The effect of an interval training course on nitric oxide levels, insulin resistance, and some blood lipid factors in type 2 diabetic male rats

Running title: The effect of an interval training course on nitric oxide

Roohollah Mohammadi Mirzaei

Faculty of Sports Sciences, Kharazmi University, Tehran, Iran. ORCID: 0000-0001-5109-8018, Dr.Mohamadi@cfu.ac.ir

Hamid Malekshahi

Department of Sports Sciences, Farhangian University, Shahid Chamran Campus, Tehran, Iran.

ORCID: 009-0008-8822-8478, h.malekshahi2017@gmail.com

Halimeh Vahdatpoor

Department of Physiology, Faculty of Sports Sciences, Hakim Sabzevari University, Sabzevar,

Iran. ORCID: 0000-0001-6364-1751, Vahdatpoor.114@gmail.com

Corresponding Author: Roohollah Mohammadi Mirzaei

Address: Department of Sports Sciences, Farhangian University, Shahid Chamran Campus,

Tehran, Iran

Email: Dr.Mohamadi@cfu.ac.ir

Tel: 09352152011-+982188085725

Abstract

Background: Diabetes is a metabolic and vascular disorder characterized by endothelial dysfunction. Physical activity, particularly intermittent exercise, may offer therapeutic benefits. This study examined the effects of six weeks of intermittent exercise on serum nitric oxide (NO), insulin resistance, and lipid profiles in male diabetic rats.

Methods: Sixty Wistar rats (8 weeks old, 200 ± 20 g) were divided into six groups (n=10): healthy control, sham, interval training, interval training + saline, diabetic control, and diabetic + interval training. Diabetes was induced via nicotinamide-streptozotocin injection. The exercise groups underwent treadmill training (5 sessions/week for 6 weeks). Post-intervention, glucose, insulin, NO, LDL, HDL, triglycerides, and cholesterol were measured.

Results: Diabetes induction significantly increased glucose and insulin resistance while reducing insulin and NO levels compared to controls (P value < 0.001). After six weeks, the diabetic + exercise group showed significant reductions in glucose and insulin resistance (P value < 0.001) and increased NO levels (P value < 0.001) versus the diabetic group. Insulin levels did not differ significantly among groups (P value = 0.11). Lipid profiles (LDL, HDL, triglycerides, cholesterol) remained unchanged (P value > 0.05).

Conclusion: Six weeks of intermittent exercise improved glucose metabolism and vascular function in diabetic rats by reducing insulin resistance and increasing nitric oxide levels, suggesting its potential as a non-pharmacological therapy for diabetes-related endothelial dysfunction.

Keywords: Interval training, Insulin resistance, Nitric oxide, Fat factors, Diabetes.

Introduction

As one of the major health problems, diabetes has affected more than 400 million people worldwide, and most of them are type 2 diabetics (1). Various studies have shown that diabetes is not only a metabolic disease but also a vascular disease due to its effects on small and large vascular beds, and the relationship between diabetes and increased prevalence of cardiovascular disease has been well proven (2). High blood pressure and type 2 diabetes seem to be the main factors that increase the risk of death from cardiovascular diseases (3). Diabetic patients are faced with endothelial functional disorders and damage. These disorders include functional changes in the endothelium (impaired vasoconstriction) and increased inflammatory activity that is related to cardiovascular diseases (4). The most active vasodilator produced by endothelial cells is nitric oxide (5). Nitric oxide is the cause of vascular dilation, it is a regulatory molecule with extensive metabolic, vascular, and cellular effects, which originates from the oxidation of gonidine group, L-arginine. It is found in almost all tissues. It is also secreted in response to stimuli such as lack of oxygen in the tissue and shear stress of endothelial cells (6). Reducing the amount of nitric oxide inhibits vascular dilation, and its amount in the blood of diabetic patients is lower than that of healthy people (7). Another characteristic of type 2 diabetes is insulin resistance, which seems to cause vascular endothelial dysfunction through increased fat, insulin, and oxidative stress. Also, increased fatty acids stimulate endothelial apoptosis. As a result, it increases vascular oxidative stress. It also reduces the amount of NO (8). Therefore, increasing NO by reducing vascular tone, reducing blood pressure, and improving insulin sensitivity has been recognized as a suitable treatment for patients with type 2 diabetes (9). Sports activity through successive changes in shear stress and its increase leads to an increase in the biological activity of NO and improvement of vascular endothelial function (10). In a study, Farzanegi showed that 30 minutes of swimming training had significantly reduced blood glucose, insulin, and NO in diabetic rats (11).

