



## JOURNAL OF PHARMACEUTICAL AND BIOMEDICAL SCIENCES

Marjani Abdoljalal et al. **Age related metabolic syndrome among Fars ethnic women in Gorgan, Iran.** *Journal of pharmaceutical and biomedical sciences (J Pharm Biomed Sci.)* 2013, May; 30(30): 929-935. (Article no 09)

The online version of this article, along with updated information and services, is located on the World Wide Web at: [www.jpbms.info](http://www.jpbms.info)

*Journal of Pharmaceutical and Biomedical Sciences (J Pharm Biomed Sci.)*, **Member journal.**  
**Committee of Publication ethics (COPE) and Journal donation project (JDP).**

*Original article*

**Age related metabolic syndrome among Fars ethnic women in Gorgan, Iran.**

**Abdoljalal Marjani<sup>1\*</sup> & Najmeh Shahini<sup>2</sup>**

<sup>1</sup>Department of Biochemistry and Biophysics, Metabolic Disorders Research Center, Gorgan Faculty of Medicine, Golestan University of Medical Sciences, Gorgan, Golestan province, Iran.

<sup>2</sup> Medical Student, Student Reaserch Committee, Golestan University of Medical Sciences, Gorgan, Iran.

**Abstract:**

*Background:* Studies have been shown that the prevalence of metabolic syndrome changes among different ethnic and sex groups. *Objective:* We determined age related prevalence of metabolic syndrome among Fars ethnic women. *Study design:* Health center based study. *Setting:* Metabolic Disorders Research Center in Gorgan. *Participants:* 160 Fars adult women. *Sampling:* randomized sampling method. *Results:* The most age distribution was in ages from 35 to 40 years (50%). There were significant differences between the mean value of waist circumferences, fasting blood glucose, body mass index and triglyceride among subjects with and without metabolic syndrome in age groups 20-24 and 25-29 years old ( $P < 0.05$ ). HDL-cholesterol and LDL-cholesterol levels were significantly decreased and increased in these age groups, respectively ( $P < 0.05$ ). There were also significant increases in fasting blood glucose and triglyceride levels among subjects with and without metabolic syndrome in age groups 30-34 and 35-40 years old ( $P < 0.05$ ). There were decreased HDL-cholesterol in age group 30-34 years old ( $P < 0.05$ ). There were significant differences in the prevalence of metabolic syndrome in age groups 20-24 and 30-34 years old ( $P < 0.05$ ).

*Conclusion:* High waist circumference, Fasting Glucose and triglyceride, and low HDL cholesterol are more effective among subjects with metabolic syndrome and, in the presence of obesity may elevate the risk of coronary heart disease. The increasing number of overweight and obese subjects in our study makes certain that metabolic syndrome begins from young age and will continue as age progresses. It seems that women require altering their life style to prevent cardiovascular complications.

**Keywords:** Gorgan, metabolic syndrome, age related, ethnic group.

**Introduction:**

The prevalence of the metabolic syndrome increases whole around the world. The metabolic syndrome is described by the clustering of hypertension, dyslipidaemia, central obesity, insulin resistance and high fasting plasma glucose<sup>[1]</sup>. In 1923, metabolic syndrome was explained for the first time by Klyn<sup>[2]</sup>. Gerald Reaven re-exhibited the concept of metabolic syndrome in 1988. He has been shown that metabolic syndrome is the clustering of hypertension, glucose intolerance; high triglycerides and low high density lipoprotein (HDL) concentration<sup>[3]</sup>. Studies have been shown that the prevalence of metabolic syndrome changes among different ethnic groups<sup>[4-6]</sup>. The genetically differences, nutritional regiment, physical exercise, age and gender may affect the prevalence of metabolic syndrome and its components<sup>[7]</sup>. Many studies have indicated that the prevalence of metabolic syndrome changes worldwide 8-24% and 7-46.5% among men and women, respectively<sup>[8-11]</sup>. The main causes of death among women are cardiovascular disease<sup>[12]</sup>. Many epidemiological studies have indicated the importance of the metabolic syndrome<sup>[13-15]</sup>. It has been shown that the metabolic

syndrome causes health problem in developed and developing countries. The prevalence of metabolic syndrome in Europe and European Americans vary nearly 20%-30% in men and women<sup>[4, 7, 9, 16-17]</sup>. It has been also revealed that the prevalence of metabolic syndrome is elevating in Asian countries<sup>[18]</sup>. It was reported that the prevalence of metabolic syndrome is the similar for men (24%) and women (23.7%) in the US, but there was an ethnic differences in the prevalence of metabolic syndrome<sup>[9]</sup>. There are a few studies on the prevalence of the metabolic syndrome among adult

