Fatigue in patients with colorectal cancer treated with capecitabine and oxaliplatin

Sá Lilian A1, Vettori Jossiane C1, Moreira Claudia LR1, Leandro Miraya S1, Tirapelli Daniela PC2, Perdoná Gleici SC3, Souza Hayala CC1, Oliveira Harley F1, Santos Fabiana N1, Rapatoni Liane1, Ribeiro Karen B1, Almeida Thiago CL1, Feres Omar2, Rocha José JRR2 and Peria Fernanda M4*

1School of Medicine of Ribeirão Preto, University of São Paulo, Brazil
2Department of Surgery and Anatomy, School of Medicine of Ribeirão Preto, University of São Paulo, Brazil
3Department of Social Medicine, School of Medicine of Ribeirão Preto, University of São Paulo, Brazil
4Department of Internal Medicine, School of Medicine of Ribeirão Preto, University of São Paulo, Brazil

Abstract

Cancer-related fatigue is characterized by feelings of extreme tiredness, which do not improve with rest and negatively affects the physical and psychosocial function of patients, significantly reducing their quality of life. The prevalence of fatigue in cancer patients undergoing chemotherapy can be as high as 90%. Colorectal cancer is one of the neoplasias with the highest incidence of cancer-related fatigue, especially among individuals older than 65 years old. To perform a comparative assessment of cancer-related fatigue in patients with colorectal cancer during the first four cycles of chemotherapy using the CAPOX (capecitabine and oxaliplatin) regimen.

From 35 individuals initially recruited, 27 responded to the FACT-F (Functional Assessment of Chronic Illness Therapy - Fatigue scale) questionnaire at the four timepoints stipulated to assess fatigue. The average age of the sample population was 61.9 (33-84) years old, and most participants were female (66.7%). Analysis of the data revealed that fatigue did not increase and that the quality of life did not decrease over the course of chemotherapy using CAPOX. Analysis of the confidence interval of the difference of the means for the qualitative variables of the FACT-F (fatigue) and FACT-G (quality of life) questionnaires revealed that gender was the only variable that exhibited a significant correlation with fatigue, as the scores of females were correlated with greater fatigue compared with the males (p= 0.03).

Fatigue did not increase in patients with colorectal cancer during the course of the first four cycles of chemotherapy using the CAPOX regimen. The results indicated that fatigue was stronger among females compared with males.

Introduction

Cancer-related fatigue is characterized by feelings of severe tiredness that do not improve with rest and significantly impair the quality of life of patients in various stages of cancer [1]. Despite being considered one of the worse symptoms and occurring frequently, fatigue is not always given adequate attention by healthcare professionals or the patients themselves [2].

Fatigue may be present at the time of diagnosis, and its prevalence exhibits a remarkable increase as the disease progresses [3,4]. The prevalence of cancer-related fatigue may be as high as 75% in untreated patients and may reach 90% among patients undergoing radiotherapy and/or chemotherapy [5,6].

The incidence and prevalence of colorectal cancer are high worldwide, and individuals older than 60 years old are the most affected. Relapse occurs in approximately 40% of the patients with non-metastatic disease at diagnosis who are subjected to curative surgery [7,8].

The literature describes several instruments for the diagnosis and measurement of fatigue intensity, among which the Functional Assessment of Cancer Therapy - Fatigue Scale (FACT-F) stands out. The FACT-F is a 40-item unidimensional questionnaire that allows detection of significant variations in the degree of fatigue and correlation of these variations with the various treatments used in cancer and their outcomes. In addition, the FACT-F has already been translated and validated for the Brazilian population [9-12].

Fatigue poses a major challenge for oncology, as few clinical studies have investigated specific treatments for this symptom. Its pathogenesis is unknown, but fatigue is assumed to be multifactorial, involving the participation of psychological, social, and physiological components. Among the latter, anemia, disorders of the hypothalamic-pituitary-adrenal axis, serotonin metabolism disorders, and increased proinflammatory cytokine activity have been investigated. Proinflammatory cytokines may represent the common denominator underlying the various pathways involved in cancer and either in fatigue development [13,14].

As colorectal tumors are one of the most prevalent forms of neoplasia and given that a high percentage of oncology patients...
experience fatigue at some point during treatment, that quality of life significantly decreases due to fatigue, and that the pathophysiology of fatigue remains poorly understood, studies aimed at assessing the impact of fatigue on oncology patients are needed to contribute to the development of possible therapies and clinical interventions.

Therefore, the aim of the present study was to assess the presence of fatigue in patients with colorectal cancer before the first four cycles of chemotherapy using the CAPOX protocol (capecitabine and oxaliplatin) by the FACT-F questionnaire and to assess the progression of this symptom over the course of subsequent chemotherapy cycles.

**Materials and methods**

From January to August of 2014, 35 patients diagnosed with colorectal cancer and subjected to treatment with the CAPOX regimen as outpatients at the chemotherapy unit of the Teaching Hospital of the Medical School of Ribeirão Preto (Hospital das Clínicas da Faculdade de Medicina de Ribeirão Preto - HC-FMRP/USP) were recruited for the present study. Only 27 participants filled out the questionnaires at the four stipulated timepoints and were included in the analysis.

To analyze the presence of fatigue at baseline and its progression during the four first cycles of chemotherapy, the participants were asked to answer all of the questions listed in FACT-F, which investigates the physical, social/family, emotional, and functional wellbeing domains, in addition to a section specifically focused on the assessment of fatigue. The participants’ total scores were then calculated.

The answers to the FACT-F items were also used to calculate the scores of FACT-G, which provides a measure of the patients’ quality of life.

