Aerobic Training Improves Symptoms of Neuropathy and Quality of Life in Patients with Diabetic Peripheral Neuropathy

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Abstract

Background: Painful symptoms of diabetic peripheral neuropathy (DPN) is common in patients with type 2 diabetes mellitus (T2DM), which can reduce the quality of life. The effect of aerobic training on neuropathic symptoms and quality of life in people with type 2 diabetes with peripheral neuropathy was investigated in the present study.

Methods: Twenty-four patients with DPN participated in this randomized controlled study. Participants were randomly assigned to the experimental or control groups. The experimental group underwent exercise training consisting of 20-45 minutes aerobic training on the treadmill at 50%-70% of heart rate reserve over three months. Before and after the experimental period, blood samples were taken in a fasting state, and Michigan Diabetic Neuropathy Score (MDNS), monofilament test and quality of life (QOL), and sleep quality (SQ) questionnaires were completed. Data analysis was performed using repeated-measures ANOVA.

Results: In the experimental group MDNS score significantly improved compared with the control condition (P<0.05). There was a significant difference between groups regarding 10 g monofilament test score (P<0.05). Moreover, QOL and SQ significantly improved in the experimental group with a significant inter-group difference (P<0.05). Hemoglobin A1c (HbA1c) level decreased by 7.4% in the experimental group, which was statistically different from the control group (P<0.05). Pearson’s correlation indicated a significant correlation between HbA1c level and monofilament score.

Conclusion: Regular aerobic training over a short period potentially improves symptoms of DPN and QOL. Lowered levels of HbA1c were correlated with improved foot sensation.

Keywords: Type 2 diabetes mellitus, Diabetic neuropathy, Exercise training, Pain, Life quality

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nerve fiber branching and reduced pain in DPN (13). In addition to these physiological effects of exercise training, it also increases the QOL in people with T2DM (14). Patients with T2DM with DPN experience more severe symptoms than those without DPN (15). Thus, the QOL consequences of exercise training in these people may be different with patients with T2DM without DPN and depends on its impact on neuropathic symptoms. Any intervention to reduce pain may have a direct impact on aspects of life QOL in this population.

**Objectives**

Therefore, we aimed to investigate the effect of regular aerobic training on symptoms of neuropathy, QOL, and sleep in patients with DPN and its relation with HbA1c.

**Methods**

**Participants and Design**

This was a randomized-controlled parallel-design study. Thirty-one patients volunteered to participate in the present study based on the inclusion criteria. Inclusion criteria were; having T2DM for more than five years, HbA1c > 6.6%, DPN diagnosis, and an inactive lifestyle. The presence of DPN was confirmed by a neurologist based on initial assessments using the Michigan Diabetic Neuropathy Instrument (scores > 3) and abnormal electrophysiological measures at lower limbs. Patients with foot ulcers and orthopedic issues were not included in this study. The participants were not engaged in any exercise program and had no experience of regular exercise participation. A complete description of the aims and procedure of the research was given to the subjects, and then they signed the written consent form.

Participants were randomly allocated to the experimental and control groups, and 12 individuals completed the trial in each group. The experimental group was familiarized with the exercise protocol over a week prior to the main experimental period. During this period, the participants performed 15-20 m walking on the treadmill with the least range of intensity to be familiarized with the exercise procedure. Then, they performed the main aerobic exercise program over 3 months. Exercise training consisted of walking or running at 50%-70% of heart rate reserve lasting 20-45 min each session with a frequency of three sessions/week. Karvonen equation was used to assess target heart rate, and heart rate was monitored using a polar heart rate monitor (PE3000, Polar Electro, Kempele, Finland). Each session included a 10-15 min warm-up at the beginning and 5-10 minutes cool down at the end. The control group was required to keep habitual physical activity and inform the researchers in case of any alteration.

**Medication and Diet**

The participants were on an oral hypoglycemic regimen which did not change during the study period. The participants were asked to keep their usual diet. They were asked to record food intake the day preceding the measurements and replicate it during the day before the final measurements. They were also asked to record a 3-day (2 weekdays and 1 weekend) dietary recall a week prior to the study period and during the last week. The calorie and macronutrients consumption were assessed to be considered as a covariate in the analysis if there was any significant difference between groups and from pre-to post-test.

