

Original Article



The Association between Food Group Consumption Patterns and Early Metabolic Syndrome Risk in Non-Diabetic Healthy People

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Conflict of Interest

The authors declare that they have no
competing interests.

ABSTRACT

We investigated the association between dietary habits/food group consumption patterns and early risk of metabolic syndrome (MetS), a main cause for metabolic disease. Study participants were recruited from the health promotion center in Dong-A University Hospital and public advertisement. Study subjects (n = 243, 21–80 years) were categorized into three groups: Super-healthy (MetS risk factor [MetS RF] = 0, n = 111), MetS-risk carriers (MetS RF = 1–2, n = 96), and MetS (MetS RF ≥ 3, n = 27). Higher regularity in dietary habits (breakfast-everyday, regular eating time, non-frequent overeating, and non-frequent eating-out) was observed in the Super-healthy group than in the MetS-risk carriers, and particularly in the MetS subjects. The relationship between food group consumption patterns and MetS-risk related parameters were investigated with adjustment for confounding factors. Fruit consumption was positively associated with HDL-cholesterol, and tended to be negatively associated with waist circumference, triglyceride, LDL-cholesterol, and insulin resistance (IR). The consumption of low-fat meats and fish, and vegetables was negatively associated with hs-CRP. Specifically, the consumption of sea-foods belonging to the low-fat fish was negatively associated with fasting glucose, hs-CRP, and interleukin (IL)-6. Anchovy/dried white baits consumption was negatively associated with fasting insulin and IR. Green-yellow vegetables consumption was negatively associated with fasting insulin, IR, and hs-CRP. On the other hand, sugars and fast-foods were positively associated with LDL-cholesterol. Additionally, fast-foods consumption was positively associated with hs-CRP and IL-6 levels. In conclusion, dietary habits/food group consumption patterns are closely associated with MetS-risk related parameters in Koreans. It may suggest useful information to educate people to properly select healthy foods for early prevention of MetS.

Keywords: Dietary habit; Food group consumption pattern; Metabolic syndrome; Inflammation

INTRODUCTION

Metabolic syndrome (MetS) is the name for a group of metabolic disorders including hyperglycemia, hypertension, dyslipidemia, and obesity, particularly abdominal obesity [1]. Uncared metabolic disorders are associated with the increased risk of various chronic diseases related to insulin resistance (IR) such as diabetes, cardiovascular diseases (CVDs), cancer, dementia, and non-alcoholic fatty liver disease [2-6]. The prevalence of MetS has been globally increasing [7-9], and the medical expenses related to the treatment of MetS

have annually increased from 3.7 trillion won in 2010 to 4.7 trillion won in 2014 (the Health Insurance Review and Assessment Service data). Therefore, it is needed to prevent and control metabolic disorders as an important health issue.

Many causes including age, heredity, and life style, potentially affect the risk of MetS [10,11]. Among them, diet is one of the important causes. Many studies have reported the associations between dietary patterns and MetS [12-17]. Westernized diet patterns (i.e., high amounts of saturated fats and simple sugars) have been associated with a higher risk of MetS [12]. In British cohort study, irregular energy intake, particularly from breakfast and between meals was associated with increased cardio-metabolic risk [13]. In addition, rapid eating habit increased the incidence of MetS risk [14]. On the other hand, Mediterranean dietary patterns (i.e., a high consumption of vegetables, fruits, whole cereals, and fish) decreased the risk of MetS [12]. Similarly, the higher meat-eating dietary pattern was associated with a higher prevalence of MetS in Korean male adults in National Cancer Center study [15], higher breakfast skipping was related to an increased risk of obesity and MetS [16], and fast eating behavior was associated with an increased presence risk of non-alcoholic fatty liver disease [17]. However, many studies examining the relationship between dietary patterns and MetS performed in Koreans were implemented with simple survey.

Therefore, this study aimed to investigate if dietary habit and food group consumption patterns are associated with early risk of MetS in non-diabetic healthy Korean adults.

MATERIALS AND METHODS

Study subjects

Study participants (aged over 20 years old) were recruited from the Health Promotion Center in Dong-A University Hospital, and through the public advertisement. Those who have any diagnosis of chronic diseases such as diabetes, CVD, coronary heart disease, stroke, cancer, and any other metabolic diseases were excluded. None of the subjects were taking antihypertensive, lipid-lowering, antidiabetic, or antithrombotic medications. After the screening, a total of 243 Korean adults were finally enrolled in this study. The study was approved by the Institutional Review Board of Dong-A University (2-1040709-AB-N-01-201310-BR-02-04). The informed written consent of all subjects was obtained.

