Published online 2020 August 18.

Research Article

The Efficacy of a Peer-Led Versus Nurse-Led Self-Management on the Subjective Well-Being of Patients with Multiple Sclerosis: A Randomized Clinical Trial

Maryam Azizi¹, Najmeh Javan¹, Seyed Mohammad Mousavi Mirzaei² and Seyyed Abolfazl Vagharseyyedin ¹,*

¹Department of Nursing, Faculty of Nursing and Midwifery, Birjand University of Medical Sciences, Birjand, Iran ²School of Medicine, Birjand University of Medical Sciences, Birjand, Iran

^{*} *Corresponding author*: Nursing and Midwifery College, Birjand University of Medical Sciences, Ayatollah Ghaffari Avenue, Postal Code: 9717853577, Birjand, Iran. Tel: +98-5632395000, Fax: +98-5632430076, Email: waghars@bums.ac.ir

Received 2020 March 28; Revised 2020 June 02; Accepted 2020 June 24.

Abstract

Background: Subjective well-being (SWB) can be a buffer against mental health problems. Patients with multiple sclerosis (MS) tend to experience low levels of subjective well-being.

Objectives: The two objectives were (1) to examine whether peer-led and nurse-led self-management training can improve SWB of patients with MS and (2) to compare the effect of these two interventions on SWB of patients with MS.

Methods: In this trial, 81 patients were randomly assigned to either nurse-led (n = 41) or peer-led (n = 40) groups. Both groups participated in six weekly-based 1.5-hr self-management sessions. The cognitive and affective components of SWB were evaluated using the Satisfaction with Life Scale and the Scale for Positive and Negative Experience, respectively.

Results: Life satisfaction increased in both groups over time (P < 0.001), although it was not significantly different between the two groups (P > 0.05). Positive and negative experiences increased in both groups over time (P < 0.05), but the increase in positive experience in the peer-led group was significantly larger than those in the nurse-led group (P < 0.05).

Conclusions: Peer-led and nurse-led self-management training, are feasible tools to enhance the cognitive component of SWB, i.e., life satisfaction. Considering the affective component of SWB, positive experience in the peer-led group improved better than in the nurse-led group over time. Therefore, if healthcare providers improve the SWB of patients with relapsing-remitting MS, the self-management intervention led by peers is preferred than nurse-led self-management training.

Keywords: Multiple Sclerosis, Nurse-Led Intervention, Peer-Led Intervention, Self-Management, Subjective Well-Being

1. Background

Multiple sclerosis (MS) is the leading cause of nontraumatic neurologic disability to affect young adults in many countries. The global prevalence of 33 per 100,000 for MS was estimated in 2013, while the prevalence was 30 per 100,000 in 2008, suggesting an increasing trend for the disease (1). A meta-analysis in 2019 reported a prevalence rate of 29.3 per 100,000 for MS in Iran (2). Although the clinical spectrum of MS is diverse, the majority of MS cases are classified as relapsing-remitting, where acute attacks or relapses are followed by partial or full periods of recovery (remission) (3).

MS and its complications can jeopardize the mental health of the patients with MS. Therefore, psychological disturbances such as depression, stress, and anxiety in these patients are anticipatable (4, 5). In line with, subjective well-being, as a buffer against mental health problems, has been frequently addressed in patients with chronic conditions such as MS (6, 7). Subjective well-being (SWB) refers to an individual's evaluation of his/her life. An evaluation of this type is both cognitive and affective. The cognitive component of SWB denotes the global judgment one makes of his/her life (i.e., life satisfaction), while the affective component refers to one's experience of many pleasant emotions and moods (namely, positive affect) and few unpleasant emotions and moods (namely, negative affect) (8).

A substantial number of patients with MS experience low levels of subjective well-being. MS-related symptoms and signs are considered as contributors to decreased SWB among patients with MS (7). Thus, it is expected that

Copyright © 2020, Jundishapur Journal of Chronic Disease Care. This is an open-access article distributed under the terms of the Creative Commons Attribution-NonCommercial 4.0 International License (http://creativecommons.org/licenses/by-nc/4.0/) which permits copy and redistribute the material just in noncommercial usages, provided the original work is properly cited. an improvement of self-management skills can lead to an enhanced SWB by enabling patients to better control of these signs and symptoms. It has been shown that self-management training for patients with chronic conditions, including Parkinson's disease and epilepsy, has led to improved SWB (9, 10).

Self-management is described as the ability of active and responsible individuals to live with the medical, role, and emotional consequences of their chronic condition within their social network in collaboration with healthcare providers (11). Self-management interventions can be categorized into two broad groups: peerled self-management interventions and nurse-led selfmanagement interventions. In peer-led interventions, the peer leader has conditions similar to those of other patients within a group and is expected to provide relevant and meaningful information, act as a role model, and guide the peers (12, 13). However, in nurse-led selfmanagement interventions, a nurse presents a set of predesigned educational materials.