Marjani Abdoljalal et al. *J Pharm Biomed Sci.* 2013, May; 30 (30): 929-935.  
Available at [jpbms.info](http://jpbms.info)

people. It has been indicated that the prevalence of the metabolic syndrome among 15 (in Japan), 12-19 (in the United States), 10-18 (in Mexico), 10-19 (in Iran) [19-22] and 12-19 (US black) years old were 1%, 6.4%, 6.5%, 10% and 4%, respectively<sup>[23]</sup>. In present study, we determined age related prevalence of the metabolic syndrome among Fars ethnic women in this area (Gorgan, North East of Caspian Sea).

### **Materials and methods:**

This present study was carried out in the Metabolic Disorders Research Center in Gorgan. 160 Fars adult females were took part (Women who speak only Persian language) that was directed to the Health Centers in Gorgan. Demographic data is collected by a questionnaire. Women with hormone replacement therapy, taking anti-diabetes and anti-hypertensive anti-lipidemic agents and active smokers were excluded. A blood sample was collected after 8-12-hours fasting. Fasting blood glucose, triglycerides, total cholesterol, LDL-cholesterol and HDL-cholesterol levels were determined in serum of all subjects. Commercial kits and spectrophotometer techniques (Model JENWAY 6105 UV / VIS) were used to assess these biochemical parameters in the Metabolic Disorders Research Center. Metabolic syndrome among women was considered if any of subjects had 3 or more of the following ATP III Criteria<sup>[24]</sup>:

1. Serum glucose level higher than 110 mg/dl.
2. Low HDL-cholesterol lower than 50 mg/dl.
3. Serum triglycerides level higher than 150 mg/dl.
4. Systolic Blood Pressure (SBP) higher than 130 mmHg and/or Diastolic Blood Pressure (DBP) higher than 85 mmHg (Hypertension).
5. Waist Circumference higher than 88 cm (Abdominal obesity).
- 6.

Weight was measured with minimal clothed, using digital scales. Height was measured with tape meter when the shoulder was in a normal position. Calculation of body mass index (BMI) was done when weight in kilograms divided by height in meters squared. BMI with 25.0-29.9 Kg/m<sup>2</sup> were arranged as overweight. Subjects with a BMI greater than 30 Kg/m<sup>2</sup> and 45 Kg/m<sup>2</sup> were specified as obese and very obese, respectively<sup>[25]</sup>. Abdominal obesity was assessed at the point halfway between the lower border of ribs and the iliac crest in a horizontal plane<sup>[26]</sup>. Blood pressure was determined in sitting position from the right hand. The results were reported in percentages and mean value. SPSS- 16 version software was used to analyze the data. Independent student t test was used to evaluate the results. P- Value lower than 0.05 was considered statistical significant.

### **Results:**

160 Fars females were included in this study. The mean age of women was 53.65±9.50 years (the age range was 20-40 years old). The mean waist circumferences, fasting blood glucose, triglyceride,

HDL-cholesterol, LDL-cholesterol levels, systolic blood pressure, diastolic blood pressure and body mass index were shown in table 1 according to age distribution among Fars ethnic group. There were no significant differences between all parameters in different age groups. The mean waist circumferences increased from age 25 to 40 years old. The mean HDL-cholesterol decreased in all age groups. Table 2 shows distribution of metabolic syndrome components among subjects with and without metabolic syndrome according to age group. The most age distribution was in ages from 35 to 40 years (50%). There were significant differences between the mean value of waist circumferences, fasting blood glucose, body mass index and triglyceride among subjects with and without metabolic syndrome in age groups 20-24 and 25-29 years old (P<0.05). The mean HDL-cholesterol and LDL-cholesterol levels were significantly decreased and increased in these age groups, respectively (P<0.05). There were also significant increases in fasting blood glucose and triglyceride levels among subjects with and without metabolic syndrome in age groups 30-34 and 35-40 years old (P<0.05). There were decreased HDL-cholesterol in age group 30-34 years old (P<0.05). There were no significant differences between other parameters in subjects with and without metabolic syndrome in different age groups (P>0.05). There were significant differences in the prevalence of metabolic syndrome in age groups 20-24 and 30-34 years old (P<0.05).

