RESEARCH ARTICLE

Open Access

Association between changes in body fat and disease progression after breast cancer surgery is moderated by menopausal status



Li-Ni Liu¹, Yung-Chang Lin², Christine Miaskowski³, Shin-Cheh Chen⁴ and Mei-Ling Chen^{2,5,6*}

Abstract

Background: Obesity is linked to poor disease outcomes in breast cancer patients. However, this link was mostly based on body weight or BMI rather than body-fat. The purpose of this study was to evaluate the relationship between body-fat gain and disease progression in Taiwanese women after breast cancer surgery and how this relationship is influenced by menopausal status.

Methods: Body fat percentage was measured 1 day before and 6 months after surgery in 131 women with stages 0–III breast cancer. Disease outcomes (metastasis and death) were assessed by chart review and telephone contact 7 to 8 years after diagnosis. These data were analyzed by multivariate Cox proportional hazard model analysis.

Results: The percentage of women with over 5% gain in body-fat was 56% for premenopausal and 42% for postmenopausal. Rates of distant metastasis and all-cause mortality were 17.6 and 9.9%, respectively over the follow-up period. Distant metastases were predicted in postmenopausal but not premenopausal women with breast cancer by increased body fat percentage (HR = 1.3, p = 0.035), after controlling other potential covariates, including disease severity, estrogen receptor expression, progesterone receptors expression, age, and exercise habit before diagnosis. Survival was not significantly associated with body-fat percentage gains.

Conclusions: Our results suggest that increased body fat percentage 6 months after breast surgery is an important predictor of distant metastasis in postmenopausal Taiwanese women with breast cancer. Clinicians may need to measure patients' body fat periodically. Our findings should be validated in studies with a longer follow-up time.

Keywords: Breast cancer, Menopausal status, Body fat, Disease progression

Background

Breast cancer is the most common female cancer in Taiwan, with more than 10,000 women diagnosed every year [1]. Breast cancer has a relatively high 5-year survival rate of 86.5% [2], compared to the survival rate of 53.3% for all cancers combined. Despite this good prognosis, most breast cancer survivors who have completed treatment are still concerned about recurrence, metastasis, or mortality [3]. Predictive factors for poor outcomes include disease stage, axillary lymph node involvement, negative expression of estrogen receptors (ER) and

progesterone receptors (PR), younger age [4], and exercise [5]. In addition, weight gain was reported to be a risk factor for poor breast cancer outcomes [6–9].

Weight gain during or after breast cancer treatment due to changes in metabolism, physical activity, and dietary intake [10] is well documented. After treatment, 34% to 96% of women with early-stage breast cancer gain 0.9–7 kg (kg) [11–15] and they do not automatically lose this extra weight when treatment ends [16, 17].

Besides gaining weight, these women experience significant increases in the percent of body fat [13, 18, 19]. However, body weight or body mass index (BMI), which can be easily measured, are used in most studies as a surrogate measure for obesity [13, 20]. Since body weight fails to discriminate between body fat and lean mass [21, 22], changes in body fat cannot be precisely ca

⁵School of Nursing, College of Medicine, Chang Gung University, No.259, Wenhua 1st Rd., Guishan Dist., Taoyuan City 33302, Taiwan, Republic of China Full list of author information is available at the end of the article



^{*} Correspondence: mechenl@mail.cgu.edu.tw

²Department of Internal Medicine, Chang Gung Memorial Hospital and Chang Gung University, Taoyuan, Taiwan

ptured by measuring body weight [10, 13, 14]. Body fat, as the main body component inducing higher amounts of circulating estrogens, interleukin 6, tumor necrosis factor alpha, retinol-binding protein-4, C-reactive protein and leptin, may be a more appropriate indicator to monitor than BMI or body weight for its impact on disease outcomes [10, 23–25]. However, no study has elucidated the effect of body fat change after diagnosis on breast cancer outcomes.

In determining the relationship between obesity and occurrence of breast cancer, several lines of evidence suggest an influential role for menopausal status. For example, obesity increases breast cancer risk after menopause but decreases this risk before menopause [13, 26, 27]. In addition, premenopausal women with breast cancer had more weight gain after treatment than did postmenopausal women [16, 28]. It is hypothesized that menopausal status may modify the effect of body fat gain on disease progression in women with breast cancer.

Therefore, this study had two purposes: (1) to evaluate the relationship between body-fat percentage gain and disease progression in Taiwanese women after breast cancer surgery; and (2) to explore the role of menopausal status in moderating the relationship between body-fat percentage change and disease progression.

