COPD AND OCCUPATION: A RETROSPECTIVE COHORT STUDY OF INDUSTRIAL WORKERS*

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The aim of this paper was to ascertain chronic obstructive pulmonary disease (COPD) prevalence among industrial workers in the Russian Federation and determine relative contribution of smoking and occupational factors to COPD.

We recruited 1,375 workers aged 30 or over. Six hundred and twenty-four of them were occupationally exposed to vapours, gases, dust, and fumes (VGDF). Physical examination and baseline spirometry were performed for all the participants of the study. Those with airflow limitation of FEV₁/FVC<0.70 were considered having COPD and those with presence of cough and sputum production for at least three months in each of two consecutive years were considered having chronic bronchitis (CB), with no overlapping between these 2 groups. Data on occupational history and VGDF levels in the working area were collected from all participants. In total, 105 cases of COPD and 170 cases of CB were diagnosed in the cohort of examined workers. Occupational exposure to VGDF was twice as often present among COPD patients than among both patients with CB and the control group of healthy workers (p<0.05). More than 40 % of COPD patients were occupationally exposed to VGDF above the value of 3.0 of the occupational exposure limit (OEL) and more than 20 % to 6.0 OEL and higher. Overall odds ratio for COPD development due to occupational VGDF exposure was 5.9 (95 % CI=3.6 to 9.8, p=0.0001). Both smoking and VGDF seem to be important for the development of COPD. Analysis of the combined effect of tobacco smoking and occupational noxious particles and gases on COPD development has shown the following order of risk factors based on the strength of their influence: VGDF levels, smoking index, age, and heating microclimate. There is a statistically significant level of relationship and “dose-effect” dependence between occupational exposures to VGDF and the development of COPD. The effect of VGDF composition on the probability of COPD development was not found in the study. Results of this study were used to substantiate the inclusion of COPD into the National List of Occupational Diseases of the Russian Federation.

KEY WORDS: chronic obstructive pulmonary disease, occupational exposure, risk assessment, silica dust, smoking

Chronic obstructive pulmonary disease (COPD) is an increasing cause of chronic morbidity and mortality around the world (1). This disease, which has already affected 44 million people in Europe and is deemed the 4th leading cause of death worldwide, is likely to become the 3rd such cause by 2030 according to the predictions of the World Health Organization (3). Thus, the prognosis of CJ Murray
and AD Lopez about the global burden of COPD, made in 1996 (2), has come true.

The major cause of COPD is smoking. However, tobacco smoking is not the only cause of COPD. According to the last updates of the Global Initiative for Chronic Obstructive Lung Disease (GOLD), occupational exposure is one of the two most important risk factors for COPD (4).

It is well known that tobacco smoke and occupational exposures exert a synergistic effect and increase each other's influence (5). However, relative impacts of each of these factors are poorly understood. Therefore differentiation between the two risk factors' individual effects may be important for planning strategies for the prevention and treatment of COPD.

Strong evidence implicates occupational exposures as one of the causes of COPD (6). A significant part of the literature accumulated over the past two decades demonstrated the relationship between vapour, gases, dust, and fumes (VGDF) and the development of COPD (7-13). However its importance remains underappreciated, especially in the Russian Federation. This applies particularly to the combined effect of occupational exposure and smoking. Since COPD develops predominantly during the working age, a comprehensive analysis of this joint effect seems to be important.

The aims of the present study were to ascertain the prevalence of COPD among industrial workers in the Russian Federation, to establish the relative contributions of smoking and occupational factors, and to investigate the accuracy of the following hypotheses: (1) COPD can be caused by VGDF only, irrespective of smoking; (2) there is a "dose - effect" dependence between VGDF and COPD development; and (3) the influence of smoking and VGDF on COPD development is similar.

Results of this study were used to corroborate the inclusion of COPD into the National List of Occupational Diseases of the Russian Federation.

**METHODS**

**Study design**

The retrospective cohort study was chosen due to its capability to study the outcomes after the exposure; the ability to yield true incidence rates, values of relative risks, and other measures of association.

**Study population**

One thousand three hundred and seventy-five workers, 879 men and 496 women, aged 30 to 60 years, were selected randomly for this study during periodical medical examinations of industrial workers, residents of three largest cities of the Republic of Tatarstan between June 2005 and December 2008 (Figure 1). The data were collected from workers of five enterprises - two foundry plants, one aircraft plant, and two oil extracting enterprises. Eligible participants were those who met the following inclusion criteria: (1) voluntary informed consent, (2) working for an industrial enterprise for at least five years. Exclusion criteria were: (1) refusal to participate in the study, (2) any other respiratory disease except COPD or chronic bronchitis (CB). Six hundred and twenty-four of included subjects were occupationally exposed to VGDF: 327 of these were exposed to silica dust, 244 to nonfibrogenic dusts, and 53 to nonfibrogenic dusts with vapours of irritants and sensitizers.