Marjani Abdoljalal et al. *J Pharm Biomed Sci.* 2013, May; 30 (30): 929-935.  
Available at jpbms.info

Table 1. Distribution of metabolic syndrome components according to age group.

Age group (year)	20 - 24	25 - 29	30 - 34	35 - 40
<b>Parameters</b>				
<b>WC (mg/dl)</b>	86.82±13.47	90.46±13.47	97.95±11.27	96.28±12.99
<b>FG (mg/dl)</b>	101.77±85.15	92.24±43.36	98.87±54.43	109.03±75.81
<b>BMI (kg/m<sup>2</sup>)</b>	24.22±4.43	25.50±4.95	28.36±5.86	27.98±5.23
<b>SBP (mmHg)</b>	107.61±11.38	112.33±11.94	112.50±18.31	115.91±14.97
<b>DBP (mmHg)</b>	67.60±9.82	70.80±8.27	68.25±13.30	70.67±10.90
<b>TG (mg/dl)</b>	97.67±42.52	122.0±113.68	131.75±75.41	111.05±58.89
<b>HDL- chol (mg/dl)</b>	43.45±11.88	43.98±14.54	40.61±10.27	45.39±14.0
<b>LDL- chol (mg/dl)</b>	110.42±39.92	117.59±49.89	119.34±34.85	117.45±36.82
<b>Total n (%)</b>	28(17.50)	30(18.75)	20(12.5)	82(51.25)

WC: Waist Circumferences, FG: Fasting Glucose, BMI: Body Mass Index, SBP: Systolic Blood Pressure, DBP: Diastolic Blood Pressure, TG: triglyceride, HDL-chol: HDL-cholesterol and LDL-chol: LDL-cholesterol. P>0.05

Table 2. Distribution of metabolic syndrome components in subjects with and without metabolic syndrome according to age group

Age group (year)	20 - 24		25 - 29		30 - 34		35 - 40	
	MS	WMS	MS	WMS	MS	WMS	MS	WMS
WC (mg/dl)	100.33±9.81*	85.20±8.32	107.67±5.46*	86.16±11.22	103.14±5.58	96.06±13.99	94.97±9.41	96.62±13.81
FG (mg/dl)	244.37±249.39*	84.65±11.31	129.12±90.25*	83.02±12.51	125.0±86.95*	88.90±19.04	176.38±116.70*	91.42±48.48
BMI (kg/m <sup>2</sup> )	31.29±9.40*	23.37±2.79	30.87±5.57*	24.15±3.84	30.03±4.50	28.27±6.81	26.66±4.52	28.32±5.38
SBP (mmHg)	123.33±5.77	105.72±10.41	123.33±8.16	109.58±11.22	118.57±21.93	110.62±15.69	110.59±12.97	117.31±15.23
DBP (mmHg)	73.33±15.27	66.92±9.20	69.16±4.91	71.20±8.95	67.85±16.29	69.37±11.23	70.0±10.0	70.84±11.20
TG (mg/dl)	173.67±31.26*	88.56±33.83	324.67±93.65*	71.33±31.57	212.86±72.60*	86.0±23.96	186.65±60.30*	91.27±39.56
HDL- chol (mg/dl)	28.73±51.14*	45.21±11.19	49.50±26.82	42.60±10.03	33.71±7.36*	45.88±9.51	37.85±14.27*	47.36±13.35
LDL- chol (mg/dl)	100.93±51.19	111.56±39.52	136.65±61.75*	112.84±46.82	111.71±25.86	128.55±38.93	119.02±20.92	117.04±40.07

WC: Waist Circumferences, FG: Fasting Glucose, BMI: Body Mass Index, SBP: Systolic Blood Pressure, DBP: Diastolic Blood Pressure, TG: triglyceride, HDL-chol: HDL-cholesterol and LDL-chol: LDL-cholesterol. MS: Metabolic Syndrome, WMS: Without Metabolic Syndrome.\*P<0.05.

