Quality of Life of Women with Breast Cancer and Socio-Demographic Factors

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Abstract

Background. Breast cancer treatment is an aggressive therapy that affects the deterioration of women’s quality of life (QOL) in many areas. Knowledge about factors that influence the assessment of the QOL is of particular importance. The aim of the study was to analyse areas of the quality of life of women with breast cancer, taking into account social and demographic factors. Methods: The research was carried included 324 women with breast cancer. The research was carried out using a diagnostic survey, the author’s questionnaire and a standardized questionnaire for measuring the QOL of women treated for breast cancer, ie the European Organization for Research and Treatment of Cancer, Quality of Life Questionnaire (EORTC) QLQ-C30 and QLQ-QLQ module BR23. Statistical analysis uses Statistica 10.0 software. The results were considered statistically significant when the calculated probability met the inequality of $p < 0.05$. Results: A lower QoL was found in women with diagnosed breast cancer. With age, the QoL score of respondents in the area of physical, sexual and hair loss decreased, and the following symptoms intensified: pain, insomnia, lack of appetite and shoulder ailments. The age of the respondents positively correlated with the image of their own body and social functioning. Higher QOL (higher values of functional scales and lower intensity of symptoms) were found in patients in relationships (in the scope of: cognitive functioning, future prospects), with higher education (in the scope of: physical, emotional, cognitive and sexual functioning) and declaring very good situation financial (in the scope of: physical, emotional, cognitive functioning, performing social roles). Conclusions: Age, marital status, education and financial situation influenced the QOL of women with breast cancer. In the care of women with breast cancer, attention should be paid to the individualisation of the therapeutic process, with particular emphasis on psychotherapy and support by social services.

Keywords: Breast cancer- quality of life- socio-demographic factors

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Introduction

On a global scale, the incidence of breast cancer is assessed as one of the most common cancer diseases of women. The most current data indicate 1,700,000 new diagnoses of breast cancer per year. About 25% of oncological diagnoses among women are breast cancer (Ginsburg et al., 2017; Ferlay et al., 2015). In the world over the last 25 years, an increase of around 30% in the incidence rates in both developed and developing countries has been found (Harbeck and Gnant, 2017; Siegel et al., 2016).

Breast cancer is also a major threat to women in Poland. The incidence of this type of cancer has more than doubled during the last 30 years, and currently nearly 20,000 new cases of breast cancer are diagnosed annually in Poland (Didkowska and Wojciechowska, 2013). The diagnosis of breast cancer is associated with the need to implement long-term and aggravating treatment. Most often it includes surgery, chemotherapy, radiotherapy and causes side effects - transient or permanent, i.e. pain, fibrosis of tissues or limitation of physical activity. Oncology therapy is usually aggressive and causes fear in patients due to uncertainty about the effects of treatment and side effects. Women are afraid of disability and death, as well as the breakdown of the family. They are also often accompanied by anger, aggression, depressed mood and a sense of self-esteem (Kyrano et al., 2013). After surgical treatment due to high trauma and mental anxiety caused by total or partial loss of breasts, a “half-body / body complex” and self-esteem can occur, especially in the social aspect (Fontes et al., 2018).

Currently, medicine assumes a holistic approach to therapy, which takes into account the treatment of not only the body, but also the psyche, and the goal of treatment is to extend life as well as improve QOL (Menen, 2016). The assessment of the mental condition, emotional state of women and the acceptance of new health conditions

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by them can be made through QOL examination. In the face of alarming statistics and a significant increase in the number of women with breast cancer, knowledge about determinants affecting the QOL of women with this cancer is becoming more and more important. Their thorough knowledge and analysis can be used to take action to ensure the best possible comfort of life during illness. The results of such tests may also be used in the selection of appropriate methods of treatment and rehabilitation of patients (Pinto et al., 2011). Including the QOL assessment for all medical care positively affects the relationship between the patient and the doctor and strengthens the effectiveness of the therapy (Ahn, 2007). The results of these studies are very important because contemporary oncology is focused on getting the greatest chance for cure or long-term survival, while taking into account high QOL (Visser et al., 2006). A limited number of studies considering the relationship between QOL and socio-demographic factors of patients results in low individualization of the therapeutic process. Currently, the QOL assessment is treated as one of the endpoints of clinical trials. Due to the continuous progress of medicine and the frequently changing understanding of psychological mechanisms affecting QOL, such research should be constantly conducted.

