Latent pulmonary involvement in Crohn’s disease: biological, functional, bronchoalveolar lavage and scintigraphic studies

P Bonniere, B Wallaert, A Cortot, X Marchandise, Y Rioü, A B Tunnel, J F Colombel, C Voisin, and J C Paris

From the Clinique des maladies de l’appareil digestif, and Service associé de médecine nucléaire, Cité Hospitalière, Lille; Département de pneumologie, and Service d’explorations fonctionnelles respiratoires, Hôpital Calmette, Lille, France

SUMMARY We have investigated the following pulmonary related parameters in 22 patients with Crohn’s disease who were free of clinical pulmonary symptoms and had normal chest roentgenograms and in 25 controls: serum angiotensin converting enzyme, pulmonary function tests, bronchoalveolar lavage (lymphocyte count and subpopulations, macrophage viability and superoxide anion release by macrophages) and pulmonary scannings. Serum angiotensin converting enzyme was lower in Crohn’s disease (14.1±5.1) than in controls (25.2±4.7) (p<0.001). Twelve of 22 Crohn’s disease (54%) had a bronchoalveolar lavage lymphocytosis (>18% alveolar lymphocytes). Bronchoalveolar lavage lymphocytes subpopulations were quite variable. Twelve of 17 Crohn’s disease (71%) had an increase spontaneous and/or stimulated superoxide anion production by alveolar macrophages. Six of 12 Crohn’s disease (50%) had an increase physiologic dead space in the upper part of their lung against one of 11 controls (9%). These data suggest that most patients with Crohn’s disease have a latent pulmonary involvement.

Systemic manifestations are frequent in inflammatory bowel disease, but classically the lung was regarded as rarely involved.1 2 A few recent reports, however, have described pulmonary manifestations occurring in patients with inflammatory bowel disease: bronchial suppuration, or bronchiectasis,3-5 granulomatous lung disease,6 diffuse or localised interstitial fibrosis7-12 and sulphasalazine pneumonitis.13-21

The rarity of pulmonary clinical manifestations contrasts with striking abnormalities of pulmonary function tests described in patients with inflammatory bowel disease: reduced lung transfer factor, increased residual volume, or decreased forced expiratory volume in one second.22-26

We have attempted to determine the actual frequency of pulmonary tests and bronchoalveolar abnormalities in patients with Crohn’s disease and without clinical or radiological pulmonary abnormalities.27 28 Furthermore, we tried to clarify the pulmonary abnormalities mechanism by carrying out pulmonary scintigraphic studies in those patients.

Methods

PATIENTS Twenty two patients with Crohn’s disease were included in the study. There were 11 women and 11 men, 18–44 years old (mean age 28±7.8 years). Eleven were cigarette smokers. None had a previous history of pulmonary disease or chest radiograph abnormalities. Physical examination and chest radiographs were normal in all patients at the time of the study. The length of Crohn’s disease history at the onset of the study ranged from one month to 22 years (mean duration 41±57 months). Small bowel alone was involved in one case, colon alone in nine cases, and the localisations were multiple in 12 patients (stomach, small bowel, colon, rectum, anus). Eight patients had active anal Crohn’s disease and 10 systemic manifestations: arthralgia (five), ankylosing spondylitis (three), aphthous ulcers (two). Mean Crohn’s Disease Activity Index (CDAI)29 was...
207±86 (50–383). Six patients had quiescent disease (CDAI<150) at the time of the study. Four patients had received sulphasalazine (1·5 g–3 g/day for the previous three months to three years), one prednisolone (0·5 mg/kg body weight for the previous three weeks) and 10 metronidazole (1 g–1·5 g/day for the previous three months).

CONTROL SUBJECTS
The control group included hospital staff and other healthy volunteers. There were 25 controls subjects for bronchoalveolar lavage (13 were cigarette smokers), nine women and 16 men, 24–46 years old (mean age 34±7 years). Thirteen controls were tested for alveolar cell function studies (nine were cigarette smokers), five women and eight men, 22–46 years old (mean age 31±7 years). Eleven controls were tested for pulmonary scintigraphic studies (six were cigarette smokers), three women and eight men, 22–45 years old (mean age 29±6 years).

BRONCHOALVEOLAR LAVAGE
Informed consent was obtained from all patients. Bronchoalveolar lavage was done using a technique which has been previously described. Total number and differential cell count were determined in lavage fluid. Cellular viability of alveolar macrophages was assessed by trypan blue exclusion. The state of activation of alveolar macrophages was evaluated with a lucigenin-dependent chemiluminescence method with or without phorbolmyristate-acetate and with or without superoxide dismutase in 17 Crohn’s disease patients and 13 controls. Results are expressed as relative luminescent units per 0·5×10⁹ viable macrophages.

