Introduction/Background Cancer and its treatments can result in physical, psychological, and cognitive impairments. The Healthy Eating Active Lifestyle (HEAL) – GYN “rehabilitation” program was developed to provide intensive group lifestyle training on exercise, nutrition, sleep, social integration, and stress management via a telemedicine platform. The aim of this study was to determine the impact of such an intervention on short-term quality of life for gynecologic cancer patients.

Methodology Gynecologic cancer patients underwent experiential instruction and personalized goal setting through an entirely virtual platform. A multidisciplinary team led by a gynecologic oncologist addressed diet, physical activity, strategies for sleep and stress management, sexual health, smoking cessation and alcohol intake. The intervention was aimed to address unmet psychosocial, emotional, physical, sexual, and spiritual needs common to cancer survivors. Self-administered questionnaires with Likert scales (1-5) were utilized in a pre- and post-fashion to assess improvements in physical activity, dietary and sleep habits, and a general medical symptom questionnaire (MSQ).

Result(s) We report outcomes on the first 22 participants. The mean age was 58.8 years; 22 were Caucasian, and 7 patients were on maintenance therapy during their enrollment. There was a significant decrease in the average number of general symptoms reported by a comprehensive medical symptom questionnaire (MSQ) (36.39 vs 24.77, p<0.05). Trends towards improvement were demonstrated in eating patterns (4.59 vs 3.74, p=0.06), perceived stress (11.32 vs 10.73, p=0.28), levels of anxiety and depression (10.76 vs 7.68, p=0.07) and weight management (17.55 vs 16.79, p=0.23). Patients also reported feeling an increased sense of purpose and connection as a result of their enrollment in the program (35.57 vs 37.26, p=0.07). Effect sizes (d) within the program were mild to moderate for all evaluated dimensions. 100% of participants would “highly recommend” the program and none complained of stress or altered mood associated with online instruction.

Conclusion A telemedicine-based peri-habilitation program is feasible and well tolerated. Based on preliminary data, there are trends towards improvement in overall general medical symptoms, eating patterns, perceived stress, levels of anxiety and depression, and physical activity. These findings support continued investigation of a telemedicine-based healthy lifestyle peri-habilitive program.

Introduction/Background Women with a history of cancer often suffer with symptoms related to treatment long after their cancer diagnosis. The aim of this study is to explore the extent to which women experience menopausal symptoms following treatment, and the impact this has on their daily lives.

Methodology Women with a history of cancer were invited to participate in an online survey in March 2020. Three separate surveys were completed which focused on menopausal symptoms, sleep disturbance and anxiety. Participants were recruited through patient support groups and via social media.

Result(s) 418 women responded to the survey, with a median age of 49 (23 – 72). Breast (63%, n=265) and cervical (17%, n=71) were the most prevalent cancers. The menopausal symptoms women found most troublesome and which they would most like to be rid of were hot flushes/night sweats (31%, n=96), disturbed sleep (27%, n=85), and feeling tired or worn out (19%, n=56). Hot flushes severely interfered with women’s quality of life, with 48% reporting “severe” sleep disturbance and 28% (n=98) felt their enjoyment of life was significantly impaired as a result. Sleep was severely affected by menopausal symptoms, with 80% (n=324) of women reported waking up in the middle of the night or early morning and trouble falling asleep again. A significant number of women were using prescribed medication to aid sleep, with 30% (n=109) using medication more than once per week. Furthermore, 43% (n=150) women reported difficulty in staying awake while driving, eating or while socialising at least once a week. Questions relating to anxiety found 75% (n=273) were “unable to relax” during the past month and this was moderate to severe for 45% of women. Furthermore, a fear of dying was reported by 30% (n=186) of women.

Conclusion This survey highlights the degree to which women suffer on a day to day basis following a cancer diagnosis and living with menopausal symptoms. Sleep disturbance and hot flushes were highlighted as particularly challenging. These data demonstrate the need to develop appropriate evidence based solutions and support services to manage menopausal symptoms after cancer treatment.

Introduction/Background The growing number of cancer survivors has led to increased focus on follow-up care. Resent...
follow-up strategies tend to shift towards personalised care with focus on patient education and self-management. There are limited data on gynaecological cancer survivors’ self-management abilities. This study aimed to assess the relationship between self-management skills and adherence to current follow-up guidelines among gynaecological cancer survivors in the Netherlands, Norway, and Denmark. We also assessed the relationship between adherence to follow-up and use of additional healthcare services.

Methodology A qualitative study was conducted with the aim of understanding the recovery process and return to daily activity in women treated for cervical cancer. Interviews were undertaken face-to-face and via the telephone using a semi-structured interview schedule. Inductive thematic analysis was used to derive a thematic framework from the data.

