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Systematic Review

Supportive Care of Breast Cancer Patients in Iran: A Systematic Review

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Abstract

Context: As breast cancer treatment going forward, need for supportive strategies grows. That creates an important call to summarize what has been done regionally.

Objectives: In this study, we systematically reviewed articles that proceeded rehabilitation and supportive care in breast cancer patients in Iran to present a research map of rehabilitation research in the past 10 years in Iran.

Data Sources: All articles published from January 2006 to October 2015 were included. All of the breast cancer studies in Iran were searched in 3 English (Web of Science, PubMed, and Scopus) and 2 Persian databases (SID and IranMedex).

Study Selection: All papers related to rehabilitation in breast cancer were included and categorized into 5 subgroups including qualitative, instrument, lymphedema, interventional, and observational studies. Three reviewers (two surgeons and an epidemiologist) screened the primary search and divided it into subgroups.

Data Extraction: Two reviewers used a checklist to critically appraise the full text of the selected articles. The necessary information of retrieved articles was extracted and recorded in the designed data extraction spreadsheet in Excel software.

Results: A total of 194 articles (102 in English and 92 in Persian) were assessed for eligibility of inclusion in the review, of which, 121 were excluded, and 73 studies were kept. The included studies consisted of 14 on qualitative design, 5 studies in the translation and validation of research instruments, 7 articles in the field of lymphedema, 20 articles about different intervention modalities on breast cancer patients (including education, social status, psychological, exercise, etc.), and 27 observational studies about anxiety, depression, quality of life, sexual function, emotional distress, complementary medicine, lifestyle, etc.

Conclusions: Most of the reviewed studies insisted on a prevalence of physical, psychological, functional, and spiritual problems of breast cancer survivors and their caregivers. Designing a mega project to offer a palliative and rehabilitation service package according to the needs of Iranian patients may become a priority in their health care system.

Keywords: Breast Cancer, Supportive Care, Iran, Systematic Review

1. Context

Breast cancer (BC) treatment exposes patients and their caregivers to a series of challenges that make coping with the situation a struggle. At the same time, the burden of the disease impacts society at large (1). Younger women with BC are more likely to experience psychosocial and menopause-related concerns, weight gain, and physical inactivity (2). The broad spectrum of psychosocial issues experienced by women with BC can be categorized into 4 groups, including (1) dealing with cancer, (2) the importance of caring, (3) the aftermath of cancer, and (4) fertility and infertility (3).

Owing to improvements in cancer detection and treatment, BC is increasingly becoming a chronic illness with longer survival years. Therefore, for BC rehabilitation to be comprehensive and effective, patients and their caregivers should be taught how to incorporate health-promoting habits in their lives by redesigning their lifestyles (4). There is strong evidence that BC patients and survivors can benefit enormously from exercise interventions in terms of quality of life (QoL), cardiorespiratory fitness, physical functioning, and fatigue (5, 6). For instance, yoga practice has been shown to enhance health and alleviates some treatment-related side effects for patients recovering from BC (7). In total, acknowledging practitioners and clinicians may help to improve a BC prognosis through recommending exercise, anticipating the physiological effects on cancer (8).

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2. Objectives

According to the published research, BC patients need various services during and after treatment in order to maintain a better QoL. It seems that most of these aspects are unclear in Iranian women. In this study, we systematically reviewed articles that begin with rehabilitation in BC patients in Iran. Presenting the research map of rehabilitation activity in the past 10 years in Iran may provide a useful perspective for future research to establish the efficient and beneficial services for developing the QoL of these patients.

3. Methods

This study is part of a big project to study the different aspects of BC in Iran. All of the published articles about BC in Iran, within a defined time, were included in the study. Then they were divided into 5 subgroups, and a specific process of systematic review was followed in each subgroup. Details of the methodology are as follow:

4. Data Sources

All articles published from January 2006 to October 2015 were included. To achieve the most comprehensive medical electronic databases, keywords were extracted from the medical subject headings (MeSH) of PubMed. Studies were searched in English and Persian databases. English online international electronic databases consisted of Web of Science, PubMed, and Scopus. The English search formula was "breast cancer" OR "breast carcinoma" OR "breast tumor" OR "breast neoplasm" AND "Iran". In total, 1986 English abstracts were included.

Persian databases consisted of SID and IranMedex, the most comprehensive national electronic databases, with the most coverage of Iranian public health and medical journals. It was not possible using a combined formula in a Persian search; so, keywords were searched separately (Saratan-e-Sineh, Saratan-e-Pestan, breast tumor, breast cancer, breast carcinoma, breast neoplasm and Iran). Then, they were combined with each other. In total, 1345 Persian abstracts were retrieved.

5. Study Selection

Three reviewers (two surgeons and one epidemiologist) screened the primary search and divided it into subgroups; 50 articles were randomly selected for validation of the reviewers' agreement assessment. They were instructed to decrease their disagreements in article classification. The trained experts divided the articles into 5 subgroups (epidemiology and risk factors, genetic, prevention, diagnosis and treatment, and rehabilitation), although some of them were allocated to more than one

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group. Being classified according to titles and abstracts, the duplicated and unrelated articles were excluded. Totally, 763 English and 572 Persian abstracts were divided into 5 subgroups to find and assess the full-texts. Irrelevant studies were evaluated again by the team supervisor. So, the final number of eligible articles for systematic review in 5 subgroups was 1646 abstracts. The full text of abstracts was found, and in some occasions, a letter was sent to the author to get the necessary information. Two reviewers used a checklist to critically appraise the full text of the selected articles. If there was any disagreement, the reviewers would discuss the eligibility of the articles and make a decision about it.

6. Data Extraction

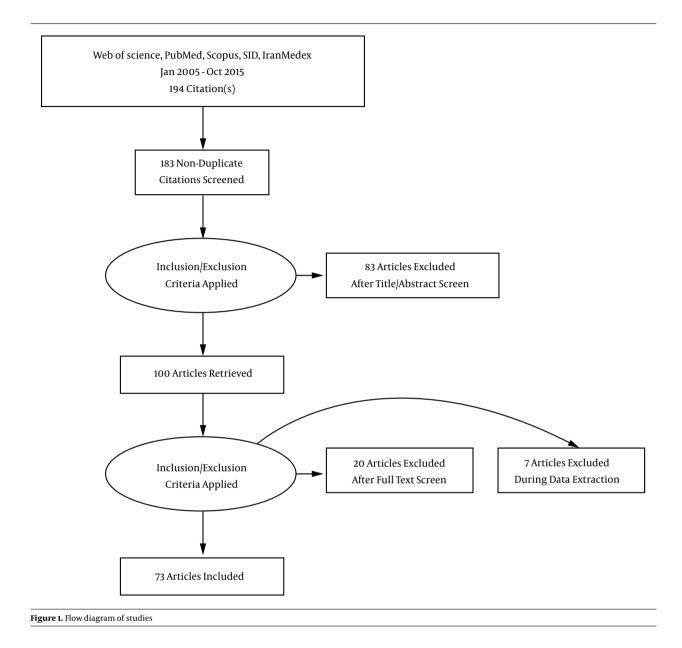
The necessary information of retrieved articles was extracted and recorded in the designed data extraction spreadsheet in Excel software. Datasheet consisted of general information (title, the place of study, study year, journal name, and publication year), methodological information (study design, sample size, data source, studied population, measurement tools, sampling method), and results of the study (main outcome, effect size, and measurement tools). All of the articles were extracted by two people. A representative of each subgroup organized the two extracted forms into one sheet.

