

# Genetic variants in *FAM13A* and *IREB2* are associated with the susceptibility to COPD in a Chinese rural population: a case-control study

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**Background:** Genome-wide association studies identified several genomic regions associated with the risk of chronic obstructive pulmonary disease (COPD), including the 4q22 and 15q25 regions. These regions contain the *FAM13A* and *IREB2* genes, which have been associated with COPD but data are lacking for Chinese patients. The objective of the study was to identify new genetic variants in the *FAM13A* and *IREB2* associated with COPD in Northwestern China.

**Methods:** This was a case-control study performed in the Ningxia Hui Autonomous Region between January 2014 and December 2016. Patients were grouped as COPD and controls based on  $FEV_1/FVC < 70\%$ . Seven tag single-nucleotide polymorphisms (SNPs) in the *FAM13A* and *IREB2* genes were genotyped using the Agena MassARRAY platform. Logistic regression was used to determine the association between SNPs and COPD risk.

**Results:** rs17014601 in *FAM13A* was significantly associated with COPD in the additive (odds ratio [OR]=1.36, 95% confidence interval [CI]: 1.11–1.67,  $P=0.003$ ), heterozygote (OR=1.76, 95% CI: 1.33–2.32,  $P=0.0001$ ), and dominant (OR=1.67, 95% CI: 1.28–2.18,  $P=0.0001$ ) models. Stratified analyses indicated that the risk was higher in never smokers. rs16969858 in *IREB2* was significantly associated with COPD but in the univariate analysis only, and the multivariate analysis did not show any association.

**Conclusion:** The results suggest that the new variant rs17014601 in the *FAM13A* gene was significantly associated with COPD risk in a Chinese rural population. Additional studies are required to confirm the role of this variant in COPD development and progression.

**Keywords:** *FAM13A*, *IREB2*, chronic obstructive pulmonary disease, single-nucleotide polymorphism

## Introduction

Chronic obstructive pulmonary disease (COPD) is a major cause of morbidity and mortality worldwide. It is characterized by persistent respiratory symptoms and airflow limitation.<sup>1</sup> In People's Republic of China, the prevalence of COPD in individuals  $\geq 40$  years of age is estimated at 8.2%<sup>2</sup> or varying from 5% to 13% in different provinces/cities.<sup>3</sup> Cigarette smoking is considered as the most important risk factor, but genetic characteristics play an important role in the susceptibility to COPD. Genome-wide association studies (GWAS) identified several genomic regions associated with higher COPD risk. Some GWAS loci are located in the *FAM13A* gene on chromosome 4q22 and in the 15q25 locus, which includes the *IREB2* gene.<sup>4,5</sup>

GWAS showed that variants in *FAM13A* (family with sequence similarity 13, member A) were associated with  $FEV_1/FVC$  and COPD.<sup>6–8</sup> *FAM13A* was initially

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considered as a signal transduction gene because of the RhoGAP functional domain in the exon region,<sup>9</sup> but it is now known to be associated with  $\beta$ -catenin signaling, which is typically activated during injury repair and tissue regeneration.<sup>10,11</sup> Hypoxia commonly accompanies COPD and enhances *FAM13A* expression.<sup>9</sup> In addition, Kim et al<sup>12</sup> showed that *FAM13A* SNPs associated with a higher risk of COPD were also associated with an increased *FAM13A* expression in the lungs, suggesting a possible causative association with pathological changes in the lung. Corvol et al<sup>13</sup> showed that the association of the *FAM13A* gene with pulmonary function parameters ( $FEV_1$ %predicted and  $FEV_1$ /FVC) was observed in different independent cohorts, suggesting that *FAM13A* is associated with a specific phenotype of COPD. Furthermore, Choo et al<sup>14</sup> demonstrated an association between the CTGA diplotype in *FAM13A* and the emphysema phenotype of COPD, and Jiang et al<sup>11</sup> provided the basis for the role of *FAM13A* in the development of emphysema. A recent study by Corvol et al<sup>15</sup> showed that *FAM13A* and airway epithelial-mesenchymal transition (EMT) are closely associated in cystic fibrosis. EMT is also thought to play an important role in airway remodeling in COPD.<sup>16</sup> Taken together, these results strongly suggest that *FAM13A* is involved in the etiology of lung diseases and COPD.

Iron is found in cigarette smoke<sup>17</sup> and has been shown to disrupt the lung homeostasis, making lung tissues more susceptible to damage from any cause.<sup>18</sup> *IREB2* is a gene that is translated into the iron regulatory protein 2 (IRP2), which plays a key role in iron homeostasis. *IREB2* is in strong linkage disequilibrium with nicotine receptor genes (*CHRNA3* and 5).<sup>5</sup> *IREB2* expression is increased in the lungs of patients with COPD.<sup>19</sup> IRP2 regulates cellular iron homeostasis and mitochondrial function.<sup>20,21</sup> It is reported that some *IREB2* variants may affect COPD in the presence of high levels of iron due to cigarette smoke exposure.<sup>18</sup> Therefore, there could be some association between *IREB2* and respiratory conditions such as COPD.

The association between single-nucleotide polymorphisms (SNPs) in *FAM13A* and *IREB2* and the risk of COPD is still unclear, although some GWAS loci have been reported.<sup>4,5</sup> We aimed to identify new genetic variants associated with COPD in People's Republic of China. The aim of the present case-control study was to examine the association between tag SNPs<sup>22</sup> in *FAM13A* and *IREB2* and COPD risk. In addition, this is the first study evaluating the effect of genetic factors on the pathogenesis of COPD in the Ningxia Hui Autonomous Region (Northwest China).

