

Effects of the Use of the Provider Resilience Mobile Application in Reducing Compassion Fatigue in Oncology Nursing

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Background: Oncology nurses have increased exposure to the prolonged illness, tragedy, loss, and premature death of patients. As a result, they are at higher risk for developing compassion fatigue.

Objectives: The aim of this study was to examine if use of the Provider Resilience mobile application (PRMA) will improve oncology nurses' professional quality of life.

Methods: The quasiexperimental design was comprised of a longitudinal approach to evaluate the effect of an intervention program, PRMA, on professional quality of life between two nonrandomized groups (intervention and control) using pre- and post-tests in a sample of oncology RNs.

Findings: The findings of this study demonstrated no significant relationships between the intervention and control groups on secondary traumatic stress, compassion satisfaction, and burnout among oncology nurses.

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Oncology nurses are at particular risk for a type of secondary traumatic stress (STS) known as compassion fatigue (CF). Nurses with compassion fatigue have been reported to experience psychological, emotional, and, eventually, physical symptoms. Without intervention, CF can lead to nurses changing jobs or careers (Boyle, 2011). This study explores the effect of a convenient, low-cost, and accessible mobile application on professional quality of life of oncology nurses at risk for CF.

Literature Review

A review of the professional literature provided descriptions of CF among nurses and other members of helping professions. In Table 1, comparisons between CF and professional

burnout illustrate the particular predisposition of nurses to CF related to the nature of their professional caring roles.

Potter et al. (2010) defined CF as “the traumatization of helpers through their effort at helping others” and as “a relational source of stress that also weighs heavily on oncology nurses” (p. E56). Several other definitions exist, and the nursing community has yet to reach a formal definition (Boyle, 2011). However, other definitions of CF share common descriptions of physical exhaustion and emotional distress resulting from close contact with patients and families. Nurses' internalization of this suffering, combined with the inability to alleviate it, contributes to feelings of professional futility or self-blame. If these scenarios repeat without alleviation or attention, nurses begin to experience emotional detachment from patients and apathy toward the patient and family.

TABLE 1. Comparison of Burnout and CF

Variable	Burnout	CF
Onset	Gradual, over time	Sudden, acute onset
Causes	Response to work or environmental stressors (e.g., staffing, workload, inadequate supplies or resources)	Consequences of caring for patients, inability to change course or painful scenario
Symptoms	Exhaustion, diminished morale, added coping demands, and lack of perceived goal achievement	Gastrointestinal distress, headache, sleep problems, and lethargy
Outcomes	Decreased empathy; withdrawal; may leave nursing, change position, or transfer	Nurses continue to give, which results in an imbalance of empathy and objectivity.

CF—compassion fatigue

Note. Based on information from Boyle, 2011; Sansó et al., 2015.

Similarities exist between CF and professional burnout, namely exhaustion, diminished morale, and lack of perceived goal achievement. Professional burnout arises from frustration when personal goals are not met, whereas CF evolves from perceived failure of rescue caretaking strategies (Valent, 2002). Nurses are particularly vulnerable to CF because they are partners, rather than observers, in patients' healthcare journeys (Boyle, 2011). Oncology nurses have the added stress of working with a patient base in which survivorship is not guaranteed, contributing to "failure" of rescue caretaking strategies. In addition, oncology nurses working in mixed-model structures (serving curative and end-of-life patients) experience value conflicts that are confusing and taxing (Fillion et al., 2009).

Coetzee and Klopper (2010) suggested that, because CF has not been formally defined within nursing practice, the phenomenon has not been explored, described, or explained in a manner that would allow nurses to identify and combat CF effectively. However, long-term sequelae of unrecognized CF can be severe and lead to career changes. Radziewicz (2001) discussed the various domains in which CF symptoms are manifested. Symptoms are physical, behavioral, and spiritual, with spiritual being at the furthest end of the spectrum and the most severe and damaging (see Figure 1).

