Evaluation of heart rate reserve and high-sensitivity C-reactive protein in individuals with and without metabolic syndrome in Isfahan, Iran

Yosef Khaledi⁽¹⁾, Esmaeil Aghababaei⁽¹⁾, <u>Masoumeh Sadeghi</u>⁽²⁾, Mohammad Hashemi⁽³⁾, Hamid Sanei⁽³⁾

Abstract

BACKGROUND: Lack of heart rate increase proportionate to exercise causes poor prognosis. Moreover, inflammatory factors such as C-reactive protein (CRP) are associated with atherosclerosis. The current study compared these two indices in individuals with and without metabolic syndrome in Isfahan, Iran.

METHODS: This study was performed on 203 people without and 123 patients with metabolic syndrome who were randomly selected from the participants of the Isfahan Cohort Study. The demographic data, waist circumference, blood pressure, height, and weight of the participants were recorded. Moreover, serum tr`viglyceride (TG), fasting blood sugar (FBS), total cholesterol, high density lipoprotein (HDL), low density lipoprotein (LDL), and high-sensitivity CRP (hs-CRP) levels were measured. Exercise test was carried out according to the Bruce standard protocol and heart rate reserve (HRR) was determined and recorded. The age-adjusted data was analyzed using generalized linear regression and student's t-test in SPSS₁₅.

RESULTS: The mean ages of participants without and with metabolic syndrome were 54.16 ± 8.61 and 54.29 ± 7.6 years, respectively. The corresponding values for mean LDL levels were 116.17 ± 24.04 and 120.12 ± 29.55 mg/dl. TG levels were 140.38 ± 61.65 and 259.99 ± 184.49 mg/dl for subjects without and with the metabolic syndrome, respectively. The mean FBS levels were 81.81 ± 9.90 mg/dl in the participants without the syndrome and 107.13 ± 48.46 mg/dl in those with metabolic syndrome. The mean systolic blood pressure was 116.06 ± 13.69 mmHg in persons without metabolic syndrome and 130.73 ± 15.15 mmHg in patients with the syndrome. The values for mean diastolic levels in the two groups were 76.52 ± 6.69 and 82.84 ± 8.7 mmHg, respectively. While the two groups were not significantly different in terms of HRR (P = 0.27), hs-CRP levels in the metabolic syndrome group was significantly higher than the other group (P = 0.02).

CONCLUSION: We failed to establish a relationship between HRR and the metabolic syndrome. However, the observed relationship between metabolic syndrome and hs-CRP level, which is an inflammatory factor, indicates elevated levels of hs-CRP in patients with metabolic syndrome.

Keywords: Metabolic Syndrome, Exercise Test, Heart Rate Reserve, High-Sensitivity C-Reactive Protein.

ARYA Atherosclerosis Journal 2012, 8(2): 70-75

Date of submission: 30 Apr 2011, Date of acceptance: 7 Jun 2012

Introduction

In 1988, Reaven introduced metabolic syndrome as a set of risk factors, basically including insulin resistance, hypertension (HTN), dyslipidemia, and other metabolic disorders, which increase the risk of cardiovascular diseases.¹ Therefore, scientists all around the world focused on identification of the syndrome to prevent, treat, and lower the risk of cardiovascular diseases.^{2,3} In patients with metabolic syndrome, exercise test is helpful in early diagnosis of cardiovascular diseases and also prediction of the risk

of mortality and occurrence of cardiovascular events. 4-6 Exercise test is an affordable, low-risk method which provides valuable information for physicians. Heart rate reserve (HRR) is one of the important findings of the exercise test. HRR is defined as increased heart rate as a result of increased activity. HRR values lower than or equal to 85% are considered to be normal and desirable and thus associated with poor prognosis. 7-16

High-sensitivity C-reactive protein (hs-CRP) is an inflammatory factor, known to be related to the

¹⁻ Resident, Isfahan Cardiovascular Research Center, Isfahan Cardiovascular Research Institute, Isfahan University of Medical Sciences, Isfahan, Iran.

²⁻ Associate Professor, Cardiac Rehabilitation Research Center, Isfahan Cardiovascular Research Institute, Isfahan University of Medical Sciences, Isfahan, Iran.