### Discussion:

The results of present study indicated that metabolic syndrome components increased as age progresses. Waist circumference and HDL-cholesterol (Lower than 50 mg/dl) were increased and decreased among different age groups, respectively (Table 1). Many studies have revealed that metabolic syndrome is more common in women than men<sup>[27]</sup>. Kazan et al. indicated that the prevalence

elevated as age increased in women than men<sup>[28]</sup>. Sex seems to have an important role in elevation of prevalence of risk factors like obesity, high lipid levels and low HDL cholesterol level among females<sup>[29]</sup>. Our study has been shown that Waist

circumference and BMI increased from age 25 to 40 years old (Table 1). Some other studies have shown that women in different age groups indicated higher waist circumference and blood glucose and low HDL cholesterol when compared to males<sup>[30]</sup>. This means that sex and age have an important effect on the prevalence of cardiovascular risk factors in subjects with metabolic syndrome. Study in Turkey showed that the prevalence of metabolic syndrome increased from ages 30 to 80 years old<sup>[31]</sup>. Study of Ford revealed that the prevalence was elevated for women among US people aged from 20 to 60 years old. They have shown that there was an association between a higher prevalence of metabolic syndrome and older age groups<sup>[32]</sup>. Study of Park et al. indicated that the prevalence of the metabolic syndrome increases among women from ages 20 to 70 years old<sup>[33]</sup>. Although age-related differences in the mean value of metabolic syndrome components were not significant overall in different age groups, but they were significant for the components of high fasting glucose, high triglyceride (significant in all age groups) and high waist circumference (significant in age groups 20-24 and 25-29 years old) among women with metabolic syndrome. Our study has been shown that Hyperglycemia, hypertriglyceridemia, high waist circumference and low HDL cholesterol were particularly common amongst the woman subjects. The present study indicated that the most metabolic syndrome components found in age group 20-24 years old (4 components of metabolic syndrome). This means that young people is particularly of interest, due to the fact that the risk of cardiovascular disease elevates as age progresses. Study has revealed that the causes of metabolic syndrome are associated with obesity, lack of physical activity, high LDL-cholesterol diets, aging and genetic factors<sup>[34]</sup>. Studies have shown that low HDL cholesterol was seen among females with metabolic syndrome<sup>[35-36]</sup>.

Many studies have revealed that obesity is accompanied with waist circumference greater than 88 cm which is a significant prognosis for CVD in different age groups<sup>[37-39]</sup>. It has been shown that Obesity, glucose metabolism abnormality, hypertension and dyslipidemia are known as a risk factor for CVD leading to fatal outcome. These risk factors are the components of metabolic syndrome, therefore it is important to determine the components of metabolic syndrome to prevent the early onset of CVD <sup>[40-42]</sup>. Study among US women younger than 50 years have been shown that the annual death rate from CVD is more than breast cancer<sup>[43]</sup>. In other studies in the United States have indicated that the epidemiological increase of overweight and obesity is associated with an elevated prevalence of early onset CVD and type 2 diabetes mellitus<sup>[44-45]</sup>. Studies have shown that these epidemic changes may in relation with lifestyles alterations<sup>[46,47]</sup>. Abdominal obesity was the common metabolic syndrome component in our study groups which is in agreement with other study groups in the United States<sup>[48]</sup>. It has been shown that there is an association between the metabolic syndrome and elevation of age and BMI<sup>[33,49-50]</sup>. Other studies have indicated that BMI and age are the most important predictors of metabolic syndrome for women and men<sup>[48]</sup>. Studies

among different populations indicated that waist circumference alone<sup>[51]</sup> or with body mass index<sup>[52]</sup> is a good predictors for type 2 diabetes mellitus. Report of a WHO Expert Committee revealed that waist circumference is the most effective and easiest anthropometric index to be utilized for measurement of fatness and fat location <sup>[53]</sup>. Study of Ford et al.<sup>[38]</sup> indicated that waist circumference is a better predictor than body mass index of the metabolic syndrome, diabetes, cardiovascular disease. There is a relationship between abdominal obesity and the elevation of portal free fatty acid levels which causes hyperinsulinemia<sup>[54-55]</sup>. The hyperinsulinemia is associated with cardiovascular disease risk factors<sup>[56]</sup>. In our study the mean BMI varied from 24.22±4.43 kg/m<sup>2</sup> to 28.36±5.86 kg/m<sup>2</sup>, which is the same to women in northeastern Iran, Golestan Province (28.6 kg/m<sup>2</sup>)<sup>[57]</sup>. Women in our study group were overweight and obese. The most females with metabolic syndrome were obese. Obesity is associated with the elevation of the risk of dyslipidemia, type 2 diabetes mellitus and hypertension. Obesity is an important predictor of coronary heart and cardiovascular disease<sup>[58]</sup>. Low HDL-cholesterol level is associated with increased serum triglycerides and remnant lipoproteins<sup>[59-60]</sup>. There are association between low HDL cholesterol and high LDL-cholesterol level<sup>[61]</sup>. Low HDL-cholesterol level is associated with insulin resistance and metabolic risk factors<sup>[62]</sup>. Our results indicates that the mean value of HDL cholesterol in our study group were lower than 50

Marjani Abdoljalal et al. *J Pharm Biomed Sci.* 2013, May; 30 (30): 929-935.  
Available at [jpbms.info](http://jpbms.info)

mg/dl. There are contrary relationship between high HDL cholesterol and the decreased risk of coronary heart disease [63].

