Survivorship in Danish women surgically treated for gynecological cancer – The impact of postoperative rehabilitation

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ABSTRACT

Objective: As cancer treatments improve and the number of survivors increase, rehabilitation and survivorship issues have become an increasingly important aspect of cancer care. Presently there is little evidence describing how to address patients survivorship needs. On this ground a cancer rehabilitation program that matches the needs of women surgically treated for early stage gynecological malignancies and their relatives, was developed and tested.

Methods: The rehabilitation program was conducted by a multidisciplinary team and consisted of lectures, physical training and group support sessions. The program consisted of four three-hour sessions held weekly over four weeks. The program was evaluated by two questionnaires completed by the patients, the Short Form 36 and Sense of Coherence, pre and post intervention.

Results: Twenty-one courses were held with 371 participants: 217 patients (participation rate 35%) and 154 relatives. The patients’ age ranged between 22-85 years. Of these, 61% had been operated for localized stages of endometrial cancer, and 27% for local stages of cervical cancer. The questionnaires demonstrated an increase in all elements of the physical and mental health components during the study period. However, general health was the only component that surpassed the level recorded for healthy women.

Conclusions: The evaluation demonstrated that the patients experienced a positive personal outcome due to participation in the program. This finding was however not fully retrieved in the questionnaire results. Informal feedback revealed that the program provided clinicians with significant patient perspectives on survivorship issues in gynecological cancer care. Most patients completed their surgical treatment within a few days, and were considered clinically cured. To these patients and their relatives, the rehabilitation program represents a combined supportive self-care and specialized shared care intervention, which is feasible in combination with the pre-existing follow-up.

Key Words: Group sessions, Physical training, Women, Relatives, Clinical pathway

1. INTRODUCTION

In this article, we present the prospective, long term evaluation of a postoperative rehabilitation program designed to match the needs of women surgically treated for early stage gynecological cancer. The evaluation applies a patient perspective in terms of the participants’ general health, coping capacity, and personal outcome of attending. In addition, the organizational and clinical experiences of running the program over a decade will be described.

Long-term survivorship issues of cancer patients are a well
recognised concern. As cancer treatment improves and inpatient hospitalisation shortens, the postoperative rehabilitation therefore constitutes an increasingly important multidisciplinary element of the overall cancer treatment trajectory. Generally, women who are treated for localised gynecological cancer diseases with surgery alone have a very good prognosis. It is anticipated that for these cancer survivors and their relatives, going through a successful postoperative rehabilitation program will support their future general health and health-related quality of life.

Rehabilitation is defined as a goal-oriented, cooperative process involving an individual, her relatives, and professionals over a specific period of time. The concept originates from The World Health Organisation’s International Classification of Functioning, Disability, and Health, and combines biological, psychological, and cultural perspectives on health in relation to rehabilitation. Within this theoretical framework, the individual woman is at the same time a physical body, an individual person, and a participant.

Due to the general increase in incidence and survival of cancer diseases in the Western world, cancer survivorship issues have been subject to an increasing focus in the organisation of treatment and care, during the last decade. In relation to cancer, rehabilitation aims to reduce the physical, psychological, and social consequences of the malignant disease and its treatment for patients as well as their families. In gynecological cancers, prognosis and treatment trajectory depends on the type and stage of disease and pre-existing comorbidities. The rehabilitation process can therefore be heterogeneous as survivorship needs vary substantially according to type and stage, and from one woman to another. The rising body of evidence however demonstrates a substantial need for post treatment rehabilitation and addressing female survivorship issues in general, such as psychosocial and sexual needs, femininity, infertility, fatigue and fear of recurrence.

In the Western world, women who are diagnosed with low stage gynecological malignancies have a good prognosis with a cancer specific five-year survival exceeding 90%. Still, approximately 20% of these bio-medically cured women seem to continue to perceive themselves as cancer patients, regardless of the time that has passed since surgery, of their age, and of the fact that most of them have finished follow-up visits years earlier. Furthermore, some of those who actually do experience side effects seem to neither look for, nor receive, any treatment of their condition. Thus, a lack of knowledge and a lack of health care services in relation to postoperative rehabilitation and late effects following fast-track gynaecological cancer surgery were identified. On this ground, a rehabilitative intervention for women who underwent surgery for low-stage localised gynaecological cancer diseases and their relatives was developed. The aim of the study was to prospectively evaluate the impact of postoperative rehabilitation in gynaecological cancer by measuring the participants’ general health, coping capacity, and personal outcome of attending.