³⁻ Associate Professor, Department of Cardiology, School of Medicine, Isfahan University of Medical Sciences, Isfahan, Iran. Correspondence To: Masoumeh Sadeghi, Email: sadeghimasoumeh@gmail.com

metabolic syndrome.¹⁷⁻²³ However, all patients with metabolic syndrome are not at the same risk for development of cardiovascular diseases, and several factors including race, lifestyle, and health culture are effective in this respect.^{24,25} Various studies have demonstrated different results regarding these factors. Furthermore, the effectiveness of interventional approaches in reducing the disease risk is not identical.²⁶⁻²⁹ As mentioned, the metabolic syndrome has been suggested to play an important role in development of cardiovascular diseases which are in turn a major cause of death among adults in Iran. Moreover, the prevalence of metabolic syndrome is very high in Iranian population which differs ethically and climatically with other populations studied in other countries.³⁰⁻³³ Therefore, in the current study, we compared the HRR and hs-CRP of patients with and without metabolic syndrome.

Materials and Methods

This cross-sectional study was carried out on 203 individuals without metabolic syndrome and 123 individuals with metabolic syndrome who were registered in the Isfahan Cohort Study. As a prospective cohort study, the Isfahan Cohort Study started in 2002 and will continue until the end of 2012. It used multistage cluster sampling to select people older than 34 years of age from Isfahan, Najafabad, and Arak (3 cities in central Iran). All the demographic and behavioral data, as well as the indices such as blood pressure, body mass index (BMI), and the results of routine blood tests were recorded for all participants in the first year of the study. Afterwards, the occurrence of cardiac infarction, stroke, sudden death, and hospitalization were asked via telephone calls with two-year intervals. The clinical examinations and routine blood tests were reperformed in the fifth and sixth years of the follow-up. Further details were published by Sarraf-Zadegan et al. in 2003.30

The current study used accessible sampling to select 260 individuals with metabolic syndrome and 260 individuals without metabolic syndrome from the participants of the Isfahan Cohort Study. Using the provided telephone numbers in the records of participants, the individuals were invited to take part in the study. Finally, 123 individuals with metabolic syndrome and 203 individuals without the syndrome participated in all steps of the study. They were invited to the Isfahan Cardiovascular Research Center (Isfahan, Iran) at an appropriate time based on their schedule. They then attended interviews and the objectives and methodology of the study were

explained for them. After signing a written consent, they were included in the study. The metabolic syndrome was approved according to the protocol of the National Cholesterol Education Program/Adult Treatment Panel (NCEP/ATPIII). Therefore, individuals having three or more of the following criteria were considered to have the metabolic syndrome:

- 1- waist circumference above 102 cm in men and above 88 cm in women;
 - 2- blood triglyceride (TG) level ≥ 150 mg/dl;
 - 3- fasting blood sugar (FBS) \geq 110 mg/dl;
- 4- high density lipoprotein (HDL) level ≤ 40 mg/dl in men and ≤ 50 mg/dl in women; and
- 5- systolic blood pressure (SBP) \geq 135 mmHg or diastolic blood pressure (DBP) \geq 85 mmHg.³⁴

The members of the control group were individuals without metabolic syndrome who were selected from the same population in the Isfahan Cohort Study. They were included after matching for gender and age. The exclusion criteria were not being able to perform the exercise test, the presence of an absolute contraindication for performing exercise test, for instance myocardial infarction in the recent two days, advanced heart block, acute pulmonary emboli, uncontrolled HTN, acute myocarditis, severe aortic valve stenosis, or uncompensated cardiac failure, pregnancy, and not being willing to participate.4 The participants referred to the Isfahan Cardiovascular Research Center for the examinations after 12 hours of fasting. General physical examination and blood sampling were followed by waist circumference, blood pressure, weight, and height measurements according to international standards.35 TG and total cholesterol (TC) levels were determined using the enzymatic method by an autoanalyzer (Hitachi 902). After evaluating HDL levels using heparinmagnesium precipitation,³⁶ LDL levels were calculated according to Friedewald formula.³⁷ If the TG level was above 400 mg/dl, the LDL level was directly measured using a specific kit. The fasting blood sugar (FBS) level was determined using an enzymatic method (glucose oxidase). The 12-lead electrocardiogram (ECG) was taken by a trained technician according to the World Health Organization's multinational monitoring of trends and determinants in cardiovascular disease (WHO MONICA).38 Exercise test was then carried out according to Bruce or modified Bruce protocol and the HRR value was determined and recorded in each participant's file. The level of hs-CRP was quantitatively determined using specific kits. Since the values were expressed as \geq zero, we considered levels of 0-6 mg/l as negative and above 40 mg/l as +3.