### **Conclusion:**

High waist circumference, Fasting Glucose and triglyceride, and low HDL cholesterol are more effective among subjects with metabolic syndrome and, in the presence of obesity may elevate the risk of coronary heart disease. The increasing number of overweight and obese subjects in our study makes certain that the metabolic syndrome begins from young age and will continue as age progresses. It seems that women require altering their life style to prevent cardiovascular complications.

### **References:**

1. Miranda PJ, DeFronzo RA, Califf RM and Guyton JR. Metabolic syndrome: definition, pathophysiology, and mechanisms. *Am Heart J.* 2005; 149: 33-45.
2. Kylin E. Studien ueber das Hypertonie-Hyperglykamie - Hyperurikamiesyndrom. *Zentralblatt fuer Innere Medizin* 1923; 44: 105-27.
3. Reaven G. Mand Banting Lecture. Role of insulin resistance in human disease. *Diabetes* 1988; 37: 1595-607.
4. Meigs JB, Wilson PW, Nathan DM, et al. Prevalence and characteristics of the metabolic syndrome in the San Antonio heart and Framingham offspring studies, *Diabetes* 2003 52: 2160-7.
5. Burke JP, Williams K, Gaskill SP, et al. Rapid rise in the incidence of type 2 diabetes from 1987 to 1996: Results from the San Antonio heart study, *Arch Intern Med.* 1999; 41: 1450-6.
6. King H, Zimmet P. Trends in the prevalence and incidence of diabetes: Noninsulin dependent diabetes mellitus, *World Health Stat* 1998; 41: 190-6.
7. Cameron AJ, Shaw JE and Zimmet PZ. The metabolic syndrome: prevalence in worldwide populations, *Endocrinol Metab Clin N Am.* 2004; 33: 351-75.
8. Gupta A, Gupta R, Sarna M, Rastogi S, Gupta VP and Kothari K. Prevalence of diabetes, impaired fasting glucose and insulin resistance syndrome in an urban Indian population, *Diabetes Res Clin Pract.* 2003; 61: 69-76.
9. Ford ES, Giles WH and Dietz WH. Prevalence of the metabolic syndrome among US adults: findings from the Third National Health and Nutrition Examination Survey, *JAMA.* 2002; 287: 356-9.
10. Balkau B, Vernay M, Mhamdi L, Novak M, Arondel D and Vol S, et al. The D.E.S.I.R Study Group. The incidence and persistence of the NCEP (National Cholesterol Education Program) metabolic syndrome, The French D.E.S.I.R. study. *Diabetes Metab.* 2003; 29: 526-32.
11. Ramachandran A, Snehalatha C, Satyavani K, Sivasankariand S, Vijay V. Metabolic syndrome in urban Asian Indian Adults-a population study using modified ATP III criteria., *Diabetes Res Clin Pract.* 2003; 60: 199-204.
12. Lloyd-Jones D , Adams R, Carnethon M, De SG, Ferguson TB, Flegal K, et al. Heart disease and stroke statistics-2009 update: a report from the American Heart Association Statistics Committee and Stroke Statistics Subcommittee, *Circulation* 2009; 119: 480-6.
13. Shepherd J, Cobbe SM, Ford I, Isles CG, Lorimerand Mac AR, Farlane PW, et al. Prevention of coronary heart disease with pravastatin in men with hypercholesterolemia, West of Scotland Coronary Prevention Study Group, *N Engl J Med.* 1995; 333: 1301-7.
14. Downs JR, Clearfield M, Weis S, Whitney E, Shapiroand DR, Beerc PA, et al. Primary prevention of acute coronary events with lovastatin in men and women with average cholesterol levels: results of AFCAPS/ TexCAPS. Air Force/Texas Coronary Atherosclerosis Prevention Study, *JAMA.* 1998; 279:1615-22.
15. Ballantyne CM, Olsson AG, Cook TJ, Mercuri MF, Pedersen TR and Kjekshus J. Influence of low high-density lipoprotein cholesterol and elevated triglyceride on coronary heart disease events and response to simvastatin therapy in 4S, *Circulation* 2001; 104: 3046-51.
16. Qiao Q. Comparison of different definitions of the metabolic syndrome in relation to cardiovascular mortality in European men and women, *Diabetologia* 2006; 49: 2837-46.
17. Hildrum B, Mykletun A, Hole T , Midthjell K, DahlAA. Age-specific prevalence of the metabolic syndrome defined by the International Diabetes Federation and the National Cholesterol Education Program: the Norwegian HUNT 2 study, *BMC Public Health* 2007; 7: 220.
18. Meigs JB. "Invited commentary: insulin resistance syndrome? Syndrome X? Multiple metabolic syndrome? A syndrome at all? Factor analysis reveals patterns in the fabric of correlated metabolic risk factors," *American Journal of Epidemiology* 2000; 10: 908-12.
19. Duncan GE, Li SM, Zhou XH. Prevalence and trends of a metabolic syndrome phenotype among U.S. Adolescents, 1999-2000. *Diabetes Care* 2004; 27: 2438-2443.
20. Esmailzadeh A, Mirmiran P, Azadbakht L, Etemadi A, Azizi F. High prevalence of the metabolic syndrome in Iranian adolescents. *Obesity (Silver Spring)* 2006; 14:377-82.
21. Rodriguez-Moran M, Salazar-Vazquez B, Violante R, Guerrero-Romero F: Metabolic syndrome among children and adolescents aged 10-18 years. *Diabetes Care* 2004; 27:2516-2517.
22. Saito I, Mori M, Shibata H, Hirose H, Tsujioka M, Kawabe H. Prevalence of metabolic syndrome in young men in Japan. *J Atheroscler Thromb.* 2007; 14:27-30.
23. Johnson WD, Kroon JJ, Greenway FL, Bouchard C, Ryan D, Katzmarzyk PT. Prevalence of risk factors for metabolic syndrome in adolescents: National Health and Nutrition Examination Survey (NHANES), 2001-2006. *Arch Pediatr Adolesc Med.* 2009; 163:371-7.
24. Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults. Executive Summary of the 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). *JAMA* 2001; 285: 2486-97.
25. World Health Organization. Prevention and Management of the Global Epidemic of Obesity. Report of the WHO Consultation on Obesity. WHO: Geneva, 1998 (Technical Report Series, No. 894).
26. Dalton M, Cameron AJ, Zimmet PZ, Shaw JE, Jolley D, Dunstan DW, Welborn TA. AusDiab Steering Committee. Waist circumference, waist-hip ratio and body mass index and their correlation with cardiovascular disease risk factors in Australian adults. *J Intern Med.* 2003; 254: 555-63.
27. Sarrafzadegan N, Kelishadi R, Baghaei A, Hussein SG, Malekafzali H, Mohammadifard N, et al. Metabolic syndrome: an emerging public health