2. Method

2.1 Participants

Women who underwent fast-track surgery for localised stages of any gynecological malignancy at a national centre for surgical treatment of gynaecological malignancies in Denmark were included in the study. Women who were unable to read or write Danish, suffered from mental illness, required adjuvant chemotherapy or radiation, or had an additional cancer diagnosis were excluded.

2.2 Study design

The design of the postoperative rehabilitation program was based on the existing knowledge of patient education programs. The program consisted of four sessions, held once per week with each session lasting three hours. It was facilitated by nurse specialists from within the surgical department, who gave lectures and served as coordinators. Sessions were held in facilities situated at the cancer centre. Each session provided a combination of lectures, physical training and supportive group sessions delivered by a multidisciplinary team of healthcare professionals and counsellors (see Table 1). Relatives were invited to attend the first session of the program including a session about being a relative of a person with cancer. During sessions, the participants were invited to share personal experiences and tell their illness history. The postoperative rehabilitation program was piloted between 2004 and 2006. As the preliminary results showed significant improvements in the participants’ coping capacity, vitality and physical functioning, the programs were continued. During the study period (2004-2014), a total of 21 courses were held. With the exception of a few small alterations, the courses maintained their original structure in providing a combination of information, physical training, and supportive group sessions (see Table 1).

2.3 Assessment tools

The postoperative rehabilitation program represented a complex intervention. It was therefore monitored quantitatively in terms of the effect on general health and coping capacity before and one year after attending, as well as qualitatively by investigating the personal impact of participation. Baseline evaluations via questionnaires took place upon enrolment into the program, with secondary evaluation one year later.
Two questionnaires that were previously validated in a Danish context were used to survey the study period. To minimize recall bias, the evaluation took place prospectively. The Short Form 36 (SF-36) questionnaire was chosen to survey general health, because it estimates self-assessed general health status in adult individuals. It is a 36-item questionnaire, where health status is defined within eight sub-areas: “physical functioning”, “limitations due to physical conditions”, “pain”, “general health”, “energy/tiredness”, “social functioning”, “limitations due to mental conditions”, and “mental well-being”.\(^{13}\) Scores range from 0 to 100, in which higher scores represent better health status. To survey the individual coping capacity, the Sense of Coherence questionnaire “SOC” was applied.\(^{14}\) This questionnaire evaluates personal resources for coping with difficult life situations in terms of being able to manage tension, to find solutions, and to mobilize personal resources. It is a 29-item questionnaire, each of which contains seven possible answers, and has a range of the means from 100 to 165, with higher scores representing better coping. Both questionnaires were filled out in two questionnaire rounds by each participant. The first time in relation to the first session, and second time one year later, where the questionnaires were sent to the respondents by mail.

### 2.4 Data, statistics and text analyses

Data entry was carried out by a data manager, and the statistical software package Stata 10 was used in the analyses.\(^{15}\) To match the purpose, data are reported descriptively in numbers, percentages, mean values, medians and range. Data were analysed comparatively, using paired t-tests to establish pre- and post differences between the study groups. Since one of the purposes of the study was to measure the difference between time points, the findings were presented with 95% confidence intervals (CI). In a single indicator, where the overlapping in CI was marginal, a two-sided p-value for the t-distribution was applied to determine significance. After having completed the postoperative rehabilitation program, the women filled in a 12-item, open-ended course evaluation form, which addressed two main focus areas: the personal outcome of participating in the program, and the content, timing, and facilities. These patient statements underwent thematic text analysis.

### 2.5 Ethics

The study was approved by the Danish Data Protection Agency, file no. 2001-41-1603 & 2004-41-4223. According to the Danish Regional Committee on Biomedical Research Ethics, the study needed no further approvals. All procedures were in accordance with the ethical standards of the Danish Regional Committee on Biomedical Research Ethics and the 1964 Helsinki declaration. The women received both oral and written information about the postoperative rehabilitation program during their hospitalization and in relation to final diagnosis, and were invited to participate by mail, one to three months after their surgery.

