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The Impact of Varying Circuit Resistance Training Intensity on Apolipoproteins in Men with Obesity

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ABSTRACT

Introduction: This study aimed to investigate the impact of 12 weeks of circuit resistance training at varying intensities on Apolipoproteins A (Apo A), B (Apo B), and J (Apo J) in men with obesity.

Materials & Methods: A total of 44 young men with obesity, aged 23-32 years, were divided into four groups: control (n=11), low-intensity (n=11), moderate-intensity (n=11), and high-intensity circuit resistance training (n=11) groups. The training program spanned 12 weeks and was conducted with three sessions per week. Blood samples were collected 72 hours before and 72 hours after the 12-week training programs. These blood samples were transferred to specialized plasma test tubes containing EDTA and were then centrifuged at 10 rpm for 10 minutes.

Findings: A significant decrease in Apo B levels (P < 0.0001) and an increase in Apo A levels (P < 0.05) were observed in the low-intensity training group compared to the control group. Additionally, a significant decrease in Apo J levels was observed in all exercise training groups compared to the control group (P < 0.05).

Discussion & Conclusion: The findings of this study suggest that circuit resistance training, particularly at lower intensities, may lead to a reduction in the risk of cardiovascular diseases in obese individuals through the modification of Apo A, Apo B, and Apo J levels.

Keywords: Circuit Resistance Training, Obese Men, Apolipoprotein, Training Intensity

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Introduction

Obesity has profound implications for human health, and it is linked to a range of diseases, including type 2 diabetes, lipid disorders, cardiovascular ailments, and various types of cancer. Ultimately, obesity is associated with reduced life expectancy and premature mortality, imposing a substantial burden on public health (1).

Insulin resistance stands as significant risk factor for type 2 diabetes, with notable abnormalities in fat and plasma lipoproteins, including reduced HDL cholesterol and elevated triglyceride levels. These changes are linked to increased hepatic triglyceride secretion. Apolipoproteins, a group of proteins located at the outermost layer of lipoproteins, play a pivotal role in regulating lipid transport and lipoprotein metabolism (2).

Emerging research suggests that two key apolipoproteins, apolipoprotein A (Apo A) and apolipoprotein B (Apo B), which are major components of HDL-C and LDL-C, respectively, may offer a more accurate assessment of coronary heart disease risk than conventional lipid markers (3). Multiple studies have demonstrated Apo B's pivotal role in the structure of LDL-C and VLDL-C, with its interaction with LDL-C receptors playing a crucial role in their uptake by peripheral cells and liver (4).Plasma concentration is influenced by Apo B's hepatic secretion and catabolism and is a determining factor in LDL-C levels plasma density (5). Human studies have consistently revealed a significant inverse relationship between HDL-C levels, particularly Apo A, and the risk of atherosclerosis (6). The Apo B/Apo A-1 ratio, when compared to LDL-C, provides a more reliable indicator of coronary artery disease risk (7). LDL-C alone is insufficient indicator as an predicting cardiovascular risk (8). In essence, Apo A, the primary protein associated with HDL-C, possesses wellknown antioxidant properties (9, 10). Together with Apo A, HDL-C plays a critical role in the reverse transfer of cholesterol from lipid-rich macrophages to the liver, preventing the accumulation of cholesterol esters in macrophages and the formation of foam cells (9,10).