Definition of MetS

MetS was defined by a combination of modified National Cholesterol Education Program-Adult Treatment Panel (NCEP-ATP) III criteria [18], the American Diabetes Association and the Korean society for the study of obesity. MetS risk factors (MetS RFs) include waist circumference (WC) ≥ 90 cm (male) or ≥ 85 cm (female); systolic blood pressure (BP) ≥ 130 mmHg or diastolic BP ≥ 85 mmHg; fasting blood glucose ≥ 100 mg/dL; high-density lipoprotein cholesterol (HDL-C) < 40 mg/dL (male), < 50 mg/dL (female); and triglyceride (TG) ≥ 150 mg/dL. The subjects were divided into 3 groups according to the numbers of MetS RFs: those who have no MetS RFs as 'Super-healthy'; those who have 1 or 2 MetS RFs as 'MetS risk carrier'; those who have 3 or more MetS RFs, as 'MetS.'

Basic information and diet survey

Basic information including demographics, medical history, family history, and physical activity of subjects was collected through a questionnaire interviewed by a registered

dietitian. Diet survey consists of usual dietary habits, a semi-quantitative food frequency questionnaire (FFQ), and a 3-day diet record with 24-hour recall method (2 weekdays and 1 weekend) by face to face interview. FFQ consists of 32 questions based on Korean National Health and Nutrition Survey form, and results were expressed as a sum of weekly food consumption based on 1 serving size. Briefly, 32 questions were composed of grains (cooked rice, noodle, bread/sponge cake/cake/crackers/biscuits, rice cake/potato/corn), meat and fishes (red meat/pork/chicken, low fat fishes, squid/shrimp, crab/shellfish/oyster, anchovy/dried white bait, eggs, medium fat fishes, tofu/black bean, hams, high fat fishes, rib/sausage/eel), vegetables (green-yellow, white, young radish/radish leaves, kimchi/diced radish kimchi, seaweeds), fats (nuts, plant oils, mayonnaise/butter/creams/bacon), milk products (milk, yoghurt, ice cream), fruits (fresh fruit/fresh juice), sugars (sugar/syrup/sweetened drink/candy/chocolate), and others (fast-foods, instant noodle, Korean stews, coffee/tea). Energy intake and nutrient content from 3-day diet records were estimated using the Computer Aided Nutritional analysis program (CAN-pro 4.0; the Korean Nutrition Society, Seoul, Korea).

Anthropometric measurements, BP and blood collection

Anthropometric measurements were taken with subjects wearing light clothes except shoes. Height, body weight, and fat and muscle mass were measured by automatic body composition analyzer (N20; AIIA Communication Inc., Seongnam, Korea) without metallic materials. Body mass index (BMI) was calculated as body weight divided by height in square meters (kg/m^2). WC was measured by measuring tape. BP was measured at the seated subjects' arm after a rest, using an automatic BP monitor (HEM-7220; OMRON, Matsusaka, Japan). Blood samples were collected in plain and ethylenediaminetetraacetic acid (EDTA)-treated tubes in the morning after an 8-hour fast. Then, samples were separated into serum and plasma, respectively using a centrifuge and stored at -80°C before analysis.

Lipid profiles and glycemic parameters

Serum total cholesterol (TC) and TG levels were measured with kits on a Hitachi 7150 auto-analyzer (Hitachi Ltd., Tokyo, Japan). After precipitation of serum chylomicrons with dextran sulfate magnesium, the concentrations of low-density lipoprotein cholesterol (LDL-C) and HDL-C in the supernatants were enzymatically measured. Fasting serum glucose levels were measured using a glucose oxidase method with a Beckman Glucose analyzer (Beckman Instruments, Irvine, CA, USA). Insulin and C-peptide levels were measured by radioimmunoassays with commercial kits (ImmunoNucleo Corporation, Stillwater, MN, USA). IR was calculated with the homeostasis model assessment (HOMA) using the following equation:

$$\text{HOMA-IR} = (\text{fasting insulin } [\mu\text{IU}/\text{mL}] \times \text{fasting glucose } [\text{mmol}/\text{L}]) / 22.5$$

It developed by Matthews et al. [19] and Choi et al. [20]. Hemoglobin A1c (Hb_{A1c}) was measured by a glycosylated hemoglobin analyzer (SD A1cCare™; SD Biosensor Inc., Suwon, Korea).

