

LOW DENSITY LIPOPROTEINS ACT AS NEGATIVE ACUTE PHASE PROTEINS IN THE METABOLIC SYNDROME

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Abstract

Background: We tried to understand whether or not low density lipoproteins (LDL) act as negative acute phase proteins (APP) in the metabolic syndrome.

Methods: Patients with plasma triglycerides values lower than 60 mg/dL were collected into the first, lower than 100 mg/dL into the second, lower than 150 mg/dL into the third, lower than 200 mg/dL into the fourth, and 200 mg/dL and greater into the fifth groups, respectively.

Results: The study included 875 cases (370 males and 505 females), totally. Although the mean age, male ratio, smoking, body mass index (BMI), fasting plasma glucose (FPG), white coat hypertension (WCH), hypertension (HT), diabetes mellitus (DM), chronic obstructive pulmonary disease (COPD), coronary heart disease (CHD), and chronic renal disease (CRD) increased parallel to the increased plasma triglycerides values, continuously ($p < 0.05$ nearly in all steps), the mean LDL values increased just up to plasma triglycerides values of 200 mg/dL, but then decreased, significantly (137.5 versus 129.0 mg/dL, $p = 0.020$).

Conclusions: The increased plasma triglycerides values may be one of the most sensitive parameters of the metabolic syndrome that is characterized with disseminated endothelial damage, inflammation, fibrosis, accelerated atherosclerosis, end-organ insufficiencies, early aging, and premature death. Although the mean age, male ratio, smoking, BMI, FPG, WCH, HT, DM, COPD, CHD, and CRD increased parallel to the increased plasma triglycerides values, continuously, LDL was the only metabolic parameter that increased just up to the plasma triglycerides values of 200 mg/dL but then decreased, significantly. So LDL acts as negative APP in the metabolic syndrome.

Key words: Low density lipoproteins, triglycerides, acute phase proteins, metabolic syndrome

Introduction

Chronic endothelial damage may be the most common sort of vasculitis, and the leading cause of early aging and premature death in the human being (1-4). Much higher blood pressure (BP) of the afferent vasculature may be the major underlying mechanism by inducing recurrent injuries on vascular endothelium. Probably, whole afferent vasculature including capillaries are predominantly involved in the process. Therefore the term of venosclerosis is not as famous as atherosclerosis in the medical literature. Because of the chronic endothelial damage, inflammation, edema, and fibrosis, vascular walls thicken, their lumens narrow, and they lose their elastic nature which reduces blood flow to terminal organs, and increases systolic BP further. Some of the well-known components of the inflammatory process are physical inactivity, animal-rich diet, overweight, smoking, alcohol, hypertriglyceridemia, hyperbetalipoproteinemia, dyslipidemia, impaired fasting glucose, impaired glucose tolerance, white coat hypertension (WCH), rheumatologic disorders, chronic infections, and cancers for the development of terminal endpoints including obesity, hypertension (HT), diabetes mellitus (DM), cirrhosis, peripheral artery disease (PAD), chronic obstructive pulmonary disease (COPD), coronary heart disease (CHD), chronic renal disease (CRD), mesenteric ischemia, osteoporosis, stroke, early aging, and premature death (5-10). Although early withdrawal of the predisposing factors may delay terminal consequences, after development of HT, DM, cirrhosis, COPD, CRD, CHD, PAD, mesenteric ischemia, osteoporosis, stroke, and aging, endothelial changes cannot be reversed completely due to their fibrotic nature. Up to now, the predisposing factors and terminal endpoints were researched under the titles of metabolic syndrome, aging syndrome, or accelerated endothelial damage syndrome in the medicine, extensively (11-14). Although its normal limits could not be determined clearly yet, increased plasma triglycerides may be one of the most sensitive parameters of the metabolic syndrome (15-17). Due to the growing evidence about the strong association between higher plasma triglycerides values and prevalence of CHD, Adult Treatment Panel (ATP) III adopts lower cutpoints for triglycerides abnormalities than did ATP II (18, 19). Although ATP II determined the normal plasma triglycerides value as lower than 200 mg/dL in 1994 (19), World Health Organisation in 1999 (20) and ATP III in 2001 reduced their normal limit as lower than 150 mg/dL (18). Although these cutpoints are usually used to define limits of the metabolic syndrome, there are still suspicions about the safest value of plasma triglycerides in the medical literature (16, 17). Though the absolute sensitivity of plasma triglycerides in the metabolic syndrome, role of low density lipoproteins (LDL) is suspicious (21). We tried to understand whether or not LDL act as negative acute phase proteins (APP) in the metabolic syndrome.

