ORIGINAL ARTICLE

Identification of new susceptibility loci for gastric non-cardia adenocarcinoma: pooled results from two Chinese genome-wide association studies

Zhaoming Wang,1,2,3 Juncheng Dai,3 Nan Hu,1 Xiaoping Miao,4 Christian C Abnet,1 Ming Yang,5 Neal D Freedman,1 Jinfei Chen,6 Laurie Burdette,1,2 Xun Zhu,3 Charles C Chung,1,2 Chuanli Ren,3,7 Sanford M Dawsey,1 Meilin Wang,8 Ti Ding,9 Jiangbo Du,3 Yu-Tang Gao,10 Rong Zhong,4 Carol Giffen,11 Wenting Pan,5 Woon-Puay Koh,12 Ningbing Dai,3 Linda M Liao,1 Caiwang Yan,3 You-Lin Qiao,13 Yue Jiang,3 Xiao-Ou Shu,14 Jiaping Chen,3 Chaoyu Wang,1 Hongxia Ma,3 Hua Su,1 Zhendong Zhang,8 Lemin Wang,1 Chen Wu,15 Yong-Bing Xiang,10 Zhibin Hu,3 Jian-Min Yuan,16,17 Lu Xie,3 Wei Zheng,14 Dongxin Lin,15 Stephen J Chanock,1 Yongyong Shi,18 Alisa M Goldstein,1 Guangfu Jin,3 Philip R Taylor,1 Hongbing Shen3

ABSTRACT

Objective Although several genome-wide association studies (GWAS) of non-cardia gastric cancer have been published, more novel association signals could be exploited by combining individual studies together, which will further elucidate the genetic susceptibility of non-cardia gastric cancer.

Design We conducted a meta-analysis of two published Chinese GWAS studies (2031 non-cardia gastric cancer cases and 4970 cancer-free controls) and followed by genotyping of additional 3564 cases and 4637 controls in two stages.

Results The overall meta-analysis revealed two new association signals. The first was a novel locus at 5q14.3 and marked by rs7712641 (per-allele OR=0.84, 95% CI 0.80 to 0.88; p=1.21×10^{-10}). This single-nucleotide polymorphism (SNP) marker maps to the intron of the long non-coding RNA, lnc-POLR3G-4 (XLOC_004464), which we observed has lower expression in non-cardia gastric tumour compared with matched normal tissue (P_{wilcoxon signed-rank}=7.20×10^{-4}). We also identified a new signal at the 1q22 locus, rs80142782 (per-allele OR=0.62; 95% CI 0.56 to 0.69; p=1.71×10^{-17}), which was independent of the previously reported SNP at the same locus, rs4072037 (per-allele OR=0.74; 95% CI 0.69 to 0.79; p=6.28×10^{-11}). Analysis of the new SNP conditioned on the known SNP showed that the new SNP remained genome-wide significant (P_{conditional}=3.47×10^{-6}). Interestingly, rs80142782 has a minor allele frequency of 0.05 in East Asians but is monomorphic in both European and African populations.

Conclusion These findings add new evidence for inherited genetic susceptibility to non-cardia gastric cancer and provide further clues to its aetiology in the Han Chinese population.

INTRODUCTION

Globally, gastric cancer remains the third-leading cause of cancer death in both sexes,1 with more than half of gastric cancer cases worldwide occurring in East Asia, predominantly in China. Most cases of gastric cancer are sporadic,2 and its aetiology is related to both genetic susceptibility and....

CrossMark
Significance of this study

How might it impact on clinical practice in the foreseeable future?

▸ The results from this GWAS meta-analysis will improve our understanding of the aetiology of non-cardia gastric cancer, which will further shed light on risk prediction and early detection of this malignancy.

epidemiological risk factors such as age, sex, Helicobacter pylori infection, family history, excessive salt intake and tobacco smoking. Anatomically, gastric cancer is classified into cardia gastric cancer and non-cardia gastric cancer, which are characterised by distinct risk factors and clinical features.  