Both self-management mentioned above interventions have shown promising results, including increased improved quality of life, decreased depression, and enhanced psychological well-being in patients with chronic conditions (14-16). Nevertheless, there is no research, as yet, to investigate the effects of these two methods on SWB of patients with MS.

2. Objectives

Therefore, the current study was designed with two aims: (1) to examine whether peer-led and nurse-led selfmanagement training can improve SWB of patients with MS and (2) to compare the effect of these two interventions on SWB of patients with MS.

3. Methods

This randomized clinical trial was conducted at the Comprehensive Center for the Treatment of Special Diseases affiliated to Birjand University of Medical Sciences (BUMS) in 2018.

Taking a type I error = 0.05 and a type II error of 0.2, and considering the results of previous research (17) (M1 = 215.77, M2 = 232.57, S1 = 25.55, and S2 = 25.62), the desired sample size was calculated as 37 subjects per group. An attrition rate of 15% was factored, whereby the sample size increased to 42 subjects per group.

Potential patients were approached during their usual clinic visits at the Comprehensive Center for the Treatment of Special Diseases. Patients were eligible if they had the inclusion criteria, i.e., ability to read and write in Persian; willingness to participate in the study; age between 20 to 45 years; a minimum one-year history of MS confirmed by the neurologist; high likelihood of continuing to present MS clinically as remitting-relapsing during the study period based on the neurologist's opinion; obtaining a score between 0 and 5.5 on the Expanded Disability Status Scale (EDSS); and lack of any acute or chronic physical, mental, and psychiatric disorders that interfere with self-management. Failing to attend more than two sessions, being hospitalized during the intervention, or unwillingness to continue participation were considered as exclusion criteria.

The study was explained to eligible patients, and written informed consent forms were signed by the interested patients.

Initially, 120 patients were assessed for eligibility of whom 84 were selected. A demographics form, the Satisfaction with Life Scale (SWLS), and the Scale for Positive and Negative Experience (SPANE) were completed for all the participants by a researcher assistant. The participants were subsequently assigned to the study groups via simple randomization using computer-generated random numbers. For this purpose, 84 selected participants were assigned a number from 1 to 84 of which a list of 42 random numbers was created. The participants with these random numbers were allocated to the peer-led group, and the other participants were assigned to the nurse-led group (Figure 1). Recruitment was performed from March to May 2018.

3.1. Intervention

In the nursing-led group, the patients were divided into two sub-groups of 10 and two sub-groups of 11 members. Afterward, the prepared self-management contents (Table 1) were presented by the same experienced nurse in six sessions, each lasted for about 1.5 hours held every week. The self-management intervention contents were prepared using credible sources, including books (18-20) and a valid website (21), and were subsequently confirmed by two neurologists independently.

In the peer-led group, the leader was selected according to the following criteria: willingness to play the assigned tasks, acceptable social communication skills, ability to manage the sessions, a minimum five-year history of MS, and a proper educational level (Bachelor's degree or above). Considering these criteria and according to the neurologist's suggestion, a patient from among the MS patients referring to the Comprehensive Center for the Treatment of Special Diseases was appointed as the leader of the group.



Figure 1. The Consort diagram of the study

Three training sessions were held for the leader, comprising of the following issues: an introduction to the objectives and nature of the peer education sessions, essential points to communicate with group members, useful strategies to help keep the discussion focused on the presented topics, and a brief review of the prepared contents. These sessions were held in the presence of the two researchers under the supervision of the neurologist. In the next step, the participants in the peer-led group also were divided into two sub-groups of 10 and two subgroups of 11 members. The prepared self-management contents (Table 1) were presented by the peer leader for each group through six weekly-held sessions, each lasting for approximately 1.5 hours.

In both the nurse-led and peer-led groups, participants were assigned into sub-groups to have maximum homo-

Session	Detail
Session 1	Familiarity of the researchers with the participants and the participants with each other, gaining the trust of the participants, determining the roles and responsibilities of the participants and researchers, overviewing self-management and its techniques
Session 2	Overviewing MS, its course, and therapeutic options; introducing self-management strategies for MS-induced fatigue
Session 3	Providing self-management strategies for establishing a balance between activities and rest
Session 4	Providing self-management strategies for managing ataxia in MS
Session 5	Providing self-management strategies for stress, anxiety, and depression
Session 6	Providing self-management advice on nutrition, summarizing the sessions, clarifying potential ambiguities raised by the participants, completing research instruments

geneity in terms of educational level. Also, one of the researchers was present as a supervisor in both groups. Both interventions were conducted over a period of 6 weeks during the months from July to August 2018.