Material and Methods

The study was performed in the Internal Medicine Polyclinic of the Dumlupinar University between August 2005 and March 2007. Consecutive patients at and above the age of 15 years were included. Their medical histories including HT, DM, COPD, and already used medications were learned, and a routine check up procedure including fasting plasma glucose (FPG), triglycerides, and LDL was performed. Current daily smokers with six pack-months and cases with a history of three pack-years were accepted as smokers. Patients with devastating illnesses including type 1 DM, malignancies, acute or chronic renal failure, chronic liver diseases, hyper- or hypothyroidism, and heart failure were excluded to avoid their possible effects on weight. Additionally, anti-hyperlipidemic drugs, metformin, and/or acarbose users were excluded to avoid their possible effects on blood lipid profiles and/or body weight (22, 23). Body mass index (BMI) of each case was calculated by the measurements of the same physician instead of verbal expressions. Weight in kilograms is divided by height in meters squared (18). Cases with an overnight FPG value of 126 mg/dL or greater on two occasions or already using antidiabetic medications were defined as diabetics (18). An oral glucose tolerance test with 75-gram glucose was performed in cases with a FPG value between 110 and 126 mg/dL, and diagnosis of cases with a 2-hour plasma glucose value of 200 mg/dL or greater is DM (18). CRD is diagnosed with a persistently elevated serum creatinine level of 1.3 mg/dL in males and 1.2 mg/dL in females. Additionally, office blood pressure (OBP) was checked after a 5 minute rest in seated position with a mercury sphygmomanometer on three visits, and no smoking was permitted during the previous 2 hours. A 10-day twice daily measurement of blood pressure at home (HBP) was obtained in all cases, even in the normotensives in the office due to the risk of masked HT after a 10 minute education session about proper BP measurement techniques (24). An additional 24-hour ambulatory blood pressure monitoring was not needed due to its similar effectivity with the HBP measurements (3). Eventually, HT is defined as a mean BP of 135/85 mmHg or greater on HBP measurements, and WCH as an OBP of 140/90 mmHg or greater but a mean HBP measurement of lower than 135/85 mmHg (24). An exercise electrocardiogram is performed just in cases with an abnormal electrocardiogram and/or angina pectoris. Coronary angiography is taken just for the exercise electrocardiogram positive cases. So CHD is diagnosed either angiographically or with the Doppler echocardiographic findings as the already developed movement disorders in the cardiac walls. The spirometric pulmonary function tests were performed in required cases after the physical examination, and the criterion for diagnosis of COPD is post-bronchodilator forced expiratory volume in one second/forced vital capacity of less than 70% (25). Eventually, patients with plasma triglycerides values lower than 100 mg/dL were collected into the first, lower than 150 mg/dL into the second, lower than 200 mg/dL into the third, and 200 mg/dL and higher into the fourth groups, respectively. The male ratio, mean age, BMI, FPG, triglycerides, and LDL, and prevalence

of smoking, WCH, HT, DM, COPD, CHD, and CRD were detected in each group and compared in between. Mann-Whitney U test, Independent-Samples T test, and comparison of proportions were used as the methods of statistical analyses.

