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. 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 1996–2001.4 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. 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). and have shown important roles for these enzymes in the metabolism of alcohol in the body. 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. 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. The AA ALDH2 genotype results in higher blood acetaldehyde levels after alcohol intake, leading to facial flushing, palpitations and nausea. 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. 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
AA and AG genotypes have about 200 and 100 times higher ethanol oxidizing capacity, respectively, than the GG ADH2 genotype,\textsuperscript{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,\textsuperscript{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).

\section*{METHODS}

\subsection*{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.

\subsection*{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.\textsuperscript{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: 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,\textsuperscript{16} and the last two measurements were averaged for analysis.

Blood samples were collected following 8–10h 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.\textsuperscript{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 spectrometry.

\subsection*{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,\textsuperscript{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 × (SBP + DBP × 2) and hypertension defined as use of antihypertensive medication or SBP/DBP > 140/90 mmHg.

\subsection*{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.

\subsection*{Statistical analysis}

We used genetic polymorphism of alcohol metabolism enzyme (ADH2 and ALDH2 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),\textsuperscript{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.\textsuperscript{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 48.8%, 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.

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 ADH2) 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...
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 + GG 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 pressure and no relationship between ALDH2 and blood pressure. As ALDH2 has been shown to be associated with blood pressure through affecting
which was in accordance with previous studies showing that the ADH2 genotype and found no associations (table not shown), confounding effect of ADH2 genotype, we analyzed the association between ADH2 genotype and blood pressure in alcohol drinkers with the GG ADH2 genotype had a higher blood pressure than those with the AA or AG genotype.16 Moreover, the authors also indicated that those with the ADH2 GG genotype had a significantly increased risk of hypertension (1.40 (1.01, 1.96), see Table 3).

The current study employed genetic polymorphism of an alcohol metabolism enzyme as an instrumental variable to obtain an unbiased estimate of the association between alcohol drinking and blood pressure. However, the strength of the current study is limited by the fact that the sample size was small, the null association in this study would 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 heavy drinkers (which enabled a detailed analysis of gene–environment interactions 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 ethanol 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 estimate of the association between alcohol drinking and blood pressure.
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 women drinkers and younger drinkers, are warranted to confirm our results.

CONFLICT OF INTEREST
The authors declare no conflict of interest

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