

Interactions between single nucleotide polymorphism of *SERPINA1* gene and smoking in association with COPD: a case–control study

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Background: *SERPINA1* gene has been implicated in the pathogenesis of chronic obstructive pulmonary disease (COPD), while smoking is a known risk factor for COPD. Little is known on the effect of *SERPINA1* gene and its interaction with smoking in the Chinese population. In this study, the effect of *SERPINA1* gene polymorphisms on COPD risk and its interaction with smoking status has been investigated.

Method: A total of 120 COPD patients and 481 healthy controls were recruited at The Armed Police Corps Hospital. Data on demographic variables, smoking status, history of occupational dust exposure, and allergies were collected. Genotyping for single nucleotide polymorphism's (SNP) rs1243160, rs2854254, and rs8004738 was performed in all participants.

Results: SNP rs8004738 genotype was associated with a significantly higher risk for COPD (odds ratio (OR) = 1.835, 95% confidence interval (CI): 1.002–3.360), whereas SNPs rs1243160 and rs2854254 did not exhibit such an association. Smoking habit also significantly increased the risk for COPD (OR = 2.306, 95% CI: 1.537–3.459). On stepwise logistic regression analysis, advanced age, smoking, and SNP rs8004738 variant were associated with increased risk for COPD, while female gender and higher educational status decreased the risk. On additive interaction analysis, a significant interactive effect of SNP rs8004738 and smoking was observed in this population (relative excess risk due to interaction = 0.478; attributable proportion due to interaction (AP) = 0.123; S = 1.197).

Conclusion: SNP rs8004738 of *SERPINA1* gene significantly interacted with smoking status and was associated with a higher risk for COPD in the Chinese population.

Keywords: chronic obstructive pulmonary disease (COPD), single nucleotide polymorphism (SNP), smoking, *SERPINA1* polymorphism

Introduction

Chronic obstructive pulmonary disease (COPD) is characterized by a progressive decrease in pulmonary ventilation due to obstruction of small airways. Multiple genetic and environmental factors are implicated in the causation of COPD. Genome-wide association studies have implicated genetic polymorphisms including that of *SERPINA1* gene in its pathogenesis. *SERPINA1* gene, located on chromosome 14q32.1, encodes for a serine protease inhibitor (alpha 1 antitrypsin, AAT), which targets elastase, plasmin, thrombin, trypsin, chymotrypsin, and plasminogen activator, and has been shown to play a role in the development of COPD.¹ A null mutation of *SERPINA1* gene was shown to significantly increase the risk of severe emphysema.² Single nucleotide polymorphisms (SNPs) of *SERPINA1* have also linked to a heightened susceptibility to COPD.^{3,4} However, majority of these studies were performed in Caucasians, definitive

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evidence of such an association is yet to be demonstrated in Asian population.^{5,6}

An earlier study that investigated potential associations of different SNPs found that only rs8004738 allele frequency was significantly different between COPD and control after adjusting for confounders.⁷ Smoking is a leading environmental risk factor for COPD with 2–4 times higher risk in smokers compared to that in nonsmokers.^{8,9} As there is a high prevalence of smokers in China,¹⁰ smoking would continue to be a predominant risk factor for COPD. Of note, a synergistic effect between genetic polymorphisms and environmental factors has also been implicated in the pathogenesis of COPD.^{11,12} Therefore, this study aimed to investigate rs8004738, which only has not been studied in Chinese population and other two nearby SNPs, rs1243160 and rs2854254, were also selected to assess their potential interactions with smoking in the association with the risk of COPD.

Materials and methods

Study participants

Patients diagnosed with COPD and admitted to The Armed Police Corps Hospital between September 2014 and September 2015 were eligible for participation. Patients who fulfilled the diagnostic criteria for COPD laid down in the GOLD (Global initiative for chronic Obstructive Lung Disease)

Guidelines 2015 were included.¹³ The criteria included 1) cough, expectoration, and/or wheeze for >3 months in a year; 2) $FEV_1/FVC < 70\%$ and $FEV_1 < 80\%$ on lung function tests after administration of bronchodilator drug.

The exclusion criteria included 1) patients unable to perform lung function tests; 2) history of bronchial asthma, interstitial lung disease, bronchiectasis, pulmonary embolism, lung cancer, active tuberculosis, or any type of chest surgery or mental disease which would affect lung function. A total of 676 patients met the inclusion criteria, whereas 556 patients met the exclusion criteria. A total of 120 patients were included in the case group for final analysis.

