



RESEARCH ARTICLE

Open Access

Prevalence of metabolic syndrome and its correlated factors among children and adolescents of Ahvaz aged 10 – 19

Homeira Rashidi¹, Seyed Peyman Payami^{1*}, Seyed Mahmoud Latifi¹, Majid Karandish², Armaghan Moravej Aleali¹, Majid Aminzadeh¹, Kuroush Riahi¹ and Marzieh Ghasemi¹

Abstract

Background: Population-based studies for prevalence of metabolic syndrome (M.S) in children and adolescents are relatively rare. The aim of this study was to assess the Prevalence of M.S and correlated factors among children and adolescents aged 10 to 19 years in Ahvaz.

Methods: In this descriptive-analytical population- based study, 2246 children and adolescents, 10–19 years old (1113 male and 1133 female) in Ahvaz, were evaluated. Anthropometry, biochemical measurement and blood pressure (BP) were assessed. Modified ATP III criteria 2005 were used for M.S. definition. Center for disease and Control preventions (CDC) percentile were applied to define cut off points of waist circumference and BP.

Results: Prevalence of M.S. was 9% (95% CI: 8-10%) with prevalence in male 11% (95% CI: 10-12%) and female 7% (95% CI 6-8%). Among individuals with M.S, triglyceride (TG) and decreased high density lipoprotein (HDL) cholesterol levels were the most common components (33.5% and 24.1%, respectively). Prevalence of M.S was higher in overweight persons comparing to participants with at risk and normal weight group (in male: 24.1%, 14.3% and 9.9% respectively $P = 0.0001$), (in female: 22.6%, 18.3% and 4.5% respectively $P = 0.0001$). Among the correlated factors of M.S age ($P = 0.0006$), sex and BMI ($P = 0.0001$) had significant differences between subjects with and without M.S. whereas there was no significant difference between two groups in ethnicity, history of breast fed, birth weight neonatal ICU admission, maternal history(GDM, gestational HTN, Parity) and family history of HTN, obesity and DM ($P > 0.05$).

Conclusion: This study shows high prevalence of M.S in Children and Adolescents in south west of Iran (Ahvaz) especially in overweight persons.

Keywords: Metabolic syndrome, Prevalence, Children, Adolescents, Cardiovascular risk factor, Iran

Introduction

Since early 1960s, an association between obesity and elevated level of triglycerides (TG), decreased level of high-density lipoprotein (HDL) cholesterol, hyperinsulinemia, impaired glucose tolerance, high blood pressure (BP) and cardiovascular disease, (CVD) has been discussed. For first time it was introduced in 1988 by Reaven et al. as metabolic syndrome (M.S) [1]. M.S is the risk factor of all cause and C.V. mortality in adults with and without type 2 diabetes [2]. According to some studies, M.S could be

originated from embryonic period [3,4]. Although the prevalence of M.S and its risk factors have been widely studied in adults, limited studies are available in children and adolescents so that there is no clear definition of M.S in this age group. Therefore more studies are required to provide a better definition of M.S in children and adolescents. Currently, the same risk factors of adults are used in this age group based on age and gender percentiles from global or rational data or study specific distributions. Based on limited studies 4.2% of the American children and adolescents who participated in the NHANES III 1988-1994 had M.S. [5]. That increased to 6.2% in NHANES 1999–2000 [6]. As the prevalence of overweight in American children and adolescents increase [7], the prevalence of M.S is

* Correspondence: peyman_payami1@yahoo.com

¹Health research Institute, Diabetes Research Center, Ahvaz Jundishapur University of Medical Sciences, Ahvaz 61357-15794, Iran
Full list of author information is available at the end of the article

likely to rise, such that above study showed that 29% and 32% of overweight subjects had M.S., respectively. As well, a recent study in Tehran revealed that the prevalence of overweight in Iranian children and adolescents was high with an estimated prevalence of 21% [8]. On the other hand, in another study in Tehran, the prevalence of M.S in children and adolescents was estimated about 10% based on modified ATP III criteria [9]. These studies indicated that the prevalence of M.S is significantly higher in more obese subjects. Recently, the prevalence of M.S has been reported among Mexican [10], Tunisian [11], Indian [12] and Chinese [13] children and adolescents. Different genetic, racial [14] and environmental factors like lifestyle, change of lifestyle from rural to urban, physical activity and socioeconomic conditions play significant role in the prevalence of M.S. Therefore, this study was designed to determine the prevalence of M.S and its risk factors among children and adolescents of Ahvaz aged 10–19 years old.