Many other risk factors increase cardiovascular complications in these patients. The most common of these factors are increased levels of TG, decreased levels of HDL, increased levels of LDL, obesity, weight gain, and high blood pressure, which lead to atherosclerosis (12).

HIIT (High-intensity interval training) is an effective training method in cardiovascular rehabilitation, which has been confirmed to have positive effects on health and disease. In this training model (13) and compared to continuous training, HIIT has a greater effect on improving parameters related to cardiovascular health (14). Despite the cardiovascular protective effects of HIIT and its role in the management of diabetes, information about its cellular mechanisms is incomplete and sometimes contradictory, and more studies are needed on how HIIT affects, and in this regard, in this study, the effect of 6 weeks of HIIT on some Factors related to cardiovascular diseases including hyperglycemia, insulin resistance, nitric oxide and some lipid factors were investigated in diabetic rats.

Methods

Analysis method

The present experimental study used 60 male Wistar rats (8 weeks old, 200 ± 20 g), obtained from the Ahvaz Physiology Research Center. The tested animals were kept in groups of five in polycarbonate racks and had free access to standard food and water. The ambient temperature was 22 ± 3 centigrade, the light-dark cycle was 12:12 hours, and the humidity was $55.6\pm4\%$. After one week familiarization with the environment, all animals were introduced to running on the treadmill for one week (10 minutes at a speed of 10 meters per minute and five days a week). Then the rats were randomly divided into six healthy and diabetic groups. 1) Healthy groups: These groups included 40 rats, which were randomly divided into four groups: control, intermittent exercise, sham, and intermittent exercise + saline. 2) Diabetic groups: This group included 20 rats. They were randomly divided into two groups: diabetes and diabetes + interval training.

To induce type 2 diabetes, first, nicotinamide (120 mg/kg) was injected intraperitoneally into the rats, and 15 minutes later, a single dose of streptozotocin (STZ) (60 mg/kg) dissolved in normal citrate buffer 1/ 0 M was injected intraperitoneally (IP) into the animal. Then, to ensure that the animal became diabetic, the level of blood sugar increase was evaluated 72 hours after STZ injection using a glucometer. The rats whose fasting glucose was more than 250 mg/dl were considered diabetic. The reason for the risk of hypoglycemia is caused by STZ. Rats received a 10% glucose solution after 6 hours of STZ administration until 24 hours later.

Exercise protocol

One week after the induction of diabetes, the rats in the exercise intervention group performed intermittent exercise on the treadmill for 6 weeks and 5 days a week. Before starting the main exercises and for familiarization, the rats started running on the treadmill for 10-15 minutes at a speed of 5-7 meters per minute with a zero-degree slope for two consecutive days. 2 days after the familiarization exercises, the main intermittent exercises began and the rats' performed activities on a special treadmill for 6 weeks. The interval training program was such that the speed of the training program in the first week started from 10 meters per minute. From the second to the sixth week, the weekly training speed increased by 2 meters per minute, so that it reached 20 meters per minute in the last week. The duration of training in the periodic group increased daily from the first to the sixth week, so that the duration of the activity increased from 15 minutes on the first day to 40 minutes on the sixth week. The interval training group performed the training for the specified duration in two sessions in the first week, four sessions in the second to fourth weeks, and six sessions in the fifth to sixth weeks. 4 minutes of exercise and 1 minute of active rest) was considered active, running at a speed of 3 meters per minute (Table 1). At the beginning of each training session, each training group ran for 3 minutes at a speed of 1 m/min to warm up, and then to reach the desired speed, 2 m/min was added to the speed of the treadmill, and in At the end of the training, they ran for 3 minutes at a speed of 1 meter per minute to cool down. Rats in the intermittent training group were encouraged to continue running in all training sessions using a weak electric shock (intensity 0.5 mA) that did not cause much stress in the animal (15).