After medical examination, 22 workers were excluded from the study because four of them had been diagnosed with pneumoconiosis, and 18 with asthma. We compared three groups of workers: patients with COPD (N=105), those with CB (N=170), and a reference group of healthy workers with no signs of COPD/CB (N=1,100) (Figure 1).

**Figure 1 Selection of the study population**

A hierarchical population sample was selected as follows: all subjects exposed to occupational hazards in the Republic of Tatarstan (n=230,136); a total number of workers, aged 30 to 60 years, examined at the University Clinic of Occupational Medicine during 2005-2008 (n=9,859), workers who agreed to take part in the study (n=1,397), subjects who had vapour, gases, dust and fumes (VGDF) in their working area (n=624), i.e. 10 excluded people (4 with pneumoconiosis, 6 with asthma), 71 with chronic bronchitis (CB), and 83 with chronic obstructive pulmonary disease (COPD); subjects without any VGDF in their working area (n=773), i.e. 12 were excluded due to asthma, 99 with CB, and 20 with COPD. The total number of selected population was 1,375 people.
Definition of exposures

Data on the smoking status and occupational exposures were collected from all study participants. Subjects who had smoked a minimum of 100 cigarettes since they had started smoking were regarded as current smokers. Subjects who had smoked a minimum of 100 cigarettes in their lifetime but did not smoke at the time of the study were regarded as former smokers. Subjects who had never smoked or smoked less than 100 cigarettes were regarded as never smokers (14).

Workers who were regarded as both current and former smokers were included in the calculation of the smoking index. The average time elapsed since the former smokers quit smoking was 3.1 years.

Occupational exposure to VGDF was categorised into three groups: (1) silica dust, i.e. dust containing 10 or more percent of silica; (2) nonfibrogenic dusts; and (3) nonfibrogenic dust simultaneously with vapours of irritants and sensitizers. Occupational exposure limits (OEL) for each kind of VGDF were established by the Russian Federation regulations.

Definition of outcomes

We used common definitions for COPD/CB, used in the last GOLD revision (15). Therefore, we considered COPD as “a common preventable and treatable disease, characterised by persistent airflow limitation that is usually progressive and associated with an enhanced chronic inflammatory response in the airways and the lung to noxious particles or gases”. We confirmed the presence of persistent airflow limitation using spirometry. Subjects with forced expiratory volume in 1 second and forced vital capacity ratio (FEV1/FVC) value of less than 0.7 were regarded as COPD patients. The stages of COPD were also determined according to GOLD criteria.

To evaluate chronic bronchitis, we used the GOLD (2011) diagnosis criteria: the presence of cough and sputum production for at least three months in each of two consecutive years (15).

Therefore, there was no overlapping between COPD and CB groups.

Questionnaires

We used the modified European Society for Coal and Steel (ECSC) questionnaire (16) for the pre-assessment of physical health status. We determined the degree of tobacco dependence using the modified Fagerstrom test (17). Both instruments were translated into Russian.

Spirometric measurements

Physical examination and baseline spirometry were performed for all the participants of the study. We used a portable computerised spirometer (Spirolab III, MIR, Italy). All spirometric measurements were performed by only one of the authors (MNN), respecting the American Thoracic Society (ATS)/European Respiratory Society (ERS) guidelines (18). We calculated the FEV1/FVC and the percentage of the predicted values for FVC (FVC % predicted) and FEV1 (FEV1 % predicted) using published reference values for Europeans (19).

Study participants with value of FEV1/FVC of less than 0.7 were examined with post-bronchodilator test, which was performed according to the ATS / ERS guidelines (18) 15 minutes after the administration of 400 micrograms of salbutamol. The increase in FEV1 by more than 15 % (or 200 mL) from baseline was regarded as reversible obstruction.

Occupational exposure assessment

We evaluated occupational exposures in two ways. First, we assessed the workers’ lifetime occupational exposures to VGDF directly by self-reports during the periodical medical examination. Subjects who answered affirmatively to at least one of the following two questions i.e.: “Have you ever worked for five years or more in any dusty job?” and/or “Have you ever been exposed (for five years or more) to gas or chemical fumes in your work?” were regarded as exposed to VGDF.

