

Association of Visceral Fat and Risk Factors for Metabolic Syndrome in Children and Adolescents

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Purpose: Visceral fat (VF) is closely associated with many metabolic risk factors and is also known to be a strong predictive factor for severe metabolic complications in adults. But there are only a few studies concerning the association of VF and risk factors for metabolic syndrome (MS) in children and adolescents. In our study, we emphasized the association of VF [measured by VF computed tomography (VFCT)] and risk factors for metabolic syndrome in children and adolescents. **Materials and Methods:** The subjects were outpatients aged 6 to 18 years who underwent VFCT in the family medicine of The Catholic University of Korea from January 2005 to August 2009. There were 82 patients in total (42 children, 40 adolescents). Height, weight, blood pressure (BP), blood tests, body composition analysis and VF were measured. The three groups were also classified by metabolic score. **Results:** In children, only high density lipoprotein cholesterol (HDL-C) showed a statistically significant difference, while in adolescents, triglyceride, HDL-C, BP, body mass index (BMI), waist circumference (WC) and VFA showed statistically significant differences. In terms of VFA, fasting glucose, BP, BMI, basal metabolic rate (BMR) and WC showed statistically significant differences. BMI showed a statistically significant difference in terms of BP, BMR, WC, VFA and HDL-C. **Conclusion:** There is a need to acknowledge the statistically significant associations of VF and risk factors for MS in children and adolescents. Screening tests for BP, cholesterol, fasting glucose and WC should be given in clinics for children and adolescents so that MS can be detected and its risk factors treated early.

Key Words: Children, adolescents, metabolic syndrome, visceral fat

INTRODUCTION

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In recent years, the prevalence of overweight or obese children and adolescents has been rapidly increasing.¹ The rate of increase exceeds even that of adults.² A 10-fold increase (1.7→17.9%) for boys and a 4.5-fold increase (2.4→10.9%) for girls was reported in Seoul between 1979 and 2002.³ Obese children and adolescents are more likely to develop into obese adults,^{4,5} and also have a high risk of

affliction with associated metabolic complications as they grow to adults. For instance, Type II diabetes mellitus, hypertension, dyslipidemia and atherosclerosis can develop.⁶⁻⁸ This fact causes excessive personal and social cost, and is the reason why more aggressive intervention and countermeasures must be implemented to control metabolic syndrome (MS) in Korean children and adolescents.^{9,10} Until now, there has been insufficient information concerning diagnostic criteria for MS in children and adolescents. In children and adolescents, physical as well as physiological changes are taking place. Blood pressure (BP), body mass index (BMI) and blood cholesterol levels vary according to age, which makes it hard to set a cut off value for diagnostic criteria. Fortunately, in 2007, the International Diabetes Federation reported the diagnostic criteria for MS in children and adolescents,¹¹ and this criterion is being used along with the variation for children and adolescents of National Cholesterol Education Program (NCEP) Adult Treatment Panel (ATP) III criteria. VF is closely associated with many metabolic risk factors and is also known to be a strong predictive factor of severe metabolic complications in adults.¹²⁻¹⁷ However, few studies have been done concerning the association of VF and risk factors for MS in children and adolescents. In our study, we emphasized the association between VF [measured by VF computed tomography (VFCT)] and risk factors for MS in children and adolescents.

MATERIALS AND METHODS

Materials

All subjects gave written informed consent before taking part in this study. The subjects were outpatients, aged 6 to 18 years, who underwent VFCT in the family medicine clinic of The Catholic University of Korea located in Seoul and Gyeonggi province from January 2005 to August 2009. There were 82 patients in total (42 children, 40 adolescents). Height, weight, BP, blood tests, body composition analysis and VF were measured. The three groups of children and adolescents were then classified by metabolic scores. The study was approved by The Catholic University of Korea's ethics committee.

