Impact of SNP-SNP interactions of DNA repair gene ERCC5 and metabolic gene GSTP1 on gastric cancer/atrophic gastritis risk in a Chinese population

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Abstract

AIM
To investigate the interactions of the DNA repair gene excision repair cross complementing group 5 (ERCC5) and the metabolic gene glutathione S-transferase pi 1 (GSTP1) and their effects on atrophic gastritis (AG) and gastric cancer (GC) risk.
METHODS

Seven ERCC5 single nucleotide polymorphisms (SNPs) (rs1047768, rs2094258, rs2228959, rs4150291, rs4150383, rs751402, and rs873601) and GSTP1 SNP rs1695 were detected using the Sequenom MassARRAY platform in 450 GC patients, 634 AG cases, and 621 healthy control subjects in a Chinese population.

RESULTS

Two pairwise combinations (ERCC5 rs2094258 and rs873601 with GSTP1 rs1695) influenced AG risk ($P_{interaction} = 0.008$ and $0.043$, respectively), and the ERCC5 rs2094258-GSTP1 rs1695 SNP pair demonstrated an antagonistic effect, while ERCC5 rs873601-GSTP1 rs1695 showed a synergistic effect on AG risk (OR = 0.51 and 1.79, respectively). No pairwise combinations were observed in relation to GC risk. There were no cumulative effects among the pairwise interactions (ERCC5 rs2094258 and rs873601 with GSTP1 rs1695) on AG susceptibility ($P_{trend} > 0.05$). When the modification effect of Helicobacter pylori (H. pylori) infection was evaluated, the cumulative effect of one of the aforementioned pairwise interactions (ERCC5 rs873601-GSTP1 rs1695) was associated with an increased AG risk in the case of negative H. pylori status ($P_{trend} = 0.043$).

CONCLUSION

There is a multifarious interaction between the DNA repair gene ERCC5 SNPs (rs2094258 and rs873601) and the metabolic gene GSTP1 rs1695, which may form the basis for various inter-individual susceptibilities to AG.

Key words: Excision repair cross complementing group 5; Glutathione S-transferase pi1; Atrophic gastritis; Gastric cancer; Single nucleotide polymorphisms

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Core tip: We detected seven excision repair cross complementing group 5 (ERCC5) single nucleotide polymorphisms (SNPs) and a glutathione S-transferase pi1 (GSTP1) SNP using the Sequenom MassARRAY platform in a Chinese population and used them to investigate their interactions and their effects on atrophic gastritis and gastric cancer risk. The results showed a multifarious interaction between the DNA repair gene ERCC5 SNPs (rs2094258 and rs873601) and the metabolic gene GSTP1 rs1695. In addition, the cumulative effect of one pairwise interaction (ERCC5 rs873601-GSTP1 rs1695) was associated with an increased atrophic gastritis risk in the case of negative H. pylori status when the modification effect of H. pylori infection was evaluated.

INTRODUCTION

In light of the study of gastric cancer (GC) pathogenesis, there is increasing evidence to suggest that the interactions between various inherited susceptibility genes may affect the risk of GC development in individuals[1]. Single nucleotide polymorphisms (SNPs), as one of the most general forms of genetic variation, play a key role in predicting cancer risk in individuals and are widely applied to study tumor incidence and prognostic evaluation. However, they are inadequately utilized for studies of various genes in intricate diseases such as cancer[2], and the presently investigated polymorphisms for each single gene may not entirely reveal a definite phenotype[3]. Some studies have shown that interactions among genes are more significant than solitary genes in determining cancer susceptibility[4]. Numerous epidemiological studies have shown that inherited polymorphisms involved in xenobiotic metabolism and DNA repair are related to GC[5,6]. These genes are acknowledged as risk-modifier indicators, especially those whose allelic polymorphisms are accountable for the repair of oxidative stress induced DNA damage and/or the impaired metabolism of exogenous carcinogens. Excision repair cross complementing group 5 (ERCC5) is a critical element of the nucleotide excision repair (NER) pathway, and the ERCC5 gene is mapped to a region on chromosome 13q33 and comprises 15 exons[7]. It encodes a structure-specific endonuclease that has multiple functions during NER[7]. Its main role is to identify and shear damage to the DNA chain 3’ terminus[8]. Its gene mutation may lead to abnormal cell proliferation and differentiation and increased cancer susceptibility. SNPs of ERCC5 linked with GC susceptibility have been reported, including rs2094258, rs751402, rs2296147, rs1047768, rs873601, rs2227869, and rs17655[6,9-15]. We previously analyzed six SNPs of the ERCC5 gene in 2686 subjects from northern China and found that the selected polymorphisms of the ERCC5 gene were not significantly associated with atrophic gastritis (AG)/GC risk[16]. Glutathione S-transferase (GST) is an important member of the phase II metabolic enzymes, including GSTM1, GSTT1, and glutathione S-transferase pi1 (GSTP1)[17], which can affect detoxification processes and increase individual susceptibility to cancers[18]. The GSTP1 Ile105Val polymorphism produces the amino acid replacement of Ile (105) with Val via the change of A (Ile) to G (Val) in exon 5, which diminishes enzyme catalytic activity[19] and indirectly stimulates DNA repair and protection of the cell genome[17,20]. Our previous study also identified SNP rs1695 in GSTP1, which appears to drastically change the susceptibility of individuals to
GC\[^9\]. This finding is consistent with previous studies\[^{10,11}\].