In the third step, the extracted data should be analyzed. Because of the variety in the field of studies, the results were categorized according to their study design, except for specific fields such as lymphedema and validation of instruments. Thus, the rehabilitation subgroup reviewed articles were categorized into 5 groups, including qualitative, instrument, lymphedema, interventional, and observational studies. Interventional studies were divided into educational, social, psychological, exercise, and other interventions. Observational studies consisted of anxiety and depression, QoL, sexual status, emotional distress, complementary medicine, and social studies.

7. Results

In total, 194 articles, consisting of 102 English and 92 Persian, were considered eligible for study in this group, of which 11 studies were duplicated and removed. From 183 studies, 83 studies were irrelevant and excluded. In the next step, more than 27 studies were excluded based on full text. The reasons for exclusion were as follows:

Nine articles were just abstracts, 7 were not specifically related to BC, 7 were review studies, 2 articles were not related to the Iranian population, and 2 studies were on animals (Figure 1). The reviewed articles were categorized into 5 groups according to their main themes. The results of the papers in each group were as follow:



7.1. Qualitative

As qualitative studies must be analyzed differently in the systematic review, we are just reporting an overview of them. In total, 14 studies were conducted by qualitative design. All of these studies (9-20), except two (21, 22), were done on BC patients. A study was on BC caregivers. Different purposes were assumed for these studies. The assumed purposes were patients' needs (14, 18), caregivers experiences (22), their religious experience (10, 16), their experience with BC and symptoms (12, 13, 17), coping with BC and treatment side effects (9, 19, 20), their perceptions of life (15) and role of social and cultural factors in a patient's life (11), and the patient's husband's perception toward their sexual life (21). Different extracted articles in this theme have been presented in Table 1.

7.2. Instrument

Articles in which different instruments had been translated and validated were categorized in another theme. In sum, we found 5 studies. All of them except one had studied BC patients (23-26). The other one had been conducted on the BC patient's caregivers (27). Most of the studies had validated an instrument in the field of QoL. FACT-B, mini MAC scale, need questionnaire, QoL index-cancer scales, and QLQ-C30 had been validated in these studies (Table 2).

Study	Sample Size	Age, Mean \pm SD	Age Range	Outcome of Interest	Main Extracted Themes
Taleghani et al. (19)	19	N/A	31-25	Coping with BC	Religious approach, thinking about the disease, accepting the disease, social and cultural factors, support sources
Taleghani (20)	45	N/A	31 - 56	Patient's experiences in adjusting to the disease	Perceived threat to life, living with the disease with tolerance, religious aspects, and barriers to efforts leading to health, will to recover, supportive dimensions, increase in endurance, and inhibitors and facilitators of tolerance
Harandy et al. (10)	39	N/A	30 - 87	Patient's experience with religious context	Religiosity does not prevent Iranian wome from seeking medical care
Harandy et al. (11)	39	N/A	30 - 87	Correlated factors of health-related quality of life	Fatigue, pain, and lymphedema are the most common complaints
Sajadian et al. (17)	51	48.4 ± 10.5	25 - 72	Patient's experience with BC	Importance of spirituality and family support, especially husband and children, during the diagnosis and treatment; chemotherapy as the worst experience
Nasrabadi et al. (15)	23	N/A	N/A	Patient's perception of life	Cancer as a kind of divine test, a very bitter and debilitating experience, chemotherap as the most difficult experience of cancer, a continuous struggle
Joulaee et al. (12)	13	N/A	34 - 67	Patient's experience with BC	Negative aspects: losing something important, uncertainty, living with fear, emotional confusion, needing support, positive aspects: new aspects of life
Moradian et al. (14)	30	$42 \pm N/A$	19 - 59	Patients' needs	Treatment costs, psychological distress
Nasiri et al. (21)	18	51.5 ± 10.47	33 - 70	Patient's husband perceptions	Altered sexual relations; sexual abstinence avoidance, or restraint; attempt to normalize relationship
Fouladi et al. (9)	20	N/A	33 - 71	Coping with mastectomy	Loss and death contest, reconstruction of evaluation system, reactions and troubles after a loss, health, reorganization and compatibility with changes
Khakbazan et al. (13)	27	$42.8 \pm N/A$	26 - 71	Patient's experience with symptoms	Symptom recognition, confronting the fea of cancer, labeling symptoms, interactive understanding
Sadati et al. (16)	8	N/A	N/A	Patient's experience with religious context	Fatalism, hope, and empowerment
Taleghani et al. (18)	19	N/A	34 - 60	needs of patients	Information, beliefs, and skills
Hashemi-Ghasemabadi et al. (22)	23	$37.5 \pm N/A$	20 - 69	Caregivers	Being involved in a new situation, abandoned in the role, infinite absence, perceived inefficiency

7.3. Lymphedema

Seven studies (28-34) were conducted on BC patients, who suffered from lymphedema (Table 3). Although some of these studies were looking at lymphedema treatment methods, we tried to discuss the rehabilitation aspects because of their importance in BC QoL and rehabilitation of which 3 were interventional (29-31), and the remaining 4 were observational (28, 32, 33). Two kinds of intervention were used in this field; 2 of them were a combination of complex decongestive therapy (CDT) vs. CDT + intermittent pneumatic compression (29, 31), and they both reported the significant effects of this method on BC-related lymphedema. The third interventional study evaluated the effect of a home-based rehabilitation education on edema reduction (30). It also had a significant effect on lymphedema reduction.

7.4. Interventional Studies

In total, we found 20 articles, which looked at different intervention modalities on BC patients. These studies were conducted on distinct fields including education, social status, psychological, exercise, and other aspects. Those fields have been reported in Table 4.