Second, we collected the data on current occupation, occupational history, and occupational exposure for all the study participants (N=1,375): the employers provided us with all these information from the workplace certification cards, established by the Russian Federation regulations.

We combined different types of VGDF into three groups, as was described above, due to a large variety of occupational hazards in the foundry plants, where 15 to 36 various chemical substances could simultaneously be present in the working area. Occupational exposure to VGDF in the aircraft and oil extracting enterprises, i.e. nonfibrogenic dusts or organic solvent vapours in values of less than OEL, were present at a much smaller number of workplaces.

We took into account the influence of occupational exposures other than VGDF. We considered heating microclimate and excessive physical activity as two
additional risk factors for occupational respiratory
diseases because of their effects on the breathing
rate. The information on the presence of overheating
and excessive physical activity in the workplace
were also drawn from workplace certification
cards.

Occupational Exposure Limit values of VGDF
were accepted according to the National hygienic
standards.

Data Analysis

Analyses were conducted using the R statistical
system (version 2.11.1) software (20). Prevalence of
COPD and CB was estimated for the entire cohort by
age, gender, education level, smoking status, and
occupational exposures. The difference between
variables was evaluated by Student’s t-test for
continuous data and chi-square test for categorical
data. The main epidemiological criteria traditionally
used as measures of association, i.e. odds ratio (OR),
attributable risk (AR), and population attributable risk
(PAR) were also calculated. The relationship between
the influence of smoking and occupational factors for
COPD development was analysed by Cochran-
Mantel-Haenszel test (CMH-test) of data stratification,
with smoking being an additional affecting factor.

The logistic model of regression analysis was used
for the same raw data to evaluate the combined effect
of occupational exposures and smoking. Generalised
additive model (GAM) (21) with binomial family was
used to model the COPD probability; the model can
include interactions between independent variables.
The nonlinear dependence on smoking index was
modeled using the smoothing term (22). “MGCV”
package for R was used to fit the model.

To model the response with ordered values,
Ordered Logistic Regression (23) was used. Model
was fitted using the MASS R package to estimate

Table 1 General characteristics of the studied cohort

<table>
<thead>
<tr>
<th></th>
<th>COPD (n=105)</th>
<th>CB (n=170)</th>
<th>Reference group (n=1100)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age / year, mean ± SD</td>
<td>51.7±8.4 †</td>
<td>49.5±8.1 †</td>
<td>47.5±9.9</td>
</tr>
<tr>
<td>Women, n (%)</td>
<td>3 (2.9) †</td>
<td>46 (26.7) †</td>
<td>447 (40.6)</td>
</tr>
<tr>
<td>Education level, n (%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Secondary</td>
<td>99 (94.3)*</td>
<td>152 (89.4) †</td>
<td>930 (84.5)</td>
</tr>
<tr>
<td>Higher</td>
<td>6 (5.7) †</td>
<td>18 (10.6)*</td>
<td>170 (15.5)</td>
</tr>
<tr>
<td>Level of income, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Low</td>
<td>16 (15.2) †</td>
<td>13 (7.6)</td>
<td>60 (5.5)</td>
</tr>
<tr>
<td>Middle</td>
<td>83 (79.0)</td>
<td>103 (60.6)*</td>
<td>910 (82.7)</td>
</tr>
<tr>
<td>High</td>
<td>6 (5.7)*</td>
<td>54 (31.8)*</td>
<td>130 (11.8)</td>
</tr>
<tr>
<td>Smoking status, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Nonsmokers</td>
<td>20 (19.1) †</td>
<td>43 (25.3) †</td>
<td>713 (64.8)</td>
</tr>
<tr>
<td>Smokers</td>
<td>75 (71.4) †</td>
<td>113 (66.5) †</td>
<td>292 (26.5)</td>
</tr>
<tr>
<td>Former smokers</td>
<td>10 (9.5)</td>
<td>14 (8.2)</td>
<td>95 (8.6)</td>
</tr>
<tr>
<td>Smoking index / pack-years, n (%):</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Less than 20</td>
<td>55 (33.3) †</td>
<td>60 (35.3) †</td>
<td>231 (21.0)</td>
</tr>
<tr>
<td>20 or more</td>
<td>50 (47.6) †</td>
<td>67 (39.4) *</td>
<td>156 (14.2)</td>
</tr>
<tr>
<td>Occupational exposure to VGDF, n (%)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Low (&lt;OEL to 3.0 OEL)</td>
<td>85 (80.9) †</td>
<td>71 (41.8)</td>
<td>458 (41.6)</td>
</tr>
<tr>
<td>Medium (3.1 OEL to 6 OEL)</td>
<td>46 (43.8) †</td>
<td>35 (20.6) †</td>
<td>106 (9.6)</td>
</tr>
<tr>
<td>High (&gt;6.0 OEL)</td>
<td>21 (20) †</td>
<td>0</td>
<td>27 (2.5)</td>
</tr>
</tbody>
</table>

