

# Binge Drinking and Its Relation to Metabolic Syndrome in Korean Adult Men

Original  
Article

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**Background:** It is reported that heavy drinking increases the risk of metabolic syndrome. But there have been few studies on the relationship between the intensity of drinking and metabolic syndrome when drinking the same amount of alcohol. This study aimed to assess the relationship between the frequency of binge drinking and metabolic syndrome in Korean adult men.

**Methods:** From the database of the 4th and 5th Korea National Health and Nutrition Examination Survey conducted in 2007–2010, data of 8,305 adult men ( $\geq 19$  years of age) was included in this analysis. Cross-sectional relationship between the frequency of binge drinking and metabolic syndrome was investigated adjusting for pure alcohol consumed per day.

**Results:** Adjusting for various confounders including pure alcohol consumed per day, the adjusted odds ratio for metabolic syndrome in those in higher frequency (more than 1/wk) binge drinking group was 1.62 (95% confidence interval, 1.30 to 2.03;  $P$  for trend =  $<0.001$ ) compared to those in the non-binge drinking group. Through analysis of the relationship between pure alcohol consumed per day and metabolic syndrome, it was found that pure alcohol consumed per day had a positive relation to metabolic syndrome in the higher frequency binge drinking group ( $P$  for trend = 0.041). The relationship was inverse in the non-binge drinking group ( $P$  for trend = 0.002).

**Conclusion:** Our study found a positive relationship between frequency of binge drinking and metabolic syndrome in adult men. And the effect of drinking on metabolic syndrome may depend on the frequency of binge drinking. Further studies are required to confirm this association.

**Keywords:** Binge Drinking; Metabolic Syndrome; Asian Continental Ancestry Group; Men

## INTRODUCTION

Metabolic syndrome is a combination of metabolic risk factors and increases the risk of heart attack or stroke and the mortality from related diseases such as diabetes and

hypertension.<sup>1,2)</sup> The prevalence of metabolic syndrome has been increasing worldwide over the past few years.<sup>3)</sup> According to the Korea National Health and Nutrition Examination Survey, the prevalence of metabolic syndrome in Korea was 24.1% in 2005.<sup>4)</sup>

There have been many studies on the relationship between drinking and metabolic syndrome. It was reported that heavy drinking increases the risk of metabolic syndrome.<sup>5)</sup> On the other hand, moderate drinking is reported to lower the risk of metabolic syndrome.<sup>6)</sup> But there have been few studies on the relationship between the intensity of drinking and metabolic syndrome. In terms of the distinction between frequent drinking of smaller quantities of alcohol versus occasional drinking of large quantities of alcohol, it has not been reported which increases the risk of metabolic syndrome more.

The level of alcohol consumption per person in Korea was

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8.1 L per year in 2005, lower than Organization for Economic Cooperation and Development average of 9.5 L per year.<sup>7)</sup> However, Korea has the highest rate of binge drinking (defined as drinking more than five glasses at once) among the countries that provided data on global alcohol drinking patterns published by the World Health Organization (WHO) in 2005.<sup>8)</sup> Therefore, the authors aimed to investigate the relationship between the quantity of alcohol consumed per drinking episode and metabolic syndrome in Korean adult men.

## METHODS

### 1. Subjects

From the database of the 4th and 5th Korea National Health and Nutrition Examination Survey (KNHANES) conducted in 2007–2010, data of 10,882 adult men ( $\geq 19$  years of age) was included in this analysis. We excluded the subjects whose independent variables (the questionnaires about alcohol consumption) were missing ( $n = 917$ ). We also excluded the subjects whose dependent variables (data of triglyceride, fasting glucose, waist circumference, systolic blood pressure, diastolic blood pressure, high density lipoprotein [HDL] cholesterol) were missing ( $n = 430$ ) and the subjects whose confounding variables (questionnaires about smoking, exercise, marital status, income, working status, insurance, education, height, weight) were missing ( $n = 1,230$ ). Following these exclusions, we included 8,305 subjects in the analysis.