The aim of the study was to analyse areas of the quality of life of women with breast cancer, taking into account social and demographic factors and their potential influence on the QoL.

Materials and Methods

Place of research, characteristics of the studied population, the process of data collection

The research was carried out in Father B. Markiewicz Podkarpackie Oncological Centre in Brzozów (Poland). The study included a group of 350 women with histopathologically confirmed breast cancer. Due to the lack of questionnaires in the questionnaire, 324 correctly completed and complete research tools were accepted for analysis. Before the examination, all patients were informed about their purpose, assured about confidentiality, anonymity and voluntary participation in the research. In addition, the examined women were informed about the possibility of resigning from the study at every stage. The study included women who had histopathologically confirmed diagnosis of breast cancer, agreed to participate in the study and were treated at Father B. Markiewicz Podkarpackie Oncological Centre in Brzozów. Criteria disqualifying a patient to participate in the study were the following factors: the occurrence of cancer other than breast cancer in the last 5 years, palliative treatment and the patient’s lack of consent for participation in the study. The consent of the director of the Father B. Markiewicz Podkarpackie Oncological Centre in Brzozów. The project also received a positive opinion of the Bioethic Committee.

Research tools

The research was carried out by means of a diagnostic survey using the author’s questionnaire and a standardized questionnaire for measuring the quality of life of women treated for breast cancer, i.e. European Organization for Research and Treatment of Cancer, Quality of Life Questionnaire (EORTC) QLQ-C30 (quality of life questionnaire) and the QLQ-BR23 module (breast cancer) (Aaronson et al., 1993). The EORTC approval was obtained for the use of the QLQ-C30 and QLQ-BR23 questionnaires. In Poland, an assessment of the validity and reliability of the QLQ-C30 questionnaire and its version of BR 23 was carried out, which confirmed the validity of their use during QOL assessment of breast cancer patients (Zawiszas et al., 2007). This fact was decisive when choosing a standardized research tool. The QLQ-C30 questionnaire is used to determine QOL and the sense of (health) state, assess the functioning in the physical, emotional and social dimensions of people with a novel. The QLQ-C30 survey consists of a general scale assessing the state of health and quality of life, 5 functional scales, 3 symptomatic scales and 6 single points (questions) also determining the intensity of symptoms. There are 30 questions in the survey. The answers to most of them are defined on a 4-point scale, i.e. at all (1), a little (2), significantly (3), very (4), which assess the severity of the analyzed parameters. In the last two questions (numbers 29, 30), the patients assessed their health and general QOL on a scale from 1-7 (1 - very bad / bad, 7 - excellent / perfect). The EORTC QLQ-BR23 scale is used to determine the quality of life of women with breast cancer and is a complementary module for the QLQ-C30. The questionnaire contains 23 questions to be answered in a 4-step scale, i.e., at all (1), a little (2), significantly (3), very (4). For each question, the patient chose one answer. EORTC QLQ-C30 and QLQ-BR23 questionnaires were developed statistically in accordance with the EORTC guidelines. For each patient, the raw ratio and linear transformation were calculated in order to obtain the value of the coefficient (score), whose value for both scales and individual symptoms could range from 0 to 100. In the case of functional scales, their higher coefficient corresponds to a better one (higher) level of functioning, while the higher the ratio (score) for symptomatic scales and individual symptoms, the greater the severity of the symptom and the sick one feels worse.

Statistical analysis

Statistica 10.0 (StatSoft Inc., 2011) was used for statistical analysis. The basic measures of descriptive statistics were calculated: the arithmetic mean (M), the median (Me) and the standard deviation (SD). Conformity of the distribution of quantitative variables with the normal distribution was studied using the Shapiro-Wilk test. In the situation when the assumptions regarding the use of parametric methods to verify statistical hypotheses were not used, nonparametric methods were used. The following tests were used: Mann-Whitney U test, Kruskal-Wallis test (together with Dunn post-hoc test) and Spearman’s rank correlation coefficient. The significance level $\alpha = 0.05$ was assumed. The results were considered statistically significant when the calculated probability $p$ was satisfied by the inequality of $p < 0.05$. 