Methods

Height and weight were measured with the patient wearing light clothes or only a gown. We measured up to 0.1 kg, 0.1

cm respectively. Waist circumference (WC) was measured while the patient exhaled slightly, with no pressure applied, and then we measured the area between the lowest rib and the upper part of the iliac crest. BP was measured according to the recommendations from the American Heart Association (AHA). The patients relaxed for more than 10 minutes and while they were sitting we measured twice, each measurement 5 minutes apart. The final BP was the mean value of these two measurements. The patients did not eat anything for 12 hours prior to the blood tests. After withdrawing 15 mL of blood, it was sent to the laboratory for fasting glucose, total cholesterol, triglyceride, high density lipoprotein cholesterol (HDL-C) etc. Body composition analysis was made using Biospace Corp. In Body 4.0 body fat analyzer. VFCT was referred to the radiologist. The lower abdomen of each subject was covered with lead protectors in order to protect them from radioactive exposure. VFCT was done around the 4th lumbar vertebra area where subcutaneous and visceral fat were measured.

Definition

The definition of the MS in children and adolescents was made by the corrected criteria of the NCEP ATP III. If three or more criteria were satisfied, we considered it MS in children and adolescents (Table 1).¹⁸

Statistical analysis

According to NCEP ATP III criteria, the MS score was classified as 0-1, 2-3 and 4-5. We compared fasting blood glucose, total blood cholesterol, blood triglyceride, blood HDL-C, systolic and diastolic BP, basal metabolic rate (BMR), WC, subcutaneous fat, VF and total fat area. All results were denoted as a mean \pm S.D. A t-test for independent samples was used to assess the differences between two groups. The three groups were classified by MS score and analyzed by an ANOVA test. Visceral fat area (VFA), BMI and other risk factors were analyzed by the Pearson correlation. Statistical analysis of the data was done by Statistical Package for Social Science (SPSS) 15.0 for Windows (SPSS Inc, Chicago, IL, USA). We considered a *p* value of less than 0.05 to be statistically significant.

RESULTS

Patients and baseline characteristics

There were 82 subjects in total (children: 42, adolescents:

Table 1. Modified Criteria of Metabolic Syndrome in Children and Adolescents

Criteria	Adults & Adolescents (NCEP ATP III)	Children
Triglyceride (mg/dL)	≥ 150	≥ 110
HDL-C (mg/dL)		
Boys	< 40	≤ 40
Girls	< 50	≤ 40
Fasting glucose (mg/dL)	≥ 110	≥ 110
Waist circumference (cm)		
Boys	≥ 102	≥ 90th% for age, sex
Girls	≥ 88	≥ 90th% for age, sex
Blood pressure (mmHg)	≥ 130/85	≥ 90th% for age, sex, height

NCEP ATP III, National Cholesterol Educational Program Adult Treatment Panel III; HDL-C, high density lipoprotein cholesterol.

Table 2. Characteristics of the Study Population (n = 82)

	Boys (n = 42)		Girls (n = 40)	
	Children (n = 27)	Adolescents (n = 15)	Children (n = 15)	Adolescents (n = 25)
Age (yrs)	10.1 ± 1.5	15.5 ± 1.6	9.1 ± 1.8	15.5 ± 1.7
Fasting glucose (mg/dL)	96.1 ± 9.8	103.9 ± 9.1	93.9 ± 8.7	95.3 ± 17.0
Triglyceride (mg/dL)	137.7 ± 66.5	152.6 ± 51.6	152.7 ± 86.6	135.9 ± 60.6
HDL-C (mg/dL)	45.6 ± 12.6	40.0 ± 8.5	43.4 ± 6.8	43.5 ± 9.0
Systolic BP (mmHg)	120.9 ± 11.7	138.6 ± 10.4	116.5 ± 11.3	128.1 ± 15.9
Diastolic BP (mmHg)	73.4 ± 8.4	77.1 ± 6.8	69.5 ± 10.0	77.6 ± 10.0
BMI (kg/m ²)	27.1 ± 3.4	31.4 ± 4.5	26.1 ± 5.8	32.4 ± 7.7
%Body fat	38.6 ± 4.7	33.1 ± 5.3	37.0 ± 4.9	42.3 ± 7.7
BMR (kcal)	1211.3 ± 185.2	1868.3 ± 446.2	1214.0 ± 309.4	1505.1 ± 312.8
Waist circumference (cm)	90.9 ± 6.6	97.5 ± 7.9	80.7 ± 10.5	94.1 ± 9.5
Visceral fat area (cm ²)	75.2 ± 26.0	87.6 ± 30.8	61.9 ± 15.2	89.4 ± 42.5
Subcutaneous fat area (cm ²)	213.6 ± 67.0	272.9 ± 84.6	197.4 ± 100.8	336.1 ± 144.5
Total abdominal fat area (cm ²)	292.2 ± 86.4	376.0 ± 84.6	263.0 ± 109.7	441.9 ± 198.8

BP, blood pressure; HDL-C, high density lipoprotein cholesterol; BMI, body mass index; BMR, basal metabolic rate.