Materials and methods

This descriptive-analytic study performed in Ahvaz, capital city of Khuzestan province (South West of Iran) in 2009–2011. Of 25 health centers, 6 ones were selected randomly by multi-steps cluster sampling method. Informed written consent was obtained from subjects aged 10–19 and parents of subjects under the age of 19. A questionnaire including age, sex, ethnicity, birth weight, history of breast feeding, neonatal ICU admission, history of pregnancy in participant's mother [gestational diabetes mellitus (GDM), gestational hypertension and parity number], family history of diabetes mellitus (DM), hypertension (HTN) and obesity was filled for each subject. In this study after excluding subjects taking medication that would affect serum lipids, blood pressure and carbohydrate metabolism and subjects with history of chronic disease, of heart, lung, kidney and liver and chronic diarrhea and hospitalization during three months ago, individual with full relevant data were included. Blood pressure was measured twice at least 30 minutes apart by an appropriate mercury sphygmomanometer after 15 minutes rest in a sitting position. The mean of the two readings was recorded as individual blood pressure.

Waist circumference was measured at the midpoint of the lowest rib and iliac crest over light clothing at the end of exhalation using a tape measure. Heights and weights were measured, using a tape measure in a standing position with bare feet and seca scales with minimum possible clothing, respectively. Body mass index (BMI) [weight (Kg)/Height (m)²] was calculated for each subject. Blood samples were drawn after a 12-hour overnight fasting. About 30–45 minutes after sampling, the samples were centrifuged by 2500–3000 rpm for 10 minutes; sera were stored in refrigerator and then sent to laboratory.

Fasting blood sugar (FBS), TG, total cholesterol and HDL cholesterol were measured by enzyme-calorimetric method using Pars Azmoon kits (with biotechnical instruments type BT-3000 Germany). Modified ATP III criteria 2005 [5,6] were used to define M.S as follows:

- 1- Abdominal obesity (waist circumference \geq the age and sex specific 90th percentile using C.D.C percentiles)
- 2- Elevated BP (systolic and/or diastolic blood pressure \geq the age and sex -specific 90th percentile using C.D.C percentiles except for 18 and 19 years old subjects, for whom the cut off values of \geq 130 and/or \geq 85 mmHg for systolic and diastolic blood pressure were used, respectively.
- 3- HDL-cholesterol \leq 40
- 4- TG \geq 110 mg/dl
- 5- FBS \geq 100mg/dl

Subjects with 3 or more characteristics of the above components were categorized as M.S.

Overweight, at risk for Overweight and normal weight were defined based on the study specific percentile curves of BMI for age and sex as \geq 95th percentile, \geq 85 to $<$ 95th percentile and $<$ 85th percentile, respectively.

Descriptive statistics were used to prepare graphs and tables. Chi -Square test was applied to compare ratios and multivariate logistic regression analysis was used to determine the potential determinants of M .S.

$P < 0.05$ was considered as significant. All data were analyzed by SPSS software version 19.

Results

A total of 2246 children and adolescents (1120 boys, 1139 girls) aged 10–19 participated in our investigation. Baseline characteristics of study population according to sex is shown in Table 1. According to the modified ATP III criteria, the prevalence of M.S was 9% (95% CI: 0.08-0.10). The prevalence of MS in boys [11% (95% CI: 10-12%)] was significantly higher than girls [7% (95% CI: 6-8%)], ($P=0.001$) (Table 2). In both gender, the prevalence of MS was different between age groups. The prevalence of MS decreases approximately after the age of 10 to 11 while it increases after the age 14 and highest rate was observed at ages 16 to 17. In boys, the relationship between the prevalence of M.S and increase of age was insignificant ($P = 0.403$) but it was significant in girls ($P = 0.010$) (Figure 1).

