The role of WWOX polymorphisms on COPD susceptibility and pulmonary function traits in Chinese: a case-control study and family-based analysis

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Single nucleotide polymorphisms (SNPs) in the WW domain containing oxidoreductase (WWOX) gene were recently identified to be quantitative trait loci for lung function and thus likely to be susceptible biomarkers for COPD. However, the associations between WWOX SNPs and COPD risk are still unclear. Here, by conducting a two-center case-control study including 1511 COPD cases and 1677 controls and a family-based analysis comprising 95 nuclear pedigrees, we tested the associations between five SNPs that are rs10220974C>T, rs3764340C>G, rs12918952G>A, rs383362G>T, rs12828G>A of WWOX and COPD risk as well as the hereditary inclination of these loci among COPD families. We found that the SNP rs383362G>T was significantly associated with an increased risk of COPD in a T allele-number dependent-manner (OR = 1.30, 95%CI = 1.11-1.52). The T allele was more prone to over transmit to sick children and sibs than the G allele (Z = 2.900, P = 0.004). Moreover, the forced expiratory volume in one second/forced vital capacity (FEV1/FVC), FEV1/predicted-FEV1 and annual FEV1 also significantly decreased in the rs383362T carriers compared to the rs383362GG carriers. For other SNPs, no significant association was observed for COPD and pulmonary function. Taken together, our data demonstrated that the SNP rs383362G>T of WWOX plays a role in COPD inheritance.

Chronic obstructive pulmonary disease (COPD) is one of the most strikingly increasing lung diseases characterized by incompletely reversible airflow obstruction. During 1990–2010, COPD ranked the forth and the third causes of death in American and in Chinese, respectively. The values are still ongoing now and no doubt that COPD will seriously influence the life quality of and life health of patients and cause high economic burden on patients as well as the society.

The pathogenesis of COPD is complex involving both environmental and genetic factors. Environmental factors such as tobacco smoking and air pollution can cause a series of sophisticated biological reactions like oxidative stress to induce COPD development, while genetic variants can regulate the expression or function of such molecules participating in above reactions and thus determinate COPD susceptibility. Tobacco smoking can bring a large amount of oxygen free radical into lung and trigger oxidative stress, which directly damages the lung tissue to participate in the pathological courses of COPD. In response to the damage, the organism activates the antioxidant system to resist the harm of oxygen free radical such as up-expression of a series of antioxidant enzymes in airway epithelial cells. Due to different genetic background, different smokers show significantly different expression or function of these enzymes. In this condition, genetic variants located in these antioxidant enzymes might alter individuals' susceptibility to develop COPD.
Table 1. Genotype distributions of WWOX SNPs and their associations with COPD risk in the southern Chinese population. Abbreviation: MAF, minor allele frequency; OR_rhom, heterozygote versus wild-genotype homozygote; OR_homo, Variant homozygote versus wild-type homozygote; OR heter, OR_homo, OR_rhom, OR_homo, calculated by the additive, dominant and recessive model, respectively; 3′-UTR, 3′-untranslated region. *Wild-type homozygote/heterozygote/variant homozygote. The observed genotype frequencies among the controls all matched the Hardy-Weinberg equilibrium in the control subjects (P > 0.05 for all). P values of a two-sided χ² test for genotype distribution between the cases and controls. 4M AF of the variant allele. Data were calculated by the unconditional logistic regression, adjusted for age, sex, pack-years smoked, biomass as fuels and sample source region.