40). The mean age of the children was 10.1 ± 1.5 yrs for boys and 9.1 ± 1.8 yrs for girls. The mean age of the adolescents was 15.5 ± 1.6 yrs for boys and 15.5 ± 1.7 yrs for girls. Fasting blood glucose (FBG) was 96.1 ± 9.8 mg/dL (boys) and 93.9 ± 8.7 mg/dL (girls) in children, and 103.9 ± 9.1 mg/dL (boys) and 95.3 ± 17.0 mg/dL (girls) in adolescents. Blood triglyceride levels were 137.7 ± 66.5 mg/dL (boys) and 152.7 ± 86.6 mg/dL (girls) in children, and 152.6 ± 51.6 mg/dL (boys) and 135.9 ± 60.6 mg/dL (girls) in adolescents. Blood HDL-C was 45.6 ± 12.6 mg/dL (boys) and 43.4 ± 6.8 mg/dL (girls) in children, and 40.0 ± 8.5 mg/dL (boys) and 43.5 ± 9.0 mg/dL (girls) in adolescents. Systolic BP was 120.9 ± 11.7 mmHg (boys) and 116.5 ± 11.3 mmHg (girls) in children, and 138.6 ± 10.4 mmHg (boys) and 128.1 ± 15.9 mmHg (girls) in adolescents. BMR was 1211.3 ± 185.2 kcal

(boys) and 1214.0 ± 309.4 kcal (girls) in children, and 1868.3 ± 446.2 kcal (boys) and 1505.1 ± 312.8 kcal (girls) in adolescents. WC was 90.9 ± 6.6 cm (boys) and 80.7 ± 10.5 cm (girls) in children, and 97.5 ± 7.9 cm (boys) and 94.1 ± 9.5 cm (girls) in adolescents. VFA was 75.2 ± 26.0 cm² (boys) and 61.9 ± 15.2 cm² (girls) in children, and 87.6 ± 30.3 cm² (boys) and 89.4 ± 42.5 cm² (girls) in adolescents. BMI was 27.1 ± 3.4 (kg/m²) (boys) and 26.1 ± 5.8 (kg/m²) (girls) in children, and 31.4 ± 4.5 (kg/m²) (boys) and 32.4 ± 7.7 (kg/m²) (girls) in adolescents. Percentage body fat (%BF) was 38.6 ± 4.7% (boys) and 37.0 ± 4.9% (girls) in children, and 33.1 ± 5.3% (boys) and 42.3 ± 7.7% (girls) in adolescents. Subcutaneous fat area (SCFA) was 213.6 ± 67.0 cm² (boys) and 197.4 ± 100.8 cm² (girls) in children, and 272.9 ± 84.6 cm² (boys) and 336.1 ± 144.5 cm² (girls) in adolescents. The

Table 3. Correlation between Metabolic Syndrome Score and Obesity Related Factors