### 3. Results

During the study period, a number of 21 postoperative rehabilitation programs had a total of 371 participants: 217 patients (in the following referred to as women) and 154 relatives. As 618 women had been invited, the participation rate was on average 35%, with age distribution from 22 years to 85 years (median 58). The relative participation rate was 71% (see Table 2), comprising a majority of male partners (77%). Thus, participants per course were on average ten female patients of varying age, and seven relatives.

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**Table 1.** Overview of the elements in the survivorship care program (duration 3 hrs 30 min)

<table>
<thead>
<tr>
<th>Session Elements</th>
<th>Activity: Relaxation and exercise (Physiotherapist)</th>
<th>Group session - mixed (Nurse specialist)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Session no. 1</td>
<td>Gynecological cancer - the diseases and their treatment (Chief surgeon)</td>
<td>Being close to a person with cancer - for relatives (Nurse specialist)</td>
</tr>
<tr>
<td>Group session</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Session no. 2</td>
<td>Activity: The abdominals and the pelvic floor - training after surgery (Physiotherapist)</td>
<td>Cancer and life (Psychotherapist)</td>
</tr>
<tr>
<td>Group session</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Session no. 3</td>
<td>Activity: Five rhythms - workout for body and soul (Body therapist)</td>
<td>Identity, body image and sexuality (Sexologist)</td>
</tr>
<tr>
<td>Group session</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Session no. 4</td>
<td>Activity: Five rhythms - workout for body and soul (Body therapist)</td>
<td>Towards a new everyday life (Nurse specialist)</td>
</tr>
<tr>
<td>Group session</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
The majority of the women had undergone fast-track surgery for localised stages of endometrial cancer (61%) or cervical cancer (27%) (see Table 2). At the beginning of the study period, surgery was performed by laparotomy with either abdominal hysterectomy and bilateral salpingo-oophorectomy, or abdominal radical hysterectomy and pelvic lymph node resection. Pelvic lymph node resection in patients with high risk endometrial cancer was introduced in 2008. Since 2012 laparoscopy with or without robotic assistance are the main treatment modalities in both endometrial and cervical cancer.

### Table 2. Distribution of participants and diagnosis 2005-2014

<table>
<thead>
<tr>
<th>Distribution of participants</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Invited patients</td>
<td>618</td>
</tr>
<tr>
<td>Participating patients</td>
<td>217</td>
</tr>
<tr>
<td>Participating relatives</td>
<td>154</td>
</tr>
<tr>
<td>Participants total</td>
<td>371</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Distribution of diagnosis</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ovarian cancer</td>
<td>9</td>
</tr>
<tr>
<td>Endometrial cancer</td>
<td>133</td>
</tr>
<tr>
<td>Cervical cancer</td>
<td>59</td>
</tr>
<tr>
<td>Vulva cancer</td>
<td>7</td>
</tr>
<tr>
<td>Other gyn. cancer</td>
<td>9</td>
</tr>
</tbody>
</table>

### 3.1 Questionnaire results

A total of 202 pairs of SF36 and SOC questionnaires were collected in two questionnaire rounds. Subsequently three pairs of questionnaires were excluded due to > 10% incomplete answers. Of the remaining 199 pairs of questionnaires, 107 (54%) were answered in relation to the first session and 92 (46%) were answered after one year. The distribution in age and diagnoses in the respondent group proved similar to the total group of participants.

Regarding the individual coping capacity (SOC), mean ratio scores demonstrated an increase during the study period. However the increase proved non-significant measured by 95% CI (see Table 3).

Regarding the self-assessed general health status (SF 36), increase in all elements of both physical and mental health took place during the study period. In the physical component, this increase was only significant measured by 95% CI in relation to Physical Role, whereas the improvements of Physical functioning, Bodily Pain and General Health were non-significant. In the mental component the increase was significant in relation to Vitality, Social Functioning, and Emotional Role, whereas a two-sided p-value for the t-distribution in Mental Health on 0.27 demonstrated that this improvement was actually non-significant (see Table 3).

### 3.2 Personal evaluations

Between 2008 and 2013, evaluation forms were collected from 127 of 191 women (response rate 64%). The thematic analysis of the personal outcome of attending revealed that on a personal level the programme had initiated rehabilitation. One woman exemplified: “The cancer diagnosis has been a great sorrow, but at the same time my cure has been a great miracle – the rehabilitation programme has made it possible for me to cope with both aspects.” The personal outcome addressed health promotion in a broad sense: “I feel more healthy and normal after having attended the course.” The women stressed that future patients should be encouraged to participate: “My first reaction [to the invitation] was that I was doing well, but in retrospect I realize that I actually really needed this.”

