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Quality of Life among Long-Term Survivors of Advanced Stage Ovarian Cancer: A Cross-Sectional Approach

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Abstract

Purpose—Long-term survival of women with advanced-stage ovarian cancer is relatively rare. Little is known about quality of life (QOL) and survivorship concerns of these women. Here, we describe QOL of women with advanced-stage ovarian cancer surviving for 8.5 years or longer and compare women with 0–1 recurrence to those with multiple recurrences.

Methods—Participants ($n=56$) recruited from 5 academic medical centers and the Ovarian Cancer Research Fund Alliance completed surveys regarding QOL (FACT-O), mood (CESD), social support (SPS), physical activity (IPAQ-SF), diet, and clinical characteristics. Median survival was 14.0 years (range 8.8–33.3).

Results—QOL and psychological adjustment of long-term survivors was relatively good, with mean FACT-G scores (multiple recurrences: 80.81 ± 13.95 ; 0–1 recurrence: 89.05 ± 10.80) above norms for healthy community samples (80.1 ± 18.1). Survivors with multiple recurrences reported more compromised QOL in domains of physical and emotional well-being ($p < .05$), and endorsed a variety of physical and emotional concerns compared to survivors with 0–1 recurrence. Difficulties in sexual functioning were common in both groups. Almost half (43%) of the survivors reported low levels of physical activity.

Conclusions—Overall, women with advanced-stage ovarian cancer who have survived at least 8.5 years report good QOL and psychological adjustment. QOL of survivors with multiple recurrences is somewhat impaired compared to those with 0–1 recurrence. Limitations include a possible bias towards participation by healthier survivors, thus under-representing the level of compromise in long-term survivors. Health care practitioners should be alert to psychosocial issues faced by these long-term survivors to provide interventions that enhance QOL.

Keywords

Ovarian cancer; long-term survivor; psychosocial; quality of life; mood; lifestyle

Introduction

Although the majority of advanced-stage ovarian cancer patients have poor survival, a subset of patients live for many years after diagnosis. Women with stage III disease have 5- and 10-year survival rates of 36% and 23%, respectively, with rates of 17% and 8% for stage IV disease [1]. Clinical predictors of long-term survival have recently been described [2]; however, quality of life (QOL) and survivorship concerns of long-term advanced-stage survivors have been minimally characterized, thus little is known about the needs of this population.

Studies of ovarian cancer survivors within the first few years post-diagnosis frequently describe elevated distress, depression, anxiety, and sexual concerns in this population [3]. These negative sequelae are compounded for survivors experiencing physical complications and treatment side effects. In contrast, some survivors report personal growth and strengthened relationships post-diagnosis [3]. Prior studies of long-term adjustment of ovarian cancer survivors predominantly included early-stage survivors, or women surviving for at least three or five years without recurrence [4–6]. In one study of women with advanced stage non-recurrent disease averaging 6 years post-diagnosis, 64% of survivors reported mental health at or above medical outpatient norms, and most (71.4%) reported a strong sense of life purpose [7]. However, a subset of these survivors (28.6%) reported feeling depressed and 45.2% reported substantial anxiety [7].

The 2016 Institute of Medicine report on the state of ovarian cancer research highlighted the need for improved patient care across the continuum of survivorship [8]. Although ovarian cancer patients have been surviving longer [9, 10] there has been minimal characterization of QOL needs of advanced-stage ovarian cancer patients surviving for 8.5 years or longer post-diagnosis. Furthermore, there is substantial heterogeneity in disease course among long-term survivors [2]. Although some live for many years recurrence-free and disease-free, others have multiple recurrences, and may have long-lasting intermittent treatment. From a clinical perspective, long-term survivors with a single recurrence tend to be regarded similarly as those with no recurrence in terms of prognosis and clinical management [2]. This study thus examined QOL, survivorship concerns, and lifestyle factors (e.g. exercise) among long-term (8.5+ years) survivors of advanced-stage epithelial ovarian cancer, and compared women with 0–1 recurrences to those with multiple recurrences. We hypothesized that survivors with multiple recurrences would have poorer QOL, more survivorship concerns, higher levels of distress, poorer well-being and relationships, and be less physically active than those with 0–1 recurrence.

Methods

Participants

Women with epithelial ovarian, peritoneal, or fallopian tube cancer were recruited from Cedars-Sinai Medical Center, Memorial Sloan Kettering Cancer Center, University of Iowa Holden Comprehensive Cancer Center, Stephenson Cancer Center at the University of Oklahoma Health Sciences Center, University of Texas MD Anderson Cancer Center (MDACC), and the Ovarian Cancer Research Fund Alliance (OCRFA). The study was

approved by the IRB of each academic site. Potentially eligible participants from medical centers were screened for eligibility and enrolled at a clinic visit if interested in participation. OCRFA recruited participants via online advertisements; these survivors contacted MDACC, and, if eligible, were enrolled by MDACC by phone. Surveys were completed at one time-point by internet or by hard copy if the patient was reluctant to use the internet-based-survey platform. Patients were eligible if they a) spoke English and b) were at least 8.5 years from diagnosis with stage III–IV epithelial ovarian cancer. All participants provided written informed consent.

The precise cut-point for designating a woman as a long-term survivor of ovarian cancer is currently not specifically defined; however, examination of conditional survival rates in the general vicinity of 10 years (e.g., between 8.5 and 10 years) do not show clear differences [11]. We performed a sensitivity analysis to examine cutoffs of 8.5, 9, and 10 years, but changing this cutoff did not alter the pattern of results. In the present study, therefore, a cutoff of 8.5 years was employed to maximize sample size while preserving relative homogeneity of outcomes.

