The Effect of Education on Hemodialysis Patients' Fatigue and Self-Care

Eğitimin Hemodiyaliz Hastalarında Yorgunluk ve Öz Bakım Üzerine Etkisi

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Abstract

Aim: The study was done to evaluate the effect of education on the fatigue and self-care of patients receiving hemodialysis treatment.

Method: The study sample consisted of 70 patients, randomly selected (35 controls and 35 intervention groups). The study datas were obtained by using the Patient Identification Form, Piper Fatigue Scale and Self-Care Agency Scale. Individual training on the content supporting coping with fatigue and self-care power was given to patients in the intervention group. Piper Fatigue Scale and Self-Care Agency Scale were reapplied to all patients at the 3th and 6th months following the completion of the training sessions.

Results: In the pre-training evaluation, %88.6 of the intervention group and 91.4% of the control group experienced fatigue. The mean self-care score of the the intervention group was 86.20 and the control group was 79.11. In addition, there was no significant difference between the intervention and control groups in terms of their fatigue and self-care scores. In the evaluations at the third and sixth months, the control group's total fatigue and self-care scores had not changed (high fatigue scores and low self-care scores). Post intervention, the fatigue scores of the intervention group decreased while self-care scores increased, with a moderately significant negative correlation found between the fatigue and self-care score.

Conclusion: Results supported that individual training given to hemodialysis patients decreased fatigue scores and increased self-care scores.

Key Words: Hemodialysis; Fatigue; Self-Care; Education; Nursing.

Özet

Amaç: Bu araştırma, eğitimin hemodiyaliz tedavisi alan hastaların yorgunluk ve öz bakımlarına etkisini değerlendirmek amacıyla yapılmıştır.

Yöntem: Araştırma örneklemi rastgele seçilen (35 kontrol ve 35 müdahale grubu) 70 hastadan oluşmaktadır. Araştırma verileri Hasta Tanıtım Formu, Piper Yorgunluk Ölçeği, Öz-Bakım Gücü Ölçeği uygulanarak elde edildi. Müdahale grubundaki hastalara, yorgunlukla başetme ve öz bakım gücünü destekleyen içerikte bireysel eğitim verildi. Eğitimlerin tamamlanmasının ardından 3. ve 6. aylarda tüm hastalara Piper Yorgunluk Ölçeği ve Öz Bakım Gücü Ölçeği tekrar uygulandı.

Bulgular: Eğitim öncesi değerlendirmede müdahale grubundaki hastaların %88,6'sının, kontrol grubundaki hastaların yaşadığı, grubundaki %91.4'ünün vorgunluk müdahale hastaların öz-bakım puanı ortalamalarının 86,20, kontrol grubundaki hastaların öz bakım puanı ortalamalarının 79,11 olduğu, yorgunluk yaşama ve öz-bakım puanları açısından gruplar arasında farklılık olmadığı belirlendi. Üçüncü ve altıncı aylarda yapılan değerlendirmelerde kontrol grubunun toplam yorgunluk ve öz bakım puanları değişmedi (yüksek yorgunluk puanları ve düşük öz bakım puanları). Müdahale sonrası, müdahale grubunun yorgunluk puanları azalırken, öz bakım puanları artarken, yorgunluk ve öz bakım puanı arasında orta derecede anlamlı bir negatif korelasyon bulundu.

Sonuç: Çalışmanın sonuçları, hemodiyaliz hastalarına verilen bireysel eğitimin yorgunluk puanlarını azalttığını ve öz bakım puanlarını artırdığını desteklemektedir.

Anahtar Kelimeler: Hemodiyaliz; Yorgunluk; Öz Bakım; Eğitim; Hemşirelik.

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INTRODUCTION

Chronic kidney failure (CKD) is a chronic and progressive disease in which uremia result from a decrease in the patient's glomerular filtration rate (GFR). CKD in the world varies 14,9 % (1). In Turkey, the prevalence of CKD was 15.7 % (2).