| SNP       | Location in gene | Case* (n = 1025) | Control* (n = 1061) | P   | MAF*   | OR_rhom (95%CI) | OR_homo (95%CI) | OR heter (95%CI) | OR hom (95%CI) | OR rec (95%CI) |
|-----------|------------------|------------------|---------------------|-----|-------|---------------|----------------|----------------|--------------|-------------|
| rs10220974C > T | Promoter                | 748/252/25      | 806/231/24         | 0.290 | 0.147 | 0.132         | 1.20 (0.97–1.48) | 1.02 (0.57–1.83) | 1.13 (0.95–1.35) | 1.18 (0.96–1.45) |
| rs3764340C > G | Exon                    | 832/186/7       | 837/214/10         | 0.390 | 0.098 | 0.110         | 0.71 (0.26–1.94) | 0.91 (0.74–1.12) | 0.73 (0.27–1.97) |
| rs12918952G > A | Exon                    | 884/137/4       | 910/144/7          | 0.688 | 0.071 | 0.075         | 0.95 (0.78–1.13) | 0.98 (0.77–1.29) | 1.00 (0.77–1.29) | 0.66 (0.19–2.28) |
| rs383362G > T | 3′-UTR                  | 755/259/11      | 834/220/7          | 0.025 | 0.137 | 0.110         | 0.42 (0.20–0.90) | 2.23 (0.84–5.93) | 1.28 (1.06–1.56) | 1.28 (1.04–1.57) |
| rs12828G > A | 3′-UTR                  | 474/463/88      | 473/489/99         | 0.692 | 0.312 | 0.324         | 0.94 (0.78–1.13) | 0.96 (0.84–1.10) | 0.94 (0.79–1.12) | 0.98 (0.72–1.34) |
Table 3 shows results from stratification analysis of the association between the SNP rs383362G > T and COPD risk, as well as the interaction between the SNP and variables on COPD risk. The rs383362G > T significantly interacted with biomass as fuels on increasing COPD risk ($P = 0.021$), because there was an intuitionistic
FVC (GG, TG, TT: 2.714 ± 0.728) to sick children and sibs under the additive genetic model (Z single-marker analysis indicated that the T risk allele of rs383362G further exerted significantly excessive transmission from parents to affected offspring under the additive genetic model (Z = 2.550, P = 0.011), so other haplotypes did not (P > 0.05 for all). In addition, the logistic regression model showed that the rs383362T allele and T-C haplotype conferred significantly increased risk of COPD compared to the rs383362G allele (OR = 2.04, 95%CI = 1.41–2.95) and G-C haplotype (OR = 1.64, 95%CI = 1.11–2.41), respectively.

### Possible function of the rs383362G > T polymorphism by bioinformatics analysis

The SNP rs383362G > T is located in 3′- untranslated region (3′-UTR) of WWOX that may affect binding ability of potentially microRNA or RNA-binding protein and thus regulates WWOX expression. By querying the SNPexp database (http://app3.titan.uio.no/biotools/tool.php?app=snpexp) with regard to the CHB population (i.e., a Chinese population), the mean WWOX expression was consistently lower in 8 cases of lymphoblastoid cells with the rs383362TG genotype than that in 37 cases with the rs383362GG genotype using different expression microarray platforms. However, the difference was not significant, which may be due to the limited sample size. Also, by polling the SNPinfo website (http://snpinfo.niehs.nih.gov/snpfunc.html), the transversion of G to T may influence four microRNAs' binding ability to WWOX 3′-UTR. Interestingly, two microRNAs among the four microRNAs,
hsa-miR-134 and hsa-miR-758 were experimentally proved to be translational regulators of WWOX (http://www.genecards.org/cgi-bin/carddisp.pl?gene=WWOX).

**Discussion**

In the current study, based on a two-stage case-control and a family-based study, we found that the T genotypes of WWOX SNP rs383362G>T were significantly associated with an increased risk of COPD and decreased pulmonary function traits in Chinese. The T allele exerted significantly excessive transmission from parents to sick offspring. The genotypes also interacted with biomass as fuels on increasing COPD risk. Bioinformatics analysis further showed that the SNP has a possible regulation on WWOX expression.

COPD is a well-established disease involving both genetic and environmental factors. Multiple association studies especially the GWASs, have reported an abundant of SNPs to be susceptible loci for COPD or pulmonary function. These SNPs majorly belong to genes that involve the major pathogenic mechanisms of COPD, such as inflammation\(^\text{22,23}\), oxidative stress\(^\text{15,24}\), DNA damage\(^\text{25,26}\) and epithelial-mesenchymal transition (EMT)\(^\text{26,27}\). However, most of these SNPs are considered to be lack of function and non-causal variant with regard to their genomic location in gene introns or intergenic region. An effective strategy to discover causal variants was fine-mapping based on the results from GWASs. Fine mapping majorly analyzes those putatively function variants considering their location in cis-acting element of genes that harbor susceptible loci of disease. Based on this strategy, we found a 3’-UTR SNP rs383362G>T of WWOX, a gene that has been reported to harbor susceptible loci of pulmonary function by GWAS\(^\text{15,16}\), was associated with COPD risk and pulmonary function traits. The rs383362T risk allele was more prone to over transmitted to sick children and sibs. This SNP got Bioinformatics data support to have potentiality on affecting WWOX expression. Taken together, all these evidences suggested the rs383362G>T polymorphism to be a susceptible loci for COPD in Chinese.

