Association of CHRNA5-A3-B4 Variation with Esophageal Squamous Cell Carcinoma Risk and Smoking Behaviors in a Chinese Population

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Abstract

Background: CHRNA5-A3-B4, the gene cluster encoding nicotinic acetylcholine receptor subunits, is associated with lung cancer risk and smoking behaviors in people of European descent. Because cigarette smoking is also a major risk factor for esophageal squamous cell carcinoma (ESCC), we investigated the associations between variants in CHRNA5-A3-B4 and ESCC risk, as well as smoking behaviors, in a Chinese population.

Methods: A case-control study of 866 ESCC patients and 952 healthy controls was performed to study the association of polymorphisms (rs667282 and rs3743073) in CHRNA5-A3-B4 with cancer risk using logistic regression models. The relationships between CHRNA5-A3-B4 polymorphisms and smoking behaviors that can be quantified by cigarettes smoked per day (CPD) and pack-years of smoking were separately estimated with Kruskal-Wallis tests among all 840 smokers.

Results: CHRNA5-A3-B4 rs667282 TT/TG genotypes were associated with significantly increased risk of ESCC [adjusted odds ratio (OR) = 1.32, 95% confidence interval (CI) = 1.03 – 1.69, P = 0.029]. The increased ESCC risk was even higher among younger subjects (<60 years) [OR = 1.44, 95% CI = 1.04 – 1.98, P = 0.024]. These effects were not found in another polymorphism rs3743073. No evident association between the two polymorphisms and smoking behaviors was observed.

Conclusions: These results support the hypothesis that CHRNA5-A3-B4 is a susceptibility gene cluster for ESCC. The relationship between CHRNA5-A3-B4 and smoking behaviors in a Chinese population needs further investigation.

Introduction

Esophageal cancer is the eighth most common cancer in the world, and there were 402,300 newly diagnosed cases worldwide in 2008 [1,2]. Of the two major histological types, esophageal squamous cell carcinoma (ESCC) rather than adenocarcinoma is common worldwide [3], and it is relatively more prevalent among the East Asian population, including the Chinese. Although the exact etiology of ESCC remains to be identified, cigarette smoking has been demonstrated as the major factor that contributes to the development of ESCC in accumulating epidemiological and clinical studies [1]. We can not exclude the possibility that genes in association with smoking may modify the susceptibility to ESCC.

The nicotinic acetylcholine receptor (nAChR) belongs to the superfamily of ligand-gated ion channels, and it is activated by acetylcholine (Ach), choline and nicotine. It is involved in the regulation of nicotine and nitrosamines [4]. Nicotine in tobacco triggers the psychological and neurobiological effects that are associated with smoking consumption and addiction [5], while nitrosamines are important carcinogens in tobacco that contribute to the development of many smoking-related cancers, including lung cancer and ESCC [6]. Furthermore, the nAChR pathway is related to tumor cell proliferation, apoptosis, survival, migration, invasion, and angiogenesis [7,8], which can affect the progression of cancer patients. Currently, nAChR is thought to be an important regulator of a complex network of neurotransmitters that govern the synthesis and release of growth, angiogenic and neurotrophic factors in cancer cells, the cancer microenvironment and distinct organs [9].
Recently, genome-wide association (GWA) studies have shown three single nucleotide polymorphisms (SNPs) (rs1051730, rs16969968 and rs8034191) in the CHRNA5-A3-B4 gene cluster, which encodes the nAChR subunits to be related to lung cancer risk and smoking consumption and addiction behaviors in people of European descent [10–13]. Because these three variants are extremely rare in the Asian population, according to the HapMap database and a study of Wu et al [14], they play little roles, if any, in risk of smoking-related cancers and smoking behaviors in Chinese people. Interestingly, two independent studies found that variants rs667282 and rs3743073 of CHRNA5-A3-B4 can affect lung cancer risk in a Chinese population [14,15], similar to rs1051730, rs16969968 and rs8034191 in people of European descent. Consequently, we investigated the associations of the 2 SNPs at CHRNA5-A3-B4 (rs667282 and rs3743073) with ESCC risk and smoking behaviors in a Chinese population, and we further explored the influence of CHRNA5-A3-B4 polymorphisms on cancer progression.