### 2. Demographic and Socioeconomic Factors

We collected data on the demographic factors and health behaviors of the subjects. The demographic variables were age, sex, educational level (elementary school or less, middle school, high school, and college or more), working status (active, inactive), marital status (married, separated/widowed/divorced, single), insurance status (national health insurance, medical aid), and monthly income ( $< 1$  thousand won/mo,  $\geq 1$  thousand won/mo and  $< 3$  thousand won/mo,  $\geq 3$  thousand won/mo). The health behavioral variables included smoking (non-smoker, ex-smoker, smoker), exercise (low, moderate, or high by tertiles of metabolic equivalent of task minutes per week). The Korean version of the International Physical Activity Questionnaire

short form was used to calculate the metabolic equivalent of task minutes per week (MET-minutes per week).<sup>9)</sup> Then we classified the subjects into 3 exercise groups by tertiles of MET-minutes per week.

### 3. Assessment of Drinking

We calculated pure alcohol consumed per day using the database of the Korea National Health and Nutrition Examination Survey. We used the data of the frequency of drinking in the last 1 year and glasses of alcohol per drinking. Then we classified the subjects into nonconsumer ( $< 1$  g/d), light consumer (1–14.9 g/d), moderate consumer (15–29.9 g/d), and heavy consumer ( $\geq 30$  g/d) groups.<sup>10)</sup>

WHO defined high risk drinking as drinking over 60 g of pure alcohol a day for men, and 40 g of pure alcohol a day for women.<sup>11)</sup> According to this definition, binge drinking is defined as drinking more than 7 glasses of alcohol for men in the Korea National Health and Nutrition Examination Survey.

We grouped subjects into a non-binge drinking group and a binge drinking group. Then we classified the binge drinking group by the frequency of binge drinking according to the questionnaires of the Korea National Health and Nutrition Examination Survey (lower = about 1/mo or less, higher = more than 1/wk).

### 4. Anthropometric Measures

Body weight and height were measured using standard protocols to the nearest 0.1 kg and 0.1 cm, respectively. Waist circumference was measured at the narrowest point between the lower borders of the rib cage and the uppermost borders of the iliac crest at the end of normal expiration. Blood pressure was measured with a mercury sphygmomanometer.

### 5. Laboratory Evaluation

Antecubital vein blood samples were drawn and centrifuged after fasting (fasting time  $\geq 12$  h). These blood samples were used to evaluate total cholesterol, HDL cholesterol, triglyceride, and glucose levels.

### 6. Definition of Metabolic Syndrome

We defined metabolic syndrome according to National Cholesterol Education Program's Adult Treatment Panel III revised in 2005 (meeting more than 3 of following criteria):<sup>12)</sup>

1) abdominal obesity: waist circumference  $\geq 90$  cm;<sup>13)</sup> 2) high blood pressure: systolic blood pressure  $\geq 130$  mm Hg or diastolic blood pressure  $\geq 85$  mm Hg or taking hypertension medication; 3) fasting hyperglycemia: fasting blood sugar  $\geq 100$  mg/dL or taking diabetes medication or insulin therapy; 4) hypertriglyceridemia: triglyceride  $\geq 150$  mg/dL or taking dyslipidemia medication; 5) low HDL cholesterol: HDL cholesterol  $\leq 40$  mg/dL or taking dyslipidemia medication.

We grouped those who take dyslipidemia medication into a hypertriglyceridemia group and low HDL cholesterol group. And we grouped those who take diabetes medication or insulin into a fasting hyperglycemia group, and those who take hypertension medication into a high blood pressure group.

## 7. Analysis

We examined the characteristics of the subjects by t-test (continuous variable) and by chi-square test (categorical variable). We used multiple logistic regression analysis to examine the relationship between the frequency of binge drinking and metabolic syndrome and each component of metabolic syndrome. We examined the adjusted odds ratios and 95% confidence intervals adjusting for age in model 1. In model 2, smoking status, pure alcohol consumed per day, exercise, insurance status, education, working status, marital status, and monthly income were adjusted for in addition to model 1. In model 3, body mass index was adjusted in addition to model 2. We divided subjects into 3 groups including a non-binge drinking group, a lower binge drinking group, and a higher binge drinking group, and divided each group into 4 subgroups by pure alcohol consumed per day. Adjusting for age, smoking status, exercise, insurance status, education, working status, marital status, and monthly income, we examined the adjusted odds ratio for metabolic syndrome in those in each of the 11 subgroups compared to those in the non-binge drinking and nonconsumer subgroups. Reported probability values were 2-sided and a P-value  $< 0.05$  was considered statistically significant. Stata SE ver. 12.0 (Stata Co., College Station, TX, USA) was used in all statistical analysis with 'svy' commands to account for complex sampling design, and included sampling weights, which enabled the results to represent the entire population of adult men.