**Measurements**

Before and 48 h following the experimental period, body composition was analyzed with the participants wearing minimal clothing using the bioelectrical impedance analyzer (Inbody230. Portable, Frequency: 20 kHz, 100 kHz – Display: 240*320 Color LCD). Blood samples were taken in a fasting state to assess Hb1Ac levels at baseline and after the intervention. HbA1c level was assessed by liquid chromatography (Pishtaz, Tehran, Iran). Neuropathic symptoms were evaluated by Michigan Diabetic Neuropathic Score (MDNS), which includes 46 points clinical score and is used for DPN staging (16). QOL state was assessed by QOL questionnaire that assesses subjective feelings of patients. Quality of sleep was assessed by Petersburg Questionnaire. To assess foot sensation, 10 gr filament was applied to the dorsum of the great toe, and the patients were asked to respond YES when they felt the filament (16).

**Statistical Analysis**

Data analysis was performed using SPSS software, version 25. Normal distribution of the data was confirmed using Shapiro–Wilk test. Repeated-measures ANOVA was used to determine the intergroup differences. Pre-test and post-test values were also analyzed by paired t test, and independent t test was used to compare the groups at pre-test. Pearson's correlation coefficient was also used to assess the relationship between variables. P < 0.05 was considered statistically significant.

**Results**

Descriptive data are shown in Table 1. Data analysis indicated no significant difference between the groups at baseline (P > 0.05). Participants had mild to moderate DPN.

Repeated measures ANOVA indicated that HbA1c significantly decreased in the experimental group compared with the control group (P = 0.01). MDNS score also improved in the experimental group throughout the intervention period, which was statistically significant with the control group (P = 0.001). MDNS improvement was accompanied by foot sensation improvement, and the monofilament test score increased significantly.

**Table 1**

| Variable | Pre-Test | Post-Test | P-Value |
|----------|----------|-----------|---------|
| HbA1c    | 6.8      | 6.2       | <0.05   |
| MDNS     | 3.5      | 2.7       | <0.01   |

Descriptive data are shown in Table 1. Data analysis indicated no significant difference between the groups at baseline (P > 0.05). Participants had mild to moderate DPN.
Pearson's correlation coefficient also indicated a significant correlation between monofilament score and HbA1c (r = -0.53 and P = 0.021) and between MDNS and QOL scores (r = 0.73 and P = 0.01). Data analysis revealed that QOL and sleep significantly improved following three months of exercise compared with the control group (P = 0.03 and P = 0.01, respectively) (Table 2).

Discussion
We found that after three months of aerobic training, HbA1c levels improved, which was coupled with improvement in foot sensation, neuropathic symptoms, and QOL and sleep.

The results regarding reduced HbA1c levels following exercise training are consistent with most studies in this area (9, 17, 18). Metabolic effects of aerobic training have been previously established, especially in patients with metabolic disorders (19). Several mechanisms have been proposed; for instance, glucose uptake in muscles increases during and following exercise because of the increased muscle permeability as a result of contraction (20). In addition, GLUT4 transcription and translocation increase by exercise, which has a direct impact on insulin sensitivity and glucose uptake (20). Although very few studies have recently investigated the effectiveness of exercise on complications of diabetes, particularly DPN, this metabolic effect of exercise training could be promising in this population as hyperglycemia plays a crucial role in the pathogenesis and progression of DPN.

Another finding of the present study was the improvement in MDNS following exercise training. Despite the lack of research on patients with DPN, this finding is consistent with some studies (13, 21). Dixit and colleagues reported that 8-week exercise training improved neuropathic symptoms in people with DPN (21). Kluding et al also found that 10-week combined endurance-resistance training reduced neuropathic pain in patients with diabetes with peripheral neuropathy (13). We also observed that foot sensation assessed by the monofilament test improved by exercise training. Possible causes of improvement in neuropathic symptoms can be attributed to the improvement of underlying factors, including blood glucose and blood circulation to the peripheral nerves. We found that improvement in foot sensation was coupled with HbA1c reduction. Although we did not explore the vascular changes following exercise training in our study, it has been previously reported that patients with neuropathy can benefit from vascular effects of exercise training (22). Thus, these changes brought about by exercise training may explain improvements in neuropathic symptoms. Even though the results were consistent to some extent, this study could be comparable with the aforementioned research in some aspects. For instance, Kluding and colleagues' study was not controlled and randomized, and MNSI was applied by this research group to evaluate the state of neuropathy (13).