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**Results**

The average age of women enrolled in the study was 52.4 (SD = 13.7) years, with the youngest patient being 26 years old and the oldest 75 years old. The largest group were respondents above 60 years of age (29.4%), while the smallest group were those aged 20-30 (3.7%). The surveyed respondents were mostly city dwellers (54.3%). Most patients lived in cities up to 10,000 residents (28.1%) and cities up to 50,000 inhabitants (17.6%), and the smallest percentage were women living in cities over 50 thousand residents (8.6%). The study group was dominated by women living in a relationship (66.1%) and with the average (35.2%) and higher education (33.3%). Most of the surveyed women stayed at work in a state-owned factory (35.5%). Almost 40% of the surveyed women rated their financial situation as very good, while 30.6% as good, and 29.6% as bad. Among the women included in the study the most were those who had two children (32.7%), slightly fewer had one (21.3%) and three (13.3%) children, while the least four (8.0%) and more (4.9%). Almost 1/5 of respondents did not have children (Table 1).

In 71.3% (231) of the examined patients, surgical treatment was used, whereas in 112 women (34.6%) mastectomy was performed, whereas in 119 patients (36.7%) the surgery was performed. The time from surgery in more than a half of the respondents (54.1%) was over 1 to 2 years, in 22.9% - less than a year, for 12.6% women - over 3 years, and for 10.4% women this was above 2 to 3 years (Table 2).

**Quality of life and marital status**

A lowering of health status and QOL (M = 52.1 - unmarried women, M = 54.8 - women in relationships). There were statistically significant differences between the functional and symptomatic rooms QLQ-C30 and QLQ-BR23 depending on the marital status of the studied women (Table 4). Despite the fact that the general assessment of QOL in free and

| Table 1. Sociodemographic Characteristics of the Studied Group |
|-----------------|----------------|-----|
| No. | Characteristics          | Category                        | N  | % |
| 1   | Age (years of age)       | 20–30                          | 12  | 3.7|
|     |                             | 31–40                          | 67  | 20.6|
|     |                             | 41–50                          | 70  | 21.6|
|     |                             | 51–60                          | 80  | 24.7|
|     |                             | Over 60                        | 95  | 29.4|
|     | M (SD)                     | 52.4 (13.7)                    |     |    |
|     | Min–Max                    | 26–75                          |     |    |
| 2   | Dwelling place            | Country/Village                | 148 | 45.7|
|     |                             | Town of up to 10 thousand dwellers | 91  | 28.1|
|     |                             | Town of up to 50 thousand dwellers | 57  | 17.6|
|     |                             | Town/City of over 50 thousand dwellers | 28  | 8.6 |
| 3   | Marital status            | Single                         | 110 | 33.9|
|     |                             | In relation                    | 214 | 66.1|
| 4   | Education                 | Basic vocational               | 102 | 31.5|
|     |                             | Secondary/post gymnasium       | 114 | 35.2|
|     |                             | Higher                         | 108 | 33.3|
| 5   | Source of income          | Work on your own farm          | 31  | 9.6 |
|     |                             | Work at a state-owned factory / institution | 115 | 35.5|
|     |                             | Own business                   | 29  | 8.9 |
|     |                             | Retirement                     | 51  | 15.7|
|     |                             | Pension                        | 66  | 20.4|
|     |                             | Other                          | 32  | 9.9 |
| 6   | Financial situation       | Very good                      | 129 | 39.8|
|     |                             | Good                           | 99  | 30.6|
|     |                             | Bad                            | 96  | 29.6|
| 7   | Number of children        | 0                             | 64  | 19.8|
|     |                             | 1                             | 69  | 21.3|
|     |                             | 2                             | 106 | 32.7|
|     |                             | 3                             | 43  | 13.3|
|     |                             | 4                             | 26  | 8.0 |
|     |                             | More than 4                    | 16  | 4.9 |
in a relationship was similar, the differences in the value of individual scales indicate better functioning of women in relationships. Women remaining in relationships rated the cognitive functioning higher (M = 65.2-vacant, M = 73.0 in the compound, p = 0.020). On the other hand, in women, higher intensity of vegetative and physical symptoms was observed: nausea and vomiting (M = 42.6), pain (M = 28.6), dyspnoea (M = 24.5), lack of appetite (M = 45.2), constipation (M = 19.7), as well as insomnia (M = 44.5). The unmarried women were also more strongly affected by financial problems (M = 43.9) compared to women in relationships (M = 32.4). The shown differences were confirmed statistically Table 3).