Apolipoprotein J (Apo J), also known as clusterin, represents a disulfidelinked heterodimer protein expressed in various tissues and bodily fluids (11).Apo I is a fundamental component of high-density cholesterol and low-density cholesterol, and its function activates enzymes involved in lipoprotein metabolism (2). Apo J is found in a subset of dense HDL particles that include Apo A I and Paraoxonase (PON). It is also present in various physiological fluids, such as human plasma, urine, breast milk, semen, and cerebrospinal fluid (11). Increased levels of circulating Apo I associated with various pathological conditions, including obesity, diabetes, Alzheimer's disease, and cardiovascular disorders (12). Recent research has highlighted the close relationship between Apo I circulation and markers of cardiovascular metabolism. with documented links between Apo J and insulin resistance in humans (13). Changes in serum Apo I levels are correlated inversely improvements in insulin sensitivity in with patients tvpe 2 diabetes. suggesting that Apo I turnover reflects insulin resistance (14). Interestingly, Apo J in HDL has been linked to insulin sensitivity, while Apo J in LDL/VLDL has been associated with insulin resistance, suggesting a potential

connection between Apo J and insulin function (14). Consequently, some studies propose that Apo I may have a protective role in individuals with insulin resistance, and elevated serum Apo I levels could serve as a marker for assessing insulin resistance (15). Research has shown that oral administration of Apo I reduces atherosclerosis in Apo E-null mice and anti-inflammatory improves the properties of HDL in monkeys (16). Apo J is identified as a multifunctional protein and may serve as an effective and sensitive predictor of atherosclerotic lesions and cardiovascular disease (17).

Exercise is widely acknowledged as a key factor in preventing and treating obesity and its related ailments, significantly enhancing insulin sensitivity and mitigating associated complications (18). Aerobic exercise enhances insulin sensitivity, while resistance training increases glucose uptake in muscle tissue by augmenting muscle mass and Glut4 expression. The combined effects of these two exercise types are instrumental in both preventing and treating obesityrelated diseases. Circuit resistance training, characterized by performing multiple exercises in quick succession, offers an effective way to lose weight and increase muscle strength (19,20).

Navab et al. demonstrated that 12 weeks of combined resistance-aerobic exercise training in women with type 2 diabetes resulted in significant reductions in insulin resistance and Apo I levels (16). Given the scarcity of studies on the effects of exercise training, especially circuit resistance training at different intensities, on changes in Apo J and its precise role in obesity, this study aimed to investigate the effects of 12 weeks of circuit training resistance with varving intensities on Apo J, Apo A, and Apo B levels in obese individuals.

Materials and Methods

Given that the study focused on obese male subjects and extended over a 12-week research period, it can be categorized as both practical and quasi-experimental.

Participant Selection

The study's participants were obese male volunteers who were recruited through outreach to public, academic, and administrative institutions. To be eligible for the study, participants had meet specific criteria, which included the absence of drug and alcohol addiction, a lack of regular exercise for at least six months before the study's commencement, no history of kidney, liver, cardiovascular disease, or diabetes, and meeting specific health conditions (BMI ≥ 30 and Waist-to-Height Ratio (WHtR) > 0.5). Furthermore. each participant underwent a comprehensive medical examination to confirm their physical fitness. Prior to their involvement in the research, all participants received a detailed explanation of the study's procedures and methods. After gaining full understanding and completing a questionnaire, medical written consent was obtained from each participant. The study selected 44 volunteers within an age range of 23-32 years. All research procedures were subjected to scrutiny and approval by the Research and Ethics Committee of Urmia University (Ethics code: IR-UU1400-16), and all activities adhered to the latest revision of the Declaration of Helsinki.

Initial Assessments

During the initial session, participants' height, weight, and dietary habits (recorded via a food frequency questionnaire) were documented.

Furthermore, all participants received training on the study's various stages, the correct execution of the exercise program, and safety precautions related to the exercises.

Second Session

In the second session, participants completed appetite and physical activity questionnaires. Blood samples were collected, and a one-repetition maximum test was conducted.

Group Allocation

Participants were evenly distributed into four groups based on their

maximum strength in the onerepetition maximum test:

Control group (n=11)

Low-intensity circuit resistance training group (n=11)

Medium-intensity circuit resistance training group (n=11)

High-intensity circuit resistance training group (n=11).

This rigorous selection and grouping process ensured the participants' homogeneity, facilitating an effective study.

