Title
Single nucleotide polymorphisms of one-carbon metabolism and cancers of the esophagus, stomach, and liver in a Chinese population.

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Introduction

Upper gastrointestinal (GI) cancers are major causes of morbidity and mortality throughout the world. Based on GLOBOCAN 2012 estimates, stomach, liver, and esophageal cancers are the fifth, sixth, and eighth most common cancers, respectively, with a global incidence of approximately 2,189,829 new cancer cases (15.5% of the total), and 1,880,328 deaths (22.8% of the total) [1]. The majority of these cancer cases (1,694,874 cases, 77.4%) occur in less developed countries. China alone accounts for almost half of all incident GI cancers (1,023,072 cases, 46.7%) [1].

Continued research regarding the involvement of single nucleotide polymorphisms (SNPs) in the etiology of these three upper GI cancers has been fruitful. Of particular interest are the SNPs located within genes involved in folate metabolism [2–4]. Folate maintains DNA stability by regulating DNA biosynthesis, DNA repair and DNA methylation [5]. Neoplasms may develop when this pathway is disregulated by the depletion of micronutrients or through the incorporation of polymorphisms [5]. Several enzymes are involved in one-carbon metabolism, including the methylenetetrahydrofolate reductase (MTHFR), methionine synthase (MTR), and methionine synthase reductase (MTRR), DNA methyltransferases (DNMTs), and mitochondrial aldehyde dehy-
dihydrogenase 2 (ALDH2), MTHFR, MTR, and MTRR are involved in DNA synthesis, and the generation of S-adenosylmethionine (SAM)—a universal methyl-donor for methylation reactions. DNMTs catalyze DNA methylation and replicate methylation patterns. ALDH2 is responsible for metabolizing acetaldehyde generated during alcohol metabolism. Alcohol and acetaldehyde can inhibit folate absorption and impair DNA methylation [6].

The role of folate and one-carbon metabolism in upper GI cancers is not fully understood. Animal studies provided some evidence for an effect of low folate levels in oxidative stress, DNA methylation, and hepatocarcinogenesis [7,8]; while high folate intake can increase global DNA methylation and reduce gastric cancer risk [9,10]. Epidemiologic studies have suggested that genetic polymorphisms of genes in one-carbon metabolic pathway might modulate the risk of esophageal and gastric cancer [4]. However, published results are inconclusive and limited in terms of the number of genes/polymorphisms being investigated. Possible modification by related micronutrients and known risk factors has seldom been explored. Therefore, considering the importance of one-carbon metabolism in upper GI cancer development, we examined the associations between eight SNPs in genes in one-carbon metabolic pathway and cancers of the esophagus, stomach, and liver in a Chinese population. We also evaluated heterogeneity of the associations across different strata of plasma micronutrients (including folate, vitamin B12, and total homocysteine) and known risk factors for these cancers.

Materials and Methods

Ethics statement

This study was exempted by the institutional review board of the University of California at Los Angeles (Certified Exempt 02-248).

Study Design and Population

A detailed description of the study design has been published previously [11,12]. Briefly, this was a population-based case-control study conducted in Taixing City, Jiangsu Province, China. Eligible cases were newly diagnosed patients with pathologically or clinically confirmed esophageal cancer (between June 1 and December 30, 2000), stomach cancer (between June 1 and December 30, 2000), and liver cancer (between January 1 and June 30, 2000) reported to the Taixing CDC Tumor Registry. Other inclusion criteria including being 20 years of age or older, in stable medical condition as determined by a physician, residency in Taixing for 10 years or more, and willingness to participate. A total of 218 esophageal cancer cases, 206 stomach cancer cases, and 204 liver cancer cases participated, representing 67, 65 and 57%, respectively, of all newly diagnosed cancer patients.

Controls were randomly selected among healthy residents of Taixing City with a 2:3 frequency matching ratio to the combined case group on 5-year age categories (20–24 to 80–84), sex, and residency (village in rural township or in an urban residential block in central Taixing City). There are 23 townships (rural areas) and one central town (urban area) in Taixing City. Each rural township consists of 10–12 villages, and the central urban area consists of 10–12 residential blocks. Other inclusion criteria were the same as the cases. A total of 464 potential controls were approached, and 415 (89.4%) consented to participate.

Epidemiologic data collection

All of the recruited cases and controls completed a standard questionnaire administered by trained interviewers. Interviews took place either at the participants’ homes, in the hospitals (for cases), or in the county doctor’s office (for controls). Cancer cases were usually interviewed within 6 months of diagnosis. The questionnaire collected detailed information on demographic factors, current height and weight, dietary history, tobacco smoking history, alcohol drinking history, tea drinking habits, occupational history, family history of cancers, and physical activities.

Laboratory assays

Each study participant provided a 5-ml peripheral blood sample after their interviews. DNA was isolated from blood clots, using the phenol-chloroform method. Hepatitis B virus surface antigen (HBsAg), IgG antibodies for hepatitis C virus (HCV), and IgG antibodies for CagA-Helicobacter pylori (H. pylori) were measured by enzyme-linked immunosorbant assay (ELISA) using kits from the Reagent Company of the Shanghai Hospital for Infectious Diseases (Shanghai, China), the Shanghai Huamei Biological Company (Shanghai, China), and the Reagent Company of the Shanghai Biotechnology Industry Park (Pudong, Shanghai, China), respectively. Plasma aflatoxin B1 (AFB1)-albumin adduct levels were determined by ELISA assay, as previously described [13], using free aflatoxin (Supelec) for the aflatoxin standards. A comparison between free and bound aflatoxin standards revealed a log-linear relationship, allowing us to estimate the absolute values of the samples. Plasma folate and vitamin B12 levels were measured using a competitive radioassay with iodine 125-labeled folate and cobalt 57-labeled vitamin B12 as tracers (Quanthase II B12/folate radiobinding kit, Bio-Rad, CA). Plasma total homocysteine (tHcy) levels were measured using a commercially available chemiluminescent immunoassay system (IMMULITE 1000 Automated Analyzer, DPC, Los Angeles, CA).

We selected eight SNPs from MTHFR, MTR, MTRR, DNMT1, and ALDH2 genes, based on the following criteria: 1) SNPs which are functional or potentially functional (SNPs located in the coding, 3'- and 5'-untranslated regions); 2) SNPs previously reported to be associated with upper GI cancers; and 3) SNPs with minor allele frequency of at least 5% in the National Center for Biotechnology Information SNP database. Genotyping was performed using the TaqMan (MTR rs1805087, MTR rs1532268/rs1801394, and ALDH2 rs886205) or the SNPlex (DNMT1 rs2228612 and ALDH2 rs2238151) assay, as previously described [14], or the PCR-RFLP analysis (MTHFR rs1801133 and ALDH2 rs671) modified from previously published methods [15,16]. Genotyping call rates were over 97% for TaqMan and PCR-RFLP methods, and over 80% for the SNPlex assay. Reproducibility was 98% for the SNPlex assay (3% random duplicate samples) [17], and 100% for the TaqMan assay (10% random duplicate samples).

