Oncology Volunteers

The effect of a personal cancer history on compassion and psychological well-being

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BACKGROUND: The impact of support work on volunteers with a personal history of cancer has rarely been examined, despite the possibility that supporting distressed individuals may become a psychological burden for someone who has faced a life-threatening disease themselves.

OBJECTIVES: The purpose of this study is to compare compassion, self-compassion, self-coldness, and psychological well-being of oncology volunteers to the general population and clinical samples.

METHODS: Volunteers completed questionnaires on demographic and volunteer work–related characteristics, the Compassion Toward Others Scale, the Self-Compassion Scale, and the Psychological General Well-Being Index.

FINDINGS: Overall, volunteers indicated higher levels of self-compassion and psychological well-being and lower levels of self-coldness than clinical and community samples. Peer volunteers were less satisfied with their volunteer work and reported worse general health and psychological well-being than volunteers without a cancer history.

KEYWORDS: oncology volunteers; compassion; self-compassion; psychological well-being

SOCIAL SUPPORT PLAYS A SIGNIFICANT ROLE FOR PATIENTS adapting to a cancer diagnosis and in mitigating treatment-related distress (Lusczynska, Pawlowska, Cieslak, Knoll, & Scholz, 2013; Tedeschi & Calhoun, 2004). Cancer survivors who share similar experiences may be ideally positioned to provide information and support to individuals undergoing treatment (i.e., peers) to decrease social isolation and facilitate adaptive coping (Brunet, Love, Ramphal, & Sabiston, 2014; Pistrang, Jay, Gessler, & Barker, 2012). Peer support involves a survivor providing emotional or informational support to a patient at an earlier stage of the cancer trajectory (Pistrang, Jay, Gessler, & Barker, 2013). Because survivors possess firsthand knowledge of the experience of living with cancer, they offer a unique perspective that may be unavailable to individuals without a personal history of cancer.

Patients newly diagnosed with cancer and their families often are paired with cancer survivors and caregivers through volunteer organizations (Schulman-Green, Wagner, & McCorkle, 2015; Wagner & McCorkle, 2010). Systematic reviews of the psychosocial literature suggest that volunteer support programs are beneficial for patients with cancer who require information, stress management, and empowerment (Campbell, Phaneuf, & Deane, 2004; Hoey, Ieropoly, White, & Jefford, 2008; Macvean, White, & Sanson-Fisher, 2008). However, studies may lack rigorous research design and methodologies (Meyer, Coroiu, & Körner, 2015; Wagner & McCorkle, 2010), and a knowledge gap exists regarding the impact of the support work on volunteers themselves (Pistrang et al., 2013). The broader literature indicates that support providers experience benefits from their work (e.g., enhance self-esteem, self-efficacy, and feelings of social usefulness) (Brown, Hoye, & Nicholson, 2012). However, supporting distressed individuals also can lead to the supporters experiencing mood disturbances (Pistrang et al., 2013) and triggering the supporters’ own anxieties, such as fear of recurrence. Given the potential vulnerability of peer volunteers, healthcare providers may need to monitor volunteers for potentially negative psychosocial effects, prompted by their support of those in distress (Embuldeniya et al., 2013; Giese-Davis et al., 2006; Pistrang et al., 2012, 2013). Nurses are at the forefront of supportive cancer care but also play a leading role in the establishment, maintenance, and integration of volunteer programs into the larger network of psychosocial oncology services. Of
Compassion for others and compassion for oneself (i.e., self-compassion) are two related concepts that are important characteristics of volunteers providing direct services to patients with cancer. Compassion consists of recognizing, understanding, and feeling another’s suffering, while also acting to alleviate it (Strauss et al., 2016). Self-compassion is defined as extending kindness to oneself, seeing one’s experience as part of human experience, and being aware of painful thoughts and feelings (Neff, 2003). In contrast, self-coldness (a counterpart to self-compassion) corresponds to feelings of self-judgment, isolation from others, and overidentification with negative thoughts and emotions (Gilbert, McEwan, Matos, & Rivis, 2011). A strong imbalance between compassion toward others and toward oneself may lead oncology volunteers to become emotionally overinvolved in the lives of the patients they serve, leading to unhelpful and even harmful behavior. This behavior may include blaming patients for their problems and for their need for support and offering false reassurance and unsolicited advice (Gottlieb & Wachala, 2007). In addition, the widely demonstrated positive relationship between self-compassion and psychological well-being suggests that self-compassion can act as a coping strategy to prevent compassion fatigue and burnout (Boellinghaus, Jones, & Hutton, 2014; Klimecki & Singer, 2011). Ultimately, the mental health of volunteers is not only important for their own well-being, but also for the success of oncology volunteer programs.