### Table 3. The results from the two questionnaire rounds SOC and SF36

<table>
<thead>
<tr>
<th></th>
<th>First session (n = 107)</th>
<th>After a year (n = 92)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sense of Ccoherence (SOC)</td>
<td>Mean value</td>
<td>95 % CI</td>
</tr>
<tr>
<td></td>
<td>148</td>
<td>145 - 152</td>
</tr>
<tr>
<td>Short Form 36 (SF36)</td>
<td>Physical component</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Physical Functioning (PF)</td>
<td>84.5</td>
</tr>
<tr>
<td></td>
<td>Role Physical (RP)</td>
<td>63.2</td>
</tr>
<tr>
<td></td>
<td>Bodily Pain (BP)</td>
<td>81.4</td>
</tr>
<tr>
<td></td>
<td>General Health (GH)</td>
<td>75.9</td>
</tr>
<tr>
<td></td>
<td>Mental component</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Vitality (V)</td>
<td>63.3</td>
</tr>
<tr>
<td></td>
<td>Social Functioning (SF)</td>
<td>84.7</td>
</tr>
<tr>
<td></td>
<td>Role Emotional (RE)</td>
<td>69.4</td>
</tr>
<tr>
<td></td>
<td>Mental Health (MH)</td>
<td>77.0</td>
</tr>
</tbody>
</table>

*The two sided p-value for the t-distribution on 0.264 determined this sub-area non-significant
We find the existing number of participants appropriate, and would have preferred this topic to be dealt with on an individual bases, for instance during regular clinical follow-up.

The women estimated the ideal timing for attending to be approximately two months after their surgery, when physical recovery allowed increasing levels of physical activity, and personal reflections and psychosocial reactions still impacted daily life. Continued presence of experienced cancer nurses with insight into the various elements of the treatment trajectory was positively evaluated, as this meant that individual reactions and needs were followed up during the course. However, some specified that they found issues concerning intimacy and sexuality less suited for a group session, and would have preferred this topic to be dealt with on an individual bases, for instance during regular clinical follow-up.

3.3 Program evaluations

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4. DISCUSSION

The aim of this study was to prospectively evaluate the effect of attending a postoperative rehabilitation program for women, solely surgically treated in fast-track regimens for localised stages of gynaecological cancer. The overall results have illustrated a benefit for both patient and family through provision of specialised, postoperative rehabilitation among women, surgically treated for gynaecological cancer. Based on evaluations, postoperative rehabilitation should be initiated at the end of the postoperative period, approximately eight weeks after the surgery.

Although the description of the study group is limited, prominent characteristics are firstly the range in age distribution among the women in need, and secondly the prevalent representation of women treated for endometrial cancer (see Table 2). It represents a challenge to match the sessions to a heterogeneous group in terms of age and diagnosis, in order to make the program fully relevant to the participants. On the other hand the findings demonstrate that the need of postoperative rehabilitation does not seem to decrease with age. We find the existing number of participants appropriate, and advice to increase the number of courses rather than the number of participants, if required. Programs tailored for those women who are treated for endometrial cancer alone, or programs targeting women in a certain age, should furthermore be considered.

The questionnaire response rate was moderate of respectively 54% and 46%, resulting in quite broad confidence intervals especially in the second questionnaire round. In addition, a lack of a genuine control group made comparison difficult. The results must therefore be interpreted with caution, and further studies might include screening in a control group for problems targeted in the rehabilitation program. Still, prospective, repeated measurements of the General Health (SF36) and the Coping capacity (SOC) provided significant information concerning the nature of the healing process that women go through the first year after fast-track surgery, if they were offered specialised rehabilitation on a shared care level. As participation in the postoperative rehabilitation program was optional, the women represented those who reached out for support. It is unknown whether they also represented those who experienced the most severe problems, and worth noting that a constant level in coping capacity scores (SOC) demonstrated the presence of a quite stable and robust coping capacity among the participating women. Still, the results have indicated that the psychosocial impact of the cancer disease and treatment may exceed one year post treatment, also for those women who are fully cured and have an intact coping capacity, and despite the intervention, which sustains findings by Sekse et al.