Based on the QLQ-BR23 questionnaire, the respondents rated the body image the best, while the sexual functioning was the lowest. In the case of symptoms also assessed using the QLQ-BR23 questionnaire, the women surveyed showed the discomfort caused by hair overgrowth and the side effects of systemic treatment to the highest degree. Women remaining in relationships statistically significantly higher evaluated the future perspective (M = 37.6) in comparison to free women (M = 27.6). At the same time, patients in relationships experienced greater stress due to hair loss (M = 74.2) (Table 3).

| Table 2. Characteristics of the Examined Group in Terms of Medical Characteristics |
| No | Characteristics | Category | N | % |
|----|----------------|----------|---|---|
| 1  | Surgical treatment | Yes | 231 | 71.3 |
|    | | No | 93 | 28.7 |
| 2  | Type of an operation | Mastectomy | 112 | 34.6 |
|    | | A saving operation | 119 | 36.7 |
| 3  | Time from surgery (in years) | Less than 1 year | 53 | 22.9 |
|    | | Above up to 2 years | 125 | 54.1 |
|    | | Over 2 to 3 years | 24 | 10.4 |
|    | | Over 3 years | 29 | 12.6 |
| 4  | Chemotherapy | Yes | 278 | 85.8 |
|    | | No | 46 | 14.2 |
| 5  | Radiotherapy | Yes | 94 | 29.0 |
|    | | No | 230 | 71.0 |
| 6  | Hormone therapy | Yes | 56 | 17.3 |
|    | | No | 268 | 82.7 |

Table 3. Women's Quality of Life Assessment - Categories Related to QLQ-C30 and BR 23 and Marital Status

| Functional scales and symptoms QLQ-C30 i QLQ BR-23 | n | 110 | 214 |
|---------------------------------------------------|---|-----|-----|
| Health status and quality of life                 | 0.297 | 52.1±21.3 | 50 | 54.8±18.9 | 50 |
| Functional scales                                 |   |   |   |   |   |
| Physical functioning                              | 0.184 | 71.8±21.6 | 73.3 | 76.4±15.8 | 80 |
| Performing social roles                           | 0.424 | 71.5±25.7 | 66.7 | 75.1±21.3 | 66.7 |
| Emotional functioning                             | 0.422 | 57.7±27.6 | 66.7 | 60.9±23.5 | 66.7 |
| Cognitive functioning                             | 0.02 | 65.2±28.2 | 66.7 | 73.0±23.6 | 83.3 |
| Social functioning                                | 0.637 | 67.7±31.3 | 66.7 | 71.0±27.2 | 66.7 |
| An image of your own body                         | 0.248 | 63.9±34.7 | 66.7 | 60.4±32.1 | 66.7 |
| Sexual function                                   | 0.264 | 15.8±23.2 | 0 | 18.4±23.7 | 0 |
| Sexual satisfaction                               | 0.749 | 45.3±33.2 | 33.3 | 46.8±27.2 | 33.3 |
| The prospect of the future                        | 0.011 | 27.6±32.8 | 0 | 37.6±35.0 | 33.3 |