Aycock and Boyle (2009) surveyed national Oncology Nursing Society (ONS) members on the presence of workplace resources to combat CF. The most common reported resource (60%) was the employer-provided employee assistance program, but one can argue that this is not specifically aimed at CF for oncology nurses.

Other nurses reported simple unit-based practices to deal with staff bereavement, such as having memory walls, having quiet spaces, and sending sympathy cards to families after patient deaths have occurred. Potter, Deshields, and Rodriguez (2013) described more intense strategies involving provision of a comprehensive education program to staff

in 90-minute sessions lasting five weeks (later modified to one-day workshops).

Although these extended studies succeeded in educating nurses on CF and its risks, not much data exist to support increased professional quality of life for nurses. Studies were also small (most with fewer than 50 participants) and used convenience samplings that limit generalization of findings. No published studies could be located in which technology was used as an intervention to combat CF or burnout.

Methods

Study Design

A prospective, quasiexperimental study was conducted in a 26-bed oncology unit at a medical center in California. The quasiexperimental design was comprised of a longitudinal approach to evaluate the effect of an intervention program, the Provider Resilience mobile application (PRMA), on professional quality of life between two nonrandomized groups (intervention and control) using pre- and post-tests in a sample of oncology RNs. At baseline, the intervention and control groups received CF education. The intervention group used the PRMA for six consecutive weeks. Both groups were evaluated at baseline and after completion of the six-week intervention. The institutional review board approved the study and study flyers. Informed consent was obtained from all participants.

Sample

Participants were recruited via advertisements asking for volunteers. Inclusion criteria were being aged 18 years and older, being fluent in English, being employed full-time as an oncology nurse, and owning a smartphone. A convenience sample was comprised of 25 clinical RNs from an inpatient oncology unit who attended the CF group education session. Participants were nonrandomly allocated to the intervention or control group, with 16 in the intervention group and 9 in the control group.

Data Collection

An educational inservice was given to all staff members to define CF and raise awareness. RN investigators then

Physical

Insomnia, fatigue, head and muscle aches, gastrointestinal distress, hypertension

Behavioral

Absenteeism, poor record keeping, medication errors, poor communication with peers and patients

Spiritual

Doubt in values and belief systems, becoming angry or bitter, withdrawal from specific job or profession in general

FIGURE 1. Domains of Compassion Fatigue

Note. Based on information from Coetzee & Klopper, 2010; Radziewicz, 2001.

provided an accessible and convenient intervention to staff members in the intervention group via PRMA, which was developed by the Department of Defense to aid in alleviating CF among the social workers, nurses, and physicians who treat military service members (Bonanno, 2004). RN investigators invited staff members to download and use PRMA. PRMA use was monitored for a six-week period via tracking software, which participants downloaded onto their phones.

All participants in the control and intervention groups completed the Professional Quality of Life Scale (ProQOL) as a measurement tool pretest (prior to education and PRMA download) and post-test (following the six-week monitoring period).

Instrument

Professional Quality of Life Scale: The ProQOL 5 (<http://bit.ly/IOWPcbA>) is a 30-item instrument that uses a five-point Likert-type scale ranging from 1 (never) to 5 (very often) to assess the positive and negative quality-of-life elements that those in the helping professions experience in relation to their work (Stamm, 2010). The ProQOL 5 is comprised of three subscales: STS, compassion satisfaction, and burnout. STS and burnout are components of CF. STS addresses an individual's exposure to extreme or stressful traumatic events in the workplace (Stamm, 2010).

Burnout is characterized by feelings of hopelessness and difficulty coping with work or performing one's job effectively. Compassion satisfaction assesses the degree of pleasure derived from performing one's job effectively (Stamm, 2010). Each subscale is comprised of 10 items, which asks participants to reflect on how frequently they experienced situations in their current work setting in the past 30 days.