Table 1. Characteristics of the subjects in each group

Group Variable	Control	LT	MT	НТ
Age (years)	25 <u>+</u> 40	26.00 <u>±</u> 50	28 <u>±</u> 30	27 <u>±</u> 60
Height (cm)	174.18 <u>+</u> 4.75	178.10 <u>+</u> 6.08	174.36 <u>+</u> 4.60	175.92 <u>+</u> 5.31
Weight (kg)	95.91 <u>+</u> 9.40	98.10 <u>±</u> 10.51	96.36 <u>+</u> 8.21	94.42 <u>+</u> 8.46
BMI (kg/m ²)	31.35±0.96	30.91±0.72	31.68±0.60	30.72±0.93

Data are shown as mean \pm SD. LT; low, MT; moderate and HT; severe intensity circuit resistance training groups

Determination of 1RM

To establish the one-repetition maximum (1RM), the Berziski method was utilized with subjects from the 1RM resistance training group. We selected weights for participants that allowed them to complete a maximum of 6-8 repetitions. Subsequently, the calculated weight, taking repetitions into account, was applied to the appropriate formula (22):

 $1RM = weight / (1.0278 - 0.0278 \times reps)$

Circuit Resistance Training Protocol

The circuit resistance training protocol consisted of eight exercises targeting the upper and lower torso, which included squat, forearm exercises, chest press, knee opening, flexion, barbell head lifts, leg presses, and armpit wire exercises from These behind. exercises were conducted at varying intensities (23,24)

High-Intensity Circuit Exercise Group: 3 sets of 10 repetitions at 80% of 1RM.

Medium-Intensity Circuit Exercise Group: 3 sets of 13 repetitions at 60% of 1RM

Low-Intensity Circuit Exercise Group: 3 sets of 20 repetitions at 40% of 1RM.

Training Volume Calculation:

The training volume was calculated using the formula presented by Baechele et al (1994). RT volume = number of sets \times repetitions \times weight lifted (25).

Rest Periods

Rest intervals between exercise stations were minimized to the time required for participants to transition from one station to another (\leq 15 seconds). Rest intervals between sets were set at 2 minutes and were passive in nature (26).

Blood Sampling and Analysis

Fasting blood samples were collected 72 hours before and after a twelveweek training period from the right arm vein of the participants. The blood samples were collected in specialized plasma test tubes containing EDTA and then centrifuged at 10,000 rpm for 10 minutes. The resulting plasma was stored at -70°C. Plasma Apolipoprotein measured using was immunoturbidimetric method with a quantitative detection kit from Bionic Company, with a variation coefficient and sensitivity of 4.8% and 0.31 mg/100 ml, respectively. Apolipoprotein B was measured using the immunoturbidimetric method with quantitative detection kit from Bionik Company, featuring a variation coefficient and sensitivity of 1.2% and 20 mg/100 ml, respectively. Plasma Apol concentration was measured using an ELISA kit from Boster Biological Technology according to the manufacturer's protocols, with sensitivity < 20 of pg/mL. coefficients of variation within and between tests were 4.2% to 4.6% and 6.9% to 7.5%, respectively.

Statistical Analysis

Descriptive statistics were employed for classification and determination of dispersion indices. The Golmogorov-Smirnov test confirmed that the data followed a normal distribution. A paired t-test was used to compare pretest and post-test changes within each test group. For intergroup comparisons, a repeated two-way analysis of variance and Bonferroni post hoc test were applied. All data were expressed as mean ± standard deviation. Statistical analyses were carried out using SPSS version 22, with significance set at P < 0.05.

Results

Analysis of Apo A Values

The analysis of Apo A values in subjects before and after 12 weeks of circuit resistance training at different intensities, using a repeated measures analysis of variance test with the intergroup factor, revealed that both the time factor (P < 0.0001, F = 0.66) and the time-group interaction (P < 0.0001, F = 3.40) were statistically significant.

