Prevalence and risk factors of airflow limitation in a Mongolian population in Ulaanbaatar: Cross-sectional studies

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Abstract

The burden of chronic obstructive pulmonary disease (COPD) is expected to increase in the coming decades. In Ulaanbaatar, Mongolia, air pollution, which has been suggested to correlate with COPD, is a growing concern. However, the COPD prevalence in Ulaanbaatar is currently unknown. This study aims to estimate the prevalence of airflow limitation and investigate the association between airflow limitation and putative risk factors in the Mongolian population. Five cross-sectional studies were carried out in Ulaanbaatar. Administration of a self-completed questionnaire, body measurements, and medical examination including spirometry were performed in 746 subjects aged 40 to 79 years living in Ulaanbaatar. The age- and sex-standardized prevalence of airflow limitation in Ulaanbaatar varied widely from 4.0 to 10.9% depending on the criteria for asthma. Age, body mass index (BMI), and smoking habit were independent predictors for airflow limitation while residential area and household fuel type were not significant. In conclusion, prevalence of putative COPD was 10.0% when subjects with physician-diagnosed asthma were excluded from COPD. Older age, lower BMI, and current smoking status were putative risk factors for airflow limitation. This prevalence was consistent with reports from Asian countries.

Introduction

Chronic obstructive pulmonary disease (COPD) is characterized by persistent airflow limitation caused by a mixture of small airway disease and parenchymal destruction [1]. According to WHO estimates, COPD is the third leading cause of death globally [2]. There were 3.1 million COPD deaths in 2012, corresponding to 5% of all deaths worldwide. The morbidity and mortality of COPD, and its associated economic and social burdens, are increasing globally even in the low- and middle-income countries [3, 4]. The most evident risk factor for COPD is cigarette smoking, with COPD prevalence correlating with tobacco smoking prevalence. In developing countries, outdoor and indoor air pollution is also a major risk factor of COPD [5].
The prevalence and burden of COPD are expected to increase in the coming decades due to high smoking rates and severe air pollution in developing countries, and population aging in developed countries.

COPD prevalence varies according to survey methods, diagnostic criteria, and analytical methods. The prevalence of physiologically defined COPD in adult subjects over 40 years has been reported to be 9–10% although there are regional or methodological differences [6]. Amongst Asian countries, COPD prevalence has been reported to be 10.9% in Japan [7], 13.7% in Korea (data from the Korea National Health and Nutrition Examination Survey) [8], 8.2% and 11.4% in China [9], and 13.9% in Manila, Philippines [10].

Ulaanbaatar, the Mongolian capital, is one of the world’s worst air-polluted cities and concerns about air pollution are growing [11]. Coal and biomass fuel consumption is the major cause of indoor and outdoor air pollution in Ulaanbaatar [12], and increased coal consumption in winter is attributed to increased household use of coal-fired stoves. This increase in household coal consumption is partly due to population influx from rural areas to Ulaanbaatar, leading to the development of districts comprised of traditional tents with poor infrastructure (termed “ger” in Mongolia). People living in ger districts use smoke-rich fuel for household use, resulting in frequent smog and indoor air pollution in Ulaanbaatar. Indoor air pollution is a known important risk factor for COPD development [13–15], while other reports suggest that outdoor air pollution also affects lung function and respiratory symptoms although the effect seems to be small [1, 16–18].

The COPD prevalence in Ulaanbaatar is currently unknown. Here, we have conducted cross-sectional studies to assess the determinants and estimated prevalence of airflow limitation among Mongolian subjects aged 40–79 years living in Ulaanbaatar.

**Methods**

**Study design**

We conducted five cross-sectional studies among Mongolian adults aged 40 to 79 years at eight facilities (two hospitals and six community clinics) in Ulaanbaatar from 2012 to 2013 (May and September, 2012 and March, July, December, 2013). Participants were recruited by announcements advertising the health survey. Subjects received a self-completed questionnaire containing questions pertaining to age, gender, occupation, respiratory symptoms, and medical history. At the time of questionnaire administration, height and body weight were measured, and medical interviews, auscultations, and pulmonary function tests by respiratory specialists were conducted. Each survey included different sets of subjects and no duplicative subjects.