From the second week up to two months after the intervention completed, a weekly telephone follow-up was conducted on the implementation of the self-management practice. At the end of the sessions, the provided contents were available to all of the patients.

Patients registered at the Comprehensive Center for the Treatment of Special Diseases usually attended the center individually. In order to reduce the contact between participants and minimize the potential contamination effects, the peer-led intervention was conducted in the morning on Saturdays through Tuesdays. For participants in the four nurse-led groups, the intervention was implemented on the same days, yet in the evening. Thus, contamination effects can be assumed to be negligible.

The study scales were completed for the participants in the two groups both immediately and two months after the intervention by the same researcher assistant who completed the scales at baseline.

3.2. Study Measurements

The participants' demographic characteristics, including age, gender, marital status, educational status, occupation, insurance type, and MS duration, were collected using a demographics form.

The EDSS was completed by the neurologist to evaluate the degree of the participants' disability. This instrument is on an ordinal scale ranging from 0 indicating normal neurological status to 10, indicating death due to MS. It provides a final score, which reflects the eight functional system scores (i.e., pyramidal, cerebellar, brain stem, sensory, bowel/bladder, visual, cerebral and others) (22).

The cognitive component of SWB (i.e., life satisfaction) was assessed using the SWLS, a five-item self-report measure that assesses one's satisfaction with life and accomplishments of essential goals in life. The participants were asked to respond to items on a 7-point Likert scale ranging from 1 = strongly disagree to 7 = strongly agree. The total score of SWLS ranges from 5 to 35 (23). Previous research on patients with MS has reported a desirable reliability coefficient for the SWLS, with a Cronbach's alpha of 0.84 (7). In Iran, several studies have shown good reliability coefficients for this scale (24). In the present study, the alpha coefficient for the SWLS was 0.82.

The affective component of SWB was measured by SPANE. This scale is divided into two subscales. One subscale assesses positive feelings (SPANE-P), and the other assesses negative feelings (SPANE-N). Each sub-scale contains six items that are scored on a five-point scale ranging from 1 (very rarely or never) to 5 (very often or always). The SPANE-P and SPANE-N scores are the sum of the scores given to their respective six items, and the scores can range from 6 to 30. This study used the Persian version of SPANE, which is available on the official website of Dr. Diener, the original developer of this tool. In previous research, this scale had acceptable reliability (25). Regarding reliability, the Cronbach's alpha coefficients were 0.8 and 0.9 for the SPANE-P and SPANE-N sub-scales, respectively.

3.3. Ethical Consideration

The present study was approved by the Ethics Committee of Birjand University of Medical Sciences (BUMS) (code: Ir.bums.REC.1397.190). Also, the researchers obtained permission from BUMS and received a clinical trial code (IRCT20190305042926N1) before the eligibility of the patients with MS registered in the center was assessed.

3.4. Data Analysis

The collected data were analyzed with SPSS v.16 software (IBM Company, New York, USA). The descriptive statistics, including mean, standard deviation, and frequency, were used to describe the participants' demographic characteristics and their responses on study measures.

The data concerning gender, marital status, educational level, occupation, and insurance type were compared between the two groups using the chi-square test or Fisher's exact test. The independent t-test was applied to analyze the difference between the two groups in terms of age and the duration of MS. The EDSS scores were compared using the Mann-Whitney U test.

There was a non-normal distribution of life satisfaction mean scores in the peer-led group before the intervention and in the nurse-led group immediately after the intervention. In the nurse-led group, also, the negative experience mean scores before the intervention, and two months after the intervention were non-normally distributed (Shapiro-Wilk test, P < 0.05). The main variables in other time points were normally distributed.

In both groups, life satisfaction mean scores over time were evaluated using the Friedman test with post hoc Wilcoxon signed-rank test. The Mann-Whitney test was employed to do between-group comparisons.

Positive experience mean scores were compared between the two groups over time using repeated-measures ANOVA followed by post hoc Bonferroni correction.

In the nurse-led group, the mean score of negative experience over time was assessed using the Friedman test with post hoc Wilcoxon signed-rank test. In the peerled group, a repeated-measures ANOVA followed by post hoc Bonferroni correction was used to compare withingroup changes of negative experience over time. Finally, between-group comparisons were performed with the independent *t*-test or Mann-Whitney test.