Renal replacement therapies are needed when the glomerular filtration rate in GFR decreases below 15 ml/min./1.73m² (3,4). The most widely used (70.7%) renal replacement treatment in the world is hemodialysis (HD) (1). In Turkey, HD treatment is used with 76.93 % of patients (5).

Although HD is a life-saving treatment option, it requires many important changes in patients' life processes. HD treatment can cause patients to experience many physical, mental and social problems (2,6) with fatigue being one of the most common problems.

Fatigue is a subjective finding that includes emotional, cognitive and behavioral components, ranging from mild burnout affecting the whole body to unbearable exhaustion, preventing the person from performing his/her functions and using his normal capacity (7-10). Rates of fatigue, in patients with HD, can vary between 12 % and 97 % (7-9). Patients receiving HD treatments may experience fatigue associated with anemia (11), nutrition (8), sleep disorders (12), physical inactivity (13) and psycho-social factors (14). Fatigue can also be a factor that negatively affects individual's diet, sexual activity, family and friendship relationships, work life and leisure activities (7). Not properly managed, fatigue also may negatively affects the self-care of individuals (9,15). Self-care behaviors of patients with HD include many areas such as following the regimen, regular medication treatment compliance with HD treatment, fluid restriction. adherence to diet, communication, information and life satisfaction (16-18).

Nurses fulfill important responsibilities in assisting patients to manage their chronic diseases and nurses'effectiveness in disease management is increasing (19). By developing educational strategies, nurses can improve HD patients' adaptation to lifestyle changes, their coping with important complaints and their self-care skills (18). There are studies examining the level of fatigue in HD patients, and the relationships between fatigue (8) and exercise (9), sleep (12), anemia (20), psychosocial problems (12) and self-care and the factors affecting it (15).

However, there are no studies examining the effect of patient education on their fatigue and self-care. It is proposed that, with an effective nurse-patient education, the level of patient fatigue can be reduced, symptom control can be achieved and self-care adequacy can be increased. In this study, the effects of individual nurse-patient education, with patients who have HD, on fatigue and self-care of patients was investigated.

METHODS

Using a randomized assignment of participants to control and intervention groups, this study evaluated the effect of patient education on the fatigue and self-care of participants with. Seventy participants, aged 18 and over, were selected. Inclusion criteria included: receipt of HD treatment for at least six months, HD treatment three times a week, no diagnosed psychiatric disease, hemoglobin value greater than 10mg/dl, conscious, and willing to participate in the study. Patients meeting the sample selection criteria were then randomized to the intervention and control groups.

Initially, the Patient Identification Form, Piper Fatigue Scale, and Self-Care Agency Scale were given to the participants in both groups. The intervention group was then provided patient individual education in the form of three separate 30-minute sessions. The patient education materials developed by the researchers and included visual education materials, narration, question and answer, and discussion. In the first educational session, the following content was covered: the concept of fatigue, causes of fatigue in patients with HD,

dialysis treatment, diet and anemia. Content covered in the second education session included problems related to fatigue (such as sleep problems, stress, physical activity problems). The final education session addressed approaches to improve the participant's coping with their fatigue and approaches to further suppport self-care agency. The participants, completing all educational sessions were given a written educational booklet.

Data collection instruments were then administered to both groups three and six months after the individual educational sessions were completed. The following instruments were used to gather data as part of this research: 1) Patient Information form, 2) Piper Fatigue Scale (Piper et al, 1998), and 3) the Self-Care Agency Scale (Kearney & Fleischer, 1979).

- 1) **Patient Identification Form:** This instrument, based on related literature, consisted of questions about HD treatment and participants' sociodemographic information such as age, gender, education level, and comorbid diseases, self-care activities and medications.
- 2) Piper Fatigue Scale: The scale developed by Piper and colleagues (1998) consists of a total of 22 items. Responses for each item was scored between 0-10.The patient's perceptions about fatigue in the scale were evaluated in four sub-dimensions (behavior/ violence sub-dimension, sensory sub-dimension and cognitive/mental sub-dimension). dimension scores were obtained by dividing the number of items by the total sumof all items in that sub-dimension. The total fatigue score was derived by summing the scores of the 22 items and dividing them by the number of items. High scores obtained from the scale indicated that perceived fatigue level was high (21). The validity and reliability study of the scale for the Turkish society was conducted (22) and the Cronbach's alpha was 0.94. In the current study, the Cronbach's alpha was 0.98.
- 3) **Self-Care Agency Scale:** This insturment was developed Kearney and Fleischer (1979) to

determine the ability and power of people to take care of themselves (23). The scale is based on four characteristics, namely active or passive response to situations, motivation, knowledge of health practices and self- esteem, and includes a total of 35 statements.

