

ORIGINAL ARTICLE

Effect of alcohol and aldehyde dehydrogenase gene polymorphisms on alcohol-associated hypertension: the Guangzhou Biobank Cohort Study

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The effects of alcohol dehydrogenase (ADH) 2 and aldehyde dehydrogenase (ALDH) 2 genotypes on the alcohol–blood pressure association are unclear. We examined the association of ADH2 or ALDH2 genotypes with blood pressure in older Chinese men. Based on the Guangzhou Biobank Cohort Study (GBCS), 4792 men with valid ADH2, ALDH2 genotypes were included, and genotyping of rs1229984 ADH2 and rs671 ALDH2 (AA, AG/GA or GG) was performed using a Sequenom Mass-Array platform. Information on socio-demographics and lifestyle factors, including alcohol use, was obtained from a questionnaire, and blood pressure was measured. Among alcohol drinkers, systolic and diastolic blood pressure (SBP and DBP) and mean arterial pressure (MAP) were highest for men with the GG ADH2 genotype (136.6, 77.9 and 97.5 mm Hg, respectively), followed by those with the (AA/AG ADH2 + GG ALDH2) genotype (133.4, 77.6 and 96.2 mm Hg, respectively) and then the (AA/AG ADH2 + AA/AG ALDH2) genotype (SBP = 132.6, DBP = 76.6 and MAP = 95.2 mm Hg) (*P* for trend ranged 0.025–0.035). After adjustment for potential confounders, as well as frequency or amount of alcohol use, men with the GG ADH2 genotype were more likely to have hypertension (odds ratio (OR) = 1.62, 95% confidence interval 1.15–2.28) as were men with the (AA/AG ADH2 + AA/AG ALDH2) genotype (OR = 1.40, 95% confidence interval 1.01–1.96) compared with men with the (AA/AG ADH2 + GG ALDH2) genotype). ADH2 or ALDH2 genotypes were unrelated to hypertension among those who never drink alcohol. ADH2 genotype influences blood pressure and risk of hypertension among *male* alcohol drinkers, suggesting that the hypertensive effect of alcohol is due to ethanol rather than acetaldehyde.

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INTRODUCTION

Alcohol consumption is increasingly popular in China.¹ Based on a WHO report, alcohol consumption per capita in pure ethanol for Chinese adults was 1.03 litres in 1970, and rose to 4.5–5.2 litres in 1996–2001.^{2–4} The increasing use of alcohol has caused a great health, financial and social burden in China, yet alcohol consumption is continuing to rise. Local research is urgently needed to provide supporting evidence for developing a policy for alcohol restriction to improve public health.

Alcohol has been linked to hypertension in previous studies, and the association was partly explained by genetic variation in alcohol metabolism.⁵ Studies in Japanese and other East Asian populations have compared the alcohol–blood pressure relationships among individuals with varying activities of alcohol-metabolizing enzymes, such as alcohol dehydrogenase (ADH) and aldehyde dehydrogenase (ALDH),^{6,7} and have shown important roles for these enzymes in the

metabolism of alcohol in the body.⁸ After alcohol intake, ethanol is firstly metabolized to acetaldehyde then to acetic acid through several mechanisms. The main enzymes involved in the metabolic process are ADH2 and ALDH2 enzymes.^{9–11} The activities of these enzymes differ with genetic polymorphisms, leading to different effects of alcohol consumption on health. The GG and AA ALDH2 genotypes encode the active and the inactive subunit, respectively.^{12,13} The AA ALDH2 genotype results in higher blood acetaldehyde levels after alcohol intake, leading to facial flushing, palpitations and nausea.^{7,14} A meta-analysis of studies using ALDH2 single-nucleotide polymorphisms as an instrumental variable for alcohol use showed that alcohol significantly increased blood pressure and risk of hypertension in individuals of AA genotype.⁵ However, this meta-analysis was designed to assess the effect of alcohol use, and specifically ethanol, on blood pressure, but the effect of the ADH2 genotype on blood pressure was not assessed. As the isozymes of ADH2 encoded by the

