of life was also lower in cases of patients receiving 6 or less numbers of chemotherapy than those with more than 6 chemotherapy (19.9 vs 20.22).

When considering the treatment plan, it was observed that the depression levels was higher in neo-adjuvant chemotherapy than in adjuvant chemotherapy patients (33.62 vs 32.19). The score of anxiety was also marked higher in neo-adjuvant chemotherapy patients than in adjuvant chemotherapy patients (49.19 vs 47.78). The score of quality of life was surprising higher than that of adjuvant chemotherapy treatment patients (21.1 vs 20.3).

Factors associated with psychological symptoms including anxiety and depression and quality of life. The results of multivariate analysis of factors associated with psychological symptoms and quality of life are displayed in Table 3. With the adjustment for demographic factors including age, relationship status, residence, education and socioeconomic status as well as clinic pathological factors such as menopausal status, current treatment status, and number of chemotherapy was significantly associated with altered appearance distress, body image, and quality of life. The patients who had higher level of family income (>1000) were more likely to have better psychological management (OR=1, 95% CI (2.83-3.06)* than patients who had lower income (<500). In case of quality of life, patients living without spouse had poorer quality of life ((OR=1.5, 95% CI (5.83-7.02) than patients living with a spouse (p<0.05). Patients with graduate level of education had better quality of life by 7.16 points than patients with primary level of education after adjusting for all other factors, and it was statistically significant (p<0.01).
5. Discussion

Oliver Chang et al., in 2014 had worked over association between socioeconomic status and altered appearance distress, body image and quality of life among breast cancer patients. Through their study they had said that socioeconomic status is significantly associated with altered appearance distress, body image, and quality of life in Korean women with breast cancer. Patients who suffer from altered appearance distress or lower body image are much more likely to experience psychosocial, physical, and functional problems than women who do not.

Booth et al presented their study impact of socioeconomic status on Stage of Cancer at Diagnosis Survival in the year 2010. The study was conducted in Ontario, Canada. They studied the impact of SES on cancer patients of Canada. They investigated that even though the government provided the residents with overall world class healthcare services, the residents’ SES did play an important role in the survival rates of the patients. This study focused on forms of cancer, such as, breast, colon, NSCL, cervical and laryngeal cancers.

Following above mentioned studies we want to describe the impact of socioeconomic characteristics on psychological problems and quality of life among 111 breast cancer patients who experienced anxiety and depression during the period of cancer treatment in West Bengal. We found that a patient’s educational level and family income were positively associated with psychological problems. As a result it affects their quality of life.

In West Bengal, most of the women after marriage occupationally are housewives. They are more depending on their husband economically rather than emotionally. In such situation when they had diagnosed as a breast cancer patient and when they see their family members devote everything for getting well soon but they are helpless. As a result their response was “I want to live for my family members not for that I want to live. I want to make them successful. But, unfortunately I cannot… due to day to day deterioration of health”. So, in this situation socioeconomic status laid an import role for developing psychological distress and that affect their quality of life. Support with numerous previous findings in our study, quality of life increased for patients who were married and for patients with more education than patients who were living alone or patients with less education (Janz et al., 2005; Ahn et al., 2009; Ashing-Giwa and Lim, 2009; Salonen et al., 2009).

Education of the practical situation as well as for cancer is very important for leading a better quality of this result had contradicted with Ashing-Giwa and Lim, 2009. They had found that physicians spend more time with affluent and educated patients than with financially and educationally deprived patients who actually need more attention and care. For people belonging to low socioeconomic statuses affording the expenses of cancer care for a prolonged period of time can prove to be a costly affair. Depending on their limited income for not only providing for daily expenditures, but also for an extra cost of medications, treatments and hospital visits can be an independent factor in development of psychological symptoms such as anxiety or depression. Also, for the breast cancer patients for whom the expenses are to be made may also feel responsible, and hence, helpless. The result of the present study was supported with Oliver Chang et al. 2014.The results of our present study indicate that socioeconomic factors could be better in dependent factors of a breast cancer patient’s development of psychological problems and quality of life than clinical characteristics.

Most cancers are related to socioeconomic status with low SES individuals more at risk. Moreover, even incidence rates for major cancers have slowed or decreased, declines to mortality have been slow in minority population than among whites (Glanz et al., 2003). One of the most potent risk factors for early disease disability and death is low socio-economic status (SES). From birth throughout life, those who are born into the lower social classes experience more and more intense stressors of all kinds, which have a cumulative toll on health risks. Lower income and poor educational and occupational attainment leads to exposure to a broad array of stressors including inadequate housing, violence, danger, lack of vital goods and services, inadequate medical facilities, poor sanitation, exposure to environmental pollutants and numerous other hazards. Leigh Ann Simmons et al., (2009) examined factors associated with self-reporting depression for low-income rural women experiencing
depressive symptoms. Women reporting depression were significantly likelier to report physical health problems, injury/illness, and more frequent physician visits. Low socio-economic status is associated with a higher prevalence of depression, but it is not yet known whether change in socio-economic status leads to a change in rates of depression.

Vincent Lorant (2007) assessed whether longitudinal change in socio-economic factors affects change of depression level, socio-economic factors based on material standard of living, education, employment status and social relationships. A lowering in material standard of living between annual waves was associated with increase in depressive symptoms and case ness of major depression. Life circumstances also influenced depression. Ceasing to cohabit with a partner increased depressive symptoms and caseness, and improvement in circumstances reduced them; the negative effects were stronger than the positive ones. The study showed a clear relationship between worsening of socio-economic circumstances and depression.

6. Conclusion
Through the findings of this study, we aim to also acquire any kind of government funding for the patients belonging to low socioeconomic statuses for better possibilities and opportunities of their treatment. This will eliminate the chances of factors such as becoming a liability for their family, or dependency on family for bearing the costs of cancer treatment, to become a factor for developing psychological symptoms.

Reference