The purpose of this study was to examine volunteer work characteristics, compassion, self-compassion, self-coldness, and psychological well-being in oncology volunteers with a personal cancer history (i.e., peer volunteers) and volunteers who have never been diagnosed with cancer. The following research questions were addressed:

- What are the levels of compassion, self-compassion, self-coldness, and psychological well-being in oncology volunteers?
- Do oncology volunteers, with or without a history of cancer, have different levels of compassion, self-compassion, self-coldness, and psychological well-being compared to general and clinical populations?
- Do peer volunteers and volunteers without a personal cancer history differ in demographic and volunteer work-related characteristics, as well as in compassion, self-compassion, self-coldness, and psychological well-being?

**Methods**

Data were collected at a well-established community cancer support program, Hope and Cope, and its cancer wellness center in Montreal, Canada, with more than 450 volunteers (340 provide direct patient service). Professional staff and a volunteer executive provide leadership and program administration. Most concern are risks when volunteers provide peer-based psychosocial care without proper support for them.

### TABLE 1.
**SAMPLE CHARACTERISTICS**

<table>
<thead>
<tr>
<th>CHARACTERISTIC</th>
<th>PEER VOLUNTEERS (N = 71)</th>
<th>VOLUNTEERS WITHOUT CANCER (N = 76)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>18–34</td>
<td>3</td>
<td>19*</td>
</tr>
<tr>
<td>35–54</td>
<td>18</td>
<td>15</td>
</tr>
<tr>
<td>55 or older</td>
<td>50</td>
<td>42*</td>
</tr>
<tr>
<td>Sex</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Female</td>
<td>57</td>
<td>66</td>
</tr>
<tr>
<td>Male</td>
<td>14</td>
<td>10</td>
</tr>
<tr>
<td>Marital status</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Married or cohabiting</td>
<td>52</td>
<td>43</td>
</tr>
<tr>
<td>Widowed</td>
<td>8</td>
<td>7</td>
</tr>
<tr>
<td>Divorced or separated</td>
<td>7</td>
<td>3</td>
</tr>
<tr>
<td>Single</td>
<td>4</td>
<td>22</td>
</tr>
<tr>
<td>Occupation</td>
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<td>29</td>
</tr>
<tr>
<td>Employed</td>
<td>30</td>
<td>33</td>
</tr>
<tr>
<td>Unemployed</td>
<td>5</td>
<td>11</td>
</tr>
<tr>
<td>On leave</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>Ethnic background</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Caucasian</td>
<td>67</td>
<td>63</td>
</tr>
<tr>
<td>Other</td>
<td>4</td>
<td>13</td>
</tr>
<tr>
<td>First language</td>
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<td></td>
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<tr>
<td>English</td>
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<td>46</td>
</tr>
<tr>
<td>French</td>
<td>9</td>
<td>10</td>
</tr>
<tr>
<td>English and French</td>
<td>2</td>
<td>1</td>
</tr>
<tr>
<td>Other</td>
<td>22</td>
<td>17</td>
</tr>
<tr>
<td>Peer mentor</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Yes</td>
<td>46</td>
<td>9**</td>
</tr>
<tr>
<td>No</td>
<td>25</td>
<td>67</td>
</tr>
</tbody>
</table>

* p ≤ 0.002; ** p ≤ 0.0004 (Bonferroni corrected)

**Note.** For volunteers without cancer, marital status was unknown for one, occupation was unknown for two, and first language was unknown for two.
ONCOLOGY VOLUNTEERS

volunteers have a personal history of cancer, and others have family members or friends diagnosed with the disease. Oncology volunteers provide many services, including mentoring, maintaining the library, transporting patients, sharing information, leading self-support groups, welcoming patients at the reception, presenting workshops, making hospital visits, providing bereavement support, conducting research, and maintaining a newsletter. Oncology volunteers are screened, selected, and assigned to roles that correspond to their interests and to the needs of the organization (Remmer, Edgar, & Rapkin, 2001). They receive a 15-hour orientation program as well as ongoing training, support, and educational opportunities.