Although some studies have found that ERCC5 SNPs and GSTP1 polymorphisms were related to GC risk, there are limited data on the effects of gene-gene interactions, and some results are equivocal\[^{12,13}\]. Additionally, given the vital impact of environmental factors on the susceptibility to GC and our previous findings regarding gene interaction and environmental factors\[^{4,14,16}\], we explored possible two-dimensional gene interactions among inherited polymorphisms in the DNA repair gene ERCC5 (rs1047768, rs2094258, rs2228959, rs4150291, rs4150383, rs751402, and rs873601) and the metabolic gene GSTP1 (rs1695), as well as the three-dimensional interactions between SNP-SNP and environmental factors in diverse stages of gastric carcinogenesis to assess the possibility of predicting GC risk and the identification of a combination of biomarkers for precancerosis and GC.

**MATERIALS AND METHODS**

**Study population**

In all, 1705 subjects were included in the present study, comprising 621 healthy controls, 634 cases of AG, and 450 cases of GC. All registered individuals originated from a Screening Program for Gastric Diseases or hospitals in Zhanhu and Shenyang of Liaoning Province, China between 2002 and 2013, as previously described\[^{16}\]. Metadata for every participant was collected using a standardized questionnaire survey and stored in a spreadsheet, including gender, age, history of illness, status of smoking, and alcohol consumption. Every participant signed a written informed consent form, according to the Declaration of Helsinki and its later revision. We collected peripheral venous blood from all participants, and experienced endoscopists simultaneously performed gastroscopic examination. All subjects received histopathological diagnosis according to the updated Sydney System\[^{15}\] and the World Health Organization criteria, independently, by two gastrointestinal pathologists. This project was approved by the Human Ethics Review Committee of China Medical University (Shenyang, China).

**SNP selection and genotyping assay**

Briefly, as described in our previous study\[^{16}\], we extracted ERCC5 genotype data from the HapMap Chinese Han Beijing population (http://www.HapMap.org). Tag SNPs were derived from pairwise linkage disequilibrium information to maximally capture ($r^2 > 0.8$) common or rare variants [minor allele frequency (MAF) > 0.05] using Haploview 4.2 (http://www.broadinstitute.org/mpg/haploview). FastSNP Search was used to predict potential SNP function. Finally, a total of seven ERCC5 SNPs (rs1047768, rs2094258, rs2228959, rs4150291, rs4150383, rs751402, and rs873601) were chosen in this study. In addition, GSTP1 rs1695 was selected according to our previous study and literature references\[^{9-11}\]. Genomic DNA was isolated from blood samples using a routine phenol-chloroform method and then diluted to working concentrations (50 ng/$\mu$L) for genotyping. Samples were placed randomly in 384-well plates and blinded for disease status. Selected SNP genotyping was performed using the Sequenom MassARRAY platform (Sequenom, San Diego, CA, United States) according to the manufacturer’s instructions\[^{16}\]. The average genotyping rate was 99.3% and the results of all duplicated samples were 100% consistent.

**Assessment of Helicobacter pylori serology**

*Helicobacter pylori* (H. pylori) immunoglobulin G levels was tested using an enzyme-linked immunosorbent assay (ELISA kit, Biohit, Helsinki, Finland) according to the manufacturer’s instructions, as previously described\[^{26}\]. *H. pylori* positivity was defined as a numerical reading exceeding 34 enzyme immune units.

**Statistical analysis**

Statistical analyses in the study were completed by applying SPSS 17.0 software (SPSS Inc., Chicago, IL, United States). We used the $\chi^2$ test to calculate the differences in demographic characteristics and genotypes between cases and controls. The two- or three-dimensional interaction effects among SNP-SNP with or without environmental factors were estimated using multivariate logistic regression models. General linear regression modeling was used to assess the trends with an increasing number of mutation genotypes in the cumulative effect. Associations were evaluated by odds ratios (ORs) and 95% confidence intervals (CIs) adjusted by sex, age, and *H. pylori* infection status except for being stratified by *H. pylori* infection status. Two-sided *P*-values < 0.05 were considered statistically significant.

