

PDF hosted at the Radboud Repository of the Radboud University Nijmegen

The following full text is a publisher's version.

For additional information about this publication click this link.

<http://hdl.handle.net/2066/48419>

Please be advised that this information was generated on 2021-01-17 and may be subject to change.

Quality-of-Life Effects of Prophylactic Salpingo-Oophorectomy Versus Gynecologic Screening Among Women at Increased Risk of Hereditary Ovarian Cancer

Joanna B. Madalinska, Judith Hollenstein, Eveline Bleiker, Marc van Beurden, Heiddis B. Valdimarsdottir, Leon F. Massuger, Katja N. Gaarenstroom, Marian J.E. Mourits, René H.M. Verheijen, Eleonora B.L. van Dorst, Hans van der Putten, Ko van der Velden, Henk Boonstra, and Neil K. Aaronson

From the Division of Psychosocial Research and Epidemiology, Department of Gynecology, The Netherlands Cancer Institute; Department of Gynecology, Vrije University Medical Center; Department of Gynecology, Academic Medical Center, Amsterdam; Department of Gynecology, University Medical Center Nijmegen, Nijmegen; Department of Gynecology, Leiden University Medical Center, Leiden; Department of Gynecology, Academic Hospital Groningen, Groningen; Department of Gynecology, University Medical Center Utrecht, Utrecht; Department of Gynecology, Academic Hospital Maastricht, Maastricht, the Netherlands; and Department of Psychiatry, Mount Sinai School of Medicine, New York, NY.

Submitted October 20, 2004; accepted April 20, 2005.

Supported by Grant No. NKI 2001-2382 from the Dutch Cancer Society.

Authors' disclosures of potential conflicts of interest are found at the end of this article.

Address reprint requests to Neil K. Aaronson, PhD, The Netherlands Cancer Institute/Antoni van Leeuwenhoek Hospital, Division of Psychosocial Research and Epidemiology, Plesmanlaan 121, 1066 CX Amsterdam, the Netherlands; e-mail: n.aaronson@nki.nl.

© 2005 by American Society of Clinical Oncology

0732-183X/05/2328-6890/\$20.00

DOI: 10.1200/JCO.2005.02.626

A B S T R A C T

Purpose

Recommendations for women at high risk of ovarian cancer include periodic gynecologic screening (GS) and prophylactic bilateral salpingo-oophorectomy (PBSO). The aim of the current study was to determine the quality-of-life (QOL) effects of PBSO versus GS.

Patients and Methods

Questionnaire data were obtained from 846 high-risk women who had participated in this nationwide, cross-sectional, observational study. Forty-four percent of the women had undergone PBSO, and 56% had opted for GS. Topics addressed by the questionnaire included generic QOL, cancer-specific distress, endocrine symptoms, and sexual functioning.

Results

No statistically significant between-group differences were observed in generic QOL (Short Form-36), with women in both the PBSO and GS groups scoring similarly to the general population. Compared with GS, PBSO was associated with fewer breast and ovarian cancer worries ($P < .001$) and more favorable cancer risk perception ($P < .05$). However, the PBSO group reported significantly more endocrine symptoms ($P < .001$) and worse sexual functioning ($P < .05$) than the GS group. Eighty-six percent of women would choose PBSO again, and 63% would recommend it to a friend with familial risk of ovarian cancer.

Conclusion

PBSO had no measurable adverse impact on generic QOL of high-risk women. The favorable effects of PBSO in terms of reduced cancer worries and low perceived cancer risk need to be weighed against the increase in endocrine and sexual symptoms. Balanced information will help clinicians and high-risk women to make informed decisions about the optimal preventive health strategy.

J Clin Oncol 23:6890-6898. © 2005 by American Society of Clinical Oncology

INTRODUCTION

Ovarian cancer is one of the most common and lethal of gynecologic malignancies. In the Netherlands, the average age-adjusted incidence and mortality rates are 13.1 and 9.6 per 100,000 women, respectively, which are comparable to the rates observed in the United States.^{1,2} A family history of ovarian cancer is considered to be one of the stron-

gest predictors of developing the disease, and it is estimated that 5% to 10% of all ovarian cancer patients have a hereditary basis.^{3,4} Female carriers of a *BRCA1* gene mutation have a lifetime ovarian cancer risk in the range of 39% to 54%.^{5,6} Women with a *BRCA2* mutation have a lower ovarian cancer risk (11% to 23%), but this is still approximately 10-fold greater than the risk of women in the general population.^{5,6}

Principal preventive health strategies for women at increased risk of ovarian cancer include periodic gynecologic screening (GS) and prophylactic bilateral salpingo-oophorectomy (PBSO), which are aimed at early cancer detection and cancer risk reduction, respectively. Although annual GS is offered as a basic surveillance strategy to high-risk women, its efficacy has yet to be established.⁷ Current techniques, such as transvaginal sonography and CA-125 serology yield a significant number of false-positive or false-negative results, leading either to unnecessary medical investigations or to undetected early-stage malignancies. Because early ovarian cancer is asymptomatic and the available techniques have not been demonstrated to be effective for early diagnosis in the general population,⁸ the majority of diagnosed ovarian cancers are characterized by advanced stages and, therefore, by a poor prognosis.⁹

In view of the uncertainty surrounding screening procedures, high-risk women may opt for surgical removal of their ovaries and fallopian tubes. PBSO reduces ovarian cancer risk in *BRCA1/2* mutation carriers by 96% and breast cancer risk by 53%.^{10,11} However, PBSO does not eliminate the risk of ovarian cancer entirely because 1% to 2% of women may develop peritoneal carcinoma.¹⁰⁻¹² Adverse effects associated with prophylactic surgery in premenopausal women are loss of fertility and immediate onset of menopause as a result of estrogen deprivation, including vasomotor symptoms and possible sexual dysfunction.^{13,14} Estrogen deprivation may also lead to higher risk of developing osteoporosis.¹⁵ To relieve climacteric symptoms, hormone replacement therapy (HRT) is often prescribed.¹⁶ However, the effectiveness of HRT in combating symptoms associated with surgically induced menopause has not yet been established.