## RESULTS

### 1. Characteristics of Subjects

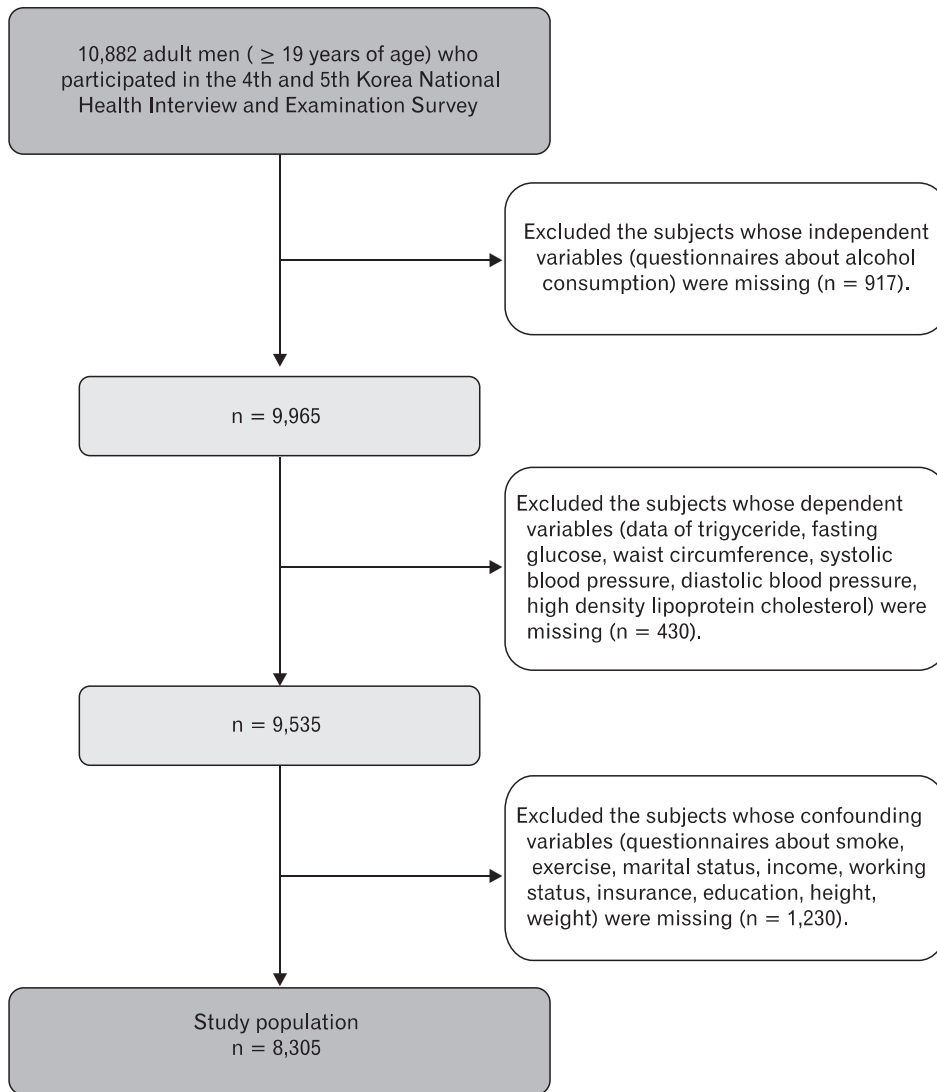
As demonstrated in Figure 1, final study population were 8,305 men. Subjects who had higher frequency of binge drinking had higher body mass index, waist circumference, triglyceride, HDL cholesterol, systolic blood pressure, diastolic blood pressure, fasting glucose, pure alcohol consumed per day, and statistically significant. Age, hypertension, diabetes, dyslipidemia treatment, smoking status, exercise, marital status, education, working status, monthly income, and insurance status were different between groups and statistically significant (Table 1).

