Methodological study to evaluate the psychometric properties of FACIT-CD in a sample of Brazilian women with cervical intraepithelial neoplasia

Cristiane Menezes Sirna Fregnani 1*, José Humberto Tavares Guerreiro Fregnani 1 and Adhemar Longatto-Filho 1,2,3,4

Abstract

Background: The occurrence of cervical intraepithelial neoplasia (CIN) is associated with changes in health-related quality of life, including psychological factors, such as fear and shame, and changes in sexuality and sexual satisfaction, such as decreased sexual desire and frequency of sexual intercourse. Personal relationships are the most affected because CIN is sexually transmitted and many women tend to blame their partner for disease transmission. The aim of this study was to evaluate the psychometric properties of the FACIT-CD questionnaire in Brazilian women diagnosed with CIN.

Methods: The properties of the FACIT-CD questionnaire were tested on a sample of 439 women seen at the Department of Prevention of Barretos Cancer Hospital, including 329 patients who were diagnosed with CIN and 110 women who were not diagnosed with the disease. The analysed parameters included internal consistency (Cronbach’s alpha), reproducibility (intraclass correlation coefficient), structural validity, convergent validity (correlation with the SF-12 and EORTC QLQ-CX24 questionnaires), discriminant validity (according to disease status, and self-rating of health), sensitivity, and responsiveness.

Results: The Cronbach alpha values of the FACIT-CD scales were higher than 0.70 with the exception of the relationship scale (0.66). The FACIT-CD reproducibility was satisfactory, with variation in the intraclass correlation coefficients ranging between 0.50 and 0.83, although the 95% confidence interval (CI) was lower than 0.40 (0.33 – 0.64) on the treatment satisfaction scale. Regarding structural validity, only one item on the physical well-being scale was not kept in the original domain. The expected correlations between the FACIT-CD and SF-12 were not confirmed, whereas the correlations between the FACIT-CD and EORTC QLQ-CX24 were confirmed. The questionnaire was able to discriminate the groups according to disease status and self-rating of health. The sensitivity was low for the relationship scale and moderate for the other scales. The responsiveness of the FACIT-CD questionnaire varied between the groups that denominate the self-perception of health as no change, improvement or worsening.

Conclusion: Our results are encouraging and indicate that the FACIT-CD questionnaire is a promising tool for the analysis of the quality of life of women with CIN.

Keywords: Cervical intraepithelial neoplasia, FACIT-CD, Psychometric properties, Human papilloma virus
Background

Human papillomavirus (HPV) infection is the most prevalent sexually transmitted disease worldwide [1]. Approximately 440 million people are estimated to have genital HPV infections worldwide [2], and approximately 10% of women will carry HPV at some point in their life [3].

Approximately 40 types of HPV can invade the mucous membranes of the upper aerodigestive tract and anogenital region of humans; these HPV types are classified as low and high risk according to their carcinogenic potential [4]. Low-grade intraepithelial lesions spontaneously regress in 60% of cases, and only 10% of cases progress to high-grade lesions. Even cervical carcinoma in situ (CIN 3) may undergo spontaneous regression to normality in one-third of women [4]. The period from HPV infection to the onset of invasive cervical cancer is estimated to extend 10 to 20 years, which makes this disease preventable using well-structured screening strategies [5].

The occurrence of cervical intraepithelial neoplasia is associated with changes in health-related quality of life (HRQoL), including psychological factors, such as fear and shame, and changes in sexuality and sexual satisfaction, such as decreased sexual desire and decreased frequency of sexual intercourse [6–8]. Such problems tend to sustain for a period of time after the treatment [9]. Anxiety, distress, concern with fertility, changes in family dynamics and work-related changes are also negative effects of CIN diagnosis and treatment [10–13]. Because this disease is sexually transmitted, many women tend to blame their partner for transmission [13, 14].

Despite the availability of instruments to objectively assess HRQoL, few instruments have investigated the impact of HPV infection in the female genital tract. The number of studies on aspects related to HRQoL in women diagnosed with cervical cancer has significantly increased. This increased interest can be justified by the magnitude of the disease, which predominantly affects young women who will live the rest of their lives with the consequences of the disease and treatment [15–18]. However, little is known about the impact of diagnosis and treatment on HRQoL in women diagnosed with precursor lesions of cervical cancer.

In 2010, Rao et al. [6] developed a tool that was designated the Functional Assessment of Chronic Illness Therapy – Cervical Dysplasia (FACIT-CD) to assess the functional, physical, and psychological characteristics of women with CIN. The questionnaire has recently been translated and adapted to Brazilian Portuguese.

The FACIT system questionnaires are easy to apply (self-applied or using interviews), require little time to complete, have adequate validity and sensitivity to detect changes, and are designed to reach a population with a level of education corresponding to the fourth year of primary school (9–10-year-old age group) [19].

The aim of this study was to evaluate the psychometric properties of the FACIT-CD questionnaire in Brazilian women diagnosed with CIN.

Methods

This methodological longitudinal study was conducted in the Department of Prevention and Oncological Gynaecology of the Barretos Cancer Hospital, Barretos, state of São Paulo, Brazil. A total of 439 women were eligible, including 329 women with a histopathological diagnosis of CIN (low or high grade) without treatment and 110 women not diagnosed with the disease. The participants attended the Department of Prevention for screening via a cervical cytology examination (Papanicolaou test). Illiterate women and women known to have psychological or psychiatric disorders that could hinder the understanding of the questionnaire and the informed consent form were excluded.

After formal agreement to participate in the study, the participants answered the questionnaires, which were applied using interviews by a single interviewer. Sociodemographic and clinical data were initially collected. Then, the FACIT-CD, EORTC QLQ-CX24, and SF-12 (version 2) questionnaires were applied; this step was considered the first stage of the study.

Among the 329 women diagnosed with CIN, the first 112 were selected to answer the FACIT-CD questionnaire a second time to assess the reproducibility of the instrument. Interviews were conducted in a second consultation 30 days after the first interview to inform the test results. Of the 112 women selected, 87 (77.7%) returned on the expected date and answered the questionnaire.