| The scale of the symptoms                         |   |   |   |   |   |
| Fatigue                                           | 0.221 | 40.5±22.2 | 33.3 | 36.0±18.8 | 33.3 |
| Nausea / vomiting                                  | 0.008 | 42.6±40.6 | 33.3 | 28.9±34.6 | 16.7 |
| Pain                                              | 0.018 | 28.6±23.1 | 33.3 | 22.0±18.9 | 16.7 |
| Shortness of breath                                | 0.002 | 24.5±27.7 | 33.3 | 14.8±21.8 | 0 |
| Insomnia                                          | 0.034 | 44.5±30.0 | 33.3 | 36.8±26.2 | 33.3 |
| Lack of appetite                                   | 0.001 | 45.2±35.4 | 33.3 | 31.3±31.0 | 33.3 |
| Constipation                                       | 0.008 | 19.7±26.4 | 0 | 12.0±20.6 | 0 |
| Diarrhea                                          | 0.176 | 13.3±21.7 | 0 | 9.3±16.3 | 0 |
| Financial problems                                | 0.004 | 43.9±34.4 | 33.3 | 32.4±30.8 | 33.3 |
| Side effects of systemic treatment                 | 0.387 | 33.1±21.8 | 28.6 | 30.6±21.0 | 28.6 |
| Ailments related to the arm                        | 0.119 | 29.3±19.7 | 27.8 | 26.2±18.9 | 22.2 |
| Ailments related to the breast                     | 0.545 | 25.2±18.9 | 25 | 23.8±18.9 | 25 |
| Hair loss                                         | 0.005 | 59.2±37.0 | 66.7 | 74.2±33.8 | 100 |

M, arithmetic mean; Me, median; SD, standard deviation; 1, higher value means a better level of functioning and quality of life (at least 0, max. 100); 2, higher value means greater severity of symptoms (at least 0, max. 100); a, b, averages marked with different letters differ statistically significantly at p <0.05.
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Table 4. Women’s Quality of Life Assessment - Categories Related to QLQ-C30 and QLQ-BR 23 and Education

| Functional scales and symptoms QLQ-C30 | QLQ-BR23 | p | Vocational | | Secondary | | Higher |
|------------------------------------|----------|---|------------|---|----------|---|----------|
| N                                  |          |   |            |   |          |   |          |
| Health status and quality of life  | 0.657    |   | 51.9±18.1  | 50 | 53.2±20.8 | 50 | 56.5±19.8 | 58.3 |
| Functional scales<sup>1</sup>       |          |   |            |   |          |   |          |
| Physical functioning               | 0.001    |   | 70.3±18.9  | 73.3| 75.1±18.5 | 80 | 78.9±15.9 | 80 |
| Performing social roles            | 0.423    |   | 71.9±21.1  | 66.7| 74.1±23.2 | 66.7| 75.5±24.3 | 66.7 |
| Emotional functioning              | 0.043    |   | 55.6±27.1  | 66.7| 60.0±23.7 | 66.7| 63.4±23.9 | 66.7 |
| Cognitive functioning              | 0.021    |   | 65.0±26.1  | 66.7| 71.9±25.7 | 83.3| 73.6±24.2 | 83.3 |
| Social functioning                 | 0.516    |   | 71.4±30.0  | 83.3| 69.0±29.7 | 66.7| 69.3±26.4 | 66.7 |
| An image of your own body          | 0.292    |   | 60.6±36.4  | 66.7| 64.7±33.0 | 66.7| 59.2±29.3 | 66.7 |
| Sexual function                    | 0        |   | 15.0±25.5  | 0  | 14.6±21.7 | 0  | 22.8±22.7 | 16.7 |
| Sexual satisfaction                | 0.393    |   | 43.2±31.8  | 33.3| 50.8±26.4 | 33.3| 43.0±28.8 | 33.3 |
| The prospect of the future         | 0.461    |   | 28.4±35.2  | 0  | 33.0±34.5 | 33.3| 31.2±32.0 | 33.3 |
| The scale of the symptoms<sup>2</sup> |          |   |            |   |          |   |          |
| Fatigue                            | 0        |   | 43.1±19.7  | 33.3| 37.4±21.6 | 33.3| 32.3±17.3 | 33.3 |
| Nausea / vomiting                  | 0.11     |   | 38.9±39.6  | 33.3| 34.4±37.3 | 16.7| 27.6±34.3 | 16.6 |
| Pain                               | 0        |   | 30.4±20.6  | 33.3| 24.1±20.6 | 16.7| 18.7±19.3 | 16.7 |
| Shortness of breath                | 0.023*   |   | 22.2±26.3  | 16.7| 19.0±24.7 | 0  | 13.3±21.4 | 0  |
| Insomnia                           | 0.005    |   | 45.8±28.1  | 33.3| 39.8±28.7 | 33.3| 33.0±25.2 | 33.3 |
| Lack of appetite                   | 0        |   | 46.4±33.6  | 33.3| 35.7±34.0 | 33.3| 26.6±29.1 | 33.3 |
| Constipation                       | 0.664    |   | 14.0±20.7  | 0  | 17.5±28.1 | 0  | 12.0±18.5 | 0  |
| Diarrhea                           | 0.81     |   | 10.8±18.3  | 0  | 11.7±19.8 | 0  | 9.6±17.1  | 0  |
| Financial problems                 | 0.057    |   | 39.9±31.8  | 33.3| 38.3±33.0 | 33.3| 30.9±32.1 | 33.3 |
| Side effects of systemic treatment | 0.002    |   | 37.2±22.3  | 33.3| 31.5±21.8 | 28.6| 25.9±18.1 | 23.8 |
| Ailments related to the arm        | 0.029    |   | 30.4±19.5  | 33.3| 31.5±21.8 | 28.6| 24.0±15.4 | 22.2 |
| Ailments related to the breast     | 0.016    |   | 29.4±21.8  | 25  | 21.7±18.5 | 16.7| 22.1±15.2 | 20.8 |
| Hair loss                          | 0.8921   |   | 67.2±36.9  | 66.7| 67.7±36.2 | 66.7| 70.8±34.1 | 83.3 |