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AA and AG genotypes have about 200 and 100 times higher ethanol oxidizing capacity, respectively, than the GG ADH2 genotype,¹⁵ the effect of alcohol on blood pressure should vary with ADH2 genotype. A previous study on men found that subjects with the GG genotype of ADH2 had higher blood pressure than those with the GA or AA genotype,¹⁶ but this study did not consider whether the association varied with ALDH2 genotype, despite the effects of ALDH2 on alcohol metabolism. Figure 1 shows the likely exposures for different combinations of ADH2 and ALDH2 genotypes. Individuals with an inactive ADH2 (GG) genotype metabolize alcohol slowly and have high ethanol exposure on alcohol use. Individuals with an inactive ALDH2 (AA or AG) genotype metabolize acetaldehyde slowly and have high acetaldehyde exposure on alcohol use. Individuals with an active ADH2 (AA or AG) genotype and an active ALDH2 (GG) genotype metabolize both ethanol and acetaldehyde quickly and have high acetic acid exposure on alcohol use. We hypothesized that in alcohol drinkers, those with an inactive ADH2 (GG) genotype have higher levels of ethanol than those with an active genotype and that this genotype may be associated with a higher risk of hypertension after alcohol drinking.

In the GBCS, we examined the associations of ADH2 and ALDH2 genotypes with blood pressure in men, and whether the associations varied with drinking status (alcohol drinker or never drinker).

METHODS

Ethics statement

All subjects had given informed consent before participating. The study has ethics approval from the Guangzhou Medical Ethics Committee of the Chinese Medical Association, Guangzhou, China.

The details of the GBCS

The GBCS is a 3-way collaboration among the Guangzhou No. 12 Hospital, the Universities of Hong Kong and Birmingham. The details have been reported elsewhere.¹⁷ Briefly, participants were recruited from the 'Guangzhou Health and Happiness Association for the Respectable Elders', a community social and welfare association unofficially aligned with the municipal government. This is a large unofficial organization with more than 150 branches throughout Guangzhou. The membership is open to Guangzhou residents aged 50 years or above for a nominal fee of 4 CNY (50 US cents) per month. Members of the Guangzhou Health and Happiness Association for the Respectable Elders account for about 7% of older permanent residents in Guangzhou. Approximately one-third of them participated in the GBCS if they were capable of consenting, ambulatory and not receiving treatment modalities that, if omitted, may result in immediate life-threatening risk, such as chemotherapy or radiotherapy for cancer, or dialysis for renal failure. As participants in the GBCS could receive a free medical examination, the response rate was very high (about 95%). About 5% of the eligible subjects refused to participate, particularly men, because of a cultural unwillingness to give blood due to an associated loss of 'shung qi' or 'life energy', or because of job commitments. The baseline examinations were conducted in three phases:

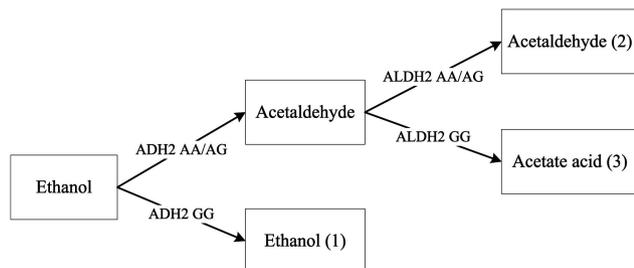


Figure 1 Effect of ADH2 and ALDH2 genotype on alcohol metabolism in human.

phase 1 from September 2003 to November 2004, phase 2 from April 2005 to May 2006 and phase 3 from September 2006 to January 2008. The second examination started in 2008 and in progress.

Computer based face-to-face interview and detailed physical examinations were done at baseline recruitment. Information on alcohol consumption was obtained in terms of frequency and usual amount per occasion for specific types of alcoholic beverage. Seated blood pressure was measured three times using an automated sphygmomanometer (Omron 705CP; Omron Corp., Kyoto, Japan), which had been previously validated by comparison with the manual mercury sphygmomanometer,¹⁸ and the last two measurements were averaged for analysis.