The responsiveness and sensitivity of the FACIT-CD questionnaire were evaluated in 228 participants with a medical indication for surgical treatment using the loop electrosurgical excisional procedure (LEEP). Of this total, 179 (78.5%) returned after treatment during the stipulated period (4–6 months) and answered the FACIT-CD questionnaire a second time and the first question of the SF-12 questionnaire (“In general, would you say your health is?”). The responses obtained to this question at baseline and after treatment allowed the creation of groups and the classification of women as having improved health, worsened health, or no change in health. Among the other participants who underwent LEEP (49 women), 7 presented with invasive carcinoma and were forwarded to the Department of Oncological Gynaecology, 12 women returned outside the period stipulated for re-application of the questionnaire, and the remaining participants did not return on the previously scheduled date.

FACIT-CD questionnaire

The FACIT-CD instrument in Brazilian Portuguese is a specific instrument to assess the HRQoL of women with
CIN and comprises 37 questions divided into five scales to assess aspects related to physical well-being (9 questions), treatment satisfaction (4 questions), general perception (7 questions), emotional well-being (11 questions), and relationships (6 questions). The scores were calculated using the specific guidelines provided by the FACIT [20]. The responses were based on experiences from the last 7 days. The answer scale is Likert, with scores ranging between 0 and 4 (a little bit to very much). A score was assigned to each scale, and the scores were summed to obtain a single value. The total score of the questionnaire ranged from 0 to 136. A higher score indicated a better HRQoL.

**EORTC QLQ-CX24 questionnaire**
The EORTC QLQ-CX24 questionnaire was developed and validated cross-culturally by the European Organization for Research and Treatment of Cancer and was used for the assessment of HRQoL in patients with cervical cancer [21]. This instrument consists of 24 questions divided into three scales of multiple items and six scales of single items, including 11 questions on symptoms (questions 31 to 37, 39, and 41 to 43), 3 questions on body image (questions 45 to 47), 4 questions on sexual/vaginal function (questions 50 to 53), 1 question on lymphedema (question 38), 1 question on peripheral neuropathy (question 40), 1 question on menopause symptoms (question 44), 1 question on sexual worry (question 48), 1 question on sexual activity (question 49), and 1 question on sexual enjoyment (question 54). The scores were calculated separately for each scale of the multiple and single items to allow the evaluation of sexuality using the questions on sexual/vaginal function, sexual activity, and sexual enjoyment [21].

**SF-12 questionnaire**
The SF-12 questionnaire is a generic instrument for the assessment of HRQoL. This questionnaire is considered a smaller version of the Medical Outcomes Study 36 – Item Short-Form Health Survey (SF-36). The main goal of developing an instrument with a reduced number of items was to provide a questionnaire that could be answered quickly and easily, which is a good option for population-based studies and health screening [22]. The questionnaire consists of 12 questions derived from the SF-36 questionnaire. In Brazil, the SF-36 questionnaire was translated into Brazilian Portuguese and validated by Ciconelli et al. in 1999 [23]. The scores were calculated using specific software provided by the Medical Outcomes Health Survey.

**Analysis of psychometric properties**
The classical psychometric properties of the FACIT-CD questionnaire were tested by assessing the internal consistency, reproducibility, structural validity, convergent and divergent validity, known-group validity, sensitivity, and responsiveness.

Cronbach’s alpha coefficient was used to test the internal consistency of the instrument, with values equal to or higher than 0.70 considered appropriate [24]. The reproducibility of the FACIT-CD was evaluated by comparing the scores obtained in the questionnaire during the first and second interviews. For this purpose, the intraclass correlation coefficient (ICC) was used. Structural validity was assessed using a confirmatory factor analysis. The oblique rotation method was used for principal component analysis, and a five-factor solution was forced, as presented in the original questionnaire. For the analysis of convergent and divergent validity, the scores generated by the FACIT-CD questionnaire were correlated with the scores generated by the SF-12 questionnaire and the scores of the scales that assessed sexuality in the EORTC QLQ-CX24 questionnaire. The Spearman correlation coefficient was used to calculate the correlations, with values higher than 0.40 considered appropriate [25]. The assumptions of correlations between the FACIT-CD, SF-12, and EORTC QLQ-CX24 scales were established a priori.

To assess the known-group validity, women without the disease were compared with women diagnosed with CIN using the Mann-Whitney test. These two groups were also assessed based on the answers to the first question of the SF-12 (“In general, would you say your health is:”). The responses were classified as excellent/very good, good, and poor/very poor and were compared using the Kruskal-Wallis test.

Sensitivity was evaluated by calculating the magnitude of the effect using the Cohen’s D, standardized response mean (SRM), and relative efficiency tests [26]. The tests were applied to the groups before and after treatment.

Responsiveness was analysed using hypotheses established a priori. For this purpose, the study groups were compared before and after treatment (LEEP). The reference statistical method most commonly used to measure the magnitude of changes in HRQoL scores is the assessment of the effect size (ES) and the SRM [27, 28], which provide useful data concerning significant changes in clinical practice [29]. The ES and SRM are defined using Cohen’s criteria, in which values up to 0.20 indicate low responsiveness, values up to 0.50 indicate moderate responsiveness, and values higher than 0.80 indicate high responsiveness [26, 30]. The level of significance was 5% in all statistical tests.

**Ethical considerations**
This study was approved by the Research Ethics Committee of the Barretos Cancer Hospital under CAAE No. 36619714.9.0000.5432, and all the women who agreed to participate in the study signed an informed consent form.
Results
The characteristics of the study sample are shown in Table 1. The mean age of the women was 35.2 ± 10.1 years; most participants had a low education level and were Caucasian, married, and worked from home. The most common cytological result was a high-grade squamous intraepithelial lesion (ASC-H), and the most common histopathological result was CIN 2/3.

Table 2 shows the descriptive statistical analysis conducted using the scores obtained in each of the scales and the corresponding Cronbach’s alpha coefficients and intra-class correlation coefficients (ICC). Only the relationship scale presented a Cronbach’s alpha coefficient smaller than 0.70, with a value of 0.66. The coefficients that evaluated the reproducibility of the FACIT-CD questionnaire scales ranged between 0.50 and 0.83; however, the lower limit of the 95% CI was smaller than 0.40 only on the treatment satisfaction scale.

In the known-group validity analysis, the comparison between the groups of women with and without a diagnosis of the disease indicated significant differences in the average scores on all FACIT-CD questionnaire scales. Considering the health status rating by each participant, the group of women who rated their health as excellent/very good had significantly higher scores on all scales compared with the groups that rated their health as good or fair/poor (Table 3).