When examined by BMI category, overweight group had a higher prevalence of MS than at risk for overweight and normal weight groups ($P = 0.0001$) (Table 3).

In this study, 46% and 5% of the participants with a waist circumference \geq 90th percentile and $<$ 90% percentile had M.S respectively ($P = 0.0001$). The mean of waist circumference between two genders showed significant

Table 1 Basic characteristics of the study population according to sex

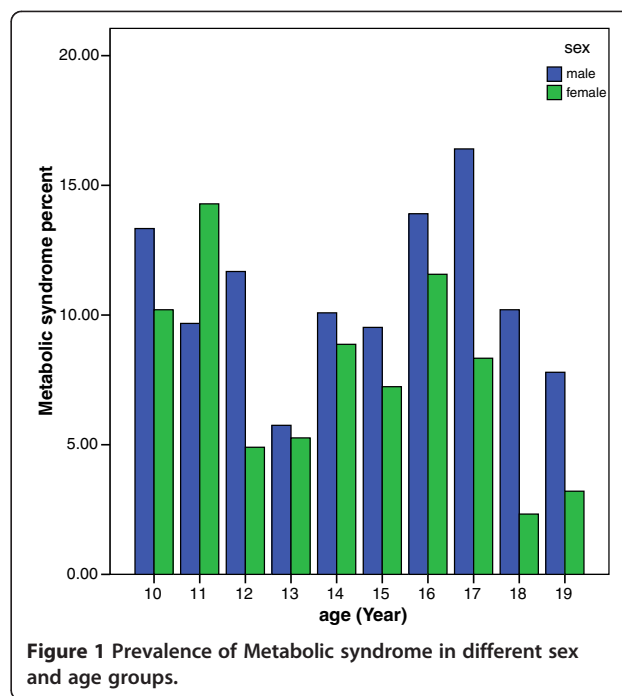
Variable	Male	Female	P value
Age (year)	14.63 ± 2.58	15.27 ± 2.74	0.0001
Height (cm)	162.83 ± 13.82	151.81 ± 9.32	0.0001
Weight (Kg)	52.39 ± 14.69	50.99 ± 12.39	0.015
BMI (Kg/m ²)	19.75 ± 4.43	20.83 ± 4.20	0.0001
Waist circumference (cm)	70.69 ± 11.23	68.19 ± 9.82	0.0001
HDL (mg/dl)	53.77 ± 12.18	55.62 ± 11.77	0.0001
TG (mg/dl)	111.62 ± 67.07	100.52 ± 57.66	0.0001
SBP (mm Hg)	106.58 ± 11.25	106.32 ± 10.99	0.574
DPB (mm Hg)	60.65 ± 10.20	64.85 ± 9.68	0.0001
FBS (mg/dl)	91.85 ± 12.45	89.16 ± 12.45	0.012

difference ($P = 0.0001$). The prevalence of each component of M.S in subjects with M.S was: abdominal obesity 10.3% elevated BP 22.1%, low HDL-cholesterol 24.1%, high TG 33.5% and high FBS 16.4% (Table 4). In subjects with M.S., 81.8% had three components of M.S, 16.7% had four and 1.5% had five ones. In the whole study population, 36.5% were normal (no components of M.S), 34.6% had one, 19.9% two, 7.3% three, 1.5% four and 0.1% had five components.

Factors that can affect the prevalence of M.S are seen in Table 5. Among the correlated factors of M.S age ($P = 0.006$), sex and BMI ($P = 0.0001$) had significant differences between subjects with and without M.S. whereas ethnicity, breast feeding, birth weight, neonatal ICU admission, maternal history (GDM, Gestational HTN, Parity no) and family history of HTN, obesity and D.M had no differences ($P > 0.05$).