The Likert scale has 5 scores: "it never defines me at all" (0 points), "it does not define me very much" (1 point), "I have no idea" (2 points), "defines me a bit" (3 points), or "defines me very much" (4 points). Nine items were scored negatively. Evaluation was based on a total of 136 points. A total score below 82 was "low", 82-120 points was "medium" and above 120 points was "high" self-care power. A high total score indicated that the individual was independent and able to perform self-care (23). In Turkey, thevalidity and reliability of the self-care agency scale (24) identified a Cronbach's alpha of 0.90.

DATA ANALYSIS

The datas analysis used SPSS 22. Percentages, t tests and ANOVAs were used with the level of error set at 0.05.

ETHICAL CONSIDERATIONS

The study was submitted to the University Ethics Committee who determined the research plan was ethically appropriate. Written permission was also obtained from the institution where the research would be conducted. All participants were informed about the study and their verbal and written consents were obtained. After the data collection process was completed, the patient education, provided to the intervention group, was also provided to the control group to assure the implementation of the equality principle.

RESULTS

As detailed in Table 1, the patients in the intervention and control groups had similar characteristics in terms of basic socio-demographic and disease characteristics (p> 0.05).

Table 1. Socio-Demographic and Disease Characteristics of Intervention and Control Group Patients

Spesifications	Intervention Group (n=35)		Control G	Control Group (n=35)		p
	Number	%	Number	%		
Candan	•	•	•		•	
Gender	10	71.4	10	540	<u> </u>	
Female	18	51.4	19	54.2	0.57	0.811
Male	17	48.6	16	45.8		
Age		_	_	ı		
31-40	7	20.0	3	8.6	3.51	
41-50	3	8.6	3	8.6		0.319
51-60	12	30.0	9	25.7	3.51	
61-70	13	37.1	20	57.1		
Education						
Literate	5	14.3	12	34.3		
Primary education	20	57.1	20	57.1	7.49	0.050
Highschool	7	20.0	3	8.6	7.48	0.058
University	3	8.6	0	0.0		
Marital Status	•	•	•			
Married	29	82.9	22	62.9	2 17	0.05
Single	6	17.1	13	37.1	3.45	0.06
Working Status						
Working	6	17.1	4	11.4		0.498
Not working	29	82.9	31	88.6	0.46	
Occupational Status	2,	02.9	31	00.0	_	
Housewife	16	45.7	19	54.3		0.657
Officer	4	11.4	1	2.9	+	
Retired	12	34.3	13	37.1	2.43	
Self-employment	2	5.7	13	2.9	2.43	
Worker	1	2.9	1	2.9	\dashv	
	1	2.9	1	2.9	_	
Income Rate	7	20.0		5.7		
Good	7	20.0	2	5.7	2.40	0.174
Mid	15	42.9	20	57.1	3.49	
Bad	13	37.1	13	37.1		
Family type				T .		
Extended family	29	82.9	23	65.7	2.69	0.101
Seed	6	17.1	12	34.3		
Disease Education	1			1		
Yes	20	57.1	19	54.3	0.58	0.810
No	15	42.9	16	47.7	3.50	3.010
Diet compliance						
Yes	20	57.1	16	45.7	0.91	0.339
No	15	42.9	19	54.3	0.91	0.339
Using drug						
User	34	97.1	32	91.4	1.00	0.303
Non-user	1	2.9	3	8.6	1.06	
Using erythropoietin		•				
User	33	94.3	32	91.4		0.643
Non-user	2	5.7	3	8.6	0.21	
Comorbidity	_					
Yes	24	68.6	26	74.3		0.597
No	11	31.4	9	25.7	0.28	
110	11	31.4)	43.1		

^{*}p<0.05

As detailed in Table 2, there was no significant difference between the patients in the intervention and control groups in terms of their descriptive characteristics such as fatigue, duration of fatigue, state of fatigue affecting their daily lives and coping behaviors (p> 0.05).