4. Results

A total of 81 patients with MS participated in this study. The age means of the patients in the peer-led, and nurseled self-management education groups were 37.7 (95% CI: 34.89 - 39.51) and 35.61 (95% CI: 32.71 - 38.51) years, respectively. There was no significant difference in the mean age between the two groups (P = 0.39). The EDSS mean scores of the peer-led and nurse-led groups were 2.35 (95% CI: 1.99 - 2.7) and 2.36 (95% CI: 1.98 - 2.75), respectively, with no significant difference according to the Mann-Whitney U test (P = 0.95). As shown in Table 2, the differences between the two groups did not reach statistical significance concerning other demographic characteristics.

Before the intervention, there was no significant difference in the mean scores of life satisfaction, positive experience, and negative experience between the two groups (P > 0.05) (Table 3).

As expressed in Table 4, the Friedman test revealed a significant increase in the scores of life satisfaction in both groups in the course of time (P < 0.001). The Wilcoxon signed-rank test showed that in both groups, life satisfaction increased significantly over time from before intervention to two months after intervention (P < 0.05). However, between-group comparisons showed no significant differences between the two groups in terms of life satisfaction mean scores at the three-time points (P > 0.05).

A repeated-measures ANOVA showed that positive experience mean scores were different between groups (F = 3.33, P < 0.001) and within groups at different time points

Characteristic Peer-Led Group (N = 40) Nurse-Led Group (N = 41) P Val (N = 41) Gender (%) 0.82 0.82 Male 8 (20) 9 (21.95) 0.82 Female 32 (80) 32 (78.05) 0.82	ue 2 ^a
Gender (%) 0.82 Male 8 (20) 9 (21.95) Female 32 (80) 32 (78.05)	a } ^a
Male 8 (20) 9 (21.95) Female 32 (80) 32 (78.05)	} ^a
Female 32 (80) 32 (78.05)	} ^a
	3 ^a
Marital status (%) 0.08	
Single 4 (10) 10 (24.4)	
Married 36 (90) 31 (75.6)	
Educational 0.6 ² status (%)	a
Primary 11 (27.5) 9 (22) and secondary school	
High 12 (30) 10 (24.4) school	
University 17 (42.5) 22 (53.7)	
Occupation (%) 0.24	b
Housewife 23 (57.5) 16 (39)	
Retired 1(2.5) 2(4.9)	
Unem- 5 (12.5) 9 (22) ployed	
Self- 0 (0) 3 (7.3) employed	
Employee 11 (27.5) 11 (26.8)	
Duration of MS 5.67 (95% CI: 4.54 - 6.75 (95% CI: 5.3 - 0.24 (years) 6.8) 8.2)	¢
Insurance type 0.15 (%)	ь
Social 17 (42.5) 25 (61) welfare	
Armed 4 (10) 1 (2.4) forces	
Health and 19 (47.5) 15 (36.6) medical service	

^aThe chi-square test

^bThe Fisher's exact test

^cThe independent *t* test

(F = 3.24, P < 0.001). Furthermore, a significant time \times group interaction was found (P = 0.01), indicating that increases in positive experience over time in the peer-led group were significantly greater than those for the nurse-led group (Table 5). In both groups, the participants reported a significantly higher positive experience immediately and two months after the intervention compared to before intervention (P < 0.05). Also, positive experience in the two groups increased from immediately after the intervention to two months after the intervention; however, the change was not significant (P > 0.05).

Variable		Peer-Led Group (N = 40)	Nurse-Led Group (N = 41)	z or t Values	P Value
Life satisfaction [Mean (95% CI)]					
Befo	ore intervention	18.55 (16.83 - 20.26)	16.87 (14.87 - 18.88)	-1.5	0.13 ^a
Imm	nediately after intervention	21.2 (18.88 - 23.51)	18.58 (17.05 - 20.12)	-1.89	0.06 ^a
Two	months after intervention	22.22 (20.1 - 24.34)	19.85 (18.48 - 21.22)	1.9	0.06 ^b
Positive experience [Mean (95% CI)]					
Befo	re intervention	16.5 (14.96 - 18.03)	14.75 (13.82 - 15.68)	1.96	0.54^{b}
Imm	nediately after intervention	19.82 (18.37 - 21.27)	15.8 (14.87 - 16.7)	4.72	<0.001 ^b
Two	months after intervention	19.77 (18.54 - 21.00)	16.36 (15.54 - 17.19)	4.64	<0.001 ^b
Negative experience [Mean (95% CI)]					
Befo	pre intervention	17.05 (15.79 - 18.3)	16.68 (15.55 - 17.8)	-0.13	0.89 ^a
Imm	nediately after intervention	15.1(13.88 - 16.31)	15.04 (14.04 - 16.05)	0.06	0.94 ^b
Two	months after intervention	14.2 (13.1 - 15.29)	13.8 (12.89 - 14.71)	-0.3	0.75 ^a