The difference between variables was evaluated by Student’s t-test for continuous data and chi-square test for categorical data.
Marked differences (* p<0.01, † - p<0.05, ‡ - p<0.001) between COPD and/or CB patients and control group.

COPD – chronic obstructive pulmonary disease
CB – chronic bronchitis
VGDF – vapour, gases, dust and fumes
OEL – occupational exposure limit
confidence intervals. We built seven predictive models using the same initial data. As a dependent variable, relation to one of the following three groups of workers was taken into account: “healthy”, “CB”, and “COPD”. The variables with discrete values were presented as the factors with fixed rates. Contributions of 14 factors in the model was estimated, i.e. sex, age, education, and income levels, VGDF levels in the working area, presence of other occupational factors, i.e. physical exertion, overheating, vibration, and noise. Besides, period of work, smoking status, smoking history, smoking index value, and the degree of nicotine dependence were estimated too. Contribution of each factor was evaluated separately. Significance of the factors was determined against the baseline, which was assumed to be the most important factor value in an importance order. The contribution of the baseline was considered as the zero value of the model to be defined.

The predictive capacity of models was evaluated by cross-validation and Receiver Operating Characteristic (ROC)-analysis. The repetitive cross-validation (bootstrap) was used: data was split into teaching (80 % of data) and validation (20 % of data) subsets, model was fitted, and the predictions on the validation subset were compared with the true values. This procedure was repeated 100 times to estimate the histograms and confidence intervals; these results were used to select the best predictive model. ROC was estimated and plotted using the ROCR package (24).

RESULTS

In total, 105 cases of COPD and 170 cases of chronic bronchitis (CB) have been diagnosed in the cohort of workers examined (Table 1). Thus, prevalence of COPD was 7.5 % for the entire cohort and 7.9 % for people older than 40; prevalence of CB was 12.1 % for the entire cohort and 13.4 % for the people over 40. Distribution of COPD patients across stages according to the GOLD criteria was as follows: stage I - 69 people (65.7 %); stage II - 29 people (27.6 %), stage III - 7 people (6.7 %). Proportion of smokers was significantly lower in healthy workers than among COPD and CB patients (p<0.05). The highest proportion of smokers was found among patients with COPD. Occupational exposure to VGDF occurred twice more often among COPD patients than among both the CB patients and the control group of healthy workers (p<0.05). More than 40 % of COPD patients were occupationally exposed to VGDF on the level of 3.0 OEL and more. The 6.0 OEL and higher levels were found in more than 20 % of COPD patients.

A strong link between exposure to VGDF at the workplace and COPD development was found (Table 2). Thus, the overall odds ratio for COPD development due to occupational VGDF exposure was 5.9 (95 % CI=3.6 to 9.8; p=0.0001).

The values of odds ratios for different types of occupational noxious particles and gases among COPD patients were highest for silica dust (OR=6.2; 95 % CI=3.6 to 10.7; p<0.0001). The same refers to other indicators of risk assessment, e.g. PAR % values for COPD patients. The difference between PAR % values for silica dust and other kinds of dust were not statistically significant (56.7 % and 49.8 %, p>0.05). The lowest value of PAR % for COPD patients was found for occupational exposure to dust with vapours of irritants and sensitizers (18.3 %). The overall value of PAR % obtained in our study was 65.3 %. This is more than three times higher than the value demonstrated in ATS statements (2003, 2010) (15, 16) dedicated to occupational burden of lung diseases.

Another kind of dependence was found for CB patients. The overall odds ratio for CB development was statistically non-significant (OR=1.0; 95 % CI=0.7 to 1.4; p=0.4). However, the statistically significant value of odds ratio for CB development was obtained for VGDF levels above 3.0 OEL. The highest value of odds ratio was reached for CB patients who had occupational exposure to dust with vapours of irritants and sensitizers (OR=2.0; 95 % CI=1.0 to 4.1; p=0.07) but not for silica dust.