Participants and Data Collection
The authors included oncology volunteers aged 18 years or older who were fluent in English and who have direct contact with patients with newly diagnosed cancer (e.g., face-to-face and telephone counseling, hospital visits, group support and workshops). Volunteers with a personal cancer diagnosis were labeled peer volunteers. Volunteers who never were diagnosed with cancer were labeled volunteers without cancer. The term oncology volunteers refers to both groups (i.e., peer volunteers and volunteers without cancer). The study protocol was approved by the ethics board of the hospital associated with the cancer support program, and informed consent was obtained from all participants included in the study. Study invitation, questionnaires, and consent forms were mailed to all eligible oncology volunteers. Participants received as many as three reminder telephone calls to complete the study materials. Data collection took place from April to June 2012.

Measures
Demographic information included age, sex, marital status, occupation, first language, ethnic background, education, and personal or family cancer history.
Volunteer work–related information included the specific domains of volunteer work at the cancer support program, the duration of the volunteering, and the average hours of volunteering per week. Volunteer work satisfaction was assessed with one item rated from 1 (not at all satisfied) to 5 (maximum satisfaction).

Compassion was measured using the total score of the 24-item Compassion Toward Others Scale (CS) (Pommier, 2010) addressing kindness versus indifference, common humanity versus disengagement, and mindfulness versus separation (α of the total sample = 0.81).

Self-compassion and self-coldness were assessed with the 26-item Self-Compassion Scale (SCS) (Neff, 2003) addressing self-kindness versus self-judgment, common humanity versus isolation, and mindfulness versus overidentification. In line with research on the psychometric properties of SCS, two composite subscale scores were computed and termed self-compassion for the mean of the three positive subscales and self-coldness for the mean of the three negative subscales (α of self-compassion = 0.87, α of self-coldness = 0.89) (Brenner, Heath, Vogel, & Créde, 2017; Coroiu et al., 2018; Gilbert et al., 2011; Pfattheicher, Geiger, Hartung, Weiss, & Schindler, 2017).

Psychological well-being was assessed with the 22-item Psychological General Well-Being Index (PGWBI) (Dupuy, 1984) capturing anxiety, depression, positive well-being, self-control, general health, and vitality. The PGWBI has good psychometric properties in clinical samples with chronic physical illness and in general population samples (α of total sample = 0.9) (Chassany et al., 2004).

Statistical Analysis
Statistical analyses were performed using IBM SPSS Statistics, version 20.0, and GraphPad Prism. Descriptive analyses were conducted on demographic characteristics and volunteer work–related information, as well as on compassion, self-compassion, self-coldness, and psychological well-being variables. Peer volunteers and volunteers without cancer were compared on outcome measures using nonparametric methods (Mann–Whitney U test and Kruskal–Wallis test, chi-square test, significance levels Bonferroni correction, Cohen’s d effect sizes).

To address the second study objective, study outcomes were compared to published clinical samples and non-student samples with similar demographic and illness-related characteristics using unpaired t tests with Bonferroni corrections for multiple comparisons. Compassion scores were compared with participant data from a Mindful Self-Compassion (MSC) Program (Pommier, 2010) because the authors could not locate a published study with CS in patients with cancer. Self-compassion and self-coldness subscale scores were compared to data from patients with cancer (Crane-Oka et al., 2012; K. Sherman, personal communication, June 7, 2017; Nakamura, Lipschitz, Kuhn, Kinney, & Donaldson, 2013; Przedziecki et al., 2013; R. Crane-Oka, personal

“The mental health of volunteers is not only important for their own well-being, but also for the success of oncology volunteer programs.”
communication, June 14, 2017; Y. Nakamura, personal communication, June 22, 2017) and community samples (Neff, Whittaker, & Karl, 2017). Psychological well-being scores (PGWBI) were compared to data from patients with colorectal cancer and community adults (Chassany et al., 2004).

**Results**

A total of 147 (43%) of 340 eligible oncology volunteers completed questionnaires. The majority of the oncology volunteers were Caucasian women aged 55 years or older with an average of 15.67 years of education (SD = 3.22, range = 7–24) (see Table 1).