Result(s) Twenty-one women who had been treated for cervical cancer aged 18-60 years and were living in the UK participated in an interview. Analysis of the data revealed themes which described participants’ experience and perceptions of treatment as a paradox; physical recovery and self-management strategies of treatment related issues; psychological coping both in the short and long-term after treatment. A key finding were the nuances in different experiences between treatment types, with physical changes perceived to be more disruptive following radical treatments, whilst psychological repercussions seemed to be significant regardless of treatment type.

Conclusion This study provides novel insight into the varied recovery experiences of those treated with surgery, chemotheraphy, and radiotherapy after cervical cancer, which can be used to improve recovery experiences for future survivors.

533 CREATSAS VAGINOPLASTY IN PATIENTS AFTER PELVIC CHEMORADIATION IN CERVICAL CANCER: A PRELIMINARY STUDY

B Segarra Vidal*, P Padilla Iserte, V Lago, MT Luis Javier, S Domingo. Hospital Universitario y Politécnico de La Fe, Gynecological Oncology Unit, València, Spain

Introduction/Background Chemoradiotherapy followed by brachytherapy is the standard treatment in locally advanced cervical cancer and vaginal stenosis is one of the most common side effects after pelvic radiation. Sexual quality is the main condition that does not improve over time in these patients.

Methodology The main objective of this study was to assess the impact of the Creatsas vaginoplasty on the improvement in the sexual life of patients with cervical cancer who have received chemoradiotherapy and have sexual dysfunction measured by The Female Sexual Function Index. The design was an observational prospective individual cohort study between January 2018 - December 2019 of patients report outcomes (PRO) using The Female Sexual Function Index.

Result(s) A total of 12 patients were evaluated and 6 patients met the inclusion criteria. Among all patients, the global scores of The Female Sexual Function Index questionnaire improved after surgery except for one patient. No patients had post-operative complications. The median of the global score before the surgery was 22.95 (IQR 19.18, 24.92), after 6 months was 26.95 (IQR 24.95, 27.23) and, after 12 months was 24.5 (IQR 23.18, 26.42). The global score had increased after the surgery (OR 1.54, 95% CI 1.03-2.32) among non-adherent survivors. No other associations (OR 1.54, 95% CI 1.03-2.32) among non-adherent survivors. Non-adherent survivors were likely to be older, treated by surgery only, Danish residents and have endometrial cancer. Over-users not only reported more follow-up visits but also used additional healthcare services, such as oncologic rehabilitation and/or oncological nurses, more frequently than adherent survivors.

Conclusion Possession of low self-management skills appears to reduce the likelihood of adhering to national guidelines for gynaecological cancer follow-up. As personalisation of care is gradually implemented, a special attention on patient education for survivors at risk of low self-management may help to ensure their adherence to recommended follow-up.

460 UNDERSTANDING THE RECOVERY PROCESS AND RETURN TO DAILY ACTIVITY IN WOMEN TREATED FOR CERVICAL CANCER

N Millet, ªE Moss*, ¹H Munt, ²H Modermott. ¹Loughborough University, School of Sport, Exercise and Health Sciences, Loughborough, UK; ²University of Leicester, Leicester Cancer Research Centre, Leicester, UK

Introduction/Background Cervical cancer is predominantly a cancer of younger women and improvements in oncological outcomes have resulted in more women surviving cervical cancer and living with the long-term effects of treatment at a younger age. Understanding the recovery process and the experience of cancer survivors is essential to increase awareness of the long-term psychological and physical needs of survivors.

Methodology A qualitative study was conducted with the aim of understanding the recovery process and return to daily activity in women treated for cervical cancer. Interviews were undertaken face-to-face and via the telephone using a semi-structured interview schedule. Inductive thematic analysis was used to derive a thematic framework from the data.

Result(s) Of 4455 invited survivors, 2428 (55%) returned the questionnaires. After excluding 992 randomly selected Danish survivors to even the distribution between countries and survivors with more than 5 years follow-up, 911 survivors were included in the study. Survivors with high self-management skills (e.g. ability to manage medical and psychosocial aspects of one’s illness) most frequently attended the recommended number of follow-up visits, non-adherent if they had fewer visits than recommended, or over-users if they had more visits than recommended. Results Twenty-one women who had been treated for cervical cancer aged 18-60 years and were living in the UK participated in an interview. Analysis of the data revealed themes which described participants’ experience and perceptions of treatment as a paradox; physical recovery and self-management strategies of treatment related issues; psychological coping both in the short and long-term after treatment. A key finding were the nuances in different experiences between treatment types, with physical changes perceived to be more disruptive following radical treatments, whilst psychological repercussions seemed to be significant regardless of treatment type.

Conclusion This study provides novel insight into the varied recovery experiences of those treated with surgery, chemotheraphy, and radiotherapy after cervical cancer, which can be used to improve recovery experiences for future survivors.