Table 3. Comparison of the Mean Life Satisfaction, Positive Experience, and Negative Exprience Scores of Peer-Led and Nurse-Led Self-Management Education in Three Time Points

^aMann-Whitney U test

^bIndependent t-test

 Table 4.
 Within- and Between-Group Comparisons Respecting Life Satisfaction

 Scores of Peer-Led and Nurse-Led Self-Management Education in Three Time Points

Time	Peer-led Group, (N = 40), Mean (95% CI)	Nurse-Led Group, (N = 41), Mean (95% CI)	Test Results ^a
Before intervention	18.55 (16.83 - 20.26)	16.87 (14.87 - 18.88)	0.13 ^a
Immediately after intervention	21.2 (18.88 - 23.51)	18.58 (17.05 - 20.12)	0.06 ^a
Two months after intervention	22.22 (20.1 - 24.34)	19.85 (18.48 - 21.22)	0.06 ^b
Test results ^c	$\chi^2 = 39.96; P < 0.001$	$\chi^2 = 23.52; P < 0.001$	-

^aMann-Whitney U test

^bIndependent t-test

^cFriedman test

Table 5. Repeated Measures ANOVA of Positive Experience in Peer-Led and Nurse-Led Self-Management Education Groups

Variable	Source	F Value	P Value
Positive experience	Group	3.33	< 0.001
	Time	33.24	< 0.001
	Group \times Time	0.69	0.01

A significant decrease was found in the mean score of negative experience in the nurse-led group over time (χ^2 = 44.27; P < 0.001). The Wilcoxon signed-rank test indicated that negative experience decreased significantly from before the intervention to two months after intervention (P < 0.05). The repeated measures ANOVA determined that the

mean score of negative experience in the peer-led group also decreased significantly over time (F (2, 78) = 30.85, P < 0.001). Post hoc tests using the Bonferroni correction revealed a significant decrease in negative experience from before intervention to immediately after the intervention as well as from before the intervention to two months after the intervention (P < 0.001). A reduction was also found in negative experience from immediately after the intervention to two months after the intervention; however, it was not significant (P = 0.06).

The results of between-group comparisons with respect to negative experience revealed no significant differences before the intervention, immediately after the intervention, and two months after the intervention (P > 0.05).

5. Discussion

The purpose of the current study was to compare the effect of a peer-led versus a nurse-led self-management intervention on SWB in patients with MS.

In this study, both peer-led and nurse-led selfmanagement interventions resulted in a significant improvement of the cognitive component of subjective well-being, i.e., life satisfaction. Previous studies have proved the effect of self-management on the participants' life satisfaction. Sahar et al. (26), for example, reported that implementation of a 12-week self-help intervention for a sample of older people in Jakarta led to significantly higher life satisfaction in the intervention group than the control group (27). In other research, 60 Iranian patients with epilepsy participated in a four-session selfmanagement training program. One month after the intervention, the mean scores of life satisfaction and its dimensions (including mental health, social functioning, worry about seizures, energy-fatigue, cognitive functioning, and medication effects) in the intervention group were significantly higher than in the control group (11). However, in Abbasi et al. (27) study, a self-management education program consisting of three sessions and a three-month follow-up for 60 people with chronic heart failure (CHF) did not result in a significant difference in the life satisfaction of intervention and control groups (28). These findings collectively suggest that the impact of self-management interventions on an individuals' life satisfaction may be informed by factors such as the disease nature and disease-related difficulties. In this line, in Ryan et al. (7) study, disease severity was a predictor for life satisfaction in patients with MS (8).

The impact of the self-management interventions in enhancing life satisfaction may be partially attributed to the potential of these interventions in improving patients' capacity to manage MS signs and symptoms more appropriately, which could have led to an improved feeling of health. Such feeling, in turn, can lead to higher life satisfaction. The Common-Sense Model of Self-Regulation (29) can help explain this finding. According to this model, when people face threats to their health, they create mental representations (beliefs) of illness threats. An individual's beliefs concerning illness threats include issues such as the disease identity, cause, controllability, duration/timeline, and consequences. Patients' beliefs about these five dimensions influence their bio-psycho-social adjustment (29, 30). In a meta-analysis of studies adopting the Common-Sense Model of Self-Regulation, Hagger et al. (28) found that illness presentations had a direct effect on disease outcomes such as patient's well-being (29).

The results of this study showed that there was no significant difference between the two groups over time, despite the improvement in life satisfaction. The two interventions may have in-built advantages of one over the other. For example, the possible empathy created among the participants of the peer-led group cab be an advantage (31, 32). The nurse-led intervention, on the other hand, is characterized by a more robust and structured education. Nevertheless, these advantages did not result in a significant difference between the two groups with respect to life satisfaction.