Marjani Abdoljalal et al. *J Pharm Biomed Sci.* 2013, May; 30 (30): 929-935.  
Available at [jpbms.info](http://jpbms.info)

- problem in Iranian women: Isfahan Healthy Heart Program. *Int J Cardiol.* 2008; 131(1): 90-6.
28. Kozan O, Oguz A, Abaci A, Erol C, Ongen Z, Temizhan A, et al. Prevalence of the metabolic syndrome among Turkish adults. *Eur J Clin Nutr.* 2007; 61(4): 548-53.
29. Hadaegh F, Zabetian A, Tohidi M, Ghasemi A, Sheikholeslami F, Azizi F. Prevalence of metabolic syndrome by the Adult Treatment Panel III, International Diabetes Federation, and World Health Organization definitions and their association with coronary heart disease in an elderly Iranian population. *Ann Acad Med Singapore* 2009; 38(2): 142-9.
30. Mukuddem-Petersen J, Snijder MB, van Dam RM, Dekker JM, Bouter LM, Stehouwer CD, et al. Sagittal abdominal diameter: no advantage compared with other anthropometric measures as a correlate of components of the metabolic syndrome in elderly from the Hoorn Study. *Am J Clin Nutr* 2006; 84(5): 995-1002.
31. Sanisoglu SY, Oktenli C, Hasimi A, et al. Prevalence of metabolic syndrome-related disorders in a large adult population in Turkey. *BMC Public Health* 2006; 6:92.
32. Ford ES, Li C, Imperatore G, Cook S. Age, sex, and ethnic variations in serum insulin concentrations among US youth: findings from the National Health and Nutrition Examination Survey 1999–2002. *Diabetes Care* 2006; 29:2605–11.
33. Park YW, Zhu S, Palaniappan L, et al. The metabolic syndrome: prevalence and associated risk factor findings in the US population from the Third National Health and Nutrition Examination Survey, 1988–1994. *Arch Intern Med.* 2003; 163:427–36.
34. Grundy SM. Small LDL, atherogenic dyslipidemia, and the metabolic syndrome. *Circulation* 1997; 95:1– 4.
35. Reddy KS, Shah P, Shrivastava U, Prabhakaran D, Joshi M, Puri SK, et al. Coronary heart disease risk factors in an industrial population of north India. *Can J Cardiol.* 1997; 13(Suppl. B):26B.
36. Krishnaswami S. Conventional risk factors for coronary artery disease in Indian patients. In: Sethi KK, editor. *Coronary Artery Disease in Indians: a Global Perspective.* Bombay: Cardiological Society of India, 1998; 73– 82.
37. Orio F Jr, Palomba S, Cascella T, Savastano S, Lombardi G, Colao A. Cardiovascular complications of obesity in adolescents. *J Endocrinol Invest.* 2007;30: 70–80.
38. Ford ES, Mokdad AH, Giles WH. Trends in waist circumference among US adults. *Obes Res.* 2003; 11: 1223–31.
39. Hillege HL, Fidler V, Diercks GF, et al. Urinary albumin excretion predicts cardiovascular and noncardiovascular mortality in general population. *Circulation* 2002; 106:1777–82.
40. Isomaa B, Almgren P, Tuomi T, et al. Cardiovascular morbidity and mortality associated with the metabolic syndrome. *Diabetes Care* 2001; 24: 683–689.
41. Meigs JB, D'Agostino RB Sr, Wilson PW, Cupples LA, Nathan DM, Singer DE. Risk variable clustering in the insulin resistance syndrome. The Framingham Offspring Study. *Diabetes* 1997; 46:1594–1600.
42. Natali A, Toschi E, Baldeweg S, et al. Clustering of insulin resistance with vascular dysfunction and lowgrade inflammation in type 2 diabetes. *Diabetes* 2006; 55:1133–40.
43. Rosamond W, Flegal K, Friday G, et al. Heart disease and stroke statistics—2007 update: a report from the American Heart Association Statistics Committee and Stroke Statistics Subcommittee. *Circulation* 2007; 115(4):e69–e171.
44. Gill H, Mugo M, Whaley-Connell A, Stump C, Sowers JR. The key role of insulin resistance in the cardiometabolic syndrome. *Am J Med Sci.* 2005; 330: 290–294.