**Subjects**

The present study included an initial 1,030 male and female community volunteers. Of these, 29 (2.8%) and 3 (0.3%) subjects were excluded due to ineligible age (< 40 or ≥ 80 years old) and lack of gender information, respectively. The remaining 998 (96.9%) were subjected to pulmonary function tests, however, the physician diagnosed that 71 (6.9%) subjects were contraindicated for spirometry due to conditions such as active tuberculosis, postpneumonectomy, or high blood pressure. Furthermore, spirometry on 24 (2.3%) subjects could not be performed due to a facility electric outage, and 57 (5.5%) subjects were excluded from analysis due to other reasons such as poor reproducibility, inappropriate spiromograms, and/or mechanical issues. Lastly, 100 (9.7%) subjects were excluded due to restrictive ventilatory impairment (FEV1/FVC > 0.7 and FVC < 80% of predicted value). Characteristics of those subjects with...
restrictive ventilatory impairment was shown in S1 Table. Consequently, 746 (72.4%) subjects were eligible for further analysis.

**Pulmonary function test**

A pre-bronchodilator pulmonary function test using the HI-105 spirometer (CHEST M.I., Inc., Tokyo, Japan) was conducted by the respiratory specialists. Spirometers were calibrated before each survey. At least three spirograms were obtained, and the highest forced expiratory volume in one second (FEV1) and forced ventilatory capacity (FVC) values were applied for diagnosis. Mongolian and Japanese respiratory specialists assessed the reproducibility and quality of spirograms (volume-time curve and flow-volume loop) and the invalid spirograms, which showed artefacts, insufficient inspiration, or blow-out, were excluded from analysis. Severity of airflow limitation was classified by the GOLD criteria (FEV1/FVC < 0.7) with the predicted FEV1 defined by GLI2012 reference equations for North East Asians [1, 19].

**Data handling and statistical analyses**

All data were anonymized, and managed as electronic data for the analysis. Prevalence of airflow limitation was age- and sex-standardized by direct standardization using Mongolian national population data published by the United Nations [19, 20]. Multiple logistic regression analyses were carried out by forced inclusion procedure for all the variables. Odds ratios (ORs) for presenting airflow limitation were adjusted for age group, sex (coded as 0 for female and 1 for male), body mass index (BMI) (coded as 0 for BMI ≥ 25.0 and 1 for BMI < 25.0), smoking status (coded as 0 for non-smokers, 1 for former smokers, and 2 for current smokers), household fuel (coded as 0 for smoke-free fuels such as gas and electricity and 1 for smoke-rich fuels such as coal, wood and dry animal dung fuel), and residential district (coded as 0 for urban areas and 1 for ger districts). The omnibus tests of model coefficient were shown to be significant (P < 0.01) for all models. Correlation coefficients (r) among predictive values were tested, and all r-values were less than 0.55, showing no multicollinearity. Statistical analyses including Welch’s t-test for parametric analysis of two groups, $\chi^2$-test for analysis of categorical data, and multiple logistic regression analysis were performed using the statistical software package JMP version 11 (SAS Institute Inc., Cary, NC, USA). P values less than 0.05 on both sides were considered to be statistically significant.

**Ethics, consent, and permissions**

The present study was approved by the Clinical Ethical Review Board of Kurume University School of Medicine. Before investigation, the participants were provided with explanations in person as to the purpose and method of the study, as well as information regarding the handling of the results. The study was carried out upon receipt of written consent.

**Results**

**Characteristics of participants**

Mongolian subjects (n = 1,030) were recruited from the general population in Ulaanbaatar. There were 746 (72.4%) eligible subjects for this study after excluding subjects who were out of the age range, unknown gender, invalid spirometry, in poor physical condition, or with restrictive ventilatory impairment. Characteristics of participants are presented in Table 1. Mean ages of males and females were similar. Male-to-female ratio was 0.53 (258: 488), and BMI, smoking status, residential district, and severity of airflow limitation according to GOLD criteria were significantly different between sexes. When subjects were compared with the general Mongolian
population of the same age group in 2012 to 2013 [20, 21], the age characteristic of this study population was slightly older with a significantly lower percentage of male subjects (34.6% of the study subjects) than the general population (47.4% of the general population) (Table 2).

Expected prevalence of airflow limitation

Based on spirometry data, the unadjusted prevalence of airflow limitation in the study subjects was 11.5% (Table 2). Severity of airflow limitation according to the GOLD criteria using the