The data obtained was adjusted for age and then analyzed by generalized linear regression and student's ttest in SPSS₁₅ (SPSS Inc., Chicago, IL, USA). P values below 0.05 were considered to be statistically significant.

In addition, hypercholesterolemia, hypertriglyceridemia, diabetes, and HTN were defined as TC > 200 mg/dl while using cholesterol lowering agents, TG > 200 mg/dl while using TG lowering agents, FBS > 126 mg/dl despite taking anti-diabetic drugs, and SBP > 140 mmHg or DBP > 90 mmHg while taking at least one anti-hypertensive drug, respectively.²⁴

Results

Among the 326 individuals who participated in this study, 56 men and 67 women had metabolic syndrome while 122 men and 81 women did not suffer from the syndrome. The youngest and oldest participants aged 35 and 82 years, respectively. The mean ages of subjects with and without metabolic syndrome were

 54.29 ± 7.6 and 54.16 ± 8.61 years, respectively.

The values obtained in the examinations for weight, waist circumference, SBP, DBP, and the laboratory findings stratified by gender and the presence of metabolic syndrome were adjusted by age and provided in table 1. As the table shows, there was a statistically significant relationship between metabolic syndrome and the values of waist circumference, weight, TG, TC, HDL, FBS, SBP, and DBP (P < 0.001). The frequency rates of dyslipidemia, diabetes, and HTN in the group with metabolic syndrome were higher than those in the group without metabolic syndrome (Table 2).

According to the indices of the exercise test, the HRR values in the group without metabolic syndrome were 75.09% and 73.17% in men and women, respectively. On the other hand, the values in the group with metabolic syndrome were 73.52% and 73.13%, respectively. The two groups were not significantly different in this respect (P = 0.27) (Table 3).

Table 1. Comparison of biochemical and clinical indices in individuals with and without metabolic syndrome

_	Without metabolic syndrome	With metabolic syndrome	P
Age (years)	54.16 ± 8.62	54.29±7.60	0.890
Waist circumference (cm)	87.97 ± 9.43	98.42 ± 8.74	< 0.001
Weight (Kg)	70.10 ± 11.08	80.35 ± 12.85	< 0.001
Triglyceride (mg/dl)	140.38 ± 61.66	259.99 ± 184.50	< 0.001
Total cholesterol (mg/dl)	201.17 ± 35.79	217.34 ± 49.60	0.003
High density lipoprotein (mg/dl)	47.91 ± 11.62	41.77 ± 8.67	< 0.001
Low density lipoprotein (mg/dl)	116.17 ± 24.05	120.12 ± 29.55	0.21
Fasting blood sugar (mg/dl)	81.81 ± 9.90	107.13 ± 48.46	< 0.001
Systolic blood pressure (mmHg)	116.06 ± 13.70	130.74 ± 15.16	< 0.001
Diastolic blood pressure (mmHg)	76.53 ± 6.70	82.84 ± 8.70	< 0.001

Values are expressed as mean \pm SD.

Table 2. Relative frequency of diabetes, dyslipidemia, and hypertension based on gender and the presence of metabolic syndrome

	Men			Women		
	With metabolic syndrome	Without metabolic syndrome	P	With metabolic syndrome	Without metabolic syndrome	P
Dyslipidemia	91.10%	63.90%	< 0.001	89.60%	63%	< 0.001
Diabetes	32.10%	1.60%	< 0.001	28.40	1.20%	< 0.001
Hypertension	50%	9%	< 0.001	32.80%	11.10%	< 0.001

Table 3. Heart rate reserve in the groups with and without metabolic syndrome

	Heart rate reserve	
Men without metabolic syndrome	$70.09\% \pm 7.571\%$	
Women without metabolic syndrome	$73.17\% \pm 7.601$	
P	0.079	
Men with metabolic syndrome	$73.52\% \pm 7.685\%$	
Women with metabolic syndrome	$73.13\% \pm 9.21\%$	
P	0.805	
P between the two groups	0.270	
Individuals with metabolic syndrome	$73.31\% \pm 8.522\%$	
Individuals without metabolic syndrome	$74.33\% \pm 7.622\%$	

Table 4	High-sensitivity	C-reactive protein	hs-CRP) levels in th	ne orouns with and	without metabolic syndrome
Table 4.	T HSH-SCHSILIVILY	C-ICACHVE DIOIEHI	L 1112-CAZI TEVEI2 III III	IE PIOUDS WITH AUG	WILLIOUS THEIRDONG SYLICIOTHE

Hs-CRP	Individuals with metabolic syndrome	Individuals without metabolic syndrome	P
Mean ± SD	18.74 ± 26.37	9.03 ± 14.94	
Median	7	4	0.019
Interquartile range	4-15	1.2-7	

The mean hs-CRP levels in the groups with and without metabolic syndrome were significantly different (26.37 \pm 18.74 vs. 14.94 \pm 9.03 mg/l; P = 0.019) (Table 4).