Healthy volunteers who underwent routine health checkup at The Armed Police Corps Hospital between September 2014 and September 2015 were recruited as controls (n=19,457). Exclusion criteria for controls was individuals suffering from any organic disease based on clinical examination, chest X-ray, laboratory investigations (complete blood counts, biochemical tests, urine or feces test, and those who were unable to undergo lung function test). Therefore, a total of 481 healthy controls were included in the final analysis (Figure 1). All participants were Han Chinese and were unrelated to each other. This study was approved by the Ethic Committee of General Hospital of Chinese People's Armed Police Forces, People's Republic

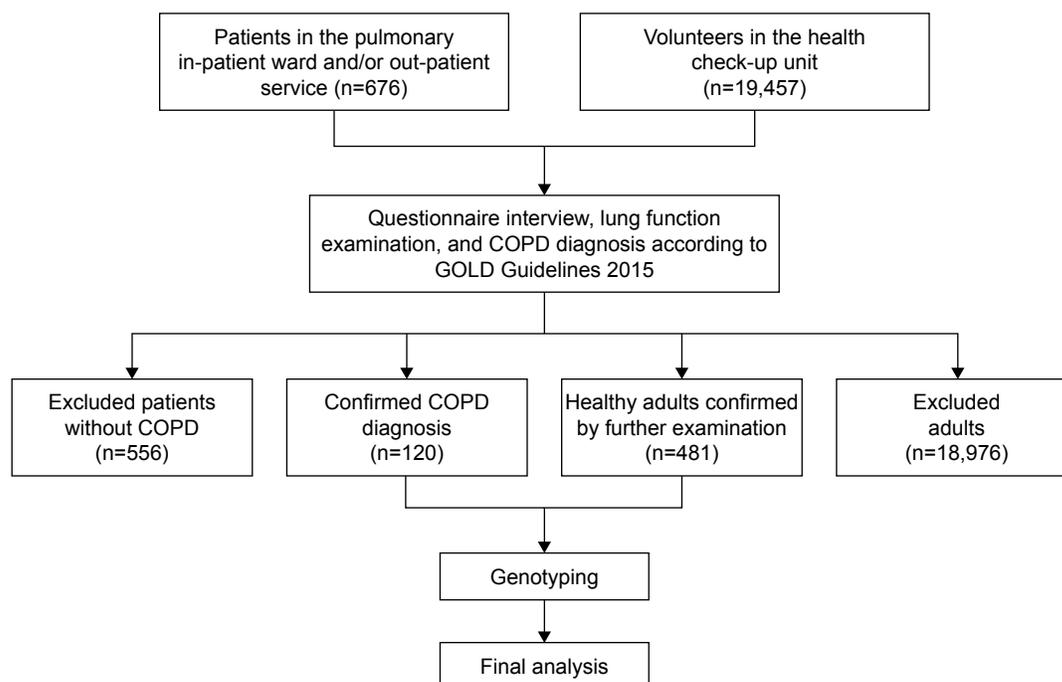


Figure 1 The flow chart of population selection.

Notes: The subjects were selected using questionnaire interview, lung function examination and COPD diagnosis according to GOLD Guidelines 2015. A total of 120 patients confirmed COPD diagnosis and 481 healthy adults confirmed by further examinations were screened for genotyping.

Abbreviation: GOLD, Global initiative for chronic Obstructive Lung Disease.

of China. Each participant signed a consent form before enrolment in this study.

Collection of demographic data

A standard questionnaire was administered to participants by trained personnel for data collection on following demographic variables: age, gender, body mass index (BMI), smoking status, education level, type of utensils and fuel used for cooking, occupational exposure to dust, and history of allergies.

Smoking index

Smoking status of participants was collected during their clinical visits. Smokers were defined as use of at least one cigarette a day, continuously for more than 1 year. Smoking index was calculated as follows:

$$\text{Smoking index} = \text{number of cigarettes/day} \times \text{years.}$$

DNA extraction, polymerase chain reaction, and genotyping

Approximately 2 mL of fasting blood was collected from each participant and stored at -80°C . DNA was extracted using DNA extraction kit (Tiangen Biotech Co. Ltd, Beijing, China). Primer pairs for each SNP were shown in Table 1. Primers were designed by Beijing Microread Genetics Co. Ltd (Beijing, China) (Table 1). Genotype was determined using SNaPshot assay (SNaPshot™ Multiplex Kit; Ukzybiotech. Ltd, Beijing, China) and sequencing was performed by Beijing Double Gene Technology Co., Ltd (Beijing, China). Polymerase chain reaction buffer system included 4.0 μL 2.5 \times buffer, 0.2 μL polymerase chain reaction primer (10 μM , F + R), 0.1 μL Taq DNA polymerase, 1.0 μL DNA templates (50–100 ng/ μL), and 4.7 μL double distilled water. Genotyping data were analyzed using software GeneMapper v3.2 (Applied Biosystems Inc., Foster City, CA, USA).

