

## Original Article

# A comparison of the prevalence of the metabolic syndrome in the United States (US) and Korea in young adults aged 20 to 39 years

Jinkyung Park PhD RD<sup>1</sup>, Jason A Mendoza MD MPH<sup>1,3</sup>, Carol E O'Neil PhD MPH RD LDN<sup>2</sup>, David C Hilmers MD EE MPH<sup>3</sup>, Yan Liu MS<sup>1</sup>, Theresa A Nicklas DrPH MPH<sup>1</sup>

<sup>1</sup>Children's Nutrition Research Center, Department of Pediatrics, Baylor College of Medicine, Houston, TX, USA

<sup>2</sup>Louisiana State University Ag Center, LSU, Baton Rouge, LA, USA

<sup>3</sup>Section of Academic General Pediatrics, Department of Pediatrics, Baylor College of Medicine, Houston, TX, USA

This study estimated and compared the prevalence of the Metabolic Syndrome and its individual components in young adults (ages 20-39 years) in the US and Korea using 2003-2004 US and 2005 Korean National Health and Nutrition Examination Survey data. The mean body mass index and rate of metabolic abnormalities in the US were significantly higher than in Korea. The prevalence of the Metabolic Syndrome in the US was nearly three times higher than in Korea using National Cholesterol Education Program-Adult Treatment Panel III and International Diabetes Federation criteria (21.6% vs. 6.9% and 23.0% vs. 6.9%,  $p < 0.001$ ). The prevalence of abdominal obesity, hyperglycemia, and hypertriglyceridemia was higher in the US while the prevalence of low high density lipoprotein-cholesterol level was higher in Korea. The rate of hypertension showed no significant difference while mean systolic blood pressure and diastolic blood pressure varied between the two countries. The proportion of subjects having at least one component of Metabolic Syndrome was similar in both countries; however, multiple abnormalities were more common in the US. These findings indicate the need for the development of race/ethnic-based norms for components of the Metabolic Syndrome and detailed analysis of the risk factors for the Metabolic Syndrome in the two countries. National health policies designed to prevent the Metabolic Syndrome, its individual abnormalities, and its complications using population-based characteristics of each nation will generate improved outcomes.

**Key Words:** National Health and Nutrition Examination Survey, the US, Korea, National Cholesterol Education Program-Adult Treatment Panel III, International Diabetes Federation, metabolic syndrome

## INTRODUCTION

The Metabolic Syndrome (MetS) is characterized by abdominal obesity, insulin resistance, hyperglycemia, hypertension, and dyslipidemia, and is a major risk factor for cardiovascular disease (CVD) and type 2 diabetes.<sup>1-4</sup> This syndrome has been recognized as a serious health problem principally in Western countries during the decade since Reaven<sup>5</sup> coined the term Syndrome X in 1988 to describe the clustering of those risk factors. However, MetS has become common in Asian countries as well.<sup>6</sup> Numerous articles have reported the prevalence of the MetS<sup>4,7-13</sup> in individual nations although few cross-sectional comparisons have been made, especially between Western and Asian countries.

Prevention, early screening, and early intervention of the MetS are recognized to be important in decreasing the morbidity and mortality associated with CVD, diabetes, and their complications.<sup>2,14,15</sup> Young adults with the MetS are especially at increased risk for CVD and type 2 diabetes.<sup>1,7,8,16-21</sup> Behavioral risk factors for the MetS and CVD,<sup>22-24</sup> such as tobacco use,<sup>25</sup> alcohol consumption, and low levels of physical activity,<sup>26-28</sup> often start during

early adulthood under the influence of social and cultural pressures. Such behaviors can contribute to the risk of the MetS and CVD<sup>14,17,18,29-31</sup> and may continue through late adulthood.<sup>19, 32-34</sup> Thus, understanding the prevalence of the MetS in young adults is essential to the formulation of strategies to prevent and treat the MetS.

Since the World Health Organization initially proposed a definition for the MetS in 1998,<sup>35</sup> several other organizations have promulgated alternative criteria.<sup>36-39</sup> The National Education Cholesterol Program Adult Treatment Panel III (NCEP-ATP III) definition, published in 2002<sup>37</sup> and partially modified in 2004<sup>40</sup>, has become most widely used worldwide because of its convenience in clinical

**Corresponding Author:** Dr. Theresa A. Nicklas, Children's Nutrition Research Center, Department of Pediatrics, Baylor College of Medicine, 1100 Bates Avenue, Houston, TX 77030-2600 USA

Tel: 713-798-7087; Fax: 713-798-7098

Email: tnicklas@bcm.tmc.edu

Manuscript received 13 March 2008. Initial review completed 24 July 2008. Revision accepted 11 August 2008.

applications.<sup>40</sup> A newer definition, proposed by the International Diabetes Federation (IDF), emphasized the importance of central adiposity as determined by race/ethnic group-specific thresholds of waist circumference.<sup>8,41</sup> Although little is known about the prevalence of central adiposity and MetS in Asian populations using the IDF criteria, the Korean Society for the Study of Obesity (KSSO)<sup>42</sup> utilizes Korean-specific waist circumference standards for IDF-defined abdominal obesity.

Representative population-based surveys are important resources for establishing national health policies. Data from the US National Health and Nutrition Examination Survey (NHANES) have been used to determine the prevalence of the MetS among three ethnicities: non-Hispanic Whites, non-Hispanic Blacks, and Mexican Americans.<sup>8,9</sup> Asians, however, were classified as 'other race' in the survey. Korea conducted a similar national survey called the Korean National Health and Nutrition Examination Survey (KNHANES). To date, few studies have compared the prevalence of the MetS in the US to Asian countries and no studies have specifically compared the prevalence in the US to that in Korea. We sought to fill this gap in order to provide insight into underlying disease mechanisms and to help guide national health strategies for the prevention and treatment of the MetS. The objectives of this study were 1) to estimate the prevalence of the MetS and its individual components using NCEP-ATP III and IDF criteria among young adult populations in the US and Korean NHANES data sets, and 2) to compare the results between the two countries.

## MATERIALS AND METHODS

### *Study population and data sources*

This study was based on data from the 2003-2004 US NHANES and 2005 KNHANES. Both were cross-sectional nationally representative surveys of the US and Korean civilian non-institutionalized populations, and used standardized protocols for all interviews and medical examinations. Subjects selected from the US NHANES were Mexican-Americans, non-Hispanic Whites, and non-Hispanic Blacks and from the KNHANES were Koreans. Because of the limited sample available in the US NHANES for subjects designated as "other race-including multiracial" and "other Hispanic," these subgroups were excluded. The subjects from both data sets were limited to men and non-pregnant women aged 20 to 39 years who participated in the medical examination of each NHANES and who had fasted at least 8 hours prior to the blood collection. Only subjects who had all five indicators of the MetS measured according to NCEP-ATP III and IDF criteria were included in the study. Accordingly, representative samples of 481 (264 male) and 1,792 (745 male) subjects from the US and Korean NHANES, respectively, were collected, and the estimated US and Korean population represented by each sample was 63.1 million and 15.3 million, respectively.

US and Korean NHANES methodologies have been described in detail previously.<sup>43-44</sup> Briefly, socio-demographic data were collected by personal interview. Anthropometric indices and blood pressure were collected by a standard protocol, and the mean of the 2nd and 3rd systolic and diastolic blood pressure (BP) read-

ings were used for this study. For both surveys, fasting blood glucose (FBG) was measured indirectly by spectrophotometry after an enzymatic reaction, and triglycerides (TG) were indirectly determined by spectrophotometry after hydrolyzation to glycerol and two stages enzymatic reactions. High density lipoprotein cholesterol (HDL-C) was indirectly measured by spectrophotometry after precipitation with sulfated alpha-cyclodextrin and a two stage enzymatic reaction. Low density lipoprotein cholesterol (LDL-C) was calculated using the Friedewald equation (serum total cholesterol - HDL-C - serum TG/5).<sup>45</sup>

### *Definition of the MetS*

The revised NCEP-ATP III<sup>40</sup> and IDF<sup>39</sup> definitions were used to identify individuals with MetS. For the NCEP-ATP III criteria, subjects who had three or more of the following criteria were defined as having the MetS: 1) abdominal obesity: waist circumference > 102 cm in men and > 88 cm in women; 2) hypertriglyceridemia: TG  $\geq$ 150 mg/dl; 3) low HDL-C: HDL-C <40 mg/dl in men and <50mg/dl in women; 4) high BP: SBP/DBP  $\geq$ 130/85 mmHg or taking antihypertensive medication; and 5) high fasting plasma glucose: FBG  $\geq$ 100 mg/dl or taking antidiabetic medication (insulin or oral agents). For the IDF criteria, one must have central adiposity: 1) abdominal obesity: waist circumference  $\geq$  94 cm in non-Hispanic White men;  $\geq$  94 cm in non-Hispanic Black men;  $\geq$  90 in Mexican-American men;  $\geq$  80 cm for all US women; the KSSO thresholds were:  $\geq$  90 cm for Korean men;  $\geq$  85 cm in Korean women; plus two or more of the following factors: 2) hypertriglyceridemia: TG  $\geq$ 150 mg/dl; 3) low HDL-C: HDL-C <40 mg/dl in men and <50mg/dl in women; 4) high BP: SBP/DBP  $\geq$ 130/85 mmHg or treatment of previously diagnosed hypertension; and 5) high fasting plasma glucose: FBG  $\geq$ 100 mg/dl or previously diagnosed type 2 diabetes.