**RESULTS**

**Demographic and geographic characteristics**

The distribution characteristics of gender, age, and *H. pylori* infection status of all participants are shown in Table 1. No significant differences were found in the gender or age distribution among the case and control groups. The study subjects consisted of 634 AG patients, 450 GC patients, and two control groups, including 620 and 535 for AG and GC cases, matched by gender and age, respectively. Additionally, there were significantly higher *H. pylori* infection rates (59.5% and 49.6%, respectively) in the AG and GC groups compared to the two matched control groups (27.1% and 26.7%, respectively, $P < 0.001$).

**Pairwise interactions between the ERCC5 SNPs and GSTP1 rs1695 polymorphism**

We primarily examined SNP-SNP two-dimensional interaction effects in the main effect analysis using a full-factor model. Two pairwise SNP combinations...
were found that could affect AG risk, but no pairwise combination was found in relation to GC risk. The results indicated that the ERCC5 rs2094258 and rs873601 polymorphisms with GSTP1 rs1695 polymorphism could engender interaction effects for AG risk ($P_{interaction} = 0.008$ and 0.043 respectively, Table 2). The ERCC5 rs2094258-GSTP1 rs1695 SNP pair demonstrated an antagonistic effect, while ERCC5 rs873601-GSTP1 rs1695 showed a synergistic effect on AG risk ($OR = 0.51$ and $1.79$, respectively, Table 2). No significant differences were observed among other SNP-SNP interactions ($P > 0.05$).

Epistatic effect of two-way interactions
We further investigated epistatic effects between pairs of ERCC5 rs2094258 and rs873601 polymorphisms with GSTP1 rs1695. For ERCC5 rs2094258 and GSTP1 rs1695, the AG/AA genotypes of rs2094258 and AA genotype of rs1695 were related to an increased risk of AG, but GA/GG genotypes of rs1695 were associated with a reduced risk of AG ($OR = 1.523$ and $0.678$, respectively). For ERCC5 rs873601 and GSTP1 rs1695, AA genotype of rs873601 resulted in a reduced risk of AG, only in the presence of AA genotype of rs1695 ($OR = 0.678$) (Table 3). These findings illustrated that ERCC5 rs2094258 and rs873601, individually, had no main effect but did display epistatic interactions with GSTP1 rs1695.

Cumulative effect of the interacting factors of ERCC5 SNPs-GSTP1 rs1695
We also investigated the cumulative effect among the interacting SNPs of ERCC5 rs2094258 and rs873601 with GSTP1 rs1695, but neither had a statistically significant relationship to AG risk ($P > 0.05$, Table 4). We further analyzed the cumulative effect of interacting SNPs modified by H. pylori. The ERCC5 rs873601-GSTP1 rs1695 SNP pair had significant differences in AG risk among the subgroups with negative H. pylori infection status ($P_{trend} = 0.043$). Moreover, AG risk was significantly reduced while one or two mutation genotypes were present ($OR = 0.66$, 95%CI: 0.37-1.16, and $OR = 0.73$, 95%CI: 0.53-1.02, respectively).

Three-dimensional analysis of the effect of interactions of ERCC5 SNPs-GSTP1 rs1695-environmental factors on AG risk
To explore the influence of environmental factors on the interaction, we further explored probable three-dimensional interactions among ERCC5 SNPs (rs2094258 and rs873601), GSTP1 rs1695, and environmental factors (smoking, alcohol consumption, and H. pylori infection status). We found no significant three-dimensional interactions with regard to AG risk ($P > 0.05$, Supplementary Table 1).

DISCUSSION
GC is an outcome of the interaction between multiple genes and environmental factors and is considered a multistep and multifactor process involving different carcinogen metabolic and DNA repair pathways[19,20]. Currently, researchers are concentrating more on the gene-gene interaction effect rather than a single-gene effect. In this study, we examined the possible interaction effect of DNA repair gene ERCC5 SNPs and the metabolic detoxification gene GSTP1 polymorphism. We first found new two-pair SNP interactions among ERCC5 SNPs and the GSTP1 polymorphism (ERCC5 rs2094258-GSTP1 rs1695 and ERCC5 rs873601-GSTP1 rs1695), which could alter the susceptibility to AG compared to host genetic effects alone. Moreover, the cumulative effect resulting from two-way interaction of ERCC5 rs873601-GSTP1 rs1695 was shown to differ in a stratified analysis of H. pylori infection status. The change from no cumulative effect to significant difference in AG risk in the case of negative H. pylori status indicated that H. pylori infection status could modify the cumulative effect mentioned above for the interacting SNPs. Genetic polymorphisms may explain partial individual deviations in disease risk, but a more multifarious condition involving numerous gene-gene interactions and gene-environment characteristics must