Materials and Methods

Ethics Statement

The study is in compliance with the Helsinki Declaration, and was approved by the Ethics Committees of Qilu Hospital of Shandong University (approval number: KYLL2010058). At recruitment, all participants gave written informed consent.

Study Sample

This study consisted of 866 ESCC patients and 952 healthy controls. All subjects were biologically unrelated ethnic Han Chinese from Shandong Province in China. Patients who were newly diagnosed with histologically confirmed primary ESCC were recruited from Qilu Hospital of Shandong University, Shandong Tumor Hospital and Institute and the Second Hospital of Shandong University from 2010 to 2012. Controls were healthy individuals selected from the physical examination center of Qilu Hospital during the same period and matched with the patients by age and gender.

Clinical Data Collection

We collected subject data through face-to-face interviews conducted by trained medical students or clinical doctors with a pre-tested standardized questionnaire regarding age, gender, drinking status, smoking status, smoking dose, and smoking exposure. Smokers or drinkers were defined as those who had been regularly smoking or drinking for 1 year or longer in their lifetime, whereas nonsmokers or nondrinkers were defined as those who had not. ESCC patients were staged according to the American Joint Committee on Cancer (AJCC) tumor-node-metastasis (TNM) staging system whenever possible from operative specimens [16]. Because positron emission tomography and endoscopic ultrasonography were not required as a component of the pretreatment evaluation at the time of study, we staged inoperable patients, who were not applicable for the AJCC staging system, using computer tomography [17,18]. Both staging methods are accepted by oncologists [19-21].

Genotyping

Whole-blood samples were collected from all study participants. We extracted genomic DNA from the blood samples using the Tiangen Biotech kit (DP319; Tiangen Biotech (Beijing) Co., Ltd., Beijing China). Rs667282 and rs3743073 were genotyped by the 5’ nuclease cleavage assay (TaqMan method) obtained from Applied Biosystems (assay ID: C__25648144_10, respectively), which uses two allele-specific TaqMan MGB probes and a PCR primer pair to detect the specific SNP target. Following the manufacturer’s instructions, PCR amplifications were conducted in a 96-well Applied Biosystems 7500 Real Time PCR System, and allelic discrimination was performed using the SDS 1.4 software.

Statistical Analysis

The genotype frequencies of each SNP were tested for deviation from the Hardy–Weinberg equilibrium using a goodness-of-fit chi-square test. This method was used to determine whether the samples were from a population and representative. The association of each SNP with ESCC risk was estimated using a logistic regression model with adjustment for age, sex, smoking status, and drinking status. Associations between the 2 SNP genotype groups with the histological grade and clinical stage of ESCC were evaluated using the chi-square test. The Kruskal–Wallis test was used to compare the difference of cigarettes per day (CPD) and pack-years of smoking. Individuals who denied ever smoking were excluded from this analysis, because they may have never had sufficient exposure to cigarette smoking to become addicted [11,22]. Statistical analyses were performed using SPSS (version 20) and Stata (version 12.0). P<0.05 was the criterion of statistical significance, and all statistical tests were two-sided.

Results

Baseline Characteristics of the Study Population

The frequency distributions of the baseline characteristics and clinical features of the study population are presented in Table 1. There were no significant differences between cases and controls in terms of age or gender distribution (P=0.86 and 0.75, respectively). In contrast, smoking and drinking were significantly more frequent in cancer patients compared to control subjects (both P<0.001). Among the ESCC patients, the fractions of histological grades from well to poor were 21.2%, 48.7% and 30.0%, respectively. Approximately 52.3% and 47.7% of ESCC patients were at the early clinical stage (I – II) and advanced stage (III – IV), respectively.