M, arithmetic mean; Me, median; SD, standard deviation; *, no statistically significant differences were found in the post hoc analysis; 1, higher value means a better level of functioning and quality of life (at least 0, max. 100); 2, higher value means greater severity of symptoms (at least 0, max. 100); a,b, averages marked with different letters differ statistically significantly at p <0.05.

**Quality of life and education**

Another variable significantly affecting the assessment of the health status and QOL of the surveyed women was education (Table 4). The values of QLQ scales were assessed in three groups of respondents: the first - declaring vocational education, the second - with secondary and third - with higher education. A general analysis of the health status and QOL of women diagnosed with breast cancer indicated a higher score in the case of women with secondary and higher education compared to women with vocational education, but this was not statistically significant. For some functional and symptomatic scale values (QLQ-C30) statistically significant differences were found. Women with higher education higher rated physical functioning (M = 78.9), emotional (M = 63.4) and cognitive (M = 73.6) compared to those with secondary and vocational education. Performed tests of multiple comparisons indicated that in the case of the respondents’ scale with vocational education differed statistically significantly in terms of physical functioning (p = 0.001), emotional (p = 0.043) and cognitive (p = 0.021) from women with higher education. The level of education differentiated QOL in terms of the assessment of the body image, sexual satisfaction, future prospects and hair loss, as a side effect of the applied therapy. In the studied groups, the area of sexual functioning and the prospect of the future were assessed low. Sexual functioning was rated the lowest by women with vocational education (M = 15.0) and average (M = 14.6). There were statistically significant differences between the arithmetic mean values for the scale - sexual functioning in the group of women with higher education and secondary and vocational education (p = 0.000). Sexual satisfaction among women who were sexually active was assessed at an average level (Table 4).

Respondents with the lowest level of education more often indicated the nuisance of symptoms associated with the disease. Statistically significant differences were found between women with vocational education and the group of women with higher education in the arithmetic mean values calculated for the following symptoms: fatigue (p = 0.000), pain (p = 0.000), insomnia (p = 0.005) and lack of appetite (p = 0.000). In the case of other scales,
the existing differences were not confirmed statistically (Table 4).

Side effects of the applied therapy (p = 0.002) and arm ailments (p = 0.029) were most seldom indicated by women with higher education, and the observed differences between arithmetic means for this scale between the group of women with higher education and occupational level achieved statistical significance. Breast ailments mostly concerned women with vocational education (the highest arithmetic mean), and the difference between this group and the group of women with secondary education was confirmed statistically (p = 0.016) (Table 5).