### Table 1. Characteristics of the study subjects with valid spirometry.

| Characteristics               | Male        | Female       | Statistical analysis |
|-------------------------------|-------------|--------------|----------------------|
| Age (Mean ± SD)               | 54.1 ± 10.7 | 54.1 ± 9.5   | N.S.                 |
| BMI (kg/m²) (Mean ± SD)       | 25.8 ± 4.0  | 27.9 ± 4.9   | P < 0.001            |
| Obese and overweight (BMI ≥ 25.0) | 128 49.6   | 347 71.1    | P < 0.001            |
| Normal and underweight (BMI < 25.0) | 130 50.4   | 141 28.9    |                       |
| Smoking status                |             |              |                      |
| Never smoker                  | 88 34.1     | 416 85.2    | P < 0.001            |
| Former smoker                 | 32 12.4     | 21 4.3      |                       |
| Current smoker                | 138 53.5    | 51 10.5     |                       |
| Household fuel                |             |              |                      |
| Smoke-free                    | 89 34.5     | 199 40.8    | N.S.                 |
| Smoke-rich                    | 169 65.5    | 289 59.2    |                       |
| Residential district          |             |              |                      |
| Urban area                    | 100 38.8    | 256 52.5    | P < 0.001            |
| Ger district                  | 158 61.2    | 232 47.5    |                       |
| Airflow limitation            |             |              |                      |
| Stage I                       | 18 7.0      | 10 2.0      | P < 0.005            |
| Stage II                      | 19 7.4      | 33 6.8      |                       |
| Stage III + IV                | 4 1.6       | 2 0.4       |                       |
| None                          | 217 84.1    | 443 90.8    |                       |
| Total (n = 746)               | 258 34.6    | 488 65.4    |                       |

### Table 2. Prevalence of airflow limitation in study population standardized by Mongolian population.

| Characteristic | Study subject | Mongolian population (40–79 years old) | Prevalence of airflow limitation (%) |
|----------------|---------------|----------------------------------------|-------------------------------------|
|                | N              | %                                      | N (%)                               |
| Age group      | N (%)         | N (×10^5) % | N (×10^5) % | Male | Female | Overall | Crude | Standardized |
| 40–49          | 267 35.8      | 178 50.6  | 186 47.5  | 6 6.3 | 8 4.7   | 14 5.2 | 5.5   |
| 50–59          | 264 35.4      | 106 30.1  | 121 31.0  | 15 17.2 | 15 8.5  | 30 11.4 | 12.6  |
| 60–69          | 146 19.6      | 46 13.1   | 54 13.8   | 12 27.9 | 12 11.7 | 24 16.4 | 19.1  |
| 70–79          | 69 9.2        | 22 6.3    | 30 7.7    | 8 25.0 | 10 27.0 | 18 26.1 | 26.2  |
| Total          | 746 100.0     | 352 100.0 | 392 100.0 | 12.4 |
| Sex            |               |            |            |       |        |        |       |
| Female         | 488 65.4      | -          | 392 52.7   | - 45 | 9.2    | 86 11.5 | 8.5   |
| Male           | 258 34.6      | 352 47.3   | -          | 41 15.9 | -    |        | 13.6  |

**Age- and sex-standardized prevalence** 10.9

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GLI 2012 reference equations for North East Asians was: mild, 43.9% (n = 18, male) and 22.2% (n = 10, female); moderate, 46.3% (n = 19, male) and 73.3% (n = 33, female); and severe or higher, 9.8% (n = 4, male) and 4.4% (n = 2, female). No females were classified as having very severe airflow limitation. Crude and sex-standardized prevalence of airflow limitation were increased with increasing age, and crude and age-standardized prevalence of airflow limitation in males were higher than in females (Table 2). Age- and sex-standardized prevalence of airflow limitation was 10.9%. The prevalence of airflow limitation stratified by subgroup is shown in Table 3. Multiple logistic regression analyses to estimate odds ratios for presenting airflow limitation were performed using age, sex, BMI, smoking status, household fuel type, and residential district as predictive variables. Crude ORs were significant higher in older subjects, males, the lower BMI group (< 25), and in current smokers. Of these, an older age, a lower BMI, and current smoking status maintained significance after adjustment (Table 3). The prevalence for subjects living in ger districts was higher than those in urban area but OR was not significant.

### Differentiation of asthma from putative COPD

To assess the possibility of asthma, the study population was stratified by the number of affirmative responses to four asthma-related questions; self-reported history of asthma, history of wheezing in the past 12 months, frequent or occasional wheezes, and physician-diagnosed asthma (Table 4). The number of subjects with physician-diagnosed asthma was 17 (4 male; 13 female) (data not shown). There were 8 subjects (2 male; 6 female) with physician-diagnosed asthma also had airflow limitation. The crude prevalence of airflow limitation ranged from 4.3% to 11.5% depending on the number of affirmative responses to asthma-related questions. The prevalence ranged from 4.0% to 10.9% when they were standardized by age and sex. When subjects with physician-diagnosed asthma were excluded from putative COPD, the