There were 3 papers in the education field (37, 40, 53). Two, out of the 3 studies, were controlled trials (40, 53), and the 1 remaining was a trial without a control group (37). All 3 studies had a sample size of more than 50. One study found that a life skill educational program was associated with improvements in somatization disorders, depressive and anxiety symptoms, disorders of social functioning, and sleep disorders (53). Two other studies had looked at the effects of self-care education on QoL and mental health (37, 40). The results showed that self-care educa-

Study	Sample Size	Age, Mean \pm SD	Age Range	Validated Instrument	Main Result
Safaee et al. (25)	132	48.61 ± 11.22	NA	QLQ-C30	Cronbach's alpha: fitted in all subscales except fatigue (0.65), pain (0.69) and nausea and vomiting (0.66). Convergent validity correlation: > 0.40 for all subscales. Item discriminant validity: significant difference in all subscales except for item 4 of the physical functioning
Khanjari et al. (26)	166	40.7±13.1	18 - 75	Caregiver quality of life index-cancer scale	Cronbach's alpha: 0.72 - 0.90
Ghaffari et al. (24)	160	44.6 ± 12.63	NA	Self-assessed support needs questionnaire for BC cases	Cronbach's alpha for all items: 0.83, stability of test 0.78, Cronbach's alpha of the first factor: 0.90
Patoo et al. (23)	300	$43.34\pm\mathrm{NA}$	23 - 72	Functional assessment of cancer therapy-breast (FACT-B)	Cronbach's alpha (total): 0.92, Cronbach's alpha (subscales): 0.63 to 0.93, significant concurrent and discriminant validity was fitted.
Patoo et al. (27)	320	NA	NA	Mini-mental adjustment to cancer scale (mini-MAC scale)	Cronbach's alpha: 0.84

Study	Study Design	Sample Size	Age, Mean \pm SD	Intervention	Outcome	Main Extracted Themes
Haddad et al. (28)	Cross sectional	355	NA	-	Lymphedema	Mean prevalence of lymphedema: 17%
Haghighat et al. (29)	Trial	112, INT: 56, CON: 56	INT: 53.4 \pm 11.4, CON: 52.7 \pm 10.8	CDT vs. CDT + IPC	Edema volume	CDT was more effective than CDT + IPC, acute phase: P = 0.036, maintenance phase P = 0.167
Khosh-Nazar (30)	Trial (before- after)	16	53 ± 9.5	Education (SLD + self-care + exercise)	Edema volume	Decrease of lymphedema P < 0.001
Moattari et al. (31)	Trial (before- after)	21	50.38 ± 9.92	CDT + IPC	Edema volume	Decrease in lymphedema, P < 0.001
Haghighat et al. (32)	Cross sectional	137	53.5 ± 10	-	Edema volume in CDT and predictive factors	Initial lymphedema volume and the duration of lymphedema are predictors of outcome (P = 0.003 and 0.002, respectively
Hemmati et al. (33)	Cross sectional	170	NA	-	Lymphedema and related factors	Correlation of lymphedema and BMI (P = 0.02) and involved lymph nodes (P = 0.0001)
Haghighat et al. (34)	Case- control	410, case: 123, CON: 287	Total: 49 ± 10.9, case: 50.6 ± 11.4, CON: 48.4 ± 10.6	-	Lymphedema and correlated risk factors	High BMI (OR: 1.09; 95% CI, 1.05 - 1.15), No. o involved lymph nodes (OR: 1.15; 95% CI, 1.08 - 1.21), and a longer period after surgery (OF 1.01; 95% CI, 1.01 - 1.02) are associated with an increased risk of lymphedema

Abbreviations: CDT, complex decongestive therapy; IPC, intermittent pneumatic compression; SLD, self lymphatic drainage.

tional intervention was associated with the significant improvement of mental health (37); however, the other study reported non-significant changes in QoL (40).

Three studies looked at the effects of social intervention on the QoL of BC patients (39, 41, 42). In all 3 studies, the intervention was participation in a peer support group. They studied the intervention effect on QoL. In total, social intervention had a significant effect on vitality, mental health, and the total score of QoL.

We found 8 studies that had investigated the effects of psychological interventions on BC patients. All 8 studies were done on BC patients. Five out of 8 studies had a sample size between 20 and 30 (43, 45, 50, 51, 54). Two studies had sample sizes of more than 50 (38, 44). A variety of psychological interventions were used; a complete list of interventions is presented in Table 4. Three different kinds of outcomes were assessed including QoL (43, 44, 52, 54), body image (38, 45, 51), and depression (50). Interestingly, all psychological interventions led to improvement in the selected outcomes.

Two papers had used exercise as an intervention (36, 46). One of them had measured physiological indices (36) and the other one measured QoL (46). Exercise had a significant effect on physiological outcomes (36) related to cardiopulmonary fitness. The other study reported no significant effect on QoL (46).

One study looked at exercise plus ginger supplementation on cardiopulmonary and IL-10. Results of the study showed that this intervention had a significant effect on the selected outcomes (47). One study looked at the effect of ginger supplement on nausea and the result showed a significant effect of ginger on nausea (49). Also, there was a study about the effect of massage therapy on nausea and its result showed no significant effect (48). Finally, one study had looked at the effect of a rehabilitation intervention on QoL and result showed significant improvement in all subscales of QoL (35).

7.5. Observational Studies

In total, 27 studies had been conducted by observational design. All of these studies except 4 were performed on BC patients. Three of the mentioned studies were on BC caregivers and the 4th was on cancer patients. Different purposes were assumed for these studies. The main purposes of these studies were anxiety (55, 56), depression (55-57), QoL (58-64), sexual status (65-67), emotional distress (68), caregiver's QoL (69-71), complementary use status (72, 73), QoL status relationship with lifestyle (74), QoL relationship with nutritional status (75), QoL status along with sense of coherence (76), the relationship between quality of sleep and with spiritual well-being (77), the relationship between hope, body esteem, and mental health (78), QoL status relationship with spiritual well-being (79), and social (80, 81). Also, different themes were extracted that are presented in Table 5. In all of these groups of articles, the frequency of outcome, its range of variation during a time interval, and some of their correlated factors in a specified sample size had been reported.

8. Discussion

The aim of this study was presenting all of the articles that had studied Iranian BC patients and looked at outcomes related to HRQoL. In total, we found 73 studies. As studies were so varied, we divided them to 5 parts, including qualitative, instrument, lymphedema, interventional, and observational studies. Interventional and observational studies contained different structures.

8.1. Qualitative Studies

Though we present the results of the qualitative studies, the conclusions of these studies were not presented because the method for reviewing these kinds of studies is different than in quantitative studies (72). Thus, there is just a report of them. Qualitative studies were conducted on BC patients and also on their caregivers. Such different themes were extracted that a discussion on these themes is beyond the scope of this article. In these studies, different goals were followed, including general experience and perspective, religious experience, ways of coping, their needs, and the role of other factors in that experience (9-22). Only two of 14 articles were about husbands (21) and caregivers (22). Unfortunately, none of the qualitative studies have led to a quantitative guideline for future studies. Developing some native quantitative instruments for the evaluation of patients' needs or experiences is suggested to provide applicable guidelines for the promotion of rehabilitation services.