	Children (n = 42)				Adolescents (n = 40)			
	MS Score				MS Score			
	0 - 1 (n = 2)	2 - 3 (n = 29)	4 - 5 (n = 11)	p value	0 - 1 (n = 7)	2 - 3 (n = 24)	4 - 5 (n = 9)	p value
Fasting glucose (mg/dL)	97.5 ± 14.8	93.6 ± 8.0	99.3 ± 11.4	0.224	91.1 ± 5.1	97.4 ± 13.2	107.2 ± 21.0	0.086
Triglyceride (mg/dL)	93.0 ± 4.2	134.3 ± 73.9	175.3 ± 71.2	0.181	94.6 ± 36.7	138.1 ± 48.9	189.9 ± 59.2	0.002*
HDL-C (mg/dL)	35.0 ± 5.7	42.2 ± 9.6	53.5 ± 9.8	0.003*	54.0 ± 9.6	40.9 ± 6.7	36.6 ± 4.3	0.000*
Systolic BP (mmHg)	126.0 ± 12.7	118.0 ± 10.6	121.7 ± 14.2	0.477	120.7 ± 12.3	131.0 ± 15.2	143.8 ± 5.3	0.005*
Diastolic BP (mmHg)	63.5 ± 2.1	72.1 ± 8.8	73.5 ± 10.1	0.369	71.9 ± 8.4	77.0 ± 8.0	82.9 ± 9.0	0.038*
BMI (kg/m ²)	24.3 ± 0.8	27.0 ± 4.8	26.6 ± 3.5	0.698	27.2 ± 4.2	31.8 ± 5.2	36.2 ± 9.2	0.022*
%Body fat	36.2 ± 3.2	38.0 ± 4.5	38.6 ± 5.8	0.806	34.5 ± 5.8	38.8 ± 7.9	42.5 ± 9.5	0.151
BMR (kcal)	1202.5 ± 64.3	1227.7 ± 256.8	1173.1 ± 189.0	0.810	1466.9 ± 344.3	1653.1 ± 384.3	1745.6 ± 493.7	0.392
Waist circumference (cm)	79.7 ± 2.3	87.0 ± 10.1	89.4 ± 8.1	0.408	86.3 ± 8.4	96.6 ± 8.4	99.0 ± 7.1	0.008*
Visceral fat area (cm ²)	85.4 ± 43.1	69.9 ± 23.6	69.1 ± 21.4	0.659	70.3 ± 17.9	81.9 ± 28.1	121.4 ± 54.1	0.008*
Subcutaneous fat area (cm ²)	162.9 ± 16.9	208.9 ± 78.2	213.2 ± 92.6	0.719	228.6 ± 91.6	322.9 ± 132.5	349.8 ± 122.7	0.140
Total abdominal fat area (cm ²)	248.3 ± 59.8	283.8 ± 93.8	282.4 ± 108.8	0.883	314.9 ± 113.3	414.8 ± 167.6	503.2 ± 207.5	0.102

MS, metabolic syndrome; BMI, body mass index; BP, blood pressure; HDL-C, high density lipoprotein cholesterol; BMR, basal metabolic rate.

*The mean difference is significant at the 0.05 level.

total abdominal fat area (TAFE) was 292.2 ± 86.4 cm² (boys) and 263.0 ± 109.7 cm² (girls) in children, and 376.0 ± 84.6 cm² (boys) and 441.9 ± 198.8 cm² (girls) in adolescents (Table 2).

Comparisons between groups by MS risk factors

In our study we classified the 82 subjects according to the NCEP ATP III criteria and the scores for MS were classified using 0-1, 2-3 and 4-5.

1) FBG, %BF, BMR, SCFA and TAFE: these factors did not show statistically significant differences between each group in children and adolescents ($p < 0.05$).

2) Blood triglyceride, systolic and diastolic BP, BMI, WC and VFA: these factors did not show statistically significant differences between groups in children, but did show statistically significant differences in adolescents ($p < 0.05$).

3) Blood HDL-C: this factor showed statistically significant differences between groups in both children and adolescents.

Correlations between VFA, BMI by metabolic risk factor

1) BMI: At a p value of 0.01, there were statistically significant differences in systolic and diastolic BP, %BF, BMR, WC, VFA, SCFA and TAFE. At a p value of 0.05, there were statistically significant differences in HDL-C.

2) VFA: At the p value of 0.01 there were a statistically significant differences in fasting glucose, systolic and diastolic BP, BMI, %BF, BMR, WC, SCFA and TAFE. At the p value of 0.05, there were statistically significant differences in HDL-C.

DISCUSSION

It is known that obesity in children and adolescents is related to high prevalence of metabolic complications.¹⁹⁻²¹ In 2004 Chang, et al.²² reported that the prevalence of MS in children and adolescents was 37.5% (boys: 38.7%, girls: 35.2%) by the corrected criteria of NCEP ATP III. Considering that the prevalence of obesity in children and adolescents in the United States is 28.7%, we believe the time, method and criteria of the study is critical in Korea.^{23,24}

There have not been many studies concerning the

Table 4. Correlation between Body Mass Index, Visceral Fat Area and Obesity Related Factors