| Characteristics | Lower BMI Group (< 25) | Higher BMI Group (≥ 25) |
|-----------------|------------------------|-------------------------|
| Age group       |                         |                         |
| 40–49           | 5.2 (5.5) 1.00 (ref)     | 5.5 (ref) 1.00 (ref)     |
| 50–59           | 11.4 (12.6) 2.32 (1.22–4.60) | 19.1 (3.56) 1.80–7.28 |
| 60–69           | 16.4 (19.0) 3.56 (1.80–7.28) | 7.0 (4.06) 2.02–8.45 |
| 70–79           | 26.1 (26.2) 6.38 (3.00–13.86) | 8.66 (8.66) 3.89–19.81 |
| Sex             | Male (Female)           | Male (Female)           |
| Male            | 15.9 (9.2) 13.6 (8.5)    | 1.86 (1.18–2.93) 1.13 (0.63–2.03) |
| Male            | 13.6 (8.5) 1.86 (1.18–2.93) | 1.13 (0.63–2.03)   |
| BMI             | < 25.0 (> 25.0)         | < 25.0 (> 25.0)         |
| < 25.0          | 16.6 (8.6) 14.8 (8.2)    | 2.11 (1.34–3.32) 2.05 (1.27–3.32) |
| > 25.0          | 8.5 (1.00 (ref) 1.00 (ref) |
| Smoking status  | Never                  | Former                  |
| Never           | 9.7 (8.5) 1.00 (ref)     | 9.4 (6.1) 0.97 0.32–2.34 |
| Current         | 16.9 (14.8) 1.89 (1.16–3.05) | 1.91 (1.02–3.56) |
| Household fuel  | Smoke-rich (Smoke-free) | Smoke-rich (Smoke-free) |
| Smoke-rich      | 11.4 (11.8) 8.0 (9.0)    | 0.96 (0.61–1.53) 0.79 (0.45–1.39) |
| Residential area| Ger district (Urban area)| Ger district (Urban area)  |
| Ger district    | 13.1 (9.8) 12.4 (8.4)    | 1.38 (0.88–2.19) 1.55 (0.88–2.77) |

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crude prevalence of putative COPD was 10.5% (age- and sex-adjusted prevalence: 10.0%) in the study population.

**Discussion**

This is the first study on the prevalence of airflow limitation in Mongolia. We investigated lung function in Mongolian subjects aged 40 to 79 years living in Ulaanbaatar and found that the crude prevalence of airflow limitation was 11.5%. Multiple logistic regression analysis revealed that an older age, lower BMI, and current smoking status were independent predictive variables for airflow limitation. In this study, COPD may be confounded by the presence of asthma as post-bronchodilator spirometry was not conducted. When subjects with airflow limitation were stratified according to different criteria for asthma, putative COPD prevalence varied between 4.3 to 11.5%.

The prevalence of COPD shows interstudy variation depending on the diagnostic criteria. Halbert et al. reported that the pooled prevalence of 37 estimates for COPD was 7.6% (95% Confidence interval (CI): 6.0–9.5) [6]. Of these 37 estimates, the differences based on criteria were relatively large, for example, spirometric criteria indicated 9.2% prevalence while patient-reported diagnosis resulted in 4.9%. The most popular criteria using spirometric results were based on the GOLD criteria [1, 6]. The crude prevalence of airflow limitation (GOLD stage I (FEV1/FVC < 0.7)) in the present study was 11.5% when the possibility of asthma was not considered and 10.5% when subjects with physician-diagnosed asthma were excluded from putative COPD (Table 4). Furthermore, the age- and sex-standardized prevalence excluding subjects with physician-diagnosed asthma was 10.0%. When criterion used in BOLD study (GOLD stage II or higher (FEV1/FVC < 0.7 and FEV1 < 80% of predicted value)) was employed, the crude prevalence of airflow limitation was 7.8% and standardized prevalence was 7.3% (S2 Table) [6]. Age- and sex-standardized prevalence excluding physician-diagnosed asthma was 6.6% (S3 Table). Our results are comparable with other prevalence studies [6–9].