Statistical analyses

Data analyses were performed using SPSS software, ver 17.0 (SPSS Inc., Chicago, IL, USA). Quantitative data are

presented as mean \pm standard deviation. Independent *t*-test was used to compare the differences between age. Between-group differences with respect to baseline demographic variables and SNP genotypes were assessed by chi-squared test (χ^2). Further, Hardy–Weinberg genetic equilibrium test was performed. Stepwise logistic regression analysis was performed to identify independent risk factors. Odds ratios (ORs) and confidence intervals (CIs) were presented by multivariable logistic regression analysis. Additive interactions between SNP genotypes and smoking habit in association with COPD was accessed using following equations:

$$\text{Relative excess risk due to interaction (RERI)} = OR_{AB} - OR_{A\bar{B}} - OR_{\bar{A}B} + 1;$$

$$\text{Attributable proportion due to interaction (AP)} = \text{RERI}/OR_{AB};$$

$$\text{Synergy index (S)} = (OR_{AB} - 1)/[(OR_{A\bar{B}} - 1) + (OR_{\bar{A}B} - 1)].$$

A value of $P < 0.05$ was considered statistically significant.

Results

Baseline characteristics of study participants

COPD group consisted of 92 males and 28 females (mean age: 65.3 ± 11.3 years); the control group consisted of 322 males and 159 females (mean age: 49.3 ± 10.4 years) (Table 2). Statistically significant between-group differences were observed with respect to age and gender distribution ($P < 0.001$ and $P = 0.040$, respectively). Smoking status, BMI, and educational level were also significantly different between the two groups (higher rate for smokers, higher BMI, and lower educational level in the COPD group; $P < 0.001$ for all). No significant difference was observed with respect to type of utensils and fuel used for cooking, and for occupational exposure to dust, or history of allergy (Table 2).

Risk factors for COPD

On stepwise logistic regression analysis, advanced age (OR: 1.121, 95% CI: 1.088–1.155, $P < 0.001$), current smoking status (OR: 1.011, 95% CI: 1.006–1.016, $P < 0.001$), and mutation at rs8004738 of *SERPINA1* (OR: 3.948, 95% CI: 1.688–9.232, $P = 0.002$) was significantly associated with a higher risk for COPD (Table 3). Female gender (OR: 0.348, 95% CI: 0.154–0.787, $P = 0.011$) and higher educational status (OR: 0.323, 95% CI: 0.186–0.561, $P < 0.001$) showed a significant negative association with COPD (Table 3).

Table 1 SNPs and primers design

SNPs	Primers	
rs8004738	Forward	5'-GCAGCAACCCTCAGAGTCCT-3'
	Reverse	5'-ACTGGGCTGTGGTTGAGGTC-3'
rs1243160	Forward	5'-TGACCTGGGACAGGAATCG-3'
	Reverse	5'-GCACCTCAGCAGGCGGATA-3'
rs2854254	Forward	5'-TGACCTGGGACAGGAATCG-3'
	Reverse	5'-GCACCTCAGCAGGCGGATA-3'

Abbreviation: SNPs, single nucleotide polymorphisms.

Table 2 Baseline characteristics of subjects by study group

Characteristics	Category	COPD group	Control group	P-value
Age (years)		65.3±11.3	49.3±10.4	<0.001
Gender	Male	92	322	0.040
	Female	28	159	
Smokers	Yes	65	163	<0.001
	No	55	318	
BMI (kg/m ²)	<24	79	220	<0.001
	24–28	40	189	
	>28	1	72	
Educational status	Less than secondary	27	47	<0.001
	Senior high to college	51	184	
	University or higher	42	250	
Material for cooking	Liquefied gas or natural gas	89	389	0.103
	Firewood or grass	31	92	
Material for heating	Heating installation or electricity	87	387	0.056
	Coal	33	94	
History of occupational exposure to dust	Yes	15	46	0.341
	No	105	435	
History of allergy	Yes	14	92	0.055
	No	106	389	