Bonferroni Test Results

The results of the Bonferroni test indicated that there was no statistically significant difference between the changes in Apo A values of subjects in the control group and the low-intensity training group (P =0.181). However, significant differences were observed when comparing the control group with the moderate training group (P < 0.0001) and the high-intensity training group 0.0001). Additionally. significant difference was found when comparing the low-intensity training group with the high-intensity training group (P < 0.0001) and the moderateintensity training group (P < 0.0001).

However, no statistically significant difference was detected between the moderate-intensity training group and the high-intensity group (P = 0.99).

Intra-Group Changes in Apo A Values

An examination of intra-group changes in Apo A values using a paired t-test revealed that the changes following 12 weeks of training were statistically significant in the control group (P = 0.008), the moderate intensity group (P = 0.001), the high-intensity group (P < 0.0001), and the low-intensity training group (P = 0.001). Please refer to Figure 1 for a visual representation of these changes.

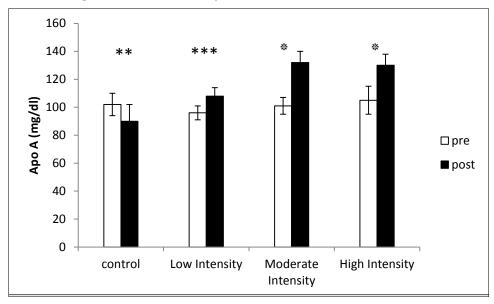


Figure 1. Apo A values of subjects before and after 12 weeks of circuit resistance training with different intensities. * sign indicates a significant difference within the group. ** sign indicates a significant difference between the control group and other high and medium intensity groups. *** sign shows a significant difference between low intensity groups and high and medium intensity groups.

Analysis of Apo B Values

An analysis of Apo B values in subjects before and after 12 weeks of circuit resistance training with different intensities, employing a repeated measures analysis of variance test with the intergroup factor, revealed that both the time factor (P < 0.0001, F1.40 = 30.9) and the time-group interaction (P < 0.0001, F3.40 = 8.8) were statistically significant.

Bonferroni Posthoc Test Results

The results of the Bonferroni posthoc test indicated that there was no statistically significant difference between the changes in Apo B values of subjects in the control group and the low-intensity exercise group (P =0.481). However. significant differences were observed when comparing the control group with the moderate exercise groups (P < 0.0001) and the high-intensity group (P <0.0001). Additionally, a significant difference was found when comparing the low-intensity training groups with the high-intensity training group (P = 0.044) and the moderate-intensity group (P = 0.022). However, no statistically significant difference was detected between the moderateintensity training group and the highintensity group (P = 0.99).

Intra-Group Changes in Apo B Values

An examination of intra-group changes in Apo B values using a paired t-test revealed that the changes after 12 weeks of training were not statistically significant in the control group (P = 0.363) and the low-intensity training

group (P = 0.076). However, these changes were statistically significant in the medium-intensity group (P = 0.006) and the high-intensity group (P <0.0001). Please refer to Figure 2 for a visual representation of these changes.

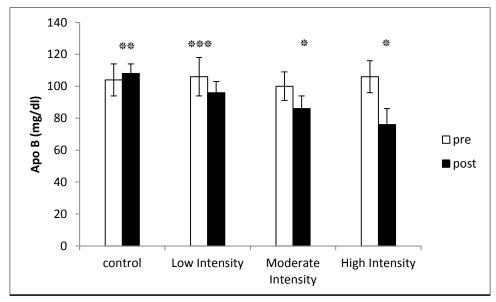


Figure 2. Apo B values of subjects before and after 12 weeks of circuit resistance training with different intensities .The * sign indicates a significant difference within the group .The ** sign indicates a significant difference between the control group and other high and medium intensity groups . the mark *** Shows a significant difference between low intensity groups and high and medium intensity groups.