## Materials and methods

### Study design and population

The COPD screening and early intervention project in the Ningxia Hui Autonomous Region was funded by the Ningxia government and aimed to carry out a prospective investigation between January 2014 and December 2016 to acquire data on COPD in the Ningxia Hui Autonomous Region. The investigation was approved by the ethics committee of the General Hospital of Ningxia Medical University. All participants provided a written informed consent.

A total of 6,130 participants  $\geq 40$  years of age and from a single township volunteered to participate in the study. They were local farmers, and at least three generations of their families were Han.

The participants were grouped as COPD and controls. COPD was defined as post-bronchodilator  $FEV_1$ /FVC  $< 70\%$  and with chronic respiratory airway symptoms including dyspnea, chronic cough, sputum production, or wheezing. The exclusion criteria were as follows: 1) history of other respiratory diseases such as bronchial asthma, pulmonary tuberculosis, interstitial lung disease, or lung cancer or 2) unable to perform the lung function tests for any reason. The controls were with normal pulmonary function ( $FEV_1$ /FVC  $> 70\%$ ) and had no known medical illnesses or family disorders.

All volunteers underwent blood tests and completed questionnaires. The participants were interviewed by trained interviewers using standardized questionnaires about the risk factors of COPD. All pulmonary function measurements were performed using portable spirometers (MicroLab Spirometer, MD Spiro, Lewiston, ME, USA) and according to the guidelines of the American Thoracic Society.<sup>23</sup> Peripheral blood samples (2 mL) were collected in EDTA Vacutainer tubes for DNA extraction.

### SNP selection and genotyping

We selected seven tag SNPs in the *FAM13A* and *IREB2* genes using the Genome Variation Server database (<http://gvs.gs.washington.edu/GVS147/>) and the National Center for Biotechnology Information database (<http://www.ncbi.nlm.nih.gov/projects/SNP>) and based on the following criteria: tag SNPs in the CHB and Asian database selected by the Haploview 4.2 software (Broad Institute, Cambridge, MA, USA)<sup>24</sup> with Hardy-Weinberg equilibrium (HWE)  $P$ -value  $\geq 0.05$ , a minor allele frequency  $\geq 0.05$ , and  $r^2 \geq 0.8$ .

Peripheral blood leukocyte DNA was extracted using a DNA extraction kit (Promega, Madison, WI, USA). DNA



concentration was determined using a NanoDrop 1000 (Thermo Fisher Scientific, Waltham, MA, USA). Genotyping was done by CapitalBio Corporation (Beijing, People's Republic of China) using a MassARRAY platform (Agena Biosciences Inc., San Diego, CA, USA), according to the manufacturer's protocol. Multiplex reaction primers were designed using the MassARRAY Assay Design Tools on the Agena official website (<https://agenacx.com/online-tools/>) (Table 1). Mass determination was carried out using a MALDI-TOF mass spectrometer (Agena Biosciences Inc.), and the MassARRAY Type 4.0 software (Agena Biosciences Inc.) was used for data acquisition.

## Covariate assessment

Demographic characteristics and COPD risk factors (age, sex, height, weight, smoking status, age at first cigarette, number of cigarettes/day, cooking and heating with coal stove, family history of lung disease, and childhood history of respiratory disease) were obtained using a questionnaire designed for this specific population. All participants were personally interviewed by trained interviewers.

The smoking status was defined as follows:<sup>25</sup> subjects who had smoked >20 packs of cigarettes in a lifetime or 1 cigarette/d for a year were regarded as ever smokers (current or former); otherwise, they were classified as never smokers. Smokers who were still smoking at the time of the interview were considered as current smokers, and those who had quit (for at least 30 days before the interview) were former smokers.<sup>26</sup> Pack-years were calculated in smokers by multiplying the average number of cigarettes smoked per day by the number of years of smoking and by dividing by 20 cigarettes/pack. Participants were classified as follows: 0, 0–20 and >20 pack-years. Cooking and heating with coal stove meant that the participants were using coal as domestic fuel. Body mass index (BMI) was calculated by dividing the weight (in kilograms) by the squared height (in meters).

## Statistical analysis

Continuous data were presented as mean  $\pm$  standard deviation, and categorical data were presented as frequency or percentage. The differences in the distributions of demographic characteristics, selected variables, and genotypes between the two groups were analyzed with Student's *t*-test or the chi-square test, as appropriate. HWE for each SNP was tested using the chi-square test in the control group.

Unconditional logistic regression analyses without or with adjustment for the covariates (age, sex, BMI, smoking status, pack-years, coal consumption, pulmonary problems in childhood, and family history of pulmonary diseases) were used to estimate the odds ratio (OR) and 95% confidence interval (95% CI) for evaluating the effect of each SNP on COPD risk. Then, genetic variants were assessed using different genetic models (additive, dominant, and recessive). The subgroup analyses according to the abovementioned covariates were performed using stratified models.

All analyses were performed with R (<http://www.R-project.org>, The R Foundation) and Empowerstats (<http://www.empowerstats.com>, X&Y Solutions, Inc., Boston, MA, USA). Two-sided *P*-values <0.05 were considered statistically significant.