<table>
<thead>
<tr>
<th>MEASURE</th>
<th>ONCOLOGY VOLUNTEERS (N = 147)</th>
<th>BREAST CANCER (N = 48)</th>
<th>BREAST CANCER (N = 279)</th>
<th>VARIOUS CANCERS (N = 297)</th>
<th>COMMUNITY ADULTS (N = 1,594)</th>
<th>MSC PROGRAM (N = 24)</th>
<th>COMMUNITY ADULTS (N = 1,453)</th>
<th>COLORECTAL CANCER (N = 321)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>NA*</td>
<td>65.6</td>
<td>53.4</td>
<td>52.6</td>
<td>36</td>
<td>51.2</td>
<td>39.9</td>
<td>63.6</td>
</tr>
<tr>
<td>CS total</td>
<td>4.4</td>
<td>0.5</td>
<td>3.27</td>
<td>0.64**</td>
<td>3.19</td>
<td>0.78**</td>
<td>3</td>
<td>0.76**</td>
</tr>
<tr>
<td>SCS total</td>
<td>3.77</td>
<td>0.65</td>
<td>3.37</td>
<td>0.74*</td>
<td>3.29</td>
<td>0.65**</td>
<td>3.27</td>
<td>0.65**</td>
</tr>
<tr>
<td>Self-compassion subscale</td>
<td>3.71</td>
<td>0.68</td>
<td>3.36</td>
<td>0.73</td>
<td>3.29</td>
<td>0.65**</td>
<td>3.27</td>
<td>0.65**</td>
</tr>
<tr>
<td>Self-coldness subscale</td>
<td>2.29</td>
<td>0.73</td>
<td>3.18</td>
<td>0.93**</td>
<td>3.21</td>
<td>0.77**</td>
<td>2.87</td>
<td>0.97**</td>
</tr>
<tr>
<td>PGWBI total</td>
<td>79.65</td>
<td>10.96</td>
<td>76.87</td>
<td>14.56</td>
<td>89.08</td>
<td>14.62</td>
<td>76.47</td>
<td>13.91</td>
</tr>
<tr>
<td>(No) anxiety</td>
<td>76.87</td>
<td>14.56</td>
<td>76.87</td>
<td>14.56</td>
<td>89.08</td>
<td>14.62</td>
<td>76.47</td>
<td>13.91</td>
</tr>
<tr>
<td>(No) depressed mood</td>
<td>89.08</td>
<td>14.62</td>
<td>89.08</td>
<td>14.62</td>
<td>89.08</td>
<td>14.62</td>
<td>89.08</td>
<td>14.62</td>
</tr>
<tr>
<td>Positive well-being</td>
<td>76.47</td>
<td>13.91</td>
<td>76.47</td>
<td>13.91</td>
<td>76.47</td>
<td>13.91</td>
<td>76.47</td>
<td>13.91</td>
</tr>
</tbody>
</table>

* p ≤ 0.002; ** p ≤ 0.0004 (Bonferroni corrected)

* Crane-Okada et al., 2012; 1–32 years (X = 9.8 years) post–breast cancer diagnosis, all women, 79% Caucasian, 58% without a partner, Mindful Movement Program intervention (baseline assessment)

* Przezdziecki et al., 2013; 88% were more than 2 years post–breast cancer diagnosis, all women, 72% in a committed relationship

* Nakamura et al., 2013; more than 5 months post–active treatment, 77% women, relationship status not reported, baseline data of intervention study for sleep disturbances

* Neff et al., 2013; 18% women, 77% Caucasian, relationship status not reported, recruited from Mechanical Turk

* Neff & Germer, 2013; control group participants of an MSC program (baseline assessment), 72% women, 84% Caucasian, relationship status not reported

* Chassany et al., 2014; community adults were 52% women, 67% in a committed relationship, and patients with colorectal cancer were 58% women, relationship status not reported

* To protect study participants, the authors were only permitted to assess age ranges, as reported in Table 1.

CS—Compassion Toward Others Scale; MSC—Mindful Self-Compassion; NA—not applicable; PGWBI—Psychological General Well-Being Index; SCS—Self-Compassion Scale

Note. All categories except oncology volunteers were comparative samples.

Note. For CS total, scores range from 1–5, with greater scores indicating higher levels of compassion. For SCS total and the SCS self-compassion subscale, scores range from 1–5, with greater scores indicating higher levels of self-compassion. For the SCS self-coldness subscale, scores range from 1–5, with greater scores indicating higher levels of self-coldness. For PGWBI total and subscales, scores range from 0–100, with greater scores indicating higher levels of psychological well-being.
Other ethnic backgrounds included Asian, African American, and Hispanic.