Recently, several genome-wide association studies (GWAS) of gastric adenocarcinoma were conducted in East Asians. Notable findings include single-nucleotide polymorphism (SNP) markers mapping to 8q24.3, for both an intronic SNP (rs2976392) and an exonic SNP (rs2294008) in the Prostate Stem Cell Antigen gene (PSCA); and two markers rs2075570 and rs2070803 near the Mucin 1 gene (MUC1) on 1q22.9. The findings on 1q22 locus and gastric cancer risk were further replicated in several follow-up studies and additional evidence pointed to the non-synonymous SNP rs4072037, as the functional variant underlying the observed association. In addition, 3q13.31 marked by rs9841504, 5p13.1 marked by rs13361707 or rs10074991 and 6p21.1 marked by rs2294693 were reported to be associated with non-cardia gastric adenocarcinoma in China, whereas 10q23 marked by rs2274223, a functional variant underlying the observed association.  

For the Nanjing and Beijing GWAS, individuals were derived from four prospective cohort studies and one large case–control study as reported in Abnet et al. In addition, all subjects used in replication in the original paper were subsequently genotyped using the Illumina 660W-Quad microarray; this included scanning of 725 additional gastric cancer cases and 608 additional controls. The current analysis included all 1025 non-cardia gastric cancer cases and 2697 controls from the NCI Upper Gastrointestinal Cancer GWAS.  

For the Nanjing and Beijing GWAS, individuals were derived from separate case–control studies conducted in Nanjing (565 cases and 1162 controls) and Beijing (468 cases and 1123 controls), as previously reported in Shi et al, where individuals were genotyped using the Affymetrix Genome-Wide Human SNP Array (V6.0).

Repetition samples

The first-stage replication including 1145 cases and 2253 controls were derived from Jiangsu Province. The second-stage replication included 2419 cases and 2384 controls, which were derived from Beijing, Hubei and Shandong province.

Gastric cancer cases tested for expression of the lncRNA associated with the GWAS SNPs came from our UGI Cancer Genetic Studies (URL: http://dceg.cancer.gov/about/staff-directory/biographies/O-Z/taylor-philip). Genotypes for all these cases are known because they were also all participants in our previous GWAS.  

All study individuals provided informed consent and both the institutional review boards of NCI and Nanjing Medical University approved all procedures and all experiments, which were conducted in accordance with the approved guidelines.

SNP selection and replication genotyping

The meta-analysis included 6223 896 SNPs based on the intersection of the three imputed datasets. Individual SNPs for the stage 2A replication were selected based on the following criteria: (1) SNPs with INFO score of ≥0.5; (2) MAF in control set ≥0.01; (3) p value for HWE in control set >1.0×10⁻⁴ in each set; (4) I² >1.0×10⁻⁴ and I² ≤75% in meta-analysis; (5) linkage disequilibrium (LD) pruning: included only one SNP from any region or allele frequency matching for adenine/thymine (A/T) and guanine/cytosine (G/C) SNPs. We implemented a 4 Mb sliding window to impute across the genome, resulting in 744 windows. A pre-phasing strategy with SHAPEIT software (V2) was adopted to improve the imputation performance. The phased haplotypes from SHAPEIT were fed directly into IMPUTE2.

Genotype imputation

In addition to the quality control procedures performed in the previous primary publications for previous GWAS, SNPs with call rate of <95%, p value for Hardy–Weinberg Equilibrium (HWE) in controls ≤1.0×10⁻⁶ or minor allele frequency (MAF) of <1% in controls were further removed before imputation. Imputation was conducted separately for the NCI (Illumina 660W) and the Nanjing+Beijing (Affymetrix 6.0) scan data taking all populations in the 1000 Genomes Project Phase I (V3) as the reference set and using IMPUTE2 software (V2.2.2), which automatically finds haplotypes from the best matching population from the entire reference set to do the imputation.