### Table 1 Baseline characteristics of the subjects n (%)

| Variable                      | AG vs CON | AG vs CON | GC vs CON | GC vs CON |
|-------------------------------|-----------|-----------|-----------|-----------|
|                               | CON       | AG        | CON       | GC        |
|                               | n = 620   | n = 634   | n = 535   | n = 450   |
| Gender                        |           |           |           |           |
| Male                          | 362 (58.4)| 358 (56.5)| 363 (67.9)| 298 (66.2)|
| Female                        | 258 (41.6)| 276 (43.5)| 172 (32.1)| 152 (33.8)|
| Age                           |           |           |           |           |
| Mean ± SD                     | 54.7 ± 9.1| 54.8 ± 9.0| 55.6 ± 9.2| 56.3 ± 10.1|
| Median                        | 54        | 55        | 56        | 57        |
| Range                         | 17-85     | 16-82     | 17-85     | 26-84     |
| H. pylori infection status    |           |           |           |           |
| Positive                      | 168 (27.1)| 137 (59.5)| 143 (26.7)| 223 (49.6)|
| Negative                      | 452 (72.9)| 257 (40.5)| 392 (73.5)| 227 (50.4)|

AG: Atrophic gastritis; GC: Gastric cancer; CON: Controls.
| Gene | Genotype | Number of participants | GSTP1 rs1695 |
|------|----------|------------------------|-------------|
|      |          |                        | AA + GG     | AA + GA | GG |
| AG vs CON (rs = 634 vs 620) | TT | No. of cases/controls | 231/200 | 124/116 | 338/307 | 17/9 |
|      | OR (95%CI) | 1 (Ref.) | 0.93 (0.68-1.27) | 1 (Ref.) | 1.72 (0.75-3.91) |
|      | TC + CC | No. of cases/controls | 177/188 | 102/116 | 270/289 | 9/15 |
|      | OR (95%CI) | 0.82 (0.62-1.08) | 0.76 (0.55-1.06) | 0.85 (0.68-1.07) | 0.55 (0.24-1.26) |
|      | P = 0.317 | Interaction index = 0.88 |
|      | TT + TC | No. of cases/controls | 376/345 | 207/214 | 559/538 | 24/21 |
|      | OR (95%CI) | 1 (Ref.) | 0.89 (0.70-1.13) | 1 (Ref.) | 1.10 (0.61-2.00) |
|      | CC | No. of cases/controls | 32/43 | 19/18 | 49/38 | 2/3 |
|      | OR (95%CI) | 0.68 (0.42-1.10) | 0.97 (0.50-1.87) | 0.81 (0.55-1.23) | 0.64 (0.31-1.36) |
|      | P = 0.296 |
|      | ERCC5 rs1047768 | GG | No. of cases/controls | 132/162 | 93/84 | 214/234 | 11/12 |
|      | OR (95%CI) | 1 (Ref.) | 1.36 (0.94-1.98) | 1 (Ref.) | 1.00 (0.43-2.32) |
|      | GA + AA | No. of cases/controls | 276/226 | 133/148 | 394/362 | 15/12 |
|      | OR (95%CI) | 1.50 (1.12-2.00) | 1.10 (0.79-1.53) | 1.19 (0.94-1.50) | 1.37 (0.63-2.99) |
|      | P = 0.008 |
|      | GG + GA | No. of cases/controls | 337/328 | 195/204 | 508/510 | 24/22 |
|      | OR (95%CI) | 1 (Ref.) | 0.93 (0.73-1.19) | 1 (Ref.) | 1.10 (0.61-1.98) |
|      | AA | No. of cases/controls | 71/60 | 31/28 | 100/86 | 2/2 |
|      | OR (95%CI) | 1.15 (0.79-1.68) | 1.08 (0.63-1.84) | 1.17 (0.85-1.60) | 1.00 (0.14-7.15) |
|      | P = 0.594 |
|      | CC + CA | No. of cases/controls | 408/383 | 224/231 | 606/590 | 26/24 |
|      | OR (95%CI) | 1 (Ref.) | 0.91 (0.72-1.15) | 1 (Ref.) | 1.06 (0.60-1.86) |
|      | AA | No. of cases/controls | 0/5 | 2/1 | 2/6 | 0/0 |
|      | OR (95%CI) | 1.88 (0.17-20.79) | 1.88 (0.17-20.79) | 3.22 (0.07-120.14) | 1.00 (0.72-1.31) |
|      | P = NA |
|      | ERCC5 rs2228959 | CC | No. of cases/controls | 371/346 | 198/215 | 548/539 | 21/22 |
|      | OR (95%CI) | 1 (Ref.) | 0.86 (0.67-1.09) | 1 (Ref.) | 0.94 (0.51-1.73) |
|      | CA + AA | No. of cases/controls | 37/42 | 28/17 | 60/57 | 5/2 |
|      | OR (95%CI) | 0.82 (0.52-1.31) | 1.54 (0.83-2.86) | 1.04 (0.71-1.52) | 2.46 (0.48-12.73) |
|      | P = 0.103 |
|      | CC + CA | No. of cases/controls | 408/383 | 224/231 | 606/590 | 26/24 |