## Results

### Characteristics of the participants

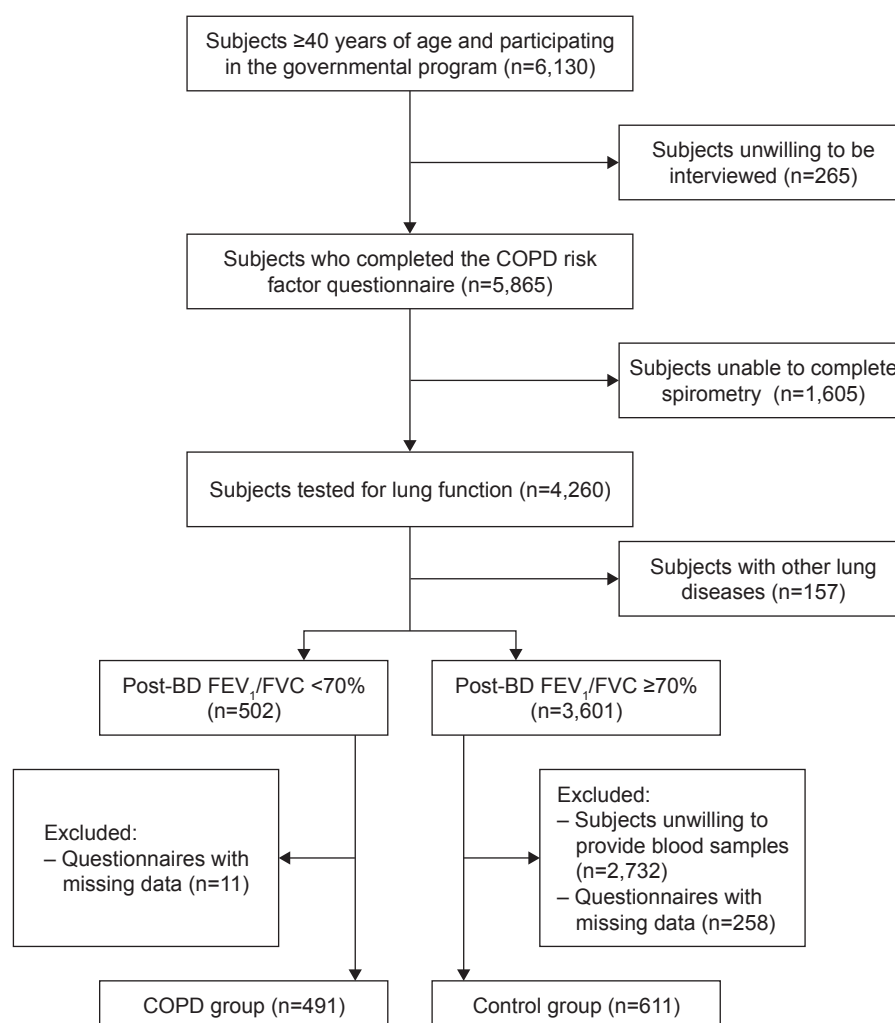
Figure 1 presents the study flowchart; 491 patients with COPD and 611 controls were included. Their characteristics are presented in Table 2. There were no differences in age, sex, coal use, and childhood pulmonary problems between the two groups (all *P*>0.05). Compared with the control group, the COPD group had significantly more smokers (*P*<0.0001), more pack-years smoked (*P*<0.0001), higher family history of pulmonary diseases (*P*=0.03), and lower BMI (*P*<0.0001). Among the patients with COPD, 25.9%, 45.8%, 21.6%, and 6.7% were classified as GOLD stages I, II,

**Table 1** SNPs and primers

SNP ID	First PCR primer	Second PCR primer
rs17014601	5'-ACGTTGGATGCTCAAAACATAAAGTGCAAC-3'	5'-ACGTTGGATGTACCTCCCCAGTTGGCAAG-3'
rs16996144	5'-ACGTTGGATGAAACCTTTGACTCTGGCCTC-3'	5'-ACGTTGGATGTCCTGCAGATCATAGAGGAC-3'
rs1870339	5'-ACGTTGGATGGTTAAGACCTATTCAACTTCC-3'	5'-ACGTTGGATGGAGTTATACTGTAAAACACAC-3'
rs2009746	5'-ACGTTGGATGGAGCACAGAAGTATAAAATC-3'	5'-ACGTTGGATGAAACGCTCCTGTGAAATAAC-3'
rs16969858	5'-ACGTTGGATGAGTGAAGAAGGATTATTG-3'	5'-ACGTTGGATGTCTGGGCCACCACCATTTC-3'
rs2656065	5'-ACGTTGGATGCTTAGTGTGTGTATTTCCC-3'	5'-ACGTTGGATGGATAACTGTAATCCTTTTTC-3'
rs3743079	5'-ACGTTGGATGCTGCCATGGTCCATCTTCAT-3'	5'-ACGTTGGATGCCTTAGTCTAACTGCAAGGG-3'

**Abbreviation:** SNPs, single-nucleotide polymorphisms.





**Figure 1** Study flowchart.

**Abbreviation:** COPD, chronic obstructive pulmonary disease.

III, and IV, respectively, according to airflow limitation ( $FEV_1\%$ pre levels).

## Association between SNPs and COPD susceptibility

The information about the selected SNPs is shown in Table 3. All of the SNPs were distributed within the parameters of HWE for the control population ( $P>0.05$ ). The call rates during the genotyping of these SNPs were all above 95%.

The frequencies of the SNPs in the COPD and control groups are shown in Table 4. The frequencies of rs17014601 T>C and rs16969858 C>T were significantly different between the two groups ( $P=0.0005$  and  $P=0.03$ , respectively). When considering the rs17014601 SNP in the *FAM13A* gene, compared with the TT genotype, the CT genotype was associated with an increased risk of COPD (non-adjusted OR=1.65, 95% CI=1.28–2.13). The association

was stable in the age- and sex-adjusted analysis (adjusted OR=1.66, 95% CI=1.29–2.15), and even after adjusting for more covariates (age, sex, BMI, smoking status, pack-years smoked, coal use, family history of lung disease, and childhood history of respiratory disease) (adjusted OR=1.76, 95% CI=1.33–2.32).