TABLE 2. Sample Characteristics By Group

Characteristic	Intervention (N = 16)	Control (N = 9)	p
Age (years)			0.43
18–35	10	4	
36 or older	6	5	
Gender			0.36
Female	11	8	
Male	5	1	
Years as an RN			0.38
0–5	10	4	
6 or more	6	5	
Education			0.31
Associate of Science	2	3	
Bachelor of Science	13	5	
Master of Science	1	1	
Certifications			1
Yes	8	5	
No	8	4	
Oncology experience (years)			0.67
0–5	11	5	
6 or more	5	4	

Note. Statistical analyses were performed by chi-square test and Fisher's exact test.

TABLE 3. Levels of ProQOL Subscales for Secondary Traumatic Stress, Compassion Satisfaction, and Burnout By Group

Characteristic	Intervention (N = 16)	Control (N = 9)	p
Pretest stress level			1
Low	7	4	
Average	9	5	
High	–	–	
Post-test stress level			0.35
Low	10	6	
Average	6	2	
High	–	1	
Pretest compassion level			0.66
Low	–	–	
Average	6	2	
High	10	7	
Post-test compassion level			1
Low	–	–	
Average	7	4	
High	9	5	
Pretest burnout level			1
Low	10	5	
Average	6	4	
High	–	–	
Post-test burnout level			1
Low	8	4	
Average	8	5	
High	–	–	

ProQOL—Professional Quality of Life Scale

Note. Statistical analyses were performed by chi-square test and Fisher's exact test.

Higher scores on the compassion satisfaction subscale reflect greater work satisfaction, whereas higher scores on the burnout and STS subscales reflect greater difficulty related to feelings of job effectiveness (Stamm, 2010). Construct validity has been previously established (Stamm, 2010). Reliabilities (Cronbach alpha) are reported to range from 0.75–0.9 across all three subscales (Neville & Cole, 2013; Young, Derr, Cicchillo, & Bressler, 2011).

Provider Resilience mobile application: The goal of PRMA is to serve as a tool to increase provider resilience in the face of the demands on healthcare providers' personal resources. PRMA provides the following:

- Psychoeducation about the risks of burnout, CF, and STS
- A method to evaluate healthcare providers' level of burnout, CF, compassion satisfaction, and STS using a standardized assessment and a visual analog scale, which can be used to track symptoms and document change
- Tools that serve as reminders to healthcare providers to engage in self-care and brief interventions to increase resilience and reduce burnout

The resilience rating is based on the last ProQOL score, burnout rating, resilience builders and killers responses, and the last self-recorded vacation day.

The burnout rating is found through the burnout visual analog scale, which allows participants to rate themselves

TABLE 4. Mean Pre- and Post-Test Measurements of ProQOL Subscales for Secondary Traumatic Stress, Compassion Satisfaction, and Burnout in the Intervention (N = 16) and Control (N = 9) Groups

Group	Pretest			Post-Test		
	\bar{X}	95% CI	p	\bar{X}	95% CI	p
ProQOL secondary traumatic stress						
Intervention	23.06	[18.97, 31.13]	0.56	21.75	[18.5, 25]	0.5
Control	25	[18.87, 31.13]		23.78	[17.22, 30.33]	
ProQOL compassion satisfaction						
Intervention	42.63	[40.74, 44.51]	0.54	41.19	[38.29, 44.09]	0.51
Control	41.44	[36.75, 46.14]		42.78	[38.15, 47.4]	
ProQOL burnout						
Intervention	20.25	[18.08, 22.42]	0.47	21.38	[18.77, 23.98]	0.89
Control	21.67	[17.47, 25.86]		21.67	[17.64, 25.7]	

CI—confidence interval; ProQOL—Professional Quality of Life Scale
 Note. Statistical analyses were performed by independent samples t test.
 Note. High levels of secondary traumatic stress and burnout indicate higher stress levels and risk factors for compassion fatigue. Higher compassion satisfaction scores indicate lower risk factors for compassion fatigue.

on 10 affective domains, including happy, trapped, satisfied, preoccupied, connected, worn out, caring, on edge, valued, and traumatized. The rating gives the participant a simple way to track his or her feelings in each of these areas over time.