Discussion

In the current study, we evaluated two important factors, i.e. HRR and hs-CRP, for predicting the risk of cardiovascular diseases in individuals with and without metabolic syndrome who attended the Isfahan Cohort Study. HRR values were not significantly different between the two groups and the two sexes. However, the hs-CRP level was significantly higher in metabolic syndrome group. Similarly, Regitz-Zagrosek stated that the impact of gender differences on the prognosis of metabolic syndrome was not definitely confirmed.³⁹ Moreover, the impact of gender differences on HRR in normal people is not definitely defined.³⁹ On the contrary, Nilsson et al. stated that HRR is lower only in women with metabolic syndrome.⁴⁰

In a study, it was shown that the increase in the hs-CRP level in obese individuals was 4.6 folds more than non-obese people. It was detected that the mean log of hs-CRP elevated progressively as the components of metabolic syndrome increased. Hs-CRP level was also reported to be evidently higher in individuals having three or more components of the syndrome compared with those having one component of the syndrome. Another study however, suggested that only abdominal obesity was associated with significant increase in hs-CRP levels. 18

Investigations on the relationship between inflammation and metabolic syndrome revealed that individuals with the syndrome had a significantly higher level of hs-CRP in comparison with individuals without the syndrome. Several factors may contribute to increased levels of hs-CRP in the metabolic syndrome. One of these factors could be the effect of adipocytes, tumor necrosis factor (TNF), interleukin 6 (IL-6), and adiponectin on the hs-CRP level in obese people. Statistically significant relationships have also been reported between obesity and inflammatory markers. A previous study demonstrated higher levels of visceral obesity to be associated with elevated levels of CRP, TNF, and IL-6.21

Furthermore, increased levels of CRP have been found to be related with insulin resistance and the metabolic syndrome.²² Moreover, a study on American women reported higher CRP levels in women with one component of the syndrome compared to those without any components.²³

In the current study, the mean level of hs-CRP in the group with metabolic syndrome was significantly higher than that in the group without the syndrome (P = 0.019). This finding is in agreement with the results reported in other studies.

Therefore, the results of this study suggested that compared to HRR, hs-CRP level had a more powerful relationship with metabolic syndrome in our studied population. Such a finding would be helpful for physicians and health planners in making more accurate decisions. Moreover, this study could serve as a foundation for further studies in our community on people with metabolic syndrome, the relationship between the syndrome and cardiovascular diseases, and also the predictive values of the above-mentioned indices.

Acknowledgements

The authors wish to thank the staff of the Isfahan Cardiovascular Research Center, particularly the staff of the exercise test, surveillance, and analysis units, who kindly collaborated in performing this study.

Conflict of Interests

Authors have no conflict of interests.

References

- **1.** Reaven GM. Banting lecture 1988. Role of insulin resistance in human disease. Diabetes 1988; 37(12): 1595-607.
- **2.** Ford ES, Giles WH, Dietz WH. Prevalence of the metabolic syndrome among US adults: findings from the third National Health and Nutrition Examination Survey. JAMA 2002; 287(3): 356-9.
- **3.** Ozanne SE, Hales CN. Early programming of glucose-insulin metabolism. Trends Endocrinol Metab 2002; 13(9): 368-73.
- 4. Chaitman BR. Exercise stress testing. In: Libby P, Braunwald E, Editors. Braunwald's Heart Disease: A