Five genetic models (additive, heterozygote, homozygote, dominant, and recessive) were used to analyze the associations between the variants and risk of COPD with and without adjustments for covariates (Table 5). The rs17014601 T>C SNP fitted the additive model, showing a significantly increased risk of COPD in the presence of a C allele, in a dose-dependent manner, and after adjustment for age, sex, and other covariates (adjusted OR=1.36, 95% CI=1.11–1.67,  $P=0.0028$ ). The rs17014601 CT genotype was associated with an increased risk of COPD based on the heterozygote model (adjusted OR=1.76; 95% CI=1.33–2.32;  $P=0.0001$ )



**Table 2** Characteristics of the subjects

Variables	Cases (n=491)	Controls (n=611)	P-value
Age (years) (mean $\pm$ SD)	69.0 $\pm$ 7.4	68.1 $\pm$ 7.5	0.0660
Male subjects, n (%)	347 (70.7)	410 (67.1)	0.2040
Body mass index (kg/m <sup>2</sup> ) (mean $\pm$ SD)	22.6 $\pm$ 3.5	23.5 $\pm$ 3.4	<0.0001
Smoking status at baseline, n (%)			<0.0001
Current	182 (37.1)	125 (20.6)	
Former	90 (18.3)	165 (27.2)	
Never	219 (44.6)	316 (52.2)	
Pack-years smoked (mean $\pm$ SD)	21.0 $\pm$ 26.7	11.7 $\pm$ 19.1	<0.0001
Coal consumption, n (%)	404 (82.3)	492 (80.5)	0.4570
Pulmonary problems in childhood, n (%)	38 (7.8)	51 (8.4)	0.7290
Family history of pulmonary diseases, n (%)	42 (8.6)	32 (5.3)	0.0280
Lung function (mean $\pm$ SD)			
FEV <sub>1</sub> (L)	1.57 $\pm$ 0.59	2.21 $\pm$ 0.56	<0.0001
FEV <sub>1</sub> , %predicted (%)	64.0 $\pm$ 21.6	85.7 $\pm$ 13.4	<0.0001
FVC (L)	2.57 $\pm$ 0.85	2.78 $\pm$ 0.72	<0.0001
FEV <sub>1</sub> /FVC (%)	60.7 $\pm$ 8.3	79.5 $\pm$ 5.4	<0.0001
GOLD stage			
1	127 (25.9)		
2	225 (45.8)		
3	106 (21.6)		
4	33 (6.7)		

and dominant model (adjusted OR=1.67; 95% CI=1.28–2.18;  $P=0.0001$ ).

No associations between the other SNPs and COPD were observed in the multivariate logistic regression analyses and genetic models.

## Stratification analysis

The association between variant genotypes and COPD risk was further evaluated using analyses stratified by age, sex, BMI, smoking status, pack-years smoked, coal use, family history of lung disease, and childhood history of respiratory disease (Table 6). The FAM13A rs17014601 CT genotype was associated with an increased COPD risk

in never smokers (adjusted OR=1.97, 95% CI=1.05–3.72,  $P=0.0356$ ) and 0 pack-years smoked (adjusted OR=2.06, 95% CI=1.44–2.96,  $P=0.0001$ ), when compared to the respective reference groups.

Stratified analysis of association between the rs16969858 SNP and COPD risk provided no statistically significant result (Table S1). There was no effect of the rs16969858 SNP genotype in different smoking status (Table S2). There was no effect of the rs16969858 SNP genotype in different smoking status according to the dominant and recessive models (Table S3). There was no effect of the rs17014601 SNP in different smoking statuses (Table S4), nor according to the dominant and recessive models (Table S5).

## Discussion

In the present population-based case-control study, we investigated the potential associations of FAM13A rs17014601 T>C, rs16996144 G>A, and rs1870339 C>G and IREB2 rs2009746 A>G, rs16969858 C>T, rs2656065 G>A, and rs3743079 C>T SNPs with COPD susceptibility in a Chinese rural population. We found that FAM13A rs17014601 had an independent effect on the COPD risk. The C allele in the FAM13A rs17014601 was significantly associated with COPD risk or occurrence in a Chinese rural population.

FAM13A on chromosome 4q22 has been consistently associated with COPD by GWAS. Among many tested SNPs, the rs7671167 SNP in FAM13A is most highly associated with COPD in Caucasians<sup>7,27,28</sup> and Asians, especially in Chinese.<sup>29</sup> Indeed, Xie et al<sup>29</sup> confirmed that the FAM13A rs7671167 SNP was associated with COPD risk in a Chinese Han population and that rs7671167 was related to lung function decline. In addition, Guo et al<sup>30</sup> reported that the frequency of the rs2869967 C allele was significantly increased in Chinese patients with COPD,<sup>30</sup> and Wang et al<sup>31</sup> showed that five FAM13A SNPs (rs7671167, rs2869966,