BMI was significantly lower in subjects with airflow limitation than those with normal lung function (28.1 ± 5.0 (Normal) vs. 25.7 ± 2.9 (Airflow limitation), P < 0.001). Due to the cross-sectional design of the present study, it is not clear whether the low BMI is a risk factor or secondary to airflow limitation, although low BMI has been reported to be a risk factor for COPD development [22, 23]. The present study also showed that lower BMI was one of the independent predictive variables for presenting airflow limitation, suggesting that low BMI is strongly associated with COPD development. Smoking and second-hand smoke have been reported to be the most important risk factors for COPD development [1, 24, 25]. Smoking rates in Asian

| Diagnosis criteria for asthma | Possible asthma [n (%)] | Putative COPD [n (%)] | Prevalence of putative COPD in study population (%) |
|------------------------------|-------------------------|-----------------------|--------------------------------------------------|
| None of four criteria affirmative | 54 (62.8) | 32 (37.2) | 4.3 (Age- and sex-standardized prevalence: 4.0) |
| Any one of four criteria affirmative | 36 (41.9) | 50 (58.1) | 6.7 (6.2) |
| Any two of four criteria affirmative | 16 (18.6) | 70 (81.4) | 9.4 (8.7) |
| Any three of four criteria affirmative | 4 (4.7) | 82 (95.3) | 11.0 (10.4) |
| Four of four criteria affirmative | 0 (0.0) | 86 (100.0) | 11.5 (10.9) |
| Physician-diagnosed asthma | 8 (9.3) | 78 (90.7) | 10.5 (10.0) |

**Criteria for asthma diagnosis:** self-reported history of asthma; history of wheezing in the past 12 months; frequent or occasional wheezes; physician-diagnosed asthma

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countries are high especially for males [26], and in the present study (Table 1). Crude OR for airflow limitation was significantly higher in males (Table 3), but significance was lost when OR was adjusted, reflecting the high smoking prevalence in male subjects. A current smoking status was found to be a significant risk factor for airflow limitation even after the adjustment (Table 3). Smoking cessation or repeated attempts at smoking cessation has been reported to be effective in preventing lung function decline and decreasing the prevalence of respiratory symptoms even with subsequent cessation failure [27–31]. In contrast to current smokers, former smoking was not significant risk factor for airflow limitation although the number of former smokers in the present study was small (Table 3). Therefore, we can speculate that smoking cessation is effective in Mongolia for preventing lung function deterioration. Furthermore, for the subjects with airflow limitation of GOLD stage II or higher, both current and former smoking were not significant risk factors for airflow limitation (S4 Table). These results suggest that smoking does not have large impact on the risk for moderate to severe airflow limitation (GOLD stage II or higher) but for mild (GOLD stage I) airflow limitation in this population. It is consistent with the higher prevalence of mild airflow limitation (GOLD stage I) in male which showed higher smoking prevalence than in female. In general, people using smoke-rich fuel such as wood and coal for household use were exposed to indoor air pollution. Exhausted smoke from gers is a primary cause of ambient air pollution in ger districts. Thus, people living in ger districts are exposed to ambient air pollution for longer periods than people living in urban area although air pollution covers urban areas as well, depending on time and direction of the wind. Since indoor and outdoor air pollution is reported to be a cause of airflow limitation [1, 13–18], we included the residential district and household fuel type as independent variables in the multivariate analysis. Here, residential area and household fuel type was not a significant predictor of airflow limitation (Table 3). Since actual exposure to air pollution was not measured in the present study, subject responses accurately reflect exposure to air pollution cannot be validated. The prevalence of airflow limitation in ger districts was higher than that in urban areas but did not reach significance (Table 3). This result suggests that the effect of air pollution on airflow limitation is relatively small in the present study. Our previous study suggested that subjects living in ger districts with ventilatory impairment showed significantly higher prevalence of respiratory symptoms, and decreased health status in the cold season (unpublished data). Therefore, we can speculate that the effect of air pollution was one of the risk factor for the exacerbation of symptoms and health status rather than for the decline in lung function.

Since patients do not present themselves to the physician for examination until they have severe symptoms or significant impairment, COPD is likely underestimated [32]. We assessed the multivariate model for predicting airflow limitation using respiratory symptoms as independent variables to encourage subjects with certain symptoms to consult a physician for early detection of COPD. In this study, subjects presenting with the symptom “usually cough up sputum first thing in the morning” were liable to develop airflow limitation even after adjustment (S1 Fig). Our result is consistent with a report by Kessler et al., where the fluctuation of COPD symptoms over the day in COPD patients and all COPD-related symptoms including phlegm were most predominant upon waking in the morning [33]. Encouraging subjects who suffer respiratory symptoms in the morning to consult a physician would be effective for early diagnosis of COPD and reduction of its associated burdens.