Statistical analysis

We used Pearson’s chi-square test for Hardy-Weinberg equilibrium (HWE) for the distributions of genotype frequencies of the eight SNPs in the controls only. Testing for HWE among the controls is a commonly used preliminary quality-control method in genetic association studies to identify systematic genotyping errors in unrelated individuals. We analyzed each SNP-cancer association under co-dominant, log-linear, dominant, and recessive genetic models, using unconditional logistic regression models to calculate odds ratios (ORs) and 95% confidence intervals (CIs). Models included age-matched categories, sex, residency (urban/rural), education (illiteracy/primary school/higher than middle school), body mass index (BMI, continuous), smoking pack-years (continuous), alcohol consumption frequency (never/occasionally/often/everyday), H. pylori infection (stomach cancer; negative/
positive), HBsAg status (liver cancer; negative/positive) and plasma AFB1-albumin adduct levels in quintiles (liver cancer; estimated quintile: <222.7, 222.7–344.2, 344.2–442.6, 442.6–588.3, and >588.5 fmol/mg). To adjust for residual confounding effects from age, we also included the deviation of each person’s age from the mean age in each age category [18]. We caution that a number of adjustment variables may be affected by genetic variations, as these variables occur afterward. At best, our estimates are for direct genotype effects, and otherwise may be over-adjusted or confounded by uncontrolled factors that affect both the adjustment variables and the outcomes [19]. Thus, we checked estimates for direct genotype effects against estimates adjusted only for age and sex.

We further conducted stratified analyses to check heterogeneity across strata of micronutrients or modifiable risk factors, including plasma micronutrients (folate, vitamin B12, and tHcy), smoking status, alcohol consumption, H. pylori infection (stomach cancer), HBsAg status (liver cancer), and plasma AFB1 levels (liver cancer). We used estimated median levels in controls to dichotomize plasma levels of folate (12.76 nmol/l), vitamin B12 (228.88 pmol/l), tHcy (9.5 pmol/l), and AFB1 (388.95 fmol/mg). We used the dominant genetic model, which assumed that the effect of the variant allele is dominant if the ratio of the ORs comparing variant allele homozygotes to heterozygotes was smaller than that comparing heterozygotes to common allele homozygotes; otherwise we used the recessive genetic model. We assessed heterogeneity across strata using likelihood ratio tests by comparing models with and without product terms. To reduce the risks of multiple-comparison artefacts and sparsedata bias, we used a semi-Bayes (SB) shrinkage (penalized-likelihood) method to estimate genotype coefficients [20]; the odds-ratio estimates we report are the antilogs of these coefficients. Shrinkage estimation has been recommended extensively as an alternative superior to Bonferroni in the statistical literature for eliminating multiple-testing artefacts in comparative studies [21–24]. In shrinkage estimation, instead of changing the alpha level, we regress (‘shrink’) the estimates toward zero to a degree proportional to their estimated variances and inversely proportional to the prior variances v. The prior variance plays a role analogous to the adjusted α-level, in that smaller values correspond to more stringent rejection/detection criteria, with α = 0 and v = 0 being the lower limits of adjustment at which rejection of the null becomes impossible. At the other extreme, no adjustment occurs when using the original value of α or a huge (effectively infinite) value for v.

In our study, we assigned a prior variance of 0.50, and a prior median OR = 1 (no association) which results in a 95% prior probability of falling within the interval 0.25, 4. This pulls the observed associations toward the null to the degree that would result if there had been a previous null experiment observing 4/ 8 cases total and it had been merged with the current data [20,25]. When differing stratum-specific SNP effects were allowed, such as in stratified analyses, the prior variance was reduced to 0.25, which corresponds to a variance of 0.50 for the coefficient of the stratum-SNP product (interaction). For each SB posterior estimate, we further provide the directional (one-sided) SB P-values, which equal the posterior probability that the point estimate is on the wrong side of the null under the fitted model and the shrinkage priors [26,27].

To summarize the associations of the 8 SNPs for each of the three upper GI cancers, we constructed a polygenic risk score (PRS) [28]. The PRS was calculated as the weighted sum of the risk genotype (under either dominant or recessive model as in the stratified analyses) counts, where the weight for each SNP was determined by the semi-Bayes log OR of its association with each cancer in the fully adjusted model. PRS was only estimated among those with complete genotype data on all of the 8 SNPs, which include 126 esophageal cancer cases, 125 stomach cancer cases, 142 liver cancer cases, and 287 controls. The range (maximum minus minimum) of PRS for each cancer was divided into three equally spaced categories; these ranges were 0.11 to 2.05 for esophageal cancer, 0 to 1.91 for stomach cancer, and 0 to 1.40 for liver cancer. Data analyses were performed using SAS 9.1.3 (SAS Institute, Cary, NC).

Results

Compared to population controls, cancer cases tended to be smokers, had lower BMI, and lower education levels (Table 1). Esophageal and stomach cancer cases were older than the controls, while liver cancer cases were the youngest. Liver cancer patients had the highest male-to-female ratio of 3.53, and were most likely to consume alcohol; esophageal cancer patients drank more frequently than the other cancer cases and controls in this study. For risk factors specific to each cancer site, we did not observe differing frequency of H. pylori infection between stomach cancer patients and controls. Compared with controls, liver cancer patients showed a higher percentage of HBsAg positive (65 vs. 25%), anti-HCV positive (9 vs. 3%), and had higher plasma AFB1-albumin adduct levels (30 vs. 20% in the 5th quintile).

Table 2 presents the SB odds-ratio estimates (SBOR) for each SNP-cancer association of the eight SNPs; Table S1, S2, S3, S4, S5, S6 shows stratified associations and Figure 1 summarizes selected results. Genotype distributions among controls appeared compatible with Hardy-Weinberg equilibrium, except possibly for DNMT1 rs2228612, which had P = 0.010, below the traditional alpha level of 0.05, but larger than the Bonferroni-adjusted alpha level of 0.05/8 = 0.006 (testing all eight SNPs). However, we note that matching may bias controls away from equilibrium if the matching factors are associated with both the SNPs and cancer.