Psychosocial Assessments

Quality of Life—Quality of Life was measured using the *Functional Assessment of Cancer Therapy-Ovarian* (FACT-O), a validated 38-item questionnaire consisting of physical, social, functional, and emotional well-being subscales (comprising the FACT-G) plus a disease-specific subscale measuring ovarian cancer-specific concerns [12, 13]. An additional FACT-Spiritual subscale was also included. Lower scores on the FACT indicate greater impairment.

Depressed Mood—Depressed Mood was measured using the *Center for Epidemiological Studies Depression Scale* (CES-D), a well-validated 20-item measure. Scores of 16 and above indicate clinically significant levels of depressed mood [14].

Social Support—Social Support was measured by the *Social Provisions Scale*, a 24-item self-report scale measuring extent to which social relationships are perceived as supportive. Items are rated on a 4-point scale with higher scores indicating greater support [15]. The 7-item *Abbreviated Dyadic Adjustment Scale* (ADAS) was used to measure marital adjustment and satisfaction [16].

Psychological Well-Being—Psychological Well-Being was assessed using four subscales from the Ryff *Psychological Well-Being Scales* (PWBS) [17]. Each scale consists of 7 items rated on a 7-point scale, with higher scores indicating greater well-being.

Physical Activity—Physical Activity was measured with the 7-item *International Physical Activity Questionnaire Short-Form* (IPAQ-SF), which assesses physical activity over the past 7 days [18].

Each survey above has been extensively validated.

Demographic and Clinical Characteristics—Demographic and clinical information was provided by self-report. Upper body obesity was measured using the *waist-to-hip ratio* [19] with a tape measure and DVD provided for instructions. Body mass index (BMI) was categorized into underweight (< 18.5), normal weight (18.5 and < 25), overweight (25 and < 30) and obese/morbidly obese (≥ 30).

Descriptive assessments—Changes in social support and marital relationships after cancer were assessed with 9 descriptive survey items designed for this study asking about changes in quality of relationships with friends and partners and changes in sexual health since diagnosis. Two additional questions addressed change in diet and exercise. Alternative and complementary treatment use was assessed with 6 survey items regarding use of complementary methods since diagnosis. (See Supplement 2)

Statistical Analyses

Stata v14.1 (College Station, TX) was used to analyze data. All distributions were examined for outliers and assumptions of non-normality. Descriptive statistics were used to examine dependent variables. ANOVAs and the Wilcoxon rank-sum test were conducted to compare differences between survivor groups on continuous measures. Fisher's exact test and Chi squared tests were used for categorical measures. To affirm the validity of dividing survivors into two groups based on number of recurrences, preliminary analyses were performed comparing the primary QOL outcome variable, FACT-O total scores, between survivors with 0 recurrences, one recurrence, and multiple recurrences. A 3-group analysis of variance (ANOVA) confirmed that the groups were significantly different ($p=0.045$). Post-hoc tests indicated that survivors with no recurrences had significantly higher FACT-O scores (mean=125.18±15.23) than those with multiple recurrences (mean=113.56±17.35, $p=0.026$), but were not significantly different from survivors with only one recurrence (mean=125.91±13.15, $p=0.89$). Moreover, FACT-O scores of survivors with one recurrence were lower than means of those with multiple recurrences ($p=0.057$), but this did not reach significance, likely due to the small sample size. As 3–7 points on the FACT represents a clinically significant difference in QOL [13], these data suggest relative similarity between survivors with 0 and 1 recurrence along with clinically significant differences between these survivors and those with multiple recurrences, thus supporting the use of a two group analytic strategy (i.e., 0–1 recurrence vs. multiple recurrences) for testing the present hypotheses. Thus, analyses compared survivors in two groups: 40 women reporting 0–1 recurrence and 16 women reporting two or more recurrences or persistent disease, categorized as multiple recurrences. A p -value of 0.05 was used for statistical significance.

Results

Participant Characteristics

Of the potentially eligible patients at referring sites, 6 refused and 8 were not approached because of health issues. Of the 84 women who indicated interest, 51 were from OCFRA and 33 were from consortium sites. Two were ineligible, 5 withdrew, and 21 did not complete surveys. (Figure 1). The final sample included 56 long-term survivors with advanced-stage disease at diagnosis (34 from OCFRA and 22 from clinical sites). Of these

women, 91.1% (n=51) were at least 10 years post-diagnosis (median survival time 14.0 years (range 8.8–33.3)).

Mean age of participants was 65.5 (range: 40.2–85.9) years. Respondents were predominantly white, non-Hispanic, and well-educated; more than half (55.4%) had completed college and/or graduate/professional school. About two thirds (66.1%) were married or living with a partner. Half of the participants (50.0%, n=28) reported no recurrences, 21.4% (n=12) reported one recurrence, 25.0% (n=14) reported multiple recurrences, and 3.6% (n=2) reported persistent disease. Treatment status included 73.2% (n=41) of survivors not receiving treatment, 21.4% (n=12) currently receiving treatment, and 5.4% (n=3) not specified (Table 1–Table 2).