Occupational risk assessment calculated separately for nonsmoking and smoking workers showed that the odds ratios for occupational COPD were significantly higher for nonsmokers. For the smoking workers, smoking was a major risk factor for COPD development (Table 3). Thus, the odds ratio for occupational COPD for non-smokers was almost seven times higher than that of smokers and their attributable risk for occupational COPD was more than 95 % compared to 65 % for smokers. Total risk of COPD development for smoking workers due to the influence of both major risk factors for COPD, i.e. smoking and VGDF, was also calculated. As Table 3 shows, the overall value was about four times higher than the risk for non-smoking workers. It is rather interesting that the risk
of COPD from occupational VGDF exposures established for non-smoking workers was comparable with that obtained for smokers who were not in contact with VGDF in the working area.

As regards the second hypothesis, there is a quite clear and statistically significant "dose - effect" dependence between the level of occupational exposures and COPD development (Table 2). The risk assessment for occupational COPD revealed a regular and statistically significant risk increase when working conditions deteriorated.

Cochran-Mantel-Haenszel test of data stratification with smoking as an additional affecting factor showed the relationship between the influence of smoking and occupational factors. Higher values of odds ratios of COPD development (see above) were obtained for non-smoking workers (Table 4). The difference caused by the smoking status was statistically significant for workers with occupational exposure to silica dust and both to dust and irritants (p=0.032 and 0.012, respectively). No significant difference between smokers and nonsmokers was found for

**Table 2** The risk of COPD/CB development from occupational exposures to VGDF, depending on their type and levels

<table>
<thead>
<tr>
<th>Exposure Type</th>
<th>OR (95% CI)</th>
<th>p-level</th>
<th>AR / %</th>
<th>PAR / %</th>
</tr>
</thead>
<tbody>
<tr>
<td>COPD Overall</td>
<td>5.9 (3.6 to 9.8)</td>
<td>0.0001</td>
<td>80.7</td>
<td>65.3</td>
</tr>
<tr>
<td>Low levels of VGDF *</td>
<td>1.7 (0.9 to 3.4)</td>
<td>0.07</td>
<td>42.4</td>
<td>20.1</td>
</tr>
<tr>
<td>Medium levels of VGDF †</td>
<td>13.9 (7.9 to 24.5)</td>
<td>&lt;0.0001</td>
<td>90.0</td>
<td>62.7</td>
</tr>
<tr>
<td>High levels of VGDF ‡</td>
<td>24.9 (12.1 to 51.5)</td>
<td>&lt;0.0001</td>
<td>93.1</td>
<td>47.7</td>
</tr>
<tr>
<td>Silica dust, i.e. dust containing 10 % or &gt;10 % silica</td>
<td>6.2 (3.6 to 10.7)</td>
<td>&lt;0.0001</td>
<td>81.3</td>
<td>56.7</td>
</tr>
<tr>
<td>Nonfibrogenic dusts</td>
<td>5.7 (3.2 to 10.2)</td>
<td>&lt;0.0001</td>
<td>80.0</td>
<td>49.8</td>
</tr>
<tr>
<td>Nonfibrogenic dust simultaneously with vapours of irritants and sensitizers</td>
<td>5.5 (2.1 to 14.6)</td>
<td>0.002</td>
<td>79.3</td>
<td>18.3</td>
</tr>
<tr>
<td>CB Contact with VGDF, including:</td>
<td>1.0 (0.7 to 1.4)</td>
<td>0.4</td>
<td>0.45</td>
<td>0.19</td>
</tr>
<tr>
<td>Low levels of VGDF *</td>
<td>0.7 (0.5 to 1.1)</td>
<td>0.1</td>
<td>&lt;0</td>
<td>&lt;0</td>
</tr>
<tr>
<td>Medium levels of VGDF †</td>
<td>2.1 (1.4 to 3.3)</td>
<td>0.0005</td>
<td>45.4</td>
<td>11.4</td>
</tr>
<tr>
<td>High levels of VGDF ‡</td>
<td>No data</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Silica dust, i.e. dust containing 10 % or &gt;10 % silica</td>
<td>1.0 (0.7 to 1.5)</td>
<td>0.8</td>
<td>25</td>
<td>0.34</td>
</tr>
<tr>
<td>Nonfibrogenic dusts</td>
<td>0.8 (0.5 to 1.3)</td>
<td>0.4</td>
<td>&lt;0</td>
<td>&lt;0</td>
</tr>
<tr>
<td>Nonfibrogenic dust simultaneously with vapours of irritants and sensitizers</td>
<td>2.0 (1.0 to 4.1)</td>
<td>0.07</td>
<td>48.3</td>
<td>11.6</td>
</tr>
</tbody>
</table>

Levels of VGDF in the working area: *<OEL to 3.0 OEL; †from 3.1 OEL to 6 OEL, ‡>6.0 OEL.