We have previously reported positive associations of the T allele of MTHFR rs1801133 with stomach and liver cancer [11,12]. In the present analysis, these associations remained apparent after confounder adjustment and SB shrinkage (any T versus C/C, fully adjusted SBOR: 1.79, 95% posterior limits: 1.18, 2.71 for stomach cancer; SBOR: 1.51, 95% posterior limits: 0.98, 2.32 for liver cancer). In stratified SB analyses, the association between MTHFR rs1801133 and stomach cancer appeared stronger among individuals who had lower plasma folate levels, higher plasma vitamin B12 or tHcy levels, and among smokers (Figure 1). There was no clear association of MTHFR rs1801133 with esophageal cancer (Table 2 and Tables S1, S2, S3, S4, S5).

While there was no clear overall association between SNPs in MTR and MTRR and any cancer in main effect analyses (Table 2), heterogeneity of association was suggested in stratified analyses on alcohol consumption, including associations of MTR rs1805087 with liver cancer (homogeneity P = 0.021), and MTRR rs1801394 with both esophageal (homogeneity P = 0.005) and stomach cancer (homogeneity P = 0.004). While G allele carriers of MTR rs1805087 were inversely associated with liver cancer among non-drinkers (SBOR: 0.57, 95% posterior limits: 0.31, 0.84), they were positively associated with liver cancer among drinkers (SBOR: 1.48, 95% posterior limits: 0.85, 2.57) (Figure 1). Similarly, G allele carriers of MTRR rs1801394 were inversely associated with esophageal and stomach cancer among non-drinkers (SBOR: 0.39, 95% posterior limits: 0.37, 0.94 for esophageal cancer; SBOR: 0.49, 95% posterior limits: 0.30, 0.79 for stomach cancer) but positively associated with cancer among...
Table 1. Distribution for selected demographic and health characteristics comparing esophageal, stomach, and liver cancer cases with controls.

|                               | Controls | Esophageal Cancer | Stomach Cancer | Liver Cancer |
|-------------------------------|----------|------------------|----------------|--------------|
| (N, %)                        | (N = 218) | (N = 218)        | (N = 206)      | (N = 204)    |
| Age*, mean ± SD               | 60.6 ± 9.6 | 60.6 ± 9.6       | 62.8 ± 9.8     | 53.9 ± 13.0  |
| Sex*                          | 128 (30.8) | 77 (35.3)        | 68 (33.0)      | 45 (22.1)    |
| BMI, mean ± SD                | 22.4 ± 2.6 | 21.9 ± 2.8       | 21.4 ± 2.7     | 21.3 ± 2.7   |
| Education                     | Illiterate | 73 (17.6)        | 83 (38.6)      | 66 (32.0)    |
| Middle school or higher       | 200 (48.2) | 31 (14.4)        | 33 (16.0)      | 83 (40.7)    |
| Smoking                       | Never     | 217 (52.4)       | 94 (44.6)      | 85 (43.3)    |
| Alcohol drinking              | Occasionally | 72 (17.5)       | 18 (8.5)       | 31 (15.4)    |
| H. pylori CagA status         | +         | 251 (68.8)       | 130 (64.7)     | 25 (13.0)    |
| HBsAg status                  | +         | 102 (24.6)       | 132 (64.7)     | <0.001       |
| Anti-HCV status               | +         | 12 (2.9)         | 18 (9.0)       | 0.001        |
| Plasma AFB1-albumin adduct    | 1         | 75 (19.9)        | 26 (14.4)      | 0.054        |
| levels (quintile)             | 2         | 76 (20.2)        | 36 (19.9)      |              |
|                               | 3         | 75 (19.9)        | 36 (19.9)      |              |
drinkers (SBOR: 1.56, 95% posterior limits: 0.95, 2.56 for esophageal cancer; SBOR: 1.39, 95% posterior limits: 0.83, 2.32 for stomach cancer) (Figure 1).

For DNMT1 polymorphism, rs2228612 was inversely associated with esophageal cancer in the dominant genetic model (any G versus A/A, SBOR: 0.60, 95% posterior limits: 0.39, 0.94) (Table 2). Among three ALDH2 SNPs, rs671 was associated with esophageal cancer in the recessive genetic model (A/A versus any G, SBOR: 1.76, 95% posterior limits: 0.96, 3.24). In stratified adjusted analyses, ALDH2 rs671 appeared associated with esophageal cancer among individuals with lower plasma folate levels (A/A versus any G, SBOR: 2.12, 95% posterior limits: 1.01, 4.44) (Figure 1). The ALDH2 rs2238151 appeared inversely associated with liver cancer when comparing T allele carriers to those with the C/C genotype (age and sex-adjusted SBOR: 0.47, 95% posterior limits: 0.24, 0.92). While we did not find associations between ALDH2 rs886205 and cancer susceptibility in main effect analyses, stratum-specific SBOR suggested that ALDH2 rs886205 was positively associated with stomach cancer among participants with higher plasma vitamin B12 levels (SBOR: 1.87, 95% posterior limits: 1.09, 3.20) (Figure 1).

Except for analysis on single SNP models, we also did joint SNPs analysis by including all of the 8 SNPs in a model (Table 3). The results from joint SNPs analysis suggested similar associations as in the single SNP models, but the 95% posterior intervals were wider.

The analysis on PRS suggested roughly a doubling of odds for esophageal and liver cancers among individuals in the highest PRS category compared to those in the lowest category (SBOR: 2.06; 95% posterior limits: 1.13, 3.77 for esophageal cancer and SBOR: 2.09, 95% posterior limits: 1.05, 4.17 for liver cancer), with somewhat less consistency across categories for stomach cancer. In the continuous PRS analysis, the results suggested a doubling of odds for these three upper GI cancers with one unit (in log OR) increase of PRS (Table 4). We caution however that PRS analyses do not account for the score construction from the data, and thus may overestimate effects and underestimate variability in the resulting estimates.

**Discussion**

We examined the associations between eight SNPs in genes involved in the one-carbon metabolic pathway and susceptibility of esophageal, stomach, and liver cancers in a Chinese population. After applying SB shrinkage methods and controlling for potential confounders, we observed that any T genotype of MTHFR rs1801133 was positively associated with both stomach and liver cancer. We also found an inverse association between the variant G allele of DNMT1 rs2228612 and esophageal cancer. In addition, our study suggested potential OR variations across strata of alcohol consumption, including associations of MTR rs1801394 with esophageal and stomach cancer, and MTR rs1805087 with liver cancer. The odds for upper GI cancers were roughly doubled for Chinese participants with one unit (in log OR) increase of PRS.