Inflammatory markers

High-sensitivity C-reactive protein (hs-CRP) was measured by hs-CRP-latex (II) X2 kit (Seiken Laboratories, Tokyo, Japan). Plasma tumor necrosis factor (TNF)- α and interleukin (IL)-6 were analyzed by Quantikine HS ELISA Kits (R & D Systems, Minneapolis, MN, USA). The color reaction results were measured by iMark™ microplate absorbance reader (Bio-Rad Laboratories, Hercules, CA, USA) at 490 nm.

Statistical analysis

The statistical analysis was carried out with SPSS version 23.0 (SPSS Inc., Chicago, IL, USA). To evaluate differences among the groups, 1-way analysis of variance (ANOVA) with post hoc Bonferroni test and general linear model were used. Pearson and partial correlations were used to describe the association among dietary consumption patterns and MetS RFs. Results were expressed as mean \pm standard error, percentages, or correlation co-efficient. A 2-tailed p value under 0.05 was considered significant.

RESULTS

Basic characteristics and biochemical parameters and dietary habits according to MetS risk status

Table 1 presents basic and biochemical characteristics and dietary habits of study subjects according to MetS risk status. As the number of MetS RF increased, people were older, and more obese, particularly had higher WC, and showed dyslipidemia for example, higher levels of triglyceride, LDL-C, TC, and lower levels of HDL-C. Regarding glycemic parameters, fasting levels of glucose, insulin, C-peptide, HOMA-IR, and Hb_{A1C} were increased when the number of MetS RF increased. As the number of MetS RF increased, the proportions of males and current alcohol drinkers were increased, but no significant differences were observed in

Table 1. Basic and biochemical characteristics and dietary habits according to MetS risk status

Parameters	Super-healthy (n = 111)	MetS risk carriers (n = 96)	MetS (n = 27)	p value*
Age, yr	43.20 \pm 1.15	46.80 \pm 1.49	51.60 \pm 2.84	0.010 [†]
BMI, kg/m ²	21.90 \pm 0.20	25.40 \pm 0.31	27.60 \pm 0.55	< 0.001
WC, cm	74.20 \pm 0.62	86.70 \pm 0.92	93.10 \pm 1.74	< 0.001
Systolic BP, mmHg	110.20 \pm 0.84	121.10 \pm 1.50	133.00 \pm 2.55	< 0.001
Diastolic BP, mmHg	70.00 \pm 0.56	77.40 \pm 1.01	82.20 \pm 1.97	< 0.001
Triglyceride, mg/dL [‡]	65.50 \pm 2.48	107.50 \pm 6.06	195.20 \pm 20.60	< 0.001
HDL-C, mg/dL	68.00 \pm 1.17	58.60 \pm 1.37	47.70 \pm 2.68	< 0.001
LDL-C, mg/dL [‡]	114.20 \pm 2.45	125.70 \pm 3.30	130.30 \pm 7.50	0.020
TC, mg/dL	186.80 \pm 2.42	196.90 \pm 3.67	204.90 \pm 8.41	0.022
Glucose, mg/dL [‡]	84.70 \pm 0.68	95.70 \pm 2.03	105.10 \pm 3.60	< 0.001
Hb _{A1C} , % [‡]	5.20 \pm 0.03	5.51 \pm 0.05	5.74 \pm 0.12	< 0.001
Insulin, μ U/mL ^{‡§}	7.10 \pm 1.21	11.90 \pm 1.86	23.60 \pm 5.15	< 0.001
HOMA-IR ^{‡§}	1.51 \pm 0.29	3.28 \pm 0.73	6.33 \pm 1.48	< 0.001
C-peptide, ng/mL [¶]	1.74 \pm 0.22	2.36 \pm 0.22	4.09 \pm 0.68	< 0.001
hs-CRP, mg/dL [‡]	0.59 \pm 0.13	1.52 \pm 0.50	1.79 \pm 0.42	< 0.001
IL-6, pg/mL [‡]	1.11 \pm 0.16	1.31 \pm 0.19	1.70 \pm 0.44	0.015
TNF- α , pg/mL [‡]	0.96 \pm 0.05	1.27 \pm 0.19	1.43 \pm 0.22	0.043
Proportion				
Male	16 (14.4)	33 (34.4)	12 (44.4)	< 0.001
Current smoker	7 (6.3)	11 (11.5)	1 (3.7)	NS
Current drinker	29 (26.1)	40 (41.7)	13 (48.1)	0.027
Dietary habit				
Breakfast everyday	59 (53.2)	51 (53.1)	13 (48.1)	< 0.050
Regular eating time	74 (66.0)	54 (56.2)	11 (40.7)	< 0.001
Frequent overeating	7 (6.3)	8 (8.3)	4 (14.8)	< 0.050
Frequent eating-out	14 (12.6)	16 (16.7)	5 (18.5)	< 0.050
Regular exercise	36 (32.4)	33 (34.4)	9 (33.3)	NS