Almost all oncology volunteers had experiences related to cancer. About half of the sample (n = 71, 48%) had a cancer diagnosis (i.e., peer volunteers), and most had a family member (n = 99, 67%) or friend (n = 52, 35%) with cancer. The most common cancer diagnosis among peer volunteers was breast cancer (n = 25, 35%), followed by hematologic cancers (n = 11, 16%). On average, the most recent cancer diagnosis and treatment completion date occurred 10.9 (SD = 6.6, range = 1–32) and 8.25 (SD = 5.58, range = 0–26) years prior to study enrollment, respectively. Participants had been active volunteers at the community cancer support program for an average of six years (X̄ = 6.27, SD = 6.29, range = 0.5–31), volunteering for four hours a week (X̄ = 4.01, SD = 2.59, range = 1–15). Study participants reported high satisfaction with their volunteer work (X̄ = 4.35, SD = 0.68, range = 2–5).

Data Analysis

To provide context, the current findings were compared to those from clinical and community studies. The authors identified one study examining compassion using CS, three studies reporting on self-compassion using SCS, and two studies on psychological well-being using PGWBI. The current authors contacted authors who conducted studies using SCS but did not report subscale scores for self-compassion and self-coldness in their published reports.

Information about these comparative samples (e.g., age, gender, diagnosis, intervention program) is presented in Table 2. No significant differences were found between CS scores in the current study’s volunteer sample and a community sample of participants of an MSC program (Neff & Germer, 2013). The current study’s oncology volunteers reported significantly higher self-compassion subscale scores than breast cancer survivors and community adults did, but they did not differ significantly from baseline data of patients with breast cancer who were enrolled in a Mindful Movement Program intervention. Self-coldness subscale scores were significantly lower in oncology volunteers than in all identifiable samples of patients with breast cancer and community adults. The current study participants reported more psychological well-being (PGWBI) than a general population sample and a sample of patients with colorectal cancer.

When comparing volunteers without cancer to peer volunteers, the latter were significantly older and were more likely to...
act as mentors for patients with cancer and their loved ones. In addition, peer volunteers were less satisfied with their volunteering job (X = 4.2, SD = 0.69 versus X = 4.49, SD = 0.64; U[df] = 2,067.5 [145], p < 0.01) and had served as volunteers for a longer time (X = 7.61 years, SD = 6.01 versus X = 5.05 years, SD = 6.34; U[df] = 1,699 [141], p < 0.001). Finally, peer volunteers did not differ from volunteers without cancer regarding compassion, self-compassion, and self-coldness. However, peer volunteers felt less psychological well-being and reported worse general health (see Table 3).

Discussion
The purpose of the current study was to examine demographics, work-related characteristics, and levels of compassion, self-compassion, self-coldness, and psychological well-being in volunteers at an oncology support program consisting of individuals with and without a personal history of cancer. The authors also compared the sample's CS, SCS, and PWBGI scores to published findings for patients with cancer and the general population (i.e., North American community samples).

The results suggest that volunteer work within a community cancer support program can be experienced as highly satisfactory, with participants in the current study completing, on average, more than six years of service. Mirrieeles et al. (2017) evaluated a peer support program for patients with breast cancer and also found high satisfaction and a sense of accomplishment in support providers. Satisfaction with the volunteer experience in the current sample also was similar to that in a study by Remmer et al. (2001), who examined the same cancer support program more than 16 years ago. Despite overall high satisfaction, peer volunteers felt significantly less satisfied with their volunteer work when compared to their colleagues without a personal history of cancer. This difference may be attributed to the nature of the volunteer work, but it is also possible that volunteers without cancer can keep some emotional distance from their work while feeling less burdened or less discouraged by the limitations of the support they can provide.

Overall, the current sample of oncology volunteers fared well when compared to community samples and patients with cancer. Study participants reported less anxiety and depression and more psychological well-being, self-control, and vitality than a large community sample (Dupuy, 1984). Although both samples included similar proportions of married or cohabiting individuals (n = 95.6% of oncology volunteers and n = 974.67% of community adults), the older age of the study participants may have contributed to these findings; 92 participants (63%) were at least 55 years of age, and the mean age in the comparison sample was 40 years. Higher life satisfaction and better psychological well-being have been reported in studies of participants from Western countries (Carr, Friedman, Corman, & Schwartz, 2014; Steptoe, Deaton, & Stone, 2015; Stone, Schwartz, Broderick, & Deaton, 2010; Van Willigen, 2000). Oncology volunteers (Wittenberg et al., 2010), older adults (Steptoe et al., 2015; Van Willigen, 2000), and those who are married (Carr et al., 2014; Wittenberg et al., 2010). In addition, participants in the current study experienced less self-coldness and more self-compassion than clinical and nonclinical samples, regardless of whether they had a personal history of cancer. However, the cross-sectional design does not allow for determining causality. The current study’s results suggest that oncology volunteers are a self-selected group of individuals who dealt well enough with adversity related to their cancer experience to be capable of lending a helping hand to others in need. It is also possible that providing support to others may further enhance one’s experience of meaning in life (Frazier et al., 2013) and increase feelings of compassion and self-compassion (Konrath & Brown, 2013). Finally, volunteer work can provide additional opportunities for sharing of emotions, social support, and cognitive and emotional processing of the cancer experience, and it can promote posttraumatic growth (Calhoun & Tedeschi, 2006; Cohen & Numa, 2011).