Table 2 Prevalence of M.S in different sex and age groups

Age (year)	N	No. of M.S (%) Total	N	No. of M.S (%) (Male)	N	No. of M.S (%) (Female)
10	94	11 (11.7)	45	6 (13.3)	49	5 (10.2)
11	208	24 (11.5)	124	12 (9.7)	84	12 (14.3)
12	239	21 (8.8)	137	16 (11.7)	102	5 (4.9)
13	163	9 (5.5)	87	5 (5.7)	76	4 (5.3)
14	243	23 (9.5)	119	12 (10.1)	124	11 (8.9)
15	299	25 (8.4)	147	14 (9.5)	152	11 (7.2)
16	272	35 (12.9)	151	21 (13.9)	121	14 (11.6)
17	236	30 (12.7)	128	21 (16.4)	108	9 (8.3)
18	227	13 (5.7)	98	10 (10.2)	129	3 (2.3)
19	265	12 (4.5)	77	6 (7.8)	188	6 (3.2)
Total	2246	203 (9)	1113	123 (11.1)	1133	80 (7.1)
P Value		0.0006		0.0403		



Discussion

In this study, the prevalence of M.S among children and adolescents aged 10–19 was 9% based on the modified ATPIII 2005 criteria which was significantly higher in boys than girls ($P = 0.001$). The prevalence of M.S in our study was similar to the study of Esmailzadeh et al. [9] in Tehran based on same criteria. In that report, prevalence of M.S was not significantly different between boys and girls and study-specific distributions to define thresholds of risk factors were used. In another study in Tehran by Chiti et al. [15], the prevalence of M.S among children and adolescents aged 10–19 was 9.5% based on the same criteria which was higher in boys than girls. The result of that study was consistent with ours. Kelishadi et al. [16] conducted another study in students aged 6–18 in Isfahan that showed the

Table 3 Prevalence of metabolic syndrome in different body weight groups

	BMI status (percentile)	No. of subjects (%)	Subjects with M.S (%)	P value
Total	Normal (<85)	1911 (85.1)	7.2	0.0001
	At risk (85-95)	227 (10.1)	16.3	
	Overweight (≥ 95)	108 (4.8)	26.9	
Male	Normal (<85)	946 (85.1)	9.9	0.003
	At risk (85-95)	112 (10.1)	14.3	
	Overweight (≥ 95)	54 (4.9)	24.1	
Female	Normal (<85)	946 (85.1)	4.5	0.0001
	At risk (85-95)	115 (10.2)	18.3	
	Overweight (≥ 95)	54 (4.8)	29.6	

Table 4 Prevalence of individual component of metabolic syndrome according to age and sex among subjects with metabolic syndrome

Age (year)	Abdominal obesity (%)	TG \geq 110 (%)	HDL \leq 40 (%)	BP \geq 130/85 (%)	FBS \geq 100 (%)
Total population					
10	9.7	34	18.1	36.2	21.3
11	11.4	34.6	17.3	34.6	19.2
12	10.3	35.2	20.9	24.3	18.6
13	9.4	37.7	19	16	14.2
14	10.2	33.3	27.2	21.4	25.1
15	9.6	36.7	27.8	26.1	11.2
16	10.7	31.8	30.5	36.4	15.4
17	10	40.6	28	25.4	15
18	11.1	26.3	21.1	3.1	12.5
19	10	26.1	23.1	3.8	14.9
Total	10.3	33.5	24.1	22.1	16.4
Males					
10	11.1	31.1	11.1	24.4	22.2
11	9.7	33.9	15.3	33.9	18.5
12	11.1	34.8	19.7	35	21.3
13	9.3	37.2	14.9	16.1	17.4
14	9.5	34.5	26.1	24.4	30.3
15	9.7	40.3	31.3	27.9	13.9
16	10.6	36.1	37.1	35.8	15.6
17	9.6	48.4	32.8	24.2	15.6
18	11.5	32	35.7	5.1	17.5
19	10.4	36.8	36.4	3.9	23.7
Total	10.2	37.1	27.1	25	19.1
Females					
10	8.3	36.7	24.5	46.9	20.4
11	14.1	35.7	20.2	35.7	20.2
12	9.2	35.6	22.5	9.8	14.9
13	9.5	38.2	23.7	15.8	10.5
14	10.8	23.3	27.2	18.5	20.2
15	9.5	33.3	24.3	38.2	8.7
16	10.7	26.7	22.3	37.2	15
17	10.5	31.1	22.2	26.9	14.2
18	10.9	22	10.1	1.6	8.7
19	9.8	21.6	17.6	3.7	11.4
Total	10.3	29.9	21.1	19.3	13.6

TG: Triglyceride, HDL-D: high density lipoprotein cholesterol, BP: blood pressure, FBS: fasting blood sugar.

prevalence of M.S as 14%, indicating a much difference with above- mentioned studies.