First, genomic coordinates for National Center for Biotechnology Information (NCBI) human genome Build 36 were converted to those of NCBI human genome Build 37 using the UCSC liftOver tool. The few loci for which coordinates could not be converted were also excluded from imputation. Second, the strand of the inference data was aligned with the 1000 Genomes Project data by simple allele state comparison or allele frequency matching for adenine/thymine (A/T) and guanine/cytosine (G/C) SNPs. We implemented a 4 Mb sliding window to impute across the genome, resulting in 744 windows. A pre-phasing strategy with SHAPEIT software (V2) was adopted to improve the imputation performance. The phased haplotypes from SHAPEIT were fed directly into IMPUTE2.
cases and 2384 controls) (see online supplementary table S3). Further information on primers and probes are available on request. For quality control purposes: (1) case and control samples were mixed on each plate; (2) genotyping was performed blind to case/control status and (3) two water controls were used in each plate as blank controls.

Quantitative reverse transcription-PCR

Total RNA was extracted from each patient’s matched frozen tumour and normal surgical resection tissues using All Prep DNA/RNA/Protein kit (QIAGEN) in accordance with the manufacturer’s instructions. RNA quality and quantity were determined using the RNA Nano Chip/Agilent 2100 Bioanalyzer (Agilent Technologies). Reverse transcription of RNA was done by adding 0.2–2 μg total RNA, 1 μL of oligo(dT)12–18 (500 μg/mL), 1 μL (200 units) of SuperScript II reverse transcriptase, 1 μL (2 units) of E-coli RNase and 1 μL of 10 mmol/L deoxynucleotide triphosphate (Invitrogen) in total volume of 20 μL. All real-time PCRs (RT-PCRs) were done using an ABI 7300 Sequence Detection System. Primer and probe for the target gene and the internal control gene (GAPDH, glyceraldehyde-phosphate dehydrogenase) were designed and ordered from ABI (Assay ID: ABBXUX; Part Number: 44411114). A singleplex reaction mix was prepared according to the manufacturer’s protocol of ‘Assays-on-Demand Gene Expression Products’, including 10 μL Taqman Universal PCR Master Mix, No AmpErase UNG (2X), 1 μL of 20X Assays on-Demand Gene Expression Assay Mix (all Gene Expression assays have a 6-carboxyfluorescein (FAM) reporter dye at the 5’ end of the TaqMan minor groove binder (MGB) probe and a non-fluorescent quencher at the 3’ end of the probe) and 9 μL of cDNA (1000 ng) diluted in RNase-free water to a total volume of 20 μL. Each sample for the gene was run in triplicate and the expression level was averaged over all runs. The thermal cycling conditions included an initial denaturation step at 95°C for 10 min, 40 cycles at 95°C for 15 s and 60°C for 1 min.

Statistical analysis

Association testing was performed using SNPTTEST software (V2.2), with adjustment for age, sex and study variables for NCI. Two eigenvectors (ev4 and ev8) were significantly associated with case status (p<0.05) in the baseline model (not including SNP effects) which was adjusted for age, sex, study and all top 10 eigenvectors, and therefore these two significant eigenvectors were also included to adjust for population stratification in final association models. For Nanjing and Beijing, age, sex, smoking and alcohol consumption were adjusted in baseline models. Three eigenvectors (ev1, ev4 and ev9) for Nanjing and five eigenvectors (ev1, ev3, ev7, ev9 and ev10) for Beijing were adjusted for population stratification separately, which were also significantly associated with case status (p<0.05). In each replication study, we adjusted for gender, age, smoking and alcohol consumption only.

For the meta-analysis we used the meta-module implemented in genotype utilities (GLU) (see URLs). Strand flipping was handled by comparing alleles either with direct matching or with reverse complement matching. For A/T or G/C SNPs, strand matching was based on allele frequency checking. The fixed-effects inverse variance method was used to combine the $\beta$ estimates and SEs from each GWAS scan as well as the replication stages. The $p$ value for heterogeneity was calculated using Cochran’s Q, which is distributed as a $\chi^2$ statistic with (n–1) degrees of freedom, where n is the number of sets included in the meta-analysis. $I^2$ was calculated as $100\% \times (Q-(n-1))/Q$. Data analysis and management was performed with GLU or PLINK (see URLs).

PLINK was also used for the conditional haplotype analysis. Wilcoxon signed-rank test (R package) was applied to the tumour/normal paired quantitative RT-PCR data to assess the RNA expression level.