- Textbook of Cardiovascular Medicine.PhiladelphIA: Saunders/Elsevier, 2008. p. 197-205.
- 5. Gibbons RJ, Balady GJ, Bricker JT, Chaitman BR, Fletcher GF, Froelicher VF, et al. ACC/AHA 2002 guideline update for exercise testing: summary article. A report of the American College of Cardiology/American Heart Association Task Force on Practice Guidelines (Committee to Update the 1997 Exercise Testing Guidelines). J Am Coll Cardiol 2002; 40(8): 1531-40.
- **6.** Vivekananthan DP, Blackstone EH, Pothier CE, Lauer MS. Heart rate recovery after exercise is a predictor of mortality, independent of the angiographic severity of coronary disease. J Am Coll Cardiol 2003; 42(5): 831-8.
- 7. Froelicher VF, Myers J. Exercise and the heart. 5th ed. Philadelphia: Saunders; 2006.
- 8. Elhendy A, Mahoney DW, Khandheria BK, Burger K, Pellikka PA. Prognostic significance of impairment of heart rate response to exercise: impact of left ventricular function and myocardial ischemia. J Am Coll Cardiol 2003; 42(5): 823-30.
- **9.** Jouven X, Empana JP, Schwartz PJ, Desnos M, Courbon D, Ducimetiere P. Heart-rate profile during exercise as a predictor of sudden death. N Engl J Med 2005; 352(19): 1951-8.
- **10.** Wei M, Kampert JB, Barlow CE, Nichaman MZ, Gibbons LW, Paffenbarger RS, et al. Relationship between low cardiorespiratory fitness and mortality in normal-weight, overweight, and obese men. JAMA 1999; 282(16): 1547-53.
- **11.** Mora S, Redberg RF, Cui Y, Whiteman MK, Flaws JA, Sharrett AR, et al. Ability of exercise testing to predict cardiovascular and all-cause death in asymptomatic women: a 20-year follow-up of the lipid research clinics prevalence study. JAMA 2003; 290(12): 1600-7.
- **12.** Chaitman BR. Abnormal heart rate responses to exercise predict increased long-term mortality regardless of coronary disease extent: the question is why? J Am Coll Cardiol 2003; 42(5): 839-41.
- **13.** Morise AP. Heart rate recovery: predictor of risk today and target of therapy tomorrow? Circulation 2004; 110(18): 2778-80.
- **14.** Fletcher GF, Balady GJ, Amsterdam EA, Chaitman B, Eckel R, Fleg J, et al. Exercise standards for testing and training: a statement for healthcare professionals from the American Heart Association. Circulation 2001; 104(14): 1694-740.
- **15.** Mark DB, Lauer MS. Exercise capacity: the prognostic variable that doesn't get enough respect. Circulation 2003; 108(13): 1534-6.
- **16.** Falcone C, Buzzi MP, Klersy C, Schwartz PJ. Rapid heart rate increase at onset of exercise predicts adverse cardiac events in patients with coronary artery disease. Circulation 2005; 112(13): 1959-64.

- **17.** Huffman FG, Gomez GP, Zarini GG. Metabolic syndrome and high-sensitivity C-reactive protein in Cubans. Ethn Dis 2009; 19(2): 115-20.
- **18.** Nakamura H, Ito H, Egami Y, Kaji Y, Maruyama T, Koike G, et al. Waist circumference is the main determinant of elevated C-reactive protein in metabolic syndrome. Diabetes Res Clin Pract 2008; 79(2): 330-6.
- 19. Bahceci M, Gokalp D, Bahceci S, Tuzcu A, Atmaca S, Arikan S. The correlation between adiposity and adiponectin, tumor necrosis factor alpha, interleukin-6 and high sensitivity C-reactive protein levels. Is adipocyte size associated with inflammation in adults? J Endocrinol Invest 2007; 30(3): 210-4.
- 20. Dandona P, Aljada A, Chaudhuri A, Mohanty P, Garg R. Metabolic syndrome: a comprehensive perspective based on interactions between obesity, diabetes, and inflammation. Circulation 2005; 111(11): 1448-54.
- **21.** Park HS, Park JY, Yu R. Relationship of obesity and visceral adiposity with serum concentrations of CRP, TNF-alpha and IL-6. Diabetes Res Clin Pract 2005; 69(1): 29-35.
- **22.** Lee WY, Park JS, Noh SY, Rhee EJ, Sung KC, Kim BS, et al. C-reactive protein concentrations are related to insulin resistance and metabolic syndrome as defined by the ATP III report. Int J Cardiol 2004; 97(1): 101-6.
- **23.** Ridker PM, Buring JE, Cook NR, Rifai N. C-reactive protein, the metabolic syndrome, and risk of incident cardiovascular events: an 8-year follow-up of 14 719 initially healthy American women. Circulation 2003; 107(3): 391-7.
- **24.** Grundy SM, Brewer HB, Cleeman JI, Smith SC, 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): 433-8.
- 25. Hunt KJ, Resendez RG, Williams K, Haffner SM, Stern MP. National Cholesterol Education Program versus World Health Organization metabolic syndrome in relation to all-cause and cardiovascular mortality in the San Antonio Heart Study. Circulation 2004; 110(10): 1251-7.
- **26.** McNeill AM, Rosamond WD, Girman CJ, Golden SH, Schmidt MI, East HE, et al. The metabolic syndrome and 11-year risk of incident cardiovascular disease in the atherosclerosis risk in communities study. Diabetes Care 2005; 28(2): 385-90.
- 27. Onat A, Ceyhan K, Basar O, Erer B, Toprak S, Sansoy V. Metabolic syndrome: major impact on coronary risk in a population with low cholesterol levelsa prospective and cross-sectional evaluation. Atherosclerosis 2002; 165(2): 285-92.
- **28.** Girman CJ, Rhodes T, Mercuri M, Pyorala K, Kjekshus J, Pedersen TR, et al. The metabolic syndrome and risk of major coronary events in the