Other studies in different cities of Iran [17-19], not population- based, could not reflect the whole society. Population - based epidemiologic studies in other countries have shown various prevalence rates. In the U.S, the prevalence of M.S among children and adolescents has

been reported 3.1% to 12.7% with different definitions and cut off points [20]. These studies have revealed that the prevalence of M.S. is increasing by time in this age group. The investigation of Herrabi et al. [11] in Tunisia showed that the prevalence of M.S. is 4% in urban regions based on the modified -ATPIII criteria. Overweight and obesity were the most common components of the syndrome. In

Table 5 Risk factors of metabolic syndrome among children and adolescences

	B	S.E	P value	OR	Confidence interval (CI)
Sex (M/F)	0.643	0.178	0.000	1.903	1.341-2.700
Ethnicity (Arab/Fars)	-0.019	0.174	0.914	0.981	0.698-1.380
Birth weight	0.207	0.139	0.135	1.230	0.937-1.614
Neonatal nutrition (breastfeeding/other)	0.003	0.617	0.991	1.003	0.630-1.596
ICU Admission	0.356	0.508	0.483	1.428	0.528-3.863
Family history					
Obesity	0.087	0.183	0.647	1.087	0.760-1.555
Diabetes	0.430	0.223	0.054	1.523	0.993-2.380
HTN	0.278	0.226	0.204	1.333	0.856-2.076
Maternal history					
Parity no.	0.045	0.042	0.286	1.046	0.963-1.136
GDM	-0.305	0.508	0.548	0.737	0.273-1.994
Gestational HTN	0.362	0.360	0.314	1.437	0.709-2.910
BMI					
BMI 85-95/BMI<85	0.970	0.243	0.001	2.639	1.640-4.248
BMI≥95/BMI<85	1.645	0.303	0.001	5.180	2.859-9.387
Age (Year)					
11/10	-0.381	0.429	0.374	0.683	0.295-1.583
12/10	-0.326	0.415	0.432	0.722	0.320-1.628
13/10	-0.945	0.490	0.054	0.389	0.149-1.017
14/10	-0.556	0.426	0.192	0.573	0.249-1.322
15/10	-0.672	0.420	0.109	0.511	0.224-1.163
16/10	-0.244	0.403	0.546	0.784	0.356-1.728
17/10	-0.493	0.422	0.243	0.611	0.267-1.397
18/10	-1.196	0.473	0.011	0.302	0.120-0.763
19/10	-1.875	0.519	0.000	0.153	0.056-0.424

S.E: Standard error.

B: coefficient of variable.

O.R: Odd ratio.

another study in north India, Singh et al. [13] reported the prevalence of M.S to be 4.2% using modified ATP III criteria. That study again confirmed that the prevalence of M.S is higher in at risk for overweight and overweight persons. A study in Mexico by Rodriges -Moran et al. [11] using modified ATP III criteria showed prevalence of M.S. to be 6.5% while it was 4.5% by WHO criteria. As well, an investigation in China by Liet of [13] using the criteria proposed by Ferranti et al. [20], reported the prevalence of M.S to be 3.7%. According to that study, it is estimated that 3 million children and adolescents have M.S in China. The lack of a comprehensive and clear definition of M.S in children and adolescents could explain to some extent. The difference of prevalence rates reported in various studies throughout the world. On the other hand, the factors like various prevalence of overweight, race, genetic factors, change in lifestyle from rural to urban, physical activity level and socioeconomic condition could be considered.

Like other studies in Iran [8,19,21], our investigation showed a higher prevalence of M.S. among at risk for overweight and overweight children and adolescents in both genders.

On the other hand, the prevalence of overweight in Iranian children has been reported to be up to 8%. The change of nutritional style in Iranian children and adolescents is probably an important reason for the uprising problem of overweight and M.S. As well, some evidences are available that indicates decrease in physical activity of Iranian children and adolescents [22,23]. Moreover, recent epidemiological studies show a parallel increase in type 2 diabetes mellitus in children/adolescents and obesity [24,25]. Unfortunately, the rise in prevalence of M.S. and type 2 D.M in children and adolescence lead to an increase in correlated complications in young adults, including early C.V.D [20].

