Presence of acute phase response in coal workers' pneumoconiosis

G Fernandez Rego, G Ocio Achaerandio, V Gonzalez Cuervo, C Rodriguez Menendez, C Martinez Gonzalez, C Alvarez Alvarez

Abstract
To evaluate the role of personal factors in pneumoconiosis, several acute phase proteins were studied in 62 coal miners without acute illnesses and classified as having no pneumoconiosis (n = 19), simple pneumoconiosis (n = 23), or complicated pneumoconiosis with progressive massive fibrosis (n = 20). Groups were similar for age, years of work at high risk jobs, chronic bronchitis, and forced expiratory volume in one second (FEV1). C-reactive protein concentration was significantly higher in the simple and complicated pneumoconiosis groups in comparison with the no pneumoconiosis group. The C-reactive protein concentration was above the upper normal value in 12 (27.9%) out of 43 cases with simple and complicated pneumoconiosis. On the other hand only one case of no pneumoconiosis was above the upper normal range (5.3%), a significant difference taking into account a stratified analysis for chronic bronchitis. Fibrinogen concentration was significantly increased in the simple pneumoconiosis group compared with the no pneumoconiosis group. The value of fibrinogen was above the upper normal limit in 17 out of the 43 cases with pneumoconiosis (39.5%) by contrast with two cases in the no pneumoconiosis group (10.5%). No significant differences in a1-antitrypsin and ceruloplasmin concentrations were found between groups. In conclusion, an alteration in some acute phase proteins related to pneumoconiosis was found in miners. This could be used as a marker of disease activity and personal response against the pathogenic agent.

It is generally agreed that quantitative exposure to coal dust is the most important factor in the development of pneumoconiosis. The greater the exposure to coal dust, the greater the probability of contracting the disease. It should, therefore, be possible to predict the number of cases of pneumoconiosis in a group of exposed workers to within a certain degree of accuracy.

For individual cases, however, owing to several personal factors, precise prediction is not yet possible. Some of these personal factors have been discussed in previous reports. Rasche et al found an increase in C-reactive protein (C-RP) in a group of coal miners considered as predisposed to pneumoconiosis. Nevertheless, in an earlier report, they had concluded that the increase in C-RP in miners was due to the presence of chronic bronchitis and not to pneumoconiosis. An increase in a1-antitrypsin (a1-AT) was found in miners with progressive massive fibrosis compared with controls by Hahon et al. Others, however, report no differences.

Increases in bronchoalveolar lavage (BAL) cellularity and gallium-67 uptake have been found in pneumoconiosis. An active process, similar in some aspects to an inflammation, might be the cause. If this is the case, an alteration in acute phase reaction might be expected. To determine if this were so, C-RP, fibrinogen, ceruloplasmin, and a1-AT were measured in coal miners exposed to similar occupational risk and suffering different types of pneumoconiosis.

Materials and methods
This study included 62 consecutive coal miners, (minimum age 50) sent to the clinic for periodic check ups between January and June 1988. They had no acute illnesses, and were not taking steroids or other anti-inflammatory drugs.

Subjects were interviewed by trained personnel. Case histories were completed and details about working conditions were obtained. To evaluate exposure risk, the factor considered was number of years of work in high risk jobs (face workers and drillers). The definition of ex-smoker was met if
smoking had stopped three months before the test. Chronic bronchitis was defined in the standard way.\(^9\) Chest radiographs were obtained and were read by three trained experts using the 1980 International Labour Office classification.\(^10\) Three groups were established—namely, no pneumoconiosis (profusion less than 1/1), simple pneumoconiosis, and complicated pneumoconiosis with progressive massive fibrosis. The simple and complicated pneumoconiosis groups form the pneumoconiosis group. Spirometric studies were performed using a Vitalograph spirometer. At least three trials were completed and the results were selected using standard criteria.\(^11\) Table 1 summarises the group characteristics. Blood samples were analysed by the following methods: C-reactive protein by an immunoturbidimetric test using goat antibodies (Boehringer, Mannheim); fibrinogen by the Clauss method;\(^12\) ceruloplasmin by radial immunodiffusion (Mancini, Behring Institute); and \(\alpha_1\)-antitrypsin by radial immunodiffusion (Mancini, Bio-Merieux).

Differences between means of the groups were compared by two tailed \(t\) test. For C-RP the Mann-Whitney \(U\) test was employed. A stratified analysis for chronic bronchitis (Mantel-Haenszel test) was applied to compare the number of C-RP and fibrinogen values above the upper normal limit in the pneumoconiosis and no pneumoconiosis groups. Proportions of abnormally high values in these two groups were compared with an appropriate test for unpaired data.