Blood samples were collected following 8–10 h overnight fasting. Blood samples for DNA extraction were available in phase 3 at baseline and in phases 1 and 2 at follow-up. DNA extraction was performed at the laboratory center of the GBCS in Guangzhou 12th Hospital.¹⁹ DNA was either extracted at baseline from fresh blood using a standard phenol–chloroform extraction procedure or was extracted from blood or buffy coat previously stored at -80°C using a standard magnetic bead extraction procedure. single-nucleotide polymorphism analysis was performed using a Sequenom Mass-Array platform, which amplifies DNA with specific primers, deactivates remaining nucleotide triphosphates by phosphatase treatment, performs a single base primer extension step and identifies allele-specific extension products using mass spectroscopy.

Outcomes

After resting for 5 min, seated SBP and DBP was measured three times using an automated sphygmomanometer (Omron 705CP), which had been previously validated by comparison with the manual mercury sphygmomanometer,¹⁸ and at least 1 min rest interval was required between each measurement. The last two measurements were averaged for data analysis in the present study. Outcomes considered were SBP, DBP, pulse pressure, defined as SBP–DBP, MAP defined as $1/3 \times (\text{SBP} + \text{DBP} \times 2)$ and hypertension defined as use of antihypertensive medication or SBP/DBP $\geq 140/90$ mm Hg.

Exposures

Exposures considered were rs671 ALDH2 genotype, rs1229984 ADH2 genotype and their combination, as set out in Figure 1, which shows how ADH2 and ALDH2 were combined to represent ethanol, acetaldehyde or acetic acid exposure.

Statistical analysis

We used genetic polymorphism of alcohol metabolism enzyme (ALDH2 and ADH2 genotypes) as the instrumental variable to obtain an unbiased estimation of the effect of alcohol on blood pressure. As genetic differences are determined before birth, these are unlikely to be associated with exogenous exposures or environmental factors. Thus the estimation might not be confounded by common hypertensive risk factors, such as smoking or sodium intake, as in other observational studies. Multivariable linear regression was used to assess differences in continuous variables between genotypes with adjustment for potential confounders, giving adjusted means and 95% confidence intervals, as well as *P*-value for trend. Logistic regression was used to assess the association of ADH2 and ALDH2 polymorphism with hypertension with adjustment for potential confounders. We assessed whether the association of ADH2 with the outcomes varied with ALDH2 from the *P*-value for interaction. We similarly assessed whether the association of ADH2, ALDH2 or the combined ADH2/ALDH2 genotype with the outcomes varied with drinking status. Potential confounders were age, occupation (manual, nonmanual, others), education, International Physical Activity Questionnaire physical activity (active, moderate, and inactive),²⁰ smoking (never, former and current), body mass index, fasting plasma glucose and total and high-density lipoprotein cholesterol. We excluded former drinkers when assessing the association of ADH2 and ALDH2 genotypes with blood pressure to reduce the 'abstainer error', as former drinkers might have quit drinking because of ill-health including hypertension, and received antihypertensive medication.²¹ Statistical significance was defined as a two-sided *P*-value < 0.05 . Data analysis was performed using STATA 10.1 (Stata Corp LP, College Station, TX, USA).

RESULTS

There were 4792 men with valid ADH2 and ALDH2 genotypes and complete information about all potential confounders. The AA, AG and GG genotype frequencies of ADH2 and ALDH2 were 52.7%, 40.2% and 7.1%, and 8.6%, 41.6% and 49.8%, respectively. The Hardy–Weinberg equilibrium test showed no variance between the population under study and the general population ($P=0.77$ for ADH2 and 0.99 for ALDH2 genotypes).

Table 1 shows that the ADH2 genotype was unrelated to drinking status or frequency, or to other risk factors, such as smoking, physical activity, education, occupation, age, body mass index, total and high-density lipoprotein-cholesterol, triglycerides and fasting glucose. Current drinking and higher alcohol use (≥ 140 g per week) were more common among men with the GG ADH2 genotype ($P=0.050$ and 0.026, respectively) than men with the other ADH2 genotypes. Current drinking, frequent alcohol drinking (≥ 5 per week) and higher alcohol use (≥ 140 g per week) were all more common among men with the GG than the AA or AG ALDH2 genotypes (all $P<0.001$). Men with the AA ALDH2 genotype were more likely to have had manual occupations ($P=0.040$) and lower total and

high-density lipoprotein-cholesterol ($P=0.036$ and 0.005, respectively).