Thus far, only four studies have explicitly focused on quality-of-life (QOL) effects associated with PBSO or PBSO versus GS.¹⁷⁻²⁰ Several studies¹⁷⁻¹⁹ have reported beneficial effects of PBSO on cancer-specific distress (eg, cancer worries and anxiety) and perceived cancer risk but adverse effects on sexual functioning and vasomotor symptoms. In these studies, generic QOL was not affected by prophylactic surgery, with oophorectomized women reporting similar levels of QOL as women in the general population.^{17,18} The only study that has compared the QOL effects of PBSO and GS yielded somewhat conflicting results.²⁰ Oophorectomized women reported significantly worse generic QOL than women in the GS group; however, no comparisons with the general population were provided. Additionally, PBSO was not found to relieve cancer-specific distress or to worsen sexual functioning. Although this latter study was the first to provide a comparison of psychosocial effects of PBSO and GS, its results may not be generalizable to the entire population of high-risk women. It was a single-center study with a small sample size (PBSO, $n = 29$; GS, $n = 28$), and not all statistical analyses controlled for possible confounding medical variables (eg, DNA status and history of

breast cancer). In this report, we present the results of a nationwide, multicenter, cross-sectional, observational study that was conducted to determine possible differences in the generic and condition-specific QOL effects of PBSO versus GS.

PATIENTS AND METHODS

Sample and Procedures

Study participants were recruited from the gynecology departments of eight hospitals in the Netherlands. Women were eligible for enrollment if they were between 30 and 75 years of age, came from a hereditary breast or ovarian cancer family, and had sought advice from a gynecologist on preventive measures at one of the eight participating gynecology clinics between 1996 and 2001. Patients were excluded from participation if they had undergone oophorectomy because of any suspicious changes in the ovaries as detected by medical examination, including both benign and malignant conditions; if oophorectomy was performed as adjuvant treatment for breast cancer; or if they had terminal cancer or any other severe medical comorbidity.

Eligible women received an invitation letter by mail, an informed consent form, a questionnaire, and a postage-paid return envelope. In case of nonresponse within 2 weeks, systematic reminders by mail and telephone were used. Patients were classified as nonrespondents if they actively declined to participate by mail or telephone or if they could not be reached after multiple attempts. Age and the type of ovarian cancer prevention strategy used (PBSO *v* GS) were the only available data that could be registered for nonrespondents. The study was approved by the institutional review boards of all participating hospitals.

Measure: Generic QOL

To assess generic QOL, four of the eight subscales of the Short Form-36 (SF-36) Health Survey^{21,22} (general health perceptions, vitality, role limitations caused by emotional problems, and general mental health) and the global QOL item of the European Organisation for Research and Treatment of Cancer Quality of Life Questionnaire C30²³ were used. All raw scale scores were linearly converted to a 0 to 100 scale, with higher scores indicating higher levels of QOL.^{24,25} Cronbach's α coefficients in the present sample for the four SF-36 scales ranged from .80 to .86.

Measure: Condition-Specific QOL

Condition-specific QOL included measures of cancer-specific distress (intrusive thoughts, cancer worries, and anxiety), cancer risk perception, endocrine symptoms, and sexual functioning.

Cancer-specific distress: Intrusive thoughts, cancer worries, and postoperative anxiety. The seven-item intrusion subscale of the Impact of Events Scale (IES)^{26,27} measures the frequency of intrusive thoughts experienced because of a specific stressor, which was defined in the present study as an increased risk of developing breast or ovarian cancer. A higher sum score (range, 0 to 35) corresponds to more distress (Cronbach's $\alpha = .90$). The recommended cutoff sum score for identifying persons likely to meet criteria for post-traumatic stress syndrome (PTSD) is 20.²⁸

Five Likert-type items, adapted from Lerman,²⁹ were used to assess worries about breast and ovarian cancer. These included the frequency of ovarian and breast cancer worries (two items), the impact of cancer worries on mood and daily functioning (two items), and the frequency of worries about the possible cancer risk

in family members (1 = rarely or never, 2 = sometimes, 3 = often, 4 = all the time). These five items were summed to create a cancer worry scale (possible range, 5 to 20), with higher scores representing more frequent worries in the last 4 weeks (Cronbach's $\alpha = .70$). Additionally, women in the PBSO group were asked to rate the extent to which PBSO reduced their anxiety about developing ovarian and breast cancer, with response choices varying on a 4-point scale from not at all to very much.

Self-perceived cancer risk. Two items adapted from previous studies^{30,31} assessed patients' current perceptions of their breast cancer risk. Women were asked to rate their self-perceived risk on a scale 0% to 100%, where 0 corresponded to no risk at all and 100 corresponded to being certain about developing cancer in the future. Women in the PBSO group were also requested to estimate (retrospectively) their presurgery risk of developing breast cancer.

Endocrine symptoms and sexual functioning. The Functional Assessment of Cancer Therapy–Endocrine Symptom, an 18-item endocrine symptom scale, was used to assess menopausal symptoms.³² Occurrence of each symptom in the last 4 weeks was scored on a 5-point Likert-type scale, ranging from not at all to very much. Item scores were summed to obtain a scale score (range, 0 to 72), with lower values indicating more menopausal symptoms. The Cronbach's α in the present study was .81.