In silico bioinformatics analysis

We used GTeX (see URLs) for expression quantitative trait loci (eQTL) information for associated SNPs (see online supplementary table S5). We also searched HaploReg (V3.3) to explore potential functional annotations within Encyclopedia of DNA Elements (ENCODE) data for the genomic regions surrounding our lead SNPs (see online supplementary table S6).

RESULTS

To discover additional susceptibility alleles for non-cardia gastric cancer in the Han Chinese population, we conducted a combined analysis of two previously published GWAS10 11 after imputing the genetic data with the 1000 Genome Project data Phase I release (V3).10 The combined dataset included a total of 6 223 896 SNPs for a fixed-effects meta-analysis of 2031 cases and 4970 cancer-free controls (see online supplementary table S1). Quantile-quantile and Manhattan plots based on stage 1 meta-analysis $p$ values are shown in online supplementary figures S1 and S2, respectively. We further followed up 37 promising loci (see Methods) and genotyped them in an independent set of 1145 cases and 2253 controls in stage 2A (see online supplementary table S2). Finally, for stage 2B, we advanced five loci that were nominally significant in stage2A to a second independent set of 2419 cases and 2384 controls (see online supplementary table S3).

Based on the overall meta-analysis including two discovery GWAS scans and two replication studies, we identified two novel risk loci for non-cardia gastric cancer, the first one is rs7712641 at 5q14.3 (per-allele OR=0.84, 95% CI 0.80 to 0.88; $p=1.21\times10^{-11}$). No heterogeneity was observed across the two GWAS scans and two replication studies ($p_{het}=0.56$) (table 1). There are no protein-coding genes within the 1 Mb of the associated SNP (chr5: 88 346 298–88 459 630; hg19) (figure 1A). However, rs7712641 is located in the intron of lnc-POLR3G-4 (XLOC_004464), a long non-coding RNA (lncRNA) which is poorly characterised. To explore the possible effect of the SNP marker on the lncRNA, lnc-POLR3G-4, we extracted total RNA from 75 matched gastric non-cardia adenocarcinoma and adjacent normal tissue pairs and performed a quantitative RT-PCR analysis to measure its expression abundance. We found that expression differed between tumour and normal tissues ($p_{\text{Wilcoxon signed-rank}}=7.2\times10^{-4}$); with the majority of pairs (50 of 73) showing lower expression in tumour compared with normal tissue (figure 2). These data provided preliminary evidence that this lncRNA could function in a manner resembling a tumour suppressor gene. However, the association between rs7712641 and expression of lnc-POLR3G-4 in normal tissue was negative ($p=0.99$), which does not support the notion of a functional role for this SNP. More functional studies are warranted to clarify the complicated phenomena.

As anticipated, the current study, which included samples from these previous GWAS reports,10 11 also replicated the association with rs4072037 (table 1; per-allele OR=0.74; 95% CI 0.69 to 0.79; $p=6.28\times10^{-17}$) at 1q22. However, we also identified a second strong signal in this region and by doing so established an independent, new genome-wide significant SNP rs80142782 (table 1 and figure 1B; per-allele OR=0.62; 95%
CI 0.56 to 0.69; p=1.71×10^{-19}. Based on the evidence at hand, it seems that rs80142782 is likely an independent primary signal (rs4072037 as a secondary signal) at this locus. This evidence includes the following: (1) both SNPs are about 323 kb apart and have moderately low (r^2=0.3) pair-wise LD in 1000 Genome s Project data for Asians; (2) rs80142782 conditioned on rs4072037 remained genome-wide significant (P\textsubscript{conditional}=3.47×10^{-6}, see online supplementary table S5), although rs4072037 conditioned on rs80142782 did not (P\textsubscript{conditional}=2.95×10^{-6}, see online supplementary table S4); (3) we used the haplotype inference method implemented in the plink haplotype test — chap option. For rs4072037 and rs80142782, there are three inferred haplotypes with frequencies greater than 1% from all four possible ones. The two models compared in the conditional haplotype likelihood ratio test are the null model: {CC} {CT} {TT} and the alternative model: {CC} {CT} {TT}, where each {set} allows a unique effect. The conditional haplotype analysis demonstrated that the effect size of haplotype CT differed from that of TT risk, it is also interesting to note that GTEx data show that rs4072037 is an eQTL for several neighbouring genes (including \textit{THBS3}, \textit{GBP1}, \textit{GBA} and \textit{RP11-263K19.4}) in other tissues (see online supplementary table S5). The rs80142782 may act on the \textit{ASHH} gene based on its close proximity. \textit{ASHH} encodes a member of the trihorax group of transcriptional activators and functions as an epigenetic regulator by histone methylation (H3K4 methyltransferase) and is frequently altered in lung cancer tumours and cell lines,24 25 oesophageal squamous cell carcinoma tumour tissue23 and colorectal cancer cell lines.8 It was also implicated in inflammatory autoimmunity disease.26