Methods

Design, setting, and sample

This is a prospective observational study. The sample for this study was drawn from a longitudinal study that examined symptoms after surgery in newly diagnosed breast cancer patients who were treated at Chang Gung Memorial Hospital (Linkuo and Taipei), a large teaching hospital treating cancer patients from all parts of Taiwan. Patients' inclusion criteria have been published [29]. In brief, women with stage 0-III breast cancer who underwent unilateral breast cancer surgery were included. Women were excluded if they had breast cancer surgery on both sides, had distant metastasis at diagnosis, and/or had a defibrillator implanted. Of the 239 women invited to participate between July 2005 and September 2006, 200 provided written informed consent and were enrolled. In the original study, demographic and exercise habit data were acquired at baseline (the day prior breast surgery). Treatment information was collected at 6 months after breast surgery by reviewing medical charts. Anthropometric measures were assessed at baseline and 6 months after surgery. In 2013 (i.e., 7 to 8 years after diagnosis), we further determined these women's disease outcomes (distant metastasis and all-cause mortality) by reviewing medical charts. If the patient did not return to the hospital for more than 3 months and her survival status was not documented in medical chart, a telephone call was made to the patient or her primary caregiver to confirm the survival and metastasis status. Among these 200 women, 131

(65.5%) had complete data for both anthropometric measures and disease progression outcomes. The data from these 131 women were analyzed in this study. For those who had missing data (n = 69), 63 had missing data on body fat information at 6 months, 10 on distant metastasis information, and 6 on all-cause mortality information. Patients who had missing data on anthropometric measures was mainly because they did not come to the hospital for anthropometric measures if no treatment or follow-up was scheduled at 6 months after surgery. Compared to those with valid data, those with missing data were found to be older and had higher percentages of not receiving any chemotherapy or radiotherapy. Other variables (including initial BW, initial BMI, initial fat percentage, exercise habit, the distribution of disease stage, ER/PR status, hormonal therapy, distant metastasis, and allcause mortality) were comparable between those with valid data and those with missing data (p > 0.05). This study was approved by the Institutional Review Board of Chang Gung Memorial Hospital.

Measurements

Anthropometry

Participants' height and weight were measured using a digital electronic professional scale with height rod (Super-View Medical: HW686, Taiwan). BMI is calculated as weight (in kg) divided by height (in meters (m)) squared. Body fat percentage is calculated as body fat (in kg) divided by body weight (in kg). Body fat was measured with a portable hand-to-foot tetrapolar RJL Quantum-X Bioelectrical Impedance Analyzer (BIA) machine (RJL Systems, Clifton Township, MI, USA). BIA has been reported as easy to use, free from radiation exposure, and having good agreement in detecting body composition compared to dual X-ray absorptiometry (DXA) [30]. Given that different body tissues have different levels of electrical conductivity, it will not be detected in all tissues when a small alternating current (500 microamps at a single frequency of 50 kHz) is applied through the body. The extent to which the applied current encounters resistance or impedance is related to the type and amount of tissue through which the current passes. For example, fat tissue has higher impedance than muscle tissue [31, 32]. Based on this property, body composition can be estimated with the software Cyprus 2.7. To obtain the BIA measurement, patients removed their shoes and socks from the foot of their non-operated side and lay in a supine position on an examination table with their legs apart and arms abducted from the body. Four selfadhesive spot electrodes were placed on the dorsal surface of the hand and foot of the non-operated side. To minimize measurement error, patients were told not to eat or drink too much and to refrain from alcohol intake

and exercise for 8 h before this measurement [32]. Anthropometric measurements were made by a trained research assistant.

Disease progression

Data were collected from medical chart review and telephone interviews with patients from April to June 2013 about two disease-progression events: distant organ metastasis induced by breast cancer and all-cause mortality. Approximately 15% of the disease progression data were obtained through telephone interviews. For distant metastasis or death, time to the event was defined as months between the date of cancer diagnosis and date of the event.

Covariates

Demographic and disease/treatment information were collected using a researcher-developed form. Women were considered menopausal if they reported that their menses had stopped for more than 6 months [33]. Disease stage was re-categorized as a binary variable, disease severity: low severity (stage 0-II) and high severity (stage III). The exercise habit was a yes/no categorical variable assessed at baseline. Patients were asked whether they exercised on a regular basis during the past month. Covariates included for analysis were disease stage, ER status, PR status, age, and exercise habit. Treatment modality was not included because it was highly associated with disease stage.

Statistical analysis

The distribution of study variables was described by means, standard deviations (SD), and percentages. Changes in anthropometric variables were defined by subtracting their values before surgery from those 6 months post-surgery. Differences in demographic, exercise habit, anthropometric, disease, treatment, and disease progression characteristics between premenopausal and postmenopausal groups were evaluated by t-test for continuous variables and chi-square test for categorical variables. The effect of body fat percentage change and its interaction with menopausal status on disease progression were determined by multivariate Cox proportional hazard model analysis. The main effect of body fat percentage change adjusted for 5 covariates (disease stage, age, ER, PR, and exercise habit) was first tested. Then, the overall interaction effect of body fat percentage changes and menopausal status was examined. When a significant interaction was found, subgroup analysis was then performed for pre- and post-menopausal groups separately. All analyses were performed using SPSS software, version 19.0 (Chicago, IL). Significance levels were set at 0.05 for two-tailed tests.