Week	Exercise duration (minutes)	Training speed (meters per minute)	Type of rest
First	10	2×7.7	Active
Second	12	4×5	Active
Third	14	4×6.25	Active
Fourth	16	4×7.5	Active
Fifth	18	6×5.83	Active
Sixth	20	6×6.66	Active

 Table 1. Intermittent exercise implementation protocol

Blood sampling and laboratory analysis

Twenty-four hours after the final training session, rats from both the healthy and diabetic groups were anesthetized via intraperitoneal injection of ketamine-xylazine. Blood samples were then collected via cardiac puncture and transferred into Falcon tubes. After coagulation, the blood

samples were centrifuged at 4000 rpm for 15 minutes to separate the serum. The serum was then aliquoted and stored at -70° C until further analysis for the target variables.

To check the NO index, the Promega (Promega Corp. USA) kit was used. To measure the amount of insulin from the ELISA kit (Diacolon, France), the amount of glucose was calculated using the photometric method (Pars Azmoun, Iran), and the insulin resistance was calculated using the homeostasis model evaluation method according to the following formula (11):

HOMA-IR = [Glucose] \times [Insulin] / 22.5 \times 18

The measurement of lipid profile (high-density lipoprotein, triglyceride, and total cholesterol) was done by the enzymatic method and using commercial biochemistry kits prepared by Pars Azmoun - Iran, and low-density lipoprotein was calculated using the equation of Friedwald et al.

Statistical method

In the descriptive statistics section, the dispersion indices of mean and standard deviation were used. In the inferential statistics section, the Shapiro-Wilk test was used to determine the normality of the data distribution, and the homogeneity of the variances was measured with the Lunn test. To determine the significance of the difference between the variables and their interaction, one-way analysis of variance and Tukey's post hoc test were used. The findings were analyzed at the 95% confidence interval, and statistical analysis of the data was done using SPSS software.

Results

The average serum levels of glucose, insulin, LDL, HDL, TG, and cholesterol of the subjects are presented in Table 2. ANOVA test results showed that after six weeks, a significant difference was observed in the blood glucose level of the rats. In order to investigate the difference between the groups, the follow-up test also showed that the blood glucose of the diabetes + interval training group had decreased significantly compared to the diabetes group after six weeks (P value=0.001). Also, the amount of blood glucose in the interval training group was significantly reduced compared to the control group (P value=0.001). However, no significant difference was observed between the two intermittent training groups and the intermittent training + saline group (P value>0.55). Regarding lipid profile indicators, ANOVA test results showed that after six weeks, there was no significant difference between the levels of LDL, HDL, TG and cholesterol of the groups (respectively: (P value>0.12, P value>0.059, P value>0 /059 , P value>0.84, (Table 2)). The results of the ANOVA test regarding the insulin index also showed that after six weeks, no significant difference was observed between the groups (P value=011). A significant difference was observed between the groups (P value=0.001) (Table 2).

The ANOVA test results showed that after six weeks, there was a significant difference between the groups regarding the insulin resistance index (P value=0.001). According to the follow-up test, the insulin resistance of the diabetes + exercise group was significantly reduced compared to the diabetes group (P value=0.001). Also, a significant difference was observed between the insulin resistance of the diabetes group and the control, sham, intermittent exercise + saline, and intermittent exercise groups (P value=0.001). However, there was no significant difference between the insulin resistance of other groups (P value=0.001). (Figure 1).

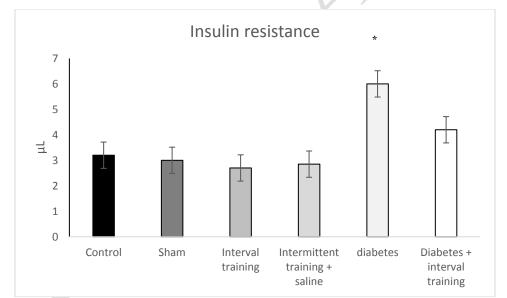
Also, the results of the ANOVA test regarding the NO index showed that after six weeks, there was a significant difference between the groups (P value=0.001). According to the follow-up test, the NO index in the diabetes + exercise group had increased significantly compared to the diabetes group (P value=0.003). Also, a significant difference was observed between the amount of NO in the diabetes group, the interval training + saline group, and the interval training group (P

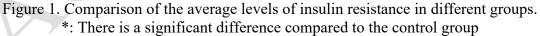
value=0.001). A significant difference was also observed between the amount of NO in the control group and the diabetes, intermittent exercise, and diabetes + intermittent exercise groups (P value=0.001). However, there was no significant difference between NO in other groups (P value>0.05) (Figure 2).