Low expression of WWOX has been reported to be an inducement of human diseases such as lung cancer\(^\text{28-30}\). The SNP rs383362G>T is located in the 3’-UTR of WWOX, the transversion of G to T might influence four microRNAs’ binding ability to WWOX mRNA as bioinformatics analysis shown. Of the four SNPs, the hsa-miR-134 is an identified microRNA to regulate WWOX expression\(^\text{31}\). Meaningfully, expression data in lymphoblastic cells of CHB from microarray platforms showed that the rs383362T variants have lower WWOX expression than the rs383362GG genotype. These indicated a conjectural mechanism underlying the effect of rs383362G>T to COPD susceptibility. The SNP also interacted with biomass as fuels. Biomass exposure is an important risk factor of COPD, from which smoke can produce environmental stimuluses and induce oxidative damage\(^\text{32}\). WWOX can be downregulated by such exposures\(^\text{33}\). Therefore, subjects with T genotypes are more vulnerable to suffer oxidative damage and thus predispose to develop COPD.

Associations between the WWOX SNPs and several human diseases have been reported including thyroid carcinoma\(^\text{34}\), prostate cancer\(^\text{35,36}\), breast cancer\(^\text{37}\), and lung cancer\(^\text{26}\). Now, we identified the WWOX SNP rs383362G>T to be associated with COPD risk. Meanwhile, the adverse effect driven by the rs383362T allele was increased by a slight in patients with GOLD stage III than those with GOLD stage II or I. The effect was not significant in...
stage IV, which may be due to the limited sample size. Furthermore, the SNP rs383362G > T was also correlated with FEV1/FVC and FEV1/predicted-FEV1, both of which are essential on COPD diagnosis. The rs383362T variants also caused more annual average decline in FEV1 than the rs383362GG genotype. Decline in FEV1 is a marker for airflow obstruction and serves as primary symptom of both developing COPD and exacerbation. These correlations suggested that the SNP has direct effect on pulmonary function, further supporting its association with COPD risk. Taken together, The SNPs might be not only a genetic biomarker for COPD onset but also a predictor for COPD severity.

Some limitations may affect the validity of current study. First, the portion of GOLD stage in our study was a little inconsistent with the parent population as suggested with respect to the high frequency of stage I COPD in the current study, suggesting a serious selection bias in the current study that influenced the external validity. Second, some important confounders such as passive smoking were not concerned in this study, which might cause confounding bias and thus affect internal validity. Third, the relatively small sample size of the nuclear pedigrees in family based analysis might also cause error results. However, the family-based analysis effectively controlled some confounding biases. Because consistent associations were observed in both the case-control study and family based analysis, and the SNP got Bioinformatics data support to be functional, it is conceivable that our finding was not achieved by chance.

To conclude, our study identified a putatively functional SNP rs383362G > T of WWOX that confers an increased risk of COPD, the rs383362T risk allele was more prone to over transmitted from parents to sick offspring. This SNP might be a genetic biomarker to predict risk of COPD in Chinese.