### 2. The Risk of Metabolic Syndrome According to the Frequency of Binge Drinking

Table 2 presents the results. In model 1, the adjusted odds ratio for metabolic syndrome in those who had higher frequency (more than 1/wk) of binge drinking was 1.94 (95% confidence interval [CI], 1.67 to 2.26; P for trend =  $< 0.001$ ). In model 2, the adjusted odds ratio for metabolic syndrome in those who had a higher frequency (more than 1/wk) of binge drinking was 1.62 (95% CI, 1.30 to 2.03; P for trend =  $< 0.001$ ). Among the components of metabolic syndrome, abdominal obesity, high blood pressure, fasting hyperglycemia, hypertriglyceridemia had a positive association with the frequency of binge drinking (P for trend = 0.048, 0.003,  $< 0.001$ ,  $< 0.001$ , respectively). In model 3, the adjusted odds ratio for metabolic syndrome in those who had higher frequency (than 1/wk) of binge drinking was 1.56 (95% CI, 1.23 to 1.98; P for trend =  $< 0.001$ ). Among the components of metabolic syndrome, high blood pressure, fasting hyperglycemia, and hypertriglyceridemia had a positive association with the frequency of binge drinking (P for trend = 0.012,  $< 0.001$ , 0.001, respectively).

### 3. The Risk of Metabolic Syndrome According to the Frequency of Binge Drinking and Daily Pure Alcohol Consumed per Day

We examined the adjusted odds ratio for metabolic syndrome in those in each subgroup of the frequency of binge drinking and pure alcohol consumed per day. Table 3 presents the results. We could not calculate the odds ratio in those in the non-binge drinking and heavy consumer groups, because there were only



**Figure 1.** The study population framework. 2007–2010 The Fourth and Fifth Korean National Health and Nutritional Examination Survey.

2 subjects in the subgroups. In each group of light consumer and moderate consumer, the frequency of binge drinking had a positive association with metabolic syndrome ( $P$  for trend =  $<0.001$ ,  $0.018$ , respectively). In the heavy consumer group, the odds ratio for metabolic syndrome was  $2.10$  (95% CI,  $0.94$  to  $4.69$ ) in the lower binge drinking group, and the odds ratio was  $1.99$  (95% CI,  $1.64$  to  $2.42$ ) in the higher binge drinking group. In the higher frequency binge drinking group, pure alcohol consumed per day had a positive association with metabolic syndrome ( $P$  for trend =  $0.041$ ). In the non-binge drinking group, light consumers had a lower risk for metabolic syndrome compared to nonconsumers or moderate consumers.

## DISCUSSION

We found that the frequency of binge drinking is positively associated with metabolic syndrome in adult men after adjusting for pure alcohol consumed per day and other possible confounding factors. We also found that pure alcohol consumed per day was positively associated with a higher risk of metabolic syndrome in the higher frequency binge drinking group, while an inverse association was observed in the non-binge drinking group. In other words, the effect of drinking on metabolic syndrome may depend on the frequency of binge drinking.

Another study was conducted to examine the relationship between the intensity of alcohol drinking and metabolic syndrome.<sup>14)</sup> In that study, the intensity of drinking had a linear relationship with