In one-carbon metabolism, MTHFR irreversibly catalyzes the conversion of 5,10-methylenetetrahydrofolate (5,10-methyleneTHF) to 5-methyltetrahydrofolate (5-methylTHF). The 5,10-methyleneTHF is essential in purine and thymidilate synthesis, and 5-methylTHF is a co-substrate for remethylation of homocysteine to methionine, which is further converted to SAM for methylation reactions [5]. The MTHFR C677T (rs1801133) polymorphism, which results in an alanine to valine substitution, leads to reduced MTHFR enzyme activity [29], decreased 5-
|                | Esophageal Cancer | Stomach Cancer | Liver Cancer |
|----------------|-------------------|----------------|-------------|
|                | Ca/Co             | SBOR¹ (95% posterior limit) | One-sided P² | Ca/Co | SBOR¹ (95% posterior limit) | One-sided P² | Ca/Co | SBOR¹ (95% posterior limit) | One-sided P² |
| **MTHFR**      |                   |                 |             |       |                 |             |       |                 |             |
| rs1801333      |                   |                 |             |       |                 |             |       |                 |             |
| CC             | 65/135            | 1.00            | 1.00        | 50/135 | 1.00            | 1.00        | 50/135 | 1.00            | 1.00        |
| CT             | 105/199           | 1.15 (0.79, 1.65) | 0.24 | 1.19 (0.80, 1.77) | 0.20 | 106/199 | 1.38 (0.94, 2.02) | 0.050 | 1.53 (1.00, 2.33) | 0.024 |
| TT             | 32/57             | 1.13 (0.69, 1.86) | 0.32 | 1.35 (0.78, 2.32) | 0.14 | 38/57 | 1.66 (1.02, 2.71) | 0.021 | 2.26 (1.30, 3.91) | 0.002 |
| P² trend       |                   | 0.50            | 0.21        |       | 0.021          | 0.001       |       | 0.12            | 0.24        |
| Dominant       |                   | 1.15 (0.81, 1.64) | 0.22 | 1.25 (0.85, 1.84) | 0.13 |       | 1.49 (1.03, 2.16) | 0.018 | 1.79 (1.18, 2.71) | 0.003 |
| Recessive      |                   | 1.05 (0.67, 1.66) | 0.41 | 1.23 (0.74, 2.03) | 0.21 |       | 1.38 (0.90, 2.14) | 0.072 | 1.78 (1.09, 2.91) | 0.011 |
| **MTR**        |                   |                 |             |       |                 |             |       |                 |             |
| rs1805087      |                   |                 |             |       |                 |             |       |                 |             |
| A/A            | 161/310           | 1.00            | 1.00        | 142/310 | 1.00            | 1.00        | 149/310 | 1.00            | 1.00        |
| A/G            | 35/76             | 0.90 (0.58, 1.37) | 0.31 | 0.79 (0.50, 1.27) | 0.17 | 46/76 | 1.31 (0.87, 1.96) | 0.096 | 1.16 (0.74, 1.83) | 0.26 |
| G/G            | 4/0               | –               | 1/0         | –     | –               | 1/0         |       | –               | –           |
| P² trend       |                   | 0.50            | 0.21        |       | 0.021          | 0.001       |       | 0.12            | 0.24        |
| Dominant       |                   | 1.15 (0.81, 1.64) | 0.22 | 1.25 (0.85, 1.84) | 0.13 |       | 1.49 (1.03, 2.16) | 0.018 | 1.79 (1.18, 2.71) | 0.003 |
| Recessive      |                   | 1.05 (0.66, 1.67) | 0.41 | 1.23 (0.74, 2.03) | 0.21 |       | 1.38 (0.90, 2.14) | 0.072 | 1.78 (1.09, 2.91) | 0.011 |
| **MTRR**       |                   |                 |             |       |                 |             |       |                 |             |
| rs1532268      |                   |                 |             |       |                 |             |       |                 |             |
| C/C            | 145/283           | 1.00            | 1.00        | 144/283 | 1.00            | 1.00        | 149/283 | 1.00            | 1.00        |
| C/T            | 47/94             | 0.99 (0.67, 1.46) | 0.47 | 1.16 (0.76, 1.78) | 0.25 | 44/94 | 0.96 (0.65, 1.43) | 0.42 | 0.98 (0.63, 1.53) | 0.47 |
| T/T            | 11/11             | 1.80 (0.86, 3.80) | 0.060 | 1.96 (0.89, 4.31) | 0.047 | 5/11 | 1.08 (0.46, 2.56) | 0.43 | 1.05 (0.43, 2.57) | 0.46 |
| P² trend       |                   | 0.27            | 0.070       |       | 0.98            | 0.97        |       | 0.14            | 0.61        |
| Dominant       |                   | 1.11 (0.77, 1.61) | 0.29 | 1.32 (0.88, 1.98) | 0.089 |       | 0.98 (0.67, 1.64) | 0.45 | 1.00 (0.65, 1.52) | 0.49 |
| Recessive      |                   | 1.81 (0.86, 3.80) | 0.059 | 1.91 (0.87, 4.19) | 0.052 |       | 1.09 (0.46, 2.58) | 0.43 | 1.06 (0.43, 2.57) | 0.45 |
| rs1801394      |                   |                 |             |       |                 |             |       |                 |             |
| A/A            | 117/204           | 1.00            | 1.00        | 119/204 | 1.00            | 1.00        | 114/204 | 1.00            | 1.00        |
| A/G            | 74/149            | 0.88 (0.62, 1.25) | 0.24 | 0.92 (0.63, 1.35) | 0.34 | 63/149 | 0.75 (0.53, 1.08) | 0.063 | 0.74 (0.49, 1.11) | 0.070 |

Table 2. Associations between SNPs of MTHFR, MTR, MTRR, DNMT1, and ALDH2 genes and cancers of the esophagus, stomach, and liver.
### Table 2. Cont.