Quality of life and financial situation

In the study population, differences in functional scale and symptom scale were observed depending on the financial situation of the women surveyed (Table 5). Overall health and QOL were rated the respondents who were in a very good and bad financial situation (M = 65.6). In the analysis of emotional and cognitive functioning of women diagnosed with breast cancer, significant differences were found between respondents who were in a very good and bad financial situations.
Table 6. Spearman’s Rank Correlation Coefficients between QLQ-C30 and OLQ-BR23 and the Age of the Subjects

| Functional and symptomatiuc scales | R Spearman | p    |
|-----------------------------------|------------|------|
| QLQ-C30                           |            |      |
| Health status and quality of life | -0.19      | 0.034|
| Physical functioning              | -0.17      | 0.002|
| Performing social roles           | -0.05      | 0.325|
| Emotional functioning             | -0.05      | 0.382|
| Cognitive functioning             | -0.03      | 0.595|
| Social functioning                | 0.11       | 0.04 |
| Fatigue                           | 0.09       | 0.089|
| Nausea / vomiting                 | 0.04       | 0.474|
| Pain                              | 0.12       | 0.033|
| Shortness of breath               | 0.09       | 0.118|
| Insomnia                          | 0.11       | 0.046|
| Lack of appetite                  | 0.17       | 0.002|
| Constipation                      | 0.03       | 0.597|
| Diarrhea                          | 0.01       | 0.778|
| Financial problems                | 0.1        | 0.067|
| QLQ-BR23                          |            |      |
| An image of your own body         | 0.22       | 0    |
| Sexual function                   | -0.37      | 0    |
| Sexual satisfaction               | -0.14      | 0.175|
| The prospect of the future        | 0.03       | 0.549|
| Side effects of systemic treatment| 0.06       | 0.248|
| Ailments related to the arm       | 0.16       | 0.003|
| Ailments related to the breast    | 0.09       | 0.108|
| Hair loss                         | -0.21      | 0.004|

Quality of life and age

A relation was found between the general assessment of health and QOL (R = -0.19, p = 0.034) and the age of the surveyed women. With age, the QOL of the respondents decreased. There were negative and significant correlations in the case of physical functioning (R = -0.17, p = 0.002), sexual (R = -0.37, p = 0.000) and hair loss (R = -0.21, p = 0.004). It draws attention to the clearest, with average strength correlation between age and sexual functioning. The age of respondents positively correlated with social functioning (R = 0.11, p = 0.040), body image (R = 0.22, p = 0.000) and with the occurrence of pain (R = 0.12, p = 0.033), insomnia (R = 0.11, p = 0.046), no appetite (R = 0.17, p = 0.002) and shoulder discomfort (R = 0.16, p = 0.003). Detailed results are provided in Table 6.

Discussion

QOL is a subjective concept and assessed from the patient’s perspective. Oncological treatment is performed on the application of effective methods of therapy, while ensuring high QOL. Understanding the factors determining QOL of women with breast cancer can indicate the directions of activities to ensure adequate comfort of life for sick women. Among the determinants influencing QOL, socio-demographic factors change.

The study evaluated QOL of women diagnosed with breast cancer depending on the following socio-demographic factors: age, marital status, education, and financial situation. The collected data show that with age, the global assessment of health and QOL of patients decreased. This dependence also occurred in the case of physical and sexual functioning. In turn, older women had a better QOL in the field of social functioning and they rated the body image more highly. With age, the occurrence of symptoms associated with the disease, such as pain, insomnia, lack of appetite and ailments of the arm on the operated side increased. The presented research results indicate that for younger women it is particularly important to assess the body image, which seems to be more critical than in the case of older women. The attention is also paid to the different type of support that should be directed to women of all ages. Younger patients, due to low assessment of the sphere of social functioning and low assessment of their own body image, need more support in this area. However, for older women, help is needed to increase their physical and sexual functioning.

The literature presents various results regarding the relationship between QOL and the age of patients. Sio et al., (2014) stated that younger women performed better in the area of physical activity, while older patients showed a good general QOL and higher assessed the body image, which largely coincides with the results presented in this paper. Huang et al., (2017) analyzing the impact of selected socio-demographic factors on QOL among 225 women with breast cancer, stated that QOL...
was not age-dependent. In turn, a study by Wöckel et al., (2017) showed that the QOL of young women waiting for treatment was lower than the older ones, and the treatment applied could still have a negative effect on the quality of life. In the studies of Nancy et al., (2005), the relationship between the applied therapy (chemotherapy, surgical treatment) and QOL of women with breast cancer was examined and the socio-demographic factors were analyzed to influence the existing relationships. One of the conclusions obtained as a result of these studies was the fact that QOL of women depended not only on the type of treatment, but also on the age of the subjects and their level of education, which should be taken into account in the therapeutic process.