Results

The study included 875 cases (370 males and 505 females), totally. Although the mean age, male ratio, smoking, BMI, FPG, WCH, HT, DM, COPD, CHD, and CRD increased parallel to the increased triglycerides values, continuously ($p < 0.05$ nearly in all steps), the mean LDL values increased just up to plasma triglycerides values of 200 mg/dL but then decreased, significantly (137.5 versus 129.0 mg/dL, $p = 0.020$) (Table 1 - next page).

Discussion

Excess weight may lead to both structural and functional abnormalities of many organs of the body. Adipose tissues produce leptin, tumor necrosis factor- α , plasminogen activator inhibitor-1, and adiponectin-like cytokines acting as acute phase reactants in the plasma (26, 27). Excess weight-induced chronic low-grade vascular endothelial inflammation may play a significant role in the pathogenesis of accelerated atherosclerosis in the whole body (1, 2). Additionally, excess weight may cause an increased blood volume as well as an increased cardiac output thought to be the result of increased oxygen need of the excessive fat tissue. The prolonged increase in the blood volume may lead to myocardial hypertrophy terminating with a decreased cardiac compliance. Combination of these cardiovascular risk factors will eventually terminate with increased left ventricular stroke work and risks of arrhythmias, cardiac failure, and sudden cardiac death. Similarly, the prevalence of CHD and stroke increased parallel to the increased BMI values in the other studies (28, 29), and risk of death from all causes including cancers increased throughout the range of moderate to severe weight excess in all age groups (30). The relationship between excess weight, elevated BP, and plasma triglycerides is described in the metabolic syndrome (15), and clinical manifestations of the syndrome include obesity, dyslipidemia, HT, insulin resistance, and proinflammatory and prothrombotic states (13). Similarly, prevalence of smoking (42.2% versus 28.4%, $p < 0.01$), excess weight (83.6% versus 70.6%, $p < 0.01$), DM (16.3% versus 10.3%, $p < 0.05$), and HT (23.2% versus 11.2%, $p < 0.001$) were all higher in the hypertriglyceridemia group in the other study (31). On the other hand, the prevalence of increased LDL cases were similar both in the hypertriglyceridemia (200 mg/dL and higher) and control groups (18.9% versus 16.3%, $p > 0.05$, respectively) in the above study (31). Similarly, plasma LDL values increased just up to the plasma triglycerides values of 200 mg/dL but then decreased significantly in the present study, too ($p < 0.05$ for all).

Smoking may be found among one of the most common causes of vasculitis all over the world. It causes a chronic inflammatory process on the vascular endothelium, probably depending on the concentration of smoke that terminates with an accelerated atherosclerosis, end-organ insufficiencies, early aging, and premature death. Therefore smoking has to be included among the major components of the metabolic syndrome. Strong and terminal atherosclerotic effects of smoking are the most obviously seen in Buerger's disease (thromboangiitis obliterans). It is an obliterative vasculitis characterized by inflammatory changes in the small and medium-sized arteries and veins, and it has never been reported in the absence of smoking in the medical literature. Although the well-known strong atherosclerotic effects of smoking, smoking in the human being and nicotine administration in animals may be associated with decreased BMI values (32). Evidence revealed an increased energy expenditure during smoking both on rest and light physical activity (33), and nicotine supplied by patch after smoking cessation decreased caloric intake in a dose-related manner (34). According to an animal study, nicotine may lengthen intermeal time and decrease amount of meal eaten (35). Additionally, the mean BMI seems to be the highest in the former, the lowest in the current and medium in never smokers (36). Smoking may be associated with a postcessation weight gain (37). Similarly, although CHD was detected with similar prevalence in both genders, prevalence of smoking and COPD were higher in males against the higher BMI, LDL, triglycerides, WCH, HT, and DM in females (38). The result may show both the strong atherosclerotic and weight decreasing roles of smoking (39). Similarly, the incidence of a myocardial infarction is increased six-fold in women and three-fold in men who smoke 20 cigarettes per day (40). In another definition, smoking may be more dangerous for women probably due to the associated higher BMI and its consequences in them. Parallel to the above results, the proportion of smokers is consistently higher in men in the literature (23). So smoking is probably a powerful atherosclerotic risk factor with some suppressor effects on appetite. Smoking-induced weight loss may be related with the smoking-induced chronic vascular endothelial inflammation all over the body, since loss of appetite is one of the main symptoms of the disseminated inflammations in the body. Physicians can even understand healing of the patients via their normalizing appetite. Several toxic substances found in cigarette smoke get into the circulation by means of the respiratory tract, and cause a vascular endothelial inflammation until their clearance from the circulation. But due to the repeated smoking habit of individuals, the clearance process never terminates. So the patients become ill with loss of appetite, permanently. In another explanation, smoking-induced weight loss is an indicator of being ill instead of being healthy (34-36). After smoking cessation, normal appetite comes back with a prominent weight gain but the returned weight is the patients' physiological weight, actually.