**Table 3** Characteristics of the SNPs in this study

SNP no	Gene	Loc	Category	Call rate (%)	MAF in the control group			GENO	P-value HWE
					Allele	1000G	Our		
rs17014601	FAM13A	4q22.1	Intron	97.1	C	0.1599	0.2698	49/229/328	0.1248
rs16996144	FAM13A	4q22.1	Intron	97.8	A	0.4948	0.4562	125/302/178	0.9738
rs1870339	FAM13A	4q22.1	Intron	98.2	G	0.3822	0.2882	55/241/313	0.1108
rs2009746	IREB2	15q25.1	Intron	97.4	G	0.2308	0.1203	10/125/469	0.0633
rs16969858	IREB2	15q25.1	Intron	98.1	T	0.0172	0.1018	6/112/491	0.9488
rs2656065	IREB2	15q25.1	Intron	97.7	A	0.3552	0.2273	30/216/360	0.9617
rs3743079	IREB2	15q25.1	3' UTR	98.0	T	0.1486	0.1310	12/135/460	0.3134

**Abbreviations:** SNPs, single-nucleotide polymorphisms; MAF, minor allele frequency; 1000G, 1000 genomes data; HWE, Hardy-Weinberg equilibrium; Loc, location; GENO, genotype; Our, data in our study.



**Table 4** Association between FAM13A and IREB2 SNPs and risk of COPD

Genotype	Genotype frequency		P-value <sup>a</sup>	Non-adjusted model OR (95% CI)	Adjusted OR (95% CI)	
	Cases n (%)	Controls n (%)			Model I <sup>b</sup>	Model II <sup>c</sup>
Total n	N=491	N=611				
rs17014601 T>C	n=479	n=596	0.0005			
TT	203 (42.4)	321 (53.9)		1.00 (ref)	1.00 (ref)	1.00 (ref)
CT	232 (48.4)	222 (37.3)		1.65 (1.28–2.13)	1.66 (1.29–2.15)	1.76 (1.33–2.32)
CC	44 (9.2)	53 (8.9)		1.31 (0.85–2.03)	1.33 (0.86–2.07)	1.34 (0.83–2.17)
rs16996144 G>A	n=480	n=594	0.4585			
GG	125 (26.0)	174 (29.3)		1.00 (ref)	1.00 (ref)	1.00 (ref)
AG	253 (52.7)	294 (49.5)		1.20 (0.90–1.59)	1.20 (0.90–1.59)	1.15 (0.84–1.57)
AA	102 (21.3)	126 (21.2)		1.13 (0.80–1.60)	1.12 (0.79–1.59)	1.22 (0.83–1.78)
rs1870339 C>G	n=484	n=597	0.3349			
CC	255 (52.7)	304 (50.9)		1.00 (ref)	1.00 (ref)	1.00 (ref)
GC	181 (37.4)	232 (38.9)		0.93 (0.72–1.20)	0.92 (0.71–1.19)	0.92 (0.70–1.22)
GG	48 (9.9)	61 (10.2)		0.94 (0.62–1.42)	0.93 (0.62–1.41)	1.01 (0.65–1.57)
rs2009746 A>G	n=485	n=603	0.3930			
AA	371 (76.5)	456 (75.6)		1.00 (ref)	1.00 (ref)	1.00 (ref)
GA	101 (20.8)	127 (21.1)		0.98 (0.73–1.31)	0.98 (0.73–1.32)	0.93 (0.67–1.28)
GG	13 (2.7)	20 (3.3)		0.80 (0.39–1.63)	0.79 (0.39–1.62)	0.96 (0.45–2.07)
rs16969858 C>T	n=483	n=596	0.0305			
CC	390 (80.8)	479 (80.4)		1.00 (ref)	1.00 (ref)	1.00 (ref)
TC	88 (18.2)	111 (18.6)		0.97 (0.71–1.33)	0.97 (0.71–1.32)	0.96 (0.69–1.35)
TT	5 (1.0)	6 (1.0)		1.02 (0.31–3.38)	0.99 (0.30–3.28)	1.17 (0.34–4.02)
rs2656065 G>A	n=470	n=590	0.5889			
GG	292 (62.1)	348 (59.0)		1.00 (ref)	1.00 (ref)	1.00 (ref)
AG	155 (33.0)	211 (35.8)		0.88 (0.68–1.13)	0.88 (0.68–1.14)	0.87 (0.66–1.15)
AA	23 (4.9)	31 (5.3)		0.88 (0.50–1.55)	0.86 (0.49–1.51)	0.88 (0.48–1.64)
rs3743079 C>T	n=466	n=581	0.5210			
CC	345 (74.0)	443 (76.3)		1.00 (ref)	1.00 (ref)	1.00 (ref)
TC	106 (22.8)	125 (21.5)		1.09 (0.81–1.46)	1.11 (0.82–1.49)	1.14 (0.83–1.57)
TT	15 (3.2)	13 (2.2)		1.48 (0.70–3.16)	1.50 (0.70–3.19)	1.20 (0.52–2.74)

**Notes:** <sup>a</sup>P-values of a two-sided  $\chi^2$  test,  $\alpha=0.05$ . <sup>b</sup>Adjusted for age and sex. <sup>c</sup>Adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; COPD, chronic obstructive pulmonary disease; OR, odds ratio; CI, confidence interval; BMI, body mass index.

rs2869967, rs2045517, and rs6830970) were associated with the FEV<sub>1</sub>/FVC ratio in all subjects and that rs6830970 was associated with the FEV<sub>1</sub>/FVC ratio in the COPD subset. Furthermore, the FAM13A locus is apparently not influenced by smoking.<sup>4</sup> van der Plaats et al<sup>32</sup> suggested that the FAM13A

loci (including rs6849143) are significantly associated with lung function measurements in never smokers.