	Mean \pm SD	BMI		VFA	
		Pearson correlation	<i>p</i> value	Pearson correlation	<i>p</i> value
Fasting glucose (mg/dL)	96.9 \pm 12.5	- 0.075	0.501	0.310 [†]	0.005
Triglyceride (mg/dL)	142.6 \pm 65.8	0.111	0.322	0.187	0.093
HDL-C (mg/dL)	43.6 \pm 9.9	- 0.230*	0.038	- 0.276*	0.012
Systolic BP (mmHg)	125.5 \pm 14.7	0.442 [†]	0.000	0.335 [†]	0.002
Diastolic BP (mmHg)	74.7 \pm 9.3	0.329 [†]	0.003	0.361 [†]	0.001
BMI (kg/m ²)	29.3 \pm 6.1	1	-	0.618 [†]	0.000
%Body fat	38.4 \pm 6.6	0.536 [†]	0.000	0.372 [†]	0.001
BMR (kcal)	1421.5 \pm 390.9	0.616 [†]	0.000	0.516 [†]	0.000
Waist circumference (cm)	91.2 \pm 10.1	0.616 [†]	0.000	0.471 [†]	0.000
Visceral fat area (cm ²)	79.4 \pm 32.6	0.618 [†]	0.000	1	-
Subcutaneous fat area (cm ²)	258.9 \pm 117.8	0.736 [†]	0.000	0.574 [†]	0.000
Total abdominal fat area (cm ²)	347.8 \pm 155.3	0.760 [†]	0.000	0.722 [†]	0.000

BMI, body mass index; VFA, visceral fat area; HDL-C, high density lipoprotein cholesterol; BP, blood pressure; BMR, basal metabolic rate.

*Correlation is significant at the 0.05 level.

[†]Correlation is significant at the 0.01 level.

relationship between VF and MS in children and adolescents. In 2008, Kim and Park²⁵ reported in one study that there is a relationship between abdominal fat and cardiovascular risk factors and metabolic risk factors in obese Korean children and adolescents. VF was found to be independently associated with BP, blood triglyceride, blood HDL-C, FBG and homeostasis model assessment (HOMA) score. Also, Druet, et al.²⁶ reported that VF was independently related to BP, blood triglyceride and blood HDL-C in obese children and adolescents. Thus, VF has been identified as a severe risk factor for MS.

In our study of 82 subjects, we first classified them as children or adolescents and then classified the risk factor scores in groups 0-1, 2-3 and 4-5 according to NCEP ATP III criteria. In children, only HDL-C showed statistically significant differences between three groups, while in adolescents, blood triglyceride, HDL-C, systolic and diastolic BP, BMI, WC and VFA all showed statistically significant differences. MS score elevation in children and adolescents indicates significant increases in the risk factors described above. The severity of MS is related to metabolic complications, including cardiovascular complications. Our study had the same results as other studies. In Korean studies, the risk of cardiovascular disease was higher when more than one metabolic risk factor was involved.²⁷ In our study, we compared the associations between VF (measured by VFCT) and BMI between risk factors for MS by groups. VFA was statistically significant in relation to fasting glucose, systolic and diastolic BP, BMI, %BF, BMR, WC, SCFA, TAFA

and HDL-C, while BMI was related to systolic and diastolic BP, %BF, BMR, WC, VFA, SCFA, TAFA and HDL-C. The aforementioned associations between VF and risk factors for MS suggest the need for abdominal fat measurement.²⁸ But there are some groups who oppose VFCT due to radioactive exposure.

Our study had two important limitations. First of all, diagnosis of MS in children and adolescents is not standardized, because children and adolescents continue to grow, making it difficult to set a normal value and to have long-term follow-up. Also, there is little interest in MS in children and adolescents worldwide, including in the Republic of Korea. Second, we enrolled only a total of 82 subjects because it was difficult to find subjects due to fear of radioactive exposure from VFCT. Safety and verification of radioactive exposure should be studied.

According to AHA, as the prevalence of MS increases in children and adolescents, severe metabolic complications also increase. In clinics for children and adolescents, screening tests for BP, serum cholesterol, serum glucose and abdominal circumference should be undertaken. Thus, early detection and follow-up of risk factors for MS can be possible. There is a strong need to set a cut-off value for diagnostic criteria of MS risk factors in children and adolescents.

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