### *Statistical analysis*

Statistical analyses were performed using SAS (version 9.1.3, SAS Institute, Inc., Cary NC, USA) and SAS-callable SUDAAN (version 9.0.3, Research Triangle Institute, Research Triangle Park, NC, USA). A four-stage stratified systematic sampling method was used for sampling in the US and Korean NHANES. For the Korean NHANES, a population-based random sample covering 39,060 persons in 12,283 households across 600 national districts was selected. Data in all statistical analyses for our study were weighted to account for the complex sampling design of US and Korean NHANES which were multistage, stratified, unequally weighted, or clustered. Appropriate statistical sampling weights from each national dataset were selected as specified by each respective survey. Estimated frequencies of categorical variables and mean and standard error of continuous variables were calculated by the PROC DESCRIPT and PROC CROSSTAB procedures of SUDAAN with each national sample weight. Finally, data from the two countries were merged into a single file by creating a new variable that identified the original data set and treated this variable as stratification. T-tests with the NEST statement in the DESCRIPT procedure of SUDAAN were performed to

**TABLE 1-1.** Weighted average values of the factors used to diagnose the metabolic syndrome: the US NHANES 2003-04, Korea NHANES 2005.

	N <sup>†</sup>	WC <sup>‡</sup>	FBG <sup>§</sup>	TG <sup>¶</sup>	HDL-C <sup>††</sup>	SBP <sup>**</sup>	DBP <sup>§§</sup>
<i>Male, 20-29yrs</i>							
US	139 (15.5m)	94.9±1.1	94.1±1.5	142.1±14.6	48.4±1.3	117.9±0.9	67.3±1.3
Korea	264 (3.7m)	79.7±0.6***	86.3±0.6***	113.9±5.2	44.5±0.7**	114.9±0.7*	76.2±0.7***
<i>Male, 30-39yrs</i>							
US	125 (17.6m)	100.5±1.8	97.4±1.0	145.4±11.4	48.8±1.6	118.3±1.5	74.1±1.9
Korea	481 (4.2m)	83.1±0.4***	90.7±0.5***	158.8±8.3	41.6±0.4***	116.5±0.6	79.2±0.6*
<i>Male, Total</i>							
US	264 (33.1m)	97.9±1.0	95.9±0.9	143.8±8.6	48.7±1.1	118.1±1.1	70.9±0.8
Korea	745 (8.0m)	81.5±0.4***	88.6±0.4***	137.8±5.1	43.0±0.4***	115.7±0.5*	77.8±0.5***
<i>Female, 20-29yrs</i>							
US	98 (12.5m)	89.8±1.3	88.9±0.6	101.9±6.0	55.2±1.8	108.9±0.8	66.4±0.9
Korea	375 (3.5m)	72.0±0.6***	84.4±0.5***	83.8±2.6**	49.4±0.6**	103.9±0.6***	69.1±0.6*
<i>Female, 30-39yrs</i>							
US	119 (17.5m)	94.5±1.9	92.6±1.3	130.5±11.6	57.1±1.8	111.0±1.0	69.4±0.8
Korea	672 (3.9m)	74.9±0.4***	88.8±0.8*	90.1±2.2***	48.4±0.5***	105.8±0.6***	70.8±0.5
<i>Female, Total</i>							
US	217 (29.9m)	92.6±1.1	91.1±0.8	118.6±7.5	56.3±1.4	110.1±0.8	68.1±0.7
Korea	1,047 (7.3m)	73.5±0.4***	86.7±0.5***	87.1±1.7***	48.9±0.4***	104.9±0.4***	70.0±0.4*
<i>Country, Total</i>							
US	481 (63.1m)	95.4±0.7	93.6±0.8	131.9±7.0	52.3±0.9	114.3±0.8	69.6±0.7
Korea	1,792 (15.3m)	77.7±0.3***	87.7±0.4***	113.5±2.9*	45.8±0.3***	110.5±0.4***	74.1±0.4***

Data are mean ± SE.

\*  $p < 0.05$ , \*\*  $p < 0.01$ , \*\*\*  $p < 0.001$  show significant differences between the US and Korea.

<sup>†</sup> N, sample size (estimated population represented in millions).

<sup>‡</sup> WC, Waist Circumference (cm) <sup>§</sup> FBG, Fasting Blood Sugar (mg/dl).

<sup>¶</sup> TG, Triglycerides (mg/dl) <sup>††</sup> HDL-C, High Density Lipoprotein Cholesterol (mg/dl).

<sup>\*\*</sup> SBP, Systolic Blood Pressure (mmHg) <sup>§§</sup> DBP, Diastolic Blood Pressure (mmHg).

compare stratum means and percentages between the two countries. All statistics were presented as mean ± standard error (SE) for continuous variables and frequency percentage (SE) for all categorical measures. A  $p$  value of  $< 0.05$  was considered statistically significant.

## RESULTS

In the US data, non-Hispanic Whites comprised the highest proportion of those surveyed (73.2%); Mexican-Americans and non-Hispanic Blacks constituted 12.9% and 13.9%, respectively (data not shown). Table 1-1 and table 1-2 show BMI and metabolic characteristics of young adults 20-39 years of age from both NHANES. The mean waist circumference of US males and females was significantly higher by 15 and 20 cm, respectively, than those of Koreans (both genders,  $p < 0.001$ ). US subjects had a significantly higher FBG than Koreans subjects ( $p < 0.001$ ). Triglycerides in US females were higher than in Korean females ( $p < 0.001$ ), but there was no significant difference in males. Koreans had significantly lower concentrations of HDL-C than Americans ( $p < 0.001$ ). The average SBP of Americans was higher than that of Koreans overall ( $p < 0.001$ ) except for males aged 30 to 39 years, whereas DBP in both US age groups was significantly lower than Koreans ( $p < 0.05$  to  $p < 0.001$ ).

The mean BMI, LDL-C, TC, and pulse pressure of US subjects was significantly higher than that of Koreans (all,  $p < 0.001$ ).

The estimated prevalence of the MetS and abnormalities of individual components as defined by the IDF and NCEP-ATP III are shown in table 2. The estimated prevalence of the MetS in the US (21.6%) was nearly three times higher than in Korea (6.9%) using NCEP-ATP III criteria ( $p < 0.001$ ) and likewise was more than three fold higher in the US (23.0%) versus Korea (6.9%) using IDF criteria ( $p < 0.001$ ). The difference was more pronounced in females than in males. Americans had a significantly higher prevalence of abdominal obesity than Koreans using both criteria (overall,  $p < 0.001$ ). The prevalence of abdominal obesity using the NCEP-ATP III criteria in US males 30-39 years of age (39.9%) was nearly 30 times higher than in Koreans (1.5%), and the highest rate using the IDF abdominal obesity definition was in the US female 30-39 age group (81.2%). The estimated prevalence of elevated FBG in the US was also higher than in Korea ( $p < 0.001$ ), and the difference was markedly greater among males than females.

Hypertriglyceridemia did not show a significant difference between subjects combining both genders in the US and Korea; Korean males 30-39 years had significantly

**TABLE 1-2.** Weighted average values of other CVD risk factors related to the metabolic syndrome: the US NHANES 2003-04, Korea NHANES 2005.