8.2. Instruments Validation

This review introduced 5 Iranian validated questionnaires for evaluating QoL (QLQ-C30), the caregiver QoL index-cancer scale, self-assessed support needs questionnaire for breast cancer cases, mini-mental adjustment to cancer scale (Mini-MAC scale), and functional assessment of cancer therapy-breast (FACT-B) (23-27). It is possible that some instruments have been validated before this time interval or might have been published in some references that have not been included in this study. The development of more valid and reliable instruments in different aspects of cancer rehabilitation can present a more accurate measurement of problems and provide more effective interventions for solving them. Thus, it seems that introducing native and national valid instruments should be considered in future researches.

8.3. Lymphedema

Breast cancer patients often experience lymphedema, which could compromise their QoL and interfere with their daily activities (65, 73). A mean prevalence of 17.5% was reported for Iranian BC patients according to one study (28). The prevalence of BC-related lymphedema had been reported at 6% to 70%, depending on the kind of surgery (64). Despite a lot of studies about lymphedema in the world, it still has many unknown aspects (82). In Iran, lymphedema treatment started 13 years ago. During this time, only a few clinics have been involved in treatment. So, very limited research is accessible. Among 7 studies presented in this systematic review, 3 clinical trials and 4 observational studies have been published. Risk factors of lymphedema have been studied in 2013. It showed that body mass index (BMI) and the numeration of involved lymph nodes were significantly related to BC-related lymphedema incidence (33). The standard method for controlling lymphedema is CDT or complete decongestive therapy (82). CDT results have been evaluated in limited studies. Though outcomes and methods of studies reviewed were completely different, 2 studies indicated that a combined decongestive therapy and pneumatic compression pump reduce the BC-related lymphedema (29, 31). In a study, it was shown that initial lymphedema volume and duration of lymphedema were important predictive factors for edema volume reduction following decongestive therapy (32). Also, a home-based rehabilitation that contains drainage, exercise, and behavioral practice education was reported significantly effective in reducing BC-related mild lymphedema (30). Presented articles insisted on the valuable effects of CDT in controlling lymphedema. Definitely, more studies are warranted to study the epidemiology, risk factors, prognostic factors, and effective modalities for diagnosis and treatment of lymphedema in the Iranian population.

8.4. Interventional Studies

Reviewed studies revealed that educational programs could increase patients' QoL and mental health (37, 40, 53). The results of reviewed studies were similar to other studies, which indicated self-care educational programs have an important role in promoting QoL and decreasing mental distress after a BC diagnosis (56). A self-care concept or life skill education consisted of different parts. In the reviewed studies, the details of interventions were somehow less than expected (63), so more studies with exact definitions of interventions are warranted. Finally, all 3 studies used traditional educational programs. As technology is integrating into our lives very quickly, it is suggested to conduct a technology-based educational program for BC patients, who often have difficulty to attend in individual classes. We found 3 studies in social part. In the reviewed article, all social interventions were about the effects of participation in a peer support group (39, 41, 42). One study indicated that this kind of intervention has a significant effect on all aspects of QoL (41); however, other studies reported its significant effect on vitality and mental health (39). Regarding the positive effect of this kind of interventions in improving QoL, designed clinical trials with various social interventions and distinct outcomes are necessary to give better insight into the effects of that on BC.

Studies categorized in psychological groups had used different interventions. Two studies conducted mindfulness-based interventions, and the results showed that these kinds of psychological modalities promote QoL and decrease fatigue (52, 54). Also, innovative relaxation therapies, like the Benson relaxation method, were used and claimed that these methods improved the QoL, although their effect size had not been reported clearly (43). Spiritual therapy was among the other psychological interventions for improving the QoL (45). In addition, one researcher studied the effect of Ellis's rational emotive behavior therapy (REBT) intervention on body image outcomes. The results showed that the intervention changed body image favorably (38). Also, in another study, existential group therapy was compared with a reality group therapy, and the results revealed that existential therapy was more effective than other group therapy methods in improving body image (45). Generally, as there were huge differences in intervention types, we could not have a consensus; however, studies show that psychological interventions have an important role in enhancing QoL and body image after a BC diagnosis. Comparing their effects in controlled clinical trials can individualize the treatments and present the most effective methods in each patient.

Two studies looked at the effect of exercise intervention on BC outcomes. In one study, it was revealed that exercise could improve cardiopulmonary measures (36). Another study assessed the effect of exercise on QoL (46). The results of this study reported no significant effect of exercise on QoL. In another study, the effect of a combined exercise and ginger supplement on IL-10 and cardiopulmonary indices was compared (47). The results of the current study reveal that this combined exercise was associated with improvement in cardiopulmonary and immunity indices. According to McNeely, larger trials with a greater focus on study quality, adverse effects, and long-term benefits of exercise are needed to introduce more beneficial rehabilitation methods in BC patients (5). Some of conflicting results show that we need more studies to define proper exercise programs for BC patients and survivors in different steps of treatment.

Decreasing chemotherapy inducing nausea has been evaluated in different researches. Ginger supplements (49) and massage therapy (48) have been introduced as effective interventions for decreasing nausea in 2 studies. Only one article had studied the application of a rehabilitation program on QoL and it had resulted in a significant effect on outcome of interest (35).

8.5. Observational Studies

8.5.1. Anxiety and Depression

According to our results, 3 studies were focused on anxiety and depression as main outcomes. Vahdaninia et al. (55) evaluated anxiety and depression, using the hospital anxiety and depression scale before treatment, 3 months after initial treatment, and 1 year after the completion of treatment. According to the findings, anxiety improved over time. Hadi et al. (56) evaluated anxiety and depression, using the symptom checklist-90 revised (SCL-90R) in BC cases and control (healthy) groups. The mean score of anger subscales was significantly higher in the control group compared to cases. Younger patients had higher anger mean scores. Education and tumor size correlated significantly with anxiety. Mashhadi et al. (57) evaluated depression, using the Beck Depression Inventory (BDI) in cancer patients. Findings showed a significantly higher prevalence of depression in patients with BC. According to Vahdaninia et al. (55), depression improved over time. In Hadi et al.'s study (56), depression was not significantly different between the 2 age groups, and higher depression mean scores were found among younger patients. These 3 studies used different instruments with different methods, but the common limitation was a disability to differentiate the anxiety and depression caused by cancer or other factors. We need more ecological studies to determine the anxiety and depression status in cancer patients and general population.