Results

Patients' characteristics

Among the 131 women in our sample, 86 (65.6%) were premenopausal and 45 (34.4%) were postmenopausal at diagnosis. Their average age was 46.9 years. Most patients did not exercise regularly (72.5%) and had stage II or earlier disease (80.9%), with 69.5% ER positive and 55.0% PR positive. More than half of the women received breast modified radical mastectomy (MRM) surgery (56.5%). Most women received adjuvant chemotherapy (83.2%) while 55.0% received radiotherapy and 63.4% received hormone therapy. Compared to premenopausal women, postmenopausal women were older (56.1 vs. 42.1 years), more of them were PR negative (66.3 vs. 33.3%), and fewer received radiotherapy (40.0 vs. 62.8%) (Table 1).

Anthropometric measures by menopausal status

The mean BMI before surgery was 23.1 kg/m² for premenopausal women and 24.5 kg/m² for postmenopausal women. The mean body fat before surgery was 17.9 kg for premenopausal women and 20.0 kg for postmenopausal women. For BMI, both groups had similar percentages (around 31%) for >5% gain at 6 months after surgery. However, the percentages of >5% gain in body fat were higher than that of BMI for both premenopausal (56%) and postmenopausal (42%) groups. Premenopausal and postmenopausal groups did not differ significantly in the percentages of >5% gain for all the anthropometric variables (Table 2).

Effect of body fat percentage change on disease progression

The median period between diagnosis and chart review was 7.6 years (range = 7-8.1). During this period, 23 (17.6%) patients developed distant organ metastasis and 13 (9.9%) died. The censored rates for distant organ metastasis and for all-cause mortality were 82.4 and 90.1%, respectively. Before examining the effect of body fat percentage change, we tested the impact of BMI change on study outcomes and found that after adjusting the covariates, no significant main effect of BMI change on both distant metastasis (B = 0.35, p = 0.058) and allcause mortality (B = 0.11, p = 0.066). We then examined the effect of body fat percentage change on study outcomes and also found no significant effect on distant metastasis (B = 0.11, p = 0.063) or all-cause mortality (B = 0.13, p = 0.104). However, disease stage was found to be a significant predictive factor for metastasis and mortality (p < 0.001) (Table 3).

Interaction effect of body fat percentage change and menopausal status on disease progression

Similar to the analysis of main effect, we first checked the interaction effect between BMI change and menopausal

Table 1 Demographic, exercise habit, and disease/treatment information by menopausal status in women with breast cancer

Characteristic	Total	Premenopausal	Postmenopausal	t/χ^2	p
	(n = 131)	(n = 86)	(n = 45)		
	mean ± SD	mean ± SD	mean ± SD		
	n (%)	n (%)	n (%)		
Age (years)	46.9 ± 9.5	42.1 ± 7.0	56.1 ± 6.3	11.2	< 0.001
Exercise habit before diagnosis				3.6	0.056
No	95 (72.5%)	67 (77.9%)	28 (62.2%)		
Yes	36 (27.5%)	19 (22.1%)	17 (37.8%)		
Type of surgery				12.6	< 0.001
BCT	57 (43.5%)	47 (54.7%)	10 (22.2%)		
MRM	74 (56.5%)	39 (45.3%)	35 (77.8%)		
AJCC staging				0.8	0.664
Stage 0/ I	50 (38.2%)	35 (40.7%)	15 (33.3%)		
Stage II	56 (42.7%)	36 (41.9%)	20 (44.4%)		
Stage III	25 (19.1%)	15 (17.4%)	10 (22.3%)		
Estrogen recepto	or			2.9	0.089
Positive	91 (69.5%)	64 (74.4%)	27 (60.0%)		
Negative	40 (30.5%)	22 (25.6%)	18 (40.0%)		
Progesterone rec	eptor			13.0	< 0.001
Positive	72 (55.0%)	57 (66.3%)	15 (33.3%)		
Negative	59 (45.0%)	29 (33.7%)	30 (66.7%)		
Chemotherapy				0.5	0.478
Yes	109 (83.2%)	73 (84.9%)	36 (80.0%)		
No	22 (16.8%)	13 (15.1%)	9 (20.0%)		
Radiotherapy				6.2	0.013
Yes	72 (55.0%)	54 (62.8%)	18 (40.0%)		
No	59 (45.0%)	32 (37.2%)	27 (60.0%)		
Hormone therapy				3.0	0.085
Yes	83 (63.4%)	59 (68.6%)	24 (53.3%)		
No	48 (36.6%)	27 (31.4%)	21 (46.7%)		
Distant organ metastasis				1.0	0.310
Yes	23 (17.6%)	13 (15.1%)	10 (22.2%)		
No	108 (82.4%)	73 (84.9%)	35 (77.8%)		
All-cause death				0.9	0.345
Yes	13 (9.9%)	7 (8.1%)	6 (13.3%)		
No	118 (90.1%)	79 (91.9%)	39 (86.7%)		