	BMI <sup>†</sup>	LDL-C <sup>‡</sup>	TC <sup>§</sup>	LDL:HDL	TC:HDL	Pulse pressure <sup>¶</sup>
<i>Male, 20-29yrs</i>						
US	27.0±0.4	102.7±2.4	176.7±3.1	2.2±0.1	3.9±0.1	50.6±1.6
Korea	23.3±0.2***	105.6±1.9	172.9±2.3	2.5±0.1*	4.0±0.1	38.6±0.6***
<i>Male, 30-39yrs</i>						
US	28.8±0.7	120.5±4.4	196.7±4.1	2.6±0.2	4.3±0.2	44.2±1.0
Korea	24.2±0.2***	110.2±2.1*	183.6±1.7**	2.7±0.1	4.6±0.1	37.3±0.4***
<i>Male, Total</i>						
US	28.0±0.3	112.2±2.6	187.3±2.4	2.4±0.1	4.1±0.1	47.2±0.8
Korea	23.8±0.1***	108.1±1.4	178.6±1.4**	2.6±0.0	4.3±0.0	37.9±0.4***
<i>Female, 20-29yrs</i>						
US	26.5±0.3	99.2±2.9	174.8±3.6	2.0±0.1	3.4±0.1	42.5±1.0
Korea	21.6±0.2***	98.0±1.4	164.2±1.7**	2.1±0.0	3.4±0.0	34.8±0.5***
<i>Female, 30-39yrs</i>						
US	29.1±0.9	115.9±3.1	200.9±4.9	2.2±0.1	3.8±0.2	41.6±0.5
Korea	22.6±0.1***	105.4±1.1**	171.8±1.4***	2.3±0.0	3.7±0.0	34.9±0.4***
<i>Female, Total</i>						
US	28.1±0.5	108.9±2.7	190.0±3.8	2.1±0.1	3.6±0.1	42.0±0.5
Korea	22.1±0.1***	101.9±0.9*	168.2±1.2***	2.2±0.0	3.6±0.0	34.9±0.3***
<i>Country, Total</i>						
US	28.0±0.3	110.6±1.8	188.6±2.2	2.3±0.1	3.9±0.1	44.7±0.4
Korea	23.0±0.1***	105.1±0.9**	173.6±1.0***	2.4±0.0	4.0±0.0	36.5±0.3***

Data are mean ± SE.

\*  $p < 0.05$ , \*\*  $p < 0.01$ , \*\*\*  $p < 0.001$  show significant differences between the US and Korea.

<sup>†</sup> BMI, Body Mass Index ( $\text{kg}/\text{m}^2$ ).

<sup>‡</sup> LDL-C, Low Density Lipoprotein-Cholesterol (mg/dl), LDL-C = [total cholesterol level - HDL - (triglyceride level ÷ 5)].

<sup>§</sup> TC, Total Cholesterol (mg/dl). <sup>¶</sup> pulse pressure = SBP-DBP.

higher TG than the US participants ( $p < 0.05$ ), whereas US females had a significantly higher prevalence in both age groups (both,  $p < 0.01$ ). The prevalence of low HDL-C levels in Korea was significantly higher than in the US ( $p < 0.001$ ), and the difference was most apparent in males 30-39 years of age and in females. There was no statistical difference in the prevalence of high BP between subjects combining both genders in the US and Korea. Stratified analyses of BP revealed that there was no difference between males in the two countries while among females aged 30-39 years ( $p < 0.05$ ), Americans had significantly higher rates of hypertension than Koreans.

Using the NCEP-ATP III definition, the proportion of subjects meeting at least one criterion was not different between the two nations; however, there was a difference using the IDF definition. Based on the IDF criteria, the proportion of individuals with more than one metabolic abnormality, which must include at least abdominal obesity, was significantly higher in the US than in Korea overall (64.4% and 14.2%, respectively,  $p < 0.001$ ) (Table 3). US subjects had a higher frequency of two or three component abnormalities than Koreans ( $p < 0.001$ ), and the frequency of Americans with more than four components was also more prevalent in males and females ages 30-39 years ( $p < 0.05$ ) than in Koreans.

## DISCUSSION

This study utilized the 2003-2004 US NHANES and the 2005 KNHANES to assess the prevalence of the MetS and abnormalities in its component measures using two definitions. The results that we obtained differed markedly depending on the definition of the MetS that was applied. This can be attributed to the individual focus of each definition; i.e., ATP-III is concerned with CVD risk factors, and IDF with central obesity. The proportion of non-Hispanic Whites, Mexican Americans, and non-Hispanic Blacks in our US sample was similar to recent US Census figures for the US population as a whole.<sup>46</sup> The prevalence of the MetS using either the NCEP-ATP III or IDF criteria was approximately three times higher among US young adults (21.6% and 23.0%, respectively) than among Koreans (6.9% with both definitions). Using census data, this translates into approximately 14 million Americans and 1 million Korean young adults who meet the definition of MetS. There has been a steady increase in the prevalence of the MetS in the US 20-39 year age group. The current US figures have risen from NHANES III 1988-1994 (male 15.7%, female 10.8%) and NHANES 1999-2000 (male 16.5%, female 19.1%).<sup>32</sup> In comparison to our prevalence estimates of the MetS in young Korean adults (6.9%), a study of Korean workers aged 30 – 60 years old showed a prevalence of 7% and 13% using two versions of the NCEP-ATP III criteria.<sup>47</sup>

**TABLE 2.** Estimated prevalence of metabolic syndrome and individual components by NCEP/ATPIII (2004) and IDF (2005) criteria: the US NHANES 2003-04, Korea NHANES 2005.

	MetS †	Abdominal obesity	Hyperglycemia	Hypertriglyceridemia	Low HDL-C	High BP
<b>NCEP/ATPIII criteria</b>						
<i>Male, 20-29yrs</i>						
US	16.6 (4.4)	25.4 (3.3)	19.1 (4.0)	25.2 (5.4)	23.8 (5.2)	19.1 (4.0)
Korea	4.7 (1.3)*	3.2 (1.4)***	2.8 (1.1)***	20.5 (3.2)	30.1 (3.4)	19.1 (2.7)
<i>Male, 30-39yrs</i>						
US	28.2 (4.6)	39.9 (5.7)	31.4 (2.8)	27.5 (3.0)	23.2 (5.3)	32.9 (5.4)
Korea	14.4 (1.6)**	1.5 (0.6)***	13.3 (1.5)***	37.0 (2.7)*	48.2 (2.4)***	29.0 (2.2)
<i>Male, Total</i>						
US	22.8 (3.3)	33.1 (3.0)	25.6 (2.6)	26.4 (2.6)	23.5 (4.4)	26.4 (3.3)
Korea	9.8 (1.1)***	2.3 (0.8)***	8.4 (1.0)***	29.3 (2.1)	39.7 (2.2)**	24.4 (1.7)
<i>Female, 20-29yrs</i>						
US	18.6 (3.0)	44.2 (3.4)	9.8 (2.1)	18.4 (4.0)	41.2 (6.2)	6.1 (2.6)
Korea	1.2 (0.5)***	5.2 (1.3)***	1.9 (0.9)***	6.7 (1.4)**	57.1 (3.1)*	2.6 (0.9)
<i>Female, 30-39yrs</i>						
US	21.5 (4.5)	60.8 (6.7)	12.3 (3.3)	24.4 (4.1)	36.7 (5.6)	11.9 (2.8)
Korea	5.9 (1.0)**	8.3 (1.3)***	7.3 (1.0)	10.2 (1.3)**	59.1 (2.3)***	6.0 (1.0)*
<i>Female, Total</i>						
US	20.3 (2.8)	53.9 (3.8)	11.2 (2.3)	21.9 (3.5)	38.6 (4.4)	9.5 (1.7)
Korea	3.7 (0.6)***	6.9 (0.9)***	4.8 (0.7)**	8.5 (1.0)***	58.2 (1.9)***	4.4 (0.6)**
<i>Country, Total</i>						
US	21.6 (2.4)	43.0 (1.8)	18.8 (2.1)	24.3 (2.4)	30.6 (2.9)	18.4 (2.1)
Korea	6.9 (0.6)***	4.5 (0.7)***	6.7 (0.6)***	19.4 (1.3)	48.6 (1.5)***	14.8 (1.0)
<b>IDF criteria</b>						
<i>Male, 20-29yrs</i>						
US	17.5 (4.5)	47.3 (4.9)	19.1 (4.0)	25.2 (5.4)	23.8 (5.2)	19.1 (4.1)
Korea	5.3 (1.5)*	11.6 (2.2)***	3.4 (1.2)***	20.5 (3.2)	30.1 (3.4)	19.1 (2.7)
<i>Male, 30-39yrs</i>						
US	32.7 (4.4)	62.5 (6.0)	31.4 (2.8)	27.5 (3.0)	23.2 (5.3)	32.9 (5.4)
Korea	14.6 (1.7)***	21.8 (2.0)***	13.3 (1.5)***	37.0 (2.7)*	48.2 (2.4)***	29.0 (2.2)
<i>Male, Total</i>						
US	25.6 (3.4)	55.4 (3.9)	25.6 (2.6)	26.4 (2.6)	23.5 (4.4)	26.4 (3.3)
Korea	10.2 (1.1)***	17.0 (1.5)***	8.7 (1.0)***	29.3 (2.1)	39.7 (2.2)**	24.4 (1.7)
<i>Female, 20-29yrs</i>						
US	17.8 (3.5)	65.0 (4.0)	9.8 (2.1)	18.4 (4.0)	41.2 (6.2)	6.1 (2.6)
Korea	0.7 (0.4)***	8.8 (1.8)***	2.2 (0.9)**	6.7 (1.4)**	57.1 (3.1)*	2.6 (0.9)
<i>Female, 30-39yrs</i>						
US	21.8 (4.4)	81.2 (3.2)	12.3 (3.3)	24.4 (4.1)	36.7 (5.6)	11.9 (2.8)
Korea	5.5 (1.0)***	13.1 (1.6)***	7.4 (1.0)	10.2 (1.3)**	59.1 (2.3)***	6.0 (1.0)*
<i>Female, Total</i>						
US	20.2 (2.8)	74.4 (2.4)	11.2 (2.3)	21.9 (3.5)	38.6 (4.4)	9.5 (1.7)
Korea	3.2 (0.5)***	11.0 (1.2)***	4.9 (0.7)**	8.5 (1.0)***	58.2 (1.9)***	4.4 (0.6)**
<i>Country Total</i>						
US	23.0 (2.4)	64.4 (2.4)	18.8 (2.1)	24.3 (2.4)	30.6 (2.9)	18.4 (2.1)
Korea	6.9 (0.6)***	14.2 (1.0)***	6.9 (0.6)***	19.4 (1.3)	48.6 (1.5)***	14.8 (1.0)