|                      | Esophageal Cancer | Stomach Cancer | Liver Cancer |
|----------------------|-------------------|----------------|-------------|
|                      | SBOR¹ (95% posterior limit) | One-sided P* | SBOR² (95% posterior limit) | One-sided P* | SBOR¹ (95% posterior limit) | One-sided P* |
| Ca/Co                |                      |               | Ca/Co       |                      | Ca/Co       |                      |
| G/G                  | 10/25              | 0.81 (0.42, 1.59) | 0.27        | 0.95 (0.47, 1.93) | 0.44        | 9/25                  | 0.76 (0.38, 1.50) | 0.21 | 0.77 (0.37, 1.60) | 0.24 | 13/25 | 0.93 (0.49, 1.79) | 0.41 | 1.39 (0.70, 2.78) | 0.17 |
| P_{trend}            | 0.36               | 0.70          | 0.10        | 0.13               | 0.30        | 0.07                  |
| Dominant             | 0.86 (0.61, 1.21)  | 0.20          | 0.92 (0.64, 1.34) | 0.33 | 0.74 (0.52, 1.05) | 0.046 | 0.73 (0.49, 1.07) | 0.055 | 0.80 (0.56, 1.13) | 0.10 | 0.83 (0.56, 1.19) | 0.19 |
| Recessive            | 0.85 (0.44, 1.64)  | 0.31          | 0.97 (0.48, 1.96) | 0.47 | 0.82 (0.41, 1.62) | 0.29 | 0.84 (0.41, 1.74) | 0.32 | 1.00 (0.54, 1.88) | 0.50 | 1.53 (0.78, 3.03) | 0.11 |
| DNMT1                |                    |               |             |                   |             |                       |
| rs2228612            |                    |               |             |                   |             |                       |
| A/A                  | 52/100             | 1.00          | 43/100      | 1.00               | 48/100      | 1.00                  | 1.00 |
| A/G                  | 56/200             | 0.58 (0.38, 0.88) | 0.006 | 0.52 (0.33, 0.82) | 0.003 | 72/200 | 0.84 (0.55, 1.28) | 0.21 | 0.82 (0.51, 1.31) | 0.20 | 74/200 | 0.79 (0.52, 1.19) | 0.13 | 0.79 (0.49, 1.28) | 0.17 |
| G/G                  | 29/57              | 1.03 (0.61, 1.73) | 0.46        | 0.96 (0.55, 1.68) | 0.44        | 28/57 | 1.11 (0.65, 1.89) | 0.36 | 1.12 (0.62, 2.04) | 0.35 | 36/57 | 1.33 (0.80, 2.22) | 0.13 | 1.35 (0.76, 2.42) | 0.15 |
| P_{trend}            | 0.57               | 0.37          | 0.88        | 0.88               | 0.42        | 0.42                  |
| Dominant             | 0.67 (0.45, 1.00)  | 0.026         | 0.60 (0.39, 0.94) | 0.012 | 0.90 (0.59, 1.36) | 0.31 | 0.89 (0.56, 1.31) | 0.31 | 0.92 (0.62, 1.37) | 0.34 | 0.93 (0.58, 1.50) | 0.39 |
| Recessive            | 1.38 (0.86, 2.21)  | 0.092         | 1.35 (0.81, 2.25) | 0.13 | 1.22 (0.76, 1.97) | 0.20 | 1.26 (0.74, 2.15) | 0.20 | 1.53 (0.97, 2.40) | 0.033 | 1.54 (0.92, 2.59) | 0.051 |
| ALDH2                |                    |               |             |                   |             |                       |
| rs671                |                    |               |             |                   |             |                       |
| G/G                  | 118/213            | 1.00          | 108/213     | 1.00               | 109/213     | 1.00                  | 1.00 |
| A/G                  | 60/160             | 0.70 (0.49, 1.01) | 0.029 | 0.76 (0.51, 1.13) | 0.088 | 76/160 | 0.97 (0.68, 1.38) | 0.44 | 1.06 (0.71, 1.57) | 0.39 | 74/160 | 0.91 (0.64, 1.30) | 0.31 | 1.05 (0.70, 1.58) | 0.41 |
| A/A                  | 25/20              | 1.97 (1.11, 3.49) | 0.010 | 1.61 (0.86, 3.00) | 0.066 | 12/20 | 1.13 (0.58, 2.21) | 0.36 | 0.99 (0.48, 2.03) | 0.49 | 10/20 | 1.03 (0.51, 2.06) | 0.47 | 0.96 (0.45, 2.06) | 0.46 |
| P_{trend}            | 0.49               | 0.67          | 0.88        | 0.88               | 0.77        | 0.94                  |
| Dominant             | 0.88 (0.63, 1.24)  | 0.24          | 0.90 (0.62, 1.31) | 0.29 | 1.00 (0.71, 1.40) | 0.49 | 1.05 (0.71, 1.54) | 0.41 | 0.93 (0.66, 1.31) | 0.33 | 1.04 (0.70, 1.54) | 0.43 |
| Recessive            | 2.20 (1.25, 3.86)  | 0.003         | 1.76 (0.96, 3.24) | 0.035 | 1.14 (0.59, 2.21) | 0.35 | 0.97 (0.48, 1.97) | 0.47 | 1.06 (0.53, 2.10) | 0.44 | 0.95 (0.45, 2.01) | 0.44 |
| rs2238151            |                    |               |             |                   |             |                       |
| C/C                  | 173/293            | 1.00          | 151/293     | 1.00               | 166/293     | 1.00                  | 1.00 |
| C/T                  | 11/34              | 0.66 (0.35, 1.22) | 0.093 | 0.66 (0.34, 1.29) | 0.11 | 18/34 | 1.08 (0.61, 1.89) | 0.40 | 1.21 (0.65, 2.23) | 0.28 | 7/34 | 0.44 (0.22, 0.87) | 0.009 | 0.51 (0.23, 0.945) | 1.11 |

SNPs in One-Carbon Metabolism and Upper GI Cancers

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|                       | Esophageal Cancer |                       | Stomach Cancer |                       | Liver Cancer |                       |                       |                       |
|-----------------------|-------------------|-----------------------|----------------|-----------------------|--------------|-----------------------|-----------------------|-----------------------|
|                       | Ca/Co             | One-sided \( P \)     | Ca/Co          | One-sided \( P \)   | Ca/Co        | One-sided \( P \)   | Ca/Co        | One-sided \( P \)   |
| T/T                   | 0/1               | –                     | 0/1            | –                     | 1/1          | 1.15 (0.33, 4.03)    | 0.41          | 1.43 (0.39, 5.33)    |
| \( P \)\(_{\text{trend}} \) | –                 | –                     | –              | –                     | 0.045        | 0.29                  |             |                       |
| Dominant              | 0.64 (0.35, 1.19) | 0.079                 | 0.65 (0.33, 1.28) | 0.11                  | 1.05 (0.60, 1.83) | 0.43                  | 1.20 (0.65, 2.21) | 0.29                  |
| Recessive             | –                 | –                     | –              | –                     | 1.16 (0.33, 4.07) | 0.41                  | 1.44 (0.39, 5.37) | 0.29                  |
| rs886205 G/G          | 173/315           | 1.00                  | 150/315        | 1.00                  | 149/315      | 1.00                  | 149/315      | 1.00                  |
| G/A                   | 29/69             | 0.81 (0.51, 1.26)     | 0.17           | 0.78 (0.48, 1.28)     | 0.17         | 1.05 (0.60, 1.83)    | 0.43         | 1.43 (0.39, 5.33)    |
| A/A                   | 2/4               | 1.07 (0.36, 3.18)     | 0.45           | 1.00 (0.33, 3.08)     | 0.50         | 1.10 (0.37, 3.30)    | 0.43         | 1.16 (0.33, 4.07)    |
| \( P \)\(_{\text{trend}} \) | 0.45              | 0.39                  | 0.32           | 0.098                 | 0.47         | 0.14                  |             |                       |
| Dominant              | 0.82 (0.53, 1.28) | 0.19                  | 0.79 (0.49, 1.29) | 0.17                  | 1.23 (0.82, 1.86) | 0.16                  | 1.48 (0.93, 2.34) | 0.047                 |
| Recessive             | 1.08 (0.36, 3.23) | 0.44                  | 1.02 (0.33, 3.13) | 0.49                  | 1.08 (0.36, 3.24) | 0.44                  | 1.16 (0.38, 3.55) | 0.40                  |