As detailed in Table 3, no statistical difference was identified between the pre-education fatigue scores of the patients in either the intervention or the control groups (p>0.05). However, the posteducation fatigue scores of the intervention group

were significantly lower than those of the control group at both three and six months. There was also a statistically significant difference between the preand post-education total fatigue score averages of the intervention group at the third and sixth months with the scores decreasing significantly after the education. In the control group, there was no statistically significant difference between the preand post-education total fatigue score averages with the fatigue scores continuing to be high.

Table 2. Fatigue Characteristics of Patients Before Education

Characteristics		Intervention Group (n=35)		Control Group (n=35)		p
		%		%		
Fatigue						
Available	31	88.6	32	91.4	0.159	0.690
Absent	4	11.4	3	8.6	0.139	
The state of fatigue affecting daily life						
Affecting	27	87.0	29	90.6	1.30	0.105
Non-affecting	4	13.0	3	9.4	1.50	
Taking support to cope fatigue						
Taking	1	3.2	2	6.3	2.25	0.08
Non-taking	30	96.7	30	93.7	2.23	
Time to experience fatigue						
For weeks	9	31.4	6	18.7	1.29	0.204
For months	22	68.6	26	81.3		
Cause of fatigue						
Hemodialysis	26	80.0	18	56.2	5.85	0.022*
Chronic renal failure	5	20.0	14	43.8	3.63	
Initiatives to reduce fatigue						
Sleeping	14	45.1	13	40.6	0.17	0.700
Resting	10	32.2	13	40.6		
Do nothing	7	25.7	6	22.8		
Complaints other than fatigue						
Hypotension	13	42.9	13	40.6	1.13	0.200
Muscle cramps	11	37.1	15	46.8		
Not	7	20.0	4	12.5		

^{*}p<0.05

Table 3. Fatigue Scores of Individuals in Intervention and Control Groups in Pre-and Post-Training Follow-up

Total fatigue scores	Before training	Third month after training	Sixth month after training	F	p
	X±SD	X±SD	X±SD		
Intervention group	7.24±1.62	4.84±1.04	4.67±0.92	266.40	0.001*
Control group	7.71±1.35	7.80±1.55	7.75±1.42	0.40	0.669
t	1.30	9.35	10.73		
p	0.198	0.001*	0.001*		_

^{*}p<0.05

As detailed in Table 4, There was no statistical difference between the pre-education self-care scores of the patients in the intervention and control groups (p>0.05).

There was no statistically significant difference between the pre- and post-education control group self-care score averages and the self-care, with posteducation self-care scores remaining low.

Self-care scores of the intervention group were significantly higher than those of the control group at post-education measurements. Furthermore, there was a statistically significant difference between the pre- and posteducation self-care score averages of the patients in the intervention group with the self-care scores of the patients increasing significantly.

An adddition finding was that of a significant, negative and medium-level relationship between the fatigue and self-care scores of the patients in both the intervention and control groups. It was also found that self-care decreased with increasing fatigue level in both groups.

DISCUSSION

Chronic diseases negatively affect the self-care activities of individuals. It is important for patients, experiencing HD, to develop self-care strategies related to fatigue (9,25-28).

Akın et al. (2013) found that only 9.5% of patients with HD were independent in terms of their self-care (29). In contrast, some studies found self-cares to be moderate in patients with HD (24,30,31). Research findings have supported that patient education

results in an increase in the self-care score of patients with HD (16,32,33).

In this study, self-care scores of the intervention and control groups were similar to the results in the literature. While self-care scores continued at a low level in the control group after education, a significant increase was observed in the scores of the intervention group. This result supports that individual patient education can increase patient self-care scores.