8.5.2. Quality of Life

The largest group of studies evaluated QoL. QLQ-C30 is the most current questionnaire to measure QoL in Iranian studies. These studies focused on patients and their caregiver's QoL and it is relationship with various variables. Tirgari et al. (58) pointed out that a mastectomy in BC participants induced a low mood state and QoL. Safaee et al. (59) mentioned that all symptoms' scores had a reverse association with QoL except loss of appetite and diarrhea. Pourhoseingholi et al. (64) introduced tumor grade, employment, financial status, menopausal status, and dyspnea as the major predictors of patients' QoL. Didehdar Ardebil et al. (60) studied HRQoL, using the Persian version of FACT-B. Frequency of depression symptoms was reported in 50% of the subjects. Significant correlation was observed between depression and overall HRQoL and between different modalities of treatments. Qol of patients treated by chemotherapy was significantly better than radiotherapy. Also, Hatam et al. (61) and Kiadaliri and Bastani (63) both reported a decrease in HRQoL due to chemotherapy with TAC more than a FAC regime. Musarezaie et al. (62) mentioned that QoL is inversely correlated with the no. of chemo sessions and educational status. Mohammadi et al. (74) reported that healthy eating practices were associated with higher scores on social, role, cognitive, and emotional scales, global QoL, and reduced symptoms of financial difficulties. Also, physical activity was significantly correlated with emotional and cognitive scales. These researchers in their other papers (75) reported significant correlations between nutritional status parameters and quality of life subscales: physical, emotional, cognitive, global QoL, symptoms scales fatigue, nausea/vomiting, pain, dyspnea, insomnia, appetite loss, constipation and diarrhea. Rohani et al. (76) reported that women with BC tended to score worse on physical and role scales, along with fatigue and financial difficulties, during the first 6 months when compared with healthy women. The sense of coherence (SOC) and baseline scores on several dimensions of HRQoL were the most important predictors of HRQoL changes. Jafari et al. (79) reported a significant, positive correlation between global QoL and spiritual well-being. Exercise can be an effective strategy to improve QoL in women with BC. It seems that most of the studies observe the effect of BC and its treatment modalities on QoL fluctuations. None of them have suggested a protocol for improving the QoL during the diagnosis and treatment processes. It is not clear if measuring the QoL index is the main problem or not? Is the measurement instrument appropriate in our population? It seems that special demographic characteristics, beliefs, and behaviors of our patients should be considered in new research instruments in order to suggest more applicable and useful interventions.

According to previous studies, exercise reduces fasting insulin levels in BC survivors. This may be due to exerciseinduced reductions in body weight. Practitioners and clinicians may better help BC prognosis be improved through exercise, anticipating physiological effects on cancer (8). Future research is necessary to determine optimal exercise modes and parameters (6). According to the mentioned articles' results, patient's QoL are affected by several factors. Therefore, we need more proportional attention to patients according their chemotherapy plan. Psychological consultation, exercise programs, nutritional advice, and educational programs may be helpful to improve patients' QoL.

Caregivers' QoL showed a negative correlation with depression (69). Family caregivers of BC patients are reported to be affected by psychological impacts for up to 6 months after the diagnosis (70). Caregiver's QoL improves over time (71). Family members as main caregivers should be empowered in order to prepare strong support for cancer patients in all steps. It seems that more studies and educational programs should be designed in future research.

Khoramirad et al. (77) showed the relationship between quality of sleep with spiritual well-being and religious activities; even though it was not significant. Heidari et al. (78) reported a significant relationship between body esteem with hope and mental health. There was a positive correlation between hope and mental health too.

8.5.3. Sexual Status

Garrusi and Faezee (65) studied BC patients' sexual status in 2008. She showed that desire was diminished in 70.6% of the women. The number of instances of coitus decreased in approximately 15% of the participants. About 30% of women reported lubrication and arousal problems, and 72% reported decreased orgasms. Sexual satisfaction did not have a significant correlation with orgasm. Harirchi et al. (66) reported pre-treatment and post-treatment sexual dysfunction rates to be 52% and 84%, respectively, indicating a significant deterioration in sexual function among BC patients. A younger age, reception of endocrine therapy, and poor sexual function at pretreatment were the most significant contributing factors to post-treatment sexual disorders. Safarinejad et al. (67) mentioned the significant prevalence of lubrication problems, satisfaction disorder, desire disorder, and arousal disorder in BC patients compared with healthy controls. Patients receiving hormone therapy were more likely to experience sexual dysfunctions; therefore, this group of patients needed more attention and sexual therapy during and after treatment. It seems that there has been limited research in the study of frequent sexual dysfunctions in BC patients, but no intervention has been tried or suggested to solve this common problem.

8.5.4. Emotional Distress

Saeedi-Saedi et al. (68) reported that 39% of the BC patients had severe emotional distress, which was significantly associated with lower functional status. Taking care of children, anxiety, fear, difficulty taking bath and wearing clothes, family issues, fever, and nasal dryness were the most common contributors to emotional distress. Designing some longitudinal research or clinical trials to evaluate important interventions to manage these unpleasant symptoms is necessary. Yoga practice has been shown to improve health and helps alleviate some treatment-related side effects in BC survivors (7).

8.5.5. Complementary Medicine

Montazeri et al. reported a significant association between depression and the use of complementary medicine (72). The use of complementary medicine among Iranian patients is not common; however, it is correlated with fear, anxiety, and mental distress (73). Leggett et al. reported evidence of different qualities to support that Guarana and Ganoderma lucidum may improve fatigue, while glutamine may be effective in improving oral mucositis symptoms. Overall, current evidence does not provide definitive recommendations regarding the effectiveness of complementary or alternative medicine in women with BC (83). Considering that there is a lack of effective chemotherapy-induced peripheral neuropathy interventions, natural products and complementary therapies merit further investigation (84). We need more trials to approve the effectiveness of complementary medicine in the Iranian population.

8.5.6. Social

Azarkish et al. (80) mentioned that older patients and those with longer work experience were less likely to return to work, while women with no pain, surgery scar, or lymphedema after the BC treatment were more likely to return to work. Moghaddam Tabrizi et al. reported a significant relationship between general health and a healthpromoting lifestyle (81). "Return to life" is an important goal in BC patients, which has not been studied in Iranian research. Fortunately, at present in Iran, most of the diagnosis and treatment modalities are accessible. It can lead to early diagnosis and higher survival in these patients; so, the frequency of survivors is increasing. Designing some projects to define a useful protocol to provide a high QoL for these survivors and a return to familial and social roles may be an important health priority.