AJCC American Joint Committee on Cancer, SD standard deviation, BCT Breast Conserving Therapy, MRM Modified Radical Mastectomy

status and no significant interaction was found for both distant metastasis (B = 0.62, p = 0.071) and all-cause mortality (B = 0.53, p = 0.126). However, after adjusting for covariates, the interaction effects of body-fat percentage change and menopausal status on both distant metastasis (B = 0.29, p = 0.012) and all-cause mortality (B = 0.32, p = 0.033) were significant (Table 4). The effect of body fat percentage change was then examined for each menopausal group separately. For premenopausal

women (n=86), body-fat percentage change was not associated with either distant metastasis or all-cause mortality. However, for postmenopausal women, higher gain in body-fat percentage was associated with higher risk of disease metastasis (HR = 1.3, p=0.035) and marginally associated with higher risk of all-cause mortality (HR = 1.5, p=0.091). For covariate effects, disease stage was consistently found to be a significant predictor on metastasis and mortality for both pre- and postmenopausal women (p<0.001). Increased in age was associated with lower risk of disease metastasis (HR = 0.9, p=0.029) in pre-menopausal women only (Table 5).

Discussion

Our study contributes to knowledge on the effect of obesity on breast cancer outcomes by clarifying that the effect of weight gain previously reported [6–8] is due to changes in body fat composition after breast cancer surgery. After adjusting for the effects of covariates, we found that increased body-fat percentage predicted distant metastasis in postmenopausal women after breast cancer surgery, but not in premenopausal women.

Many previous studies have demonstrated the association between obesity and risk of disease recurrence [11, 34] or mortality [6, 13, 35, 36] in breast cancer women. However, still other studies failed to find this association [7, 37]. The way obesity was measured in these studies included weight/ BMI, waist-hip ratio, or weight gain after diagnosis. None of these studies used the direct measure of body fat to represent obesity. In addition, none of these studies considered the potential role of menopausal status. In the current study, we directly examined the relationship between bodyfat percentage change and the disease progression. When menopausal status was not considered in the model, the relationship between body-fat percentage gain and the metastasis/mortality was only marginal which did not reach statistical significance of 0.05. However, when breast cancer women were divided into pre- and post-menopausal groups, the effect of body fat percentage gain was found to be significant in predicting metastasis in postmenopausal women but not in premenopausal women. Therefore, it is suspected that the inconsistent findings in previous studies may be due to indirect measures of obesity and lack of consideration of menopausal status.

Our findings suggest that menopausal status moderates the link between body-fat gain and disease metastasis in breast cancer women. Gain in body-fat percentage after breast cancer surgery was a predictive factor only for postmenopausal women. This finding may be explained by different sources of estrogen production in premenopausal and postmenopausal women. For premenopausal women, the ovaries are the main source of estrogen production, whereas adipose tissue serves as the extragonadal source of estrogen for postmenopausal

Table 2 Changes in anthropometric variables by menopausal status in women with breast cancer (N = 131)

	Premenopausal ($n = 86$)			Postmenopausal $(n = 45)$				
Anthropometry	Presurgery (M ± SD)	Post- surgery ^a (M ± SD)	Gain >5% n (%)	Presurgery (M ± SD)	Post-surgery ^a (M ± SD)	Gain >5% n (%)	χ²	р
Weight (kg)	56.5 ± 8.2	58.3 ± 8.2	27 (31.4%)	58.1 ± 9.2	59.1 ± 8.3	14 (31.1%)	< 0.1	0.973
BMI (kg/m^2)	23.1 ± 3.0	23.8 ± 3.1	27 (31.4%)	24.5 ± 3.3	24.9 ± 2.9	14 (31.1%)	< 0.1	0.973
Body fat (kg)	17.9 ± 5.8	19.1 ± 6.0	48 (55.8%)	20.0 ± 6.1	20.4 ± 5.5	19 (42.2%)	2.2	0.139
Body fat (%)	31.0 ± 6.4	32.1 ± 6.3	36 (41.9%)	33.7 ± 6.3	33.9 ± 5.6	14 (31.3%)	1.4	0.229

BMI body mass index, M mean, SD standard deviation a Six months after surgery

women [38]. In postmenopausal women, adipose tissue contains aromatase that can convert androgen to estrogen [38]. Therefore, increased body fat in these women may increase estrogen concentrations, consequently leading to a poorer outcome [39]. In addition, obese postmenopausal women have lower serum levels of sex hormone-binding globulin (SHBG) than obese premenopausal women [10]. SHBG can bind and inhibit estradiol, a kind of estrogen-like hormone. Lower SHBG levels will result in increased levels of circulating unbound estrogen that may promote tumor progression [10].