The Sexual Activity Questionnaire (SAQ)³³ was used to measure sexual functioning. The SAQ consists of the following three scales: pleasure (six items on desire, enjoyment, satisfaction, and current frequency of activities), discomfort (two items on vaginal dryness and pain and discomfort during penetration), and habit (frequency of sexual activity compared with the usual level). Lower scores represent poorer sexual functioning. In the present sample, Cronbach's α coefficients for pleasure and discomfort scales were .82 and .77, respectively. The SAQ was introduced during the course of the study and, thus, was administered to only a subset of women ($n = 513$) from five study centers.

Measure: Satisfaction With Preventive Health Strategies

A series of single items was used to assess the level of satisfaction with or regrets about the decision to undergo PBSO or GS. On a 5-point scale, varying from completely disagree to completely agree, women were asked to indicate their level of agreement or disagreement with the following statements: (1) I am satisfied with the decision I have made, and (2) I have regrets about the decision I have made. Women who chose agree or completely agree as their response to the first or second statement were considered as being satisfied with their decision on the preventive health option or as having regrets about it, respectively. Additionally, women were asked two questions about whether or not they would choose to undergo the same preventive health strategy again and about which preventive option they would recommend to a friend in a similar situation.

Medical and Sociodemographic Data

Medical data were obtained from two sources (a questionnaire and hospital medical records). In case of discrepancies between self-reported and medical record data, the latter were considered as the primary information source.

The questionnaire contained a series of questions on reproductive history, personal history of cancer and recent treatment for cancer, prevalence of ovarian and breast cancer among relatives, prophylactic ovarian and breast surgery, and use of HRT. Menopausal status was determined through a series of questions

on menstrual history and symptoms during the 6 months preceding PBSO or at the present moment for women who had opted for GS. Premenopause was defined as regular menstrual periods, perimenopause was defined as irregular periods, and postmenopause was defined as complete cessation of menstrual periods for at least 1 year. Women who had had PBSO were classified as postmenopausal. Additionally, clinical variables, such as DNA status, type of prophylactic ovarian or breast surgery, possible use of HRT, history of (breast) cancer, its stage at diagnosis, and cancer treatment, were retrieved from the medical records. Sociodemographic variables (age, marital status, education, and employment) were obtained from the questionnaire.

Statistical Analysis

Descriptive statistics (frequencies, means, and standard deviations [SD]) were generated to characterize the sample in terms of sociodemographics and medical variables. Student's *t* tests and χ^2 tests were used to explore potential differences in the background characteristics of women who had undergone PBSO and women who had opted for GS.

To test for the statistical significance of group differences in generic and condition-specific QOL, we used one-way analysis of covariance (ANCOVA) controlling for possible confounders (age, *BRCA1/2* status, parity, history of breast cancer, and prophylactic mastectomy [PM]). To examine the magnitude of differences between the PBSO and GS groups, effect sizes based on differences between mean scores divided by the pooled SD were calculated. In accordance with Cohen,³⁴ effect sizes of 0.20, 0.50, and 0.80 were considered small, medium, and large, respectively. Using ANCOVA, we also investigated possible differences in the SF-36 mean scale scores between the participating high-risk women and women of similar age from the general Dutch population. The SF-36 general population normative data were based on the sample reported earlier by Aaronson et al.²²

A multivariate logistic regression analysis was conducted to investigate the effects of the type of ovarian cancer prevention (GS *v* PBSO) on the odds of the presence of cancer worries, when controlling for the potential confounders. Separate items of the cancer worry scale were dichotomized (eg, worried *v* not worried), with the original categories (sometimes, often, and all the time) describing the frequency of worries and their impact on mood and functioning collapsed into one category (worried). The purpose of this analysis was to determine which specific aspects of distress contributed the most to a PBSO-GS difference.

Within the PBSO group, we also examined whether menopausal status (premenopausal *v* postmenopausal) at the time of ovarian surgery had a significant impact on the current levels of QOL. Additionally, in an ANCOVA model, we controlled for the time since surgery and current HRT use.

All statistical analyses were carried out using SPSS version 11.5.0 (SPSS Inc, Chicago, IL). Because of multiple testing, the significance level was set at $P \leq .01$. *P* values between .01 and .05 were considered to be marginally significant. All statistical tests were two sided.

RESULTS

Study Sample

On the basis of the hospital census data (Fig 1), we identified 1,205 high-risk patients who were potentially