Regarding the structural validity of the FACIT-CD questionnaire (Table 4), the factor components were similar to those of the original questionnaire. The only exception was question GP5 (“I am bothered by side effects of treatment”); although this question belonged to the physical well-being domain in the original questionnaire, it presented higher factor loading in the emotional well-being domain.

The convergent analysis results of the FACIT-CD questionnaire are shown in Table 5. The correlation

| Variable          | Description | Diagnosed with CIN | Not diagnosed with CIN |
|-------------------|-------------|--------------------|------------------------|
| Age (Mean age)    |             | 35.2               | 48.5                   |
| Years of study    | ≤ 8 years   | 175 (53.2%)        | 58 (52.7%)             |
|                   | > 8 years   | 154 (46.8%)        | 52 (47.3%)             |
| Race              | Caucasian   | 239 (72.6%)        | 82 (74.6%)             |
|                   | Mixed       | 46 (14%)           | 13 (11.8%)             |
|                   | Black       | 42 (12.8%)         | 11 (10%)               |
|                   | Asian       | 2 (0.6%)           | 4 (3.6%)               |
| Marital status    | Married/cohabitating | 175 (53.2%) | 77 (70%)           |
|                   | Single      | 103 (31.3%)        | 13 (11.8%)             |
|                   | Separated/Divorced | 37 (11.2%) | 14 (12.7%)         |
|                   | Widow       | 14 (4.3%)          | 6 (5.5%)               |
| Occupation        | Works from home | 87 (26.4%) | 4 (31%)             |
|                   | Housewife   | 43 (13.1%)         | 16 (14.4%)             |
|                   | Rural worker | 17 (5.2%)          | 2 (1.8%)               |
|                   | Saleswoman  | 17 (5.2%)          | 2 (1.8%)               |
|                   | Other       | 165 (50.1%)        | 56 (51%)               |
| Cytological result| NILM        | 12 (3.6%)          | 106 (96.4%)            |
|                   | ASCUS       | 23 (7%)            | 4 (3.6%)               |
|                   | ASCH        | 148 (45%)          | –                      |
|                   | AGC         | 5 (1.5%)           | –                      |
|                   | LSIL        | 54 (16.4%)         | –                      |
|                   | HSIL        | 87 (26.5%)         | –                      |
| Histological results | CIN I     | 133 (40.4%)        | –                      |
|                   | CIN II/III  | 195 (59.3%)        | –                      |
|                   | Invasive cancer | 1 (0.3%)  | –                      |

NILM Negative for intraepithelial lesion or malignancy, ASCUS Atypical squamous cells of undetermined significance, ASCH Atypical squamous cells – cannot exclude HSIL, AGC Atypical Glandular Cells not otherwise specified, LSIL Low grade squamous intraepithelial lesion, HSIL High grade squamous intraepithelial lesion, CIN Cervical intraepithelial neoplasia
between the FACIT-CD and SF-12 scales was weak ($r_s < 0.40$). The correlation between the FACIT-CD and EORTC QLQ-CX24 scales was moderate ($r = 0.40–0.60$), which confirmed previously established assumptions.

Table 6 shows the sensitivity of the questionnaire to detect changes. The sensitivity of the relationship scale was considered low (ES = 0.17, SEM = 0.19). The sensitivities of the other scales that composed the FACIT-CD questionnaire were moderate (ES = 0.31–0.43; SEM = 0.29–0.52).

The results of the responsiveness analysis indicated increase in the scores of the scales among women who reported improved health (4/5 scales) (Table 7). The magnitude of the change was moderate (ES = 0.27–0.58; SEM = 0.30–0.71). In this same group, the only scale in which the scores worsened after treatment was general perceptions (18.5–17.4; $p = 0.001$). The same scale indicated worsened HRQoL scores when the sensitivity of the FACIT-CD questionnaire was evaluated.

Among women without changes in health between the assessments, the average scores remained unchanged (8.6–8.8; $p = 0.021$) and had low responsiveness (ES = 0.009; SEM = 0.10) only in the relationship scale (1/5 scales). In the other scales, the HRQoL scores improved with the exception of the general perception scale, which maintained the tendency of worsening after treatment.

Different results were found in the group of women who reported worsening of health between assessments. The decrease in the HRQoL scores was evident on the scales that assessed physical well-being and general perceptions (2/5 scale). There were no differences in the relation scale and the total FACIT-CD score. However, the treatment satisfaction and emotional well-being scales improved.

**Discussion**

To the best of our knowledge, this study is the first to validate a questionnaire (translated into Brazilian Portuguese) that measures the quality of life of women diagnosed with cervical intraepithelial neoplasia. The FACIT-CD questionnaire was developed by Rao et al. [6] in 2010. To date, no other studies have evaluated the psychometric properties of this instrument, which means that some comparisons are only exploratory.

The first test assessed the reliability of the questionnaire by analysing the internal consistency using Cronbach’s alpha coefficient. Results higher than 0.70 indicate that the items on the scales or domains are homogeneous or that they measure the same attribute. In this study, the value on the relationship scale was lower than expected (0.66). However, other authors support the hypothesis that Cronbach’s alpha values higher than 0.60 could be acceptable [31]. Despite this assumption, we believe that a value of 0.70 would be more desirable, and thus, we considered that the relationship scale did not achieve adequate internal consistency. Therefore, these results suggest that the relationship scale does not measure the same attribute because it addresses questions about the emotional

---

**Table 2** Cronbach's alpha coefficients and intraclass correlation coefficients of the FACIT-CD questionnaire

| Scale                  | Mean (SD) | Median | Minimum-maximum | Variation | Cronbach's alpha | Intraclass correlation coefficient (95% CI) |
|------------------------|-----------|--------|------------------|-----------|------------------|------------------------------------------|
| Physical well-being    | 23.4 (4.2) | 24.0   | 9–28             | 0–32      | 0.70             | 0.74 (0.62–0.82)                          |
| Treatment satisfaction | 9.7 (1.9)  | 9.0    | 3–12             | 0–16      | 0.77             | 0.50 (0.33–0.64)                          |
| General perceptions    | 18.8 (3.8) | 19.0   | 5–24             | 0–28      | 0.76             | 0.72 (0.51–0.84)                          |
| Emotional well-being   | 30.6 (7.0) | 32.0   | 5–40             | 0–44      | 0.79             | 0.76 (0.65–0.84)                          |
| Relationships          | 8.6 (2.2)  | 9.0    | 1–12             | 0–16      | 0.66             | 0.67 (0.54–0.77)                          |
| FACIT-CD               | 91.1 (11.6)| 92.0   | 59–115           | 0–136     | 0.73             | 0.83 (0.75–0.89)                          |