Analysis of Apo J Values

The analysis of Apo I values in subjects before and after 12 weeks of circuit training different resistance at employing intensities. a repeated measures analysis of variance test with the intergroup factor, revealed that both the time factor (P < 0.0001, F1.40 = 118.4) and the time-group interaction (P < 0.0001, F3.40 = 11.4) were statistically significant.

Bonferroni Test Results

The results of the Bonferroni test indicated that there were significant differences between the changes in Apo J values of subjects in the control group and the low-intensity group (P < 0.0001), the moderate-intensity group (P < 0.0001), and the high-intensity group (P < 0.0001).

Intra-Group Changes in Apo J Values

An examination of intra-group changes in Apo J values using a paired t-test revealed that the changes after 12 weeks of training were not statistically significant in the control group (P =0.132). However, these changes were statistically significant in the lowintensity training group (P < 0.0001), the moderate-intensity group (P < 0.0001), and the high-intensity group 0.0001). (P < For a visual representation of these changes, please refer to **Figure**

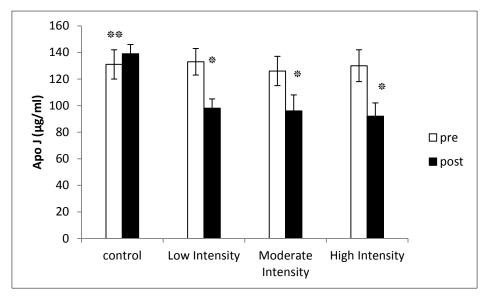


Figure 3. Apo J values of subjects before and after 12 weeks of circuit resistance training with different intensities .The * sign indicates a significant difference within the group .The ** sign indicates a significant difference between the control group and other groups.

Discussion

Insulin resistance is a significant risk factor for type 2 diabetes and is associated with disruptions in fat and plasma lipoproteins, leading to decreased HDL cholesterol, elevated triglyceride levels, and increased hepatic triglyceride secretion (27).

Our study revealed a noteworthy reduction in Apo J and Apo B levels following twelve weeks of low. medium, and high-intensity resistance training. Importantly, this decrease was more pronounced in the moderate and high-intensity training groups than in the low-intensity group, with no significant change observed in the control group. In contrast, Apo A levels displayed a significant increase following all three training intensities, with no significant change noted in the control group.

Our findings align with a prior study where women with type 2 diabetes underwent twelve weeks of combined aerobic resistance training. In that study, significant reductions in body weight, body fat percentage, and Apo J were observed post-training (15). Existing evidence suggests that Apo J levels are notably elevated individuals with conditions associated insulin resistance. such obesity, metabolic syndrome, and type 2 diabetes (14, 28). Another study post-intervention involving a combination of weight loss and thiazolidinedione treatment to enhance insulin sensitivity resulted in reduced circulating Apo J levels (14). Interestingly, Apo I levels appear to be unaffected by BMI or the degree of weight loss, but they decrease significantly in cases where patients experience weight reduction following calorie restriction (15, 23). Apo J shows no significant correlation with BMI, weight loss, leptin, lipoproteins, except for a modest association with plasma leptin (23).

Sevral studies have reported significant associations between circulating Apo J and total cholesterol and LDL cholesterol levels (21, 29). However, no definitive evidence links changes in Apo J levels with variations

in lipid profiles following exercise. Results concerning the effects of exercise on Apo B and Apo A are conflicting. For instance, one study on healthy adults found that Apo A and Apo B values remained relatively stable after six weeks of moderate and high-intensity resistance training (30). Additionally, Valente et al. (2011) reported that resistance training in conjunction with nutritional intervention led to a significant reduction in Apo B levels and a nonsignificant decrease in Apo A levels (31).

The impact of exercise intensity on apolipoproteins appears variable. The effect of different aerobic exercise intensities on changes in Apo A and has been reported Apo В statistically insignificant (32).Similarly, study anaerobic on training courses with varying showed no significant intensities changes in cardiovascular risk factors or Apo A and Apo B levels (33). However, moderate-intensity exercise was found to significantly affect the Apo B/Apo A ratio but not Apo B levels (33).