Data are means \pm SE or number (%).

MetS, metabolic syndrome; BMI, body mass index; WC, waist circumference; BP, blood pressure; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; TC, total cholesterol; Hb_{A1C}, hemoglobin A1C; HOMA-IR, homeostasis model assessment of insulin resistance index; hs-CRP, high-sensitivity C-reactive protein; IL-6, interleukin-6; TNF- α , tumor necrosis factor-alpha; NS, no statistical significance; SE, standard error; ANOVA, analysis of variance.

*Age adjusted p value; [†]Unadjusted p value; tested by 1-way ANOVA (unadjusted) or general linear model methods (age adjusted) with Bonferroni method;

[‡]Tested after log transformed due to the skewed distribution; [§]n = 147; [¶]n = 107.

the proportions of cigarette smoking and regular exercising among the 3 groups. Regarding dietary habit, super-healthy people kept higher regularity in everyday breakfast and regular eating time, but non-frequent overeating and non-frequent eating-out than the MetS risk carriers, and particularly than the MetS people.

Food group consumption per week according to MetS risk status

Table 2 shows the sum total of weekly frequency of food group consumption according to MetS risk status. No significant differences were found in the frequency of food group consumption among the 3 groups. In addition, daily intakes of total calorie and other nutrients were not significant different among the 3 groups (data not shown).

Relationship between food group consumption patterns and MetS risk related parameters

As stated above, food group consumption patterns were not significantly different according to MetS risk status. Therefore, we sub-analyzed the association between food group consumption pattern and each of MetS risk related parameters. **Table 3** shows the relationship between the food group consumption and MetS risk related parameters after adjustment for age, sex, total calorie intake, cigarette smoking, and alcohol consumption which were basic parameters and significantly different among the 3 MetS risk group. The consumptions of low-fat meats and fish ($r = -0.144$, $p = 0.032$), and vegetables total ($r = -0.188$, $p = 0.055$) were negatively associated with hs-CRP levels. Specifically, the consumption of seafoods which belong to the low-fat fish group was negatively associated with the levels of serum glucose ($r = -0.158$, $p = 0.017$), hs-CRP ($r = -0.211$, $p = 0.002$), and IL-6 ($r = -0.155$, $p = 0.021$). The consumption of anchovy and dried white baits was negatively associated with the levels of insulin ($r = -0.186$, $p = 0.027$), HOMA-IR ($r = -0.170$, $p = 0.044$), and tended toward negative association with C-peptide ($r = -0.170$, $p = 0.087$). In the vegetables group, the consumption of green-yellow vegetables was negatively associated the levels of insulin ($r = -0.170$, $p = 0.043$), HOMA-IR ($r = -0.180$, $p = 0.032$), hs-CRP ($r = -0.162$, $p = 0.015$), and tended toward negative association with glucose ($r = -0.114$, $p = 0.086$). In addition, fruits consumption was positively associated with HDL-C ($r = 0.156$, $p = 0.018$), tended to be negatively associated with WC ($r = -0.126$, $p = 0.057$), triglyceride ($r = -0.122$, $p = 0.065$), LDL-C ($r = -0.127$, $p = 0.055$), and HOMA-IR ($r = -0.157$, $p = 0.062$). On the other hand, the consumption of the sugars ($r = 0.157$, $p = 0.018$) and fast-foods ($r = 0.141$, $p = 0.033$) was positively associated with LDL-C levels. The fast-foods consumption was also positively associated inflammation markers such as hs-CRP ($r = 0.156$, $p = 0.020$) and IL-6 levels ($r = 0.132$, $p = 0.050$). However, no significant relationships were observed between milk product consumption and MetS risk related parameters.