The resilience builders and killers questionnaire is a short survey of different resilience builders and killers in which the participant may have engaged. In addition to affecting the resilience rating, this questionnaire can serve as a regular reminder of things to do—or not to do—to help the participant stay emotionally resilient.

If a participant's overall resilience rating is low, he or she may be under a great deal of work-related stress. The participant is encouraged to use some of the built-in tools, like physical exercises, videos, and daily affirmations, to help the participant reconnect with the positive impact of his or her work.

The R&R Clock is an important feature of PRMA. PRMA reminds the participant to use vacation time. The R&R Clock counts the years, months, weeks, days, hours, minutes, and seconds since the user took a day off work.

Results

Sample Characteristics

Demographic and work-related characteristics of oncology nurse participants in the total sample and by treatment group are presented in Table 2. In the total sample, on average, most oncology nurses were aged 18–35 years, were female, and had five or fewer years of experience as an RN. More

than half of all participants were ONS–certified and had five or fewer years of experience in oncology.

Bivariate analyses revealed that both groups were comparable in terms of age, gender, number of years as an RN, education level, certifications, and oncology experience. Of note, the topic of CF produced reported feelings of anxiety in one participant during the initial administration of the ProQOL. The participant chose to leave the study prior to using PRMA. The individual was given a list of referrals to follow up on potential negative results of CF. No other study participants reported negative effects of PRMA use.

According to the ProQOL suggested cutoffs, 14 of the 25 participants indicated that they had average STS at pretest, 17 reported high compassion satisfaction, and 15 reported low burnout.

At post-test, 8 participants indicated that they had low STS, 14 reported high compassion satisfaction, and 12 reported low burnout. No statistically significant differences were found when comparing levels of STS, compassion satisfaction, and burnout between intervention and control groups (see Table 3).

Differences between treatment groups for the ProQOL STS subscale from baseline (pre-test) to week 6 (post-test) were not significant. In addition, no significant differences were found in the mean scores between the intervention and control groups for compassion satisfaction and burnout subscales. Improvements in ProQOL, as assessed by arithmetic mean change scores (mean difference) in overall ProQOL subscales for STS, compassion satisfaction, and burnout, were not significantly different between the intervention and control groups (see Table 4 and Figure 2).

Discussion

The findings of the current study demonstrated no statistically significant relationships between the intervention

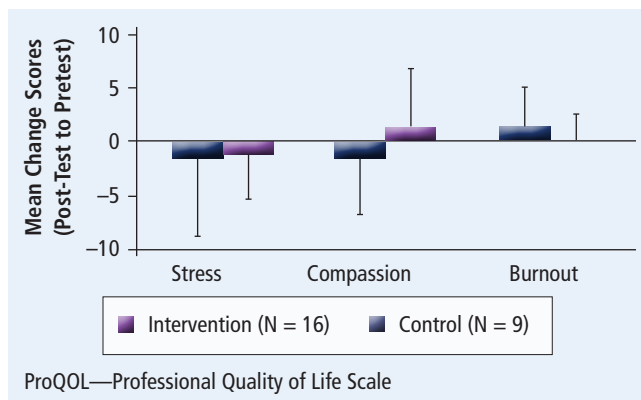


FIGURE 2. Mean Difference in ProQOL Subscale Scores

and control groups on STS, compassion satisfaction, and burnout among oncology nurses. The study investigators hypothesized an inverse relationship in the intervention group between the pre- and post-test scores for STS. Such a relationship would indicate effectiveness of PRMA in reducing nurses' stress.