Fatigue is a common problem in patients experiencing HD. In the pre-training stage, of the research, it was determined that the rate of fatigue was very high in both the control and intervention groups. In previous studies, patients with HD had a very high rate of fatigue ranging from 65.4 % (25) to 92.9 % (25-28) and approximately one third of patients had high levels of fatigue (25). Some studies have shown that, as patients' with HD general fatigue level increases, daily activities are also increasingly effected by fatigue (9,34,35). In This study identified high rates of patients indicating that fatigue affected their daily lives (intervention 77.1%; control 88.6%). This is an important finding as supports that learning coping strategies with fatigue is important for patients.

Although fatigue is an important problem in patients with HD, the literature states that patients do not generally apply effective approaches to cope with fatigue other than resting (36,37). In this study, most of the patients stated that they preferred to sleep and rest in order to cope with fatigue.

Table 4. Pre-training and After Self-Care Scores of Patients in Intervention and Control Groups

	Self-Care A			
	Intervention group	Control group		
	X±SD	X±SD	t	p
Pre-training	86.20±16.00	79.11±14.65	1.93	0.058
3rd months after training	115.40±11.13	78.68±14.47	11.89	0.001*
6th months after training	118.97±11.18	76.91±15.14	13.21	0.001*
F	133.92	1.40		
p	0.001*	0.252		

^{*}p<0.05

Both Mohamed (2014) and Patterson et al. (2013) reported that patient education reduced fatigue (37, 38). Mohamed's study (2014) identified that patient fatigue score averages in both the intervention and control groups were similar before education related to coping with fatigue, with the intervntion group's average fatigue score decreasing after the education, while the control group's average fatigue score In this study the fatigue score remained high. averages of both intervention and control groups patients before the education were found to be high. While the total fatigue scores of the intervention group decreased three and six months after the education, the fatigue scores of the control group did not change. This result supports that education is effective in decreasing fatigue levels in the patients experiencing HD.

Evaluation of fatigue and self-care in patients with HD is important to develop approaches that will contribute to reduce fatigue in these patients (9). According to Levey et al. (2007), psychosocial problems and fatigue lead to changes in individuals' daily life activities and decreased self-care (15). Akin et al. (2013) found a negative correlation between patient fatigue and self-care (29). Slesnick et al. (2015) found that self-care education reduced fatigue (39). In the current study, there was also a

negative and significant correlation between fatigue and self-care score in the pre-education evaluation. Following the patient education, a negatively insignificant correlation between these variables assists in understanding the relationship between patient education and fatigue and self-care scores.

CONCLUSION AND SUGGESTIONS

In this study, results identified that patients with HD had high levels of fatigue and low self-care scores. Withthe control group, fatigue scores were high and self-care scores were low at the first evaluation, and these results did not change at the third and sixth month evaluations. Following patient education, the patients in the intervention group had decreased fatigue scores and increased self-care scores.

These results support that patient education is important to assist patients with HD to better cope with fatigue and increase their self-care. As a result, patient eductions programs for patients with HD should be designed to assist these patients to cope with fatigue and support their self-care. It is also recommended that the organized education programming be developed by hemodialysis and/or nephrology nurses in order that self-care and fatigue symptoms, specific to patients with HD, are evaluated and addressed.

ETHICS COMMITTEE APPROVAL

Ethics committee approval: Sivas Cumhuriyet University Non-Invasive Clinical Research Ethics Committee (Date and no: 17.04.2015/ 2015-04-08)

INFORMED CONSENT

From the study participants informed consent was obtained.

CONFLICT OF INTEREST

Any financial or other interest in the study there is no conflict.

FINANCIAL SUPPORT

Any institution/organization related to the study has no financial support.

PEER REVIEW

External independent, double blind.

AUTHOR CONTRIBUTIONS

Idea and design: HTA, GD

Data collection: GD

Data analysis and interpretation: GD Preparation of the article: HTA, GD

Critical review: HTA, GD

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