Table 2. Patterns of food group consumption per week according to MetS risk status

Food group*	Super-healthy (n = 111)	MetS risk carriers (n = 96)	MetS (n = 27)	p value
Grains	18.30 ± 0.64	18.30 ± 0.62	19.10 ± 1.81	0.855
Meats and fish	24.40 ± 1.66	21.00 ± 1.31	21.00 ± 2.00	0.230
Low-fat	11.10 ± 0.98	9.81 ± 0.84	8.62 ± 1.11	0.342
Medium-fat	9.70 ± 0.88	8.69 ± 0.76	9.30 ± 1.23	0.685
High-fat	3.53 ± 0.39	2.46 ± 0.25	3.10 ± 0.94	0.112
Vegetables	26.60 ± 1.53	27.00 ± 1.83	23.80 ± 2.54	0.670
Milk products	9.54 ± 0.86	7.88 ± 0.66	9.58 ± 1.89	0.309
Fruits	6.42 ± 0.50	5.68 ± 0.48	6.31 ± 0.84	0.544
Fast-foods	1.17 ± 0.12	0.99 ± 0.14	1.07 ± 0.27	0.628

Data are means ± SE; tested by 1-way ANOVA with Bonferroni method.

MetS, metabolic syndrome; SE, standard error; ANOVA, analysis of variance.

*Calculated as sum total of 1-serving sized food consumption in each food group per week.

Comparison of adiposity and inflammation according to MetS risk status and fast-food consumption

As shown in **Table 3**, fast-foods consumption which was highly correlated with MetS and cardiovascular risk (LDL-C, IL-6, and TNF- α) was selected for sub-analysis. We sub-grouped study subjects by fast-food consumption level (the lower 50 percentiles in study subjects “lower intake group”: they consumed fast-foods less than once per week, vs. the upper 50 percentiles in study subjects “higher intake group”: they consumed fast-foods once and more per week). In addition, each of fast-foods consumption group was subdivided into super-healthy, MetS risk carriers, and MetS according to MetS RF status. Finally, study subjects were categorized into 6 sub-groups (the low intake group: super healthy [n = 61], MetS risk carrier [n = 75]; the high intake group: super healthy [n = 50], MetS risk carrier [n = 38], and MetS [n = 10]) (**Figure 1**). Interestingly, subjects with MetS were observed only in the high fast-food intake group. All subjects in the high fast-foods intake group were more obese, and had higher levels of body fat (%) and LDL-C than those in the low fast-foods intake group. Circulating levels of hs-CRP and IL-6 were also significantly higher in the high fast-foods intake group than in the low fast-foods intake group. Particularly, IL-6 levels in the high fast-foods intake group were increased as the number of MetS RF increased.

Table 3. Relationship between food group consumption patterns and MetS related risk parameters