**DISCUSSION**

Our study identified a new risk locus at 5q14.3 marked by rs7712641 which lies in the intron of a lncRNA with little known prior functional characterisation. Other lncRNAs implicated in cancers include \textit{PCA3} and \textit{PCGEM1} in prostate tumour,26 and \textit{MALAT1} in tumours of the colorectum, liver, pancreas, lung, breast and prostate.27 28 It is remarkable that a recent comprehensive transcriptome analysis nominated a total of 7942 lineage-associated or cancer-associated lncRNA genes,29 for which further functional investigations are warranted.
Overexpression or knockdown of this lncRNA may be informative in identifying target genes through analysis of differential gene expression profiles in non-cardia gastric tumour cell lines.

Our analysis also revealed an apparently stronger associated SNP (rs80142782) at 1q22 than the previously identified rs4072037. Our data indicate that the association with
Figure 2  Differential expression of Inc-POLR3G-4 between non-cardia gastric tumour and normal tissue based on a quantitative real-time PCR analysis. X axis is the log2 of tumour:normal fold change. All 75 pairs were sorted in increasing order of fold change and plotted along the Y axis. Two-thirds of pairs show lower expression in tumour compared with normal, while expression is higher in normal compared with tumour in one-third of pairs. Highlighted in the middle are 15 pairs with fold changes between 0.5 and 2 (or −1 to 1 in log2 scale). Based on all 75 pairs, the Wilcoxon signed-rank test was P=7.2×10−4 and the median tumour:normal fold change is 0.3.

rs80142782 is independent of rs4072037. Both the new locus at 5q14.3 marked by rs7712641 and the new independent signal at 1q22 marked by rs80142782 could contribute to epigenome regulation. Haploreg data show that both of these SNPs (or SNPs in high LD with them) locate to sites of multiple regulatory elements, including promoter histone marks, enhancer histone marks and DNAse hypersensitivity (see online supplementary table S6). Further functional validation studies are warranted to understand the contribution of these susceptibility alleles to gastric carcinogenesis.

Online supplementary table S7 shows the results from our meta-analysis of two GWAS scans for previously reported variants from the literature. Notably we confirmed a prior independent GWAS report of an association between multiple SNPs in PSCA at 8q24.3 and risk of non-cardia gastric cancer. We also confirmed the association for rs13361707 in PRKAA1, but there was no additional evidence to support an association for rs9841504 in PSCA or SNPs in high LD with them) locate to sites of multiple regulatory elements, including promoter histone marks, enhancer histone marks and DNAse hypersensitivity (see online supplementary table S6). Further functional validation studies are warranted to understand the contribution of these susceptibility alleles to gastric carcinogenesis.

In summary, by combining two pre-existing GWAS scans of non-cardia gastric cancer to increase the sample size for the discovery stage and adding over 8000 individuals for further replication, we identified two novel loci. In the future, additional studies are warranted with larger sample size and/or with a design that considers heterogeneity of the gastric cancer where, for example, the Cancer Genome Atlas (TCGA) recently reported four molecular subtypes for gastric cancer based on multi-omics profiling analyses.