1: Semi-Bayes odds ratio (SBOR) adjusted for age (5-year categories and deviation from stratum mean) and sex.
2: SBOR further adjusted for residency (city, rural), alcohol drinking frequency, smoking pack-years, BMI, education, \( H. pylori \) infection (in stomach cancer analyses), HBsAg (in liver cancer analyses), and plasma AFB1 levels (in liver cancer analyses), based on the assumption that genotype did not affect any of these variables.
3: One-sided semi-Bayes \( P \)-values; the posterior probability that the point estimate is on the wrong side of the null [26,27].
*: \( P \)-value from Chi-Square test for trend.
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methylTHF and an accumulation of 5,10-methyleneTHF in red blood cells [30].

Low MTHFR activity is associated with increase cancer risk due to low blood 5-methylTHF and impaired DNA methylation. Conversely, it could reduce cancer risk by increasing the availability of 5,10-methyleneTHF for normal DNA synthesis and preventing uracil misincorporation and chromosomal breakage [5]. Although evidence in support of these hypotheses is weak and inconsistent [5], an in vitro study suggested that the effect of MTHFR rs1801133 on DNA stability and methylation is site-specific and may depend on folate availability [31]. When folate supply is adequate or high, the T allele of MTHFR is associated with increased genomic DNA methylation in colon cancer cells, but decreased DNA methylation in breast cancer cells. When folate supply is limited, this variant is associated with decreased and unchanged DNA methylation in colon and breast cancer cells, respectively [31]. Uracil misincorporation is decreased in colon cancer cells expressing the MTHFR T allele, and increased in breast cancer cells expressing the same variant [31]. This site-specific difference may partly explain the difference in cancer risk associated with the MTHFR rs1801133 polymorphism [4]. In epidemiologic studies, the T allele appears to decrease the risk of colorectal and breast cancers [32,33], but increase the risk of cancers of the esophagus, stomach, liver, bladder, cervix uteri, and lung [2–4,34–36].

In the present analysis using SB shrinkage, we confirmed our previous findings of positive associations between the T allele of MTHFR rs1801133 and cancers of the stomach and liver in this Taixing population [11,12], implying that the disturbance of DNA methylation resulting from this variant plays a major role in stomach and liver carcinogenesis. Recent meta-analyses reported similar associations (T/T versus C/C, OR: 1.40, 95% CI: 1.19–1.66 for stomach cancer; OR: 1.21, 95% CI: 0.95–1.56 for liver cancer) [3,4]. In addition, Zacho et al. [4] reported a larger

**Figure 1.** Selected semi-Bayes stratum-specific associations. Selected semi-Bayes stratum-specific associations between SNPs in MTHFR, MTR, MTRR, DNMT1, ALDH2, and upper GI cancer susceptibility, by plasma levels of micronutrients (folate, vitamin B12, and homocysteine) and environmental factors (smoking, alcohol drinking, H. pylori CagA status, HBsAg status, and plasma AFB1-albumin adduct levels). Semi-Bayes adjusted ORs (SBOR) and 95% posterior limits were under dominant genetic models, except for the SBOR relating ALDH2 rs671 to esophageal cancer, where recessive genetic model was used. P* denotes P-value for homogeneity test. doi:10.1371/journal.pone.0109235.g001
Table 3. Associations between SNPs of MTHFR, MTR, MTRR, DNMT1, and ALDH2 genes and cancers of the esophagus, stomach, and liver – the comparison between results from single SNP model\(^1\) and joint SNPs model\(^2\).

|                  | Ca/Co | SBOR\(^1\) (95% posterior limit) | One-sided \(P^*\) | Ca/Co | SBOR\(^2\) (95% posterior limit) | One-sided \(P^*\) | Ca/Co | SBOR\(^1\) (95% posterior limit) | One-sided \(P^*\) | Ca/Co | SBOR\(^2\) (95% posterior limit) | One-sided \(P^*\) |
|------------------|-------|---------------------------------|------------------|-------|---------------------------------|------------------|-------|---------------------------------|------------------|-------|---------------------------------|------------------|
| **Esophageal Cancer** |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| MTHFR rs1801133  |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| CC               | 65/135| 1.00                            | 1.00             | 50/135| 1.00                            | 1.00             | 50/135| 1.00                            | 1.00             |       |                                 |                  |
| CT               | 105/199| 1.19 (0.80, 1.77)               | 0.20             | 106/199| 1.53 (1.00, 2.33)               | 0.024            | 114/199| 1.57 (1.02, 2.42)               | 0.021            | 1.34 (0.80, 2.25)               | 0.14             |
| TT               | 32/57 | 1.35 (0.78, 2.32)               | 0.14             | 38/57 | 2.26 (1.30, 3.91)               | 0.002            | 30/57 | 1.17 (0.64, 2.12)               | 0.31             | 1.07 (0.52, 2.20)               | 0.42             |
| **Stomach Cancer** |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| MTHFR rs1805087  |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| A/A              | 161/310| 1.00                           | 1.00            | 142/310| 1.16 (0.74, 1.83)               | 0.26             | 40/76 | 0.92 (0.57, 1.47)               | 0.36             | 0.94 (0.54, 1.63)               | 0.41             |
| A/G              | 35/76 | 0.79 (0.50, 1.27)               | 0.17             | 46/76 | 1.16 (0.74, 1.83)               | 0.26             | 40/76 | 0.92 (0.57, 1.47)               | 0.36             | 0.94 (0.54, 1.63)               | 0.41             |
| G/G              | 4/0   | –                              | –               | 1/0   | –                              | –               | 1/0   | –                              | –               | –                              | –               |
| **Liver Cancer** |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| MTHFR rs1532268  |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| C/C              | 145/283| 1.00                           | 1.00            | 144/283| 1.16 (0.83, 1.53)               | 0.47             | 39/94 | 0.83 (0.52, 1.32)               | 0.21             | 1.03 (0.59, 1.81)               | 0.46             |
| C/T              | 47/94 | 1.16 (0.76, 1.78)               | 0.25             | 44/94 | 0.98 (0.63, 1.53)               | 0.47             | 39/94 | 0.83 (0.52, 1.32)               | 0.21             | 1.03 (0.59, 1.81)               | 0.46             |
| T/T              | 11/11 | 1.96 (0.89, 4.31)               | 0.047            | 5/11  | 1.05 (0.43, 2.57)               | 0.46             | 4/11  | 1.09 (0.42, 2.82)               | 0.43             | 1.58 (0.54, 4.63)               | 0.20             |
| **rs1801394**    |       |                                 |                  |       |                                 |                  |       |                                 |                  |       |                                 |                  |
| A/A              | 117/204| 1.00                           | 1.00            | 119/204| 1.00                           | 1.00            | 114/204| 1.00                           | 1.00            | 1.00                           | 1.00            |
Table 3. Cont.