Food group*		Waist	SBP	DBP	TG [†]	HDL-C	LDL-C [†]	Glucose [†]	Insulin [†]	C-peptide [†]	IR [†]	hs-CRP [†]	IL-6 [†]	TNF- α [†]
Grains	r	-0.066	0.038	0.004	0.046	0.078	0.083	0.074	0.115	0.071	0.123	-0.016	0.100	0.051
	p	0.321	0.569	0.957	0.489	0.242	0.213	0.269	0.173	0.481	0.144	0.814	0.136	0.451
Meats and fish	r	-0.043	0.011	0.045	-0.014	-0.018	0.119	-0.054	-0.120	-0.133	-0.118	-0.047	-0.008	-0.071
	p	0.519	0.863	0.498	0.831	0.792	0.072	0.413	0.154	0.182	0.161	0.483	0.901	0.294
High-fat	r	-0.060	-0.027	0.072	-0.042	0.015	0.058	-0.075	-0.087	-0.087	-0.100	0.026	-0.043	0.027
	p	0.365	0.686	0.280	0.529	0.825	0.384	0.259	0.304	0.382	0.236	0.699	0.518	0.693
Medium-fat	r	0.044	0.076	0.134	-0.043	0.048	0.060	-0.009	-0.033	-0.026	-0.033	0.064	0.051	-0.101
	p	0.506	0.253	0.044	0.523	0.475	0.365	0.897	0.698	0.793	0.699	0.343	0.451	0.131
Low-fat	r	-0.086	-0.037	-0.072	0.031	-0.077	0.121	-0.052	-0.144	-0.174	-0.134	-0.144	-0.041	-0.038
	p	0.196	0.573	0.276	0.643	0.248	0.069	0.431	0.087	0.081	0.112	0.032	0.545	0.576
Seafoods among the low-fat fish														
Crab, shellfish, oyster, etc.	r	-0.115	-0.048	-0.030	-0.041	0.008	0.180	-0.158	-0.084	-0.100	-0.100	-0.211	-0.155	0.001
	p	0.084	0.473	0.649	0.539	0.907	0.006	0.017	0.320	0.317	0.237	0.002	0.021	0.992
Anchovy, dried white bait	r	-0.100	-0.073	-0.104	-0.038	0.010	0.090	-0.038	-0.186	-0.170	-0.170	-0.029	-0.061	-0.011
	p	0.133	0.272	0.117	0.568	0.881	0.177	0.569	0.027	0.087	0.044	0.665	0.368	0.865
Vegetables total														
r	-0.049	0.064	0.027	-0.056	0.032	0.037	-0.012	-0.076	0.050	-0.073	-0.188	-0.085	-0.086	
p	0.461	0.337	0.684	0.396	0.628	0.574	0.854	0.369	0.620	0.391	0.005	0.207	0.203	
Vegetable subgroup														
Green-yellow	r	-0.075	0.051	0.020	-0.057	0.012	0.029	-0.114	-0.170	-0.117	-0.180	-0.162	-0.081	-0.084
	p	0.258	0.443	0.759	0.303	0.853	0.510	0.086	0.043	0.242	0.032	0.015	0.229	0.211
Young radish, radish leaves	r	-0.090	0.093	0.052	-0.129	0.060	-0.041	-0.048	-0.219	-0.163	-0.213	-0.074	-0.046	0.075
	p	0.175	0.164	0.434	0.052	0.370	0.541	0.475	0.009	0.101	0.011	0.271	0.494	0.264
Kimchi, diced radish kimchi	r	-0.043	-0.014	0.004	0.040	0.042	0.040	0.032	0.044	0.195	0.046	-0.194	-0.081	-0.163
	p	0.515	0.831	0.957	0.550	0.526	0.553	0.634	0.600	0.049	0.589	0.004	0.228	0.015
Milk products														
r	-0.060	0.025	0.015	-0.006	0.046	0.019	-0.047	-0.064	-0.036	-0.076	0.043	-0.022	-0.058	
p	0.370	0.704	0.816	0.933	0.485	0.780	0.482	0.447	0.716	0.366	0.526	0.742	0.389	
Fruits														
r	-0.126	-0.062	-0.082	-0.122	0.156	-0.127	-0.054	-0.160	-0.098	-0.157	-0.063	-0.082	-0.055	
p	0.057	0.353	0.215	0.065	0.018	0.055	0.417	0.057	0.326	0.062	0.352	0.220	0.415	
Sugars: sugar, syrup, candy, chocolate, etc.														
r	0.108	0.065	0.004	-0.002	-0.011	0.157	-0.034	0.016	0.027	0.013	0.017	0.029	-0.013	
p	0.104	0.331	0.948	0.980	0.870	0.018	0.614	0.853	0.786	0.878	0.798	0.671	0.849	
Fast-foods														
r	0.023	0.013	-0.003	-0.004	-0.025	0.141	0.005	-0.002	0.001	-0.007	0.156	0.132	0.043	
p	0.734	0.851	0.960	0.957	0.703	0.033	0.939	0.980	0.988	0.937	0.020	0.050	0.527	

Tested by partial correlation analysis adjusted age, sex, total calorie intake, cigarette smoking, and alcohol drinking.

MetS, metabolic syndrome; SBP, systolic blood pressure; DBP, diastolic blood pressure; TG, triglyceride; HDL-C, high-density lipoprotein cholesterol; LDL-C, low-density lipoprotein cholesterol; IR, insulin resistance; hs-CRP, high-sensitivity C-reactive protein; IL-6, interleukin-6; TNF- α , tumor necrosis factor-alpha; r, correlation coefficient.

*Calculated as sum total of one-serving sized food consumption in each food group per week; [†]Tested after log-transformed.