Table 1: Characteristics features of the study cases according to the plasma triglycerides values

Variable	Lower than 60 mg/dL	p-value	Lower than 100 mg/dL	p-value	Lower than 150 mg/dL	p-value	Lower than 200 mg/dL	p-value	200 mg/dL or greater
Number of cases	84		207		235		148		201
Age (year)	<u>35.6 ± 16.4</u> (17-79)	<u>0.000</u>	<u>43.6 ± 17.5</u> (16-83)	<u>0.009</u>	<u>47.7 ± 15.3</u> (16-82)	<u>0.018</u>	<u>51.2 ± 12.6</u> (19-82)	Ns*	49.8 ± 12.3 (19-88)
Male ratio	<u>30.9%</u>	<u>0.05></u>	<u>39.1%</u>	Ns	<u>40.4%</u>	Ns	<u>43.9%</u>	<u>0.05></u>	<u>51.2%</u>
Smoking	<u>16.6%</u>	Ns	<u>21.7%</u>	Ns	<u>26.3%</u>	Ns	23.6%	<u>0.001></u>	<u>38.3%</u>
BMI† (kg/m ²)	<u>24.6 ± 5.3</u> (16.7-45.9)	<u>0.002</u>	<u>27.1 ± 5.9</u> (16.7-49.3)	<u>0.000</u>	<u>29.4 ± 6.1</u> (18.4-51.0)	Ns	<u>29.9 ± 4.8</u> (19.2-49.0)	Ns	<u>30.0 ± 5.0</u> (21.0-51.1)
FPG‡ (mg/dL)	<u>96.5 ± 35.3</u> (71-377)	<u>0.016</u>	<u>106.6 ± 48.7</u> (59-400)	Ns	<u>106.8 ± 35.1</u> (71-335)	<u>0.006</u>	<u>117.3 ± 47.8</u> (68-386)	Ns	<u>124.3 ± 55.3</u> (74-392)
Triglyceride (mg/dL)	<u>51.0 ± 7.5</u> (27-59)	<u>0.000</u>	<u>78.3 ± 10.8</u> (60-99)	<u>0.000</u>	<u>122.2 ± 14.5</u> (100-149)	<u>0.000</u>	<u>174.1 ± 14.2</u> (150-199)	<u>0.000</u>	<u>325.8 ± 160.4</u> (200-1,350)
LDL§ (mg/dL)	<u>98.6 ± 23.3</u> (56-161)	<u>0.000</u>	<u>114.6 ± 33.0</u> (31-269)	<u>0.000</u>	<u>131.1 ± 31.7</u> (56-228)	<u>0.033</u>	<u>137.5 ± 32.4</u> (50-237)	<u>0.020</u>	<u>129.0 ± 40.8</u> (10-239)
WCH	<u>17.8%</u>	<u>0.05></u>	<u>24.1%</u>	<u>0.05></u>	<u>31.0%</u>	Ns	<u>35.1%</u>	Ns	32.3%
HT**	<u>8.3%</u>	<u>0.001></u>	<u>15.9%</u>	<u>0.05></u>	<u>21.2%</u>	Ns	<u>22.2%</u>	Ns	<u>26.3%</u>
DM***	<u>2.3%</u>	<u>0.001></u>	<u>11.1%</u>	Ns	<u>13.6%</u>	Ns	<u>18.2%</u>	<u>0.05></u>	<u>24.3%</u>
COPD****	<u>4.7%</u>	<u>0.01></u>	<u>9.1%</u>	<u>0.01></u>	<u>14.0%</u>	Ns	12.8%	<u>0.05></u>	<u>18.4%</u>
CHD*****	<u>4.7%</u>	<u>0.001></u>	<u>10.1%</u>	Ns	<u>11.4%</u>	Ns	<u>14.8%</u>	Ns	11.9%
CRD*****	0.0%	Ns	1.9%	Ns	<u>0.4%</u>	<u>0.01></u>	<u>2.0%</u>	<u>0.01></u>	<u>4.9%</u>