COPD – chronic obstructive pulmonary disease
CB – chronic bronchitis
VGDF – vapour, gases, dust and fumes
OEL – occupational exposure limit

**Table 3** The risk values of chronic obstructive pulmonary disease (COPD) development among dusty trade workers and smokers not having any occupational exposures

<table>
<thead>
<tr>
<th>Exposure Type</th>
<th>OR (95% CI)</th>
<th>p-level</th>
<th>AR / %</th>
<th>PAR / %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Nonsmoking workers in dusty trades</td>
<td>22.2 (4.9 to 100.5)</td>
<td>&lt;0.0001</td>
<td>95.1</td>
<td>81.5</td>
</tr>
<tr>
<td>Smoking workers in dusty trades (risk values due to occupational exposures only)</td>
<td>3.4 (1.8 to 6.5)</td>
<td>0.0001</td>
<td>65.3</td>
<td>42.5</td>
</tr>
<tr>
<td>Smoking workers in dusty trades (overall risk from occupational and non-occupational exposures)</td>
<td>82.7 (19.9 to 342.3)</td>
<td>&lt;0.0001</td>
<td>98.3</td>
<td>95.2</td>
</tr>
<tr>
<td>Smokers not having any VGDF in the working area (risk value from smoking)</td>
<td>34.7 (7.9 to 151.5)</td>
<td>&lt;0.0001</td>
<td>96.7</td>
<td>87.0</td>
</tr>
</tbody>
</table>

OR – odds ratio
CI – confidence internal
AR – attributable risk
PAR – population attributable risk
VGDF – vapour, gases, dust and fumes
the workers who were occupationally exposed to other kinds of dust different from silica.

The logistic regression model investigated the combined effect of smoking and occupational exposures on COPD development using the same raw data. As shown in Table 5, the following factors specified in order of their significance were important for COPD: VGDF composition ($\chi^2=64.4; p<0.0001$), VGDF level ($\chi^2=63.3; p<0.0001$), smoking index ($\chi^2=43.2; p<0.0001$), age ($\chi^2=5.7; p=0.01$), and heating microclimate in the workplace ($\chi^2=5.2; p=0.02$). The contribution of all factors to the model was linear except for the smoking index.

Visual analysis of the curve of the influence of smoking index in the model of occupational COPD showed that its nonlinear dependence could be described by two linear curves: the effect of smoking index on the development of COPD in the model increased almost linearly from 0 to about 20 pack-years, and then it remained practically unchanged (Figure 2). For this reason, a new variable was introduced in the model with values of smoking index limited to the level of 20 pack-years. Assuming linear contribution of the smoking index in the model, the VGDF composition becomes insignificant for COPD development because of the counterbalancing effect of the smoking index and VGDF levels. In addition, the statistically significant interaction between two main predictors of occupational COPD (smoking index and VGDF levels) was found.

Analysis of the values of linear predictors in this model showed that the effect of smoking on COPD development in the cohort of workers studied ranged from 5 % to 40 % depending on other risk factors. The contribution of VGDF to COPD likelihood increased monotonously with the worsening of working conditions, with no changing in the likelihood of the COPD development, and was found statistically absolutely significant ($p<0.0001$). The VGDF impact was within 3 % to 5.2 % at low VGDF levels (<3.0 OEL), 33 % at middle levels (from 3.0 OEL to 6.0 OEL), and reached 44 % at high levels (>6.0 OEL).

Therefore, the degrees of smoking and VGDF influence on COPD development are probably rather similar. This suggestion was verified using seven predictive models with the same initial data. Ten from 14 studied occupational and non-occupational factors proved to be statistically significant in the models, i.e. age, gender, VGDF levels, type of VGDF, levels of irritants, physical stress, heating microclimate, smoking status, smoking index, and period of smoking. VGDF levels and the smoking index demonstrated maximal influence on CB/COPD probability. Age, gender, and levels of irritants exerted minimal but statistically significant influence.

Overall, average results of cross-validation showed high levels of reliability of predictiveness of the models on new data. The best results of cross-validation and ROC-analysis were obtained for two models, i.e. three-level generalised linear model (GLM) “(0) healthy workers & CB, (1) COPD”, and two-level GLM-model “(0) healthy workers, (1) COPD” (Figures 3, 4).