The current study has several strengths. It was a population- based study with a relatively large sample size. Moreover numerous correlated factors were assessed. One

limitation was the definition of M.S. limitations of such a definition for children and adolescents has been discussed previously [5,20,26]. Cross sectional nature of this study was the other limitation that did not allow us to make causal inference. Our study provides evidences showing a high prevalence of M.S. among Iranian children and adolescents, especially in overweight subjects. In the future, large prospective studies should be conducted to confirm the association between above-mentioned factors and M.S.

Conclusion

Among the factors correlated to M.S., age, sex and BMI were significantly different between subjects with and without M.S. whereas ethnicity, breast feeding, birth weight, neonatal ICU admission, maternal history (GDM, gestational hypertension, parity number) and family history of hypertension, obesity and diabetes mellitus did not show significant differences.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

HR, MA and MK designed the study. SPP drafted the manuscript. SML has done the statistical analysis. AMA, KR and MGh have done data collection, references search, writing and editing manuscript. All authors read and approved the final manuscript. Thanks to all authors for their support and help in this study.

Acknowledgement

This paper is issued from the research project with the registered number of D-8703. Financial support was provided by Health Research Institute, Diabetes Research Center, Ahvaz Jundishapur University of Medical Sciences. The authors would like to thank all staffs of diabetes research center, especially Miss Reshadatian, Dehghan and Hardani and Dr. Ghorbani for their help in this study.

Author details

¹Health research Institute, Diabetes Research Center, Ahvaz Jundishapur University of Medical Sciences, Ahvaz 61357-15794, Iran. ²Nutrition and Metabolic research center, Ahvaz Jundishapur University of Medical Sciences, Ahvaz, Iran.