| SNP | A/G | G/G | P<sub>bend</sub> | Dominant | Recessive |
|-----|-----|-----|-----------------|-----------|-----------|
| rs2228612 | 0.97 (0.48, 1.96) 0.47 | 1.35 (0.62, 2.92) 0.23 | 0.84 (0.41, 1.74) 0.32 | 1.09 (0.48, 2.47) 0.42 | 1.53 (0.78, 3.03) 0.11 |

| DNMT1 | A/A | A/G | G/G | P<sub>bend</sub> | Dominant | Recessive |
|-------|-----|-----|-----|-----------------|-----------|-----------|
| rs671 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 |
| rs2238151 | 1.76 (0.96, 3.24) 0.035 | 1.16 (0.56, 2.37) 0.35 | 0.97 (0.48, 1.97) 0.47 | 1.01 (0.48, 2.14) 0.49 | 0.95 (0.45, 2.01) 0.44 |

| ALDH2 | G/G | A/G | A/A | P<sub>bend</sub> | Dominant | Recessive |
|-------|-----|-----|-----|-----------------|-----------|-----------|
| rs671 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 |
| rs2238151 | 1.62 (0.97, 2.71) 0.033 | 1.00 | 1.00 | 1.00 | 1.00 | 1.00 |
### Table 3. Cont.

|                   | Esophageal Cancer | Stomach Cancer | Liver Cancer |
|-------------------|-------------------|----------------|-------------|
|                   | SBOR\(^1\) (95% posterior limit) | One-sided\(^p\) | SBOR\(^2\) (95% posterior limit) | One-sided\(^p\) | SBOR\(^1\) (95% posterior limit) | One-sided\(^p\) | SBOR\(^2\) (95% posterior limit) | One-sided\(^p\) |
| C/T               | 11/34             | 0.66 (0.34, 1.29) | 0.11         | 0.81 (0.33, 1.86) | 0.31 | 18/34 | 1.21 (0.65, 2.23) | 0.28 | 1.00 (0.45, 2.22) | 0.50 | 7/34 | 0.51 (0.23, 1.11) | 0.045 | 0.36 (0.14, 0.92) | 0.016 |
| T/T               | 0/1               | –              | –            | –              | –            | 0/1 | –              | –              | 1/1 | 1.43 (0.39, 5.33) | 0.30 | 1.00 (0.25, 4.00) | 0.50 |
| P\(_{\text{trend}}\)\(^d\) | –            | –              | –            | –              | –            | – | –              | –              | 0.29 | 0.006 |
| Dominant          | 0.65 (0.33, 1.28) | 0.11         | 0.80 (0.33, 1.84) | 0.30 | 1.20 (0.65, 2.21) | 0.29 | 1.08 (0.48, 2.35) | 0.43 | 0.58 (0.27, 1.25) | 0.082 | 0.31 (0.13, 0.75) | 0.005 |
| Recessive         | –            | –              | –            | –              | –            | – | 1.44 (0.39, 5.37) | 0.29 | 1.00 (0.25, 4.00) | 0.50 |
| rs886205          |                   |               |              |               |              |               |               |               |               |               |
| G/G               | 173/315           | 1.00         | 1.00         | 150/315         | 1.00 | 1.00         | 149/315         | 1.00 | 1.00 |
| A/G               | 20/69             | 0.78 (0.48, 1.28) | 0.17 | 1.05 (0.55, 2.00) | 0.44 | 41/69 | 1.47 (0.92, 2.34) | 0.054 | 1.76 (0.93, 3.33) | 0.041 | 40/69 | 1.38 (0.85, 2.24) | 0.096 | 1.41 (0.75, 2.66) | 0.14 |
| A/A               | 2/4               | 0.73 (0.33, 3.08) | 0.50 | 0.82 (0.25, 2.69) | 0.37 | 2/4 | 1.18 (0.39, 3.65) | 0.38 | 0.65 (0.19, 2.23) | 0.25 | 2/4 | 1.27 (0.40, 4.10) | 0.34 | 1.07 (0.31, 3.76) | 0.46 |
| P\(_{\text{trend}}\)\(^d\) | 0.37             | 0.909          | 0.37         | 0.909          | 0.37         | 0.098 | 0.31         | 0.098          | 0.31 | 0.14 | 0.17 |
| Dominant          | 0.79 (0.49, 1.29) | 0.17         | 0.99 (0.52, 1.86) | 0.48 | 1.48 (0.93, 2.34) | 0.047 | 1.38 (0.76, 2.51) | 0.15 | 1.41 (0.88, 2.28) | 0.076 | 1.34 (0.76, 2.37) | 0.16 |
| Recessive         | 1.02 (0.33, 3.13) | 0.49         | 0.85 (0.26, 2.75) | 0.39 | 1.16 (0.38, 3.55) | 0.40 | 0.70 (0.20, 2.40) | 0.28 | 1.25 (0.39, 4.03) | 0.35 | 0.93 (0.28, 3.04) | 0.45 |

1. Single SNP model: models included one SNP only; Semi-Bayes odds ratio (SBOR) adjusted for age (5-year categories and deviation from stratum mean), sex, residency (city, rural), alcohol drinking frequency, smoking pack-years, BMI, education, H. pylori infection (in stomach cancer analyses), HBsAg (in liver cancer analyses), and plasma AFB1 levels (in liver cancer analyses), based on the assumption that genotype did not affect any of these variables.
2. Joint SNPs model: models included all of the 8 SNPs; SBOR adjusted for the same set of covariates. All of the SNPs combined together were under the same genetic models (genotype-specific, log-additive, dominant, or recessive).
3. One-sided semi-Bayes \(P\)-values; the posterior probability that the point estimate is on the wrong side of the null [26,27].