*Nonsignificant (p>0.05) †Body mass index ‡Fasting plasma glucose §Low density lipoproteins

||White coat hypertension **Hypertension ***Diabetes mellitus ****Chronic obstructive pulmonary disease

*****Coronary heart disease *****Chronic renal disease

Although the several negative effects of excess weight on health, nearly three-quarters of cases above the age of 30 years have excess weight (41). The prevalence of excess weight increases by decades, particularly after the third decade, up to the eighth decade of life (41). So 30th and 70th years of age may be the breaking points of life for weight, and aging may be the major determiner factor of excess weight. Probably, partially decreased physical and mental stresses after the age of 30 years and debility and comorbid disorders-induced restrictions after the age of 70 years may be the major causes for the changes of BMI values at these ages. Interestingly, the mean age and BMI increased just up to the plasma triglycerides values of 200 and 150 mg/dL, respectively here. So smoking remained as the solitary causative factor for the hypertriglyceridemia after the plasma triglycerides values of 200 mg/dL in the present study.

Although ATP III reduced the normal limit of plasma triglycerides values as lower than 150 mg/dL in 2001 (18), much lower limits may provide additional benefit for health (16, 17). In the present study, prevalence of smoking was the highest in the highest triglycerides having group which may also indicate inflammatory roles of smoking in the metabolic syndrome, since triglycerides may actually be some acute phase reactants in the metabolic syndrome. The mean age, male ratio, smoking, BMI, FPG, WCH, HT, DM, COPD, CHD, and CRD increased parallel to the plasma triglycerides values from the first towards the fifth groups, continuously. In our opinion, significantly increased plasma triglycerides values by aging may be secondary to aging-induced decreased physical and mental stresses; those eventually terminate with onset of excess weight and many associated health problems. Interestingly, the mean age increased just up to the plasma triglycerides values of 200 mg/dL in the present study. Similarly, the mean BMI values increased just up to the plasma triglycerides values of 150 mg/dL, significantly. These trends may be due to the fact that although the borderline high triglycerides values (150-199 mg/dL) is seen together with physical inactivity and overweight, the high triglycerides (200-499 mg/dL) and very high triglycerides values (500 mg/dL or greater) may be secondary to both genetic factors and terminal consequences of the metabolic syndrome including smoking, obesity, DM, HT, COPD, cirrhosis, CRD, PAD, CHD, and stroke (18). But although the underlying causes of the high and very high plasma triglycerides values may be a little bit different, probably risks of the terminal endpoints of the metabolic syndrome do not change in them. For example, prevalence of HT, DM, COPD, and CRD were the highest in the highest triglycerides having group in the present study. Eventually, although some authors reported that lipid assessment can be simplified as the measurements of total cholesterol and high density lipoproteins (HDL) values alone (42), the present study and most of the others indicated significant relationships between plasma triglycerides and LDL values and terminal consequences of the metabolic syndrome (43).