 Table 2. Mean and standard deviation Results of glucose, insulin, LDL, HDL, TG and Cholesterol groups

Variable (Unit)	Control	Sham	Interval training	Intermittent training + saline	Diabetes	Diabetes + interval training	P value
Glucose (mg/dL)	110.11±4.64	105.74 ± 4.58	91.41±4.08	94.4±6.35	271.81±23.52	182.57 ± 12.82	0.001
Insulin (m/ml)	11.54±2.02	11.54±1.97	10.58 ± 1.81	10.77 ± 2.06	9/13±1/30	8.21±1.06	0.11
LDL (mg/dl)	37.13±3.32	36.89±5.14	34.50±4.36	34.76±4.55	39.11±2.68	36.67±3.27	0.126
HDL (mg/dl)	32.82±4.60	33.73±4.60	35.92±2.41	35.36±2.29	31.24±5.19	36.05±4.13	0.059
TG (mg/dl)	83.01±7.82	83.63±7.70	76.26±8.17	76.93±7.69	86.44±11.27	81.13±8.80	0.059
Cholesterol (mg/dl)	86.59±7.30	87.34±8.74	84.09±6.08	85.48±6.44	87.37±4.24	86.42±6.73	0.845

Repeating the letter symbol indicates a lack of significance between the two groups. Using the letter alone indicates a significance between the two groups.





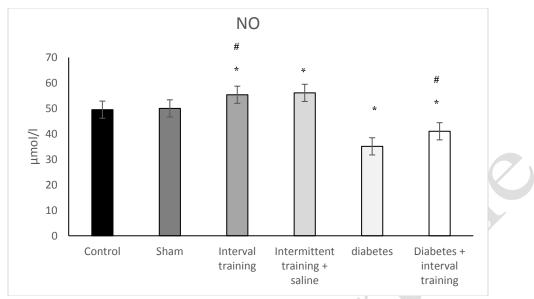


Figure 2. Comparison of mean NO serum levels in different groups. *: There is a significant difference compared to the control group #: There is a significant difference compared to the diabetes group

Discussion

The present study was conducted to investigate the effect of 6 weeks of intermittent training on nitric oxide levels, insulin resistance, and some lipid factors in type 2 diabetic male rats. The results of this study showed that the induction of diabetes caused a significant decrease in the serum levels of NO and a significant increase in the blood glucose level and insulin resistance in the diabetic group. Thus, 6 weeks of intermittent training in the diabetes group + intermittent training caused a significant increase in the serum NO level compared to the diabetes group. Also, in the present study, the blood glucose level and insulin resistance after 6 weeks of intermittent exercise in the diabetes group + intermittent exercise decreased significantly (P<008), Which is similar to the studies of Farzangi (11), Asarzadeh et al. 16) and Mohebi et al. (17) and it is consistent with the studies of Wang et al. (18) and Shad et al. (19). Probably, these different results are due to differences in the type, duration, intensity of training and different samples. Various studies have shown that exercise exercises through 1) insulin receptors, 2) protein and mRNA of glucose transporter (GLUT4), 3) increasing glycogen synthetase, protein kinase-B, hexokinase, and 4) improving internal messengers. Insulin cells and the effect on intermediate molecules in the insulin signal, such as increasing the expression of ERK2, increasing the activity of PI3K or AKt/PKB and improving the AMPK signal (20), changes in muscle composition (increasing capillary density in muscle fibers and converting muscle fibers into fast-twitch fibers) oxidation), 6) increasing the delivery of glucose to the muscle, 7) reducing the accumulation of triglycerides in the muscle cell and 8) reducing the release of fatty acids and increasing their oxidation and clearing (21), modulating insulin resistance. In the present study, the level of NO decreased significantly after induction of diabetes, but after 6 weeks of intermittent exercise, the level of NO increased significantly (P<05). Which is consistent with the studies of Mitranen et al. (22), Qardashi et al. (23), Grijava et al. (24) and Zheng et al. (25). In another study, Zhang et al. reported an increase in serum NO levels following 10 weeks of exercise. Research has shown that exercise increases myocardial NOx production, eNOS protein levels, and increases sensitivity to eNOS phosphorylation caused by insulin stimulation (26). As shown in this study, the induction of diabetes causes hyperglycemia, increase of LDL levels (in a non-significant way) and insulin resistance, each of these factors in a way causes disruption in the endothelial function of the vessels of diabetic people (27).