45. Grundy SM. Obesity, metabolic syndrome, and cardiovascular disease. *J Clin Endocrinol Metab.* 2004; 89: 2595–2600.
46. Foy CG, Foley KL, D'Agostino RB Jr, Goff DC Jr, Mayer-Davis E, Wagenknecht LE. Physical activity, insulin sensitivity, and hypertension among US adults: findings from the Insulin Resistance Atherosclerosis Study. *Am J Epidemiol.* 2006; 163:921–928.
47. Straznicki NE, Lambert EA, Lambert GW, Masuo K, Esler MD, Nestel PJ. Effects of dietary weight loss on sympathetic activity and cardiac risk factors associated with the metabolic syndrome. *J Clin Endocrinol Metab.* 2005; 90:5998–6005.
48. Ervin RB. Prevalence of metabolic syndrome among adults 20 years of age and over, by sex, age, race and ethnicity, and body mass index: United States, 2003–2006. *Natl Health Stat Report* 2009; 5:1–7.
49. Kwasniewska M, Kaleta D, Dziankowska-Zaborszczyk E, Drygas W. Healthy behaviors, lifestyle patterns and sociodemographic determinants of the metabolic syndrome. *Cent Eur J Public Health* 2009; 17:14–19.
50. Yang FY, Wahlqvist ML, Lee MS. Body mass index (BMI) as a major factor in the incidence of the metabolic syndrome and its constituents in unaffected Taiwanese from 1998 to 2002. *Asia Pac J Clin Nutr.* 2008; 17:339–51.
51. Reeder BA, Senthilselvan A, Despres JP, et al. The association of cardiovascular disease risk factors with abdominal obesity in Canada. Canadian Heart Health Surveys Research Group. *CMAJ.* 1997; 157 (suppl 1):S39-S45.
52. Wang Y, Rimm EB, Stampfer MJ, Willett WC, Hu FB. Comparison of abdominal adiposity and overall obesity in predicting risk of type 2 diabetes among men. *Am J Clin Nutr.* 2005; 81: 555-63.
53. Physical status: the use and interpretation of anthropometry. Report of a WHO Expert Committee. Geneva, World Health Organization, 1995 (WHO Technical Report Series, No. 854).
54. Foucan L, Hanley J, Deloumeaux J, Suissa S. Body mass index (BMI) and waist circumference (WC) as screening tools for cardiovascular risk factors in Guadeloupean women. *J Clin Epidemiol.* 2002; 55: 990-6.
55. Zabetian A, Hadaegh F, Azizi F. Prevalence of metabolic syndrome in Iranian adult population, concordance between the IDF with the ATP III and the WHO definitions. *Diabetes Rese Clin Pract.* 2007; 77: 251-7.
56. Schmidt MI, Watson RL, Duncan BB, et al. Clustering of dyslipidemia, hyperuricemia, diabetes, and hypertension and its association with fasting insulin and central and overall obesity in a general population. Atherosclerosis Risk in Communities Study Investigators. *Metabolism* 1996; 45: 699-706.
57. Bahrami H, Sadatsafavi M, Pourshams A, et al. Obesity and hypertension in an Iranian cohort study; Iranian women experience higher rates of obesity and hypertension than American women. *BMC Public Health* 2006; 20: 158.
58. National Institutes of Health, National Heart, Lung, and Blood Institute. Clinical guidelines on the identification, evaluation, and treatment of overweight and obesity in adults: the evidence report. *Obes Res.* 1998; 6 (suppl 2): S51-S210.
59. Phillips NR, Havel RJ, Kane JP. Levels and interrelationships of serum and lipoprotein cholesterol and triglycerides: association with adiposity and the consumption of ethanol, tobacco, and beverages containing caffeine. *Arteriosclerosis* 1981; 1: 13-24.
60. Schaefer EJ, Lamon-Fava S, Ordovas JM, et al. Factors associated with low and elevated plasma high density lipoprotein cholesterol and polipoprotein A-I levels in