Based on our study findings, we recommend that postmenopausal women with breast cancer maintain or decrease their body fat during or after treatment. However, the traditional clinical practice is to advise patients to maximize caloric intake to ensure a good physical condition against the toxic effects of adjuvant chemotherapy [10]. Based on our findings, we think this traditional diet suggestion may not be valid for all cancer types. The diet suggestion for

Table 3 Effects of body fat percentage change on distant metastasis and all-cause mortality in women with breast cance (N = 131)

,	B (± SE)	Hazard ratio (95% CI)	р
Metastasis			
Advanced disease (Stage III)	1.90 (± 0.45)	6.71 (2.78–16.17)	< .001
ER positive	0.82 (± 0.56)	2.26 (0.75-6.84)	.148
PR positive	-0.91 (± 0.52)	0.40 (0.15-1.11)	.080.
Age	-0.02 (± 0.03)	0.98 (0.93–1.03)	.372
Regular exercise	0.06 (± 0.53)	1.07 (0.38–3.00)	.905
% body fat change	0.11 (± 0.06)	1.11 (0.99–1.25)	.063
Death			
Advanced disease (Stage III)	2.97 (± 0.69)	19.49 (5.10–74.56)	< .001
ER positive	0.41 (± 0.74)	1.51 (0.36–6.38)	.576
PR positive	-0.98 (± 0.69)	0.37 (0.10-1.45)	.155
Age	0.02 (± 0.04)	1.02 (0.95-1.10)	.537
Regular exercise	-0.30 (± 0.72)	0.74 (0.18–3.05)	.676
% body fat change	0.13 (± 0.08)	1.14 (0.97–1.33)	.104

Body fat % change = body fat percentage at 6 months after surgery minus body fat percentage before surgery

ER estrogen receptor, PR progesterone receptor

breast cancer survivors may need to modify, at least for postmenopausal women. Future diet or exercise interventions that aim to facilitate body fat loss and retain lean body mass may be considered.

This study has several limitations. First, the BIA machine (RJL Systems, Clifton Township, MI, USA) used in this study may be less accurate in Asian women than in western women, who have relatively longer limbs. This difference may underestimate body-fat percentage in shorter limbed persons like Asian women [40], despite a report that the BIA machine is easy to use and has good accuracy in detecting changes in women's body composition [30]. Second, the gain in body fat percentage was defined by the difference between measures at pre-surgery and 6 months post-surgery. Some women may recover from the temporary weight or body fat gain caused by the cancer treatment.

Table 4 The interaction effect of body fat percentage change and menopausal status on distant metastasis and all-cause mortality in women with breast cancer (N = 131)

	B (± SE)	Hazard ratio	р
	D (± 3L)	(95% CI)	Ρ
Metastasis			
Advanced disease (Stage III)	2.05 (± 0.49)	7.80 (3.00–20.28)	< .001
ER positive	0.87 (± 0.58)	2.39 (0.76–7.45)	.134
PR positive	-0.75 (± 0.52)	0.47 (0.17-1.30)	.146
Age	-0.03 (± 0.03)	0.97 (0.93–1.02)	.235
Regular exercise	0.01 (± 0.52)	1.01 (0.37–2.82)	.978
Body fat % change × menopausal status	0.29 (± 0.11)	1.33 (1.06–1.66)	.012
Death			
Advanced disease (Stage III)	3.22 (± 0.79)	25.09 (5.34–117.89)	< .001
ER positive	0.41 (± 0.74)	1.50 (0.35-6.45)	.585
PR positive	-0.77 (± 0.66)	0.46 (0.13–1.70)	.246
Age	0.02 (± 0.03)	1.02 (0.96–1.09)	.555
Regular exercise	-0.32 (± 0.72)	0.73 (0.18–2.98)	.660
Body fat % change × menopausal status	0.32 (± 0.15)	1.18 (1.03–1.86)	.033

Body fat % change = body fat percentage at 6 months after surgery minus body fat percentage before surgery ER estrogen receptor, PR progesterone receptor

Table 5 Effect of body fat percentage change on distant metastasis and survival by menopausal status in women with breast cancer