**Table 1.** Characteristics of subjects according to the frequency of binge drinking\* (n = 8,305)

Characteristic	Non-binge drinking group (n = 2,582)	Binge drinking* group		P-value <sup>‡</sup>
		About ≤1 mo <sup>†</sup> (n = 2,768)	≥1/wk <sup>†</sup> (n = 2,955)	
Age (y)	56.22 ± 16.52	44.35 ± 15.36	45.73 ± 13.80	<0.001
Body mass index (kg/m <sup>2</sup> )	23.55 ± 3.18	23.98 ± 3.09	24.35 ± 3.10	<0.001
Waist circumference (cm)	84.11 ± 16.87	83.85 ± 8.78	85.49 ± 8.57	<0.001
High density lipoprotein cholesterol (mg/dL)	46.36 ± 11.03	48.33 ± 11.49	51.28 ± 12.74	<0.001
Triglyceride (mg/dL)	135.40 ± 86.74	147.72 ± 109.83	188.63 ± 180.11	<0.001
Systolic blood pressure (mm Hg)	122.39 ± 17.06	119.64 ± 15.37	123.17 ± 15.58	<0.001
Diastolic blood pressure (mm Hg)	77.81 ± 10.30	78.90 ± 10.50	82.08 ± 10.61	<0.001
Fasting glucose (mg/dL)	101.24 ± 27.97	97.33 ± 21.04	101.07 ± 25.00	<0.001
On hypertension medication (n = 1,395)	599 (23.20)	350 (12.64)	446 (15.09)	<0.001
On diabetes treatment (n = 555)	254 (9.84)	132 (4.77)	169 (5.72)	<0.001
On hyperlipidemia medication (n = 260)	103 (3.99)	72 (2.60)	85 (2.88)	0.01
Smoking				<0.001
Non-smoker (n = 1,805)	741 (28.70)	665 (24.02)	399 (13.50)	
Ex-smoker (n = 3,594)	739 (28.62)	1,208 (43.64)	1,647 (55.74)	
Current-smoker (n = 2,906)	1,102 (42.68)	895 (32.33)	909 (30.76)	
Exercise <sup>§</sup>				<0.001
1st (n = 2,723)	750 (29.05)	881 (31.83)	1,092 (36.95)	
2nd (n = 2,680)	810 (31.37)	927 (33.49)	943 (31.91)	
3rd (n = 2,902)	1,022 (39.58)	960 (34.68)	920 (31.13)	
Pure alcohol consumed per day (g/d)				<0.001
<1 (n = 2,361)	2,008 (77.77)	345 (12.46)	8 (0.27)	
1–14.9 (n = 3,283)	508 (19.67)	2,036 (73.55)	739 (25.01)	
15–29.9 (n = 1,398)	64 (2.48)	344 (12.43)	990 (33.50)	
≥30 (n = 1,263)	2 (0.08)	43 (1.55)	1,218 (41.22)	
Marital Status				<0.001
Married (n = 6,491)	2,147 (83.15)	2,019 (72.94)	2,325 (78.68)	
Separated/widowed/divorced (n = 417)	151 (5.85)	110 (3.97)	156 (5.28)	
Single (n = 1,379)	284 (11.00)	639 (23.09)	474 (16.04)	
Education				<0.001
≤Elementary (n = 1,507)	724 (28.04)	303 (10.95)	480 (16.24)	
Middle (n = 1,020)	399 (15.45)	272 (9.83)	349 (11.81)	
High (n = 3,037)	804 (31.14)	1,051 (37.97)	1,182 (40.00)	
≥College (n = 2,741)	655 (25.37)	1,142 (41.26)	944 (31.95)	
Work				<0.001
Economically active (n = 1,959)	875 (33.89)	602 (21.75)	482 (16.31)	
Economically inactive (n = 6,346)	1,707 (66.11)	2,166 (78.25)	2,473 (83.69)	
Monthly income (won)				<0.001
<1,000 (n = 1,415)	681 (26.37)	356 (12.86)	378 (12.79)	
≥1,000, <3,000 (n = 3,221)	1,001 (38.77)	1,073 (38.76)	1,147 (38.82)	
≥3,000 (n = 3,669)	900 (34.86)	1,339 (48.37)	1,430 (48.39)	
Insurance				<0.001
National health insurance (n = 8,107)	2,495 (96.63)	2,716 (98.12)	2,896 (98.00)	
Medical aid (n = 198)	87 (3.37)	52 (1.88)	59 (2.00)	

Values are presented as mean ± SD or number (%).

\*Defined as drinking more than 7 glasses of alcohol at once. <sup>†</sup>The frequency of binge drinking. <sup>‡</sup>Examined the characteristics of participants by t-test (continuous variable) and by chi-square test (categorical variable). <sup>§</sup>Used the protocol of the Korean version of the International Physical Activity Questionnaire short form to calculate metabolic equivalent of task minutes per week (MET-minutes per week), then classified into 3 groups by tertiles of MET-minutes per week. The 1st group has the lowest MET-minutes per week in average, and the 3rd group has the highest MET-minutes per week in average.

**Table 2.** The number of events and the adjusted odds ratios (95% confidence intervals) for metabolic syndrome\* and each component of metabolic syndrome\* according to the frequency of binge drinking<sup>†</sup> (n = 8,305)