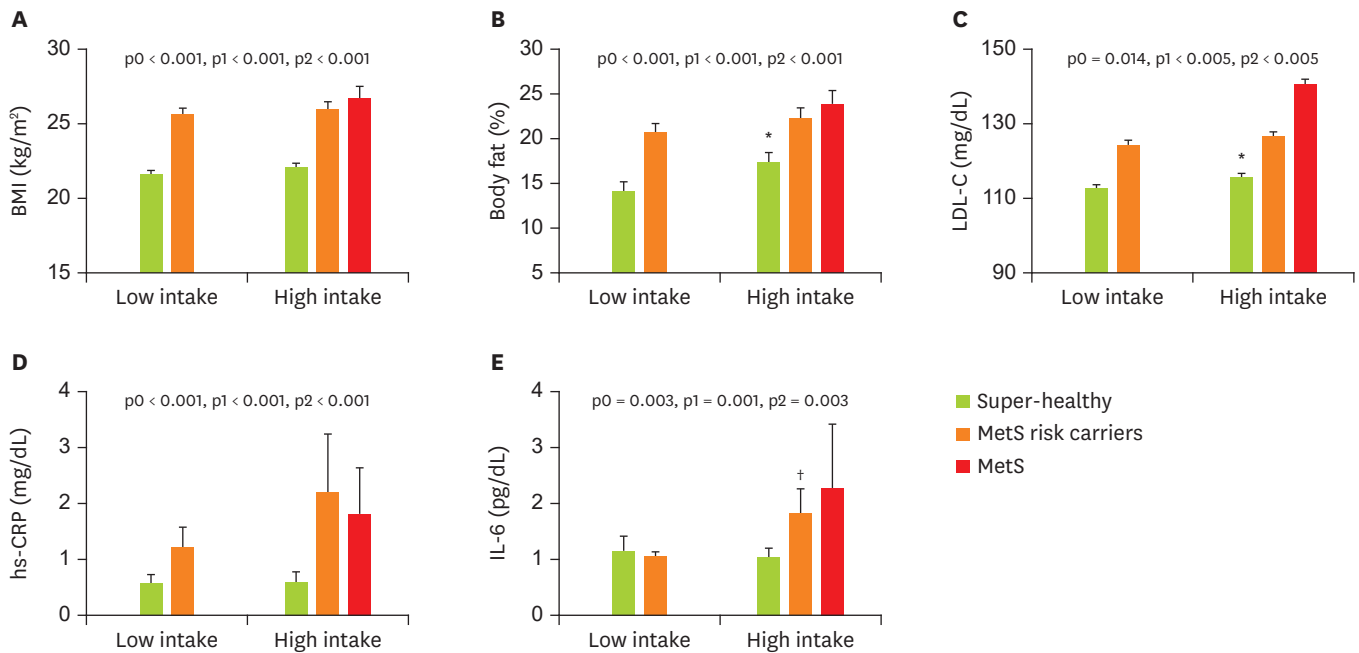


Figure 1. Adiposity and inflammation according to MetS risk status and fast-food intake level. Data are means ± SE. p₀, unadjusted p value; p₁, p value adjusted for age and sex; p₂, p value adjusted for age, sex, total calorie intake, cigarette smoking, and alcohol drinking; BMI, body mass index; LDL-C, low-density lipoprotein cholesterol; hs-CRP, high-sensitivity C-reactive protein; IL-6, interleukin-6; MetS, metabolic syndrome; SE, standard error; ANOVA, analysis of variance. *Tested after log-transformed; †Tested by 1-way ANOVA with Bonferroni method or general linear model with adjustment.

DISCUSSION

This cross-sectional study was performed to examine the association between food group consumption pattern, and early risk of MetS in non-diabetic healthy Korean adults. We identified that dietary habits and selected food consumption patterns are significantly associated with early alteration of MetS risk related parameters. This is the first study to elucidate the relationship between food consumption patterns and MetS risk status focusing on early alteration of endocrine metabolism in healthy Korean adults.

In the present study, we observed that dietary habit such as meal regularity, and the frequencies of having breakfast, overeating and eating-out were related to MetS risk. Specifically, higher proportions of irregular eating, less breakfast eating, and frequent overeating and eating out were observed in subjects with MetS RFs, particularly in those with MetS than in super-healthy people. Our results are partly in accordance with previous reports [13,21-23]. Pot et al. [13] reported that individuals with a more irregular intake of energy, especially during breakfast, had the increased cardio-metabolic risk. Odegaard et al. [21] also demonstrated that having breakfast everyday was strongly associated with the reduced risk of metabolic conditions. Shin et al. [22] showed that subjects with MetS more overeat than those without MetS. Also, people with MetS showed higher rate of eating out than those without MetS in the study by Ohta et al. [23].