Determining marital status as a differentiating variable in own studies, it was shown that people in relationships achieved significantly better results in terms of cognitive functioning and assessment of future prospects and had lower intensity of symptoms, especially vegetative, associated with the disease (nausea and vomiting, pain, shortness of breath, insomnia, lack of appetite, constipation). In addition, women in relationships have had less financial problems than single people. There were no statistically significant differences in the assessment of the body image and sexual function and sexual satisfaction between unmarried women and those in relationships. Similarly, in the studies of Parker et al., (2003), it was shown that the higher QOL had married patients. Also, studies carried out by Croft et al., (2014) showed that married women who were diagnosed with breast cancer after 5 years from the diagnosis had a higher level of optimism compared to unmarried women. In the studies of Cobo-Cuenca et al., (2018) and Acil and Cavdar (2014) it was shown that women in relationships had greater life satisfaction compared to unmarried women. Demonstrating the relationship between being in a relationship with QOL and women with breast cancer indicates the importance of family support in the fight against the disease.

In our own research, education was a factor that, although it did not result in differences in the overall health and QOL scores, influenced the value of some functional and symptomatic scales. Women declaring higher education were significantly better in the physical, emotional, cognitive and sexual areas compared to women with vocational education. At the same time, in women with lower levels of education, symptoms associated with the disease, such as fatigue, pain, insomnia, lack of appetite, breast and shoulder discomfort and side effects of systemic treatment were more pronounced than in women with higher education.

Recently, the importance of educational programs for patients diagnosed with cancer has been particularly emphasized. Shahsavari (2015) found that the implementation of such programs at the stage of breast cancer diagnosis, treatment options and continuation of therapy, as well as acceptance of the disease resulted in improved QOL of patients in the four domains of functioning: physical, psychological, social and emotional.

The results of own research carried out in Father B. Markiewicz Podkarpackie Oncological Centre in Brzozów indicate that another factor that significantly influenced QOL was the financial situation of women. The very good financial situation of the respondents was associated with a higher QOL, both global and expressed in the form of individual functional and symptomatic scales. Women who declared a very good financial situation achieved higher results in terms of physical functioning, fulfilling social roles, emotional and religious functions. As a consequence, also symptoms associated with cancer were more burdensome for women with worse financial situation compared to better-off. Almost all analyzed symptomatic scales showed a stronger intensity of symptoms in the group of women with unsatisfactory financial situation. It is noteworthy that the differences shown in the majority take on a linear character. Similar results were obtained by Huang et al., (2017) and Yan et al., (2016). In their studies, they found higher QOL in breast cancer patients who had higher income. Also in the studies of Kobayashi et al., (2008), it was shown that higher family income was positively correlated with better overall health and QOL of women. In contrast, loss of job negatively correlated with all functional scales of the QLQ-C30. Similar results in their research were obtained by Jendrian et al., (2017), in which they stated that predictors of good QOL among women diagnosed with breast cancer were among others higher monthly family income and higher professional qualifications.

The results presented by Ramadas et al., (2015) indicated that QOL of women diagnosed with breast cancer was higher in those respondents who were married and lived with their family. In these studies, it was found that women who did not have children significantly worse in the psychological area compared to patients with children.

Obtained results of own research and other authors indicate the great importance of socio-demographic factors for QOL of women diagnosed with breast cancer.

Conclusions

1. The age, marital status, education and financial situation of women diagnosed with breast cancer had an impact on their QOL.
2. In order to improve the quality of life of women, special care should be given to the elderly in the area of physical and sexual functioning and discomforts resulting from the disease, while in the group of younger women in the field of social functioning and improving the assessment of their body image.
3. Professional care that strengthens the functioning of women with breast cancer in all areas of life should be addressed especially to women who are not in relationships, with a lower level of education and in a worse financial situation.
4. A great role in caring for patients with breast cancer should be played by social monitoring and providing assistance by appropriate organizations and social services.

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