The inclusion criteria were as follows: patients diagnosed with metastatic colorectal cancer beginning first-line or adjuvant chemotherapy with the CAPOX regimen. The exclusion criteria were as follows: cognitive deficits that prevented the participants from understanding the experimental design and from answering the FACT-F questionnaire, brain metastasis, and the use of corticoids or anti-inflammatory agents.

Statistical analysis to assess the correlation between the investigated variables and scores was based on descriptive and association techniques. Linear regression analysis was performed to investigate the functional relationships among the variables.

The study was subjected to and approved by the Research Ethics Committee of HC-FMRP/USP, no. 11755/2010.

**Results**

Of the 27 individuals included in the study, 16 and 11 patients underwent adjuvant and palliative first-line chemotherapy, respectively. The average age of the sample population was 61.92 years old, varying from 33 to 84 years old. Most participants (59.26%) were 60 years old or older. No correlation was observed between the type of chemotherapy indication and the patients’ age.

Females were predominant in the present sample (66.67%), 13 participants were married (48.14%), and more than half had attended up to elementary school. Eight different occupations were mentioned by the participants, and 48.14% were retired.

Regarding the family history of cancer, the number of positive and negative answers was identical, i.e., 13, and one participant did not answer this item.

The tumor was located in the colon in 13 participants (48.15%) and in the rectum in the remainder of the sample. All of the patients were in stage III (59.3%) or IV (40.7%), and only seven exhibited distant metastases (25%).

No participant required protocol changes or used corticoids or antidepressants. Only one patient exhibited a hemoglobin concentration <10.0g/dL; this patient’s hemoglobin was 9.9 g/dL at the second chemotherapy cycle.

The average score in the fatigue questionnaire was 114.20, and the average overall score (quality of life) was 76.50. The lowest score among the four chemotherapy cycles was used for each individual. The scores obtained from the questionnaires were similar regardless of the chemotherapy cycle (Table 1), thereby indicating that fatigue did not increase over the course of the four chemotherapy cycles using the CAPOX regimen ($p=0.2574$).

The FACT-G scores remained stable (Table 2), thus indicating that the participants’ quality of life did not become worse over the course of treatment ($p=0.2411$).

The four main subscales of the FACT-F (physical wellbeing, social/family wellbeing, emotional wellbeing and functional wellbeing) and the fatigue-specific subscale were analyzed separately, but no differences were identified among the four cycles of chemotherapy.

Similarly, the FACT-F-TOI scores, which comprise all the domains in FACT-G, except for emotional wellbeing (Table 3), were also analyzed, but no difference was detected among the four cycles of chemotherapy ($p=0.6461$).

To assess the possible correlation between the scores and variables, Pearson’s coefficient was calculated. Relative to the variable age, the results indicated a weak correlation with the FACT-F and FACT-G scores.

The average FACT-F scores per tumor site were quite similar: 110 for patients with colon involvement and 117 for patients with rectal metastases (25%).
According to the results of our study, the patients’ quality of life remained constant throughout the course of treatment using the CAPOX regimen, and fatigue did not increase. We believe that these results are potentially associated with the participants’ positive response to treatment, which allowed for control of the tumor progression. The drug regimen employed is another factor that must be taken into account, as the use of oral chemotherapy (capecitabine) is associated with better quality of life compared with regimens including fluorouracil infusion, which requires hospitalization. Some studies observed better quality of life in patients using CAPOX compared to FOLFOX, a similar regimen in which the medications are administered by infusion [16].

The results of the present study, namely, the consistent quality of life and the unchanged level of fatigue, contradict the findings reported by other authors relative to other chemotherapy regimens; those authors observed correlations between systemic cancer treatment and the incidence of fatigue. According to such studies, the frequency of fatigue among oncology patients tends to increase during the course of chemotherapy and/or radiotherapy [17-20]. One study assessed 379 patients with a past history of chemotherapy for any type of cancer and found that 25% of the participants reported fatigue as the side effect with the greatest impact following the end of a cycle of chemotherapy. In addition, more than half of the sample noticed that fatigue lasted longer than any other symptom, including nausea, depression, and pain, which negatively impact the quality of life of such patients [21].

According to the results of our study, the patients’ quality of life did not become worse during chemotherapy, which might be due to the constant level of fatigue and possibly to the tumor response to cancer treatment. These findings are of paramount clinical importance, as the main goal of palliative treatment is to relieve symptoms. Within that context, a particular concern is to maintain the patients’ quality of life as long as possible during the course of treatment and to minimize the adverse events and symptoms so that the patients might assess their treatment positively over the entire course of chemotherapy. In a recent study with 48 patients with colorectal cancer treated with...
5-fluorouracil and folinic acid in the outpatient setting, the participants reported improvement in their quality of life over the course of the experiment, which the authors attributed to their expectations relative to the end of chemotherapy [22].

Relevant results were also obtained in another study, where 78 patients with colorectal cancer subjected to an average of five to eight cycles of chemotherapy using 5-fluorouracil and eniluracil per oral route assessed their quality of life. At the end of treatment, significant improvements in the patients’ overall quality of life were observed, in addition to specific improvements in the emotional domain compared with the assessment before treatment [23].

One additional study also reported satisfactory quality of life in patients with colorectal cancer subjected to adjuvant therapy. The results indicated a reduction in anxiety and improvements of physical, emotional, and social function over time [24].

One should bear in mind that cancer diagnosis and treatment exert a direct effect on the quality of life of patients with colorectal cancer. In addition, aspects such as an appropriate nutritional and cardiovascular state, practice of exercise, and the biopsychosocial structure of patients also contribute to their quality of life [25].

These data point to the difficulties in the collection and organization of the data used in the present study.

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