There are several limitations in this study. First, whether the putative risk factors found in the present study indicates a risk of airflow limitation or its consequences is unclear due to the cross-sectional design of this study. Second, in the present study, postbronchodilator spirometry to distinguish asthma from COPD could not be performed as this study was conducted as a screening. Since the biggest concern about accuracy of the estimation in this study was how
many subjects with pure asthma without COPD were included in the subjects with airflow limitation, we considered the possibility of bronchial asthma in Table 4 using differential diagnostic criteria. As shown in Table 4, the standardized prevalence ranged 4.0% to 10.9% corresponding the proportion of false positive ranged 63.3% to 0%. In Mongolia, asthma prevalence was reported to be 4.7% based on symptoms [34], and Viinanen et al. reported the prevalence of asthma in Ulaanbaatar to be 2.1% based on spirometry data although both studies did not show COPD prevalence [35]. It is difficult to distinguish asthma from COPD in smokers and older adults [36], as some patients may have clinical features of both asthma and COPD known as asthma-COPD overlap syndrome (ACOS). In the present study, 2.3% of subjects were diagnosed with asthma by respiratory specialists based on the symptoms and medical history (Table 4). This is comparable to asthma prevalence in Ulaanbaatar based on spirometry data [35]. Lastly, the age distribution and male-to-female ratio of subjects were biased in the present study as study subjects were older and with a lower male-to-female ratio compared to the general Mongolian population. The overall prevalence of airflow limitation in this study might be affected by the lower male-to-female ratio because the proportion of smokers was much higher in males than in females. Nonetheless, this study population covered approximately 0.1% of the Mongolian population aged 40 to 79 years [20, 21] and the direct standardization by age and sex were carried out using population data [20]. Furthermore, multivariate analyses adjusted for age, sex, and other factors potentially affecting airflow limitation were carried out. Hence, the results obtained in this study reflect the actual situation in Mongolia although generalizations are limited.

Conclusions
We investigated the prevalence of airflow limitation in subjects aged 40 to 79 in Ulaanbaatar, Mongolia. The age- and sex-standardized prevalence of putative COPD ranged from 4.0 to 10.9% depending on the asthma criteria and the age- and sex-standardized prevalence was 10.0% when patients with physician-diagnosed asthma were excluded from COPD. An older age, lower BMI, and current smoking status were independent risk factors for airflow limitation. The putative COPD prevalence in this study was consistent with other reports from Asian countries.

Supporting information
S1 Fig. Respiratory symptoms associated with airflow limitation. Multiple logistic regression analyses were carried out by a stepwise selection procedure used with forced inclusion of specific variables. Respiratory symptoms were added as additional predictive variables as follows; 1. Does the weather affect your cough? (Yes/No or No cough); 2. Have you ever coughed up sputum from your chest when you do not have a cold? (Yes/No); 3. Do you usually cough up sputum from your chest first thing in the morning? (Yes/No); 4. How frequently do you wheeze? (Occasionally or more often/Never); 5. Do you have or have you had any allergies? (Yes/No); 6. Do you suffer from any infectious disease? (Yes/No). Odds ratios were adjusted for age group, sex, BMI, smoking status, household fuel, and residential district. The vertical short line represents odds ratios; the horizontal line represents 95% confidence interval. The omnibus tests of model coefficient were shown to be significant (P < 0.01) for all models. Correlation coefficients (r) among predictive values were tested, and all r-values were less than 0.55, indicating no multicollinearity. P values less than 0.05 were considered to be statistically significant. (TIF)
S1 Table. Characteristics of the subjects with restrictive ventilatory impairment. (DOCX)

S2 Table. Prevalence of airflow limitation in study population standardized by Mongolian population when the cutoff value of airflow limitation was set at FEV1/FVC < 0.7 and FEV1 < 80% of predicted value (GOLD stage II or higher). (DOCX)

S3 Table. Prevalence of putative COPD (GOLD stage II or higher) by diagnostic category of asthma. (DOCX)

S4 Table. Factors affecting the prevalence of airflow limitation (GOLD stage II or higher). (DOCX)

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References
1. Global Initiative for Chronic Obstructive Lung Disease. Global strategy for the diagnosis, management, and prevention of chronic obstructive pulmonary disease, update 2015. http://www.goldcopd.org/uploads/users/files/GOLD_Report_2015.pdf (Accessed Jan 25, 2016)
2. World Health Organization. The top 10 causes of death (updated May 2014). (http://www.who.int/mediacentre/factsheets/fs310/en/) (Accessed Jan 25, 2016)
3. Lopez AD, Shibuya K, Rao C, Mathers CD, Hansell AL, Held LS, et al. Chronic obstructive pulmonary disease: current burden and future projections. Eur Respir J 2006; 27(2):397–412. https://doi.org/10.1183/09031936.06.00025805 PMID: 16452599
4. Mathers CD, Loncar D. Projections of global mortality and burden of disease from 2002 to 2030. PLoS Med 2006; 3(11):e442. https://doi.org/10.1371/journal.pmed.0030442 PMID: 17132052