The results also indicated that the affective component of subjective well-being, i.e., positive experience and negative experience, improved significantly over time in both groups. Alongside this, Benzo et al. (32) reported a significant positive correlation between self-management ability and positive affect and a significantly negative relationship between self-management ability and negative affect in patients with chronic obstructive pulmonary disease (COPD) (33). Based on the Common-Sense Model of Self-Regulation, researchers deduce that in patients with chronic diseases, factors such as a stronger illness identity and the perception that MS has substantial adverse effects on an individual's emotions are key predictors of psychological distress. Conversely, the more a patient feels that she/he understands the illness and can take actions to control it, the lower her/his distress will be.

In this study, positive experience in the peer-led group improved better than in the nurse-led group over time. This finding may be related to the peer-led group's benefits, such as the sense of empathy created between the peer leader and the members and, hence, the more intimate relationship established with him/her (31, 32). This can lead to a stronger social network and thus provide more effective social support, including more emotional support among peer-group members compared with the nurse-led group.

Notably, in the two groups, positive experience increased from immediately to two months after the intervention; however, it was not significant. In the peer-led group, also, the improvement of negative experience from immediately to two months after the intervention was not significant. Therefore, there is a need for measures to strengthen the long-term impact of the intervention. One potential way may be to extend the intervention period. To sustain the changes, it may also be helpful to hold monthly booster meetings after the six weekly training sessions.

To the best of our knowledge, this is the first study to examine the effect of peer-led versus nurse-led selfmanagement on SWB of patients with MS. In the current research, interventions were delivered in a real-world setting by a usual provider or lay peer educator. Furthermore, as evidenced by the low rate of attrition and the high rate of session attendance, the interventions were acceptable and meaningful to patients. Thus, despite the relatively small changes in the mean scores of the main variables from baseline to the follow-up phase, these changes can be assumed as clinically beneficial.

This study has some implications for patients with MS, health care providers, and clinical managers. First of all, the importance of self-management should be emphasized to patients with relapsing-remitting MS. Also, if healthcare providers are to improve the SWB of these patients, the self-management intervention led by peers can be a useful tool. Finally, clinical managers need to support healthcare professionals who are interested in peerled self-management interventions.

However, the study has a few limitations that pose cau-

tion in generalizing the results to other MS populations. First, the participants comprised only of patients with MS of the relapsing-remitting type. Secondly, the sample was selected from a single treatment center. Future studies can be designed to examine the impact of these two selfmanagement approaches on other psychological factors such as psychological well-being and the level of hope in patients with MS.

5.1. Conclusion

Peer-led and nurse-led self-management training are feasible tools to enhance the cognitive component of SWB, i.e., life satisfaction. Considering the affective component of SWB, positive experience in the peer-led group improved better than in the nurse-led group over time. Therefore, the application of the peer-led self-management program to improve SWB in patients with relapsing-remitting MS is more effective than the nurse-led approach.

Acknowledgments

We express our gratitude to all the patients for their participation in this study. SAV conceived the initial idea for the study and helped to study design, statistical analysis, and draft the manuscript. NJ contributed to drafting the manuscript and revising the manuscript. MA helped to study design, draft the manuscript, and data collection. SMMM helped to intervention design and to draft the manuscript. All authors read and approved the final manuscript.

Footnotes

Authors' Contribution: SAV conceived the initial idea for the study and helped to study design, statistical analysis, and draft the manuscript. NJ contributed to drafting the manuscript and revising the manuscript. MA helped to study design, draft the manuscript, and data collection. SMMM helped to intervention design and to draft the manuscript. All authors read and approved the final manuscript.

ClinicalTrialRegistrationCode:IRCT20190305042926N1.

Conflict of Interests: The authors declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

Ethical Approval: This study was approved by the Ethics Committee of Birjand University of Medical Sciences (Ir.bums.REC.1397.190).

Funding/Support: This research was funded by BUMS (Grant number: 455610).

Informed Consent: The study was explained to eligible patients, and written informed consent forms were signed by the interested patients.