Received: 6 February 2013 Accepted: 8 March 2014

Published: 28 April 2014

References

1. Reaven GM: **Banting Lecture 1988. Role of insulin resistance in human disease.** *Diabetes* 1988, **37**:1595–1607.
2. Rosenzweig JL, Ferrannini E, Grundy SM, Haffner SM, Heine RJ, Horton ES, Kawamori R, Endocrine Society: **Primary prevention of cardiovascular disease and type 2 diabetes in patients at metabolic risk: an endocrine society clinical practice guideline.** *J Clin Endocrinol Metab* 2008, **10**:3671–3689.
3. Zaliūnas R, Slapikas R, Luksiene D, Slapikiene B, Statkeviciene A, Milvidaitė I, Gustiene O: **Prevalence of metabolic syndrome components in patients with acute coronary syndromes.** *Medicina (Kaunas)* 2008, **44**(3):182–188.
4. Qadan LR, Ahmed AA, Safar HA, Al-Bader MA, Ali AA: **Prevalence of metabolic syndrome in patients with clinically advanced peripheral vascular disease.** *Angiology* 2008, **59**(2):198–202.
5. Cook S, Weitzman M, Auinger P, Nguyen M, Dietz WH: **Prevalence of a metabolic syndrome phenotype in adolescents: findings from the third National Health and Nutrition Examination Survey, 1988–1994.** *Arch Pediatr Adolesc Med* 2003, **157**:821–827.
6. Duncan GE, Li SM, Zhou XH: **Prevalence and trends of a metabolic syndrome phenotype among U.S. Adolescents, 1999–2000.** *Diabetes Care* 2004, **27**(10):2438–2443.
7. Troiano RP, Flegal KM: **Overweight children and adolescents: description, epidemiology, and demographics.** *Pediatrics* 1998, **101**(3 Pt 2):497–504.
8. Mohammadpour Ahranjani B, Rashidi A, Karan Dish M, Eshraghian MR, Kalantari N: **Prevalence of overweight and obesity in adolescent tehrani student S, 2000–2001: an epidemic health problem.** *Public Health Nutr* 2004, **7**:645–8.
9. Esmaillzadeh A, Mirmiram P, Azadbakht L, Etemadi A, Azizi F: **High prevalence of the metabolic syndrome in Iranian Adolescents.** *Obesity* 2006, **14**(3):377–382.
10. 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**(10):2516–2517.
11. Harabbi I, Bouaouina M, Maatoug J, Gaha R, Ghannem H: **Prevalence of the metabolic syndrome among urban schoolchildren in Sousse, Tunisia.** *Int J Cardiol* 2009, **135**(1):130–131.
12. Singh R, Bhansali A, Sialy R, Aggarwal A: **Prevalence of metabolic syndrome in adolescents from a north Indian population.** *Diabet Med* 2007, **24**(2):195–199.
13. Li Y, Yang X, Zhai F, Kok FJ, Zhao W, Piao J, Zhang J, Cui Z, Ma G: **Prevalence of the metabolic syndrome in Chinese adolescents.** *Br J Nutr* 2008, **99**(3):565–570.
14. Henneman P, Aulchenko YS, Frants RR, van Dijk KW, Oostra BA, van Duijn CM: **Prevalence and heritability of the metabolic syndrome and its individual components in a Dutch isolate: the Erasmus Rucphen Family study.** *J Med Genet* 2008, **45**(9):572–577.
15. Chiti H, Hosseinpahan F, Mehrabi Y, Azizi F: **The prevalence of metabolic syndrome in adolescents with varying degree of body weight: Tehran Lipid And Glucose study(TLGS).** *Iran J Endocrinol Metab* 2010, **11**(6):625–637.
16. Kelishadi R, Ardalan G, Gheiratmand R, Adeli K, Delavari A, Majdzadeh R: **Paediatric metabolic syndrome and associated anthropometric indices: the CASPIAN Study.** *Acta Paediatr* 2006, **95**:1625–1634.
17. Salem Z, Vazirinejad R: **Prevalence of metabolic syndrome components among children 7–11 years old in Rafsanjan 2008.** *Iran J Nutr Sci Food Ind* 2010, **5**(2):63–71.
18. Ghargerechi R, Razzaghy AM: **Prevalence of Metabolic Syndrome in Obese Children and Adolescents Dolescents.** *Tbriz Med J* 2010, **32**(3):57–61.
19. Mirhosseini NZ, Yusoff NA, Shahar S, Parizadeh SM, Mobarhen MG, Shakeri MT: **Prevalence of the metabolic syndrome and its influencing factors among adolescent girls in Mashhad, Iran.** *Asia Pac J Clin Nutr* 2009, **18**:131–136.
20. De Ferranti SD, Osganian SK: **Epidemiology of paediatric metabolic syndrome and type 2 diabetes mellitus.** *Diab Vasc Dis Res* 2007, **4**(4):285–296.
21. Azizi F, Salehi P, Etemadi A, Zahedi-Asl S: **Prevalence of metabolic syndrome in an urban population: Tehran Lipid and Glucose Study.** *Diabetes Res Clin Pract* 2003, **61**:29–37.
22. Etemadi A, Malekzadeh R: **Definition and etiology of metabolic syndrome.** *Arch Iran Med* 2008, **11**:1–2.
23. Ghassemi H, Harrison G, Mohammad K: **An accelerated nutrition transition in Iran.** *Public Health Nutr* 2005, **5**:149–155.
24. Rosenbloom A, Arsalanian S, Brink S, Conschafer K, Jones KL, Klingensmith G, Neufeld N, White N, Fagot-Campagna A, Gahagan S, Linder B: **Type 2 diabetes in children and adolescents.** *Pediatrics* 2000, **105**(3 Pt 1):671–680.
25. Libman I, Arslanian SA: **Type II diabetes mellitus: no longer just adults.** *Pediatr Ann* 1999, **28**:589–593.
26. Mancini M: **Metabolic syndrome in children and adolescents: criteria for diagnosis.** *Diabetol Metab Syndr* 2009, **19**(1):20.

doi:10.1186/2251-6581-13-53

Cite this article as: Rashidi et al.: Prevalence of metabolic syndrome and its correlated factors among children and adolescents of Ahvaz aged 10 – 19. *Journal of Diabetes & Metabolic Disorders* 2014 **13**:53.