Data are percent (SE).

\*  $p < 0.05$ , \*\*  $p < 0.01$ , \*\*\*  $p < 0.001$ : Significant difference between the US and Korea.

† MetS, Metabolic Syndrome: NCEP/ATP III definition: participants who had three or more of the following risk determinants: abdominal obesity (WC >102 cm in male and >88 cm in female), hyperglycemia (FBG  $\geq$ 100 mg/dl or the participants who currently reported using antidiabetic medication - insulin or oral agents), hypertriglyceridemia (TG  $\geq$ 150 mg/dl), low HDL-C (HDL-C <40 mg/dl in male and <50 mg/dl in female), high BP (SBP  $\geq$ 130 mmHg or DBP  $\geq$ 85 mmHg or the participants who currently reported using antihypertensive medication). IDF definition: participants having central adiposity (WC: the US male : Mexican-American > 90 cm, non-Hispanic\_White >94 cm, non-Hispanic Black >94 cm; female : all races >80, Korean male  $\geq$ 90cm, Korean female  $\geq$ 85cm) plus two or more of the following four factors : hyperglycemia (FBG  $\geq$ 100 mg/dl or previously diagnosed type 2 diabetes, hypertriglyceridemia (TG  $\geq$ 150 mg/dl), low HDL-C (HDL-C <40 mg/dl in male and <50 mg/dl in female), high BP (SBP  $\geq$ 130 mmHg or DBP  $\geq$ 85 mmHg or treatment of previously diagnosed hypertension)

In our study, the IDF definition led to higher estimates of the prevalence of abdominal obesity than the NCEP-ATP III criteria. This is a result of the lower thresholds for central obesity used as part of the IDF criteria that are based on ethnicity. However, the difference in the prevalence of abdominal obesity did not markedly change the

overall prevalence of the MetS since the other four criteria are defined nearly identically.<sup>48</sup> The IDF definition may provide a more accurate assessment of the prevalence of abdominal obesity in the US than the NCEP-ATP III definition because it uses ethnic-based cut points for waist circumference. There is still disagreement over the