Conclusion

In summary, our results demonstrate that circuit resistance training over twelve weeks, at varying intensities (low, medium, and high), leads to a significant decrease in Apo I and Apo B levels, while significantly increasing Apo A levels in obese individuals. These improvements are particularly notable in the moderate and highintensity training groups. Consequently. regular circuit resistance training holds the potential to mitigate the risk of cardiovascular diseases in obese individuals enhancing inflammatory parameters. It serves as a non-pharmacological

treatment approach with considerable effectiveness in disease prevention.

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Authors' Contributions

M. R. and A. N. collaborated throughout all stages of manuscript development.

Conflict of Interest

The authors declare no conflicts of interest.

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References

- Marinou K, Tousoulis D, Antonopoulos AS, Stefanadi E, Stefanadis C. Obesity and cardiovascular disease: from pathophysiology to risk stratification. Int J cardiol. 2010; 138(1):3-8. doi: 10.1016/j.ijcard.2009.03.135.
- Bijeh N, Sarlak Z, Farahati S. Effects of eight weeks aerobic training on serum Apo A-1, B and lipid profile in overweight women. Sport Physiology. 2016; 7(28): 45-58.
- 3. Talebi-Garakani E. The Effect of Resistance Training Intensity on Serum ApoA-I Concentration in Streptozotocin-Induced Diabetic Rats. Iran J Endocrinol Metab. 2013;15(2):183-9.
- Faam B, Hedayati M, Azizi F. Presence of the X+ allele in apolipoprotein b gene increase the total cholesterol and apolipoprotein b concentration in tehranian people. Iran J Endocrinol Metab. 2011;12(6):588-93.
- 5. Ng TW, Watts GF, Farvid MS, Chan DC, Barrett PHR. Adipocytokines and VLDL metabolism: independent regulatory effects of adiponectin, insulin resistance, and fat compartments on VLDL apolipoprotein B-100 kinetics? Diabetes. 2005;54(3):795-802. doi: 10.2337/diabetes.54.3.795.
- Ghorbanian B, Ravassi A, Kordi MR, Hedayati M. The effects of rope training on lymphocyte ABCA1 expression, plasma

- ApoA-I and HDL-c in boy adolescents. Int J Endocrinol Metab. 2013;11(2):76. doi: 10.5812/ijem.8178.
- Mashaykhi NR, Sadrneya S, Chehrei A, Javaheri J, Ahmadlou M. The correlation between serum apo lipoprotein A1 and apo lipoprotein B with coronary artery disease and its severity. J Arak Uni Med Sci. 2013;16(6):82-9.
- Kadoglou NP, Fotiadis G, Athanasiadou Z, Vitta I, Lampropoulos S, Vrabas IS. The effects of resistance training on ApoB/ApoA-I ratio, Lp (a) and inflammatory markers in patients with type 2 diabetes. Endocrine. 2012;42(3):561-9. doi: 10.1007/s12020-012-9650-y.
- Rashidlamir A, Ghanbari-Niaki A, Saadatnia
 A. The Effect of eight weeks of wrestling
 and wrestling technique-based circuit
 training on lymphocyte ABCA1 gene
 expression and plasma apolipoprotein AI.
 World J Sport Sci. 2011;2(2):144-50.
- Malekpour-Dehkordi Z, Javadi E, Doosti M, Paknejad M, Nourbakhsh M, Yassa N, et al. The effect of Alcoholic garlic (Allium sativum) extract on ABCA1 expression in human THP-1 macrophages. Tehran Univ Med J. 2011;69(3):146-152
- 11. Trougakos IP, Gonos ES. Clusterin/apolipoprotein J in human aging and cancer. Int J Biochem Cell Biol. 2002;34(11):1430-48. doi: 10.1016/s1357-2725(02)00041-9.
- 12. Bi J, Guo A, Lai Y, Li B, Zhong J, Wu H, et al. Overexpression of clusterin correlates with tumor progression, metastasis in gastric cancer: a study on tissue microarrays. Neoplasma. 2010;57(3):191. doi: 10.4149/neo_2010_03_191.
- 13. Won JC, Park CY, Oh SW, Lee ES, Youn BS, Kim MS. Plasma clusterin (ApoJ) levels are associated with adiposity and systemic inflammation. PLoS One. 2014;9(7): e103351. doi: 10.1371/journal.pone.0103351.
- 14. Seo JA, Kang M-C, Ciaraldi TP, Kim SS, Park KS, Choe C, et al. Circulating ApoJ is closely associated with insulin resistance in human subjects. Metabolism. 2018; 78:155-66. doi: 10.1016/j.metabol.2017.09.014.
- 15. Jeon YK, Kim SS, Kim JH, Kim HJ, Kim HJ, Park JJ, et al. Combined aerobic and resistance exercise training reduces circulating apolipoprotein J levels and improves insulin resistance in postmenopausal diabetic women. Diabetes Metab 2020;44(1):103-12. I 10.4093/dmj.2018.0160.
- Navab M, Anantharamaiah G, Reddy ST, Van Lenten BJ, Wagner AC, Hama S, et al. An Oral ApoJ Peptide Renders HDL Anti-