	Premenopausal (n = 86)			Postmenopausal (n = 45)		
	B (± SE)	HR (95% CI)	р	B (± SE)	HR (95% CI)	р
Metastasis						
Advanced disease (Stage III)	1.9 (± 0.6)	6.4 (1.83–22.64)	.004	2.5 (± 1.0)	11.7 (1.55–88.07)	.017
ER positive	1.7 (± 0.9)	5.5 (0.86–35.11)	.071	$-0.3 (\pm 0.9)$	0.8 (0.13-4.59)	.781
PR positive	$-1.3 (\pm 0.8)$	0.3 (0.06–1.22)	.088	0.3 (± 0.9)	1.4 (0.21-8.70)	.749
Age	$-0.1 (\pm 0.1)$	0.9 (0.83-0.99)	.029	0.1 (± 0.1)	1.0 (0.90-1.12)	.940
Regular exercise	-0.4 (± 1.1)	0.7 (0.08–6.00)	.739	0.1 (± 0.7)	1.1 (0.24–4.57)	.945
Body fat % change	0.1 (± 0.1)	1.1 (0.95–1.21)	.278	0.3 (± 0.1)	1.3 (1.02–1.72)	.035
Death						
Advanced disease (Stage III)	3.2 (± 1.0)	24.7 (3.77–161.63)	.001	4.7 (± 1.9)	106.5 (2.78–4084.85)	.012
ER positive	2.1 (± 1.3)	8.6 (0.71–103.50)	.091	-0.9 (± 1.3)	0.4 (0.03-4.88)	.476
PR positive	$-2.0 (\pm 1.1)$	0.1 (0.02–1.14)	.067	$-0.5 (\pm 1.4)$	0.6 (0.04–10.19)	.749
Age	$-0.1 (\pm 0.1)$	0.9 (0.87–1.11)	.769	0.1 (± 0.1)	1.1 (0.92–1.29)	.298
Regular exercise	0.3 (± 1.2)	1.3(0.13-13.24)	.815	-1.1 (± 1.1)	0.3(0.04-2.75)	.312
Body fat % change	0.1 (± 0.1)	1.1(0.95-1.33)	.188	0.4 (± 0.2)	1.5(0.94-2.25)	.091

BMI body mass index, ER estrogen receptor, PR progesterone receptor, CI confidence interval

Third, 3–5% of the data on disease outcomes were missing due to patients being lost to follow-up. These patients may have had worse or better disease outcomes than study participants. Fourth, the accuracy of this study's predictive ability may have been affected by its high censoring rate (82.4% for distant metastasis; 90.1% for all-cause mortality), reflecting that time to distant metastasis or all-cause mortality was unavailable due to participants lost to follow-up or the outcome event not occurring before study end [41]. Finally, the findings of this study are limited to the women recruited from one hospital and cannot be generalized to all women with breast cancer in Taiwan.

Conclusions

This study highlights the role of increased body fat percentage after breast cancer treatment on disease progression in postmenopausal women. We suggest periodically measuring not only body weight but also body fat and monitoring changes in body fat percentage. More research is needed on the effect of changes in body fat on breast cancer survival in women with different menopausal status.

Abbreviations

BIA: Bioelectrical impedance analyzer; BMI: Body mass index; BW: Body weight; DXA: Dual X-ray absorptiometry; ER: Estrogen receptors; HR: Hazard ratio; PR: Progesterone receptors; SHBG: Sex hormone-binding globulin

Acknowledgements

The authors acknowledge the Chang Gung Memorial Hospital, Chang Gung University and the Fu Jen Catholic University, for support in collecting the data and providing the research facilities.

Funding

This study was supported by a research grant to Dr. Mei-Ling Chen from the National Science Council of Taiwan (Grant No: 94–2314-B-182-025) and a research grant to Dr. Li-Ni Liu from the Taiwan Nurses Association (Grant No: TWNA- 1022019).

Availability of data and materials

All relevant materials are provided in the manuscript.

Authors' contributions

LNL and MLC were responsible for the study conception and design, performed the data analysis, obtained funding, and supervised the study. LNL performed the data collection and was responsible for drafting of the manuscript. YCL and SCC provided administrative, technical or material support. MLC, CM, YCL, and SCC made critical revision to the paper for important intellectual content. All authors read and approved the final manuscript.

Ethics approval and consent to participate

This study was approved by the Institutional Review Board of Chang Gung Memorial Hospital (No.102-1102B). All subjects provided written informed consent prior to participation.

Consent for publication

Not applicable.

Competing interests

The authors declare no conflict of interest. The funding body had no role in the design of the study and collection, analysis, and interpretation of data and in writing the manuscript.

Publisher's Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

¹Department of Nursing, Fu Jen Catholic University, New Taipei City, Taiwan. ²Department of Internal Medicine, Chang Gung Memorial Hospital and Chang Gung University, Taoyuan, Taiwan. ³Department of Physiological Nursing, School of Nursing, University of California, San Francisco, USA. ⁴Department of Surgery, Chang Gung Memorial Hospital and Chang Gung University, Taoyuan, Taiwan. ⁵School of Nursing, College of Medicine, Chang Gung University, No.259, Wenhua 1st Rd., Guishan Dist., Taoyuan City 33302, Taiwan, Republic of China. ⁶Department of Nursing, Chang Gung University of Science and Technology, Taoyuan, Taiwan.