Association between CHRNA5-A3-B4 Polymorphisms and ESCC Risk

The genotype distribution of rs667282 and rs3743073 among cases and controls and their association with ESCC risk are shown in Table 2. The observed genotype frequencies for both polymorphisms were in Hardy-Weinberg equilibrium among the controls (P=0.33 for rs667282 and P=0.18 for rs3743073). The genotypes of rs667282 were markedly distinct in cases and controls irrespective of age, gender, smoking status and drinking status, whereas the allele distribution of rs3743073 was similar between cases and controls. We calculated the odds ratios adjusted by age, sex, drinking status and smoking status. The TT/TC genotypes of rs667282 exhibited an increased association with ESCC (adjusted OR = 1.32, 95%CI = 1.03 to 1.69, P=0.029). Rs3743073 did not show a significant association with the risk of ESCC.

We further examined the impact of CHRNA5-A3-B4 (rs667282) on ESCC risk, stratified by age, gender, drinking status and smoking status (Table 3). The increased cancer risk accompanied by rs667282 TT/TC genotypes was more notable in younger subjects (≤60 years) (OR = 1.44, 95%CI = 1.04 to 1.98, P=0.024). No significant discrepancy was observed in the stratification of gender, drinking status and smoking status.
Because the CHRNA5-A3-B4 gene cluster has been suggested to influence cancer progression, we also investigated the association of the 2 SNPs in CHRNA5-A3-B4 with the histological grade and clinical stage of ESCC patients. As shown in Table 4, we found that there was a weak association between the TT/TC genotypes and ESCC risk.

Table 1. The baseline characteristics of the study population.

| Characteristic          | ESCC subjects | Control subjects | P \(^1\) |
|-------------------------|---------------|------------------|---------|
| Age, mean (SD)          | n = 866       | n = 952          | 0.86    |
| Gender, N (%)           |               |                  |         |
| Male                    | 746 (86)      | 825 (87)         | 0.75    |
| Female                  | 120 (14)      | 127 (13)         |         |
| Smoking status, N (%)   |               |                  |         |
| Never smoker            | 322 (37)      | 656 (69)         | <0.001  |
| Current smokers         | 416 (48)      | 243 (25.5)       |         |
| Former smokers          | 128 (15)      | 53 (5.5)         |         |
| Drinking status, N (%)  |               |                  |         |
| Never drinking          | 333 (38)      | 597 (63)         | <0.001  |
| Ever drinking           | 533 (62)      | 355 (37)         |         |
| Smoking variables, mean (SD) |       |                  |         |
| Cigarettes per day      | 23.1 (11.4)   | 22.2 (11.5)      | 0.22    |
| Smoking duration in years | 32.5 (9.7)   | 31.5 (9.5)       | 0.13    |
| Pack-years              | 37.0 (20.2)   | 33.9 (19.8)      | 0.01    |
| Differentiation, N (%)  |               |                  |         |
| Well                    | 184 (21.2)    |                  |         |
| Moderate                | 422 (48.7)    |                  |         |
| Poor                    | 260 (30.0)    |                  |         |
| Clinical stage, N (%)   |               |                  |         |
| I – II                  | 453 (52.3)    |                  |         |
| III – IV                | 413 (47.7)    |                  |         |

\(^1\)Two-sided \(\chi^2\) test for the categorical variables (sex, smoker status and drinking status), and Kruskal-Wallis test for the continuous variables (age and smoking variables).

Table 2. The genotype distribution of 2 SNPs among cases and controls and their association with ESCC risk.