The proportions of T cells, helper cells and suppressor cells were determined in 10 cases using monoclonal antibodies respectively for OKT₁, OKT₄, and OKT₈ that are specific for the lymphocyte subsets.

A second bronchoalveolar lavage was repeated in five patients with initially active Crohn’s disease six months later their CDAI being <150.

PULMONARY FUNCTION TESTS
These were carried out at rest and included forced vital capacity (FVC) and forced expiratory volume in one second (FEV 1). Residual volume (RV) was quantified with the helium dilution method. Total lung capacity (TLC) was calculated from: TLC=RV+FVC. Carbon monoxide diffusing capacity (DLCO) was studied using a steady state method. The predicted values for each subject based on sex, age, and height were obtained from standard tables. All data were expressed as percentage of the predicted values. Pulmonary function was regarded as abnormal when total lung capacity, forced vital capacity and/or forced expiratory volume in one second were less than 80% of predicted values and/or when carbon monoxide diffusing capacity was less than 75% of that predicted.³²

⁶⁷Gallium (⁶⁷Ga) SCANNING
Each patient received, 72 hours before scanning, 35μ Ci/kg of body weight ⁶⁷Ga citrate intravenously. Accumulation of ⁶⁷Ga in the lung and mediastinum was compared with ⁶⁷Ga uptake in the liver and in the shoulders soft tissue.

PULMONARY SCANNING
Pulmonary scanning included: (1) a ventilatory scanning obtained in a sitting position by respiration of gaseous Xenon (1 mCi/l) which provided a steady state ventilation picture, the total lung height and a recording of lung emptying at one minute. The pulmonary gas retention was estimated by the percentage of gas remaining in the lung at one minute. The distribution of gaseous Xenon was pictured so that a ‘centre of gravity’ or ‘barycentre’ of the lung ventilation could be determined. Radiation exposure was: 8·3 mRad for lungs, 0·09 mRad for whole body. (2) a perfusion scanning after slow intravenous infusion of Technetium albumin macroaggregates in a sitting position (1 mCi). Abnormal distribution of lung perfusion was detected as well as the vertical perfusion gradient from the top to the lung basis. The distribution of macroaggregates was recorded so that a ‘centre of gravity’ of the lung perfusion could be determined. (3) the distance delta(Δ) between the two ‘centre of gravity’ was determined and expressed as percentage of the total lung height.

BLOOD TESTS
Serum angiotensin-converting enzyme assay was carried out in all patients³³ as well as beta-2-microglobulin³⁴ and circulating immune complexes by 121-I-C1q binding activity and by polyethylene glycol precipitation technique.

STATISTICAL METHODS
Results were expressed as mean value (m) ± one standard deviation. The two-tailed t test was used for comparison of data between patients and controls.

Results
All patients were free of clinical pulmonary symptoms and had normal chest radiograph findings.
Latent pulmonary involvement in Crohn's disease

Fig. 1  Cell count (lymphocytes macrophages and neutrophils) in alveolar liquid after bronchoalveolar lavage (BAL) in 25 normal volunteers (○) and 22 Crohn's disease patients (●).

1b BAL data: lymphocytes
Total bronchoalveolar lavage recovery fluid (109±86 ml), total number cells (14·1±5·8×10⁶) were identical in patients and controls according to smoking habits. Alveolar lymphocytes percentages were higher in patients (25·1±19·7%) than in controls (7·2±8%) (p<0·001) (Fig. 1).

Alveolar lymphocytes percentages were above 18% (controls mean + 2 SD) in 12 of 22 patients (54%) who were defined as having an alveolar lymphocytosis. Percentages of T3 lymphocytes were the same in patients (69·7±8·1%) and controls (64±4·5%). T4/T8 ratio were increased in three patients, normal in four and low in three (Table 1).

Five patients had a second bronchoalveolar lavage six months later showing a persistent alveolar lymphocytosis in four cases and a persistent normal bronchoalveolar lavage in one.

1b BAL data: macrophages
Cellular viability of macrophages were identical in patients and controls: 83·7±11·1% and 84·6±8·9%. Spontaneous chemiluminescence of bronchoalveolar lavage cells was higher in patients (4567±4025) than in controls (954±385 relative luminescent units) (p<0·01). Similarly phorbolmyristate-acetate induced chemiluminescence was significantly increased in patients (23824±11257 relative luminescent units) as compared with controls (14392±6213 relative luminescent units) (p<0·025). Values vary widely but were higher than mean±2 SD in 12 of 17 patients with Crohn's disease (70%) (Fig. 2). Superoxide dismutase reduced spontaneous and phorbolmyristate-acetate induced chemiluminescence of alveolar cells respectively by 70% and 91%. The extent of chemiluminescence suppression by superoxide dismutase suggests that superoxide anion is involved in its production.