Materials and Methods

Case-control study. A two-stage case-control study was conducted in a southern Chinese population and an eastern Chinese population, which have been described in previously published studies. COPD patient was defined as subjects with FEV1/FVC < 70% after inhalation of 400μg salbutamol and with some chronic airway symptoms including chronic cough, dyspnea, sputum production or wheezing. In brief, the southern Chinese population included 1025 COPD patients and 1061 age (±5 years) and sex frequency-matched controls, of which 697 cases were recruited from three communities (Liwan, Xicun and Zhanqian communities) in Guangzhou city based on annual cross-sectional surveys of COPD, 328 cases were recruited from physical examination centers of three hospitals in Guangzhou city including the Guangzhou Chest Hospital, the third Affiliated Hospital of Guangzhou Medical University and the third Affiliated Hospital of Sun Yat-sen University. All controls were selected from individuals with normal lung function (i.e., FEV1/FVC > 70%) who participated in the annual cross-sectional surveys of COPD. The eastern Chinese population comprised 486 COPD patients and 616 normal controls, of which the cases were enrolled from the physical examination center of the Second Affiliated Hospital of Soochow University and the controls were selected from a database consisting of 3,500 individuals based on a physical examination in Suzhou city. All subjects are generally unrelated Han Chinese. Moreover, a long-term follow-up on lung function monitoring was performed in part of the southern Chinese population for years as previously reported. Heretofore, 427 individuals were successfully followed up and had at least four years’ data of lung function between 2002 and 2010 with annual spirometry test. Each subject was scheduled for an interview with a structured questionnaire to provide data on age, sex, smoking status, pack-years smoked, biomass using and to donate 5 mL peripheral blood after a written informed consent had obtained. The definitions of smoking status and biomass as fuels were described in our previous publications. Briefly, those participants who had smoked < 100 cigarettes in their lifetime were defined as never smokers. Otherwise, they were classified as ever smokers. Biomass as fuels means that people use bio-crop stalks and wood as fuels. The study was approved by the institutional review boards of Guangzhou Medical University and Soochow University. The methods were carried out in accordance with the approved guidelines.

Family-based analysis. A family-based analysis was conducted in a southern Chinese population between September 2010 and September 2012. COPD diagnosis was consistent with the above COPD definition. Briefly, 148 COPD probands were originally recruited from the Fifth People's Hospital of Dongguan City. Among them, 113 were males with a mean age as 67.9 ± 12.9 year-old and 35 were females with a mean age as 65.6 ± 16.3 year-old. The probands’ immediate family members who were mostly the probands’ brothers, sisters and offspring were then asked to take a diagnostic test of COPD. To the exclusion of those who did not finish or achieve lung function test, 342 individuals were enrolled included 69 COPD cases and 273 healthy people. According to the definition of nuclear family, there were 95 nuclear families in the total population (n = 308), of which 26 (27.4%, n = 88) families had both parents, 29 (30.5%, n = 95) families had one single parent and at least one brother or sister, and 40 (42.1%, n = 125) had no available parents but at least two brothers or sisters. For these 95 COPD probands of nuclear families, 70 were males with a mean age as 67.0 ± 12.5 year-old and 25 were females with a mean age as 65.8 ± 17.9 year-old. All subjects are generally unrelated Han Chinese. Each subject provided data using a questionnaire as above described and donated 5 mL peripheral blood after a written informed consent was obtained. The analysis was approved by the institutional review boards of Guangzhou Medical University. The methods were carried out in accordance with the approved guidelines.

SNPs selection and genotyping. We selected 5 putatively functional tagSNPs that are rs10220974C > G, rs12918952G > A in exons, rs383362G > T, rs12828G > A in 3’-UTR as previously described. The 5 SNPs were genotyped using the TaqMan allelic discrimination Assay on the ABI7500 system with self-designed primers and probes as shown in Supplementary Table S2.

Statistical analysis. The chi-square test was used to assess differences in the frequency distribution of SNPs’ genotypes between the cases and controls. The PROC HAPLOTYPE procedure in SAS/Genetics software was
used to infer haplotype frequencies from the observed genotypes of SNPs. The unconditional logistic regression model without or with adjustment for these covariates, which were significantly associated with COPD risk including age, sex, pack-years smoked, biomass as fuels and sample source region, was used to estimate the odds ratio (OR) and 95% confidence interval (95%CI) to measure effect of each SNP as well as haplotype on COPD risk. The multiplicative interaction analysis was applied for assessing the interaction between each SNP and surrounding factors. The family base-association test (FBAT) software was used to perform transmission disequilibrium test (TDT) and transmission disequilibrium of patient-normal (SDT) to analysis the effect of WWOX SNPs and haplotypes on susceptibility of COPD. The one-way ANOVA test was performed to assess the effect of WWOX SNPs on pre-bronchodilator pulmonary function traits. All tests were two-sided by using the SAS software (version 9.2; SAS Institute, Cary, NC). P < 0.05 was considered to be statistically significant.

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Author Contributions
C.L.X. performed the experiments, collected the data and wrote the manuscript; X.L.C. and F.M.Q. helped to analysis the data; L.S.Z., D.W., J.S.C. and D.S.H. participated in the community survey and collected data; L.Y. and J.C.L. conceived of the study, designed the experiments and supervised all aspects of the study.

Additional Information
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