Another expected outcome would have been a positive correlation in the intervention group between pretest burnout and STS scores. This relationship would indicate effectiveness of PRMA in improving the professional QOL of participants who had reported high stress and burnout levels prior to the intervention. In addition, investigators evaluated scores to determine whether the intervention group experienced higher levels of compassion satisfaction with lower levels of stress and burnout after six-week PRMA use when compared to the control group.

Analysis

The current study was not statistically significant because no changes were seen in the pre- and post-test scores of the intervention group, control group, or sample as a whole. Changes in the numbers showed a nonsignificant relationship between pretest STS and pretest burnout, which could be argued as qualitative evidence of the relationship between STS and increased burnout in oncology nurses; conversely, nurses already suffering from burnout would be more susceptible to STS. Although the data neither support nor negate the hypothesis, not enough data demonstrated any difference between groups, and the investigators believe a larger study is warranted.

The results of this pilot feasibility study are preliminary and should be interpreted with caution. The sample was relatively small, and the post-hoc power analysis suggested low statistical power to detect significant differences. Participants were not randomized into the intervention or control group. Therefore, the difference in sample size between the two groups may have factored into the lack of differences because of the lack of power. Much of the information provided via the tracking software also focused on the technical aspects of PRMA, and such data were not related to the current study's goals. The tracking software provided information regarding the amount of time each user spent on PRMA; however, time (measured in minutes) varied greatly among participants. In the end, these data were unreliable.

In addition to statistical and software limitations, logistical limitations were also discovered. The convenience sample of oncology nurses was derived from the same unit, which enforces a policy that prohibited cell phone use during work time (excluding breaks). No feasible methods were available to measure external sources of self-care used by the participants in conjunction with PRMA, which may have influenced their pre- and post-test scores. The ProQOL, although useful in theory because of its record of validity, was ultimately a challenging instrument to implement in the current study. The tool was challenging because it did not provide enough data to power the analysis. In addition, the authors hand-scored the scales, which was time consuming.

Implications for Practice

- ▶ Educate oncology nurses regarding the high risk for compassion fatigue (CF) given the close, personal relationships they share with patients and families.
- ▶ Monitor oncology nurses' behavior; the literature reflects that unrecognized CF leads to increased sick time, physical and emotional symptoms, and the possibility of nurses leaving oncology.
- ▶ Disseminate information regarding the Provider Resilience mobile application.

Implications for Practice

Lessons may be learned from the results of this pilot study. Because PRMA has not been previously studied in oncology nurses, the results of this pilot feasibility study can be used to inform a larger one. The lack of variability in pre- and post-test data may be an indicator that this particular application does not have an effect on professional quality of life among oncology nurses. However, testing the PRMA intervention requires a larger study, with a greater number of participants and data that could be more thoroughly analyzed. Additional investigations should clarify how to access more information about the PRMA application, how it works, and what information can be extracted from the tracking software's study logs prior to initiating another study.

Conclusion

Additional studies are needed to explore the use of technology to prevent CF and burnout and build resilience. A review of the literature revealed several studies that focused on the use of mobile applications for patient care, such as health prevention (Vodopivec-Jamsek, de Jongh, Gurol-Urganci, Atun, & Car, 2012), cancer pain management (Agboola et al., 2014), and use of a mobile application as a post-traumatic stress disorder (PTSD) coach (Owen et al., 2015). At the time of print, no new publications explored the use of technology to decrease CF, burnout, and PTSD in healthcare providers.

In conclusion, the literature suggests that CF is a significant problem among oncology nurses and that few interventions are effective in addressing it. Multiple intervention styles are needed to meet the varying needs of nurses, and the current study explored whether PRMA could be included in the arsenal of tools to combat the threat of CF. The small sample size in the current pilot study limited the generalizability of the findings; therefore, a larger sample is needed to explore potential effects of a mobile application on CF and burnout in this high-risk nursing population. Further research of PRMA and other technology-based tools is needed to prevent CF from subverting the dedication of oncology nurses.

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