|      | OR (95%CI) | 1 (Ref.) | 0.91 (0.72-1.15) | 1 (Ref.) | 1.06 (0.60-1.86) |
|      | AA | No. of cases/controls | 0/5 | 2/1 | 2/6 | 0/0 |
|      | OR (95%CI) | 1.88 (0.17-20.79) | 1.88 (0.17-20.79) | 3.22 (0.07-120.14) | 1.00 (0.72-1.31) |
|      | P = NA |
|      | ERCC5 rs4150291 | AA | No. of cases/controls | 347/332 | 193/205 | 517/516 | 23/21 |
|      | OR (95%CI) | 1 (Ref.) | 0.90 (0.70-1.15) | 1 (Ref.) | 1.09 (0.60-2.00) |
|      | AT + TT | No. of cases/controls | 61/56 | 33/27 | 91/80 | 3/3 |
|      | OR (95%CI) | 1.04 (0.70-1.54) | 1.17 (0.69-1.99) | 1.14 (0.62-1.97) | 1.00 (0.20-4.97) |
|      | P = 0.067 |
|      | AA + AT | No. of cases/controls | 406/382 | 225/232 | 605/590 | 26/24 |
|      | OR (95%CI) | 1 (Ref.) | 0.91 (0.73-1.15) | 1 (Ref.) | 1.06 (0.60-1.86) |
|      | TT | No. of cases/controls | 2/6 | 1/0 | 3/6 | 0/0 |
|      | OR (95%CI) | 0.31 (0.06-1.56) | 0.49 (0.12-1.96) | 0.49 (0.12-1.96) | 0.49 (0.12-1.96) |
|      | P = 0.703 |
|      | ERCC5 rs4150383 | GG | No. of cases/controls | 365/344 | 197/202 | 539/526 | 23/20 |
|      | OR (95%CI) | 1 (Ref.) | 0.92 (0.72-1.18) | 1 (Ref.) | 1.12 (0.61-2.07) |
|      | GA + AA | No. of cases/controls | 43/44 | 29/30 | 69/70 | 3/4 |
|      | OR (95%CI) | 0.92 (0.59-1.44) | 0.91 (0.54-1.55) | 0.96 (0.68-1.37) | 0.73 (0.16-3.29) |
|      | P = 0.720 |
|      | GG + GA | No. of cases/controls | 406/387 | 226/231 | 606/594 | 26/24 |
|      | OR (95%CI) | 1 (Ref.) | 0.93 (0.74-1.17) | 1 (Ref.) | 1.06 (0.60-1.87) |
|      | AA | No. of cases/controls | 2/1 | 0/1 | 2/2 | 0/0 |
|      | OR (95%CI) | 1.91 (0.17-21.11) | NA | 0.98 (0.14-6.98) | NA |
|      | P = NA |
|      | ERCC5 rs751402 | CC | No. of cases/controls | 191/173 | 97/104 | 281/266 | 7/11 |
|      | OR (95%CI) | 1 (Ref.) | 0.85 (0.60-1.19) | 1 (Ref.) | 0.60 (0.25-1.58) |
| CT + TT | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
|--------|----------------------|------------|---------|------------------|
| OR (95%CI) | 203/198              | 0.93 (0.70-1.23) | **0.99 (0.69-1.32)** | 1.38 (0.67-2.86) |
| **P** | | | | |
| CC + CT | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 355/324              | 1.02 (0.79-1.32) | **0.96 (0.76-1.21)** | 2.84 (1.18-5.05) |
| **P** | | | | |
| TT | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 39/47               | 0.76 (0.48-1.19) | **0.87 (0.60-1.25)** | 0.95 (0.24-3.81) |
| **P** | | | | |
| ERCC5 rs873601 | GG | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 126/109              | 0.83 (0.54-1.30) | **0.89 (0.61-1.18)** | 1.14 (0.47-2.78) |
| **P** | | | | |
| GA + AA | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 282/279              | 0.87 (0.64-1.19) | **0.93 (0.72-1.19)** | 0.95 (0.49-1.76) |
| **P** | | | | |
| AA | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 321/279              | 0.69 (0.50-0.96) | **0.81 (0.62-1.06)** | 1.14 (0.47-2.78) |
| **P** | | | | |

**GC vs CON**

(n = 450 vs 535 )

| TT | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 142/162              | 1.02 (0.79-1.32) | **1.05 (0.81-1.36)** | 1.60 (0.89-3.36) |
| **P** | | | | |
| TC + CC | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 128/161              | 0.89 (0.65-1.23) | **1.05 (0.81-1.36)** | 1.86 (0.79-4.36) |
| **P** | | | | |
| TT + TC | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 247/192              | 1.02 (0.79-1.32) | **1.08 (0.81-1.36)** | 1.86 (0.69-4.86) |
| **P** | | | | |
| ERCC5 rs2094258 | GG | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 110/131              | 1.02 (0.79-1.32) | **1.03 (0.72-1.19)** | 2.87 (1.33-6.17) |
| **P** | | | | |
| GA + AA | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 160/195              | 0.89 (0.50-0.96) | **0.93 (0.62-1.39)** | 1.14 (0.47-2.78) |
| **P** | | | | |