**Table 3** Known-group validity of the FACIT-CD questionnaire

| Scale                  | Women diagnosed with CIN (N = 329) | Women not diagnosed with CIN (N = 110) | p* | Excellent/ Very Good (N = 90) | Good (N = 147) | Regular/Poor (N = 92) | p** |
|------------------------|-----------------------------------|---------------------------------------|----|--------------------------------|----------------|------------------------|----|
| Physical well-being    | 23.4 (4.2)                        | 24.8 (3.8)                            | < 0.001 | 25.2 (8.66) | 23.8 (3.9) | 21.0 (4.8) | < 0.001 |
| Treatment satisfaction | 9.7 (1.9)                         | 9.0 (0.6)                             | < 0.001 | 11.0 (9.63) | 9.7 (1.7)  | 9.2 (1.6)  | < 0.001 |
| General perceptions    | 18.8 (3.8)                        | 13.9 (2.6)                            | < 0.001 | 21.6 (8.71) | 18.9 (3.3) | 16.6 (4.3) | < 0.001 |
| Emotional well-being   | 30.6 (7.0)                        | 39.8 (0.4)                            | < 0.001 | 32.6 (9.72) | 30.2 (7.0) | 30.0 (7.3) | 0.048  |
| Relationships          | 8.6 (2.2)                         | 2.8 (0.6)                             | < 0.001 | 10.4 (9.62) | 8.4 (2.1)  | 8.0 (2.3)  | < 0.001 |
| FACIT-CD               | 91.1 (11.6)                       | 81.7 (4.9)                            | < 0.001 | 96.4 (10.26)| 91.1 (11.0) | 85.8 (11.4) | < 0.001 |

CIN cervical intraepithelial neoplasia
*p = Mann-Whitney; p** = Kruskal-Wallis
support that women receive from their partner and family combined with questions about their relationships with friends and support in case of need [8]. We believe that further studies with other populations are necessary to compare the results and to determine whether the problems will be repeated.

The second stage of the study evaluated the reproducibility of the FACIT-CD questionnaire (i.e., the consistency of the results after repetition of the measurements). Most of the studies that assessed reproducibility used a period of 14 ± 5 days [32–34]. Despite this recommendation, the treatment of intraepithelial lesions is not related to sudden changes in health status. Therefore, the period between assessments used in this study was 30 days because this time frame represented the interval between the colposcopy examination and the second medical consultation. The lower limit of the 95% CI of the ICC on the treatment satisfaction scale was lower than 0.40, indicating low reproducibility (i.e., the variability in treatment satisfaction was greater than desired). Some factors reported

### Table 4: Factor analysis of the FACIT-CD questionnaire (N = 329)

| Scale               | Item | Question                                                                 | Component 1 | Component 2 | Component 3 | Component 4 | Component 5 |
|---------------------|------|--------------------------------------------------------------------------|-------------|-------------|-------------|-------------|-------------|
| Physical well-being| CD1  | I have discomfort in my pelvic area (lower part of the stomach)          | −0.016      | −0.025      | 0.703       | 0.011       | 0.011       |
|                     | CD2  | I have pain in my pelvic area (lower part of the stomach)                | −0.028      | −0.009     | 0.701       | 0.023       | 0.100       |
|                     | CD3  | I have cramping in my pelvic area (lower part of the stomach)            | 0.027       | 0.052      | 0.572       | 0.003       | 0.004       |
|                     | Cx1  | I am bothered by discharge or bleeding from my vagina                     | 0.213       | 0.221      | 0.496       | −0.072      | −0.177      |
|                     | GP5  | I am bothered by side effects of treatment                              | 0.293       | 0.165      | 0.063       | 0.069       | −0.180      |
|                     | ES8  | I have pain or discomfort with intercourse                               | 0.065       | −0.124     | 0.680       | −0.090      | 0.118       |
|                     | CD4  | I have to limit my sexual activity because of the infection              | 0.122       | −0.050     | 0.665       | −0.049      | 0.014       |
|                     | CD5  | I worry about spreading the infection                                    | 0.390       | 0.097      | 0.403       | 0.080       | −0.133      |
| Treatment satisfaction| GR1 | I have confidence in my doctor                                           | 0.045       | 0.216      | 0.081       | 0.677       | −0.012      |
|                     | CD6  | I feel I have received the treatment that was right for me               | −0.001      | 0.245      | 0.025       | 0.764       | 0.037       |
|                     | CD7  | My doctor gave me explanations that I could understand                   | −0.042      | 0.113      | −0.115      | 0.775       | 0.201       |
|                     | CD8  | My doctor explained the possible benefits of my treatment               | 0.055       | 0.060      | −0.111      | 0.768       | 0.079       |
| General perceptions | GF1  | I am able to work (including at home)                                    | 0.091       | 0.558      | −0.012      | 0.159       | 0.195       |
|                     | GF3  | I am able to enjoy life                                                  | −0.129      | 0.768      | 0.014       | 0.083       | 0.136       |
|                     | HI11 | I am hopeful about the future                                            | 0.000       | 0.667      | 0.072       | 0.118       | 0.148       |
|                     | Sp9  | I find comfort in my faith or spiritual beliefs                          | 0.013       | 0.613      | 0.059       | 0.100       | 0.128       |
|                     | GF7  | I am content with the quality of my life right now                       | −0.166      | 0.646      | −0.252      | 0.001       | 0.116       |
|                     | CD9  | I feel that I can handle things that come up around this infection       | −0.204      | 0.563      | −0.111      | 0.255       | −0.008      |
|                     | CD10 | I have accepted that I have this infection                               | −0.359      | 0.401      | 0.038       | 0.260       | −0.037      |
| Emotional well-being| CD11 | I worry that the infection will get worse                                 | 0.487       | −0.056     | 0.274       | 0.022       | 0.012       |
|                     | CD12 | I have hidden this problem so others will not notice                     | 0.700       | 0.052      | −0.050      | 0.046       | −0.251      |
|                     | CD13 | I have concerns about my ability to become pregnant                     | 0.354       | 0.022      | 0.065       | 0.108       | 0.209       |
|                     | BMT18 | The cost of my treatment is a burden on me and my family                | 0.389       | −0.078     | 0.105       | −0.046      | 0.240       |
|                     | CD14 | I worry about other people’s attitudes towards me                        | 0.661       | −0.223     | 0.037       | 0.044       | −0.028      |
|                     | CD15 | I feel embarrassed about the infection                                   | 0.681       | −0.163     | 0.145       | −0.015      | −0.057      |
|                     | CD16 | I tend to blame myself for the infection                                 | 0.565       | −0.062     | 0.002       | −0.028      | −0.070      |
|                     | CD17 | I was careful who I told about the infection                             | 0.434       | 0.214      | 0.011       | 0.106       | −0.190      |
|                     | CD18 | I have had difficulty telling my partner/spouse about the infection      | 0.529       | 0.106      | −0.052      | −0.020      | −0.155      |
|                     | CD19 | I am frustrated by the infection                                         | 0.743       | −0.172     | 0.045       | −0.069      | 0.030       |
|                     | CD20 | I am depressed about the infection                                       | 0.651       | −0.324     | 0.061       | −0.033      | 0.127       |
| Relationships       | CD21 | I get emotional support from my partner/spouse                           | −0.072      | 0.147      | −0.015      | 0.032       | 0.721       |
|                     | CD22 | I get emotional support from family members                              | −0.121      | 0.146      | 0.043       | −0.019      | 0.722       |
|                     | GS1  | I feel close to my friends                                               | −0.065      | 0.252      | 0.030       | 0.176       | 0.372       |
|                     | HI3  | I have people to help me if I need it                                    | −0.023      | 0.302      | 0.018       | 0.185       | 0.630       |
by the study participants could justify this variability. The consultations were conducted by different physicians from the same team, which might lead to dissatisfaction or conversely a better evaluation in another consultation. The impact on the emotional factors of the patient might also influence this variable (e.g., whether the consultation was scheduled only to perform follow-up tests such as colposcopy or whether it was scheduled to inform the result of a test that would define a course of action). Emotional factors in these different instances (consultation for examination and consultation to receive laboratory test results) may explain this variability.