Association between smoking index and COPD

Compared to nonsmokers, smokers had a significantly higher risk of COPD after adjusting for age, gender, educational levels, and SNP variations (OR: 2.306, 95% CI: 1.537–3.459, $P < 0.001$) (Table 4). Subgroup analysis by smoking index showed a dose response effect of smoking on COPD. Participants who had a smoking index >800 had four times higher risk for COPD compared to people who barely smoked (OR: 4.336, 95% CI: 2.094–8.981, $P < 0.001$).

SNP genotypes

Genotype frequency of SNP rs8004738, rs2854254, and rs1243160 fulfilled the Hardy–Weinberg genetic equilibrium

Table 3 Results of stepwise logistic regression analyses showing risk factors for COPD

	β	SE	Wald	P-value	OR	95% CI
Age	0.114	0.015	56.441	<0.001	1.121	1.088–1.155
Gender (female vs male)	-1.055	0.416	6.426	0.011	0.348	0.154–0.787
Smoking status	0.101	0.003	18.260	<0.001	1.011	1.006–1.016
Educational level (high vs low)	-1.130	0.281	16.128	<0.001	0.323	0.186–0.561
rs8004738 (non-GG vs GG)	1.373	0.433	10.040	0.002	3.948	1.688–9.232

Notes: Variables selected by stepwise logistic regression. SNP rs8004738 genotypes were divided as GG and non-GG (G/A and A/A).

Abbreviations: SE, standard error; OR, odds ratio; CI, confidence interval.

test (all $P > 0.05$; data not shown). There was no difference between allele frequencies for SNP rs2854254 and rs1243160. However, there was a significant difference in frequency of rs8004738 genotype between cases and controls; a higher frequency of A allele was observed in the COPD group ($P = 0.042$) (Table 5). This indicates that mutations of A allele were associated with a higher risk of COPD (OR = 1.373, 95% CI = 1.028–1.832, $P = 0.031$).

Interaction between smoking status and rs8004738 genotype

In nonsmoking participants, carriers of either G/A or A/A had a higher risk for COPD; the difference was statistically significant (OR: 1.835, 95% CI: 1.002–3.360). Among GG carriers, smokers showed a modestly higher risk for COPD compared to nonsmokers (OR: 2.582, 95% CI: 1.303–5.117). Among smokers, with non-GG genotype, the risk of COPD was significantly higher than that of nonsmokers with GG genotype (OR: 3.895, 95% CI: 2.120–7.157), which was indicative of a significant interactive effect (RERI = 0.478; AP = 0.123; S = 1.197) (Table 6).

Discussion

In this study, we selected three SNPs, rs8004738, rs1243160, and rs2854254. rs8004738 was selected based on Chappel et al where they found it was the only SNP with a significantly different allele frequency between COPD and controls.⁷ In this study, due to the limited budget, we could

Table 4 Association between smoking index and COPD risk

Smoking index (number × year)	Case group (n, %) n=120	Control group (n, %) n=481	OR	95% CI	P-value
Nonsmoker	55	318	1.000		
Smoker	65	163	2.306	1.537–3.459	<0.001
<1	55 (45.8)	318 (82.5)	1.000		
1–400	37 (30.8)	107 (15.8)	1.999	1.249–3.201	0.004
400–800	13 (10.8)	36 (0.9)	2.088	1.041–4.187	0.035
>800	15 (12.5)	20 (0.9)	4.336	2.094–8.981	<0.001

Notes: Data presented as frequency (proportion), ORs and 95% CI by logistic regression. Confounding factors included age, gender, educational levels, and genetic variations.

Abbreviations: OR, odds ratio; CI, confidence interval.

only assess two more SNPs, therefore, we picked rs1243160 and rs2854254, which is closest to rs8004738 in genetic sequence. Our results showed that a G→A allele mutation of SNP rs8004738 in *SERPINA1* gene was associated with a higher risk for COPD. More importantly, this mutation may also interact with an individual's smoking habit. The results suggest that both risk factors may contribute synergistically to COPD. Other two SNPs, rs1243160 and rs2854254, were not associated with COPD. To our knowledge, this is the first study to investigate the interrelation between genetic susceptibility and a modifiable life style factor associated with COPD in Chinese population.