**TABLE 3.** Distribution according to the number of metabolic abnormalities: the US NHANES 2003-04, Korea NHANES 2005

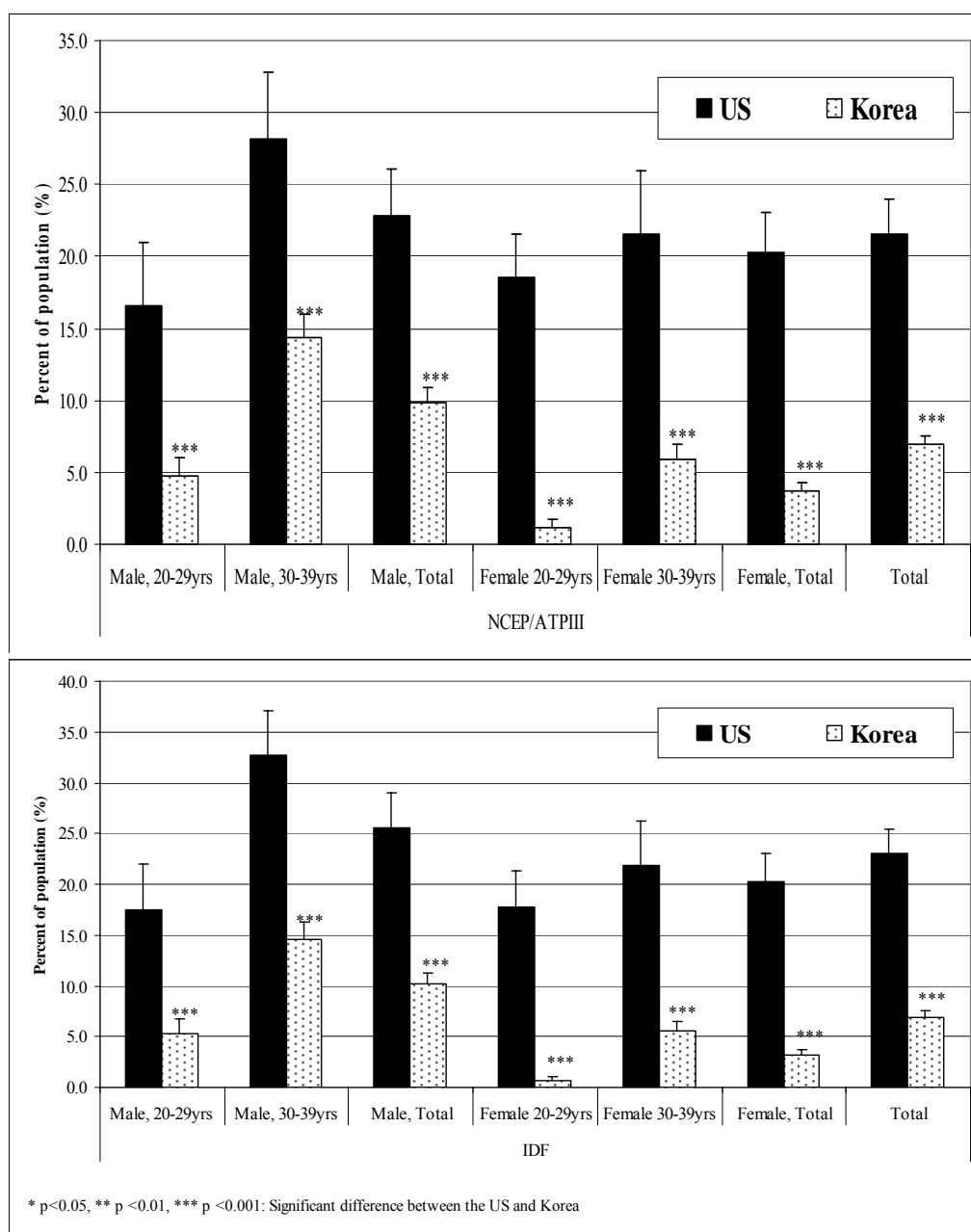
	$\geq 1$	$\geq 2$	$\geq 3$	$\geq 4$	5
<b>NCEP/ATPIII criteria</b>					
<i>Male, 20-29yrs</i>					
US	59.2 (6.4)	33.5 (4.2)	16.6 (4.4)	2.4 (1.4)	0.7 (0.7)
Korea	52.1 (3.8)	18.1 (2.7)**	4.7 (1.3)*	1.0 (0.6)	0.0 (-)
<i>Male, 30-39yrs</i>					
US	71.4 (5.2)	41.4 (4.7)	28.2 (4.6)	11.5 (3.2)	2.2 (1.5)
Korea	71.8 (2.2)	39.2 (2.5)	14.4 (1.6)**	3.7 (0.8)*	0.0 (-)
<i>Male, Total</i>					
US	65.7 (4.2)	37.7 (3.9)	22.8 (3.3)	7.3 (1.8)	1.5 (1.1)
Korea	62.6 (2.3)	29.3 (1.8)	9.8 (1.1)***	2.4 (0.5)*	0.0 (-)
<i>Female, 20-29yrs</i>					
US	63.3 (5.3)	33.6 (4.2)	18.6 (3.0)	4.3 (2.1)	0.3 (0.2)
Korea	61.6 (3.0)	10.5 (1.9)***	1.2 (0.5)***	0.2 (0.2)	0.0 (-)
<i>Female, 30-39yrs</i>					
US	73.5 (5.0)	41.2 (6.5)	21.5 (4.5)	8.9 (3.6)	1.2 (1.2)
Korea	63.9 (2.1)	19.5 (1.8)**	5.9 (1.0)**	1.2 (0.5)*	0.3 (0.2)
<i>Female, Total</i>					
US	69.2 (4.3)	38.0 (3.5)	20.3 (2.8)	7.0 (2.4)	0.7 (0.7)
Korea	62.9 (1.8)	15.3 (1.4)***	3.7 (0.6)***	0.8 (0.3)**	0.2 (0.1)
<i>Country, Total</i>					
US	67.4 (2.6)	37.9 (2.5)	21.6 (2.4)	7.1 (1.2)	1.1 (0.7)
Korea	62.7 (1.4)	22.6 (1.2)***	6.9 (0.6)***	1.6 (0.3)***	0.1 (0.1)
<b>IDF criteria</b>					
<i>Male, 20-29yrs</i>					
US	47.3 (4.9)	33.8 (3.4)	17.5 (4.5)	2.8 (1.5)	0.7 (0.7)
Korea	11.6 (2.2)***	10.3 (2.1)***	5.3 (1.5)*	2.0 (0.9)	0.7 (0.6)
<i>Male, 30-39yrs</i>					
US	62.5 (6.0)	47.4 (4.7)	32.7 (4.4)	13.5 (3.1)	2.2 (1.5)
Korea	21.8 (2.0)***	19.1 (1.8)***	14.6 (1.7)***	5.6 (1.0)*	2.6 (0.6)
<i>Male, Total</i>					
US	55.4 (3.9)	41.0 (2.9)	25.6 (3.4)	8.5 (1.7)	1.5 (1.1)
Korea	17.0 (1.5)***	15.0 (1.4)***	10.2 (1.1)***	4.0 (0.7)*	1.7 (0.4)
<i>Female, 20-29yrs</i>					
US	65.0 (4.0)	39.8 (5.7)	17.8 (3.5)	4.3 (2.1)	0.0 (-)
Korea	8.8 (1.8)***	6.5 (1.5)***	0.7 (0.4)***	0.2 (0.2)	0.0 (-)
<i>Female, 30-39yrs</i>					
US	81.2 (3.2)	47.5 (5.6)	21.8 (4.4)	8.9 (3.6)	1.2 (1.2)
Korea	13.1 (1.6)***	10.4 (1.4)***	5.5 (1.0)***	1.5 (0.5)*	0.3 (0.2)
<i>Female Total</i>					
US	74.4 (2.4)	44.3 (3.8)	20.2 (2.8)	7.0 (2.4)	0.7 (0.7)
Korea	11.0 (1.2)***	8.5 (1.1)***	3.2 (0.5)***	0.9 (0.3)*	0.2 (0.1)
<i>Country, Total</i>					
US	64.4 (2.4)	42.6 (2.6)	23.0 (2.4)	7.8 (1.2)	1.1 (0.7)
Korea	14.2 (1.0)***	11.9 (0.9)***	6.9 (0.6)***	2.5 (0.4)***	1.0 (0.2)

Data are percent (SE).

\*  $p < 0.05$ , \*\*  $p < 0.01$ , \*\*\*  $p < 0.001$  show a significant difference between the US and Korea.

appropriate cut-off point for waist circumference that defines abdominal obesity in Koreans.<sup>49</sup> Some researchers have asserted that the NCEP-ATP III-defined waist circumference criterion is not appropriate for Koreans,<sup>4, 34</sup>

while a recent study declared that the IDF definition is inferior to the NCEP-ATP III criteria for Koreans because a number of Koreans not meeting the IDF definition had more adverse metabolic profiles than others having the



**FIGURE 1.** (Estimated prevalence of metabolic syndrome in the US and Korea), the fonts of figure between NCEP/ATPIII and IDF were different

MetS by both criteria.<sup>50</sup> The DECODA (Diabetes Epidemiology Collaborative Analysis of Diagnostic Criteria in Asia) Study Group indicated that waist circumference may not be the most sensitive measurement of central obesity in Asian populations. The Group also mentioned that while using IDF criteria brought a dramatic rise in the prevalence of central obesity, the overall prevalence of the MetS did not increase greatly compared with the NCEP-ATP III definition.<sup>51</sup> This finding was corroborated in our study.

Mean waist circumference and the prevalence of abdominal obesity were significantly higher in the US than in Korea, and they were most noteworthy in the ages of 30-39 years. Similarly, Patel et al.<sup>52</sup> reported that abnormal waist circumference was considerably more prevalent among individuals with the MetS in the US compared with their Asian counterparts aged 35 years and above. However, Nestel et al.<sup>53</sup> has noted a recent increase in the

prevalence of the metabolic derangements associated with abdominal adiposity in East and Southeast Asia.

Ethnic differences in insulin sensitivity may explain the higher FBG levels and prevalence of hyperglycemia found in the US.<sup>40, 54, 55</sup> However, the DECODA Study Group indicated that lowering the cut-points for obesity and hyperglycemia did not help to identify more subjects who have three or more risk factors.<sup>53</sup> The US has a heterogeneous mixture of ethnicities, and there are many studies demonstrating different rates of insulin resistance and type 2 diabetes between races/ethnicities.<sup>55-59</sup> Additional ethnic-based standards need to be developed to evaluate elevated FBG and to modify the cutoffs appropriately.<sup>41</sup>

An increase in two factors, serum TG and obesity, accounted for much of the increased prevalence of the MetS during young adulthood.<sup>10</sup> Levels of TG's have been shown to be inversely associated with HDL-C,<sup>60</sup> and the

characteristic dyslipidemia with elevated TG and reduced HDL-C levels is regarded as a cardinal sign of insulin resistance.<sup>61</sup> However, subjects from the two countries in this study showed paradoxical lipid profiles, namely, higher TGs in the US and lower HDL-C in Korea. The high prevalence of subnormal HDL-C levels among Asians has been mentioned previously.<sup>62</sup> Decreased HDL-C among Korean adults has been associated with factors such as smoking, obesity, carbohydrate and fat intake in the diet, decreased alcohol consumption and physical activity, low education level, and family history.<sup>63-66</sup> On the other hand, other researchers have insisted that ethnic differences can be explained by genetic factors rather than environmental causes like lifestyle.<sup>60, 67</sup> Their contention that Asians are genetically predisposed to low HDL-C could also help to explain why low HDL-C was more prevalent in Korea than in the US in our study. Overall, there appears to be little difference between the two groups in cardiac risk associated with adverse lipid profiles. Both TC:HDL and LDL:HDL ratios have been used as predictors of future cardiac events,<sup>68</sup> and we saw no significant difference in these two markers between Koreans and Americans. The ethnic variation in dyslipidemias, whether of genetic or environmental origin, is a rich field for further research.