The best results were observed in the known-group validity analysis. The comparison of the groups of women with and without a diagnosis of CIN indicated significant differences in the scores on all scales. As expected, some scales showed worsening in the HRQoL scores in women without the disease. The reason for this difference was apparent in the items that composed the scales. In the scales that assessed treatment satisfaction and relationships, women without the disease responded “not at all” on various items, thereby decreasing the HRQoL scores as expected because they were not in treatment. The general perception scale evaluated items such as acceptance of infection and whether women could manage things that came up around the infection. A decrease in the HRQoL scores of women without the disease was expected for the items that composed the scale. These factors contributed to the decrease in the HRQoL scores in women without CIN compared with women with CIN based on the FACIT-CD total score. As expected, the scores of the other physical and emotional well-being scales were higher in women without the disease.

In an additional analysis, the test groups were classified based on the health status rating of each participant, with a lower score indicating a worse perception of the HRQoL. In this case, all scales showed significant differences. This analysis confirmed that the FACIT-CD questionnaire could differentiate the groups for which differences were expected.

The structural validity of the questionnaire was tested by confirmatory factor analysis. The results consistently

| Questionnaire | Scale | FACIT-CD scale | Physical well-being | General perceptions | Emotional well-being |
|---------------|-------|----------------|---------------------|---------------------|---------------------|
|               |       |                | rS (95% CI)         | rS (95% CI)         | rS (95% CI)         |
| SF-12         | Physical function | 0.20 (0.10–0.31) | NA                  | NA                  |
|               | Physical role       | 0.18 (0.08–0.28) | NA                  | NA                  |
|               | Bodily pain         | 0.16 (0.06–0.27) | NA                  | NA                  |
|               | Emotional role      | NA                | NA                  | 0.14 (0.04–0.25)   |
|               | Mental health       | NA                | NA                  | 0.38 (0.29–0.47)   |
|               | General health      | NA                | 0.32 (0.22–0.41)   | NA                  |
|               | Vitality            | NA                | 0.28 (0.17–0.37)   | NA                  |
|               | Social role         | NA                | 0.17 (0.06–0.27)   | NA                  |
|               | Physical component summary | 0.17 (0.07–0.27) | NA                  | NA                  |
|               | Mental component summary | NA              | NA                  | 0.34 (0.24–0.43)   |
| EORTC QLQ-CX24 | Sexual worry       | −0.53 (−0.61 to −0.45) | NA                  | NA                  |
|               | Sexual/vaginal function | −0.49 (−0.58 to −0.40) | NA                  | NA                  |

rs: Spearman correlation coefficient; CI: Confidence interval; NA: Not available

Table 6 Evaluation of the sensitivity of the FACIT-CD questionnaire

| Scale                  | Pre-treatment (n = 179) | Post-treatment (n = 179) | Difference between means | p*   | ES   | SRM  |
|------------------------|-------------------------|--------------------------|--------------------------|------|------|------|
|                        | Mean    | SD   | Mean    | SD   | Mean | SD   | <0.001 | 0.40 | 0.37 |
| Physical well-being    | 23.1    | 4.3  | 24.9    | 4.5  | 1.7  | 4.6  | <0.001 | 0.40 | 0.37 |
| Treatment satisfaction | 9.6     | 1.8  | 10.1    | 1.5  | 0.5  | 1.9  | <0.001 | 0.31 | 0.29 |
| General perceptions    | 18.6    | 3.7  | 17.2    | 3.3  | −1.4 | 2.7  | <0.001 | −0.37| −0.51|
| Emotional well-being   | 30.3    | 6.8  | 33.2    | 5.6  | 2.9  | 5.5  | <0.001 | 0.43 | 0.52 |
| Relationships          | 8.5     | 2.1  | 8.9     | 2.2  | 0.3  | 2.0  | <0.001 | 0.17 | 0.19 |
| FACIT-CD                | 90.2    | 11.0 | 94.5    | 10.8 | 4.2  | 9.6  | <0.001 | 0.38 | 0.44 |

SD: Standard deviation; p* = Wilcoxon; ES: Effect Size; SRM: Standardized response mean
confirmed the structure of the original questionnaire, which contained five factors. The only exception was in the fifth item of the physical well-being scale, which assessed the side effects of treatment. This item showed higher factor loading in the emotional well-being scale. The follow-up and treatment of women diagnosed with CIN have a greater emotional impact than physical impact [8]. Women who seek medical care after the diagnosis of changes in the Papanicolaou test rarely complain of physical changes but often complain of psychological changes [10–12]. This finding suggests that item GP5 ("I am bothered by side effects of treatment") is better allocated in the emotional well-being scale. On the other hand, confirmatory factor analysis is very sensitive to sample size, and its consistency requires a relatively large number of cases [35]. Therefore, an increase in the sample size may help confirm the new positioning of the variable in the model.