The questionnaire indicators concerning Physical Role, and Vitality, Social Functioning, and Emotional Role respectively were significantly improved one year after treatment. As the women from a medical perspective were cured, we found it relevant to compare these results with similar data obtained in a comparable population of Danish women. However, although Physical Role had improved significantly during the study period, it did not reach the normal values of the average female population. Likewise - with the exception of Vitality - none of the mental component indicators reached normal values in the average female population during the study period. So even though improvements in the women’s physical and mental health had taken place during the study period, and even though some were significant, the women’s health status was still below the average female population one year after their treatment.

The invited self-referral procedure ensured patient involvement. Routine screening for needs and distress as an integral part of the follow-up would facilitate this, as it might lead to increased awareness of personal needs and opportunities.
On a long term basis this might contribute to minimizing the persisting social inequity within cancer rehabilitation and survivorship.\cite{18-20}

Postoperative rehabilitation takes place first and foremost in the individual woman’s everyday life, not at the hospital. The analysis of the women’s statements showed that by recognizing and reflecting on elements of their personal illness history, and by sharing it with their relatives, the women achieved new personal insight, which also strengthened their action competencies. This aspect might be further improved if former patients were involved in the sessions.

As illustrated in the evaluation forms there are certain topics, for instance intimacy and sexuality, where there is a positive but limited outcome of attending group sessions. Since former gynecological cancer patients have a substantial symptom burden in this area, and since there is a close correlation between sexual morbidity and poorer psychological adjustment\cite{21, 22} individualized support must be available, too, in order to meet individual or complex problems concerning sexuality.

Compared with international aftercare models, the postoperative rehabilitation program represents a combined supportive self-care and specialized shared care intervention rather than complex care management.\cite{3} According to the Danish Health and Medicines Authority, approximately 70% of all cured cancer patients will experience problems in relation to survivorship, which they will be able to cope with, if they receive relevant support from relatives and healthcare professionals. Furthermore, approximately 25% may need professionally led specialized input. The remaining 5% will experience complex post-treatment problems that should be addressed by a multidisciplinary long-term intervention.\cite{23} If the patient participation rate is anticipated to reflect estimated rehabilitation needs, the recruitment to the postoperative rehabilitation program of 35% seems sufficient. In addition, teaching and spending time with the women and their relatives during sessions also provided the clinicians with increasing insights into their patients’ perspectives on survivorship and post-treatment issues.\cite{24, 25}

In the Danish National Health Service, cancer treatment and care is free and organized in politically driven regions, and the duration of hospital stays are among the lowest in the world. Within the fast-track surgical treatment regimens, patients are hospitalized for only a single or few days in relation to major surgery or even operated in day care units.\cite{26, 27} Solely surgically treated cancer patients are therefore at risk of having unmet needs when discharged, especially regarding their supportive care.\cite{27, 28} During the last couple of years, the participation rate of the programs has increased (see Table 2). Increasing rates were found especially among women undergoing surgery for low stage endometrial cancer; the group of patients with the shortest hospital stays due to introduction of minimal invasive treatment procedures.

We assume this tendency may reflect uncovered needs, as the ability to receive and process information will be compromised by the postoperative physical condition and the psychosocial impact of the diagnosis.\cite{29} If this is the case, a close consideration should be given to further integrating this type of postoperative rehabilitation in regular follow-up, which might ease the transition from hospital to self care. In order to improve long-term survival by reducing the risk of cardiovascular disease, especially in endometrial cancer, the comorbidity should be especially addressed.\cite{30} Consequently, referral to lifestyle programs including physical activity, weight loss and smoking sessions should be offered to women with endometrial cancer in their local area. It is however important to be aware that if the women experience treatment-related side effects, attendance in local lifestyle programs cannot replace specialized rehabilitation.\cite{6, 31}

5. CONCLUSIONS
Due to short hospitalisation, women who are surgically treated for gynecological cancer in fast-track programs are at risk of having substantial unmet needs when discharged. However, attending a postoperative rehabilitation program may improve their future health related quality of life and thus facilitate survivorship. Postoperative rehabilitation programs that provide information, physical training and supportive group sessions can be established with modest resource consumption. These programs represent supportive self-care and specialized after-treatment, which should be an integrated part of the follow-up. However, interventions must be monitored by screenings of individual needs, and tailored to meet predicted requirements.

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CONFLICTS OF INTEREST DISCLOSURE
Both authors declare no conflicts of interest.
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