Regarding the association between food group consumption patterns and MetS risk related parameters, the low-fat meats and fish consumption was negatively correlated with the inflammatory response in this study. Particularly, higher seafood consumption was associated with lower levels of hs-CRP and IL-6. However, contrary to our results,

Nanri et al. [24] reported that seafood consumption pattern was significantly associated with elevated hs-CRP levels in men. Precise mechanisms for the relationship between seafood intake and inflammation need to be elucidated. In addition, higher consumption of anchovy and dried white baits were associated with lower levels of insulin and HOMA-IR. Anchovy and dried white baits which belong to low-fat fish group are rich sources of dietary calcium. The Korean population study based on the Korea National Health and Nutrition Examination Survey 2008–2011 shows that calcium intake may reduce the prevalence of MetS in postmenopausal women [25]. Calcium supplementation also improved insulin sensitivity in adult obese rats [26].

Vegetable consumption was related to inflammatory response and IR in this study. Among vegetables, higher green-yellow vegetables intake was associated with lower levels of hs-CRP and HOMA-IR. Similar result was observed in the report of Nettleton et al. [27]: consumption of green leafy vegetables, whole grains, fruits, and nuts were negatively related to the levels of hs-CRP and IL-6 in a multi-ethnic population. Cook et al. [28] also found in their overweight Latino youth participants that dark green and deep yellow/orange vegetables were positively correlated with insulin sensitivity, but not with IR. In the present study, young radish and their leaves were also negatively associated with insulin levels and IR. However, there were no other reports for the relationship and the mechanisms, thereby further studies being needed. In addition, kimchi consumption was inversely associated with inflammatory responses (i.e., hs-CRP and TNF- α). It is partly in accordance with the report of Oh et al. [29] presenting that consumption of rice combined with kimchi was related to a lower risk of MetS. However, no significant changes were observed in proinflammatory cytokines after 4-week kimchi consumption in Chinese people performed by Lee et al. [30]. To elucidate the relationship between kimchi consumption and inflammation, long-term prospective observation studies are needed.

The fruits consumption was inversely associated with MetS related parameters in the current study. Fruits have plenty of nutrients and bioactive compounds, such as vitamins, minerals, antioxidants, phytochemicals, etc. Daily intake of flavanones was associated with lipid-lowering properties in type 2 diabetic patients with MetS [31]. According to the Tehran lipid glucose study, higher intake of red/purple fruits and vegetables (FVs) was related to lower body weight and abdominal fat gain after 3-year follow-up. Yellow, green, and white FVs were related to favorable lipid parameters [32]. These results are similar with our results. Park et al. [33] also exhibited that higher intake of fruits in Korean women had lower incidences of MetS. Based on our result and previous reports, fruits consumption may be beneficial for controlling lipid profiles, particularly in people exposed at early status of MetS risk.

The current study also presented that LDL-C levels were positively related with consumptions of sugary foods (chocolate, syrup, candy, chocolate, etc.) as well as fast-foods (burgers, pizzas, fries, instant noodles, etc.). Stanhope et al. [34] reported through their 2-week intervention study that consumption of high-fructose corn syrup sweetened beverages for 2 weeks increased concentrations of LDL-C, apoB, and other risk factors for CVD. Aeberli et al. [35] showed that LDL particle size was reduced after high fructose and high sucrose consumption, and more atherogenic LDL subclass distribution was observed when fructose-containing sugar sweetened beverages. Maersk et al. [36] also showed that daily intake of sucrose-sweetened soft drinks for 6-month increased ectopic fat accumulation and blood lipids. In addition, fast food consumption was significantly related to the increased serum levels of triglyceride and LDL-C in Iranian adults, particularly in middle aged adults [37]. Fast

food consumption was also associated with higher LDL-C and other MetS related parameters in the adolescents [38].

In the present, we sub-categorized study subjects according to their fast-foods intake levels (less than 1 time per week vs. 1 time and more per week), and observed higher adiposity and inflammation in higher fast-foods intake group particularly with MetS risk. Anderson et al. [39] found the odds of being obese were higher in subjects consuming fast-foods more than once a week than people who consumed less than once a week. El-Seweidy et al. [40] reported that higher expression of genes related to inflammation was induced by higher fast-food consumption.

This study has a limitation that the number of the study subjects was relatively small to be generalized to all population. Further studies with larger population and longitudinal observation are needed. Despite the limitation, these results may suggest useful information to educate people to properly select healthy foods for early prevention of MetS and CVD.

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