Several other studies have also shown that shear stress leads to the stimulation of increased expression of eNOS (28). In support of this hypothesis, it has been stated that HIIT is more effective in improving vascular function compared to moderate intensity continuous exercise training (MICT), Which is probably due to the ability of HIIT to stimulate more blood flow through the vessels supplying oxygen to the working muscles. Which increases the bioavailability of NO caused by shear stress (29). Thijssen et al. (30) confirmed this hypothesis and stated that with the increase in the intensity of sports activity, blood flow and shear stress also increase. Although the full effects of HIIT on endothelial function and NO release are not fully understood, it has been shown that HIIT can reduce the levels of Catecholamines and the density of alpha-adrenergic receptors (31). Reducing the activity of sympathetic tone and increasing the activity of parasympathetic tone has a great effect on blood pressure regulation. As previously mentioned, the increase in triglyceride (TG), decrease in HDL, and increase in LDL and cholesterol can lead to atherosclerosis (12). Diabetes, especially type 2 diabetes, is often associated with lipid metabolism disorders, and increased serum levels of fatty acids play a major role in insulin resistance (32). In this study, it was shown that the levels of LDL, TG and cholesterol increased insignificantly after the induction of diabetes and the level of HDL decreased insignificantly, but after 6 weeks of intermittent exercise in the diabetes group, the level of HDL increased insignificantly and the level of LDL, TG And cholesterol also decreased insignificantly (P<05). Which is inconsistent with the study of Niyazi et al. (33) and Ghasem Nia et al. (34) but consistent with the research of Gordon et al. (35) It can be said that the difference between the results of our research and some other researches can be attributed to the difference between the intensity, duration, training environment, type of samples, age and gender of the research samples. In general, the accumulation of excess fat from two separate main pathways causes insulin resistance, which includes the transformation of insulin signaling with cytokines secreted from adipose tissue and the damage or death of pancreatic beta cells due to the accumulation of free fatty acids. However, long-term exercise training, by reducing fat accumulation, probably improves insulin sensitivity and prevents insulin resistance (36, 37).

Conclusion

According to the results of the present study, the time and intensity of sports activity according to the type of disease and the age of the subjects can reduce blood glucose levels and factors related to cardiovascular disease (lipid profile) as well as increase NO, improve endothelial function and consequently Decrease in the occurrence of cardiovascular diseases in diabetes conditions. As a result, sports activity with appropriate intensity can be recommended as a non-pharmacological method for diabetic patients.

Acknowledgments

This study was registered in the Iranian Registry of Clinical Trials (code: IRCT2015100423002N2). The authors gratefully acknowledge all participants for their valuable contributions to this research.

Author contributions

Roohollah Mohammadi Mirzaei: Original draft, Methodology. Hamid Malekshahi: Writing, Review and Editing, Project Management. Halimeh Vahdatpoor: Methodology, Analysis.

Funding

The authors received no financial support for the research, authorship, and/or publication of this article.

Ethics approvals

Ethical approval code: (IR.HSU.REC.1400.008).

Conflict of interest

The authors declare that there is no conflict of interest regarding publication of this article

Data availability statement

The data that support the findings of this study are available from the corresponding author upon reasonable request.

References

1. Ramírez-Alarcón, K., Labraña, A. M., Victoriano, M., Meléndez-Illanes, L., & Martorell, M. Diabesity: Obesity And Type II Diabetes As A Real Health Problem In Developed Countries. Diabesity: A Multidisciplinary Approach, 2022; 1.

2. Arrieta, F., Pedro-Botet, J., Iglesias, P., Obaya, J. C., Montanez, L., Maldonado, G. F., ... & Aguilar, M. Diabetes mellitus and cardiovascular risk: an update of the recommendations of the Diabetes and Cardiovascular Disease Working Group of the Spanish Society of Diabetes. Clínica e Investigación en Arteriosclerosis (English Edition), 2022; 34(1), 36-55.

3. Nazarzadeh, M., Bidel, Z., Canoy, D., Copland, E., Bennett, D. A., Dehghan, A., Wang, J. Blood pressure-lowering treatment for prevention of major cardiovascular diseases in people with and without type 2 diabetes: an individual participant-level data meta-analysis. The Lancet Diabetes & Endocrinology, 2022; 10(9), 645-654.