5. Salvi SS, Barnes PJ. Chronic obstructive pulmonary disease in non-smokers. Lancet 2009; 374(9691):733–743. https://doi.org/10.1016/S0140-6736(09)61303-9 PMID: 19716966

6. Halbert RJ, Natoli JL, Gano A, Badamgarav E, Buist AS, Mannino DM. Global burden of COPD: systematic review and meta-analysis. Eur Respir J 2006; 28(3):523–532. https://doi.org/10.1183/09031936.06.00124605 PMID: 16611654

7. Fukuchi Y, Nishimura M, Ichinose M, Adachi M, Nagai A, Kuriyama T, et al. COPD in Japan: the Nippon COPD Epidemiology study. Respirology 2004; 9(4):458–465. https://doi.org/10.1111/j.1440-1843.2004.00637.x PMID: 15612956

8. Park H, Jung SY, Lee K, Bae WK, Lee K, Han JS, et al. Prevalence of chronic obstructive pulmonary disease in Korea using data from the fifth Korea National Health and Nutrition Examination Survey. Korean J Fam Med 2015; 36(3):128–134.

9. Zhong N, Wang C, Yao W, Chen P, Kang J, Huang S, et al. Prevalence of chronic obstructive pulmonary disease in China. A large, population-based survey. Am J Respir Crit Care Med 2007; 176(8):753–760. https://doi.org/10.1164/rccm.0612-1749OC PMID: 17575095

10. Buist AS, McBurnie MA, Vollmer WM, Gillespie S, Burney P, Mannino DM, et al. International variation in the prevalence of COPD (The BOLD Study): a population-based prevalence study. Lancet 2007; 370:741–750. https://doi.org/10.1016/S0140-6736(07)61377-4 PMID: 17765523

11. World Health Organization. Ambient (outdoor) air pollution database 2014, by country and city. http://www.who.int/quantifying_ehimpacts/national/countryprofile/aap_pm_database_may2014.xls?ua=1 (Accessed Jan 25, 2016).

12. Amarsaikhan D., Battsengel V., Nergui M., Ganzorig M., Bolor G. A study on air pollution in Ulaanbaatar city, Mongolia. J Geosci Environ Protect 2014; 2:123–128.

13. Ezzati M. Indoor air pollution and the health in developing countries. Lancet 2005; 366(9480):104–106. https://doi.org/10.1016/S0140-6736(05)66845-6 PMID: 16005317

14. Smith KR, Mehta S, Mauzezahl-Feuz M. Indoor air-pollution from household solid fuel use. In: Ezzati M, Lopez AD, Rodgers A, Murray CJL, eds. Comparative quantification of health risks: global and regional burden of disease attributable to selected major risk factors. Geneva: World Health Organization, 2004; 1435–1493, http://www.who.int/publications/crv/volume2/1435-1494.pdf (Accessed Jan 25, 2016).

15. Kurmi OP, Semple S, Simkhada P, Smith WCS, Ayres JG. COPD and chronic bronchitis risk of indoor air pollution from solid fuel: a systematic review and meta-analysis. Thorax 2010; 65:221–228. https://doi.org/10.1136/thx.2009.124644 PMID: 20335290

16. Berend N. Contribution of air pollution to COPD and small airway dysfunction. Respirology 2016; 21(2):237–244. https://doi.org/10.1111/resp.12644 PMID: 26412571

17. Hu G, Zhong N, Ran P. Air pollution and COPD in China. J Thoracic Dis 2015; 7(1):59–66.

18. Abbey DE, Burchette RJ, Knutsen SF, McDonnell WF, Lebowitz MD, Enright PL. Long-term particulate and other air pollutants and lung function in nonsmokers. Am J Respir Crit Care Med 1998; 158(1):289–298. https://doi.org/10.1164/ajrccm.158.1.9710101 PMID: 9655742

19. Quanjer PH, Stanojevic S, Cole TJ, Baur X, Hall GL, Culver BH, et al. Multi-ethnic reference values for spirometry for the 3-95-year age range: the global lung function 2012 equations. Eur Respir J 2012; 40:1324–1343. https://doi.org/10.1183/09031936.00080312 PMID: 22743675

20. United Nations Economic and Social Commission for Asia and the Pacific, ESCAP Statistical Database (2015). http://www.unescap.org/stat/data/statdb/DataExplorer.aspx (Accessed Feb 22, 2016).