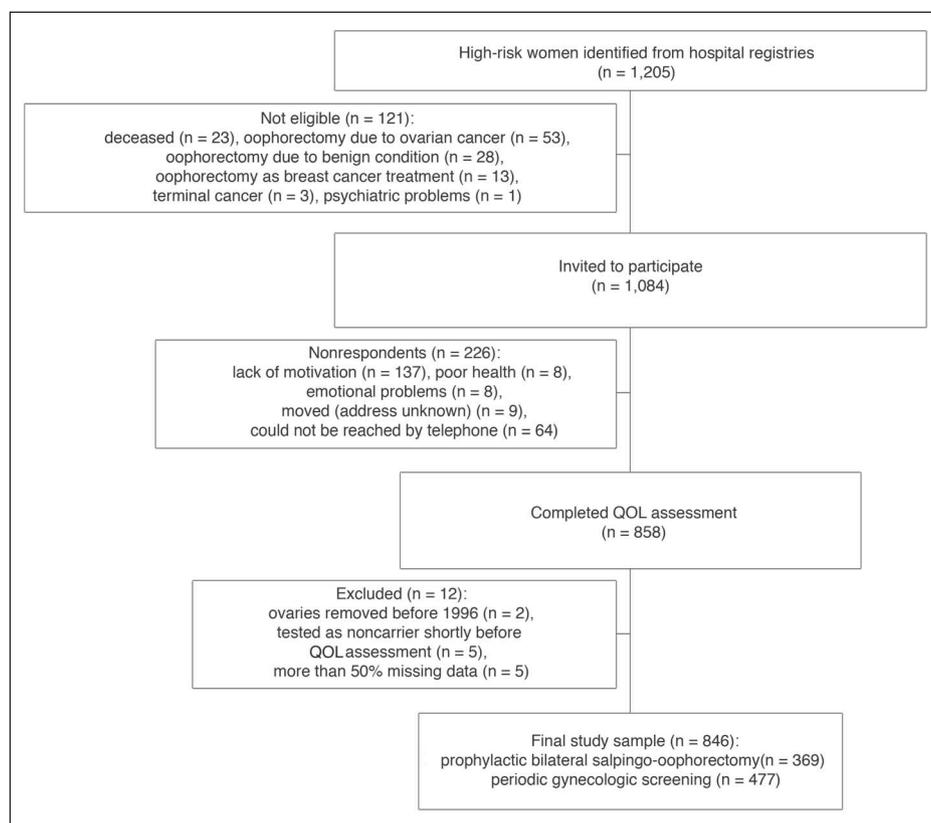


Fig 1. Flow chart for recruitment of women at increased risk for ovarian cancer in a study to evaluate quality-of-life (QOL) effects of prophylactic oophorectomy versus periodic gynecologic screening.

eligible for participation in the study. After an additional medical record audit, 121 women were excluded because of oophorectomy carried out as treatment for benign or malignant conditions ($n = 94$), death ($n = 23$), terminal cancer ($n = 3$), or severe psychiatric problems ($n = 1$). In total, 858 (79%) of 1,084 eligible women returned the questionnaires. The main reasons for nonparticipation were lack of motivation ($n = 137$), poor health ($n = 8$), and emotional problems ($n = 8$). The data of 12 women had to be excluded; five of these women reported that the questionnaire was not applicable to their present situation because their cancer risk was found not to be increased according to DNA testing, five women had a high percentage ($> 50\%$) of missing values, and two women reported having undergone an oophorectomy before 1996. There were no statistically significant differences between the respondents and nonrespondents regarding the type of ovarian cancer prevention and mean age (data not shown).

The final study sample ($n = 846$) consisted of 369 women (44%) who had undergone PBSO and 477 women (56%) who had opted for periodic GS (pelvic examination, transvaginal sonography, and CA-125 serology). Among *BRCA1/2* mutation carriers ($n = 368$), 265 women (72%) opted for PBSO, and 103 women (28%) opted for GS.

The demographic and clinical characteristics of the sample are listed in Table 1. Compared with the women in

the GS group, the women in the PBSO group were significantly older and were significantly more likely to have been diagnosed with breast cancer, to be *BRCA1/2* mutation carriers, and to have undergone (unilateral or bilateral) PM (all $P < .001$). Women with less education and women having at least one child were also more likely to undergo PBSO, although these associations only reached marginal levels of statistical significance (all $P < .05$). After PBSO, slightly more than one third of women had used HRT.

Generic and Condition-Specific QOL

Table 2 lists mean scores and SDs of the QOL measures for the PBSO and GS groups. Overall, the study respondents exhibited high levels of generic QOL as assessed by the SF-36, and no significant differences were found between the PBSO and GS groups. The SF-36 scores of both the PBSO and GS groups were, on average, not significantly different from those of similarly aged women from the general population.

There were no significant group differences in mean levels of intrusive thoughts about cancer, and similar percentages of the PBSO and GS groups (9% to 10%) reported intrusive thoughts (sum score ≥ 20) severe enough to indicate the possible presence of PTSD (Table 2). However, women who had undergone PBSO reported significantly fewer cancer worries (scale mean, 7.0; range, 5 to 14) than

Table 1. Demographic and Medical Characteristics of the Sample by Preventive Health Strategies for Ovarian Cancer: PBSO Versus GS

Characteristic	% of Patients		P
	PBSO (n = 369)	GS (n = 477)	
Age, years			< .001
Mean	49	47	< .001
SD	8	9	
30-35	1.6	11.2	
36-45	33.9	37.9	
46-55	45.5	33.3	
> 55	19.0	17.6	
Marital status			.520
Married/cohabitating	83.6	81.8	
Unmarried/without partner	16.4	18.2	
Educational level			.019
Primary school/lower level high school	22.2	18.3	
Middle level high school	49.3	44.1	
Advanced vocational/university	28.5	37.6	
Parity			.003
Null parity	11.9	19.5	
At least one child	88.1	80.5	
Current menopausal state			
Premenopausal		62.1	
Peri/postmenopausal	100*	37.9	
Menopausal state before PBSO			
Premenopausal	38.2	—	
Peri/postmenopausal	61.8	—	
Previous or present use of HRT	36.9†	5.9	< .001
DNA status			< .001
BRCA1/2 carrier	71.8	21.6	
Nonconclusive	13.3	24.9	
Not tested/other	14.9	53.5	
History of breast cancer, yes	49.3	34.0	< .001
Current use of tamoxifen, yes	5.1	3.4	.194
Prophylactic mastectomy, yes	45.5	13.2	< .001
Self-reported time since first visit to gynecologist because of high-risk status, years			.202
Mean	4.1	4.3	
SD	2.4	3.2	
Median	4.0	4.0	
Type of prophylactic oophorectomy			
Laparoscopy	80.1	—	
Laparotomy	19.9	—	
Time since PBSO, years			
Mean	2.8	—	
SD	1.9	—	
Median	2.0	—	

Abbreviations: PBSO, prophylactic bilateral salpingo-oophorectomy; GS, gynecologic screening; SD, standard deviation; HRT, hormone replacement therapy.