There was no evidence that the association of ADH2 with the outcomes varied with ALDH2, probably because of insufficient sample size (P -values ranged from 0.19 to 0.44; table not shown). Similarly, there was no evidence that the association of ADH2 or ALDH2 with the outcomes varied with drinking status; however, the association of the combined ADH2/ALDH2 group with some outcomes did vary with drinking status (SBP $P=0.035$, DBP $P=0.035$ and MAP $P=0.025$). Table 2 shows that in never drinkers, ADH2 and ALDH2 were unrelated to SBP, DBP, pulse pressure or MAP. Among current drinkers, compared with the AA or AG ADH2 genotype, men with the GG genotype had higher SBP and pulse pressure ($P<0.05$). ALDH2 genotypes were unrelated to SBP, DBP, pulse pressure or MAP. After adjusting for multiple potential confounders and frequency of drinking, SBP, DBP and MAP were highest for men with the GG ADH2 genotype, followed by those with the (AA/AG ADH2 + GG ALDH2) genotype and then the (AA/AG ADH2 + AA/AG ALDH2) genotype (P for trend ranged 0.025–0.035) (Table 2). Sensitivity analysis with additional adjustment for amount of alcohol

Table 1 Characteristics by ADH2 and ALDH2 genotypes in older Chinese men

	ADH2 (rs1229984, n = 4792)				ALDH2 (rs671, n = 4782)			
	AA	AG	GG	P	AA	AG	GG	P
Cases, %	2524 (52.7)	1927 (40.2)	341 (7.1)	0.77 ^a	409 (8.6)	1990 (41.6)	2383 (49.8)	0.99 ^a
Drinking, n (%)								
Never	1350 (53.5)	955 (49.6)	162 (47.5)	0.050	315 (77.0)	1173 (58.9)	969 (40.7)	<0.001
Former	139 (5.5)	122 (6.3)	20 (5.9)		9 (2.2)	100 (5.0)	172 (7.2)	
Current	1035 (41.0)	850 (44.1)	159 (46.6)		85 (20.8)	717 (36.0)	1242 (52.1)	
Frequency of drinking, n (%)								
Never	1350 (53.5)	955 (49.6)	162 (47.5)	0.26	315 (77.0)	1173 (58.9)	969 (40.7)	<0.001
<1 per month	537 (21.3)	415 (21.5)	82 (24.0)		70 (17.1)	432 (21.7)	536 (22.5)	
<1 per week	99 (3.9)	86 (4.5)	15 (4.4)		5 (1.2)	83 (4.2)	110 (4.6)	
1–4 per week	142 (5.6)	120 (6.2)	18 (5.3)		4 (1.0)	78 (3.9)	196 (8.2)	
≥ 5 per week	257 (10.2)	229 (11.9)	44 (12.9)		6 (1.5)	124 (6.2)	400 (16.8)	