### Table 1 Groups according to pneumoconiosis

<table>
<thead>
<tr>
<th></th>
<th>No pneumoconiosis</th>
<th>Simple pneumoconiosis</th>
<th>Complicated pneumoconiosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>No of cases</td>
<td>19</td>
<td>23</td>
<td>20</td>
</tr>
<tr>
<td>Mean age, (y (range))</td>
<td>62.9 (51-74)</td>
<td>61.5 (51-75)</td>
<td>62.2 (52-72)</td>
</tr>
<tr>
<td>Mean risk, (y (range))</td>
<td>17.3 (2-39)</td>
<td>17.4 (2-34)</td>
<td>18.4 (2-35)</td>
</tr>
<tr>
<td>Smokers</td>
<td>14</td>
<td>15</td>
<td>14</td>
</tr>
<tr>
<td>Ex-smokers</td>
<td>10</td>
<td>8</td>
<td>12</td>
</tr>
<tr>
<td>Chronic bronchitis</td>
<td>8</td>
<td>7</td>
<td>10</td>
</tr>
<tr>
<td>FEV(_1), (ml (SD))</td>
<td>2182 (634)</td>
<td>2388 (581)</td>
<td>1801 (593)</td>
</tr>
</tbody>
</table>

### Table 2 Acute phase proteins according to pneumoconiosis

<table>
<thead>
<tr>
<th>Protein</th>
<th>No pneumoconiosis</th>
<th>Simple pneumoconiosis</th>
<th>Complicated pneumoconiosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>C-Reactive protein</td>
<td>0.23 (0.34)(\dagger)</td>
<td>0.57 (0.53)*</td>
<td>1.6 (3.5)*</td>
</tr>
<tr>
<td>Fibrinogen</td>
<td>314.5 (82.3)</td>
<td>375 (101.7)*</td>
<td>371 (123.5)</td>
</tr>
<tr>
<td>Ceruloplasmin</td>
<td>84.8 (113.4)</td>
<td>65.8 (65.6)</td>
<td>52.1 (9.4)</td>
</tr>
<tr>
<td>(\alpha_1)-Antitrypsin</td>
<td>191.2 (33)</td>
<td>193.7 (35.3)</td>
<td>213.7 (93.9)</td>
</tr>
</tbody>
</table>

\(\dagger\)p < 0.05 v no pneumoconiosis.

Mean mg protein/dl (SD).

Results

Table 2 summarises the results. The concentration of C-RP was significantly increased in the simple and complicated pneumoconiosis groups compared with the no pneumoconiosis group (\(p < 0.02\)). The value of C-RP was above the upper normal value in 12 out of 43 cases with pneumoconiosis (27.9%). On the other hand, only one case in the no pneumoconiosis group was above the upper normal value (5.9%), a significant difference taking into account the stratified analysis for chronic bronchitis (\(x = 1.99; p < 0.05\)).

Fibrinogen concentration was significantly greater in the simple pneumoconiosis group in comparison with the no pneumoconiosis group (\(t = 2.08; p < 0.05\)). The value of fibrinogen was above the upper normal range in 17 cases with pneumoconiosis (39.5%) by contrast with two cases in the no pneumoconiosis group (10.5%) (\(x = 2.29; p < 0.05\)).

The number of cases with abnormally high values for ceruloplasmin and \(\alpha_1\)-AT were nine (20%) and four (10%) in the pneumoconiosis group and 4 (21%)
and one (5%) in the no pneumoconiosis group. These differences were not significant.

The figure shows the percentages of cases with values above the upper normal limit in the pneumoconiosis and no pneumoconiosis groups.

**Discussion**

In the present study, we found a non-uniform alteration in acute phase proteins in the miners studied. Fibrinogen and C-RP concentrations were increased in relation to the presence of pneumoconiosis. Concentrations of ceruloplasmin, an important serum antioxidant, were increased in some cases, but had no relation to pneumoconiosis; α₁-AT values were not significantly altered.

Other studies have examined several of these proteins with conflicting results. Hahon et al. found an increase in α₁-AT concentration in miners with progressive massive fibrosis compared with controls. These differences were not found, however, in other studies. Rasche et al. found an increase in C-RP concentration in a group of coal miners considered as predisposed to pneumoconiosis by contrast with an earlier report in which they had concluded that the increase in C-RP in miners was due to the presence of chronic bronchitis and not to pneumoconiosis. This apparent contradiction with our results may be because our cases were studied when they were in a stable situation with no signs of acute illness.

The significance of C-RP measurement for monitoring the acute phase response has recently been emphasised. This response is basically started by interleukin-1, a potent mediator that is considered to be a pathogenic factor in silicosis.

Bronchoalveolar lavage and gallium uptake studies suggest a pneumoconiotic lesion. Changes in acute phase proteins might reflect the same process at the serum level and may be useful as markers of disease activity. On the other hand the acute phase reaction could be used as a marker of individual reactivity against the pathogenic agent. Other study designs would be necessary to check this hypothesis.

In conclusion, an alteration in acute phase proteins was related to pneumoconiosis in some miners. This could be useful as a marker of disease activity, or individual reactivity against the pathogenic agent, or both.

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