- inflammatory in Mice and Monkeys and Dramatically Reduces Atherosclerosis in Apolipoprotein E–Null Mice. Arterioscler Thromb Vasc Biol. 2005;25(9):1932-7. doi: 10.1161/01.ATV.0000174589. 70190.e2.
- Yang N, Qin Q. Apolipoprotein J: a new predictor and therapeutic target in cardiovascular disease? Chin Med J. 2015;128(18):2530-4. doi: 10.4103/0366-6999.164983.
- 18. Zanuso S, Jimenez A, Pugliese G, Corigliano G, Balducci S. Exercise for the management of type 2 diabetes: a review of the evidence. Acta Diabetol. 2010;47(1):15-22. doi: 10.1007/s00592-009-0126-3.
- 19. Oliveira C, Simões M, Carvalho J, Ribeiro J. Combined exercise for people with type 2 diabetes mellitus: a systematic review. Diabetes Res Clin Pract. 2012;98(2):187-98. doi: 10.1016/j.diabres.2012.08.004.
- 20. Colberg SR, Sigal RJ, Fernhall B, Regensteiner JG, Blissmer BJ, Rubin RR, et al. Exercise and type 2 diabetes: the American College of Sports Medicine and the American Diabetes Association: joint position statement. Med Sci Sports Exerc. 2010;42(12):2282-303. doi: 10.1249/MSS.0b013e3181eeb61c.
- 21. Rull A, Martínez-Bujidos M, Pérez-Cuellar M, Pérez A, Ordóñez-Llanos J, Sánchez-Quesada JL. Increased concentration of clusterin/apolipoprotein J (apoJ) in hyperlipemic serum is paradoxically associated with decreased apoJ content in lipoproteins. Atherosclerosis. 2015;241(2):463-70. doi: 10.1016/j.atherosclerosis.2015.06.003.
- 22. Brzycki M. Strength testing—predicting a one-rep max from reps-to-fatigue. J Phys Educ Recreat Dance. 1993;64(1):88-90. doi.org/10.1080/07303084.1993.10606684.
- 23. Zanuso S, Bergamin M, Jimenez A, Pugliese G, D'Errico V, Nicolucci A, et al. Determination of metabolic equivalents during low-and high-intensity resistance exercise in healthy young subjects and patients with type 2 diabetes. Biol Sport. 2016;33(1):77. doi: 10.5604/20831862.1194124.
- Romero-Arenas S, Martínez-Pascual M, Alcaraz PE. Impact of resistance circuit training on neuromuscular, cardiorespiratory and body composition adaptations in the elderly. Aging Dis. 2013;4(5):256. doi: 10.14336/AD.2013.0400256.
- Baechle TR, Earle RW. Essentials of strength training and conditioning: Human Kinetics. 2008.
- Mukaimoto T, Ohno M. Effects of circuit low-intensity resistance exercise with slow movement on oxygen consumption during