**DISCUSSION**

Prevalence of COPD and CB obtained in this study correspond to average data in the Russian Federation and internationally (4).

High values for odds ratios of COPD development obtained in our study indicate a high level of the occupationally determined risk for COPD. This allows discussing possible inclusion of the disease in the National List of Occupational Diseases. However, a well-known statement “Chronic bronchitis is the biggest single cause of sickness absence” (25) also explains the impressive difference between COPD and CB odds ratio values shown in this study. A substantial difference between OR, AR, and PAR % values for COPD and CB call for the possible update of the classification of occupational airways diseases.
“Dose - effect” dependence between VGDF levels and COPD development (see Table 2), is another argument in favour of the possibility of COPD being developed due to VGDF only.

The odds ratio and other risk assessment criteria of COPD development for the silica dust exposed workers obtained here confirm again the importance of this risk factor (26). Insignificant difference between PAR % values for silica dust and other kinds of dust indicates the importance of all kinds of occupational noxious particles for COPD development. Further research is required for the comparative impact of occupational factors to be evaluated.

As Table 2 shows, the values of PAR % for workers exposed to low VGDF levels (<3.0 OEL) are very close to the results obtained in other studies of occupational COPD published earlier (27-29). However, our study showed that the values of risk assessment were much higher for the workers exposed to higher VGDF levels. This might result from much poorer working conditions at the foundry plants in the Russian Federation, and relatively high VGDF levels in the working area of dusty trade workers included in the cohort studied. Therefore, the effect of VGDF should be evaluated according to the levels in the working area. This may be useful for future strategies of occupational risk management.

We have also found that non-smoking dusty trade workers run the risk of developing occupational COPD six times more than smoking dusty trade workers (see Table 3). This is similar to the results described earlier by Becklake (30) in her classic epidemiological study of the prevalence of occupational COPD.
Data stratification performed by Cochran-Mantel-Haenszel test in our study showed again that non-smoking workers have higher probability of developing the disease (see Table 4) with the isolated impact of occupational risk factors for COPD. The difference due to the smoking status in the cohort studied was statistically significant for both silica dust and for dusts and irritants present simultaneously in the working area (p=0.032 and 0.012, respectively). For other types of dust, except quartz, no significant difference was found between smokers and nonsmokers. This points to a more important role of silica dust and irritants in the development of COPD.

It should be pointed out that smoking workers are influenced by both major risk factors for COPD. For non-smoking dusty trade workers, the risk of COPD is caused only by VGDF. Therefore, we evaluated the overall risk for COPD development from occupational and non-occupational exposures and the result was that the overall risk for workers influenced by both major risk factors for COPD is about four times higher than that caused by VGDF only. The studies performed by Meer et al. (31) and Blanc et al. (32) showed similar results. This makes us think about the necessity of planning preventive strategies and developing smoking cessation programs for dusty trade workers.

Similar results were obtained in our study for COPD risk due to occupational VGDF exposures for non-smoking dusty trade workers and for smokers who did not have contact with VGDF in the working area. This confirms again the well-known thesis of the two main risk factors for COPD, i.e., smoking and noxious particles and gases, whose influence seems comparable.

Analysis of the combined effect of tobacco smoking and occupational noxious particles and gases for COPD development showed the following order of risk factors based on the strength of their influence: VGDF levels, smoking index, age, and heating microclimate. The values of impact of VGDF and smoking for COPD development are similar in the regression model used.

The results of cross-validation and ROC-analysis revealed the highest level of sensitivity and specificity for two models made. The two-level GLM-model “(0) healthy workers; (1) COPD” showed the best result of cross-validation. However, the realisation of the method assuming the artificial exclusion of CB patients from the database does not allow the model for prediction to be applied.

### Table 4  *The relationship between smoking and occupational factors in the development of chronic obstructive pulmonary disease (COPD)*

<table>
<thead>
<tr>
<th></th>
<th>OR (95% CI)</th>
<th>CMH-test for homogeneity, χ²</th>
<th>p-level</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Smoking workers</td>
<td>Non-smoking workers</td>
<td></td>
</tr>
<tr>
<td>Silica dust, i.e. dust containing 10% or &gt;10% silica</td>
<td>3.5 (1.8 to 6.8)</td>
<td>20.6 (4.5 to 190.7)</td>
<td>4.61</td>
</tr>
<tr>
<td>Non-fibrogenic dusts</td>
<td>3.7 (1.9 to 7.4)</td>
<td>9.76 (1.4 to 109.4)</td>
<td>1.08</td>
</tr>
<tr>
<td>Non-fibrogenic dust simultaneously with vapours of irritants and sensitizers</td>
<td>1.75 (0.4 to 5.8)</td>
<td>32.1 (1.0 to 7.0)</td>
<td>6.28</td>
</tr>
</tbody>
</table>