Quality of Life

In addition to the group differences in total FACT-O scores described above, survivors with multiple recurrences reported significantly poorer QOL on each of the two major components of the FACT-O score, namely, the total FACT-G score and the ovarian-specific item scale. The FACT-G is useful because it can be compared to population norms. FACT-G scores of survivors with multiple recurrences (80.81 ± 13.95) were significantly poorer than those of survivors with 0–1 recurrence (89.05 ± 10.80 ; $p=0.031$). (Table 3). Although these data suggest clinical decrements in QOL among survivors with multiple recurrences, mean scores of both survivor groups approximated normative FACT-G scores of the U.S. population (80.1 ± 18.1) [13], suggesting that overall QOL was quite good. Survivors with multiple recurrences had poorer scores on the ovarian concerns subscale (32.75 ± 5.25) than those with 0–1 recurrence (36.30 ± 4.54 , $p=0.009$), indicating greater impairment from ovarian-specific issues.

FACT subscale scores of survivors with multiple recurrences were significantly lower than those of survivors with 0–1 recurrence on Physical Well-being (PWB) (0–1: 25.43 ± 2.72 vs. multiple: 23.25 ± 3.49 , $p < 0.02$) and Emotional Well-being (EWB) (0–1: 20.69 ± 2.92 vs. multiple: 17.88 ± 4.44 , $p < 0.02$) but not Functional well-being (FWB) (0–1: 23.00 ± 4.22 vs. multiple: 20.81 ± 5.60 , $p=0.18$) (Table 3). As subscale scores of 2–3 points reflect clinically significant differences [13], these decrements indicate potential clinically significant impairments in each domain. Moreover, mean EWB scores of survivors with multiple recurrences (shown above) were below normative population levels (19.9 ± 4.8) [13], suggesting poorer-than-average emotional well-being in this group. In contrast, PWB and FWB of both groups were above population norms (22.7 ± 5.4 and 18.5 ± 6.8 , respectively) [13]. Both survivor groups reported high levels of social well-being, with over 85% of each group reporting substantial emotional support from their families.

Item-specific analyses of the FACT-O were performed to examine specific concerns of long-term survivors. Compared to survivors with 0–1 recurrence, the most salient concerns of survivors with multiple recurrences were physical and emotional problems (Supplemental Table 1). Survivors with multiple recurrences endorsed more physical health symptoms such as lack of energy ($p=0.009$), having to spend time in bed ($p=0.006$), abdominal swelling ($p=0.002$), hair loss ($p=0.02$), problems with independent mobility ($p=0.007$), and bowel control ($p=0.045$). Emotionally, a greater percentage of survivors with multiple recurrences

reported losing hope in the fight against their illness ($p = 0.001$), worry that their condition would get worse ($p=0.047$), difficulty accepting their illness ($p=0.03$), and less enjoyment of things usually done for fun ($p=0.06$).

Both groups of survivors reported relatively high levels of spiritual well-being on the FACT spiritual scale (See [13] for population norms). Survivors with multiple recurrences reported greater trouble feeling peace of mind ($p=0.005$) and less ability to reach inside for comfort ($p=0.07$). Nevertheless, a majority of both groups (0–1: 60%; multiple: 56.3%) reported that their illness had strengthened their faith/spiritual beliefs and a majority of survivors (0–1: 75%; multiple: 81.3%) reported finding strength in their faith and/or spiritual beliefs (Table 3).

Psychosocial Adjustment

As seen in Table 4, both groups of long-term survivors reported relatively low levels of depressed mood (CES-D). Moreover, only 10% of survivors had CESD scores within the depressed range (> 16) and this did not differ by recurrence group ($p=0.54$). Survivors with 0–1 recurrence reported significantly higher levels of personal growth on the Psychological Well-being Scales (41.52 ± 6.32) than those with multiple recurrences (38.49 ± 4.6 , $p=0.018$); however, even the latter group approximated the mean of a community sample (38.4 ± 6.8) [20]. Mean levels of other facets of psychological well-being did not differ according to survivor group (p values > 0.22) and averaged across both groups were also within the range of a community sample (Purpose in Life: 39.54 ± 6.48 ; Environmental Mastery: 39.98 ± 6.46 ; Self-Acceptance: 37.89 ± 5.88) [20].

Social Relationships

Both survivor groups reported relatively strong perceived social support (SPS) with means of 84.75 (± 9.48) for those with 0–1 recurrence and 86.63 (± 8.55) for those with multiple recurrences. These compare favorably with norms from other populations which often range between 71.0 and 76.0 (e.g., [21, 22]). The two recurrence groups did not differ in levels of total social support, social attachment, or perceptions of dyadic adjustment (ADAS) ($n.s.$) (Table 4). A change in marital status since diagnosis was reported by 28.6% of long-term survivors (0–1: 32.5% vs. multiple: 18.75%, $n.s.$), including 5 widowed, 7 divorced or separated post-diagnosis, and two remarried. A change in relationship quality after diagnosis was reported by 34.0% of survivors (0–1: 38.8% vs. multiple: 21.4%, $n.s.$), with 9 reporting stronger relationships and 6 reporting poorer relationship quality (including emotional abandonment, spouse had an affair, left them, disease caused them to stay in an unhealthy relationship, etc.). Cancer survivor support group participation was reported by 71.4% of participants at some point after their diagnosis, 65.0% indicated that they lend support to other survivors; these proportions also did not differ by group. ($n.s.$)

Sexual Functioning

Sexual health changes after diagnosis and treatment were quite prevalent (0–1: 65% vs. multiple: 75%), with predominant concerns being decreased or absent desire, pain during intercourse, and reduced quality of orgasms. The majority of survivors (0–1: 58.9% vs. multiple: 62.5%) reported being in a relationship that could involve sex; however only about

half of these women (0–1: 52.2% vs. multiple: 50.0%) reported being sexually active in the last month. Almost two thirds of women in each group (0–1: 65.8% vs. multiple: 66.7%) reported that they were “not at all” or only “somewhat” satisfied with their sex life. Nevertheless, a majority of women (0–1: 81.5% vs. multiple: 75%) reported feeling close to their partners. These proportions did not differ by group (*n.s.*).