- and after exercise. J Sports Sci. 2012;30(1):79-90. doi: 10.1080/02640414.2011.616950.
- 27. Sarlak Z. Effects of Eight Weeks Aerobic Training on Serum Apo AI, APO B and lipid profile in Overweight Women. Sport Physiology. 2016;7(28):45-58. doi: 20.1001.1.2322164.1394.7.28.3.9.
- 28. Trougakos IP, Poulakou M, Stathatos M, Chalikia A, Melidonis A, Gonos ES. Serum levels of the senescence biomarker clusterin/apolipoprotein J increase significantly in diabetes type II and during development of coronary heart disease or at myocardial infarction. Exp Gerontol. 2002;37(10-11):1175-87. doi: 10.1016/s0531-5565(02)00139-0.
- 29. Aronis KN, Vamvini MT, Chamberland JP, Mantzoros CS. Circulating clusterin (apolipoprotein J) levels do not have any day/night variability and are positively associated with total and LDL cholesterol levels in young healthy individuals. J Clin Endocrinol Metab. 2011;96(11): doi: 10.1210/jc.2011-1555.
- 30. Sheikholeslami Vatani D, Ahmadi S, Ahmadi Dehrashid K, Gharibi F. Changes in cardiovascular risk factors and inflammatory markers of young, healthy, men after six weeks of moderate or high intensity resistance training. J Sports Med Phys Fitness. 2011;51(4):695-700.
- 31. Valente EA, Sheehy ME, Avila JJ, Gutierres JA, Delmonico MJ, Lofgren IE. The effect of the addition of resistance training to a dietary education intervention on apolipoproteins and diet quality in overweight and obese older adults. Clin Interv Aging. 2011; 6:235. doi: 10.2147/CIA.S23583.
- 32. Thompson PD, Buchner D, Piña IL, Balady GJ, Williams MA, Marcus BH, et al. Exercise and physical activity in the prevention and treatment of atherosclerotic cardiovascular disease: a statement from the Council on Clinical Cardiology (Subcommittee on Exercise, Rehabilitation, and Prevention) and the Council on Nutrition, Physical Activity, and Metabolism (Subcommittee on Physical Activity). Circulation. 2003;107(24):3109-16. 10.1161/01.CIR.0000075572.40158.77.
- 33. Behre C, Bergström G, Schmidt C. Moderate physical activity is associated with lower ApoB/ApoA-I ratios independently of other risk factors in healthy, middle-aged men. Angiology. 2010;61(8):775-9. doi: 10.1177/0003319710373746.
- 34. De Groot P, Hjeltnes N, Heijboer A, Stal W, Birkeland K. Effect of training intensity on physical capacity, lipid profile and insulin sensitivity in early rehabilitation of spinal

- cord injured individuals. Spinal Cord. 2003;41(12):673-9. doi: 10.1038/sj.sc.3101534.
- 35. Kannan Ü, Vasudevan K, Balasubramaniam K, Yerrabelli D, Shanmugavel K, John NA. Effect of exercise intensity on lipid profile in sedentary obese adults. Journal of clinical and diagnostic research: JCDR. 2014;8(7): BC08. doi: 10.7860/JCDR/2014/8519.4611.
- 36. Baharloo S, Shakeri N, Ebrahim K, Ramezani Tehrani F, Allameh Z. The effect of 12 weeks water-based Tabata training on insulin resistance, apolipoprotein A and apolipoprotein B in obese women with polycystic ovary syndrome. RJMS. 2021;28(5):11-20.