*Including women with surgically induced menopause as a result of PBSO.

†Use of HRT following PBSO.

women in the GS group (scale mean, 7.9; range, 5 to 20; $P < .001$; effect size = 0.44). The effect of PM and the interaction effect of prophylactic oophorectomy (yes *v* no) and PM (yes *v* no) were (marginally) significant ($P < .05$ and $.01$, respectively). Women who had undergone both PBSO and PM (PBSO+PM+) reported significantly lower levels of cancer worries (mean, 6.6; range, 5 to 13) compared with mastectomized women undergoing GS (mean, 8.1; range, 5 to 20). Regarding specific aspects of cancer

worries (Table 3), significantly fewer women in the PBSO group indicated being worried about their ovarian cancer risk ($P < .001$), being worried about cancer risk among their family members ($P < .05$), and that cancer worries had affected their mood ($P < .001$) and functioning ($P < .01$). Eighty-two percent and 45% of oophorectomized women reported that their anxiety about developing ovarian and breast cancer, respectively, had decreased substantially since their surgery (Table 2); for the PBSO+PM+ group,

Table 2. QOL Assessments by Preventive Health Strategies for Ovarian Cancer: PBSO Versus GS

QOL	PBSO (n = 369)		GS (n = 477)		P (PBSO v GS)	Effect Size	Population Norm (n = 487)*	
	Mean	SD†	Mean	SD†			Mean	SD
Generic QOL,‡ score								
Global health status	74.9	19.0	76.1	19.4	.51			
General health perceptions	70.3	22.4	70.9	19.7	.73		70.0	20.0
Vitality	62.7	18.7	64.0	17.2	.55		65.1	19.5
Mental health	73.7	15.9	72.9	15.7	.29		74.1	18.2
Role-emotional	75.4	37.2	79.2	33.8	.95		79.8	35.1
Condition-specific QOL								
Intrusive thoughts§								
Sum score	6.8	7.8	7.0	7.7	.37			
Patients scoring ≥ 20, %		8.9		9.6	.73			
Cancer worries,§ score	7.0	1.9	7.9	2.2	< .001	0.44		
Women reporting a large decrease in anxiety about ovarian cancer after PBSO, %		82.1						
Women reporting a large decrease in anxiety about breast cancer after PBSO, %		44.9						
Perceived breast cancer risk before PBSO (0-100), score	58.6	29.5						
Currently perceived breast cancer risk (0-100), score	29.5	28.0	39.0	28.2	< .05	0.34		
Endocrine symptoms, score	56.0	9.5	59.7	9.6	< .001	0.34		
Sexually active women, %¶		75		81	.11			
Sexual functioning,† score								
Pleasure	9.6	3.5	10.7	3.2	< .05	0.33		
Discomfort	4.4	1.7	5.1	1.4	< .05	0.45		
Habit	0.9	0.5	0.9	0.5	.73			

Abbreviations: QOL, quality of life; PBSO, prophylactic bilateral salpingo-oophorectomy; GS, gynecologic screening; SD, standard deviation; ANCOVA, analysis of covariance.

*Population norm scores were available only for the Short Form-36 scales (general health perceptions, vitality, mental health, and role-emotional). None of the comparisons between the general population scores and those of high-risk women (PBSO and GS groups) were statistically significant (all $P > .3$). All analyses were adjusted for age.

†Unadjusted means, P values, and effect sizes for the main effect. PBSO versus GS in ANCOVA. All analyses were controlled for age, DNA status, parity, history of breast cancer, and prophylactic mastectomy. Effect sizes were calculated according to the following formula: Cohen's $d = (M_{\text{PBSO}} - M_{\text{GS}}) / \sigma_{\text{pooled}}$, where $(M_{\text{PBSO}} - M_{\text{GS}})$, σ_{PBSO}^2 , and σ_{GS}^2 indicate, respectively, a difference in mean QOL scores and score variances of the PBSO and GS groups and where $\sigma_{\text{pooled}} = \sqrt{(\sigma_{\text{PBSO}}^2 + \sigma_{\text{GS}}^2) / 2}$. Effect sizes are indicated only for $P < .05$.

‡Higher scores correspond to better functioning or less symptoms.

§Higher scores indicate more intrusive thoughts or worries.

||Lower scores indicate higher levels of endocrine symptoms.

¶The Sexual Activity Questionnaire was administered to a smaller sample of women (PBSO, $n = 248$; GS, $n = 265$). Scores for sexual functioning apply only to women who reported that they had been sexually active in the last 4 weeks. Higher scores represent higher levels of sexual functioning.

these percentages were 90% and 57%, respectively. Comparable data were not available for the GS group.

Adjusting for possible confounders, the perceived risk of developing breast cancer was marginally significantly lower among women who had undergone PBSO than among women in the GS group. The effect of PM was statistically significant ($P < .001$; effect size = 0.58), with the lowest estimated risk being in the PBSO+PM+ group (mean \pm SD, 12.9 \pm 11.2) and the highest risk being in women undergoing GS only (mean \pm SD, 46.9 \pm 26.0). For the entire PBSO group, the perceived risk of breast cancer had decreased, on average, by 29.1 points on a scale 0 to 100 compared with before ovarian surgery (retrospective estimate). For women who had also undergone PM, the decrease was 51.3 points (data not shown).