Another finding of the study was the improvement in QOL and sleep quality (SQ) in this population following the intervention. This is in line with the results of another study showing that regular exercise over three months improves the QOL in elderly individuals with mild disabilities (23). Several studies have illustrated that moderate exercise training can improve sleep quantity and quality. However, little is known about its effect in

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**Table 1. Mean ± SD of Body Composition and Calorie Intake**

| Variables         | Groups                        | P Value |
|-------------------|-------------------------------|---------|
|                   | Experimental (n = 12) | Control (n = 12) |
|                   | Pre       | Post     | Pre       | Post     |
| Age (year)        | 43 ± 4    | -        | 44 ± 6    | -        | 0.49 |
| Weight (kg)       | 86 ± 15   | 85 ± 15  | 89 ± 11   | 89 ± 11  | 0.09 |
| BMI (kg/m²)       | 29 ± 5    | 28 ± 5   | 29 ± 3    | 28 ± 2   | 0.20 |
| Calorie intake (kcal) | 2352 ± 224 | 2392 ± 221 | 2437 ± 211 | 2413 ± 168 | 0.61 |

P values presented for Time * Group interactions. P is presented by baseline comparison for age.

**Table 2. Mean ± SD of Variables Throughout the Study**

| Variables   | Groups                        | P Value |
|-------------|-------------------------------|---------|
|             | Experimental (n = 12) | Control (n = 12) |
|             | Pre       | Post     | Pre       | Post     |
| MDNS        | 15.2 ± 2.3  | 12.1 ± 2.1 | 15.4 ± 2.5 | 15.3 ± 2.8 | 0.001 |
| QOL         | 54.3 ± 7.6  | 66.4 ± 8.4 | 51.8 ± 6.4 | 54.1 ± 8.2 | 0.03 |
| SQ          | 8.4 ± 2.0   | 6.5 ± 1.6 | 8.9 ± 3.2 | 8.7 ± 2.7 | 0.01 |
| Monofilament| 4.9 ± 1.5   | 5.8 ± 1.4 | 5.1 ± 1.7 | 5.0 ± 1.6 | 0.01 |
| HbA1c (%)   | 8.32 ± 1.41 | 7.74 ± 1.50 | 8.61 ± 1.43 | 8.54 ± 1.83 | 0.01 |

Abbreviations: MDNS, Michigan Diabetic Neuropathy Score; QOL, quality of life; SQ, sleep quality. P values presented for Time * Group interactions.
patients with DPN. Dixit and co-workers reported that 8 weeks of regular exercise improved the QOL in this group of patients (21). However, they did not assess whether this improved QOL was correlated with neuropathic symptoms. We observed a relationship between the changes in MDNS score and QOL in patients with DPN. We assume that one of the most important possible reasons for improved QOL and sleep with exercise training is the improvement of neuropathic symptoms. Neuropathic pain is an important factor in reducing the QOL in these patients. We also found that following three months of exercise training, the quality of sleep remarkably improved. Neuropathic symptoms in this population are aggravated at night, which could be a cause of sleep disorders and poor SQ (24). Moreover, evidence shows that exercise training improves the quality of sleep through elevating endorphin secretion and energy consumption which facilitates sleep and recovery of the body (25, 26). Thus, we speculate that exercise training can potentially bring about metabolic effects and cause physiological changes that may improve neuropathic symptoms in these patients, which may eventually result in a better QOL and sleep.

**Conclusion**

Overall, the results of the present study showed that a three-month aerobic exercise training program decreased HbA1c level and improved neuropathic symptoms, QOL and sleep in patients with DPN. There was a correlation between HbA1c and monofilament test score and also between neuropathic symptoms and QOL scores following the intervention. Moderate aerobic training may be recommended as a feasible strategy along with other therapeutic interventions for patients with diabetes and DPN.

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**Conflict of Interests**

The author declares no conflict of interest.

**Ethical Approval**

The study was approved by the ethics committee in human research (Code: BZMED.REC.1394.153). All the participants were informed about the study procedure, and written consent was obtained from each participant. The study protocol complies with the ethical guidelines of the 1975 Declaration of Helsinki. Besides, the study was registered by the Iranian Registry of Clinical Trials (identifier: IRTC2017012032066N1).

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