| Genotype group     | ESCC subjects | Control subjects | Adjusted OR (95% CI) \(^1\)  | P       |
|--------------------|---------------|------------------|--------------------------------|---------|
|                    | n = 866       | n = 952          |                                |         |
| rs667282, N (%)    |               |                  |                                |         |
| CC                 | 156 (18.0)    | 205 (21.6)       | Reference                      |         |
| TC                 | 419 (48.4)    | 457 (48.1)       | 1.29 (0.99–1.67)               | 0.062   |
| TT                 | 291 (33.6)    | 289 (30.4)       | 1.36 (1.03–1.80)               | 0.030   |
| TT/TC              | 710 (82.0)    | 746 (78.5)       | 1.32 (1.03–1.69)               | 0.029   |
| rs3743073, N (%)   |               |                  |                                |         |
| TT                 | 225 (26.1)    | 271 (28.6)       | Reference                      |         |
| TG                 | 424 (49.2)    | 453 (47.7)       | 1.15 (0.91–1.46)               | 0.249   |
| GG                 | 213 (24.7)    | 225 (23.7)       | 1.10 (0.84–1.45)               | 0.470   |
| GG/TG              | 637 (73.9)    | 678 (71.4)       | 1.13 (0.91–1.41)               | 0.267   |

Association of CHRNA5-A3-B4 Polymorphisms with Histological Grade and Clinical Stage of ESCC Patients

Because the CHRNA5-A3-B4 gene cluster has been suggested to influence cancer progression, we also investigated the association of the 2 SNPs in CHRNA5-A3-B4 with the histological grade and clinical stage of ESCC patients. As shown in Table 4, we found that there was a weak association between the TT/TC genotypes and ESCC risk.
Discussion

In this hospital-based case-control study of Chinese individuals, we found that the SNP rs667282 in CHRNA5-A3-B4, the gene cluster encoding nAChR subunits (alpha3, alpha5, and beta4), exhibited significant associations with ESCC risk. These nAChR subunits are the principle targets of nicotine, and can also bind to two nicotine-specific metabolites namely: 4-(Methylnitrosamino)-1- (3-pyridyl)-1- butanone (NNK) and N-nitrosonornicotine (NNN) [12,23], which are potent carcinogens and the effects of which on DNA are regarded as the primary cause of smoking-related cancers [24]. As the agonists for nAChRs, NNK and NNN have been found to induce many types of smoking-related cancers in laboratory animals [25,26]. Recently, Yuan et al. found that NNN exhibited a significant and important role in esophageal carcinogenesis in humans [27]. Furthermore, NNN showed a 5000× higher affinity for nAChRs compared to nicotine [28]. Thus, it is biologically plausible that variants in the gene cluster confer individuals’ susceptibility to esophageal cancer, especially ESCC.

The relationship between cancer and carcinogen-related genes has been well established in many types of cancers [29–31]. Recently, a GWA study in a Japanese population showed that two SNPs in the ADH1B and ADH1B, which encode dehydrogenases involved in alcohol metabolism, were strongly associated with ESCC risk [32]. These results indicate that ESCC may also be influenced by genetic polymorphisms of carcinogen-related genes, although ADH1B and ADH1B did not exhibit associations with ESCC in a subsequent study of Chinese subjects [33].

In the stratified analysis, associations between rs667282 genotypes and ESCC risk were stronger for younger subjects, lighter smokers or drinkers. One thing to note is the higher risk tendency for lighter smokers than heavier smokers. The pattern of higher genetic risk in the low-carcinogen exposure stratum supports the hypothesis that genotype contributes to the risk of carcinogenesis directly, rather than simply indirectly through an association with altered smoking quantity. This phenomenon has also been observed for the key SNP rs1051730 in CHRNA5-A3-B4 in lung cancers in people of European descent [34,35]. In addition, we also found that rs667282 tended to be a risk factor for advanced tumor clinical stage, although the influence was not significant. To date, studies investigating the CHRNA5-A3-B4 polymorphisms in relation to tumor clinical stage have been few; however, it has been proposed that nicotine after initiation may contribute to the progression phase of cancer development because nicotine promotes the growth of cancer cells and the proliferation of endothelial cells in vivo [36]. Furthermore, a negative correlation between CHRNA5-A3-B4 polymorphisms and lung cancer survival has been shown in many studies [37,38]. If the tendency can be confirmed by additional studies with a larger sample, rs667282 might help to accurately predict the clinical course of ESCC.