Physical Activity and Diet

With respect to exercise (IPAQ-SF), overall, 42.9% of survivors reported low levels of activity, 37.5% were moderately active, and 19.6% were highly active [18]. Exercise levels of the two survivor groups did not significantly differ, with two exceptions. First, survivors with 0–1 recurrence spent significantly fewer minutes/day (331.8 ± 209.5) and minutes/week (2322.3 ± 1466.8) sitting than those with multiple recurrences (452.3 ± 205.8 minutes/day, $p=0.036$; 3166.1 ± 1440.9 minutes/week, $p=0.036$), and second, they reported twice as many minutes of daily walking (0–1: 44.85 ± 51.41 vs. multiple: 22.19 ± 31.36 minutes, $p=0.049$) (Table 5). The majority of survivors who made changes in exercise ($n=12$) reported increasing their exercise substantially after diagnosis, including activities such as vigorous daily walking, Tai Chi, Kung Fu, stretching, yoga, and joining an exercise group. Seven survivors reported that they had decreased their exercise due to fatigue, back pain, neuropathy, and incisional hernias.

Approximately one-third (32.1%) of all participants had BMI in the overweight range and 19.6% had BMI in the obese range. Almost half of the participants had a waist-hip ratio in the high-risk range (46%) and 24% had waist-to-hip ratios in the moderate risk range. However, there were no significant differences in BMI or waist-to-hip ratio between the two groups (*n.s.*). Among all survivors, 51.8% reported changing their diet and 33.9% reported changing exercise habits since their diagnosis, with no significant differences between groups in the proportions who reported making a change (*n.s.*). Of those who made diet changes, all but four reported diet choices involving more careful food selection, such as checking food labels, eating fewer processed foods, more organic foods, vegetables, fresh fruits, lower fat, less sugar, and juicing (extraction of juice from fresh fruits and vegetables to maximize consumption of vitamins, minerals and phytonutrients). Three individuals reported persistent digestive difficulties since treatment requiring a more bland or restricted diet and one reported giving up on previous dietary changes after a recurrence.

Mind-Body or Complementary Treatments

A subset of long-term survivors used mind-body or integrative medicine treatments. Meditation was the most popular approach, with 35.2% of survivors (0–1: 40.0% vs. multiple: 21.4%) reporting having tried meditation. Of those who meditated, overall, 42.1% reported meditating 3 or more days/week, 52.6% reported meditating 1–2 days/week, and 42.1% reported meditating for more than 2 years.

Discussion

Key findings from this study indicate that QOL and psychological functioning of long-term advanced-stage ovarian cancer survivors tends to be relatively good, with mean QOL scores

commensurate with those of healthy community samples. To our knowledge, this is the first study characterizing QOL in this unique population, including contrasts between survivors with differing disease trajectories. QOL of survivors with multiple recurrences was more compromised for all domains except social and functional well-being; their decrements in emotional and physical well-being were clinically significant and their emotional well-being was below community norms. Survivors with multiple recurrences were more likely to report losing hope and worry that their condition would worsen. Even so, both groups of long-term survivors reported relatively low levels of depressed mood, and only 10% had scores in the depressed range. Both groups of women reported low levels of physical activity; almost half the long-term survivors were only minimally active, and more than half were overweight. Sexual functioning was a major concern of both groups. Social relationships were described as strong.

Ovarian cancer patients commonly report elevations in depressed mood, anxiety, and sleep disorders at the time of diagnosis [3, 23], and sustained elevations in depressed mood and sleep disorders have been observed at one year in a sizeable subset of survivors [23]. The present finding of relatively normal mood in both groups of long-term survivors is consistent with previous reports of good mental health in a majority of advanced-stage ovarian cancer survivors at 6 years post-diagnosis [7] and reported trends towards improving mental health among heterogeneous cancer patients as time since diagnosis increases [25]. The proportion of patients in our sample with clinical levels of depressive symptoms (10%) was similar to levels of depression reported in a meta-analysis of cancer patients 7 years post-diagnosis (11.6%), and was also similar to the rate of depression in healthy controls in that study (10.2%) [26]. Consistent with our finding of poorer emotional well-being among survivors with multiple recurrences, higher levels of depression have been reported in survivors with more advanced cancers, more physical symptoms, and greater loneliness [25]. Fear of recurrence or of worsening disease remains a salient concern in our long-term survivors, particularly in those who have already had a recurrence. A recent systematic review reported that a majority of long-term (> 5 years) cancer survivors experience at least moderate intensity fear of recurrence [27]. Our findings are consistent with previous reports that fear of recurrence is a common and often debilitating concern among ovarian cancer survivors [24], and one that may need health care provider attention.

The finding that QOL of long-term survivors of ovarian cancer is similar to that of the general population was surprising, and may reflect a variety of circumstances. The functional abilities of long-term survivors, particularly those with 0–1 recurrence, may be minimally compromised by cancer. These findings may also reflect support that our predominantly white, educated participants may have received from the health care system or from affiliation with resources like OCRFA.