No significant differences in the level of sexual activity were observed between the PBSO and GS groups (Table 2). However, women in the PBSO group reported marginally significantly more discomfort (vaginal dryness and dyspareunia; $P < .05$), less pleasure and satisfaction during sexual activities ($P < .05$), and significantly more endocrine symptoms ($P < .001$) than the GS group. No significant differences were observed between HRT users and nonusers after ovarian surgery in the levels of endocrine symptoms and sexual functioning (data not shown). Menopausal status at the time of PBSO (premenopausal v postmenopausal) and the time since PBSO were not significantly related to the current levels of generic and condition-specific QOL reported by oophorectomized women (data not shown).

Table 3. Multivariate OR and 95% CIs for Women Who Had Opted for GS Compared With Women Who Had Undergone PBSO by Cancer Worries in the Past 4 Weeks (worried v not worried)

Selected Item	% of Patients		OR	95% CI†	P‡
	PBSO (n = 369)*	GS (n = 477)*			
Worried about ovarian cancer	15.2	37.4	3.2	2.2 to 4.7	< .001
Worried about breast cancer	43.0	61.0	1.3	0.9 to 1.8	.14
Worries affected mood	27.9	43.3	1.7	1.3 to 2.6	< .001
Worries affected functioning	11.0	17.0	1.9	1.2 to 2.9	< .01
Worried about other family members at risk	60.8	65.9	1.4	1.0 to 1.9	< .05

Abbreviations: OR, odds ratio; GS, gynecologic screening; PBSO, prophylactic bilateral salpingo-oophorectomy.

*Unadjusted percentages.

†All analyses were controlled for age, DNA status, parity, history of breast cancer, and prophylactic mastectomy.

Satisfaction With Preventive Health Strategies

Ninety-seven percent of women who had undergone PBSO reported being satisfied with the decision they had made compared with 82% of women in the GS group ($P < .01$). Regrets about the decision on the preventive health strategy were expressed by 5% of the PBSO group and 6% of the GS group ($P > .05$). Eighty-six percent of women would choose PBSO again, and 63% would recommend it to a friend with familial risk of ovarian cancer. In the GS group, 14%, 4%, and 15% of women intended to undergo PBSO within 5 years, within 10 years, and at some unspecified time in the future, respectively. Dissatisfaction with GS was not related significantly to the intention to undergo PBSO in the future.

DISCUSSION

To our knowledge, this is the largest cross-sectional, observational study to date that describes the psychosocial issues of ovarian cancer prevention in high-risk women. The results provide a comprehensive assessment of the generic and condition-specific QOL in 846 women who had opted either for PBSO or periodic GS.

All study participants reported high levels of generic QOL that were not significantly different from the QOL levels of women in the general Dutch population. Despite the fact that PBSO is an irreversible procedure with major consequences for the bodily hormonal balance, which, in turn, may affect the level of the patients' general well-being, we found no adverse impact of PBSO on generic QOL. These results are consistent with earlier findings^{17,18} but in contrast with one study²⁰ that suggested impairments in generic QOL as a result of PBSO. The discrepant results of the latter study may be a result of methodologic issues, such as a small sample size and the lack of statistical control for possible confounding medical factors.

Our results indicate that PBSO is associated with significantly lower levels of cancer worries compared with GS,

with the fewest worries being expressed by women who had undergone both PBSO and PM. Additionally, 45% and 82% of women also indicated that PBSO had led to a large decline in anxiety about breast and ovarian cancer, respectively. As expected, the anxiety reduction was even larger for women who had undergone both prophylactic ovarian and breast surgery. Our findings are in line with other reports^{17-19,35} that have found a postsurgery reduction in cancer-specific distress but contrast with the results of Fry et al,²⁰ who found no beneficial effects of PBSO over GS on cancer worries. This discrepancy may be a result of the fact that Fry et al²⁰ used a different measure of cancer worries³⁶ than the measure used in our and other studies. Also, their sample was small, and it may not have been representative of the larger population of high-risk women. Their sample was recruited from a single center and, compared with our sample, included fewer women with a history of breast cancer (31%), more women who were premenopausal at PBSO (50%), and no women who had undergone PM.

In addition to the cancer worry scale, we also administered the intrusion subscale of the IES to assess cancer-specific distress. No significant differences were observed in the level of intrusive thoughts about breast and ovarian cancer between the PBSO and GS groups. The cancer worry scale can be viewed as a subclinical distress measure, whereas the intrusive thought subscale of the IES is intended to assess a more severe form of distress, capturing symptoms of PTSD. In our study sample, approximately 10% of all women exhibited symptoms suggesting the presence of PTSD, with breast and ovarian cancer risk as an underlying stressor. It is worth noting that, although PBSO reduces objective cancer risk, it does not eliminate high levels of cancer-specific distress in some women.

After controlling for possible confounders, a significant difference was observed in breast cancer risk perception, with the PBSO group scoring significantly lower than the GS group. A comparison between the pre- and postoperative (retrospective) assessments of perceived breast

cancer risk indicated a decrease, on average, of 29% after PBSO and 51% after both PBSO and PM. Our results suggest that high-risk women benefit both medically and psychologically from prophylactic surgery by the reduction of both their objective cancer risk and their perceived risk of developing cancer and that this benefit is the greatest among women who undergo both ovarian and breast surgeries.