Former drinkers	139 (5.5)	122 (6.3)	20 (5.9)		9 (2.2)	100 (5.0)	172 (7.2)	
Amount of alcohol use, n (%)								
Never	1350 (53.5)	955 (49.6)	162 (47.5)	0.026	315 (77.0)	1173 (58.9)	969 (40.7)	<0.001
<140 g per week	1005 (39.8)	852 (44.2)	154 (45.2)		91 (22.2)	767 (38.5)	1151 (48.3)	
≥ 140 g per week	169 (6.7)	120 (6.2)	25 (7.3)		3 (0.7)	50 (2.5)	263 (11.0)	
Smoking, n (%)								
Never	1010 (40.0)	769 (39.9)	157 (46.0)	0.09	170 (41.6)	812 (40.8)	951 (39.9)	0.74
Ever	1514 (60.0)	1158 (60.1)	184 (54.0)		239 (58.4)	1178 (59.2)	1432 (60.1)	
Physical activity, n (%)								
Inactive	216 (8.6)	126 (6.5)	33 (9.7)	0.09	39 (9.5)	160 (8.0)	178 (7.5)	0.13
Moderate	1006 (39.9)	785 (40.7)	131 (38.4)		152 (37.2)	767 (38.5)	994 (41.7)	
Active	1302 (51.6)	1016 (52.7)	177 (51.9)		218 (53.3)	1063 (53.4)	1211 (50.8)	
Education, n (%)								
Primary or below	724 (28.7)	546 (28.3)	102 (29.9)	1.00	109 (26.7)	589 (29.6)	674 (28.3)	0.29
Junior middle	769 (30.5)	587 (30.5)	106 (31.1)		121 (29.6)	600 (30.2)	734 (30.8)	
Senior middle	621 (24.6)	482 (25.0)	81 (23.8)		112 (27.4)	505 (25.4)	567 (23.8)	
College	410 (16.2)	312 (16.2)	52 (15.2)		67(16.4)	296(14.9)	408(17.1)	
Occupation, n (%)								
Manual	1285 (50.9)	931(48.3)	188(55.1)	0.12	217(53.1)	1028(51.7)	1153(48.4)	0.040
Nonmanual	973 (38.5)	768(39.9)	120(35.2)		156(38.1)	731(36.7)	970(40.7)	
Others	266 (10.5)	228 (11.8)	33 (9.7)		36 (8.8)	231 (11.6)	260 (10.9)	
Age, year (s.d.)	63.6 (6.9)	63.5 (6.9)	63.6 (6.9)	0.95	63.4 (6.9)	63.6 (7.0)	63.5 (6.9)	0.66
Body mass index, kg m ⁻² (s.d.)	23.5 (3.2)	23.5 (3.1)	23.7 (3.0)	0.56	23.5 (3.0)	23.5 (3.1)	23.5 (3.2)	0.95
Total cholesterol, mmol l ⁻¹ (s.d.)	5.59 (1.06)	5.57 (1.08)	5.52 (0.92)	0.49	5.45 (0.99)	5.57 (1.07)	5.60 (1.07)	0.036
Triglycerides, mmol l ⁻¹ (s.d.) ^b	1.43 (1.73)	1.40 (1.73)	1.40 (1.70)	0.35	1.42 (1.67)	1.40 (1.71)	1.43 (1.75)	0.35
HDL cholesterol, mmol l ⁻¹ (s.d.)	1.52 (0.37)	1.51 (0.37)	1.50 (0.35)	0.82	1.47 (0.35)	1.50 (0.37)	1.53 (0.38)	0.005
Fasting glucose, mmol l ⁻¹ (s.d.) ^b	5.59 (1.21)	5.59 (1.22)	5.57 (1.20)	0.92	5.57 (1.21)	5.58 (1.21)	5.60 (1.22)	0.68