To our knowledge, it is the first study that attempted to look at all of the different aspects of BC in Iran. The result of that study could help to draw a better framework to promote QoL of BC patients. Also, this study has some limitations. As our goal was gathering all of the studies done in Iran, some Iranian databases were searched. Despite their great contents, their search engines were somehow weak. Finally, numerous studies were selected and, then, divided into 4 parts. So, we have to discuss all QoL and rehabilitation-related studies in one article, although distinct designs were used by the included studies.

9. Conclusions

Several main problems have been demonstrated in Iranian studies about BC patients' rehabilitation. Most of them consist of the prevalence of physical, psychological, functional, and spiritual problems of BC survivors and their caregivers. More interventional trials in the fields of social aspects, emotional distress, complementary medicine, psychological and family consultations, exercise programs, nutritional plans and educational programs are necessary. Designing a mega project to offer a palliative rehabilitation service package according to the need of Iranian patients may be a priority in their health care system.

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Footnotes

Authors' Contribution: Nasim Khosravi, Najmeh Nazeri, Shahpar Haghighat, Alireza Atashi, Vahid Farajivafa, Asiie Olfatbakhsh, and Maryam Koosha participated in the literature search and data extraction. Nasim Khosravi, Najmeh Nazeri, Vahid Farajivafa, and Shahpar Haghighat analyzed the data and prepared the manuscript. All authors read and approved the final manuscript.

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Table 4. Experimental Studies Included in Rehabilitation Systematic Review	es Included in Reh	abilitation Syste	ematic Review					
Study	Field	Design	Population	Sample Size	Age, Mean \pm SD	Intervention	Purpose/Area	Main Extracted Themes
Poorkiani et al. (35)	Rehabilitation	RCT	BC survivors	66, EG: 28, CG: 30	$40.79 \pm NA/36.7 \pm N/A$	physiotherapy, education, and counseling (2-months)	QoL/QLQ-C30 and BR23	Improvement in all the scales of QoL. Potential benefit for physical, psychological and overall QoL
Rahnama et al. (36)	Exercise	Clinical trial	Post- menopausal women with BC	342	Υ Υ Υ	60 minute resistance training twice weekly (15 weeks)	VO2 max/resting heart rate/heart rate monitor belt/blood pres- sure/anth ropometric variables	Positive effects on VO ₂ max, RHR, body weight, BMI, and WHR ($P <$ 0.05) especially in postmenopausal patients. No significant effect on blood pressure
Bakhtiary et al. (37)	Education	Trial without control	BC patients	60	NA	education on self-care against chemotherapy side effects (3 weeks)	Mental health/GHQ-28	self-care significantly improved mental health ($P < 0.001$), except for 61 - 70 y age group ($P = 0.147$)
Fadaei et al. (38)	Psychological	Quasi- experimental	2 - 6 months after mas- tectomy	22	43.46 土 7.6	Consultation based on REBT method for 6 sessions (3 weeks)	Body image/Ellis rational emotive behavior therapy (REBT)	The body image score decreased in the intervention compared to control group ($P < 0.001$).
Malekpour Tehrani et al. (39)	Social	Trial	BC patients	68	44.6 土 7.5	participate in peer groups (12 weeks)	Quality of life/SF36 questionnaire	Increase in vitality score (P < 0.001) and mental health score (P < 0.001) in experimental group
Ghavam-Nasiri et al. (40)	Education	Trial	BC patients	102	46.6 ± 3.9/48.7 ± 1.23	3 sessions of self-care education	QoL/QLQ-C30	Group teaching could maintain better quality of life than individual teaching over time (P > 0.05)
Sharif et al. (41)	Social	Trial without control	BC patients	66	NA	weekly peer-lead educational programs	QoL/EORTC QLQ-30 and QLQ-BR23	significant improvements in all aspects of QoL in the intervention group (P < 0.001)
Taleghani et al. (42)	Social	Clinical trial	BC surgery for the first time, Isfahand	150	NA	Peer support group	Qol/standard instruments of Mational Medical Center and Beckman Research Institute questionnaire	The mean score of OoL were different during the second stage (P = 0.003) and 2 stages in Tehran. significant difference between 2 groups in social aspect during both stages in Tehran, spiritual aspect in Isfahan and first stage in Tehran
Salehi et al. (43)	Pscychological	Trial	Non- metastatic BC patients	25	NA	Benson relaxation technique: daily, 15 - 20 min/d at home (3 weeks)	QoL/QLQ-C-30 and QLQ-BR23	Significant improvement in QoL (P value not available)
Jafari et al. (44)	Psychological	Trial	BC patients undergo- ing radiation therapy	65	47.9 土10.6/48.1 土10.2	Spiritual therapy sessions (6 weeks)	QoL/QLQ-C30 BR-23	Increased QoL ($P < 0.001$) and all functional scales of QIQ-C30 after intervention ($P < 0.05$)
Parizadeh et al. (45)	Psychological	Semi- experimental trial	Undergone mastec- tomy	24	47.38 ± 6.3	(1) existential group therapy, (2) reality group therapy, (3) no intervention	Body- image/multidimensional body-self relation questionnaire (MBSRQ)	Significant effects were observed for appearance orientation in both intervention in groups (P = 0.039). Mean score for the existential group therapy was greater than that for the reality group therapy (P = 0.004)

Taleghani et al. (46)	Exercise	Trial	BC patient who had completed the treatment	80	AN	Exercise training/3 sessions a week, 60 minutes (8 weeks)	Quality of life/the National Medical Center and Beckman Research Institute instrument	Total mean score of the quality of life showed no significant difference before and after intervention (P = 0.29)
Fathi et al. (47)	Exercise + supple- ment	Before after	Obese BC patients	40	46.4 ± 5.5	 750 mg ginger capsules four times a day, (2) water exercises, (3) both of them, (4) none 	Cardio-pulmonary indexes and IL-10	Exercise training plus ginger upplement were associated with a decrease in IL-10, BMI, body fat percentage and increase in cardiopulmonary indexes
Bosak et al. (48)	Massage therapy	RCT	BC patients under chemo	34	νN	Massage therapy (3 sessions on consecutive days, 30 min before chemotherapy)	Nausea/visual analogue scale	Nonsignificant (P = 0.51) decrease in the severity of nausea
Ebrahimi et al. (49)	Supplement	Irial	BC patients under chemo	80	41.8 ± 8.4	250 mg ginger capsules four times a day	Nausea/visual analogue scale and some questions about severity and frequency	The severity and frequency of delayed mausea were lower in patients receiving ginger ($P < 0.01$). No complication in taking ginger capaules compared to placebo ($P = 0.50$)
Haghighi et al. (50)	Psychological	Non- randomized trial	BC patients	22	45.5 土 9/46.3 土 6.5	Logotherapy: 10 2-h sessions, a session per week.	Depression/Beck's depression inventory	Improvement in depression, P < 0.001
Izadi-Ajirlo et al. (51)	Psychological	RCT	Mastectomized BC patients	23	NA	Cognitive behavioral intervention, 12 × 90-min sessions, 2 sessions per week	Body image and self-esteem/"body image and relationships self- esteem questionnaire"	Improvements in body image and self-esteem due to cognitive behavioral group intervention (P < 0.01).
Rahmani et al. (52)	Psychological	RCT	BC patients	36	44.08 ± 3.3/43.25 ± 3.1/44.92 ± 1.8	Mindfulness-based stress reduction program (8 weekly sessions) vs. metacognition treatment (8 sessions)	Global and specific QoL/QLQ-C30 and QLQ-BR23	Mindfulness-based stress reduction treatment causes effective in QoL improvement
Shabani et al. (53)	Education	Trial	BC patients	50	46.7 ± 9.3	Life skills training classes	GHQ-28 scores	Intervention significantly reduced comarization disorders, sleep disorders, disorders of social functioning, and depressive and anxiety symptoms ($P < 0.0001$)
Rahmani et al. (54)	Psychological	RCT	BC patients	24	٧٧	Group mindfulness-based stress reduction program + conscious yoga (8 weekly group sessions)	QoL/fatigue severity scale, QLQ-C30, QLQ-BR23	Mindfulness-based stress reduction treatment is effective in improving global and specific QoL and fatigue in women with BC
Abbreviations: BC, breast cancer; CG, control group; EG, experimental group; RCT, randomized clinical trials.	st cancer; CG, contr I trials.	ol group; EG, e:	xperimental group;					