Several interesting findings concerning BP in the two countries were noted in our study. First, when genders were combined, the prevalence of high BP did not significantly differ between the two countries. The prevalence of hypertension in US adults over the last few decades has declined rapidly from 64% (NHANES 1971-1974) to 37% (NHANES 1999-2000),<sup>69</sup> which can partially explain the similar prevalence in the two countries. Secondly, there was no difference in the prevalence of high BP between the two countries despite the three-fold higher prevalence of the MetS in the US. Although BP is a criterion for the MetS, the association between hypertension and "insulin resistance" has been controversial. Some researchers have argued that there is minimal correlation between high BP and the metabolic derangements associated with hyperinsulinemia.<sup>70-73</sup> Finally, we found a higher mean SBP in the US, whereas a higher DBP in Korea, demonstrating a racial/ethnic specific variation in BP patterns. An increased difference in systolic and diastolic pressures (or "pulse pressure") has been implicated in the pathogenesis of atherosclerosis and higher rates of CVD. We found that Americans had higher pulse pressures (44.8 +/- 0.5 vs 36.5 +/- 0.3,  $p < 0.001$ ). This finding implies an earlier onset of atherosclerosis in the US and an increased risk for future coronary events.<sup>74-81</sup>

The difference in components of the MetS, such as lipid profiles and BP, between Asians and other races has not been well studied; however, there have been some reports of differences among non-Asian ethnicities in the United States.<sup>55, 82, 83</sup> Banerjee & Misra<sup>41</sup> suggested that the high BP criterion used in the diagnosis of the MetS should be altered by either changing the cutoffs for different populations or weighting the individual criteria based on their risk contribution to CVD in each population.

The proportion of subjects having at least one component of the MetS was similar in both countries. However,

the difference between subjects in the two countries grew with the number of metabolic abnormalities measured. Although Korean young adults had a lower prevalence of the MetS than Americans, the number of risk factors increased faster with age in Koreans than in the Americans. In particular, 30-39 year-old Korean females had a greater than five times increase in risk factors in comparison with their 20-29 year-old counterparts. Therefore, our study suggests that in-depth analyses by gender in each country relating to lifestyle and other contributors of the MetS over time in young adulthood are needed.

Our study had several limitations. While both surveys used similar laboratory methods, each survey used its own set of reference laboratories to analyze the samples, which may contribute to differences in results between countries. The studies were cross-sectional and therefore causal inferences cannot be made. Finally, our samples may not be representative of each country since not all participants were selected for and/or completed the required fasting phlebotomy.

In conclusion, our study reported several metabolic abnormalities in US and Korean young adults. Young adults in the US had a much higher prevalence of the MetS than in Korea; however, the prevalence of abnormalities in the individual components increases faster with age in Koreans. Most indicators of the MetS, viz. the prevalence of abdominal obesity, hyperglycemia, and hypertriglyceridemia, were higher in US young adults while the prevalence of low HDL-C levels was higher in Koreans. Cardiac risk from adverse lipid profiles did not appear to be significantly different in the two groups. The prevalence of hypertension overall did not differ, but young American adults had higher pulse pressures, which suggests earlier onset and greater prevalence of atherosclerosis. These findings also demonstrated that race/ethnic specific norms for components of the MetS such as waist circumference, FBG, and BP should be developed after these variables are correlated in outcomes-related research. This would guide health policy by quantifying CVD risk, identifying determinants of the syndrome for young adults in both countries, and encouraging young adults to make healthy lifestyle choices. While our study does not elucidate the underlying pathophysiologic origins of the Metabolic Syndrome, comparative studies between ethnic/racial groups may help find commonalities and differences which help to unlock the basis of the syndrome. Our findings also support early screening and intervention in each nation and their implementation into national health policies as the MetS causes an increase in morbidity and mortality, resulting in greatly increased US and Korean health care costs. National health policies designed to prevent the MetS, its individual abnormalities, and its complications using population-based characteristics of each nation will generate improved outcomes.

#### ACKNOWLEDGEMENT

This work (The project No: KRF-2007-357-C00122) was supported by the Korea Research Foundation Grant funded by the Korean Government (MOEHRD). This work is a publication of the United States Department of Agriculture (USDA/ARS) Children's Nutrition Research Center, Department of Pediatrics, Baylor College of Medicine in Houston, Texas and was also



funded in part with federal funds from the USDA/Agricultural Research Service under Cooperative Agreement No. 58-6250-6-003. The contents of this publication do not necessarily reflect the views or policies of the USDA, nor does the mention of trade names, commercial products, or organizations imply endorsement from the US government. Special thanks to Pamela Harris for help in preparing the manuscript.

#### AUTHOR DISCLOSURES

Jinkyung Park, Jason A. Mendoza, Carol E. O'Neil, David C. Hilmers, Yan Liu, and Theresa A. Nicklas, no conflicts of interest.

#### REFERENCES

- Muntner P, He J, Chen J, Fonseca V, Whelton PK. Prevalence of non-traditional cardiovascular disease risk factors among persons with impaired fasting glucose, impaired glucose tolerance, diabetes, and the metabolic syndrome: analysis of the Third National Health and Nutrition Examination Survey (NHANES III). *Ann Epidemiol*. 2004;14:686-95.
- Sarti C, Gallagher J. The metabolic syndrome: prevalence, CHD risk, and treatment. *J Diabet Complications*. 2006;20:121-32.
- Palomo I, Alarcon M, Moore-Carrasco R, Argiles JM. Homeostasis alterations in metabolic syndrome (review). *Int J Mol Med*. 2006;18:969-74.
- Kim MH, Kim MK, Choi BY, Shin YJ. Prevalence of the metabolic syndrome and its association with cardiovascular disease in Korea. *J Korean Med Sci*. 2004;19:195-201.
- Reaven GM. Role of insulin resistance in human disease. *Diabetes*. 1998;37:1595-1607.
- Rakugi H, Ogihara T. The metabolic syndrome in the Asian population. *Curr Hypertens Rep*. 2005;7:103-9.
- Ferreira I, Boreham CA, Twisk JW, Gallagher AM, Young IS, Murray LJ et al. Clustering of metabolic syndrome risk factors and arterial stiffness in young adults: the Northern Ireland Young Hearts Projects. *J Hypertens*. 2007;25:1009-20.
- Ford ES. Prevalence of the metabolic syndrome defined by the international diabetes federation among adults in the U.S. *Diabetes Care*. 2005;28:2745-9.
- Ford ES, Giles WH, Dietz WH. Prevalence of the metabolic syndrome among US adults. *JAMA*. 2002;287:356-9.
- Mattsson N, Ronnema T, Juonala M, Viikari JS, Raitakari OT. The prevalence of the metabolic syndrome in young adults. The Cardiovascular Risk in Young Finns Study. *J Int Med*. 2007;261:159-69.
- Moebus S, Hanisch JU, Aidelburger P, Bramlage P, Wassem J, Jöckel KH. Impact of 4 different definitions used for the assessment of the prevalence of the metabolic syndrome in primary healthcare: The German metabolic and cardiovascular risk project. *Cardiovasc Diabetol*. 2007;6:22.
- Al-Shayji IA, Akanji AO. Obesity indices and major components of metabolic syndrome in young adult Arab subjects. *Ann Nutr Metab*. 2004;48:1-7.
- Yang W, Reynolds K, Gu D, Chen J, He J. A comparison of two proposed definitions for metabolic syndrome in the Chinese adult population. *Am J Med Sci*. 2007;334:184-9.
- Bustos P, da Silva AA, Amigo H, Bettiol H, Barbieri MA. Metabolic syndrome in young adults from two socioeconomic Latin American settings. *Nutr Metab Cardiovasc Dis*. 2007;17:581-9.
- Isomaa B, Almgren P, Tuomi T, Forsen B, Lahti K, Nissen M et al. Cardiovascular morbidity and mortality associated with the metabolic syndrome. *Diabetes Care*. 2001;24:683-9.
- Akosah KO, McHugh VL, Mathiason MA, Kulkarni A, Barnhart SI. Metabolic syndrome and coronary heart disease equivalent conditions in predicting cardiovascular events in young to middle-aged adults. *J Cardiometab Syndr*. 2006;1:173-7.
- Sadeghian S, Darvish S, Salimi S, Esfehiani FA, Fallah N, Mahmoodian M et al. Metabolic syndrome: stronger association with coronary artery disease in young men in comparison with higher prevalence in young women. *Coron Artery Dis*. 2007;18:163-8.
- Lipska K, Sylaja PN, Sarma PS, Thankappan KR, Kutty VR, Vasani RS et al. Risk factors for acute ischemic stroke in young adults in South India. *J Neurol Neurosurg Psychiatry*. 2007;78:959-63.
- Tzou WS, Douglas PS, Srinivasan SR, Bond MG, Tang R, Chen W et al. Increased subclinical atherosclerosis in young adults with metabolic syndrome: the Bogalusa Heart Study. *J Am Coll Cardiol*. 2005;46:457-63.
- Li S, Chen W, Srinivasan SR, Berenson GS. Influence of metabolic syndrome on arterial stiffness and its age-related change in young adults: the Bogalusa Heart Study. *Atherosclerosis*. 2005;180:349-54.
- Bustos P, Amigo H, Arteaga A, Acosta AM, Rona RJ. Risk factors of cardiovascular disease among young adults. *Rev Med Chil*. 2003;131:973-80.
- Misra A. Risk factors for atherosclerosis in young individuals. *J Cardiovasc Risk*. 2000;7:215-29.
- Twisk JW, Kemper HC, van Mechelen W, Post GB. Tracking of risk factors for coronary heart disease over a 14-year period: a comparison between lifestyle and biologic risk factors with data from the Amsterdam Growth and Health Study. *Am J Epidemiol*. 1997;145:888-98.
- Iribarren C, Jacobs DR Jr, Slattery ML, Liu K, Sidney S, Hebert BJ et al. Epidemiology of low total plasma cholesterol concentration among young adults: the CARDIA study. *Coronary Artery Risk Development in Young Adults*. *Prev Med*. 1997;26:495-507.
- Twisk JW, van Lenthe FJ, Kemper HC, van Mechelen W. The longitudinal development of smoking behavior in men and women between 13 and 27 years and the relationship with biological risk factors for cardiovascular disease. *Ned Tijdschr Geneesk*. 1995;139:1790-3.
- Yang X, Telama R, Viikari J, Raitakari OT. Risk of obesity in relation to physical activity tracking from youth to adulthood. *Med Sci Sports Exerc*. 2006;38:919-25.
- Anderssen N, Jacobs DR Jr, Sidney S, Bild DE, Sternfeld B, Slattery ML et al. Change and secular trends in physical activity patterns in young adults: a seven-year longitudinal follow-up in the Coronary Artery Risk Development in Young Adults Study (CARDIA). *Am J Epidemiol*. 1996;143:351-62.
- Kemper HC, Post GB, Twisk JW, van Mechelen W. Lifestyle and obesity in adolescence and young adulthood: results from the Amsterdam Growth and Health Longitudinal Study (AGAHLS). *Int J Obes Relat Metab Disord*. 1999;23 Suppl 3:S34-40.
- Barbieri MA, Bettiol H, Silva AA, Cardoso VC, Simoes VM, Gutierrez MR et al. Health in early adulthood: the contribution of the 1978/79 Ribeirao Preto birth cohort. *Braz J Med Biol Res*. 2006;39:1041-55.
- Twisk JW, Kemper HCG, van Mechelen W, Post GB. Tracking of risk factors for coronary heart disease over a 14-year period: a comparison between lifestyle and biologic risk factors with data from the Amsterdam Growth and Health Study. *Am J Epidemiol*. 1997;145:888-98.