Abbreviations: ADH, alcohol dehydrogenase; ALDH, aldehyde dehydrogenase.

^aHardy–Weinberg equilibrium test.^bData was expressed as geometric means (s.d.).

Table 2 Association of ADH2 and ALDH2 genotypes with blood pressure in older Chinese men

				Adjusted means ^a (95% CI)				
				n	Systolic BP	Diastolic BP	PP	MAP
Never drinkers								
ADH2(rs1229984)				2467				
Group 1	AA			1350	132.6 (131.3, 133.9)	75.2 (74.5, 75.9)	57.4 (56.4, 58.3)	94.3 (93.5, 95.2)
Group 2	AG			955	133.0 (131.5, 134.5)	75.0 (74.2, 75.8)	57.9 (56.9, 59.0)	94.3 (93.4, 95.3)
Group 3	GG			162	131.4 (128.3, 134.5)	73.3 (71.7, 75.0)	58.0 (55.9, 60.2)	92.7 (90.7, 94.7)
P for trend					0.79	0.09	0.36	0.28
ALDH2 (rs671)				2457				
Group 1	AA			315	132.2 (129.9, 134.5)	74.5 (73.3, 75.8)	57.7 (56.0, 59.3)	93.7 (92.3, 95.2)
Group 2	AG			1173	132.9 (131.5, 134.3)	75.1 (74.3, 75.8)	57.8 (56.9, 58.8)	94.3 (93.4, 95.2)
Group 3	GG			969	132.7 (131.2, 134.1)	75.1 (74.3, 75.9)	57.5 (56.5, 58.6)	94.3 (93.4, 95.3)
P for trend					0.91	0.44	0.66	0.62
Interaction				2450				
Group 1	ADH2	ALDH2	AA + AG	1397	132.8 (131.5, 134.1)	75.1 (74.4, 75.8)	57.7 (56.8, 58.6)	94.4 (93.5, 95.2)
Group 2	ADH2	ALDH2	AA + AG	891	132.7 (131.2, 134.2)	75.2 (74.4, 76.0)	57.5 (56.4, 58.5)	94.4 (93.4, 95.4)
Group 3	ADH2	ALDH2	Any genotype	162	131.4 (128.3, 134.5)	73.3 (71.7, 75.0)	58.0 (55.9, 60.2)	92.7 (90.7, 94.7)
P for trend					0.42	0.21	0.85	0.27
Current drinkers								
ADH2 (rs1229984)				2044				
Group 1	AA			1035	133.2 (131.2, 135.1)	77.4 (76.4, 78.4)	55.8 (54.4, 57.1)	96.0 (94.8, 97.2)
Group 2	AG			850	133.2 (131.2, 135.1)	77.0 (76.1, 78.1)	56.1 (54.8, 57.5)	95.8 (94.5, 97.0)
Group 3	GG			159	136.7 (133.3, 140.1)*, #	78.0 (76.2, 79.8)	58.7 (56.4, 61.0)*, #	97.6 (95.4, 99.7)
P for trend					0.15	0.97	0.039	0.44
ALDH2 (rs671)				2044				
Group 1	AA			85	130.7 (126.0, 135.3)	76.1 (73.6, 78.5)	54.6 (51.4, 57.8)	94.3 (91.3, 97.3)
Group 2	AG			717	133.2 (131.1, 135.3)	76.7 (75.6, 77.8)	56.6 (55.1, 58.0)	95.5 (94.2, 96.9)
Group 3	GG			1242	133.5 (131.7, 135.3)	77.5 (76.6, 78.5)	56.0 (54.7, 57.2)	96.2 (95.0, 97.4)
P for trend					0.35	0.06	0.91	0.13
Interaction				2036				
Group 1	ADH2	ALDH2	AA + AG	730	132.6 (130.5, 134.7)	76.6 (75.4, 77.7)	56.0 (54.6, 57.5)	95.2 (93.9, 96.6)
Group 2	ADH2	ALDH2	AA + AG	1147	133.4 (131.6, 135.3)	77.6 (76.6, 78.6)*	55.8 (54.6, 57.1)	96.2 (95.0, 97.4)
Group 3	ADH2	ALDH2	Any genotype	159	136.6 (133.3, 140.0)*	77.9 (76.1, 79.7)	58.7 (56.4, 61.0)*, #	97.5 (95.3, 99.7)*
P for trend					0.035	0.035	0.15	0.025

Abbreviations: ADH, alcohol dehydrogenase; ALDH, aldehyde dehydrogenase, BP, blood pressure; CI, confidence interval; HDL, high-density lipoprotein; PP, pulse pressure; MAP, mean arterial blood pressure.

Compared with group 1: * $P < 0.05$. Compared with group 2: # $P < 0.05$.

^aAdjustment for age, education, occupation, smoking, physical activity, body mass index, as well as fasting total cholesterol, triglyceride, HDL cholesterol, glucose and frequency of drinking (for current drinkers only).

use instead of frequency of drinking showed similar results (Table not shown).

Table 3 shows that among ever drinkers, men with the GG ADH2 genotype had higher odds of hypertension than those with the (AA/AG ADH2 + AA/AG ALDH2) genotype after adjusting for frequency of alcohol drinking and other risk factors (OR = 1.62, 95% confidence interval 1.15–2.28). Sensitivity analysis found similar results after adjusting for amount of alcohol drinking and other risk factors (OR = 1.61 (1.14–2.27)). Compared with (AA/AG ADH2 + GG ALDH2), the adjusted OR for GG ADH2 was 1.40 (1.01, 1.96). Among men with the AA/AG ADH2 genotype, the ALDH2 genotype was not significantly associated with hypertension. (Table not shown) Among never drinkers, ADH2 or ALDH2 genotypes were not associated with hypertension.