In the present study, we found that a new FAM13A variant contributed to COPD risk in a Chinese rural population. Indeed, rs17014601 was significantly associated with

**Table 5** Association between rs17014601 and risk of COPD in different inheritance models

Model	Genotypes	Cases n (%)	Controls n (%)	Non-adjusted model		Adjustment I model		Adjustment II model	
				OR (95% CI)	P-value <sup>a</sup>	OR (95% CI)	P-value <sup>b</sup>	OR (95% CI)	P-value <sup>c</sup>
Additive				1.32 (1.10–1.59)	0.0032	1.33 (1.11–1.61)	0.0025	1.36 (1.11–1.67)	0.0028
	TT	203 (42.4)	321 (53.9)	1		1		1	
Heterozygote	CT	232 (48.4)	222 (37.3)	1.65 (1.28–2.13)	0.0001	1.66 (1.29–2.15)	0.0001	1.76 (1.33–2.32)	0.0001
Homozygote	CC	44 (9.2)	53 (8.9)	1.31 (0.85–2.03)	0.2219	1.33 (0.86–2.07)	0.1967	1.34 (0.83–2.17)	0.2245
Dominant	CT+CC	276 (57.6)	275 (46.1)	1.59 (1.25–2.02)	0.0002	1.60 (1.26–2.04)	0.0001	1.67 (1.28–2.18)	0.0001
Recessive	TT+CT	435 (90.8)	543 (91.1)	1		1		1	
	CC	44 (9.2)	53 (8.9)	1.04 (0.68–1.58)	0.8676	1.05 (0.69–1.60)	0.8230	1.03 (0.65–1.62)	0.9014

**Notes:** The Benjamini and Hochberg FDR method was used to adjust the multiple hypothesis tests, standard  $\alpha=0.05$ . <sup>a</sup>Unadjusted model. <sup>b</sup>Adjustment I model was adjusted for age and sex. <sup>c</sup>Adjustment II model was adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease.

**Abbreviations:** COPD, chronic obstructive pulmonary disease; OR, odds ratio; CI, confidence interval; FDR, false discovery rate; BMI, body mass index.



**Table 6** Stratified analysis of the association between the rs17014601 SNP and COPD risk

Factors	Cases			Controls			Adjusted <sup>a</sup> OR (95% CI)	P-value
	CC	CT	TT	CC	CT	TT		
Age (years)								
≤60	10	30	26	10	34	54	2.75 (0.79–9.51)	0.1105
>60	34	202	177	43	188	267	1.16 (0.68–1.97)	0.5951
Sex								
Male	28	259	148	38	149	215	1.19 (0.67–2.13)	0.5515
Female	16	73	55	15	73	106	2.24 (0.93–5.38)	0.0718
Smoking status								
Current	13	79	86	10	46	67	0.97 (0.31–3.02)	0.9533
Former	7	46	36	17	66	80	1.54 (0.50–4.69)	0.4506
Never	24	107	81	26	109	172	1.97 (1.05–3.72)	0.0356
Pack-years smoked								
≥20	15	94	80	14	51	71	1.50 (0.94–2.39)	0.0898
<20	5	31	40	13	62	78	0.89 (0.51–1.56)	0.6836
0	24	107	83	26	109	172	2.06 (1.44–2.96)	0.0001
BMI								
<18.5	6	23	22	3	10	21	3.34 (0.37–29.95)	0.2805
18.5–24.9	30	168	128	34	141	193	1.52 (0.85–2.73)	0.1576
≥25.0	8	48	53	16	71	107	0.75 (0.27–2.13)	0.5889
Coal consumption								
Yes	38	193	164	40	186	254	1.66 (0.98–2.81)	0.0616
No	6	39	39	13	36	67	0.96 (0.15–6.25)	0.9693
Pulmonary problems in childhood								
Yes	2	21	15	2	22	24	0.47 (0.05–4.83)	0.5254
No	42	209	187	48	200	297	1.39 (0.86–2.27)	0.1822
Family history of pulmonary diseases								
Yes	2	20	20	2	10	19	1.41 (0.12–17.12)	0.7854
No	41	211	183	51	212	302	1.36 (0.83–2.22)	0.2200

**Notes:** <sup>a</sup>Adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease. In each case, the model was not adjusted for the stratification variable.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; COPD, chronic obstructive pulmonary disease; OR, odds ratio; CI, confidence interval; BMI, body mass index.

an increased risk of COPD. In People's Republic of China, COPD is more common among rural residents compared with urban residents,<sup>2</sup> probably because of a number of environmental risk factors such as old age, smoking, coal use, infection, and low body mass index. Ningxia is an agricultural region in Northwestern China. Coal stove cooking and winter coal heating are very common. One advantage of our study is that we conducted an investigation of the possible COPD risk factors in individuals who were not yet treated for any lung disease. Then, we adjusted these associations according to known risk factors for COPD, including coal use. The association between rs17014601 and COPD remained significant after adjusting for the confounding factors. In addition, stratified analyses showed that never smokers had a higher risk of COPD in association with FAM13A rs17014601 compared with the whole cohort. This is the first report of the association between the FAM13A rs17014601 and COPD risk, but the results have to be confirmed by additional studies.