Table 4. Association between 2 SNPs genotype groups with histological grade and clinical stage of ESCC.

| Category | rs667282 | rs3743073 |
|----------|----------|-----------|
|          | CC       | TT/TC     | OR (95% CI)* | P  | TT   | GG/TG | OR (95% CI) * | P  |
| Differentiation, N (%) |          |           |            |    |       |       |            |    |
| Well     | 35       | 149       | Reference  |     | 46    | 138   | Reference  |     |
| Moderate | 76       | 346       | 1.06 (0.68–1.67) | 0.767 | 115   | 306   | 0.89 (0.60–1.32) | 0.553 |
| Poor     | 45       | 215       | 1.12 (0.69–1.83) | 0.643 | 64    | 193   | 1.01 (0.65–1.56) | 0.981 |
| Clinical stage, N (%) |          |           |            |    |       |       |            |    |
| I – II   | 90       | 363       | Reference  |     | 122   | 330   | Reference  |     |
| III – IV | 66       | 347       | 1.30 (0.78–1.85) | 0.137 | 103   | 307   | 1.10 (0.81–1.50) | 0.533 |

* CI: confidence interval; OR: odds ratio.

Table 5. Association between 2 SNPs genotype groups with cigarettes per day (CPD) and pack-years in all smoking participants.

| Genotype group | rs667282 TT/TC | CC | rs3743073 TT | GG/TG | ADH1B ADH1B | OR (95% CI) | P  |
|----------------|---------------|----|--------------|-------|--------------|-------------|----|
| N               | 666           | 23.1 (22.3–24.0) | 0.135 | 36.3 (34.7–47.8) | 0.233 |
| rs667282 CC     | 170           | 21.5 (19.9–23.2) |       | 34.6 (31.6–37.6) |       |
| rs3743073 GG/TG | 609           | 23.1 (22.2–24.0) | 0.304 | 36.0 (34.4–37.5) | 0.430 |
| rs3743073 TT    | 227           | 22.0 (20.6–23.5) |       | 35.7 (32.9–38.4) |       |

1 P values were calculated by the Kruskal-Wallis test and were two-sided. doi:10.1371/journal.pone.0067664.t005
In people of European descent, variants in the CHRNA5-A3-B4 cluster showed a strong association with smoking consumption. Recently, in a functional magnetic resonance study of smokers, a genetic variation in CHRNA5 was observed to affect reactivity to images of smoking in brain areas related to memory and habitual behavior of the hippocampus and dorsal striatum [59]. However, the relationship between CHRNA5-A3-B4 and smoking consumption was not significant in this study, which was consistent with the study by Niu et al. regarding strong lung cancer-related variants in CHRNA5-A3-B4 and smoking behaviors in Chinese people [13]. Wu et al. reported nominally significant associations between rs667282 and CPD in a study of 3,725 smokers in a Chinese population (P = 0.022) [14]. In a study using an even larger sample (8,842 smokers) in Korea, the association between variants in the CHRNA5-A3-B4 and smoking quantity was also weak (P = 0.008–0.028) [40]. It seems that this smoking association is much weaker in an Asian population. Reasons about the racial difference in the gene cluster with smoking associations are unclear. Differences in genetic and environmental backgrounds may contribute to this discrepancy.

Several limitations of this study should be mentioned. First, some data on cigarette consumption were retrospective, so there may be some recall bias. Moreover, because of the limited sample number, the effect of the variants on ESCC risk in the stratified analysis did not reach statistical significance. A large population-based study is still needed to confirm the conclusions in this study.

In summary, to the best of our knowledge, this study was the first to demonstrate that the CHRNA5-A3-B4 gene cluster was associated with increased ESCC risk. This finding reveals another case study of a gene-environment interaction, highlighting the role of carcinogen-related genes in the pathology of ESCC. Further studies are required to confirm this finding in different ethnic populations.

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Author Contributions

Conceived and designed the experiments: YW QL LH. Performed the experiments: YW HW CW YL SW. Analyzed the data: YW HW WF. Contributed reagents/materials/analysis tools: QL LF HW LH. Wrote the paper: YW LH.

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