21. United Nations Economic and Social Commission for Asia and the Pacific, ESCAP Statistical Database. Statistical yearbook for Asia and the Pacific 2014, http://www.unescap.org/sites/default/files/ESCAP-SYB2014_0.pdf (Accessed Feb 22, 2016).

22. Zhou Y, Wang D, Liu S, Zheng J, Zhong N, Ran P. The association between BMI and COPD: the results of two population-based studies in Guanzhou, China. COPD 2013; 10:1–6.

23. Hanik-Khan RI, Fleg JL, Wise RA. Body mass index and the risk of COPD. Chest 2002; 121:370–376. PMID: 11834645

24. Forey BA, Thornton AJ, Lee PN. Systematic review with meta-analysis of the epidemiological evidence relating smoking to COPD, chronic bronchitis and emphysema. BMC Pulm Med 2011; 11:36. https://doi.org/10.1186/1471-2466-11-36 PMID: 21672193

25. Laniado-Laborin R. Smoking and chronic obstructive pulmonary disease (COPD). Parallel epidemics of the 21st century. Int J Environ Res Public Health 2009; 6(1):209–224. https://doi.org/10.3390/ijerph6010209 PMID: 19440278
26. Islami F, Torre LA, Jemal A. Global trends of lung cancer mortality and smoking prevalence. Transl Lung Cancer Res 2015; 4(4):327–338. https://doi.org/10.3978/j.issn.2218-6751.2015.08.04 PMID: 26380174

27. van Schayck CP, Kaper J. Smoking and COPD: will they ever vanish into smoke? (Editorial) Primary Care Respir J 2006; 15:81–83.

28. Di Stefano A, Capelli A, Lusuardi M, Balbo P, Vecchio C, Maestrelli P, et al. Severity of airflow limitation is associated with severity of airway inflammation in smokers. Am J Respir Crit Care Med 1998; 158 (4):1277–1285. https://doi.org/10.1164/ajrccm.158.4.9802078 PMID: 9769292

29. Anthonisen NR, Connell JE, Kiley JP, Altose MD, Buist AS, et al. Effects of smoking intervention and the use of an inhaled anticholinergic bronchodilator on the rate of decline of FEV1. The Lung Health Study. JAMA 1994; 272(19):1497–1505. PMID: 7966841

30. Murray RP, Anthoniesen NR, Connell JE, Wise RA, Lindgren PG, Greene PG, et al. Effect of multiple attempts to quit smoking and relapses to smoking on pulmonary function. J Clin Epidemiol 1998; 51 (12):1317–1326. PMID: 10086826

31. Kanner RE, Connett JE, Williams DE, Buist AS. Effects of randomized assignment to a smoking cessation intervention and changes in smoking habits on respiratory symptoms in smokers with mild chronic obstructive pulmonary disease: The Lung Health Study. Am J Med 1999; 106(4):406–10.

32. van den Boom G, Rutten-van Molken MPMH, Tirumanna PRS, van Schayck CP, Folgering H, van Weel C. Association between health-related quality of life and consultation for respiratory symptoms: results from the DIMACA programme. Eur Respir J 1998; 11:67–72. PMID: 9543272

33. Kessler R, Partridge MR, Miravitlles M, Cazzola M, Vogelmeier C, Leynaud D, et al. Symptom variability in patients with severe COPD: a pan-European cross-sectional study. Eur Respir J 2011; 37:264–272. https://doi.org/10.1183/09031936.00051110 PMID: 21115606

34. Sonnambu M, Dashdemberel S, Logii N, Nakae K, Chigusa Y, Ohhira S, et al. Prevalence of asthma and allergic rhinitis among adult population in Ulaanbaatar, Mongolia. Asia Pac Allergy 2014; 4:25–31. https://doi.org/10.5415/apallergy.2014.4.1.25 PMID: 24527407

35. Viinanen A, Munhbayarlah S, Zavgee T, Narantsetseg Ts, Koskenvuo M, Helenius H, et al. Prevalence of asthma, allergic rhinoconjunctivitis and allergic sensitization in Mongolia. Allergy 2005; 60:1370–1377. https://doi.org/10.1111/j.1398-9995.2005.00877.x PMID: 16197468

36. Diagnosis of Diseases of Chronic Airflow limitation: Asthma COPD and Asthma-COPD Overlap Syndrome (ACOS). Based on the Global Strategy for Asthma Management and Prevention and the Global Strategy for the Diagnosis, Management and Prevention of Chronic Obstructive Pulmonary Disease. 2015, http://www.ginasthma.org/local/uploads/files/ACOS_2015.pdf (Accessed Jan 25, 2016).