Variable	Non-binge drinking group	Binge drinking <sup>†</sup> group		P for trend
		About ≤1 mo <sup>‡</sup>	≥1 wk <sup>‡</sup>	
Metabolic syndrome*	737 (28.54)	645 (23.30)	948 (32.08)	-
Model 1 <sup>§</sup>	1	1.24 (1.06–1.46)	1.94 (1.67–2.26)	<0.001
Model 2 <sup>  </sup>	1	1.13 (0.95–1.34)	1.62 (1.30–2.03)	<0.001
Model 3 <sup>¶</sup>	1	1.15 (0.96–1.38)	1.56 (1.23–1.98)	<0.001
Abdominal obesity*	653 (25.29)	647 (23.37)	885 (29.95)	-
Model 1 <sup>§</sup>	1	1.12 (0.96–1.31)	1.51 (1.29–1.75)	<0.001
Model 2 <sup>  </sup>	1	1.01 (0.85–1.21)	1.25 (1.00–1.56)	0.05**
Model 3 <sup>¶</sup>	1	1.04 (0.81–1.35)	1.12 (0.81–1.56)	0.48
High blood pressure*	1,243 (48.14)	1,078 (38.95)	1,514 (51.24)	-
Model 1 <sup>§</sup>	1	1.18 (1.01–1.39)	1.92 (1.66–2.23)	<0.001
Model 2 <sup>  </sup>	1	1.03 (0.86–1.23)	1.40 (1.12–1.76)	0.003
Model 3 <sup>¶</sup>	1	1.02 (0.85–1.22)	1.34 (1.06–1.69)	0.01
Fasting hyperglycemia*	908 (35.17)	782 (28.25)	1,111 (37.60)	-
Model 1 <sup>§</sup>	1	1.16 (0.98–1.38)	1.84 (1.56–2.17)	<0.001
Model 2 <sup>  </sup>	1	1.11 (0.92–1.33)	1.69 (1.34–2.14)	<0.001
Model 3 <sup>¶</sup>	1	1.11 (0.92–1.33)	1.62 (1.27–2.05)	<0.001
Hypertriglyceridemia*	867 (33.58)	997 (36.02)	1,389 (47.01)	-
Model 1 <sup>§</sup>	1	1.36 (1.19–1.56)	2.18 (1.92–2.47)	<0.001
Model 2 <sup>  </sup>	1	1.13 (0.97–1.31)	1.49 (1.21–1.83)	<0.001
Model 3 <sup>¶</sup>	1	1.12 (0.96–1.31)	1.41 (1.14–1.75)	0.00
Low HDL*	821 (31.80)	676 (24.42)	563 (19.05)	-
Model 1 <sup>§</sup>	1	0.90 (0.78–1.05)	0.65 (0.56–0.77)	<0.001
Model 2 <sup>  </sup>	1	1.04 (0.88–1.23)	0.92 (0.72–1.17)	0.51
Model 3 <sup>¶</sup>	1	1.04 (0.87–1.23)	0.86 (0.67–1.11)	0.25

Values are presented as number (%) or adjusted odds ratio (95% confidence interval).

\*Metabolic syndrome is defined by having 3 or more of the following component: high density lipoprotein cholesterol < 40 mg/dL, fasting plasma glucose ≥ 100 mg/dL or use of hypoglycemic medication, serum triglyceride ≥ 150 mg/dL, blood pressure ≥ 130 (systolic)/85 (diastolic) mm Hg or use of blood pressure medication, and waist circumference ≥ 90 cm (Asia-Pacific region). <sup>†</sup>Defined as drinking more than 7 glasses of alcohol at once. <sup>‡</sup>The frequency of binge drinking. <sup>§</sup>Statistical analysis by logistic regression with adjustment for age. <sup>||</sup>Adjusted for smoking status, pure alcohol consumed per day, exercise, insurance status, education, working status, marital status, monthly income in addition to model 1. <sup>¶</sup>Adjusted for body mass index, in addition to model 2. \*\*0.048.

fasting hyperglycemia and hypertriglyceridemia. But the association was not independent of the total amount of drinking. Thus, there was a methodological difference with our study.

It is well recognized that heavy drinking increases the risk of high blood pressure and high triglyceride and metabolic

syndrome.<sup>5,7,15)</sup> Moderate drinking is reported to lower the risk of metabolic syndrome.<sup>11)</sup> The relationship between drinking and metabolic syndrome may be a J-shape.<sup>16)</sup> There have been few studies examining the relationship between drinking and cardiovascular disease which also consider binge drinking.