Regarding the convergent and divergent validities of the FACIT-CD questionnaire, we expected to find a correlation between the SF-12 and FACIT-CD questionnaire scales. However, no correlation was found, and the values were lower than 0.40. This result may have occurred because the SF-12 is a generic questionnaire that does not specifically address the questions explored in the FACIT-CD; therefore, the purposes of the evaluations were distinct. Another study that used a generic and a specific questionnaire reported the same problem when correlating the questionnaires [33]. This analysis was also conducted using the EORTC QLQ-CX24 questionnaire, which was developed to assess the HRQoL of women with cervical cancer and could easily calculate the scores of the scales and some items separately. Therefore, for this study, only the scales that assessed sexuality were used. The results of the correlation between the scales of the FACIT-CD and EORTC QLQ-CX24 questionnaires were satisfactory. In this case, it was possible to confirm the correlation of the FACIT-CD questionnaire with other dimensions for which a correlation was already expected.

Some of the women who participated in the first stage of the study and were treated surgically (LEEP) were interviewed again 6 months after surgery. In this analysis, improvements in the scale scores were expected after treatment using the SRM and relative efficiency (ES). The goal was achieved for all scales except for the general perception scale. The scale scores improved after surgery, and the sensitivity was considered low to moderate. The general perception scale indicated deterioration in the overall score; however, it was not possible to identify which items worsened. In the present study, we used the classical test theory (CTT), which tests the validity of an instrument (i.e., the ability to measure what it proposed to measure), for the psychometric analysis of the FACIT-CD questionnaire [36]. However, future studies should conduct other analyses using the item response theory (IRT) [37], which investigates items separately. [37]

| Table 7 Analysis of responsiveness of the FACIT-CD questionnaire |
| Scale                  | Health status | n  | Pre-treatment | Post-treatment | Difference between means | ES  | SRM | p*   |
|------------------------|---------------|----|---------------|----------------|--------------------------|-----|-----|------|
|                        |               |    | Mean | SD | Mean | SD | Mean | SD | Mean | SD | 0.34 | 0.29 | 0.008 |
| Physical well-being    | No change     | 73 | 23.2 | 4.4 | 24.7 | 5.0 | 1.5  | 5.2 |       |     | <0.001 |
|                        | Improvement   | 83 | 22.9 | 4.3 | 25.5 | 3.2 | 2.5  | 4.0 | 0.58  | 0.63 | <0.001 |
|                        | Worsening     | 23 | 23.4 | 4.6 | 23.1 | 6.0 | –0.3 | 4.5 | –0.07 | –0.07 | 0.87  |
| Treatment satisfaction | No change     | 73 | 9.7  | 1.8 | 10.0 | 1.5 | 0.3  | 2.0 | 0.18  | 0.16  | 0.009 |
|                        | Improvement   | 83 | 9.5  | 1.8 | 10.2 | 1.5 | 0.7  | 1.9 | 0.43  | 0.38  | <0.001 |
|                        | Worsening     | 23 | 9.7  | 1.9 | 10.2 | 1.8 | 0.5  | 1.5 | 0.27  | 0.35  | 0.028 |
| General perceptions    | No change     | 73 | 18.5 | 4.0 | 16.6 | 3.8 | –1.9 | 3.0 | –0.48 | –0.65 | <0.001 |
|                        | Improvement   | 83 | 18.5 | 3.7 | 17.4 | 2.8 | –1.0 | 2.6 | –0.29 | –0.41 | 0.001 |
|                        | Worsening     | 23 | 19.1 | 2.7 | 18.3 | 2.5 | –0.8 | 2.2 | –0.29 | –0.36 | 0.124 |
| Emotional well-being   | No change     | 73 | 30.3 | 6.6 | 33.3 | 5.2 | 3.0  | 5.5 | 0.45  | 0.54  | <0.001 |
|                        | Improvement   | 83 | 29.8 | 7.3 | 32.9 | 6.2 | 3.1  | 5.7 | 0.42  | 0.54  | <0.001 |
|                        | Worsening     | 23 | 32.0 | 5.3 | 34.0 | 4.7 | 1.9  | 5.3 | 0.36  | 0.36  | 0.038 |
| Relationships          | No change     | 73 | 8.6  | 2.1 | 8.8  | 2.2 | 0.2  | 1.9 | 0.009 | 0.10  | 0.021 |
|                        | Improvement   | 83 | 8.4  | 2.2 | 9.0  | 2.0 | 0.6  | 2.0 | 0.27  | 0.30  | 0.003 |
|                        | Worsening     | 23 | 9.0  | 2.1 | 9.1  | 2.9 | 0.1  | 2.1 | 0.06  | 0.06  | 0.284 |
| FACIT-CD (total)       | No change     | 73 | 90.4 | 11.2 | 93.6 | 11.7 | 3.1  | 10.7 | 0.27  | 0.29  | 0.003 |
|                        | Improvement   | 83 | 89.2 | 10.9 | 95.2 | 9.9  | 5.9  | 8.4 | 0.55  | 0.71  | <0.001 |
|                        | Worsening     | 23 | 93.4 | 10.3 | 94.9 | 11.3 | 1.4  | 9.7 | 0.14  | 0.15  | 0.429 |

SD Standard deviation, p* = Wilcoxon; ES Effect size, SRM Standardized response mean
Furthermore, the responsiveness of the FACIT-CD questionnaire was evaluated using the same group in which sensitivity was measured before and after treatment. Other studies have used a methodology similar to ours to evaluate responsiveness [38–40]. However, in this case, the women were divided based on their self-reported health status. After treatment, the participants answered the FACIT-CD questionnaire and the first question of the SF-12 questionnaire (on health rating). Finally, the answers provided to this question before and after treatment were compared to allow the classification of the groups as improved, worsened, or no change in health status. In the group of 83 women who exhibited improved health, we noticed an increase in the scores of the scales, reflecting an improvement in HRQoL. The total score of the FACIT-CD indicated moderate responsiveness. Responsiveness was low in the groups of women who reported health worsening or had no changes in health status. The HRQoL scores improved even among women who reported not having good health. We believe that other health problems may have interfered with the responses and that there is no direct correlation between health worsening and the worsening of signs and symptoms resulting from CIN.