The exact biological functions of FAM13A are still unknown. The RhoGAP domain in the exon region may

be related to COPD.<sup>13</sup> Rho GTPases are key regulators of cytoskeletal dynamics involved in pulmonary endothelial barrier functions, and have been shown to be dysregulated in several lung diseases.<sup>33</sup> It is probable that genetic variations of FAM13A may affect Rho GTPases activity and the cellular pathways associated with FAM13A, hereby contributing to lung disease. The most significant SNPs in FAM13A have been found in non-coding regions downstream of the RhoGAP domain and are associated with *FAM13A* gene expression levels.<sup>7</sup> Recently, the biological function of FAM13A in emphysema development has been explored: Jiang et al<sup>11</sup> reported the expression of the FAM13A protein in airways, alveolar epithelial cells, and alveolar macrophages. FAM13A knockout mouse models are less susceptible to develop emphysema.<sup>11</sup> In vitro experiments showed that FAM13A interacts with PP2A and promotes the degradation of  $\beta$ -catenin in bronchial epithelial cells, inhibiting the activation of the Wnt pathway.<sup>10</sup> Moreover, metabolic regulation may be another mechanism by which FAM13A promotes CS-induced emphysema.<sup>34</sup>



Indeed, FAM13A promotes fatty acid oxidation (FAO) and subsequent increases in ROS, possibly by interacting with Sirtuin 1 (SIRT1) and increasing the expression of CPT1A, a key mitochondrial enzyme for the FAO pathway, thereby enhancing FAO. These findings suggest that the impact of FAM13A on COPD could be independent from smoke exposure.

A previous study in Caucasians showed that IREB2 polymorphisms had an effect of COPD susceptibility, independently from smoking.<sup>27</sup> A Russian study showed associations between the rs13180 SNP in IREB2 and COPD and lung function in the Tatar population.<sup>35</sup> A study from Poland suggests that the rs2568494 SNP in IREB2 was not associated with COPD, but with lung cancer.<sup>28</sup> In the present study, the rs16969858 SNP in the *IREB2* gene was significantly associated with COPD only in the univariate analysis, and multivariate analysis did not show any association. These discrepancies may be due to the selection of the SNPs being studied, as well as to the interactions with other environmental and genetic factors. Indeed, the rs16969858 SNP is an intronic SNP that could cause dysfunction because of differential splicing. A recent genome-wide bioinformatics study suggested that there were no differences in gene expression of *IREB2* and *FAM13A*, but that the expression of these genes was dependent on the interactions of genes.<sup>36</sup> This suggests new research avenues and expression studies in lung tissues should be performed.

There are some limitations in the present study that should be addressed. The number of COPD cases was limited. Larger scale studies are needed in different populations to validate the result. Although we were able to identify associations, we were not able to identify the causal mechanisms. This is the first study on the association between rs17014601 and COPD. The Han ethnicity was selected because it is the major ethnic group in China. Nevertheless, we agree that other ethnic groups will have to be studied. In addition, the Han populations that were enrolled in the COPD Susceptibility Study were mainly from southern China. In the present study, we included Chinese Han people from Northwest China and it is possible that there are genetic differences between South and North China.<sup>37</sup> Additional studies are still necessary to understand the genetics of COPD.

## Conclusion

The present study strongly suggests an association between the *FAM13A* gene and COPD in Chinese rural patients. Further studies are required to elucidate the functional roles of these variants, which may have an impact on the management of lung diseases.

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## Disclosure

The authors report no conflicts of interest in this work.

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## Supplementary materials

**Table S1** Stratified analysis of the association between the rs16969858 SNP and COPD risk

Factors	Cases			Controls			Adjusted <sup>a</sup> OR (95% CI)	P-value
	TT	TC	CC	TT	TC	CC		
Age (years)								
≤60	1	10	54	0	21	77	0.861 (0.350–2.119)	0.7446
>60	4	78	336	6	90	402	0.984 (0.686–1.413)	0.9315
Sex								
Male	5	66	269	3	77	322	1.051 (0.709–1.558)	0.8039
Female	0	22	121	3	34	157	0.905 (0.462–1.776)	0.7725
Smoking status								
Current	1	29	150	0	21	102	2.182 (0.969–4.914)	0.0597
Former	0	19	69	1	29	130	0.717 (0.339–1.515)	0.3828
Never	4	40	171	5	59	249	0.985 (0.619–1.568)	0.9497
Pack-years smoked								
≥20	1	35	153	1	27	107	1.299 (0.636–2.651)	0.4726
<20	1	12	64	0	25	127	0.882 (0.393–1.982)	0.7619
0	3	41	173	5	59	244	0.920 (0.583–1.453)	0.7206
BMI								
<18.5	1	4	44	1	7	27	0.12 (0.01–1.00)	0.0504
18.5–24.9	3	68	251	1	72	294	1.074 (0.725–1.592)	0.7216
≥25.0	1	16	95	4	32	157	0.916 (0.459–1.827)	0.8038
Coal consumption								
Yes	4	76	318	5	95	379	1.021 (0.709–1.469)	0.9128
No	1	12	72	1	16	100	0.771 (0.264–2.248)	0.6336
Pulmonary problems in childhood								
Yes	1	4	33	1	8	41	1.018 (0.241–4.295)	0.9803
No	4	83	355	5	103	438	1.032 (0.739–1.441)	0.8526
Family history of pulmonary diseases								
Yes	2	9	32	1	8	23	0.808 (0.214–3.056)	0.7537
No	3	79	356	5	103	455	1.026 (0.733–1.437)	0.8801

**Notes:** <sup>a</sup>Adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease. In each case, the model was not adjusted for the stratification variable.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; COPD, chronic obstructive pulmonary disease; OR, odds ratio; CI, confidence interval; BMI, body mass index.