With respect to physical activity, lower-body functional limitations have previously been reported in 52.8% of ovarian survivors [28]. Ovarian cancer survivors have reported greater difficulty performing activities including stooping, crouching, kneeling, lifting 10 pounds, walking one quarter mile, and walking up and down ten steps without rest [29], which may explain the low activity levels of our long-term survivors. Physical exercise is known to enhance physical, cognitive, and emotional functioning [30]; thus, long-term survivors may

potentially be helped by low-impact exercise or yoga programs that could address specific deficiencies, as well as by interventions that address obstacles to undertaking exercise training.

The majority of both groups of survivors reported difficulties with sexual functioning and low levels of sexual satisfaction, with sexual changes including minimal desire, pain during intercourse, and reduced quality of orgasms. These findings are consistent with previous reports of sexual health concerns among ovarian cancer survivors [4–7] and highlight the need for interventions to help long-term survivors with sexual functioning.

Both groups of survivors reported strong relationships and relatively high levels of perceived social support. Prospective longitudinal research has indicated that perceived emotional support at the time of diagnosis is related to longer survival in ovarian cancer [31]. Previous work has highlighted biological pathways by which social support may mediate survival in ovarian cancer, including angiogenesis, invasion, inflammation, cellular immunity [32, 33] and transcriptional changes in the tumor genome suggestive of less pro-inflammatory and pro-metastatic signaling [34]. Moreover, ovarian cancer survivors with high levels of psychological well-being and/or social support have lower tumor norepinephrine [35, 36], a stress hormone shown to be linked to tumor progression. These findings suggest the possibility that along with molecular and clinical determinants of long-term survival in ovarian cancer, biobehavioral factors may influence survival outcomes. Longitudinal research on trajectories of long-term survivors will help to further elucidate these issues.

Limitations

Although only a small proportion of those approached at study sites declined, some long-term survivors were not approached because of poor health. We do not know how many eligible long-term survivors did not respond to postings about the study on the OCRFA website. It is possible that only the healthiest long-term survivors volunteered, thus under-representing the level of compromise in long-term survivors as a whole. Additionally, it is possible that high levels of QOL may reflect the demographics of populations seeking care at academic medical centers or seeking support from OCRFA. We also did not assess health insurance status, which may have been able to shed light on the availability of health care to participants. As the vast majority of patients were white and educated, the demographic homogeneity may limit generalizability of findings. Because there were only 16 individuals in the multiple recurrence group, the inferences that can be drawn from their data are limited. Number of recurrences ranged from 2 to 8. It is also possible that individuals with multiple recurrences may have had additional debility due to being on treatment or maintenance chemotherapy currently or in the recent past; this may have enhanced between group differences. Because we did not collect data on time since last treatment, we were not able to examine this as a potential covariate. Thus, there may be substantial heterogeneity in the multiple recurrence group and the present data may not fully capture the impact of multiple recurrences on QOL. We also did not collect data on use of antidepressants or anti-anxiety medications, and thus do not know how such drugs may have influenced findings. The cross-sectional nature of this study does not shed light on the trajectory of QOL of survivors over time. As we did not collect data on cognitive functioning and neurotoxicity,

findings may underestimate these serious treatment side effects. Information on cancer stage, treatment, and recurrences was obtained by self-report and was not objectively verified. Although there has been some indication of concordance of self-reported and medical chart information among cancer patients [37], these studies have not examined self-reports of long-term survivors, and thus this information should be treated with caution. Finally, since long-term survival in advanced stage ovarian cancer is extremely rare, the sample size is relatively small.

Conclusions and Clinical Implications

Overall, advanced-stage ovarian cancer survivors surviving at least 8.5 years report good QOL and psychological adjustment. QOL of survivors with multiple recurrences is somewhat impaired compared to those with 0–1 recurrences. In light of these findings and the recent IOM recommendations, health care providers should be sensitive to psychosocial issues such as distress, sexual health difficulties, physical impairments, and low levels of physical activity among survivors of advanced-stage ovarian cancer from diagnosis through long-term survival. Interventions to improve QOL and to address limitations in physical activity should also be considered to help survivors with these issues.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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Research Highlights

- QOL of long-term advanced-stage ovarian cancer survivors is relatively good
- Survivors with multiple recurrences reported more compromised QOL
- Low levels of exercise were reported by 43% of long-term survivors
- Among these survivors, 52% were overweight or obese
- Sexual health concerns are common in long-term survivors