APP are a class of proteins whose plasma concentrations increase (positive APP) or decrease (negative APP) as a response to inflammation, infection, and tissue damage (44-46). In case of inflammation, infection, and tissue damage, local inflammatory cells (neutrophils and macrophages) secrete several kinds of cytokines into the blood, most notable of which are the interleukins. The liver responds by producing many positive APP. At the same time, production of many proteins is reduced. Therefore these proteins are called negative APP. Some of the well-known negative APP are albumin, transferrin, retinol-binding protein, antithrombin, and transcortin. The decrease of such proteins is also used as an indicator of inflammation. The physiological role of decreased synthesis of such proteins is generally to save amino acids for producing positive APP more effectively. Due to the decreased production of some proteins in liver during severe inflammatory conditions, production of LDL may also be suppressed. Similarly, although the mean triglycerides, fibrinogen, C-reactive protein, and glucose values were significantly higher in cases with ischemic stroke, the oxidized LDL values did not correlate with age, stroke severity, and outcome in the other study (47). Additionally, significant alterations occur in lipid metabolism and lipoprotein composition during infections, and triglycerides increase whereas HDL and LDL decrease in another study (48). Furthermore, a 10 mg/dL increase of LDL was associated with a 3% lower risk of hemorrhagic stroke in another study (49).

As a conclusion, increased plasma triglycerides values may be one of the most sensitive parameters of the metabolic syndrome that is characterized with disseminated endothelial damage, inflammation, fibrosis, accelerated atherosclerosis, end-organ insufficiencies, early aging, and premature death. Although the mean age, male ratio, smoking, BMI, FPG, WCH, HT, DM, COPD, CHD, and CRD increased parallel to the increased plasma triglycerides values, continuously, LDL was the only metabolic parameter that increased just up to the plasma triglycerides values of 200 mg/dL but then decreased, significantly. So LDL act as negative APP in the metabolic syndrome.

References

1. Widlansky ME, Gokce N, Keaney JF Jr, Vita JA. The clinical implications of endothelial dysfunction. *J Am Coll Cardiol* 2003; 42(7): 1149-1160.
2. Ridker PM. High-sensitivity C-reactive protein: potential adjunct for global risk assessment in the primary prevention of cardiovascular disease. *Circulation* 2001; 103(13): 1813-1818.
3. Helvacı MR, Seyhanlı M. What a high prevalence of white coat hypertension in society! *Intern Med* 2006; 45(10): 671-674.
4. Helvacı MR, Kaya H, Seyhanlı M, Cosar E. White coat hypertension is associated with a greater all-cause mortality. *J Health Sci* 2007; 53(2): 156-160.