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Study	Field	Study Design	Population	Total Sample Size	Age, Mean \pm SD	Outcome/Instrument	Main Results
Montazeri et al. (72)	Complementary use status	Cross sectional descriptive	BC	E	49.5 土 15.1/46.9 土 14.7	Association between QoL and the use of complementary medicine /HADS and QLQ-C30	Significant association between depression and the use of comp. med. (logistic reg.) $P = 0.04$
Montazeri et al. (73)	Complementary use status	Cross sectional	BC	625	49.5 土 15.2/46.6 土 15.1	Factors associated with the use of complementary or alternative medicine	The use of comp/alt. medicine among Iranian patients is not common; however, it is correlated with fear, anxiety and mental distress (P < 0.001 for all)
Garrusi and Faezee(65)	Sexual status	Descriptive	BC	82	49.9 土 11.8	Conceptualize sex and body image/research made questionnaire	Desire was diminished in 70.6% No. of coitus was decreased in 15% of participants. Excitement and orgasm were decreased in 72% of women
Saface et al. (59)	QoL	Cross sectional	BC	611	48.27 ± 11.42	Qol/QLQ-C30	Qol total score 64.92 \pm 24.28. The grade of tumor (P < 0.000), occupational status (P = 0.01), menopausal status (P = 0.01), financial difficulties (P = 0.03), and dyspnea (P = 0.01) were correlated with QoL
Pourhoseingholi et al. (64)	JoD	Cross sectional	BC	611	48.3 ± 11.4	Qol/QLQ-C30	Tumor grade (P < 0.001), occupational status (P = 0.01), menopausal status (P = 0.01), financial difficulties (P = 0.03), and dyspnea (P = 0.01) were correlated with QoL
Hadi et al. (56)	Anger	Case-control	R	578	Case: 48.6 ± 9.16, control: 45.4 ± 7.12	Anger and depression /symptom checklist- 90 revised (SCL-90R)	The mean anger score in BC cases was significantly lower than the control group (0.57 v, $\alpha.4$, $P=0.002$). In cases, higher mean depression and a nger scores were correlated with age ($P=0.008$ and $P=0.020$). Patients with college degrees the def to have higher anxiety scores compared with those with degrees below high school level (1.36 vs. 0.70 ; $P=0.030$). Tumor size of > 2 cm was associated with higher anxiety scores
Vahdaninia et al. (55)	Anxiety and depression	Prospective	BC	167	47.2 土 13.5	Depression and anxiety through 18 months follow-up/HADs	Anxiety and depression improved over the time ($P < 0.001$)
Kiadaliri and Bastani (63)	QoL	Cohort	BC	100	48.49 土 10.63	QoL/QLQ-C30	Adverse effects of chemo regimens after chemo: TAC (64) > FAC (68) ($P < 0.005$); improvement after 4 months: TAC (11.45 pts) > FAC (7.14 pts) $P = 0.02$
Didehdar Ardebil et al. (60)	QoL	Cross sectional descriptive	BC	60	43.81 ± 47.12	Health-related QoL/FACT-B Persian Version	Depression symptoms found in 50% of subjects; significant correlation between depression and overall HRQoI ($\beta ={177}$, P < 0.001); significant dif. Observed between different modalities of treatments: chemotherapy > radiotherapy, P = 0.006
Hatam et al. (61)	QoL	Prospective cohort	BC	100	48.49 ± 10.63	Health-related QoL/QLQ-C30	Decrease in HRQoL due to chemo with TAC > FAC, P < 0.001
Musarezaie et al. (62)	QoL	Cross sectional	BC	330	43.2 ± 5.8	QoL/SF-36 version 2	QoL is inversely correlated with the No. of chemo sessions, P< 0.05 and education