4. Yudkin JS, Eringa E, Stehouwer CDA. "Vasocrine" signalling from perivascular fat: a mechanism linking insulin resistance to vascular disease. The Lancet. 2005;365(9473):1817-20.

5. Man, A. W., Zhou, Y., Xia, N., & Li, H. Endothelial Nitric Oxide Synthase in the Perivascular Adipose Tissue. Biomedicines, 2022; 10(7), 1754.

6. Giacco F, Brownlee M. Oxidative stress and diabetic complications. Circ Res. 2010;107(9):1058-70.

7. Mehri Alvar Y, Sayeevand Z, Erfani Adab F, Heidari Moghadam R, Samavat Sharif M, Karami S. The effects of five weeks resistance training on some vascular growth factors in sedentary men. Sport Physiology. 2016;8(29):15-30.

8. Andreadi, A., Bellia, A., Di Daniele, N., Meloni, M., Lauro, R., Della-Morte, D., & Lauro, D. The molecular link between oxidative stress, insulin resistance, and type 2 diabetes: A target for new therapies against cardiovascular diseases. Current Opinion in Pharmacology, 2022; 62, 85-96.

9. Henry RM, Ferreira I, Kostense PJ, Dekker JM, Nijpels G, Heine RJ, et al. Type 2 diabetes is associated with impaired endothelium-dependent, flow-mediated dilation, but impaired glucose metabolism is not; The Hoorn Study. Atherosclerosis. 2004;174(1):49-56.

10. G Zhang, X., & Gao, F. Exercise improves vascular health: Role of mitochondria. *Free Radical Biology and Medicine*,2021; *177*, 347-359.

11. Farzanegi P. The effect of regular swim training with two different time periods on serum levels of NO, VEGF, and TGF- β 1 in diabetic male rats. mdrsjrns. 2017;20(2):37-48.

12. Stirban AO, Tschoepe D. Cardiovascular complications in diabetes: targets and interventions. Diabetes Care. 2008;31 Suppl 2:S215-21.

13. Kemi OJ, Wisloff U. High-intensity aerobic exercise training improves the heart in health and disease. J Cardiopulm Rehabil Prev. 2010;30(1):2-11.

14. Freyssin C, Verkindt C, Prieur F, Benaich P, Maunier S, Blanc P. Cardiac rehabilitation in chronic heart failure: effect of an 8-week, high-intensity interval training versus continuous training. Arch Phys Med Rehabil. 2012;93(8):1359-64.

15. Brooks GA, White TP. Determination of metabolic and heart rate responses of rats to treadmill exercise. J Appl Physiol Respir Environ Exerc Physiol. 1978;45(6):1009-15.

16. Assarzade Noushabadi M, Abedi B. Effects of combination training on insulin resistance index and some inflammatory markers in inactive men. QHMS. 2012;18(3):95-105.

17. Mohebi H, Talebi GE, Rahbarizadeh F. Effect of intensity training on plasma adiponectin concentration in male rat. 2009.

18. Wang, W., Huang, M., & Wang, J. (2021). The effect of physical exercise on blood sugar control in diabetic patients. *Revista Brasileira de Medicina do Esporte*, *27*, 311-314.

19. Shad R, Bijeh N, Fathi M. The effects of eight weeks of aerobic exercise on insulin resistance, body composition and aerobic power of overweight female students. Journal of Knowledge & Health. 2017;12(2):17-24.

20. Fízel'ová M, Cederberg H, Stančáková A, Jauhiainen R, Vangipurapu J, Kuusisto J, et al. Markers of tissue-specific insulin resistance predict the worsening of hyperglycemia, incident type 2 diabetes and cardiovascular disease. PLoS One. 2014;9(10):e109772.

21. Enomoto T, Ohashi K, Shibata R, Kambara T, Uemura Y, Yuasa D, et al. Transcriptional regulation of an insulin-sensitizing adipokine adipolin/CTRP12 in adipocytes by Krüppel-like factor 15. PLoS One. 2013;8(12):e83183.