Conclusions

Our results are encouraging and indicate that the FACIT-CD questionnaire is a promising tool for the analysis of HRQoL in women with CIN. Internal consistency and reproducibility were satisfactory. Regarding structural validity, only one item on the physical well-being scale was not kept in the original domain. The questionnaire was able to discriminate the groups according to disease status and self-rating of health. Sensitivity was low for the relationship scale, but moderate for the other scales. Responsiveness varied between the groups that denominate the self-perception of health as no change, improvement or worsening.

Abbreviations

AGC-US: Atypical Glandular Cells not otherwise specified; ASC-H: Atypical squamous cells of undetermined significance; CI: Confidence interval; CIN: Cervical intraepithelial neoplasia; CTT: Classical test theory; EORTC CLQ-CX24: The European Organization for Research and Treatment of Cancer Quality-of-Life questionnaire cervical cancer module; ES: Effect size; FACIT-CD: Functional Assessment of Chronic Illness Therapy – Cervical Dysplasia; HPV: Human papillomavirus; HRQoL: Health-related quality of life; HSIL: High grade squamous intraepithelial lesion; ICC: Intraclass correlation coefficient; IRT: Item response theory; LEEP: Loop electrosurgical excisional procedure; LSIL: Low grade squamous intraepithelial lesion; SF-12: Short-Form Health Survey; SRM: Standardized response mean

Acknowledgements

We would like to thank the staff members of the Departments of Prevention and Gynaecology Oncology from Barretos Cancer Hospital.

Funding

The postdoctoral fellowship was supported by São Paulo Research Foundation (Fundação de Amparo à Pesquisa do Estado de São Paulo - FAPESP, São Paulo, Brazil). Process number: FAPESP 2014/10158–3. The funding body had no role in the design of the study and collection, analysis and interpretation of data and in writing the manuscript.

Availability of data and materials

The datasets used and/or analysed during the current study available from the corresponding author on reasonable request.

Authors’ contributions

CMSF, ALF, JHTGF participated in the study concept and design. CMSF performed the interviews, the data collection and wrote the manuscript. ALF supervised the data collection. CMSF, JHTGF performed the analysis and interpretation of the results. CMSF, ADF, HHTGF revised the manuscript critically and approved the final manuscript.

Ethics approval and consent to participate

This study was approved by the Research Ethics Committee of the Barretos Cancer Hospital under CAAE No. 366197/14.9.0000.5432, and all the women who agreed to participate in the study signed an informed consent form.

Consent for publication

Not applicable.

Competing interests

The authors declare that they have no competing interests.

Publisher’s Note

Springer Nature remains neutral with regard to jurisdictional claims in published maps and institutional affiliations.

Author details

1Teaching and Research Institute of Barretos Cancer Hospital, Antenor Duarte Villela street, 1331. Barretos, São Paulo Zip code: 14784-400, Brazil. 2Life and Health Sciences Research Institute (ICVS), School of Health Sciences, University of Minho, 4710-057 Braga, Portugal. 3ICVS/3B, PT Government Associate Laboratory, Braga, Guimarães, Portugal. 4Laboratory of Medical Investigation (LIM) 14, FMUSP, São Paulo, Brazil.