- Scandinavian Simvastatin Survival Study (4S) and the Air Force/Texas Coronary Atherosclerosis Prevention Study (AFCAPS/TexCAPS). Am J Cardiol 2004; 93(2): 136-41.
- 29. Brinkworth GD, Noakes M, Buckley JD, Clifton PM. Weight loss improves heart rate recovery in overweight and obese men with features of the metabolic syndrome. Am Heart J 2006; 152(4): 693.e1-6.
- **30.** Sarraf-Zadegan N, Boshtam M, Malekafzali H, Bashardoost N, Sayed-Tabatabaei FA, Rafiei M, et al. Secular trends in cardiovascular mortality in Iran, with special reference to Isfahan. Acta Cardiol 1999; 54(6): 327-33.
- **31.** Sarraf-Zadegan N, Sayed-Tabatabaei FA, Bashardoost N, Maleki A, Totonchi M, Habibi HR, et al. The prevalence of coronary artery disease in an urban population in Isfahan, Iran. Acta Cardiol 1999; 54(5): 257-63.
- **32.** Sarrafzadegan N, Kelishadi R, Baghaei A, Hussein SG, Malekafzali H, Mohammadifard N, et al. Metabolic syndrome: an emerging public health problem in Iranian women: Isfahan Healthy Heart Program. Int J Cardiol 2008; 131(1): 90-6.
- 33. Sarraf Zadegan N, Baghaei A, Sadri CH, Kelishadi R, Malekafzali H, Boshtam M, et al. Isfahan healthy heart program: Evaluation of comprehensive, community-based interventions for non-communicable disease prevention. Preventionand control 2006; 2(2): 73-84.
- **34.** 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(19): 2486-97.
- **35.** World Health Organization. Obesity: Preventing and Managing the Global Epidemic: Report of a WHO Consultation on Obesity IIS microfiche library. Geneva: World Health Organization; 2000. p. 894.
- **36.** Warnick GR, Benderson J, Albers JJ. Dextran sulfate-Mg2+ precipitation procedure for quantitation of high-density-lipoprotein cholesterol. Clin Chem 1982; 28(6): 1379-88.
- **37.** Friedewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low-density lipoprotein cholesterol in plasma, without use of the preparative ultracentrifuge. Clin Chem 1972; 18(6): 499-502.
- **38.** World Health Organization. Cardiovascular Diseases Unit. Monica manual. Geneva: World Health Organization; 1990.
- **39.** Regitz-Zagrosek V, Lehmkuhl E, Mahmoodzadeh S. Gender aspects of the role of the metabolic syndrome as a risk factor for cardiovascular disease. Gend Med 2007; 4 (Suppl B): S162-77.
- **40.** Nilsson G, Hedberg P, Jonason T, Lonnberg I, Ohrvik J. Heart rate recovery is more strongly associated with the metabolic syndrome, waist circumference, and insulin sensitivity in women than in men among the elderly in the general population. Am Heart J 2007; 154(3): 460-7

How to cite this article: Khaledi Y, Aghababaei E, Sadeghi M, Hashemi M, Sanei H. Evaluation of heart rate reserve and high-sensitivity C-reactive protein in individuals with and without metabolic syndrome in Isfahan, Iran. ARYA Atherosclerosis Journal 2012; 8(2): 70-75.