31. Turhan H, Yasar AS, Basar N, Bicer A, Erbay AR, Yetkin E. et al. High prevalence of metabolic syndrome among young women with premature coronary artery disease. *Coron Artery Dis.* 2005;16:37-40.
32. Ford ES, Giles WH, Mokdad AH. Increasing prevalence of the metabolic syndrome among U.S. Adults. *Diabetes Care.* 2004;27:2444-9.
33. Lloyd-Jones DM, Liu K, Colangelo LA, Yan LL, Klein L, Loria CM et al. Consistently stable or decreased body mass index in young adulthood and longitudinal changes in metabolic syndrome components: the Coronary Artery Risk Development in Young Adults Study. *Circulation.* 2007;115:1004-11.
34. Lee WY, Park JS, Noh SY, Rhee EJ, Kim SW, Zimmet PZ. Prevalence of the metabolic syndrome among 40,698 Korean metropolitan subjects. *Diabetes Res Clin Pract.* 2004; 65:143-9.
35. Alberti KG, Zimmet PZ. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: diagnosis and classification of diabetes mellitus provisional report of a WHO consultation. *Diabet Med.* 1998;15:539-53.
36. Balkau B, Charles MA. Comment on the provisional report from the WHO consultation. European Group for the Study of Insulin Resistance (EGIR). *Diabet Med.* 1999;16:442-3.
37. Third Report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) final report. *Circulation.* 2002;106: 3143-3421.
38. Bloomgarden ZT. American Association of Clinical Endocrinologist (AACE) Consensus Conference on the Insulin Resistance Syndrome: 25-26 August 2002, Washington, DC. *Diabetes Care.* 2003;26:1297-303.
39. The IDF consensus worldwide definition of the metabolic syndrome, 2005. (<http://www.idf.org/home>).
40. Grundy SM, Cleeman JI, Daniels SR, Donato KA, Eckel RH, Franklin BA et al. Diagnosis and management of the metabolic syndrome: an American Heart Association/ National Heart, Lung, and Blood Institute Scientific Statement. *Circulation.* 2005;112:2735-52.
41. Banerjee D, Misra A. Does using ethnic specific criteria improve the usefulness of the term metabolic syndrome? Controversies and suggestions. *Int J Obes.* 2007;31:1340-9.
42. Lee SY, Park HS, Kim DJ, Han JH, Kim SM, Cho GJ et al. Appropriate waist circumference cutoff points for central obesity in Korean adults. *Diabetes Res Clin Pract.* (2006) [online publication] DOI:10.1016/j.diabres.2006.04.013.
43. Centers for Disease Control and Prevention: NHANES 2003-2004. (<http://www.cdc.gov/nchs/about/major/nhanes/nhanes03-04.htm>).
44. Korea Centers for Disease Control and Prevention. 2005 Korea National Health and Nutrition Examination Survey: The report of progression of health examination, 2005.
45. Friedewald WT, Levy RI, Fredrickson DS. Estimation of the concentration of low-density lipoprotein cholesterol in plasma, without the use of the preparative ultracentrifuge. *Clin Chem.* 1972;18:499-502.
46. United States Census 2000. (<http://www.census.gov/main/www/cen2000.html>).
47. Shiwaku K, Nogi A, Kitajima K, Anuurad E, Enkhmaa B, Yamasaki M et al. Prevalence of the metabolic syndrome using the modified ATP III definitions for workers in Japan, Korea and Mongolia. *J Occup Health.* 2005;47:126-35.
48. Ford ES, Mokdad AH, Giles WH. Trends in waist circumference among U.S. Adults. *Obes Res.* 2003;11:1223-31.
49. Kim JA, Choi CJ, Yum KS. Cut-off values of visceral fat area and waist circumference: diagnostic criteria for abdominal obesity in a Korean population. *J Korean Med Sci.* 2006;21:1048-53.
50. Yoon YS, Lee ES, Park C, Lee S, Oh SW. The new definition of metabolic syndrome by the international diabetes federation is less likely to identify metabolically abnormal but non-obese individuals than the definition by the revised national cholesterol education program: The Korea NHANES Study. *Int J Obes.* 2007;31:528-34.
51. The DECODA Study Group. Prevalence of the metabolic syndrome in populations of Asian origin Comparison of the IDF definition with the NCEP definition. *Diabetes Res Clin Pract.* 2007;76:57-67.
52. Patel A, Huang KC, Janus ED, Gill T, Neal B, Suriyawongpaisal P et al. Is a single definition of the metabolic syndrome appropriate? - A comparative study of the USA and Asia. *Atherosclerosis.* 2006;184:225-32.
53. Nestel P, Lyu R, Low LP, Sheu WH, Nitiyanant W, Saito I et al. Metabolic syndrome: recent prevalence in East and Southeast Asian populations. *Asia Pac J Clin Nutr.* 2007; 16:362-7.
54. Dickinson S, Colagiuri S, Faramus E, Petocz P, Brand-Miller JC. Postprandial hyperglycemia and reduced insulin sensitivity differ among lean young adults of different ethnicities. *J Nutr.* 2002;132:2578-83.
55. Cossrow N, Falkner B. Race/ethnic issues in obesity and obesity-related comorbidities. *J Clin Endocrinol Metab.* 2004;89:2590-4.
56. Pereira MA, Jacobs Jr DR, Van Horn L, Slattery ML, Kartashov AI, Ludwig DS. Dairy consumption, obesity, and the insulin resistance syndrome in young adults: the CARDIA Study. *JAMA.* 2002;287:2081-9.
57. Harris MI, Flegal KM, Cowie CC, Eberhardt MS, Goldstein DE, Little RR et al. Prevalence of diabetes, impaired fasting glucose, and impaired glucose tolerance in U.S. adults. *Diabetes Care.* 1998;21:518-24.
58. Brancati FL, Kao WHL, Folsom AR, Watson RL, Szklo M. Incident type 2 diabetes mellitus in African American and white adults. The Atherosclerosis Risk in Communities Study. *JAMA.* 2000;283:2253-9.
59. Park YW, Zhu S, Palaniappan L, Heshka S, Carnethon MR, Heymsfield SB. 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.
60. Tai ES, Emmanuel SC, Chew SK, Tan BY, Tan CE. Isolated low HDL cholesterol: an insulin-resistant state only in the presence of fasting hypertriglyceridemia. *Diabetes.* 1999;48:1088-92.
61. Knudsen P, Eriksson J, Lahdenperä S, Kahri J, Groop L, Taskinen MR. Changes of lipolytic enzymes cluster with insulin resistance syndrome. Botnia Study Group. *Diabetologia.* 1995;38:344-50.
62. Heng D, Ma S, Lee J J. M., Tai BC, Mak KH, Hughes K et al. Modification of the NCEP ATP III definitions of the metabolic syndrome for use in Asians identifies individuals at risk of ischemic heart disease. *Atherosclerosis.* 2006;186: 367-73.
63. Kim SM, Han JH, Park HS. Prevalence of low HDL-cholesterol levels and associated factors among Koreans. *Circ J.* 2006;70:820-6.
64. Moon SS, Lee HG, Yoon YS, Sunwoo S, Park HS. Factors associated low HDL-cholesterol in adults. *J Korean Acad Fam Med.* 2001;22:1214-23.
65. Yoon YS, Oh SW, Baik HW, Park HS, Kim WY. Alcohol consumption and the metabolic syndrome in Korean adults:

- the 1998 Korean National Health and Nutrition Examination Survey. *Am J Clin Nutr.* 2004;80:217-24.
66. Lee WY, Jung CH, Park JS, Rhee EJ, Kim SW. Effects of smoking, alcohol, exercise, education, and family history on the metabolic syndrome as defined by the ATP III. *Diabet Res Clin Pract.* 2005;67:70-7.
67. Bhalodkar NC, Blum S, Rana T, Bhalodkar A, Kitchappa R, Kim KS et al. Comparison of levels of large and small high-density lipoprotein cholesterol in Asian Indian men compared with Caucasian men in the Framingham offspring study. *Am J Cardiol.* 2004;94:1561-3.
68. Hsia SH, Pan D, Berookim P, Lee ML. Population-Based, Cross-Sectional Comparison of Lipid-Related Indexes for Symptoms of Atherosclerotic Disease. *Am J Cardiol.* 2006;98:1047-52.
69. Imperatore G, Cadwell BL, Geiss L, Saadine JB, Williams DE, Ford ES et al. Thirty-year trends in cardiovascular risk factor levels among US adults with diabetes: National Health and Nutrition Examination Surveys, 1971-2000. *Am J Epidemiol.* 2004;160:531-9.
70. Cuspidi C, Meani S, Valerio C, Sala C, Fusi V, Zanchetti A et al. Age and target organ damage in essential hypertension: role of the metabolic syndrome. *Am J Hypertens.* 2007;20:296-303.
71. Mule G, Nardi E, Cottone S, Cusimano P, Volpe V, Piazza G et al. Influence of metabolic syndrome on hypertension-related target organ damage. *J Int Med.* 2005;257:503-13.
72. 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. *Am J Epidemiol.* 2000;152:908-11.
73. Diabetes Prevention Program Research Group. Hypertension, insulin, and proinsulin in participants with impaired glucose tolerance. *Hypertension.* 2002;40:679-86.
74. Verdecchia P, Angeli F. Does brachial pulse pressure predict coronary events? *Adv Cardiol.* 2007;44:150-9.
75. Jankowski P, Bilo G, Kawecka-Jaszcz k. The pulsatile component of blood pressure—its role in the pathogenesis of atherosclerosis. *Blood Press.* 2007;16:238-45.
76. Uechi Y, Sunagawa O, Ishikawa N, Inoue T, Tamashiro M, Kamiyama T et al. Risk factors for stiffness of the wall of the thoracic aorta in patients with mild atherosclerosis. *Jpn Circ J.* 2001;65:409-13.
77. Franklin SS, Sutton-Tyrrell K, Belle SH, Weber MA, Kuller LH. The importance of pulsatile components of hypertension in predicting carotid stenosis in older adults. *J Hypertens.* 1997;15:1143-50.
78. Zanchetti A, Crepaldi G, Bond MG, Gallus GV, Veglia F, Ventura A et al. Systolic and pulse blood pressures (but not diastolic blood pressure and serum cholesterol) are associated with alterations in carotid intima-media thickness in the moderately hypercholesterolaemic hypertensive patients of the Plaque Hypertension Lipid Lowering Italian Study. PHYLLIS study group. *J Hypertens.* 2001;19:79-88.
79. Zureik M, Touboul PJ, Bonithon-Kopp C, Courbon D, Berr C, Leroux C et al. Cross-sectional and 4-year longitudinal associations between brachial pulse pressure and common carotid intima-media thickness in a general population. The EVA study. *Stroke.* 1999;30:550-5.
80. Vos LE, Oren A, Uiterwaal C, Gorissen WH, Grobbee DE, Bots ML. Adolescent blood pressure and blood pressure tracking into young adulthood are related to subclinical atherosclerosis: the Atherosclerosis Risk in Young Adults (ARYA) study. *Am J Hypertens.* 2003;16:549-55.
81. Nair GV, Waters D, Rogers W, Kowalchuk GJ, Stuckey TD, Herrington DM. Pulse pressure and coronary atherosclerosis progression in postmenopausal women. *Hypertension.* 2005;45:53-7.
82. Hall WD, Clark LT, Wenger NK, Wright JT Jr, Kumanyika SK, Watson K et al. The metabolic syndrome in African Americans: a review. *Ethn Dis.* 2003;13:414-28.
83. Burt VL, Whelton P, Roccella EJ, Brown C, Cutler JA, Higgins M et al. Prevalence of hypertension in the US adult population. Results from the Third National Health and Nutrition Examination Survey, 1988-1991. *Hypertension.* 1995;25:305-13.

## Original Article

## A comparison of the prevalence of the metabolic syndrome in the United States (US) and Korea in young adults aged 20 to 39 years

Jinkyung Park PhD RD<sup>1</sup>, Jason A Mendoza MD MPH<sup>1,3</sup>, Carol E O'Neil PhD MPH RD LDN<sup>2</sup>, David C Hilmers MD EE MPH<sup>3</sup>, Yan Liu MS<sup>1</sup>, Theresa A. Nicklas DrPH MPH<sup>1</sup>

<sup>1</sup>Children's Nutrition Research Center, Department of Pediatrics, Baylor College of Medicine, Houston, TX, USA

<sup>2</sup>Louisiana State University Ag Center, LSU, Baton Rouge, LA, USA

<sup>3</sup>Section of Academic General Pediatrics, Department of Pediatrics, Baylor College of Medicine, Houston, TX, USA

### 美國與韓國年輕成人(20-39 歲)代謝綜合症盛行率之比較

本研究采用 2003-2004 年美国 和 2005 年韩国健康和营养调查的数据来对年轻成年人(20-39 岁)中新陈代谢综合症及其个别成分的流行性作了评估和比较。美国年轻成人的平均身体质量指数和代谢异常盛行率比韩国人显著的高。采用美国胆固醇教育计划-ATPIII 和国际糖尿病联盟的标准来衡量,美国人代谢综合症盛行率比韩国人约高三倍(21.6%比 6.9%及 23.0%比 6.9%,  $p < 0.001$ )。腹部肥胖,高血糖和高甘油三酯症的盛行率在美国比较高,而低的高密度脂蛋白胆固醇的比例在韩国比较高。高血压盛行率并没有显著的差别,但两国年轻成人的平均血管收缩压和舒张压都有差异。具有至少一个代谢综合症成分的对象的比例在这两个国家是相似的,但是有多重异常性的人在美国更常见。这些发现表明:有必要开发研究基于种族/族群的代谢综合症的成分的标准,以及详细分析代谢综合症在这两个国家的危险因子。制定以每个国家的人口特征为基础的,用以预防代谢综合症以及个别异常症和并发症的国家健康政策,将产生改善效果。

**关键字:** 国家健康和营养调查、美国、韩国、美国胆固醇教育计划-ATPIII、国际糖尿病联盟、新陈代谢综合症