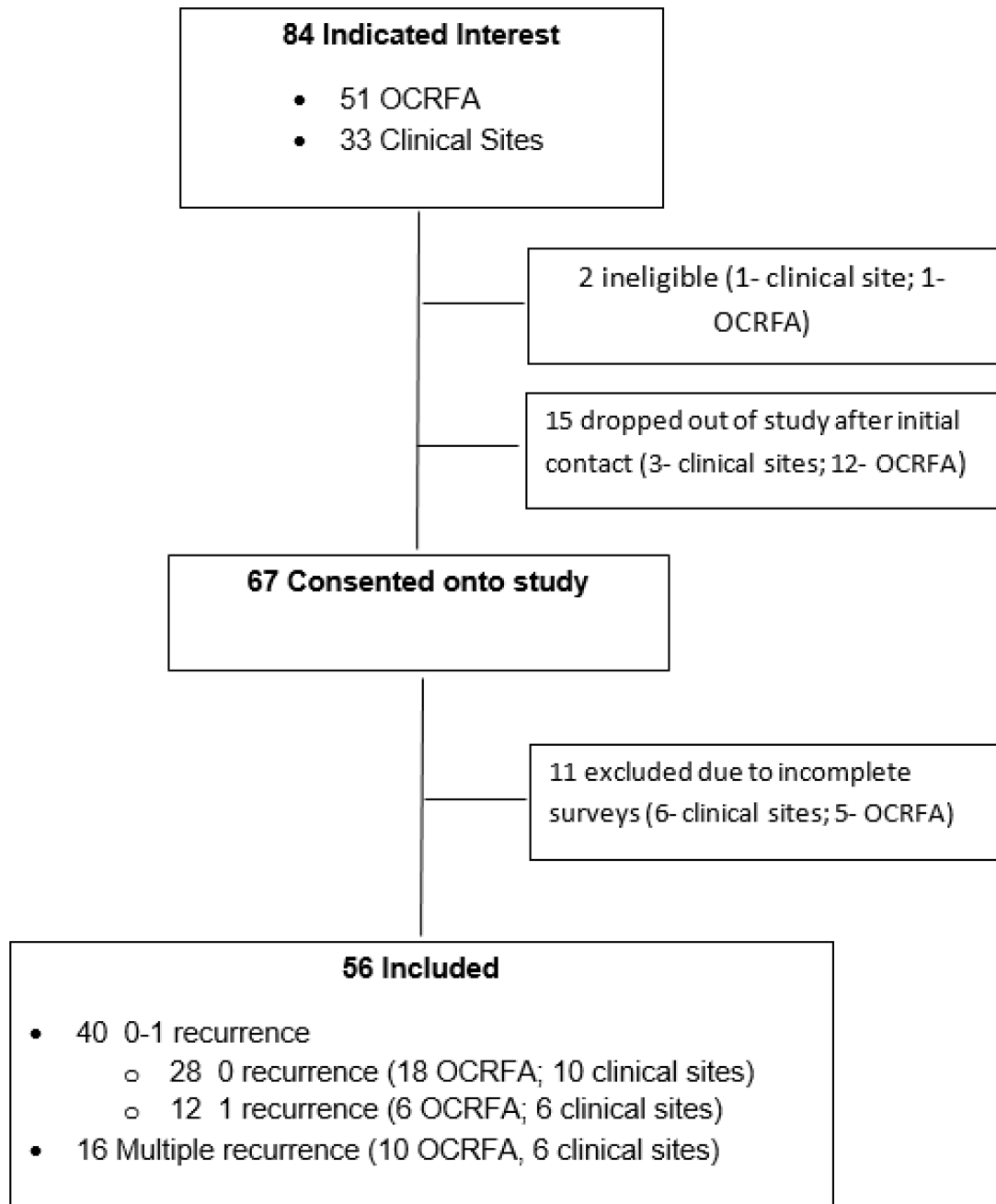


Figure 1.
Consort (Consolidated Standards of Reporting Trials) Diagram displaying patient inclusion.

Table 1

Demographics characteristics of long-term survivors by recurrence status

Characteristic	0 – 1 recurrences (n = 40)		Multiple recurrences (n = 16)		Total (n = 56)		p-value
	N	%	N	%	N	%	
Age at diagnosis							
N	40		16		56		0.394
Mean (SD)	50.5 (9.9)		48.5 (11.1)		49.9 (10.2)		
Min - Max	22.7 – 68.5		21.8 – 65.0		21.8 – 68.5		
Age at survey							
N	40		16		56		0.079
Mean (SD)	66.9 (9.2)		62.2 (10.0)		65.5 (9.6)		
Min - Max	44.1 – 85.9		40.2 – 77.1		40.2 – 85.9		
BMI							
Underweight	1	2.5	0	0	1	1.8	0.816
Normal	17	42.5	9	56.3	26	46.4	
Overweight	14	35.0	4	25	18	32.1	
Obese	8	20.0	3	18.8	11	19.6	
Waist to hip ratio							
Low risk	10	28.6	5	33.3	15	30.0	0.926
Moderate risk	9	25.7	3	20	12	24.0	
High risk	13	45.7	7	46.7	23	46.0	
Racial background							
Black	1	2.5	0	0	1	1.8	0.644
Asian	0	0.0	0	0	0	0.0	
Other	1	2.5	0	0	1	1.8	
White	38	95	15	93.8	53	94.6	
White/Native American	0	0	1	6.3	1	1.8	
White/Other	0	0.0	0	0	0	0.0	0.557
Ethnic background							