**Note:**

- \(P\)-value from Chi-Square test for trend.
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Table 4. Association between polygenetic risk score and cancers of the esophagus, stomach, and liver

| Polygenetic risk score (8 SNPs) | Esophageal cancer | Stomach cancer | Liver cancer |
|--------------------------------|-------------------|---------------|--------------|
| One-sided                        |                    |               |              |
| SBOR                            |                    |               |              |
| **P** Ca/Co SBOR                |                    |               |              |
| One-sided                        |                    | Middle        |              |
| SBOR                            |                    |               |              |
| **P** Ca/Co SBOR                |                    |               |              |
| One-sided                        |                    | Highest       |              |
| SBOR                            |                    |               |              |
| **P** Ca/Co SBOR                |                    |               |              |
| Continuous (1 unit increase in PRS) |                   |               |              |
| SBOR                            |                    |               |              |
| **P** Ca/Co SBOR                |                    |               |              |

*S*: One-sided semi-Bayes P-values; the posterior probability that the point estimate is on the wrong side of the null [26,27].

Polygenetic risk score (PRS) was calculated only among subjects with complete data on all 8 SNPs.

Association between MTHFR rs1801133 and stomach cancer among study populations without folic acid fortification (OR: 1.60, 95% CI: 1.36–1.88), as compared to those with fortification (OR: 1.15, 95% CI: 0.81–1.63), which is similar to our finding of a stronger association among individuals with lower plasma folate levels. For esophageal cancer, our data suggested an increased risk among MTHFR rs1801133 T allele carriers (any T vs. C/C, SBOR: 1.25, 95% posterior limits: 0.85, 1.84), which is consistent with findings from a meta-analysis of 19 studies (C/T versus C/C, OR: 1.47, 95% CI: 1.32–1.63; T/T versus C/C, OR: 1.69, 95% CI: 1.49–1.91) [2].

MTR and MTRR are two other important enzymes involved in one-carbon metabolism. MTR catalyzes the methylation of homocysteine to methionine. MTR A2756G (rs1805087), a common SNP leading to the substitution of aspartic acid with glycine, has been largely studied. However, no apparent associations have been observed with cancer at the following sites: lung, prostate, head and neck, bladder, esophagus, stomach, breast, or colon and rectal [37–46]. MTRR regenerates a functional MTR via reductive methylation. Two common polymorphisms, MTRR A66G (rs1801394, converts isoleucine to methionine) and C524T (rs1532268, changes serine to leucine), have been indicated to regenerate MTR less efficiently [47]. G allele carriers of MTRR rs1801394 have been associated with increased risk for hepatocellular carcinoma (HCC) [48]. Conversely, associations are inconsistent with other malignancies, including esophageal squamous cell carcinoma (ESCC), stomach cancer, and colorectal cancer [37,44,49–53]. Most studies that have investigated MTRR rs1532268 reported no associations with colorectal, gastric, breast, and lung cancer [44,51,53–56]. One should bear in mind however that apparent inconsistencies and reports of no association may only reflect expected variation in P-values (“statistical significance”) rather than any real conflicts.

Our study observed odds-ratio variation of the associations between these MTR/MTRR polymorphisms and upper GI cancers across alcohol consumption, even after conservative SB shrinkage. Alcohol consumption appeared to have modified odds-ratios relating MTR rs1805087 to liver cancer, and MTRR rs1801394 to esophageal and stomach cancer. G allele carriers of these two SNPs were positively associated with cancer among drinkers, and inversely associated with cancer among non-drinkers. Matsuo et al., observed a similar OR variation [57]: G/G genotype carriers of MTR rs1805087 showed higher colorectal cancer risk among alcohol drinkers and lower risk among non-drinkers. Although the functional effect of MTR/ MTRR polymorphisms has not been established, our results are biologically plausible as alcohol can disrupt one-carbon metabolism by inhibiting folate absorption, suppressing SAM synthesis, and impairing DNA methylation [6]. Alcohol can also cause inhibition of methionine synthase activity [6]. Therefore, it is possible that the variant allele of these two MTR/MTRR polymorphisms is protective for the upper GI cancers under the environment without alcohol exposures. However, it becomes deleterious when one-carbon metabolism is disrupted by alcohol and its metabolites.

ALDH2 is involved in alcohol metabolism by oxidizing acetaldehyde, a group 2B human carcinogen, to acetic acid. The ALDH2 rs671 polymorphism—a well-known variant that occurs exclusively in Asian populations-causes a lower catalytic efficiency of ALDH2, and hence renders lower ability to eliminate acetaldehyde [58]. The ALDH2 rs671 A allele (slow type) has been associated with increased risk of head and neck cancer, as well as esophageal cancer [59,60]. Consistent with previous findings, we observed a positive association between the A/A genotype and...
esophageal cancer in this study, and further reported a stronger association among individuals with lower plasma folate levels. Acetaldehyde also interferes with folate metabolism [6]. It is possible that the deleterious effect associated with rs671 polymorphism is more prominent under the condition of lower folate supply. Another common variation in the ALDH2 gene—rs809205 with a G to A substitution in the promoter region—has been suggested to be functional. Chou et al., reported that the promoter constructs encoded by the G allele were more active than the A allele in hepatoma cells [61]. The G allele of rs806205 was reported to be associated with increased risk of ESCC [62,63] but not with stomach [64,65] and colorectal cancer [66]. It also showed inconsistent results with head and neck cancer [67–69]. We observed a positive association between the A allele of rs806205 and stomach cancer among those with higher plasma vitamin B12 levels. Although ALDH2 rs806205 is suggested to be a functional polymorphism in hepatoma cells [61], further functionality studies are warranted. There are several limitations due to the case-control design and the multiple comparisons in this study. Because we were not able to recruit all of the identified cases and controls, selection bias may occur if participation is affected by an un-identified factor which is associated with both the SNPs and cancer. On the other hand, cancers of the esophagus, stomach, and liver are fatal, and some patients with late clinical stages at diagnosis were either too ill to participate or passed away. This selection of patients may have resulted in biased estimates if the SNPs under study are associated with disease progression. Also, we collected plasma samples after cancer diagnoses. By stratification on plasma micronutrients, we may have introduced a “collider-stratification bias” if disease states, as well as treatments and/or diet and behavior changes among cancer patients would affect the levels of these biomarkers [70]. However, given that there may be only a weak association between SNPs and plasma micronutrients in one-carbon metabolic pathway [71], we believe that the size of the bias would be small. In addition, we conducted many comparisons and subgroup analyses, which led us to employ semi-Bayes shrinkage estimation to reduce the risk of misleading results. Using these methods, in this Chinese population, several polymorphisms in the one-carbon metabolic pathway appear to be associated with esophageal, stomach, and liver cancer, with heterogeneity across strata of alcohol consumption for the odds ratios relating MTHFR/MTR/MTRR polymorphisms to these cancers, suggesting potential interactions between alcohol drinking and genes of the one-carbon metabolic pathway. Confirmation of these results and research on the underlying mechanisms are needed.

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