Participants had low mood state and QoL. The mood state was a predictor of	participants Qol. $(\mathbb{R}^{*} = 0.67; \mathbb{P} = 0.007)$ Negative correlation between depression and QoL in caregivers $(r = -0.67, \mathbb{P} = 0.01)$	OoL improves over time (adjustment). However, ratings of sense of coherence, spirituality, and negative religious coping	Sexual dysfunction was 52% before treatment and 84% after treatment; diminished sexual functioning in BC patients; post-treatment sexual disorders were associated with younger age (OR: 0.95; 95% CI, 0.93 · 0.98; P = 0.04), receiving andocrine therapy (OF: 33.4.95% CI, 137- 7.91; P = 0.007), and poor pretreatmment sexual functioning (OK: 12.3. 95% CI, 3.93 - 39.0; P < 0.0001).	57% experienced lubrication problems, 53.8% satisfaction disorders, 42.5% desire disorders, and 77% arousal disorders (all patients vs. healthy controls < 0.01). Patients receiving hormone therapy had more sexual dysfunctions ($P = 0.006$). RT+CT+HT was associated with a 6.fold increase in the risk of lubrication and satisfaction problems (adjusted OR: 6.4; 95% Cl, 4.6 -12.6, and adjusted OR: 5.7; 95% Cl, 3.4 - 11.4)	Healthy eating practices were significantly correlated with social ($r = 0.2$, $P = 0.05$), role ($r = 0.2$, $P = 0.02$), contant end endtional ($r = 0.2$, $P = 0.01$) and endtonal ($r = 0.2$, $P = 0.01$) and reduced symptoms of financial difficulties ($r = 0.2$, $P = 0.026$). Physical activity was significantly correlated with endtonal ($r = 0.2$, $P = 0.03$) scales, 0.004) and cognitive ($r = 0.2$, $P = 0.03$) scales	Significant correlations between nutritional status and QoL scales (physical: $P < 0.001$; emotional: $P = 0.026$; cognitive: $P = 0.020$; global QoL: $P = 0.026$; fatigue: $P < 0.001$; nausea/womiting: $P < 0.001$; pain: $P < 0.001$; dyspnea: $P < 0.001$; appetite loss: $P < 0.001$; constipation: $P < 0.001$; dyspnea: $P < 0.001$; dyspnea: $P < 0.001$; constipation: $P < 0.001$; diarrhea: $P = 0.003$	QoL was positively correlated with spiritual well-being ($P < 0.001$, spiritual well-being, pwell-being, pwell-being, prot), social functioning ($P < 0.001$, and arm symptoms ($P < 0.001$, were significant predictors of QoL	Patients with BC had a significantly higher prevalence of depression.
1-mood states/ the profile of mood states 2-quality of life/	FPQLI QoL and depression/caregiver quality of life index-cancer, Beck depression inventory	Qol, sense of coherence, well-being/ the caregiver Qol INDRX-CANCER; the brief religious coping scale; the spirituality perspective scale; the sense of coherence scale	Sexual function/female sexual function index (FSFI)	Physical function (PF)/SF36	 (1) Ool/QLQ-C30 BR.23, (2) eating practices/women's healthy eating and living (WHEL), (3) physical activity/international physical activity questionnaire(IPAQ) 	(1) nutritional status/patient-generated subjective global assessment(PG-SGA), (2) QoL/QLQ-C30	spiritual well-being/FACIT-Spi2, (2) QoL/QLQ-C30 amd QLQ-BR23	Depression/Beck depression inventory(BDI)
47.3 土 8.62	52.48 土 14.04	$N A \pm N A$	44.3 ± 8.6	<i>377</i> ± 6.4	47,9 ± 6.7	47.8 ± 6.7	48 土 103	45 ± 8.5
50	63	115	216	390	100	100	68	400
Mastectomised BC	BC caregivers	BC family caregivers	BC	BC	BC	BC	BC patientses under radiotherapy	Patients with cancer
Descriptive	Cross sectional, descriptive	Prospective, descriptive, correlational	Prospective	Cross sectional	Cross sectional	Cross sectional	Cross sectional	Prospective
QoL	Caregiver QoL	Caregiver QoL	Sexual status	Sexual status	QoL-relation with lifestyle	QoL-relation with nutritional status	QoL-spiritual	Depression
Tirgari et al. (58)	Heidari Gorji et al. (69)	Khanjari et al. (71)	Harirchi et al. (66)	Safarinejad et al. (67)	Mohammadi et al. (74)	Mohammadi et al. (75)	Jafari et al. (79)	Mashhadi et al. (57)

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Psychological impacts affect family caregivers of women with BC 6 months after the diagnosis as well as directly after the diagnosis	Significant correlation between GH and health-promoting lifestyle (P < 0.05)	Significantly lower scores on physical and one scales, along with fatigue and financial difficulties, during the first 6 months compared with the controls. Women had better scores on QoL ($P < 0.001$) and emotional scale ($P < 0.01$) during the same period of time. SOC ($P < 0.01$) and baseline ratings of several dimensions of HRQoL ($P <0.05$) were the most important predictors of HRQoL changes	(1) No significant correlation between global PSQI and total score on SWBS or its two subscales, (2) global PSQI score was not significantly correlated with total score of RA questionnaire ($P = 0.278$)	Significant relationship between body esteem and hope $(P < 0.01, t = 0.583)$, between body esteem and mental health ($P = 0.00_1, t = 0.472$), between hope and mental health ($P < 0.00_1, t = 0.565$)	39% of patients had sever emotional distress, which was related to their physical functioning ($P < 0.009$). Taking care of children, anxiety, fear, difficulty taking bath and warring clothes, family issues, fever, and nasal dryness were the most conmon issues related to emotional distress ($P < 0.001$)	Older patients and those with longer work experience were less likely to return to work, while patients with no pain, surgery scar, or lymphedema after the treatment were more likely to return to work
Psychological im caregivers of won the diagnosis as v diagnosis	Significant correl health-promoting	Significantly low role scales, along difficulties, durin difficulties, durin difficulties, durin difficulties, durin difficulties, durin better scores on C emotional scale (1 period of time. ST period of the st	(1) No significant correlation PSQI and total score on SWB subscales; (2) global PSQI score significantly correlated with RA questionnaire (P = 0.278)	Significant relationship bet esteem and hope ($P < 0.001$ between body esteem and n = 0.001, r = 0.472), between health ($P < 0.001$, r = 0.565)	39% of patients h distress, which w functioning (P < children, anxiety, children, anxiety, and wearing clott and wearing clott and masal drynes; issues related to € 0.001)	Older patients and thos experience were less lik while patients with no J lymphedema after the t likely to return to work
QoL/the schedule for the evaluation of individual quality of life (SEQO-DW) by interview (at two times: Th: a time close to diagnosis and T2: after 6 months	General health/health-promoting lifestyle profile II developed by Walker et al	 (1) QoL/QLQ-C30. (2) sense of coherence (SOC). (3) spiritual perspective scale, (4) religious coping +//brief religious coping (brief RCOPE) 	Relationship between quality of sleep and spiritual well-being, religious activities/pittsburgh sleep quality index (PSOI), spiritual well-being scale (SWBS), and religious activities (RA) questionnaire	Hope/herth hope index; body esteem/body esteem scale, mental health/symptom checklist 25 (SCL 25) mental health questionnaire	Emotional distress/a standard worldwide questionnaire (NCCN)	Return to work/interview
41.1 土 13.9	47.9 土 11.4	Case: 46.1 ± 9.8, control: 46.6 土 8.4	48 土 6.9	N/A ± N/A	N/A 士 10.96	44.3 土 6.7
88	262	372	80	100	82	175
BC caregivers	BC	BC	BC	Mastectomised BC	BC	BC
Descriptive and prospective	Predictive design	Longitudinal	Cross sectional	Descriptive	Cross sectional	Cross sectional
Caregiver QoL	Social	QoL along with sense of coherence	QoL-quality of sleep	QoL-hope	Emotional distress	Social
Khanjari et al. (70)	Moghaddam Tabrizi et al. (81)	Rohani et al. (76)	Khoramirad et al. (刀)	Heidari et al. (78)	Saeedi-Saedi et al. (68)	Azarkish et al. (80)