References

- Browne P, Chandraratna D, Angood C, Tremlett H, Baker C, Taylor BV, et al. Atlas of Multiple Sclerosis 2013: A growing global problem with widespread inequity. *Neurology*. 2014;83(11):1022–4. doi: 10.1212/WNL.0000000000000768. [PubMed: 25200713]. [PubMed Central: PMC4162299].
- Azami M, YektaKooshali MH, Shohani M, Khorshidi A, Mahmudi L. Correction: Epidemiology of multiple sclerosis in Iran: A systematic review and meta-analysis. *PLoS One*. 2019;**14**(7). e0219466. doi: 10.1371/journal.pone.0219466. [PubMed: 31287842]. [PubMed Central: PMC6615610].
- Yamout BI, Assaad W, Tamim H, Mrabet S, Goueider R. Epidemiology and phenotypes of multiple sclerosis in the Middle East North Africa (MENA) region. *Mult Scler J Exp Transl Clin.* 2020;6(1):2.0552173198419E+15. doi: 10.1177/2055217319841881. [PubMed: 31984137]. [PubMed Central: PMC6961141].
- Chertcoff A, Curbelo MC, Bauer J, Ferrandina F, Martinez A, Steinberg J, et al. Anxiety in Argentinian patients with multiple sclerosis: Prevalence and associated factors. *Mult Scler Relat Disord*. 2020;41:102042. doi: 10.1016/j.msard.2020.102042. [PubMed: 32163755].
- Ploughman M, Wallack EM, Chatterjee T, Kirkland MC, Curtis ME, Health L, et al. Under-treated depression negatively impacts lifestyle behaviors, participation and health-related quality of life among older people with multiple sclerosis. *Mult Scler Relat Disord*. 2020;40:101919. doi: 10.1016/j.msard.2019.101919. [PubMed: 31951860].
- Bassi M, Falautano M, Cilia S, Goretti B, Grobberio M, Pattini M, et al. Illness Perception and Well-Being Among Persons with Multiple Sclerosis and Their Caregivers. *J Clin Psychol Med Settings*. 2016;23(1):33–52. doi: 10.1007/s10880-015-9425-8. [PubMed: 26216661].
- Ryan KA, Rapport LJ, Sherman TE, Hanks RA, Lisak R, Khan O. Predictors of subjective well-being among individuals with multiple sclerosis. *Clin Neuropsychol.* 2007;21(2):239–62. doi: 10.1080/13854040600582460. [PubMed: 17455016].
- Diener E. Subjective well-being. The science of happiness and a proposal for a national index. *Am Psychol.* 2000;55(1):34–43. [PubMed: 11392863].
- Hellqvist C, Dizdar N, Hagell P, Bertero C, Sund-Levander M. Improving self-management for persons with Parkinson's disease through education focusing on management of daily life: Patients' and relatives' experience of the Swedish National Parkinson School. J Clin Nurs. 2018;27(19-20):3719–28. doi: 10.1111/jocn.14522. [PubMed: 29782061].
- Yadegary MA, Maemodan FG, Nayeri ND, Ghanjekhanlo A. The effect of self-management training on health-related quality of life in patients with epilepsy. *Epilepsy Behav.* 2015;**50**:108–12. doi: 10.1016/j.yebeh.2015.04.051. [PubMed: 26232570].
- Van de Velde D, De Zutter F, Satink T, Costa U, Janquart S, Senn D, et al. Delineating the concept of self-management in chronic conditions: a concept analysis. *BMJ Open*. 2019;9(7). e027775. doi: 10.1136/bmjopen-2018-027775. [PubMed: 31315862]. [PubMed Central: PMC6661649].
- Scott SD, Hirschinger LE, Cox KR, McCoig M, Hahn-Cover K, Epperly KM, et al. Caring for Our Own: Deploying a Systemwide Second Victim Rapid Response Team. *The Joint Commission Journal on Quality and Patient Safety*. 2010;**36**(5):233–40. doi: 10.1016/s1553-7250(10)36038-7.
- Fisher EB, Boothroyd RI, Coufal MM, Baumann LC, Mbanya JC, Rotheram-Borus MJ, et al. Peer support for self-management of diabetes improved outcomes in international settings. *Health Aff* (*Millwood*). 2012;**31**(1):130–9. doi: 10.1377/hlthaff.2011.0914. [PubMed: 22232103]. [PubMed Central: PMC4128379].

Jundishapur J Chronic Dis Care. 2020; 9(3):e103036.