**ERCC5 rs2228959 | CC | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 247/192              | 1.02 (0.79-1.32) | **1.03 (0.72-1.19)** | 2.87 (1.33-6.17) |
| **P** | | | | |

** ERCC5 rs4150291 | AA | No. of cases/controls | OR (95%CI) | P-value | Interaction index |
| OR (95%CI) | 222/276              | 1.02 (0.79-1.32) | **1.03 (0.72-1.19)** | 2.87 (1.33-6.17) |
| **P** | | | | |

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be mentioned. A combination of SNPs would produce synergistic or antagonistic effects compared to an SNP, which could change the susceptibility to disease. Individually, two ERCC5 SNPs (rs2094258 and rs873601; unpublished data) showed no effect on either AG or GC risk ($P > 0.05$). However, our findings revealed a main effect on AG risk while these polymorphisms interacted with GSTP1 rs1695 ($P_{interaction} = 0.008$ and 0.043, respectively). The pairwise ERCC5 rs2094258-GSTP1 rs1695 and ERCC5 rs873601-GSTP1 rs1695 combinations had an OR of 0.51 and 1.79, respectively, for AG risk in the above two-way interaction analysis. In all, this evidence suggests that polymorphisms harbored in ERCC5 (rs2094258 and rs873601) had a synergistic or antagonistic effect with GSTP1 rs1695, which could alter the risk of an individual towards AG. According to the potential mechanism of SNP-SNP interactions, the ERCC5 gene, as an NER pathway gene, may be responsible for repairing DNA damage from biological and environmental mutagens or regular cellular metabolism. In addition, GSTP1 as an important phase II metabolizing xenobiotic enzyme might promote DNA damage repair through an exogenous metabolic detoxification pathway. When exogenous or endogenous carcinogens cause damage to DNA, the metabolic gene GSTP1 removes some harmful substances through the detoxification effect and then promotes DNA damage repair to protect against carcinogenic progression. The DNA repair gene ERCC5 can identify and incise a DNA modification pathway. When exogenous or endogenous carcinogens cause damage to DNA, the metabolic gene GSTP1 removes some harmful substances through the detoxification effect and then promotes DNA damage repair to protect against carcinogenic progression. The DNA repair gene ERCC5 can identify and incise a DNA

| ERCC5 rs4150383 | GSTP1 rs1695 |
|-----------------|--------------|
| **OR (95%CI)**  | **OR (95%CI)** |
| 1 (Ref.)        | 1 (Ref.)     |
| 1.06 (0.64-1.74)| 0.50 (0.25-1.01)|
| $P_{interaction} = 0.060$ | $P_{interaction} = NA$ |
| **Interaction index = 0.43** | **Interaction index = NA** |

| ERCC5 rs751402 | GSTP1 rs1695 |
|-----------------|--------------|
| **OR (95%CI)**  | **OR (95%CI)** |
| 1 (Ref.)        | 1 (Ref.)     |
| 1.08 (0.65-1.81)| 0.73 (0.38-1.42)|
| $P_{interaction} = 0.409$ | $P_{interaction} = NA$ |
| **Interaction index = 0.81** | **Interaction index = 0.90** |

1 $P$ for interaction, logistic regression adjusted for gender, age, and H. pylori infection status; Statistically significant associations were highlighted in bold ($P < 0.05$). CON: Controls; AG: Atrophic gastritis; GC: Gastric cancer; NA: Not available; GSTP1: Glutathione S-transferase pi 1; ERCC5: Excision repair cross complementing group 5.
Table 3  Epistatic effect of pair-wise interacting factors on the risk of atrophic gastritis and gastric cancer

| Interacted pair-wise SNPs      | Comparison                          | Subset                  | AG vs CON | GC vs CON |
|--------------------------------|-------------------------------------|-------------------------|-----------|-----------|
|                                |                                     |                         | P value   | OR (95%CI) | P value   | OR (95%CI) |
| ERCC5 rs2094258 interacted ERCC5 rs2094258 GG vs AG + AA | GSTP1 rs1695 AA                  | 0.006                   | 1.523     | (1.125-2.062) | 0.948     | 0.989     | (0.704-1.388) |
| with GSTP1 rs1695             | GSTP1 rs1695 GA + GG                | 0.205                   | 0.766     | (0.508-1.157) | 0.820     | 0.951     | (0.619-1.462) |
|                               | ERCC5 rs2094258 GG                  | 0.148                   | 1.343     | (0.901-2.002) | 0.808     | 1.035     | (0.685-1.625) |
| ERCC5 rs873601 interacted ERCC5 rs873601 GA + GG vs GA AA | GSTP1 rs1695 AA                  | 0.014                   | 0.678     | (0.497-0.926) | 0.799     | 1.045     | (0.745-1.465) |
| with GSTP1 rs1695             | GSTP1 rs1695 GA + GG                | 0.025                   | 0.678     | (0.483-0.952) | 0.148     | 0.756     | (0.517-1.105) |
|                               | ERCC5 rs873601 GA + GG              | 0.380                   | 1.230     | (0.775-1.931) | 0.872     | 0.961     | (0.592-1.560) |
|                               | GSTP1 rs1695 GA + GG                | 0.054                   | 0.758     | (0.571-1.005) | 0.955     | 0.991     | (0.730-1.346) |
|                               | ERCC5 rs873601 AA                   | 0.226                   | 1.356     | (0.828-2.222) | 0.542     | 1.183     | (0.690-2.027) |