5. Helvaci MR, Kaya H, Yalcin A, Kuvandik G. Prevalence of white coat hypertension in underweight and overweight subjects. *Int Heart J* 2007; 48(5): 605-613.
6. Helvaci MR, Kaya H, Duru M, Yalcin A. What is the relationship between white coat hypertension and dyslipidemia? *Int Heart J* 2008; 49(1): 87-93.
7. Helvaci MR, Kaya H, Seyhanli M, Yalcin A. White coat hypertension in definition of metabolic syndrome. *Int Heart J* 2008; 49(4): 449-457.
8. Helvaci MR, Kaya H, Sevinc A, Camci C. Body weight and white coat hypertension. *Pak J Med Sci* 2009; 25(6): 916-921.
9. Helvaci MR, Sevinc A, Camci C, Yalcin A. Treatment of white coat hypertension with metformin. *Int Heart J* 2008; 49(6): 671-679.
10. Helvaci MR, Aydin Y, Gundogdu M. Smoking induced atherosclerosis in cancers. *HealthMED* 2012; 6(11): 3744-3749.
11. Eckel RH, Grundy SM, Zimmet PZ. The metabolic syndrome. *Lancet* 2005; 365(9468): 1415-1428.
12. Grundy SM, Brewer HB Jr, Cleeman JI, Smith SC Jr, Lenfant C. Definition of metabolic syndrome: Report of the National Heart, Lung, and Blood Institute/American Heart Association conference on scientific issues related to definition. *Circulation* 2004; 109(3): 433-438.
13. Tonkin AM. The metabolic syndrome(s)? *Curr Atheroscler Rep* 2004; 6(3): 165-166.
14. Franklin SS, Barboza MG, Pio JR, Wong ND. Blood pressure categories, hypertensive subtypes, and the metabolic syndrome. *J Hypertens* 2006; 24(10): 2009-2016.
15. Helvaci MR, Kaya H, Gundogdu M. Association of increased triglyceride levels in metabolic syndrome with coronary artery disease. *Pak J Med Sci* 2010; 26(3): 667-672.
16. Helvaci MR, Tonyali O, Abyad A, Pocock L. The safest value of plasma triglycerides. *World Family Med* 2019; 17(7): 22-27.
17. Helvaci MR, Abyad A, Pocock L. The lowest is the safest value of plasma triglycerides. *World Family Med* 2019; 17(10): 10-15.
18. Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) final report. *Circulation* 2002; 106(25): 3143-3421.
19. National Cholesterol Education Program. Second Report of the Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel II). *Circulation* 1994; 89(3): 1333-1445.
20. World Health Organization. Definition, Diagnosis and Classification of Diabetes Mellitus and its Complications. Report of a WHO consultation 1999.
21. Fodor JG, Tzerovska R, Dorner T, Rieder A. Do we diagnose and treat coronary heart disease differently in men and women? *Wien Med Wochenschr* 2004; 154(17-18): 423-425.
22. Helvaci MR, Kaya H, Borazan A, Ozer C, Seyhanli M, Yalcin A. Metformin and parameters of physical health. *Intern Med* 2008; 47(8): 697-703.
23. Helvaci MR, Aydin Y, Varan G, Abyad A, Pocock L. Acarbose versus metformin in the treatment of metabolic syndrome. *World Family Med* 2018; 16(5): 10-15.
24. O'Brien E, Asmar R, Beilin L, Imai Y, Mallion JM, Mancia G, et al. European Society of Hypertension recommendations for conventional, ambulatory and home blood pressure measurement. *J Hypertens* 2003; 21(5): 821-848.
25. Vestbo J, Hurd SS, Agustí AG, Jones PW, Vogelmeier C, Anzueto A, et al. Global strategy for the diagnosis, management, and prevention of chronic obstructive pulmonary disease: GOLD executive summary. *Am J Respir Crit Care Med* 2013; 187(4): 347-65.
26. Funahashi T, Nakamura T, Shimomura I, Maeda K, Kuriyama H, Takahashi M, et al. Role of adipocytokines on the pathogenesis of atherosclerosis in visceral obesity. *Intern Med* 1999; 38(2): 202-206.
27. Yudkin JS, Stehouwer CD, Emeis JJ, Coppack SW. C-reactive protein in healthy subjects: associations with obesity, insulin resistance, and endothelial dysfunction: a potential role for cytokines originating from adipose tissue? *Arterioscler Thromb Vasc Biol* 1999; 19(4): 972-978.
28. Zhou B, Wu Y, Yang J, Li Y, Zhang H, Zhao L. Overweight is an independent risk factor for cardiovascular disease in Chinese populations. *Obes Rev* 2002; 3(3): 147-156.
29. Zhou BF. Effect of body mass index on all-cause mortality and incidence of cardiovascular diseases--report for meta-analysis of prospective studies open optimal cut-off points of body mass index in Chinese adults. *Biomed Environ Sci* 2002; 15(3): 245-252.
30. Calle EE, Thun MJ, Petrelli JM, Rodriguez C, Heath CW Jr. Body-mass index and mortality in a prospective cohort of U.S. adults. *N Engl J Med* 1999; 341(15): 1097-1105.
31. Helvaci MR, Aydin LY, Maden E, Aydin Y. What is the relationship between hypertriglyceridemia and smoking? *Middle East J Age and Ageing* 2011; 8(6).
32. Grunberg NE, Greenwood MR, Collins F, Epstein LH, Hatsukami D, Niaura R, et al. National working conference on smoking and body weight. Task Force 1: Mechanisms relevant to the relations between cigarette smoking and body weight. *Health Psychol* 1992; 11: 4-9.
33. Walker JF, Collins LC, Rowell PP, Goldsmith LJ, Moffatt RJ, Stamford BA. The effect of smoking on energy expenditure and plasma catecholamine and nicotine levels during light physical activity. *Nicotine Tob Res* 1999; 1(4): 365-370.
34. Hughes JR, Hatsukami DK. Effects of three doses of transdermal nicotine on post-cessation eating, hunger and weight. *J Subst Abuse* 1997; 9: 151-159.