Characteristic	0-1 recurrences (n = 40)		Multiple recurrences (n = 16)		Total (n = 56)		p-value
	N	%	N	%	N	%	
Hispanic	3	7.7	0	0	3	5.7	
Not Hispanic	36	92.3	14	100.0	50	94.3	
Education completed							0.021
Less than high school	1	2.5	0	0	1	1.8	
High school/GED	2	5.0	0	0	2	3.6	
Some college/Technical school	18	45.0	3	18.8	21	37.5	
Four-year college or university	7	17.5	9	56.3	16	28.6	
Graduate school/Professional school	12	30.0	3	18.8	15	26.8	
Other	0	0.0	1	6.3	1	1.8	
Employment status:							0.276
Employed outside of the home	12	30.0	6	37.5	18	32.1	
Homemaker	5	12.5	3	18.8	8	14.3	
Retired	23	57.5	6	37.5	29	51.8	
Unemployed - looking for work	0	0.0	0	0	0	0.0	
Unemployed - due to disability	0	0.0	1	6.3	1	1.8	
Annual total household income							0.691
Less than \$25,000	1	2.7	0	0	1	1.9	
\$25,001 - \$50,000	14	37.8	3	20	17	32.7	
\$50,001 - \$75,000	7	18.9	4	26.7	11	21.2	
\$75,001 - \$100,000	5	13.5	2	13.3	7	13.5	
Greater than \$100,000	10	27.0	6	40	16	30.8	
Current marital status							0.495
Never married	0	0.0	1	6.3	1	1.8	
Married	23	57.5	12	75	35	62.5	
Living with partner	2	5.0	0	0	2	3.6	
Separated	1	2.5	0	0	1	1.8	
Divorced	10	25.0	2	12.5	12	21.4	
Widowed	4	10.0	1	6.3	5	8.9	

Characteristic	0-1 recurrences (n = 40)		Multiple recurrences (n = 16)		Total (n = 56)		p-value
	N	%	N	%	N	%	
Given birth to a live infant							>0.999
Yes	34	85.0	14	87.5	48	85.7	
No	6	15.0	2	12.5	8	14.3	
Smoking status							>0.999
Never	19	47.5	7	46.7	26	47.3	
Current	2	5.0	1	6.7	3	5.5	
Former	19	47.5	7	46.7	26	47.3	
Current Smoking							0.636
Everyday	1	4.8	0	0	1	3.5	
Some days	1	4.8	1	12.5	2	6.9	
Not at all	19	90.5	7	87.5	26	89.7	

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Table 2

Clinical Characteristics by Recurrence Status

Characteristic	0-1 (n = 40)		Multiple (n= 16)		Total (N = 56)		P- value
	N	%	N	%	N	%	
Time from diagnosis to survey							0.355
N	40		16		56		
Mean (SD)	16.3 (6.2)		13.7 (2.6)		15.6 (5.5)		
Min – Max	8.8 – 33.3		9.2 – 18.3		8.8 – 33.3		
What stage was your ovarian cancer at the time of diagnosis?							0.274
Stage I or II	0	0	0	0	0		
Stage III	37	92.5	13	81.3	50	89.3	
Stage IV	2	5.0	3	18.8	5	8.9	
not sure	1	2.5	0	0	1	1.8	
Grade							0.889
High grade	14	35.0	4	26.7	18	32.7	
Low grade	3	7.5	1	6.7	4	7.3	
not sure	23	57.5	10	66.7	33	60.0	
Histology							0.741
Serous	27	67.5	10	66.7	37	67.3	
Non-Serous	1	2.5	1	6.7	2	3.6	
not sure	12	30.0	4	26.7	16	29.1	
Current Treatment Status							0.005
Currently no treatment	34	85.0	7	43.8	41	73.2	
On treatment	5	12.5	7	43.8	12	21.4	
Missing	1	2.5	2	12.5	3	5.4	
Neoadjuvant chemotherapy							>0.999
Yes	2	5.0	0	0	2	3.6	
No	38	95.0	16	100	54	96.4	
Radiation at any point since diagnosis							0.395
Yes	4	10.0	3	18.8	7	12.5	

Characteristic	0-1 (n = 40)		Multiple (n = 16)		Total (N = 56)		P-value
	N	%	N	%	N	%	
No	36	90.0	13	81.3	49	87.5	0.015
Second surgery for ovarian cancer							
Yes	12	30.0	11	68.8	23	41.1	
No	28	70.0	5	31.3	33	58.9	0.475
BRCA testing							
Yes	30	75.0	14	87.5	44	78.6	
No	10	25.0	2	12.5	12	21.4	0.391
If yes, do you have a BRCA mutation?							
Yes	13	43.3	5	38.5	18	41.9	
No	16	53.3	6	46.2	22	51.2	
I'm not sure	1	3.3	2	15.4	3	7.0	<0.001
Recurrence							
Yes	12	30.0	16	100	28	50.0	
No	28	70.0	0	0	28	50.0	0.741
Other cancer							
Yes	10	25.0	5	31.3	15	26.8	
No	30	75.0	11	68.8	41	73.2	>0.999
Change in diet since diagnosis?							
Yes	21	52.5	8	50.0	29	51.8	
No	19	47.5	8	50.0	27	48.2	>0.999
Change in exercise habits since diagnosis?							
Yes	14	35.0	5	31.3	19	33.9	
No	26	65.0	11	68.8	37	66.1	