22. Tanahashi K, Akazawa N, Miyaki A, Choi Y, Ra SG, Matsubara T, et al. Aerobic exercise training decreases plasma asymmetric dimethylarginine concentrations with increase in arterial compliance in postmenopausal women. Am J Hypertens. 2014;27(3):415-21.

23. Ghardashi Afousi A, Izadi MR, Rakhshan K, Mafi F, Biglari S, Gandomkar Bagheri H. Improved brachial artery shear patterns and increased flow-mediated dilatation after low-volume high-intensity interval training in type 2 diabetes. Exp Physiol. 2018;103(9):1264-76.

24. Grijalva J, Hicks S, Zhao X, Medikayala S, Kaminski PM, Wolin MS, et al. Exercise training enhanced myocardial endothelial nitric oxide synthase (eNOS) function in diabetic Goto-Kakizaki (GK) rats. Cardiovasc Diabetol. 2008;7:34.

25. Zhang QJ, Li QX, Zhang HF, Zhang KR, Guo WY, Wang HC, et al. Swim training sensitizes myocardial response to insulin: role of Akt-dependent eNOS activation. Cardiovasc Res. 2007;75(2):369-80.

26. Adams V, Linke A, Kränkel N, Erbs S, Gielen S, Möbius-Winkler S, et al. Impact of regular physical activity on the NAD(P)H oxidase and angiotensin receptor system in patients with coronary artery disease. Circulation. 2005;111(5):555-62.

27. Yudkin JS, Eringa E, Stehouwer CD. "Vasocrine" signalling from perivascular fat: a mechanism linking insulin resistance to vascular disease. Lancet. 2005;365(9473):1817-20.

28. Kolluru GK, Sinha S, Majumder S, Muley A, Siamwala JH, Gupta R, et al. Shear stress promotes nitric oxide production in endothelial cells by sub-cellular delocalization of eNOS: A basis for shear stress mediated angiogenesis. Nitric Oxide. 2010;22(4):304-15.

29. Ramos JS, Dalleck LC, Tjonna AE, Beetham KS, Coombes JS. The impact of high-intensity interval training versus moderate-intensity continuous training on vascular function: a systematic review and meta-analysis. Sports Med. 2015;45(5):679-92.

30. Thijssen DH, Dawson EA, Black MA, Hopman MT, Cable NT, Green DJ. Brachial artery blood flow responses to different modalities of lower limb exercise. Med Sci Sports Exerc. 2009;41(5):1072-9.

31. Jung CH, Lee M, Kang YM, Lee Y, Yoon H, Kang S-W, et al. Vaspin inhibits cytokineinduced nuclear factor-kappa B activation and adhesion molecule expression via AMP-activated protein kinase activation in vascular endothelial cells. Cardiovascular diabetology. 2014;13:41.

32. Prenner SB, Chirinos JA. Arterial stiffness in diabetes mellitus. Atherosclerosis. 2015;238(2):370-9.

33. Niyazi, Arghavan, et al. High-intensity interval versus moderate-intensity continuous exercise training on glycemic control, beta cell function, and aerobic fitness in women with type 2 diabetes. Biological Research For Nursing, 2024, 26.3: 449-459.

34. Ghasemnian A, Ghaeini A, Kordi M, Hedayati M, Rami M, Ghorbanian B. EFFECT OF INTERVAL ENDURANCE TRAINING PROGRAM ON PLASMA EOTAXIN, ADIPONECTIN LEVELS, INSULIN RESISTANCE, SERUM LIPID PROFILE AND BODY COMPOSITION IN OVERWEIGHT AND OBESE ADOLESCENTS. URMIAMJ. 2013;24(6):430-40.

35. Gordon LA, Morrison EY, McGrowder DA, Young R, Fraser YT, Zamora EM, et al. Effect of exercise therapy on lipid profile and oxidative stress indicators in patients with type 2 diabetes. BMC Complement Altern Med. 2008;8:21.

36. Khademosharie M, Amiri Parsa T, Hamedinia MR, Azarnive Ma, Hosseini-Kakhk SAR. Effects of two aerobic training protocols on Vaspin, Chemerin and lipid profile in women with type 2 diabetes. BPUMS. 2014;17(4):571-81.

37. Kim ES, Im JA, Kim KC, Park JH, Suh SH, Kang ES, et al. Improved insulin sensitivity and adiponectin level after exercise training in obese Korean youth. Obesity (Silver Spring). 2007;15(12):3023-30.