35. Miyata G, Meguid MM, Varma M, Fetissov SO, Kim HJ. Nicotine alters the usual reciprocity between meal size and meal number in female rat. *Physiol Behav* 2001; 74(1-2): 169-176.
36. Laaksonen M, Rahkonen O, Prattala R. Smoking status and relative weight by educational level in Finland, 1978-1995. *Prev Med* 1998; 27(3): 431-437.
37. Froom P, Melamed S, Benbassat J. Smoking cessation and weight gain. *J Fam Pract* 1998; 46(6): 460-464.
38. Helvaci MR, Kaya H, Gundogdu M. Gender differences in coronary heart disease in Turkey. *Pak J Med Sci* 2012; 28(1): 40-44.
39. Helvaci MR, Aydin Y, Gundogdu M. Atherosclerotic effects of smoking and excess weight. *J Obes Wt Loss Ther* 2012; 2: 145.
40. Prescott E, Hippe M, Schnohr P, Hein HO, Vestbo J. Smoking and risk of myocardial infarction in women and men: longitudinal population study. *BMJ* 1998; 316(7137): 1043-1047.
41. Helvacı MR, Kaya H, Ozer C. Aging may be the major determiner factor of excess weight. *Middle East J Age and Ageing* 2008; 5(2).
42. Di Angelantonio E, Sarwar N, Perry P, Kaptoge S, Ray KK, Thompson A, et al. Major lipids, apolipoproteins, and risk of vascular disease. *JAMA* 2009; 302(18): 1993-2000.
43. Sarwar N, Sandhu MS, Ricketts SL, Butterworth AS, Di Angelantonio E, Boekholdt SM, et al. Triglyceride-mediated pathways and coronary disease: collaborative analysis of 101 studies. *Lancet* 2010; 375(9726): 1634-1639.
44. Gabay C, Kushner I. Acute-phase proteins and other systemic responses to inflammation. *N Engl J Med* 1999; 340(6): 448-454.
45. Schrödl W, Büchler R, Wendler S, Reinhold P, Muckova P, Reindl J, et al. Acute phase proteins as promising biomarkers: Perspectives and limitations for human and veterinary medicine. *Proteomics Clin Appl* 2016; 10(11): 1077-1092.
46. Wool GD, Reardon CA. The influence of acute phase proteins on murine atherosclerosis. *Curr Drug Targets* 2007; 8(11): 1203-1214.
47. Vibo R, Körv J, Roose M, Kampus P, Muda P, Zilmer K, et al. Acute phase proteins and oxidised low-density lipoprotein in association with ischemic stroke subtype, severity and outcome. *Free Radic Res* 2007; 41(3): 282-287.
48. Pirillo A, Catapano AL, Norata GD. HDL in infectious diseases and sepsis. *Handb Exp Pharmacol* 2015; 224: 483-508.
49. Ma C, Na M, Neumann S, Gao X. Low-density lipoprotein cholesterol and risk of hemorrhagic stroke: a systematic review and dose-response meta-analysis of prospective studies. *Curr Atheroscler Rep* 2019; 21(12): 52.