Sensitivity analysis with adjustment for interaction of ADH2/ALDH2 genetic polymorphism with gender, age and antihypertensive medication use also showed similar results. The interactions between ADH2/ALDH2 genetic polymorphism and these potential effect

modifiers, such as gender, age and antihypertensive medication use on blood pressure levels or hypertension, were all nonsignificant (P -values from 0.21 to 0.94).

DISCUSSION

To the best of our knowledge, this is the first study in which the association of ADH2 and ALDH2 genotypes with hypertension has been investigated in a large community-based Chinese sample. ADH2 genotype but not ALDH2 genotype was a determinant of blood pressure among alcohol drinkers; alcohol drinkers with the GG ADH2 genotype had higher blood pressure than alcohol drinkers with other genotypes, after adjusting for frequency or amount of alcohol use, suggesting that ethanol may have a key role in alcohol-induced hypertension.

Our results support studies that showed a significant interaction between ADH2 and alcohol drinking on blood pressure²² and no relationship between ALDH2 and blood pressure.^{22–24} As ALDH2 has been shown to be associated with blood pressure through affecting

Table 3 Adjusted ORs (95% confidence interval) of hypertension for ADH2 and ALDH2 polymorphisms in older Chinese men

ADH2 (rs1229984)	ALDH2 (rs671)	Never drinkers			Ever drinkers				
		n	Hypertension (%)	Model 1	n	Hypertension (%)	Model 1	Model 2 ^a	Model 3 ^a
AA + AG	AA + AG	1397	43.5	1.00	829	39.1	1.00	1.00	1.00
AA + AG	GG	891	45.5	1.09 (0.91, 1.31)	1309	43.9	1.21 (1.01, 1.46)*	1.15 (0.95, 1.39)	1.15 (0.96, 1.40)
GG	Any type	162	42.0	0.84 (0.59, 1.21)	179	51.4	1.66 (1.18, 2.34)**	1.62 (1.15, 2.28)**	1.61 (1.14, 2.27)**
P-value			0.56	0.33	0.005	0.008	0.020	0.021	
P for trend			0.75	0.98	0.001	0.002	0.008	0.008	

Abbreviations: ADH, alcohol dehydrogenase; ALDH, aldehyde dehydrogenase; OR, odds ratio; HDL, high-density lipoprotein.

Model 1: adjusting for age, education, occupation, smoking, physical activity, body mass index, as well as fasting total cholesterol, triglyceride, HDL cholesterol and glucose. Model 2: based on model 1, additionally adjusting for frequency of drinking. Model 3: based on model 1, additionally adjusted for the amount of alcohol use (< or ≥140 g per week).

*; $P < 0.05$; **, $P < 0.01$.

^aCompared with (AA/AG ADH2 + GG ALDH2), OR in GG ADH2 = 1.40 (1.01, 1.96), $P = 0.045$ in model 2 and 1.40 (1.00, 1.94), $P = 0.049$ in model 3.

alcohol drinking behavior,⁵ the association between ALDH2 and blood pressure became statistically nonsignificant after taking into account the frequency or amount of alcohol consumption in our study. As the ALDH2 genotype influences the blood acetaldehyde level, whereas the ADH2 genotype influences the blood ethanol level after alcohol drinking as in Figure 1, our results suggest that the hypertensive effect of alcohol drinking is due to ethanol rather than acetaldehyde. This finding is consistent with some but not all previous studies. In an earlier study on 898 middle-aged male Japanese, no association was found between blood pressure and ADH2 or ALDH2 genotypes after adjusting for age, body mass index and volume of alcohol consumption.⁶ However, as the sample size was small, the null association in this study should be interpreted with caution. Another study of 335 Japanese showed that the relationship between alcohol and blood pressure was significantly stronger in those with the GG ADH2 genotype than those with the AA or AG genotype.¹⁶ Moreover, the authors also indicated that those with the ADH2 GG genotype were likely to develop hypertension because of excessive alcohol use. However, in our study, we have also controlled for frequency or amount of alcohol use in the model and found that older male drinkers with the GG ADH2 genotype had a higher blood pressure than alcohol drinkers with other genotypes. To remove the confounding effect of ADH2 genotype, we analyzed the association between ALDH2 genotypes and blood pressure in men with the AA/AG ADH2 genotype and found no associations (table not shown), which was in accordance with previous studies^{6,16} showing that ADH2 genotype, but not ALDH2 genotype, modifies the association of alcohol with blood pressure. However, in a study of excessive male drinkers (> 300 g of ethanol per week), the prevalence of the ADH2 AA genotype was significantly greater in those with higher SBP,²² suggesting that the ADH2 AA genotype but not GG genotype weakens the relationship between alcohol and blood pressure. However, participants in this study were hospital workers, included only a few heavy drinkers and had a mean age of 46 years, and the frequency or amount of alcohol use was not adjusted in the model. A confounding effect from excessive alcohol use could not be ruled out.