All tests were adjusted by age, sex, and H. pylori infection. Statistically significant associations were highlighted in bold (P < 0.05). GC: Gastric cancer; AG: Atrophic gastritis; CON: Controls; ERCC5: Excision repair cross complementing group 5; GSTP1: Glutathione S-transferase pi 1.

which embody the effect of multipart interaction\[4\]. Multiple studies have revealed a relationship between epistasis and cancer risk\[25,26\]. Our previous findings also indicated epistasis by combining individual SNPs, which had no effect on disease risk at a single locus\[24,14\]. In the present study, for ERCC5 rs2094258 and GSTP1 rs1695, the AG/AA genotype of rs2094258 and AA genotypes of rs1695 were related to an increased risk of AG, but GA/GG genotypes of rs1695a were associated with a reduced risk of AG. For ERCC5 rs873601 and GSTP1 rs1695, AA genotype of rs873601 resulted in a reduced risk of AG, only in the presence of AA genotype of rs1695. Thus, there is still little direct evidence to reveal a specific functional association among the polymorphisms of ERCC5 and GSTP1. In light of previous research findings, we hypothesized an interaction effect between the DNA repair gene and xenobiotic metabolism gene by various signal pathways, and our discoveries regarding the interactions of the DNA repair ERCC5 gene and xenobiotic metabolism GSTP1 gene pathways may reveal the above assumption. Further synthetic and functional research on these two gene pathways will be performed to assess the interaction effect of susceptibility genes that directly affect gastric carcinogenesis.

Gastric carcinogenesis is also affected by environmental factors, in addition to genetic factors. H. pylori is considered a class I carcinogen by the World Health Organization and displays carcinogenic effects mediated by poisonous components\[27,28\]. Our previous studies have shown that the GSTP1 Val/Val genotype with smoking, alcohol consumption, or H. pylori IgG (+) could considerably increase AG and GC risk, and the NER SNPs (XPA rs2808668, DDB2 rs326222, rs3781619, rs830083, and XPC rs2607775) had interactive effects with alcohol consumption and smoking on AG or GC risk. In the present study, the cumulative effect was observed to be changed in subgroups with negative H. pylori infection status, which could imply an effect modification by H. pylori infection. Moreover, AG risk was significantly reduced, while one or two mutation genotypes were present (OR = 0.66 and 0.73, respectively). This suggested that H. pylori should be eliminated first for positive patients, which may be beneficial for reducing susceptibility to AG. However, we further analyzed the effect of three-dimensional interactions of the ERCC5 SNPs (rs2094258 and rs873601)-GSTP1 rs1695-environmental factors on the risk of AG, but no interaction effect was observed among them in terms of AG risk. This may be because our present sample size was relatively small, and some genotypes were scarce. In addition, there was a significant difference in H. pylori infection rates between the case groups and two matched control groups (P < 0.001). Although we performed all tests with an adjustment for H. pylori infection status except for those stratified by H. pylori, it still may be a limitation of the study. Nevertheless, it suggested that environmental factors were indispensable, although they cannot cause cancer or precancerous alone. Genetic susceptibility may play a vital role in gastric carcinogenesis. Further large-sample and comprehensive study of the function of environmental factors in SNP-SNP interactions of ERCC5 and GSTP1 is necessary and may partially remedy probable false-negative results in the study.

The present study has several limitations. First, even if our study included a relatively large sample, prospective studies consisting of larger-scale sample and multicenter surveys are necessary to validate the results of SNP-SNP interaction effects shown here. Second, since we only included one metabolic gene polymorphism (GSTP1 rs1695) of the GSTs in this study, further studies should involve other functional tagSNPs, such as GSTM1 and/or GSTT1, which should participate in SNP-SNP interactions between the DNA repair gene and xenobiotic metabolism gene pathways. In addition, the functions and mechanisms of the mentioned SNPs of the ERCC5 gene and GSTP1 gene pathways were not investigated and will require additional functional and molecular experiments to clarify.