Marjani Abdoljalal et al. *J Pharm Biomed Sci.* 2013, May; 30 (30): 929-935.  
Available at [jpbms.info](http://jpbms.info)

the Framingham Offspring Study. *J Lipid Res.* 1994; 35: 871-82.  
61. Austin MA, Hokanson JE, Edwards KL. Hypertriglyceridemia as a cardiovascular risk factor. *Am J Cardiol.* 1998; 81: 7B-12B.  
62. Wilson PWF, D'Agostino RB, Levy D, Belanger AM, Silbershatz H, Kannel WB. Prediction of coronary heart disease using risk factor categories. *Circulation* 1998; 97: 1837- 47.  
63. Vega GL, Grundy SM. Hypoalphalipoproteinemia (low high density lipoprotein) as a risk factor for coronary heart disease. *Curr Opin Lipidol.* 1996; 7: 209-16.

Conflict of interest: - **Author has not declared any conflict of interest.**

Source of funding: - **None.**

---

**\*Correspondence address:**

**Abdoljalal Marjani,**

(Associated Prof. in Biochemistry), Gorgan Faculty of Medicine, Department of Biochemistry and Biophysics, Metabolic Disorders Research Center, Gorgan Faculty of Medicine, Golestan University of Medical Sciences, Gorgan, Golestan province, .Iran. Tel & Fax: +98(171)4421651 & 4440225.

Submit your next  
Manuscript to:-

**Journal of  
Pharmaceutical and  
Biomedical Sciences  
(JPBMS)**

**An international  
member journal of  
COPE, World Health  
organization (WHO)-  
HINARI Access and  
JDP and take benefit of:**

- Convenient online submission with persistent Authors support
- Thorough peer review.
- High visibility and citation of article with readers /authors across the boundaries.
- Immediate publication on acceptance.
- Inclusion in COPE, HINARI Access (WHO) JDP, CAS, DOAJ, NLM catalog, Google Scholar and many more.....

Submit your manuscript  
at: [www.jpbms.info](http://www.jpbms.info).

Copyright © 2013 Marjani Abdoljalal et al. This is an open access article distributed under the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.