**URLS**

- **GTEx**, http://www.gtexportal.org/
- **GTEx MUC1 expression**, http://www.gtexportal.org/home/gene/MUC1
- **GLU**, https://code.google.com/p/glucogenetics/
- **IMPUTE2 software** (V2.2.2), https://mathgen.stats.ox.ac.uk/impute/impute_v2.html
- **LocusZoom**, http://csg.sph.umich.edu/locuszoom/
- **PLINK**, http://pngu.mgh.harvard.edu/~ purcell/plink/
- **SHAPEIT software** (V2), https://mathgen.stats.ox.ac.uk/genetics_software/shapeit/shapeit.html
- **SNPTEST software** (V2.2), https://mathgen.stats.ox.ac.uk/genetics_software/snptest/snptest.html
- **UCSC liftOver tool**, http://hgdownload.cse.ucsc.edu/downloads.html

**Author affiliations**

1Division of Cancer Epidemiology and Genetics, National Cancer Institute, National Institutes of Health, Bethesda, Maryland, USA
2Cancer Genomics Research Laboratory, Leidos Biomedical Research Inc., Gaithersburg, Maryland, USA
3Department of Epidemiology and Biostatistics, Beijing Kadoorie Key Lab of Cancer Biostatistics, Prevention and Treatment, Collaborative Innovation Center for Cancer Medicine, School of Public Health, Nanjing Medical University, Nanjing, People’s Republic of China
4Department of Epidemiology and Biostatistics, and the Ministry of Education Key Lab of Environment and Health, School of Public Health, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, People’s Republic of China
5State Key Laboratory of Chemical Resource Engineering, Beijing Laboratory of Biomedical Materials, College of Life Science and Technology, and Beijing University of Chemical Technology, Beijing, People’s Republic of China
6Department of Oncology, Nanjing First Hospital, Nanjing Medical University, Nanjing, People’s Republic of China
7Clinical Medical Testing Laboratory, Northern Jiangsu People’s Hospital and Clinical Medical College of Yangzhou University, Yangzhou, People’s Republic of China
8Department of Genetic Toxicology, the Key Laboratory of Modern Toxicology of Ministry of Education, School of Public Health, Nanjing Medical University, Nanjing, People’s Republic of China
9Shanxi Cancer Hospital, Taiyuan, Shanxi, P.R. China
10Shanghai Cancer Institute, Shanghai, People’s Republic of China
11Information Management Services Inc., Silver Spring, Maryland, USA
12Duke-NUS Graduate Medical School, Singapore, and Saw Swee Hock School of Public Health, National University of Singapore, Singapore
13Department of Epidemiology, Cancer Institute and Hospital Chinese Academy of Medical Sciences, Beijing, People’s Republic of China
14Department of Medicine and Vanderbilt-Ingram Cancer Center, Vanderbilt University, Nashville, Tennessee, USA
15Department of Etiology & Carcinogenesis, Cancer Institute and Hospital Chinese Academy of Medical Sciences, Beijing, People’s Republic of China
16Division of Cancer Control and Population Sciences, University of Pittsburgh Cancer Institute; Pittsburgh, Pennsylvania, USA
17Department of Epidemiology, Graduate School of Public Health, University of Pittsburgh, Pittsburgh, Pennsylvania, USA
18Key Laboratory for the Genetics of Developmental and Neuropsychiatric Disorders, Ministry of Education, Bio-X Institutes, Shanghai Jiao Tong University, Shanghai, People’s Republic of China

**Contributors**

HBS, PRT, GT, AMG, YS, SIC, ZW, JCD, NH, XM, CCA, MY, NDF and JFC organised and designed the study; ZW, JCD, HBS, PRT, AMG and SJT wrote the first draft of the manuscript; ZW and JCD contributed to the design and execution of statistical analyses; HBS, PRT, GT, AMG, YS, SIC, ZW, JCD, NH, XM, CCA, MY, NDF and JFC contributed to the writing of the manuscript; ZW, JCD, HBS, GT, ZH and LB conducted and supervised the genotyping of samples; HS, LW and NH conducted the quantitative real-time PCR experiments; MY, Zl, CIC, CR, SMD, MW, TD, JBD, etc.
REFERENCES