In conclusion, we found for the first time that two pairwise interacting DNA repair gene ERCC5 SNPs (rs2094258 and rs873601) and metabolic gene GSTP1 rs1695 polymorphism combinations were related to increased or reduced AG risk. Moreover, the results
also demonstrated a significant difference in the cumulative effect in the *H. pylori*-negative subgroup on AG risk. The conclusions inferred from the present study about the effect of interactions between genetic polymorphisms may be conductive to proposing further studies to discover gene-gene interactions between DNA repair genes with xenobiotic metabolic gene pathways in gastric carcinogenesis.

### ARTICLE HIGHLIGHTS

**Research background**

Previous studies suggested that the interactions between various inherited susceptibility genes may affect carcinogenesis in individuals. Single nucleotide polymorphisms (SNPs) are widely applied to the research of tumor incidence and prognostic evaluation.

**Research motivation**

We aimed to assess gene interactions amongst inherited polymorphisms between DNA repair gene excision repair cross complementing group 5 (*ERCC5*) SNPs and glutathione S-transferase pi1 (*GSTP1*) rs1695 to explore their possibility of predicting gastric cancer (GC) risk and identify combination biomarkers for precancerosis and GC.

**Research objectives**

The objective was to investigate the impact of interactions of the DNA repair gene *ERCC5* with metabolic gene *GSTP1* on atrophic gastritis (AG) and GC risk.

**Research methods**

Seven *ERCC5* SNPs (rs1047768, rs2094258, rs2228959, rs4150291, rs4150383, rs751402, and rs873601) and *GSTP1* rs1695 SNP were detected using the Sequenom MassARRAY platform in 450 GC patients, 634 AG cases, and 621 healthy control subjects in a Chinese population.

**Research results**

Two pairwise combinations (*ERCC5* rs2094258 and rs873601 with *GSTP1* rs1695) influenced AG risk, and the *ERCC5* rs2094258-*GSTP1* rs1695 SNP pair demonstrated an antagonistic effect while *ERCC5* rs873601-*GSTP1* rs1695 shown a synergistic effect on AG risk. When the effect modification of *Helicobacter pylori* (*H. pylori*) infection was evaluated, the cumulative effect of one aforementioned pair-way interaction (*ERCC5* rs873601-*GSTP1* rs1695) showed a risk in the case

### Table 4 Cumulative effect of the interacting factors of *ERCC5* SNPs-*GSTP1* rs1696 on atrophic gastritis risk

| No. of interacting genotypes | Total population | *H. pylori*-negative subpopulation | *H. pylori*-positive subpopulation |
|------------------------------|------------------|-----------------------------------|----------------------------------|
|                              | Cases/controls   | OR (95%CI)                        | Cases/controls   | OR (95%CI) | Cases/controls | OR (95%CI) |
| ERCC5 rs2094258-*GSTP1* rs1696 |                  |                                   |                    |            |                  |            |
| 0                            | 132/162          | 0.81 (0.54-1.23)                  | 57/115            | 1 (Ref.)   | 75/47           | 1 (Ref.)   |
| 1                            | 369/310          | 1.48 (1.11-1.98)                  | 216/79            | 0.125      | 216/79          | 0.017      |
| 2                            | 133/148          | 1.05 (0.74-1.49)                  | 47/106            | 0.655      | 86/42           | 0.363      |
|                              | *P* = 0.582      |                                   |                    |            | *P* = 0.702     |            |
| ERCC5 rs873601-*GSTP1* rs1695 |                  |                                   |                    |            |                  |            |
| 0                            | 321/279          | 0.73 (0.57-0.94)                  | 136/206           | 1 (Ref.)   | 183/73          | 1 (Ref.)   |
| 1                            | 254/286          | 0.670 (0.60-1.40)                 | 99/201            | 0.061      | 155/85          | 0.090      |
| 2                            | 59/55            | 0.911 (0.67-1.26)                 | 20/45             | 0.150      | 39/10           | 0.241      |
|                              | *P* = 0.156      |                                   |                    |            | *P* = 0.043     |            |

1 Adjusted by sex, age, and *H. pylori* infection; 2 Adjusted by sex and age. Statistically significant associations were highlighted in bold (*P* < 0.05). CON: Controls; AG: Atrophic gastritis; *ERCC5*: Excision repair cross complementing group 5; *GSTP1*: Glutathione S-transferase pi 1.
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Research conclusions

DNA repair gene ERCC5 SNPs (rs2094258 and rs873601) and metabolic gene GSTP1 rs1695 polymorphism combinations were related to an increased or reduced AG risk. Moreover, the results also demonstrated a significant difference in the cumulative effect on AG risk in the H. pylori-negative subgroup.

Research perspectives

The interaction effects between genetic polymorphisms may be conductive to proposing further studies to discover gene-gene interactions between DNA repair genes and xenobiotic metabolic genes in gastric carcinogenesis.
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