**Table 3.** The adjusted\* odds ratios (95% confidence intervals) of metabolic syndrome<sup>†</sup> according to the frequency of binge drinking<sup>‡</sup> and daily pure alcohol consumed per day (n = 8,305)

Variable	Non-binge drinking group	Binge drinking <sup>‡</sup> group		P for trend
		About ≤1 mo <sup>§</sup>	≥1 wk <sup>§</sup>	
Nonconsumer <sup>  </sup>	1	1.07 (0.83–1.38)	1.39 (0.41–4.68)	0.25
Light consumer <sup>  </sup>	0.57 (0.40–0.81)	1.10 (0.89–1.35)	1.44 (1.03–2.01)	<0.001
Moderate consumer <sup>  </sup>	0.72 (0.39–1.32)	1.17 (0.91–1.51)	1.65 (1.36–1.99)	0.02
Heavy consumer <sup>  </sup>	- <sup>¶</sup>	2.10 (0.94–4.69)	1.99 (1.64–2.42)	- <sup>¶</sup>
P for trend	0.002	0.05**	0.04	

Values are presented as adjusted odds ratio (95% confidence interval).

\*Statistical analysis by logistic regression with adjustment for age, smoking status, exercise, insurance status, education, working status, marital status, monthly income. <sup>†</sup>Metabolic syndrome is defined by having 3 or more of the following component: high density lipoprotein cholesterol < 40 mg/dL, fasting plasma glucose ≥ 100 mg/dL or use of hypoglycemic medication, serum triglyceride ≥ 150 mg/dL, blood pressure ≥ 130 (systolic)/85 (diastolic) mm Hg or use of blood pressure medication, and waist circumference ≥ 90 cm (Asia-Pacific region).

<sup>‡</sup>Defined as drinking more than 7 glasses of alcohol at once. <sup>§</sup>The frequency of binge drinking. <sup>||</sup>Grouped by pure alcohol consumed per day into nonconsumer (<1 g/d), light consumer (≥1 g/d, <15 g/d), moderate consumer (≥15 g/d, <30 g/d), heavy consumer (≥30 g/d). <sup>¶</sup> Could not calculate the adjusted odds ratio because there were only 2 subjects in the subgroup. And could not calculate because there were only 2 groups in heavy consumer. \*\*0.052.

Interestingly, in our study, pure alcohol consumed per day had a protective effect on metabolic syndrome in the non-binge drinking group. In addition, in the highest binge drinking group, pure alcohol consumed per day had a positive association with metabolic syndrome, suggesting that not only the amount of alcohol consumed but also binge drinking is associated with cardiovascular disease.

Alcohol affects the onset of metabolic syndrome in various ways. Alcohol suppresses lipid oxidation from adipose tissue and enhances fat deposition, preferentially in the abdominal area, and can lead to increases triglyceride in the blood stream.<sup>17)</sup> It is reported that heavy drinking causes enhancement of insulin resistance and type 2 diabetes.<sup>18)</sup> And it is suggested that repeated drinking and withdrawal symptoms increases the risk of hypertension.<sup>19,20)</sup> On the other hand, alcohol suppresses the removal of HDL cholesterol in the blood stream, and reduces vitality and concentration of CETP (cholesterol ester transfer protein) thus decreasing the transformation of HDL of cholesterol ester to materials causing atherosclerosis.<sup>7,21)</sup> These effects are not fully understood and need further investigation. We can only suggest that these effects are intensified when drinking over a certain amount of alcohol at one time. as Another mechanism may be at work, as reported in one study in which binge drinking induced insulin resistance by disrupting hypothalamic insulin

action in rats,<sup>22)</sup> suggesting that the same mechanism may be at work in humans. More studies are required to examine the underlying mechanism.

Our Study has some limitations. First, daily food intake is reported to be significantly associated with metabolic syndrome. However, the questionnaire about food intake on the KNHANES was a single 24-hour dietary recall which is not an optimal way to assess daily food intake. We did not adjust for daily food intake using that data. Second, the cross-sectional nature of this study prevented us from determining an exact cause-and-effect relationship. Third, with regard to definition of binge drinking, an error may have occurred due to indirect measurement. But the cause-effect relationship between alcohol drinking and metabolic syndrome is already recognized. In spite of these limitations, our study is important since it is the first study to examine the relationship between the intensity of alcohol drinking and metabolic syndrome after adjusting for several confounders including pure alcohol consumed per day in Korea.

In conclusion, our study found a significant relationship between the frequency of binge drinking and metabolic syndrome in adult men. And the effect of drinking on metabolic syndrome may depend on the frequency of binge drinking. Further studies are required to confirm this association.

## CONFLICT OF INTEREST

No potential conflict of interest relevant to this article was reported.