The most important reported mutation of gene *SERPINA1* was the null mutation, which is known to cause a severe deficiency of AAT² and enhances the susceptibility of the lungs to potential damage caused by proteases. The cross-sectional studies involving the heterozygotes for of this mutation (PiMZ) also suggested a two times higher risk for COPD.³ In this study, we did not examine the Pi genotype or the level of AAT, which is also our limitation. A previous study from Hong Kong demonstrated a very low prevalence

of PiMZ mutation in Chinese population,¹⁴ indicating that this mutation has limited effect on the development of COPD in Chinese. Besides this mutation, other polymorphisms of *SERPINA1* gene were also reported to be associated with emphysema or COPD.^{15,16} Gene polymorphism associated with COPD or poor lung function does not infer association with altered serum AAT level. Thun et al did not find association of rs8004738 with serum AAT level conditioned for M allele mutation.¹⁷ Therefore, whether SNP found in our study is an independent factor associated with serum AAT levels needs to be determined. Based on our results, SNP rs8004738 was associated with higher risk for COPD, which is consistent with previous findings in Caucasian population.⁷ However, a previous study in a Japanese population demonstrated a lack of association between rs8004738 homozygotes and risk for emphysema.¹⁸ The disparity between their results and ours are probably due to the difference in disease category and ethnic background. Future studies with larger sample size and simultaneously tested serum AAT levels can further elucidate the role of rs8004738 in the development of COPD.

The potential interaction between smoking and genetic mutations in the causation of COPD has been reported. De Jong et al reported that in utero and adult exposure to tobacco modified the protective effect of glutathione S transferase omega gene polymorphism on lung function.¹² Furthermore, a study conducted on European population revealed a significant interaction between *SERPINA1* PiMZ genotype and occupational exposure to vapors, dust, gases, and fumes in association with decrease in lung function.¹¹ In this study, we also observed that the interaction between SNP rs8004738 and smoking contributed significantly to the risk of COPD in Chinese. The findings highlight the need for a ban on use of tobacco and concerted efforts at health education for management and prevention of COPD in China. Taken together, these data provided scientific evidence on the synergistic effects of genetic and environmental factors on the development of COPD.

Table 5 SNP genotypes frequencies in COPD case and control groups

SNP	Genotypes	Genotype frequency		P-value
		COPD	Control	
rs8004738	G/G	40	221	0.042
	G/A	59	187	
	A/A	21	73	
	G:A	139:101	629:333	
rs2854254	G/G	94	381	0.930
	G/A	22	85	
	A/A	4	15	
	G:A	210:30	847:115	
rs1243160	C/C	116	464	0.774
	C/T	4	15	
	T/T	0	2	
	C:T	236:4	237:3	

Abbreviation: SNP, single nucleotide polymorphism.

Table 6 Interactions between smoking and rs8004738 genotypes in association with chronic obstructive pulmonary disease

Risk factors		Case	Control	OR	95% CI	P-value	RERI	AP	S
Smoking	Genotype								
No	GG	18	150	1.000			0.478	0.123	1.197
Yes	GG	22	71	2.582	1.303–5.117	0.005			
No	Non-GG	37	168	1.835	1.002–3.360	0.047			
Yes	Non-GG	43	92	3.895	2.120–7.157	<0.001			

Note: SNP rs8004738 genotypes were divided as GG and non-GG (G/A and A/A).

Abbreviations: RERI, relative excess risk due to interaction; AP, attributable proportion due to interaction; S, synergy index; OR, odds ratio.

Our study also ruled out the potential association of SNP rs1243160 and rs2854254 with the risk of COPD. The association of either rs1243160 or rs2854254 with COPD has not been reported earlier. It remained possible that the lack of association of rs2854254 and rs1243160 was due to our limited sample size. Therefore, further study with large sample size is required to further elucidate the role of these two SNPs.

The case–control study design and retrospective nature of the analysis are key study limitations. There is a lack of data on serum AAT levels and limited information of participant's with respect to exposure to passive smoking.

Conclusion

This study demonstrated that SNP rs8004738 in *SERPINA1* gene significantly interacted with smoking status, and that both in association lead to an increased risk for COPD in the Chinese population. These results increased our knowledge regarding the interaction between genetic and environmental factors contributing to COPD. It emphasizes the need for educating COPD patients on quitting tobacco during COPD management.

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Disclosure

The authors report no conflicts of interest in this work.

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