As expected, PBSO was associated with more endocrine symptoms and worse sexual functioning than GS. The use of HRT had relatively limited impact on the level of menopausal and sexual symptoms in the PBSO group (detailed data on this issue will be reported in a separate article). Although the PBSO and GS groups included comparable numbers of sexually active women, prophylactic surgery was associated with more discomfort and less pleasure and satisfaction during sex. Post-surgery increase in levels of menopausal symptoms and declines in sexual functioning caused by estrogen deprivation symptoms (eg, vaginal dryness, dyspareunia, and vasomotor symptoms) have also been reported in other studies.^{17,18,35} However, no significant PBSO-GS differences in sexual functioning were detected in the study by Fry et al²⁰ using the same measure of sexual functioning. This may be because of their smaller sample size and lower rates of sexually active women (66.1%) compared with our investigation.

Our findings suggest that the beneficial QOL effects of PBSO may outweigh the adverse effects because almost all women who had undergone PBSO reported being highly satisfied with the procedure. These findings are in line with previous studies.^{18,19,37-39} The vast majority of women in the PBSO group would undergo surgery again, whereas less than two thirds of women undergoing GS would choose screening again. Almost one third of women in the GS group expressed the intention to undergo PBSO in the future. These results suggest that high-risk women may perceive GS as only a temporary preventive health strategy.

Given its multicenter nature and the relatively high response rate, we believe that the study sample was representative of high-risk women in the Netherlands. However, some possible limitations of our study should be noted. First, we did not include measures of perceived anxiety reduction or a retrospective report of changes in self-perceived cancer risk in the GS group. Given the cross-sectional study design and the longitudinal nature of screening itself, there was no clear reference point in time for the GS group that would be comparable to that for women who had undergone PBSO. Data from our ongoing, longitudinal study will be able to inform this issue. Second,

because of the cross-sectional design of the study, possible changes in QOL over time induced by prophylactic treatment or screening could not be assessed prospectively. A prospective, multicenter study is currently being conducted to obtain a more thorough picture of the QOL and symptom experience over time of high-risk women who opt for PBSO versus GS. Third, women who had undergone PBSO or GS may come from slightly different populations regarding their objective risk of developing ovarian cancer because more than half of the GS group did not have DNA testing. Although we controlled for known risk factors in our analyses, statistical adjustments for confounding factors may not have entirely ruled out possible selection bias resulting from nonrandomized comparison groups. However, given the known benefits of PBSO for ovarian cancer risk reduction and the unknown efficacy of the current GS techniques in early ovarian cancer detection, a randomized trial is not feasible and would not be ethical.

In conclusion, this study has documented both beneficial and adverse QOL effects associated with the two major health strategies for ovarian cancer in high-risk women. Physicians should discuss both the pros and cons of PBSO and GS with high-risk women seeking medical advice about their risk management. Among the benefits, reduced cancer worries after PBSO should be emphasized. The likely increase of climacteric and sexual symptoms, which may not be alleviated by the postsurgical use of HRT, should be included in discussions of adverse effects of PBSO. Balanced information will help clinicians and high-risk women to make informed decisions about the optimal preventive health strategy. Finally, our results indicate that a minority of oophorectomized women may experience high levels of distress after prophylactic treatment. Such women should be identified in a timely manner, and they should be offered (additional) psychosocial care after PBSO.

Acknowledgment

We thank all women who participated in this study, Miranda Gerritsma and Esther Janssen for their logistical and administrative support of the project, and the nursing and administrative staff members in the participating hospitals who helped run the study.

Authors' Disclosures of Potential Conflicts of Interest

The authors indicated no potential conflicts of interest.

REFERENCES

1. Visser O, Siesling S, van Dijk J: Incidence of cancer in the Netherlands 1999/2000: Eleventh report of the Netherlands Cancer Registry. Utrecht, the Netherlands, Association of Comprehensive Cancer Centres, 2003

2. Ries LAG, Eisner MP, Kosary CL, et al (eds): SEER Cancer Statistics Review, 1975-2001. Bethesda, MD, National Cancer Institute, 2004

3. Whittemore AS, Gong G, Iltis J: Prevalence and contribution of BRCA1 mutations in breast cancer and ovarian cancer: Results from three U.S. population-based case-control studies of ovarian cancer. *Am J Hum Genet* 60:496-504, 1997

4. Claus EB, Schildkraut JM, Thompson WD, et al: The genetic attributable risk of breast and ovarian cancer. *Cancer* 77:2318-2324, 1996

5. Antoniou A, Pharoah PD, Narod S, et al: Average risks of breast and ovarian cancer associated with BRCA1 or BRCA2 mutations detected in case series unselected for family

history: A combined analysis of 22 studies. *Am J Hum Genet* 72:1117-1130, 2003

6. King MC, Marks JH, Mandell JB: Breast and ovarian cancer risks due to inherited mutations in BRCA1 and BRCA2. *Science* 302:643-646, 2003

7. Bell R, Petticrew M, Sheldon T: The performance of screening tests for ovarian cancer: Results of a systematic review. *Br J Obstet Gynaecol* 105:1136-1147, 1998

8. Jacobs IJ, Skates SJ, MacDonald N, et al: Screening for ovarian cancer: A pilot randomised controlled trial. *Lancet* 353:1207-1210, 1999

9. Modugno F: Ovarian cancer and high-risk women-implications for prevention, screening, and early detection. *Gynecol Oncol* 91:15-31, 2003

10. Rebbeck TR, Lynch HT, Neuhausen SL, et al: Prophylactic oophorectomy in carriers of BRCA1 or BRCA2 mutations. *N Engl J Med* 346:1616-1622, 2002

11. Kauff ND, Satagopan JM, Robson ME, et al: Risk-reducing salpingo-oophorectomy in women with a BRCA1 or BRCA2 mutation. *N Engl J Med* 346:1609-1615, 2002