Received: 13 June 2016 Accepted: 29 November 2017 Published online: 18 December 2017

References

- Taiwan Cancer Registry: Breast cancer incidence rates in Taiwan. http://tcr. cph.ntu.edu.tw/main.php?Page=N2 (2011). Accessed 5 Feb 2015.
- Taiwan Cancer Registry: Breast cancer survival rates in Taiwan. http://tcr.cph. ntu.edu.tw/uploadimages/Survival_96_100.pdf (2011). Accessed 05 Feb 2015.
- van den Beuken-van Everdingen MH, Peters ML, de Rijke JM, Schouten HC, van Kleef M, Patijn J. Concerns of former breast cancer patients about disease recurrence: a validation and prevalence study. Psychooncology. 2008;17:1137–45. 10.1002/pon.1340.
- Liedtke C, Rody A, Gluz O, Baumann K, Beyer D, Kohls EB, et al. The prognostic impact of age in different molecular subtypes of breast cancer. Breast Cancer Res Treat. 2015;152:667–73. 10.1007/s10549-015-3491-3.
- Dieli-Conwright CM, Orozco BZ. Exercise after breast cancer treatment: current perspectives. Breast Cancer (Dove Med Press). 2015;7:353–62. 10. 2147/BCTT.S82039.
- Kroenke CH, Chen WY, Rosner B, Holmes MD. Weight, weight gain, and survival after breast cancer diagnosis. J Clin Oncol. 2005;23:1370–8.
- Chen X, Lu W, Zheng W, Gu K, Chen Z, Zheng Y, et al. Obesity and weight change in relation to breast cancer survival. Breast Cancer Res Treat. 2010; 122:823–33. 10.1007/s10549-009-0708-3.
- Thivat E, Therondel S, Lapirot O, Abrial C, Gimbergues P, Gadea E, et al. Weight change during chemotherapy changes the prognosis in non metastatic breast cancer for the worse. BMC Cancer. 2010;10:648. 10.1186/ 1471-2407-10-648.
- Bradshaw PT, Ibrahim JG, Stevens J, Cleveland R, Abrahamson PE, Satia JA, et al. Postdiagnosis change in bodyweight and survival after breast cancer diagnosis. Epidemiology. 2012;23:320–7. 10.1097/EDE.0b013e31824596a1.
- Champ CE, Volek JS, Siglin J, Jin L, Simone NL. Weight gain, metabolic syndrome, and breast cancer recurrence: are dietary recommendations supported by the data? Int J Breast Cancer. 2012;2012:506868. 10.1155/ 2012/506868.
- Caan BJ, Emond JA, Natarajan L, Castillo A, Gunderson EP, Habel L, et al. Post-diagnosis weight gain and breast cancer recurrence in women with early stage breast cancer. Breast Cancer Res Treat. 2006;99:47–57.
- 12. Liu LN, Wen FH, Miaskowski C, Lin YC, Wang JS, Jeng C, et al. Weight change trajectory in women with breast cancer receiving chemotherapy and the effect of different regimens. J Clin Nurs. 2014;23:2757–68. 10. 1111/jocn.12521.
- James FR, Wootton S, Jackson A, Wiseman M, Copson ER, Cutress RI. Obesity in breast cancer—what is the risk factor? Eur J Cancer. 2015;51:705–20. 10. 1016/j.ejca.2015.01.057.
- Sheean PM, Hoskins K, Stolley M. Body composition changes in females treated for breast cancer: a review of the evidence. Breast Cancer Res Treat. 2012;135:663–80. 10.1007/s10549-012-2200-8.
- Caan BJ, Kwan ML, Shu XO, Pierce JP, Patterson RE, Nechuta SJ, et al. Weight change and survival after breast cancer in the after breast cancer pooling project. Cancer Epidemiol Biomark Prev. 2012;21:1260–71. 10.1158/ 1055-9965.epi-12-0306.
- Makari-Judson G, Judson CH, Mertens WC. Longitudinal patterns of weight gain after breast cancer diagnosis: observations beyond the first year. Breast J. 2007;13:258–65.
- Saquib N, Flatt SW, Natarajan L, Thomson CA, Bardwell WA, Caan B, et al. Weight gain and recovery of pre-cancer weight after breast cancer treatments: evidence from the women's healthy eating and living (WHEL) study. Breast Cancer Res Treat. 2007;105:177–86. 10.1007/s10549-006-9442-2.
- Liu LN. Body weight, body fat, and other body compositions in patients with breast cancer after chemotherapy: the long-term change and related factors.
 2010. Unpublished PHD dissertation, Chang Gung University, Taiwan.
- Vance V, Mourtzakis M, McCargar L, Hanning R. Weight gain in breast cancer survivors: prevalence, pattern and health consequences. Obes Rev. 2011;12:282–94. 10.1111/j.1467-789X.2010.00805.x.
- 20. Wang C, Hou XH, Zhang ML, Bao YQ, Zou YH, Zhong WH, et al. Comparison of body mass index with body fat percentage in the evaluation of obesity