Table 3

Fact-O Total and Subscale scores by recurrence status

Score	Recurrences	N	Mean	SD	Med	Min	Max	p-value
Physical Well Being Subscale Score	0 – 1	40	25.43	2.72	26	15	28	0.016
	Multiple	16	23.25	3.49	23	16	28	
Social Family Well Being Subscale Score	0 – 1	39	19.77	4.38	20	7	28	0.730
	Multiple	16	18.88	4.84	20.5	6	24	
Emotional Well Being Subscale Score	0 – 1	39	20.69	2.92	21	11	24	0.019
	Multiple	16	17.88	4.44	20	9	24	
Functional Well Being Subscale Score	0 – 1	40	23.00	4.22	24	12	28	0.184
	Multiple	16	20.81	5.60	22.5	11	28	
Additional Concerns Subscale Score	0 – 1	40	36.30	4.54	38	22	44	0.009
	Multiple	16	32.75	5.25	33	20	42	
Total Fact-G score	0 – 1	39	89.05	10.80	90	62	104	0.031
	Multiple	16	80.81	13.95	85	56	101	
Total Fact-O	0 – 1	39	125.38	14.51	129	84	148	0.017
	Multiple	16	113.56	17.35	115.5	79	143	
Trial Outcome Index Score	0 – 1	40	84.72	9.75	87.5	56	99	0.013
	Multiple	16	76.81	11.83	77.5	52	97	
FACT Spiritual Well-being Peace Scale Score	0 – 1	40	26.35	4.96	26.5	15	32	0.603
	Multiple	16	25.31	6.06	26.5	9	32	
FACT Spiritual Well-being Faith Scale Score	0 – 1	40	12.03	4.40	13	2	16	0.941
	Multiple	16	12.13	3.77	12	5	16	
FACT Spiritual Well-being Total Score	0 – 1	40	38.38	8.13	39.64	19	48	0.669
	Multiple	16	37.44	8.80	37	19	48	

Table 4

Psychosocial Scales by Recurrence Status

Item	Recurrences	N	Mean	SD	Med	Min	Max	p-value
Center for Epidemiologic Studies Depression Scale	0 – 1	40	7.17	6.46	5	0	22	0.542
	Multiple	16	7.38	2.58	8.5	2	10	
Psychological Well-Being: Environmental Mastery	0 – 1	40	40.58	6.14	40	29	49	0.344
	Multiple	16	38.50	7.18	39	21	47	
Psychological Well-Being: Personal Growth	0 – 1	40	41.52	6.32	43	23	48	0.018
	Multiple	16	38.49	4.6	38	30	44	
Psychological Well-Being: Purpose in Life	0 – 1	40	40.10	6.85	41.5	21	49	0.217
	Multiple	16	38.13	5.38	38	27	46	
Psychological Well-Being: Self-Acceptance	0 – 1	40	37.65	6.34	40	22	46	0.920
	Multiple	16	38.50	4.68	39.5	28	44	
Social Provisions Scale: Total Social Support	0 – 1	40	84.75	9.48	87	61	96	0.531
	Multiple	16	86.63	8.55	89.5	69	96	
Social Provisions Scale: Attachment subscale	0 – 1	40	14.15	2.23	15	9	16	0.765
	Multiple	16	14.44	1.90	15.5	11	16	
ADAS Total Score	0 – 1	27	23.96	5.91	25	7	33	0.820
	Multiple	13	23.54	4.48	23	15	31	

Table 5

International Physical Activity Questionnaire by Recurrence status

IPAQ item	Status	N	Mean	SD	Min	Max	p-value
Vigorous Activity Minutes Per Day	0-1	40	38.63	69.64	0	300	0.338
	Multiple	16	23.44	61.23	0	240	
Vigorous Activity minutes last 7 days	0-1	38	114.47	187.76	0	720	0.503
	Multiple	14	123.21	322.48	0	1200	
Moderate Activity Minutes Per Day	0-1	40	44.00	68.73	0	300	0.921
	Multiple	16	47.50	76.29	0	240	
Moderate Activity minutes last 7 days	0-1	38	181.45	281.95	0	1200	0.965
	Multiple	14	213.57	351.56	0	1200	
Walking Minutes Per Day	0-1	40	44.85	51.41	0	240	0.049
	Multiple	16	22.19	31.36	0	120	
Walking Minutes Last 7 Days	0-1	38	232.45	312.59	0	1680	0.153
	Multiple	14	133.57	216.5	0	840	
Sitting Minutes Per Day	0-1	34	331.76	209.54	120	1200	0.036
	Multiple	13	452.31	205.84	180	900	
Sitting Minutes Last 7 Days	0-1	34	2322.35	1466.76	840	8400	0.036
	Multiple	13	3166.15	1440.87	1260	6300	
Vigorous and Moderate Minutes combined (last 7 days)	0-1	38	295.92	383.81	0	1200	0.627
	Multiple	14	336.79	649.3	0	2400	
Total minutes of activity (last 7 days)	0-1	38	528.37	637.04	0	2840	0.508
	Multiple	14	470.36	697.24	0	2580	
Met Score for walking	0-1	38	767.08	1031.54	0	5544	0.153
	Multiple	14	440.79	714.44	0	2772	
Met Score for moderate activity	0-1	38	725.79	1127.82	0	4800	0.965
	Multiple	14	854.29	1406.24	0	4800	
Met Score for vigorous activity	0-1	38	915.79	1502.05	0	5760	0.503
	Multiple	14	985.71	2579.81	0	9600	
Total Physical Activity Score	0-1	38	2408.66	2911.29	0	11944	0.402
	Multiple	14	2280.79	3957.3	0	14994	

IPAQ item	Status	N	Mean	SD	Min	Max	p-value
Low Activity ^a	0-1	15	40.48%				0.391
Total both groups	Multiple	9	56.25%				
Moderately Activity	0-1	17	40.48%				
Total both groups	Multiple	4	25.00%				
High Activity	0-1	8	19.04%				
Total both groups	Multiple	3	18.75%				

^aResponses are categorized into three levels of physical activity: **low** (< moderate); **moderate** (3 or more days/week of vigorous activity of at least 20 minutes/day or 5 or more days/week of moderate activity, or 30 minutes/day walking) and **high** (vigorous activity on at least 3 days or 7 days/week of any combination of walking, moderate or vigorous intensity activities.) [14]