12. Piver MS, Jishi MF, Tsukada Y, et al: Primary peritoneal carcinoma after prophylactic oophorectomy in women with a family history of ovarian cancer: A report of the Gilda Radner Familial Ovarian Cancer Registry. *Cancer* 71:2751-2755, 1993

13. Bachmann GA: Vasomotor flushes in menopausal women. *Am J Obstet Gynecol* 180:S312-S316, 1999

14. Shifren JL, Nahum R, Mazer NA: Incidence of sexual dysfunction in surgically menopausal women. *Menopause* 5:189-190, 1998

15. Rossouw JE, Anderson GL, Prentice RL, et al: Risks and benefits of estrogen plus progestin in healthy postmenopausal women: Principal results from the Women's Health Initiative randomized controlled trial. *JAMA* 288:321-333, 2002

16. MacLennan A, Lester S, Moore V: Oral oestrogen replacement therapy versus placebo for hot flushes [Cochrane Database System Review]. Oxford, England, Cochrane Library, issue 3, 2004

17. Elit L, Esplen MJ, Butler K, et al: Quality of life and psychosexual adjustment after prophylactic oophorectomy for a family history of ovarian cancer. *Fam Cancer* 1:149-156, 2001

18. Robson M, Hensley M, Barakat R, et al: Quality of life in women at risk for ovarian cancer who have undergone risk-reducing oophorectomy. *Gynecol Oncol* 89:281-287, 2003

19. Tiller K, Meiser B, Butow P, et al: Psychological impact of prophylactic oophorectomy in women at increased risk of developing ovarian cancer: A prospective study. *Gynecol Oncol* 86:212-219, 2002

20. Fry A, Busby-Earle C, Rush R, et al: Prophylactic oophorectomy versus screening: Psychosocial outcomes in women at increased risk of ovarian cancer. *Psychooncology* 10:231-241, 2001

21. Ware JE, Sherbourne CD: The MOS 36-item short form health survey (SF-36): I. Conceptual framework and item selection. *Med Care* 30:473-483, 1992

22. Aaronson NK, Muller M, Cohen PD, et al: Translation, validation, and norming of the Dutch language version of the SF-36 Health Survey in community and chronic disease populations. *J Clin Epidemiol* 51:1055-1068, 1998

23. Aaronson NK, Ahmedzai S, Bergman B, et al: The European Organization for Research and Treatment of Cancer QLQ-C30: A quality-of-life instrument for use in international clinical trials in oncology. *J Natl Cancer Inst* 85:365-376, 1993

24. Ware JE, Snow KK, Kosinski M, et al: SF-36 Health Survey Manual and Interpretation Guide. Boston, MA, New England Medical Center, The Health Institute, 1993

25. Fayers P, Aaronson NK, Bjordal K, et al: EORTC QLQ-C30 Scoring Manual. Brussels, Belgium, European Organization for Research and Treatment of Cancer Study Group on Quality of Life, 1995

26. Horowitz M, Wilner N, Alvarez W: Impact of Event Scale: A measure of subjective stress. *Psychosom Med* 41:209-218, 1979

27. Brom D, Kleber RJ: De schok verwerkingslijst. *Ned Tijdschr Psychol* 40:164-168, 1985

28. Horowitz MJ: Stress response syndromes and their treatment., in Goldberger L, Breznitz S (eds): *Handbook of Stress: Theoretical and Clinical Aspects*. New York, NY, The Free Press, 1982, pp 711-732

29. Lerman C, Daly M, Masny A, et al: Attitudes about genetic testing for breast-ovarian cancer susceptibility. *J Clin Oncol* 12:843-850, 1994

30. Valdimarsdottir HB, Bovbjerg DH, Kash KM, et al: Psychological distress in women with a familial risk of breast cancer. *Psychooncology* 4:133-141, 1995

31. Lerman C, Lustbader E, Rimer B, et al: Effects of individualized breast cancer risk counseling: A randomized trial. *J Natl Cancer Inst* 87:286-292, 1995

32. Fallowfield LJ, Leaity SK, Howell A, et al: Assessment of quality of life in women undergoing hormonal therapy for breast cancer: Validation of an endocrine symptom subscale for the FACT-B. *Breast Cancer Res Treat* 55:189-199, 1999

33. Thirlaway K, Fallowfield L, Cuzick J: The Sexual Activity Questionnaire: A measure of women's sexual functioning. *Qual Life Res* 5:81-90, 1996

34. Cohen J: *Statistical Power Analysis for the Behavioral Sciences* (ed 2). Hillsdale, NJ, Lawrence Erlbaum Associates, 1988

35. Hallowell N: You don't want to lose your ovaries because you think 'I might become a man': Women's perceptions of prophylactic surgery as a cancer risk management option. *Psychooncology* 7:263-275, 1998

36. Watson M, Duvivier V, Wade Walsh M, et al: Family history of breast cancer: What do women understand and recall about their genetic risk? *J Med Genet* 35:731-738, 1998

37. Meiser B, Tiller K, Gleeson MA, et al: Psychological impact of prophylactic oophorectomy in women at increased risk for ovarian cancer. *Psychooncology* 9:496-503, 2000

38. Hallowell N: A qualitative study of the information needs of high-risk women undergoing prophylactic oophorectomy. *Psychooncology* 9:486-495, 2000

39. Swisher EM, Babb S, Whelan A, et al: Prophylactic oophorectomy and ovarian cancer surveillance: Patient perceptions and satisfaction. *J Reprod Med* 46:87-94, 2001