- in Chinese. Biomed Environ Sci. 2010;23:173–9. 10.1016/S0895-3988(10)60049-9.
- Ode JJ, Pivarnik JM, Reeves MJ, Knous JL. Body mass index as a predictor of percent fat in college athletes and nonathletes. Med Sci Sports Exerc. 2007; 39:403–9. 10.1249/01.mss.0000247008.19127.3e.
- Romero-Corral A, Somers VK, Sierra-Johnson J, Thomas RJ, Collazo-Clavell ML, Korinek J, et al. Accuracy of body mass index in diagnosing obesity in the adult general population. Int J Obes. 2008;32:959–66. 10.1038/ijo.2008.11.
- Cirillo D, Rachiglio AM, la Montagna R, Giordano A, Normanno N. Leptin signaling in breast cancer: an overview. J Cell Biochem. 2008; 105:956–64. 10.1002/jcb.21911.
- 24. Mantovani A, Allavena P, Sica A, Balkwill F. Cancer-related inflammation. Nature. 2008;454:436–44. 10.1038/nature07205.
- Park J, Morley TS, Kim M, Clegg DJ, Scherer PE. Obesity and cancermechanisms underlying tumour progression and recurrence. Nat Rev Endocrinol. 2014;10:455–65. 10.1038/nrendo.2014.94.
- Dowsett M, Folkerd E. Reduced progesterone levels explain the reduced risk of breast cancer in obese premenopausal women: a new hypothesis. Breast Cancer Res Treat. 2015;149:1–4. 10.1007/s10549-014-3211-4.
- Reeves GK, Pirie K, Beral V, Green J, Spencer E, Bull D. Cancer incidence and mortality in relation to body mass index in the million women study: cohort study. BMJ. 2007;335:1134.
- Heideman WH, Russell NS, Gundy C, Rookus MA, Voskuil DW. The frequency, magnitude and timing of post-diagnosis body weight gain in Dutch breast cancer survivors. Eur J Cancer. 2009;45:119–26. 10.1016/j.ejca.2008.09.003.
- Liu LN, Miaskowski C, Wang JS, Chen SC, Chen ML. Accuracy of body mass index to determine obesity in women with breast cancer: an observational study of Taiwanese sample. Int J Nurs Stud. 2010;47:994–1000. 10.1016/j. ijnurstu.2010.01.002.
- 30. Thomson R, Brinkworth GD, Buckley JD, Noakes M, Clifton PM. Good agreement between bioelectrical impedance and dual-energy X-ray absorptiometry for estimating changes in body composition during weight loss in overweight young women. Clin Nutr. 2007;26:771–7.
- Going S, Nichols J, Loftin M, Stewart D, Lohman T, Tuuri G, et al. Validation of bioelectrical impedance analysis (BIA) for estimation of body composition in black, white and Hispanic adolescent girls. Int J Body Compos Res. 2006;4:161–7.
- The Department of Nutrition and Food Sciences: Bioelectrical impedance analysis. http://nutrition.uvm.edu/bodycomp/bia/bia-toc.html (2006). Accessed 16 Dec 2014.
- 33. Rossi R, Nuzzo A, Origliani G, Modena MG. Metabolic syndrome affects cardiovascular risk profile and response to treatment in hypertensive postmenopausal women. Hypertension. 2008;52:865–72. 10.1161/ HYPERTENSIONAHA.108.110478.
- Camoriano JK, Loprinzi CL, Ingle JN, Therneau TM, Krook JE, Veeder MH. Weight change in women treated with adjuvant therapy or observed following mastectomy for node-positive breast cancer. J Clin Oncol. 1990;8:1327–34.
- Nichols HB, Trentham-Dietz A, Egan KM, Titus-Ernstoff L, Holmes MD, Bersch AJ, et al. Body mass index before and after breast cancer diagnosis: associations with all-cause, breast cancer, and cardiovascular disease mortality. Cancer Epidemiol Biomark Prev. 2009;18:1403–9.
- Kwan ML, John EM, Caan BJ, Lee VS, Bernstein L, Cheng I, et al. Obesity and mortality after breast cancer by race/ethnicity: the California breast cancer survivorship consortium. Am J Epidemiol. 2014;179:95–111.
- Goodwin PJ, Panzarella T, Boyd NF. Weight gain in women with localized breast cancer: a descriptive study. Breast Cancer Res Treat. 1988;11:59–66.
- Cleary MP, Grossmann ME. Minireview: obesity and breast cancer: the estrogen connection. Endocrinology. 2009;150:2537–42. 10.1210/en.2009-0070.
- Rock CL, Flatt SW, Laughlin GA, Gold EB, Thomson CA, Natarajan L, et al. Reproductive steroid hormones and recurrence-free survival in women with a history of breast cancer. Cancer Epidemiol Biomark Prev. 2008;17:614–20. 10.1158/1055-9965.EPI-07-0761.
- Snijder MB, Kuyf BE, Deurenberg P. Effect of body build on the validity of predicted body fat from body mass index and bioelectrical impedance. Ann Nutr Metab. 1999;43:277–85.
- Prinja S, Gupta N, Verma R. Censoring in clinical trials: review of survival analysis techniques. Indian J Community Med. 2010;35:217–21. 10.4103/ 0970-0218.66859.