## REFERENCES

1. Zimmet P, Alberti KG, Serrano Rios M. A New International Diabetes Federation (IDF) worldwide definition of the metabolic syndrome: the rationale and the results. *Rev Esp Cardiol (Engl Ed)* 2005;58:1371-5.
2. Gupta AK, Dahlof B, Sever PS, Poulter NR; Anglo-Scandinavian Cardiac Outcomes Trial-Blood Pressure Lowering Arm Investigators. Metabolic syndrome, independent of its components, is a risk factor for stroke and death but not for coronary heart disease among hypertensive patients in the ASCOT-BPLA. *Diabetes Care* 2010;33:1647-51.
3. Lim S, Shin H, Song JH, Kwak SH, Kang SM, Won Yoon J, et al. Increasing prevalence of metabolic syndrome in Korea: the Korean National Health and Nutrition Examination Survey for 1998-2007. *Diabetes Care* 2011;34:1323-8.
4. Korean Centers for Disease Control and Prevention. Korea National Health and Nutrition Examination Survey 2005 [Internet]. Cheongwon: Korean Centers for Disease Control and Prevention [cited 2013 Aug 16]. Available from: <http://knhanes.cdc.go.kr/knhanes/index.do>.
5. Zhu S, St-Onge MP, Heshka S, Heymsfield SB. Lifestyle behaviors associated with lower risk of having the metabolic syndrome. *Metabolism* 2004;53:1503-11.
6. Rosell M, De Faire U, Hellenius ML. Low prevalence of the metabolic syndrome in wine drinkers: is it the alcohol beverage or the lifestyle? *Eur J Clin Nutr* 2003;57:227-34.
7. Rimm EB, Williams P, Fosher K, Criqui M, Stampfer MJ. Moderate alcohol intake and lower risk of coronary heart disease: meta-analysis of effects on lipids and haemostatic factors. *BMJ* 1999;319:1523-8.
8. World Health Organization, Abuse S. Global status report on alcohol 2004. Geneva: World Health Organization; 2004.
9. Oh JY, Yang YJ, Kim BS, Kang JH. Validity and reliability of Korean version of International Physical Activity Questionnaire (IPAQ) short form. *J Korean Acad Fam Med* 2007;28:532-41.
10. Corrao G, Rubbiati L, Bagnardi V, Zambon A, Poikolainen K. Alcohol and coronary heart disease: a meta-analysis. *Addiction* 2000;95:1505-23.
11. World Health Organization. International guide for monitoring alcohol consumption and related harm. Geneva: World Health Organization; 2000.
12. 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.
13. 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* 2007;75:72-80.
14. Fan AZ, Russell M, Dorn J, Freudenheim JL, Nochajski T, Hovey K, et al. Lifetime alcohol drinking pattern is related to the prevalence of metabolic syndrome. The Western New York Health Study (WNYHS). *Eur J Epidemiol* 2006;21:129-38.
15. 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.
16. Oh SW. Effects of alcohol on obesity and metabolic syndrome. *Korean J Obe* 2009;18:1-7.
17. Suter PM. Is alcohol consumption a risk factor for weight gain and obesity? *Crit Rev Clin Lab Sci* 2005;42:197-227.
18. Xu D, Dhillon AS, Abelman A, Croft K, Peters TJ, Palmer TN. Alcohol-related diols cause acute insulin resistance in vivo. *Metabolism* 1998;47:1180-6.
19. Coca A, Aguilera MT, De la Sierra A, Sanchez M, Picado MJ, Lluch MM, et al. Chronic alcohol intake induces reversible disturbances on cellular Na<sup>+</sup> metabolism in humans: its relationship with changes in blood pressure. *Alcohol Clin Exp Res* 1992;16:714-20.
20. Randin D, Vollenweider P, Tappy L, Jequier E, Nicod P, Scherrer U. Suppression of alcohol-induced hypertension by dexamethasone. *N Engl J Med* 1995;332:1733-7.
21. Savolainen MJ, Kesaniemi YA. Effects of alcohol on



lipoproteins in relation to coronary heart disease. *Curr Opin Lipidol* 1995;6:243-50.

22. Lindtner C, Scherer T, Zielinski E, Filatova N, Fasshauer M,

Tonks NK, et al. Binge drinking induces whole-body insulin resistance by impairing hypothalamic insulin action. *Sci Transl Med* 2013;5:170ra14.