**Table S2** Effect of the rs16969858 SNP genotype in different smoking statuses

Smoking status	Genotype	Case n=483	Control n=596	Crude OR (95% CI)	Adjustment I model OR (95% CI)	Adjustment II model OR (95% CI)
Never	CC	171	249	1	1	1
Never	TC	40	59	0.971 (0.622–1.518)	0.945 (0.603–1.482)	0.874 (0.554–1.379)
Never	TT	4	5	1.146 (0.303–4.331)	1.213 (0.319–4.620)	1.377 (0.356–5.322)
Former	CC	69	130	0.760 (0.535–1.08)	0.747 (0.515–1.083)	0.682 (0.466–0.998)
Former	TC	19	29	0.939 (0.510–1.729)	0.935 (0.498–1.753)	0.918 (0.485–1.735)
Former	TT	0	1	0.0 (0.000, Inf)	0.000 (0.000, Inf)	0.000 (0.000, Inf)
Current	CC	150	102	2.107 (1.532–2.897)	2.049 (1.478–2.841)	2.043 (1.456–0.866)
Current	TC	29	21	1.979 (1.092–3.586)	1.896 (1.034–3.476)	1.748 (0.938–3.255)
Current	TT	1	0	1116446.183 (0.000, Inf)	844979.151 (0.000, Inf)	751029.731 (0.000, Inf)
P-value for interaction				0.6694	0.6999	0.6244

**Notes:** Adjustment I model was adjusted for age and sex. Adjustment II model was adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease. Inf, the sample size is too small.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; OR, odds ratio; CI, confidence interval; BMI, body mass index.



**Table S3** Effect of the rs16969858 SNP according to different smoking statuses and the dominant and recessive models

Smoking status	Genotype	Case n=483	Control n=596	Crude OR (95% CI)	Adjustment I model OR (95% CI)	Adjustment II model OR (95% CI)
Never	CC	171	249	1	1	1
Never	TC+TT	44	64	0.985 (0.640–1.515)	0.965 (0.625–1.489)	0.907 (0.585–1.408)
Former	CC	69	130	0.760 (0.535–1.08)	0.746 (0.514–1.082)	0.681 (0.465–0.997)
Former	TC+TT	19	30	0.907 (0.495–1.665)	0.906 (0.485–1.693)	0.893 (0.474–1.683)
Current	CC	150	102	2.107 (1.532–2.897)	2.048 (1.477–2.840)	2.041 (1.455–2.864)
Current	TC+TT	30	21	2.047 (1.134–3.696)	1.952 (1.068–3.566)	1.799 (0.970–3.336)
P-value for interaction				0.8718	0.8258	0.6148

**Notes:** Adjustment I model was adjusted for age and sex. Adjustment II model was adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease. rs16969858 C>T: The sample size of TT is too small.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; OR, odds ratio; CI, confidence interval; BMI, body mass index.

**Table S4** Effect of the rs17014601 SNP in different smoking statuses

Smoking status	Genotype	Case n=479	Control n=596	Crude OR (95% CI)	Adjustment I model OR (95% CI)	Adjustment II model OR (95% CI)
Never	TT	81	172	1	1	1
Never	CT	107	109	2.084 (1.432–3.034)	2.094 (1.438–3.051)	1.948 (1.328–2.858)
Never	CC	24	26	1.960 (1.06–3.624)	1.987 (1.074–3.677)	1.896 (1.012–3.554)
Former	TT	36	80	0.956 (0.595–1.535)	0.906 (0.557–1.474)	0.804 (0.490–1.319)
Former	CT	46	66	1.480 (0.934–2.344)	1.398 (0.871–2.243)	1.305 (0.807–2.109)
Former	CC	7	17	0.874 (0.349–2.192)	0.827 (0.327–2.095)	0.753 (0.292–1.944)
Current	TT	86	67	2.726 (1.801–4.125)	2.626 (1.728–3.992)	2.512 (1.626–3.882)
Current	CT	79	46	3.647 (2.327–5.716)	3.533 (2.245–5.558)	3.445 (2.173–5.461)
Current	CC	13	10	2.760 (1.162–6.561)	2.690 (1.128–6.416)	2.638 (1.087–6.401)
P-value for interaction				0.4450	0.4385	0.6186

**Notes:** Adjustment I model was adjusted for age and sex. Adjustment II model was adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; OR, odds ratio; CI, confidence interval; BMI, body mass index.

**Table S5** Effect of the rs17014601 SNP in different smoking statuses according to the dominant and recessive models

Smoking status	Genotype	Case n=479	Control n=596	Crude OR (95% CI)	Adjustment I model OR (95% CI)	Adjustment II model OR (95% CI)
Never	TT	81	172	1	1	1
Never	CT+CC	131	135	2.061 (1.442–2.945)	2.073 (1.449–2.965)	1.937 (1.345–2.789)
Former	TT	36	80	0.956 (0.595–1.535)	0.908 (0.558–1.477)	0.805 (0.491–1.322)
Former	CT+CC	53	83	1.356 (0.878–2.093)	1.283 (0.819–2.011)	1.195 (0.757–1.887)
Current	TT	86	67	2.726 (1.801–4.125)	2.627 (1.728–3.993)	2.515 (1.628–3.887)
Current	CT+CC	92	56	3.489 (2.282–5.333)	3.385 (2.205–5.197)	3.306 (2.136–5.117)
P-value for interaction				0.2256	0.2231	0.4110

**Notes:** Adjustment I model was adjusted for age and sex. Adjustment II model was adjusted for age, sex, BMI, smoking status, pack-years smoked, coal consumption, family history of lung disease, and childhood history of respiratory disease.

**Abbreviations:** SNPs, single-nucleotide polymorphisms; OR, odds ratio; CI, confidence interval; BMI, body mass index.

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