When comparing volunteers with and without a history of cancer, the participants did not differ significantly regarding compassion, self-compassion, self-coldness, and several indicators of psychological well-being. However, cancer survivors in the current sample reported worse general health than their colleagues without a personal history of cancer. This result is not surprising given the older age of the peer volunteer group and their medical history. Still, the lower overall psychological well-being of peer volunteers may not have been exclusively driven by age and medical history, because peer volunteers also reported less positive well-being, less vitality, and more symptoms of anxiety (after correcting for type I error, these differences were not statistically significant). Because most cancer survivors live with a lifelong risk of recurrence associated with psychological distress, the current findings speak to the need to be sensitive to potential support needs of those who volunteer to provide support.

Limitations and Future Directions
The results of the current study may be limited by self-selection bias, given the study participation rate of 44%. Oncology volunteers who were less satisfied and experienced lower compassion, self-compassion, or psychological well-being may not have responded to the study invitation. Nevertheless, this is one of the few studies addressing psychological characteristics of support providers in the context of cancer care. Future research may benefit from also assessing the perceived quality and quantity of

IMPLICATIONS FOR PRACTICE
- Include peer volunteers in oncology volunteer programs providing supportive cancer care.
- Monitor peer volunteer staff for signs of distress, such as feeling tense, discouraged, self-judgmental, and concerned about one’s own health.
- Incorporate training aimed at self-care and recognizing signs of compassion fatigue and burnout.
the peer support in support recipients compared to providers; it has been shown that the psychological state of both parties could influence the effectiveness and perception of the support interaction (Embuleiniya et al., 2013; Giese-Davis et al., 2006; Pistrang et al., 2013; Pistrang, Solomons, & Barker, 1999). It would also be valuable to extend this work toward examining the satisfaction, well-being, and compassion of healthcare professionals, particularly nurses who interact with patients most closely and are major supportive care providers in oncology. This may further the authors’ insight into the challenges and protective factors across different groups that are facing the task of supporting individuals and families touched by a life-threatening medical condition.

**Implications for Practice**

As leaders in hospitals and care centers, nurses play an important role in the training and management of oncology volunteers. The results of the current study confirm the importance of proper psychoeducation and training for oncology volunteers. Volunteer program coordinators and nurses working alongside peer volunteers can be particularly sensitive to the well-being of oncology volunteers with a personal history of cancer. Training for these positions may include content specifically tailored to the potential needs of peer volunteers, such as modules on self-care. Exercises aimed at increasing self-compassion and decreasing self-coldness may be appropriate because this may be a protective factor for oncology volunteers (Körner et al., 2015). Continued formal or informal monitoring of peer volunteer well-being also may prove beneficial. Nurses are an integral part of supportive cancer care and could play a leading role in this movement. Oncology nurses must advance care while being sensitive to volunteers’ well-being, as well as practicing self-care and self-compassion (see Figure 1) and addressing their own signs of early burnout.

**Conclusion**

Oncology volunteers, including those with a personal history of cancer, indicated higher levels of self-compassion and psychological well-being than patients with cancer and even community samples. This speaks for including peer volunteers when expanding volunteer programs as part of supportive care services in oncology and when establishing such programs where they do not exist yet.

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QUESTIONS FOR DISCUSSION

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Journal club programs can help to increase your ability to evaluate the literature and translate those research findings to clinical practice, education, administration, and research. Use the following questions to start the discussion at your next journal club meeting.

Does this study’s results (measuring compassion, self-compassion, self-coldness, and psychological well-being) match your perceptions of oncology volunteers that support your practice?

Would you keep or revise this study design to better understand and support oncology volunteers supporting your practice?

If revising the design, what would you revise?

For your facility or practice setting, how would you design a training program for oncology volunteers, focusing on self-care and recognition of compassion fatigue and burnout?

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