CMH-test = Cochran-Mantel-Haenszel test

### Table 5  *The significance of risk factors to the development of chronic obstructive pulmonary disease (COPD) in a model of multiple logistic regression*

<table>
<thead>
<tr>
<th></th>
<th>With initial values of smoking index</th>
<th>With transformed values of smoking index</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>χ²</td>
<td>p</td>
</tr>
<tr>
<td>Type of VGDF</td>
<td>64.5</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Level of VGDF</td>
<td>63.3</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Smoking index</td>
<td>43.2</td>
<td>&lt;0.0001</td>
</tr>
<tr>
<td>Age</td>
<td>5.7</td>
<td>&lt;0.0200</td>
</tr>
<tr>
<td>Heating microclimate</td>
<td>5.2</td>
<td>&lt;0.0230</td>
</tr>
</tbody>
</table>

VGDF – vapour, gases, dust and fumes
Nevertheless, high analytical capacity of this model provides the application of its predictors as the assessment criteria for medical examination of workers. The three-level GLM-model “(0) healthy workers & CB; (1) COPD” showed the second best results of cross-validation. All of the results of cross-validation and ROC-analysis fully support the conclusion made.

CONCLUSIONS

The differentiation between the various effects of inhaled noxious particles and gases seems to be complex. But this task is particularly challenging for the Russian Federation because of a high number of smokers in the country. Therefore, a comprehensive assessment of contribution of the two leading risk factors for COPD is of special interest for occupational medicine. Future investigations of occupational COPD seem to be important for developing prevention strategies.

REFERENCES


Sažetak

KRONIČNA OPSTRUKTIVNA PLUĆNA BOLEST I RAD: RETROSPEKTIVA ISTRAŽIVANJA PROVEDENOG NA KOHORTI INDUSTRIJSKIH RADNIKA

Cilj ovog rada bio je potvrditi prevalenciju kronične opstruktivne plućne bolesti (engl. chronic obstructive pulmonary disease, COPD) među industrijskim radnici u Ruskoj Federaciji i utvrditi relativni doprinos pušenja i profesionalnih čimbenika razvoju COPD-a.

Odabrali smo 1.375 radnika u dobi od 30 godina i starijih. Šeststo dvadeset i četiri radnika bila su izložena parama, plinovima, prašini i dimovima (engl. vapours, gases, dust, and fumes – VGDF) na radu. Svi su radnici bili podvrgnuti fizikalnom pregledu i temeljnoj spirometriji. Za radnike koji su imali smanjen protok zraka (FEV1/FVC<0,70) smatralo se da imaju COPD, a za one koji su imali kašalj i pojačan sputum barem 3 mjeseca tijekom dvije uzastopne godine smatralo se da bolju od kroničnog bronhitisa (CB). Izmjenu ovdje dvije skupine nije bilo preklapanja. Od svih su sudionika prikupljeni podaci o radnoj anamnezi i razinama izloženosti VGDF-u na radu. Profesionalna izloženost VGDF-u bila je dvostruko češće prisutna kod radnika s COPD-om nego kod radnika s CB-om i kod kontrolne skupine zdravih radnika (p<0,05). Više od 40 % bolesnika s COPD-om bilo je profesionalno izloženo VGDF-u iznad 3,0 OEL (engl. occupational exposure limit – granica profesionalne izloženosti), a više od 20 % granici od 6,0 OEL i više. Ukupni omjer izgleda (engl. odds ratio – OR) za razvoj KOPB-a zbog izloženosti VGDF-u bio je 5,9 (95 %-tni CI=3,6 do 9,8; p=0,0001). Čini se da su i pušenje i VGDF važni za razvoj COPD-a. Analiza sjedinjenog učinka pušenja i profesionalne izloženosti štetnim česticama i plinovima na razvoj COPD-a pokazala je ovakav redoslijed čimbenika rizika prema jačini njihove utjecaja: razine VGDF-a, indeks pušenja, i profesionalne izloženosti štetnim česticama i plinovima na razvoj COPD-a.

KLJUČNE RIJEČI: COPD, filterska prašina, procjena rizika, profesionalna izloženost, pušenje

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