Received: 6 November 2016 Accepted: 8 October 2017

Published online: 16 October 2017

References

1. Trottier H, Mahmud S, Prado JC, Sobrinho JS, Costa MC, Rohan TE, Villa LL, Franco EL. Type-specific duration of human papillomavirus infection: implications for human papillomavirus screening and vaccination. J Infect Dis. 2008;197(10):1436–47.
2. Initiative for vaccine research (IVR) - viral cancer - HPV. [http://www.who.int/mediacentre/factsheets/fs380/en/]. Accessed 20 Aug 2016.
3. Castellsagué X, Sanjosé S, Agudo A, Louie KS, Bruni L, Muñoz J, Díaz M, Irvin K, García O, Albero G, et al. HPV and cervical cancer in the world - 2007 report. Vaccine. 2007/25(Suppl 3):C1–230.
4. Creasman WT. Preinvasive disease of the cervix. In: PJ DS, Creasman WT, editors. Clinical gynecologic oncolgy. Philadelphia: Elsevier; 2007. p. 1–36.
5. WHO. Comprehensive cervical cancer control: a guide to essential practice. Geneva: WHO Press; 2006.
6. Rao D, Gela N, Daley EM, Kattazhom R, Rodriguez G, Cella D. Developing a measure of health-related quality of life for women with cervical dysplasia resulting from human papillomavirus infection. Int J STD AIDS. 2010;21(10):697–701.
7. Serati M, Salvatore S, Cattoni E, Zaninato M, Mauri S, Sestito G, Corni A, Ghezzi F, Bolli P. The impact of the loop electrosurgical excisional procedure for cervical intraepithelial lesions on female sexual function. J Sex Med. 2010(7):2267–72.
8. Dominak-Felden G, Cohet C, Atrux-Tallau S, Gilet H, Tristram A, Fandet A. Impact of human papillomavirus-related genital diseases on quality of life and psychosocial wellbeing: results of an observational, health-related quality of life study in the UK. BMC Public Health. 2013;13:1065.
9. Chow KM, Chan CW, Choi KC, Shiu AT, Cheng KK, Ip WY, Wong CM. Psychometric properties of the Chinese version of sexual function after
10. Kola S, Walsh JC. Patients’ psychological reactions to colposcopy and LLETZ treatment for cervical intraepithelial neoplasia. Eur J Obstet Gynecol Reprod Biol. 2009;146(1):96–9.
11. Wang SM, Shi JF, Pang P, Song P, Qiao YL. Impact of human papillomavirus-related lesions on quality of life: a multicenter hospital-based study of women in mainland China. Int J Gynecol Cancer. 2011;21(1):182–8.
12. Pirotta M, Ung L, Stein A, Conway EL, Mast TC, Fairley CK, Garland S. The psychosocial burden of human papillomavirus related disease and screening interventions. Sex Transm Infect. 2009;85(7):508–13.
13. Jach R, Posadzka E, Silvisik A, Zajac K, Kubicka-Turek M, Wojtowicz-Gryzb A, Huras H, Miletic-Taras T, Hossaw V, Pytymik K. The influence of surgery of cervical intraepithelial neoplasia (CIN) and cervical carcinoma on quality of life. Przegląd Lekarski. 2012;69(9):647–50.
14. Thangarajah F, Eirizmann T, Bergauer F, Patzke J, Schmidt-Petruschkat S, Theune M, Engel K, Puppe J, Richters L, Mallmann P, et al. Cervical screening program and the psychological impact of an abnormal pap smear: a self-assessment questionnaire study of 590 patients. Arch Gynecol Obstet. 2016;293(2):391–8.
15. Jensen PT, Groenvold M, Linnov A, Petersen MA, Machin D. Early-stage cervical carcinoma, radical hysterectomy, and sexual function. A longitudinal study. Cancer. 2004;100(1):97–106.
16. Wenzel I, Delaba I, Habbal R, Khilman BC, Fierlough D, Krebs LU, Antson-Culver H, Berkowitz R, Aziz N. Quality of life in long-term cervical cancer survivors. Gynecol Oncol. 2005;97(2):310–7.
17. Barnas E, Skret-Magierlo J, Skret A, Bidzinski M. The quality of life of women treated for cervical cancer. Eur J Oncol Nurs. 2012;16(1):59–63.
18. Stein KD, Syrija KL, Andrykowski MA. Physical and psychological long-term and late effects of cancer. Cancer. 2008;112(S11):2577–92.
19. Webster K, Cella D, Yost K. The functional assessment of chronic illness therapy (FACT) measurement system: properties, applications, and interpretation. Health Qual Life Outcomes. 2003;1:79.
20. Functional assessment of chronic illness therapy. [http://www.factorg.org/FACTOrganQuestionnaires]. Accessed 5 Oct 2015.
21. Geirmler ER, Kujanic Vlasic K, Waldenstrom AC, Duric VM, Jensen PT, Singer S, Chie W, Nordin A, Bjelic Radusic V, Wydra D. The European Organization for Research and Treatment of cancer (EORTC) quality-of-life questionnaire cervical cancer module: EORTC QLQ-CX24. Cancer. 2006;107(8):1812–22.
22. Campolina AG, Ciconelli RM. O sf-36 eo desenvolvimento de novas medidas de avaliaç ão de qualidade de vida. Acta Reumatol Port. 2008;33(2):127–33.
23. Ciconelli RM, Ferraz MB, Santos W, Meinão I, Quaresma MR. Tradução para a língua Portuguesa e validação do questionário genérico de avaliaç ão de qualidade de vida sf-36 (Brasil SF-36). Rev Bras Reumatol 1999, 39:143-150.
24. Cohen J. Statistical power analysis for the behavioral sciences. 2nd ed. New Jersey: Hillsdale; 1988.
25. Marquis P, Chassany O, Abetz L. A comprehensive strategy for the interpretation of quality-of-life data based on existing methods. Value Health. 2004;7(1):93–104.
26. Barrett B, Brown R, Mundt M. Comparison of anchor-based and distributional approaches in estimating important difference in common cold. Quality Life Res. 2008;17(1):75–85.
27. Jacobson NS, Truax P. Clinical significance: a statistical approach to defining meaningful change in psychotherapy research. J Consult Clin Psychol. 1991;59(1):12–9.
28. Husted JA, Cook RJ, Farewell VT, Gladman DD. Methods for assessing responsiveness: a critical review and recommendations. J Clin Epidemiol. 2000;53(5):459–68.
29. Hair JF, Anderson RE, Tatham RL, Black WC. Multivariate data analysis. 5th ed. New Jersey: Prentice Hall; 2006.
30. Tenwee CB, Bot SD, de Boer MR, van der Windt DA, Knol DL, Dekker J, Bouter LM, de Vet HC. Quality criteria were proposed for measurement properties of health status questionnaires. J Clin Epidemiol. 2007;60(1):34–42.
31. Fregnani CM, Fregnani JH, Dias de Oliveira Latoreiro Mdo R, de Almeida AM. Evaluation of the psychometric properties of the functional assessment of cancer therapy-cervix questionnaire in Brazil. PLoS One. 2013;8(10):e77947.
32. Chow KM, Chan CWH, WKW S, DYP L. Translation and validation of tools for assessing health-related quality of life and male sexual function in Hong Kong Chinese patients during transitional cancer survivorship. Supportive Care. 2017;25(7):2187–93.
33. Nunnally JC. Psychometric theory. New York: McGraw-Hill; 1978.
34. Pasquali L. Psicometria: Teoria e aplicaç ões. Brasilia: UNB; 1997.
35. Embretson SE, Reise SP. Item response theory for psychologists. NJ: Lawrence Erlbaum; 2000.
36. Schmidt S, Riel R, Frances A, Lorente Garin JA, Bonfill X, Martinez-Zapata MJ, Monéès Suarez-Varela M, dela Cruz J, Empananza J, Sanchez MJ, et al. Badder cancer index: cross-cultural adaptation into Spanish and psychometric evaluation. Health Qual Life Outcomes. 2014;12:20.
37. Cella D, Jensen SE, Webster K, Hongyan D, Lai JS, Rosen S, Tallman MS, Yount S. Measuring health-related quality of life in leukemia: the functional assessment of cancer therapy–leukemia (FACT-Leu) questionnaire. Value Health. 2012;15(8):1051–8.
38. Uwer L, Rotonda C, Guillemain F, Miny J, Kaminsky MC, Mercier M, Tournier-Rangeard L, Leonard I, Montcuquet P, Rauch P, et al. Responsiveness of EORTC QLQ-C30, QLQ-REC38 and FACT-C quality of life questionnaires in patients with colorectal cancer. Health Qual Life Outcomes. 2011;9:70.