In the present study, we found that the ADH2 GG genotype was associated with hypertension in alcohol drinkers after adjustment for frequency of alcohol use. Hence, the relationship between ADH2 and hypertension could not be fully explained by different levels of alcohol consumption. Ethanol accumulation might be considered a key pathway. The possible mechanisms by which ethanol increases blood

pressure include: (a) a direct effect on the central nervous system, by interfering with central inhibitory pathways, which control the vasomotor center or increasing sympathetic activity,²⁵ (b) activation of the rennin-angiotensin system by ethanol,²⁶ (c) aortic endothelial oxidative injury and downregulation of the nitric oxide generating system,²⁷ and (d) magnesium deficiency.²⁸ However, the actual effect of ethanol on blood pressure is unclear. Further experimental studies are warranted to examine the mechanism underlying the relationship between blood ethanol concentration and blood pressure, taking into account different ADH2 genotypes.

Several limitations of this study should be noted. First of all, as subjects in the present study were older Chinese people from Southern China, most of them were light to moderate drinkers. A 2001 World Health Organization sponsored survey of 24 992 Chinese residents across five areas of China showed an annual ethanol consumption of 4.5 l (s.d. = 10.9) among adults aged 15 years and older, which was low compared with Western countries: by comparison, annual alcohol use was about 8.6 l per person in 2001 in Europe.²⁹ Moreover, people living in Southern China have lower levels of alcohol use than those in the North.² Given the limited number of subjects in the present study who were heavy alcohol users, whether the effect of heavy drinking on hypertension varies with ADH2 genotype could not be assessed. Secondly, levels of alcohol use among Chinese women were too low, so we could not examine the associations among women. Thirdly, this is not a fully population-representative study, and participants included were homogenous Southern Chinese from one city. Earlier studies have shown different distributions of ADH2 genotypes in different ethnic populations, such as Chinese, Japanese, Brazilians and African Americans.^{8,30,31} Further studies are needed to examine whether ADH2 genotypes modify the effect of alcohol on blood pressure in other ethnic groups. The strengths of the present study include a large sample size (which enabled a detailed analysis of gene-environment interactions by assessing the alcohol-blood pressure relationship in participants who were or were not alcohol drinkers), the focus on a community-based sample with light to moderate level of alcohol use, and the adjustment of many potential confounding factors. Moreover, the current study, for the first time systematically considered ethanol metabolism in human body, including both acetaldehyde and acetic acid, and clearly showed that current drinkers with an inactive ADH2 genotype had a significantly increased risk of hypertension. The current study employed genetic polymorphism of an alcohol metabolism enzyme as an instrumental variable to obtain an unbiased

estimation of the effect of alcohol on blood pressure, which was unlikely to be confounded by other risk factors, such as smoking or sodium intake, as in other observational studies. This study provided the suggestive causal inference that the hypertensive effect of alcohol drinking might be due to ethanol accumulation.

In conclusion, we found that the ADH2 genotype but not the ALDH2 genotype was associated with blood pressure and hypertension in current drinkers. However, in never drinkers, no association between ADH2 or ALDH2 genotypes and blood pressure was found, indicating a gene-environment interaction effect on alcohol-induced hypertension. Among current alcohol drinkers, individuals with a GG ADH2 genotype had a higher risk of hypertension than those with the AA or AG ADH2 genotype, suggesting a key role for ethanol in alcohol-induced hypertension. Further studies in a larger sample, including woman drinkers and younger drinkers, are warranted to confirm our results.

CONFLICT OF INTEREST

The authors declare no conflict of interest

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