- Rhee H, McQuillan BE, Belyea MJ. Evaluation of a peer-led asthma self-management program and benefits of the program for adolescent peer leaders. *Respir Care*. 2012;57(12):2082–9. doi: 10.4187/respcare.01488. [PubMed: 22710616]. [PubMed Central: PMC3521097].
- Baker E, Fatoye F. Patient perceived impact of nurse-led selfmanagement interventions for COPD: A systematic review of qualitative research. *Int J Nurs Stud.* 2019;91:22–34. doi: 10.1016/j.ijnurstu.2018.12.004. [PubMed: 30669076].
- Shen Q, Zhu H, Jiang G, Liu X. Nurse-Led Self-Management Educational Intervention Improves Symptoms of Patients With Functional Constipation. West J Nurs Res. 2018;40(6):874–88. doi: 10.1177/0193945917701128. [PubMed: 28395593].
- Burschka JM, Keune PM, Oy UH, Oschmann P, Kuhn P. Mindfulnessbased interventions in multiple sclerosis: beneficial effects of Tai Chi on balance, coordination, fatigue and depression. *BMC Neurol.* 2014;14:165. doi: 10.1186/s12883-014-0165-4. [PubMed: 25145392]. [PubMed Central: PMC4236646].
- Halper J, Holland NJ. Comprehensive nursing care in multiple sclerosis. 3th ed. NY: Springer Publishing Company; 2010.
- 19. Schapiro RT. Managing the Symptoms of Multiple Sclerosis. 6th ed. New York: Demos Medical Publishing; 2014.
- Smeltzer SC, Bare B, Hinkle J, Cheever K. Brunner & Suddarth's Textbook of Medical-Surgical Nursing. 13th ed. Philadelphia: Lippincott William & Wilkins; 2014.
- 21. *MS Trust information service*. Hertfordshire: Multiple sclerosis trust; 2020, [cited 2020 Jun 15]. Available from: https://www.mstrust.org. uk.
- 22. Kurtzke JF. Rating neurologic impairment in multiple sclerosis: an expanded disability status scale (EDSS). *Neurology*. 1983;**33**(11):1444–52. doi: 10.1212/wnl.33.11.1444. [PubMed: 6685237].
- Diener E, Emmons RA, Larsen RJ, Griffin S. The Satisfaction With Life Scale. J Pers Assess. 1985;49(1):71–5. doi: 10.1207/s15327752jpa4901_13. [PubMed: 16367493].
- Salimi A. Social-Emotional Loneliness and Life Satisfaction. Procedia Soc Behav Sci. 2011;29:292–5. doi: 10.1016/j.sbspro.2011.11.241.
- Rahm T, Heise E, Schuldt M. Measuring the frequency of emotions-validation of the Scale of Positive and Negative Experience (SPANE) in Germany. *PLoS One*. 2017;**12**(2). e0171288. doi: 10.1371/journal.pone.0171288. [PubMed: 28178328]. [PubMed Central:

PMC5298234].

- Sahar J, Riasmini NM, Kusumawati DN, Erawati E. Improved Health Status and Life Satisfaction among Older People following Self-Help Group Intervention in Jakarta. *Curr Gerontol Geriatr Res.* 2017;2017:3879067. doi: 10.1155/2017/3879067. [PubMed: 29225621]. [PubMed Central: PMC5684573].
- Abbasi A, Ghezeljeh TN, Farahani MA. Effect of the self-management education program on the quality of life in people with chronic heart failure: a randomized controlled trial. *Electron Physician*. 2018;10(7):7028-37. doi: 10.19082/7028. [PubMed: 30128093]. [PubMed Central: PMC6092147].
- Hagger MS, Koch S, Chatzisarantis NLD, Orbell S. The common sense model of self-regulation: Meta-analysis and test of a process model. *Psychol Bull.* 2017;143(11):1117–54. doi: 10.1037/bul0000118. [PubMed: 28805401].
- Leventhal H, Phillips LA, Burns E. The Common-Sense Model of Self-Regulation (CSM): a dynamic framework for understanding illness self-management. *J Behav Med*. 2016;**39**(6):935–46. doi: 10.1007/s10865-016-9782-2. [PubMed: 27515801].
- Seymour JE, Almack K, Kennedy S, Froggatt K. Peer education for advance care planning: volunteers' perspectives on training and community engagement activities. *Health Expect.* 2013;16(1):43-55. doi: 10.1111/j.1369-7625.2011.00688.x. [PubMed: 21615641]. [PubMed Central: PMC3607415].
- Huber HB, Carter EW, Lopano SE, Stankiewicz KC. Using Structural Analysis to Inform Peer Support Arrangements for High School Students With Severe Disabilities. *Am J Intellect Dev Disabil.* 2018;**123**(2):119–39. doi: 10.1352/1944-7558-123.2.119. [PubMed: 29480778].
- Benzo RP, Abascal-Bolado B, Dulohery MM. Self-management and quality of life in chronic obstructive pulmonary disease (COPD): The mediating effects of positive affect. *Patient Educ Couns*. 2016;**99**(4):617– 23. doi: 10.1016/j.pec.2015.10.031. [PubMed: 26632024]. [PubMed Central: PMC4808334].
- Sawicki GS, Sellers DE, Robinson WM. Associations between illness perceptions and health-related quality of life in adults with cystic fibrosis. J Psychosom Res. 2011;70(2):161–7. doi: 10.1016/j.jpsychores.2010.06.005. [PubMed: 21262419]. [PubMed Central: PMC3052880].