2 Pulmonary function tests
All but one patient had normal carbonic acid gas and oxygen blood levels (pO2) at rest. Tests were abnormal in 15 of 21 patients. Total lung capacity, forced vital capacity and forced expiratory volume in one second were decreased in two cases associated with low diffusing capacity in one of the two.

Table 1  Alveolar lymphocytes subpopulations collected by bronchoalveolar lavage in five controls and 10 Crohn's disease patients. Alveolar lymphocytes are expressed as percentage of total alveolar cell count. Lymphocyte subpopulations are expressed as percentage of total number of alveolar lymphocytes.

<table>
<thead>
<tr>
<th>Controls (n=5)</th>
<th>Alveolar lymphocytes</th>
<th>OKT4⁺</th>
<th>OKT4⁻</th>
<th>OKT8⁺</th>
<th>OKT8⁻</th>
<th>OKT4⁺/OKT8⁺</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>11·5±6⁺</td>
<td>64±4·5</td>
<td>44±2·7</td>
<td>30±1·3</td>
<td>1·5±0·15</td>
<td></td>
</tr>
<tr>
<td>Crohn's disease</td>
<td>11·5±6⁺</td>
<td>64±4·5</td>
<td>44±2·7</td>
<td>30±1·3</td>
<td>1·5±0·15</td>
<td></td>
</tr>
<tr>
<td>1</td>
<td>44</td>
<td>57</td>
<td>42</td>
<td>32</td>
<td>1·9</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>11</td>
<td>77</td>
<td>45</td>
<td>32</td>
<td>1·6</td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>18</td>
<td>69</td>
<td>40</td>
<td>26</td>
<td>1·5</td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>25</td>
<td>76</td>
<td>43</td>
<td>29</td>
<td>1·48</td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>46</td>
<td>66</td>
<td>57</td>
<td>13</td>
<td>4·3</td>
<td></td>
</tr>
<tr>
<td>6</td>
<td>79</td>
<td>74</td>
<td>50</td>
<td>20</td>
<td>2·5</td>
<td></td>
</tr>
<tr>
<td>7</td>
<td>37</td>
<td>71</td>
<td>50</td>
<td>17</td>
<td>2·9</td>
<td></td>
</tr>
<tr>
<td>8</td>
<td>10</td>
<td>83</td>
<td>42</td>
<td>39</td>
<td>1·07</td>
<td></td>
</tr>
<tr>
<td>9</td>
<td>5</td>
<td>68</td>
<td>17</td>
<td>54</td>
<td>0·31</td>
<td></td>
</tr>
<tr>
<td>10</td>
<td>33</td>
<td>56</td>
<td>19·2</td>
<td>43·5</td>
<td>0·44</td>
<td></td>
</tr>
</tbody>
</table>

*: Results expressed as mean±SD
Reduced carbon monoxide diffusing capacity without restrictive ventilatory disorder was found in 10 cases. It was associated with lowered Pao2 in one and with increased residual volume and/or low forced expiratory volume in one second suggesting an obstructive ventilatory disorder in two. If carbon monoxide diffusing capacity is corrected according to blood haemoglobin levels,35 however, only three patients with an obstructive or restrictive syndrome have actually a reduced carbon monoxide diffusing capacity. Two patients had normal forced vital capacity and decreased forced expiratory volume in one second and residual volume. Pulmonary function tests were done again six months later in seven patients with initial abnormalities which were found basically unchanged.

367Ga scanning
No pulmonary or mediastinal uptake was observed in nine tested patients.

4 Pulmonary scanning
The percentage of gaseous Xenon remaining in lungs at one minute was identical in Crohn's disease patients and controls: 8.3±8.6 and 4.8±3 (Fig. 3). Pulmonary perfusion scanning showed a decreased perfusion of higher portion of lung associated with an hyperperfusion of lung bases. This resulted in a significant increase of the distance delta between both centres of gravity of lung perfusion and ventilation: 13±3% in Crohn's disease patients versus 8.7±4.3 in controls (p<0.05) (Fig. 3).

5 Blood tests
Serum angiotensin converting enzyme levels were significantly lower in patients (14.1±5.1 u/ml) than in controls (25.2±4.7 u/ml) (p<0.001). Serum β2-microglobulin levels were significantly higher in patients (3.4±1.6 mg/l) than in controls (2.1±0.9 mg/l), (p<0.001). Circulating immune complexes were present in three of seven patients with lymphocyte alveolitis, and in one patient with normal bronchoalveolar lavage.

6 Correlation of BAL results with clinical parameters, pulmonary scanning and function tests and biological data
(Table 2)
There was no apparent correlation between abnormal bronchoalveolar lavage and systemic disorders, activity of intestinal disease, or drug treatment. Patients with alveolitis have a longer (although not significant) Crohn's disease duration (61±70 months) than those without (26.8±25 months). Alveolar lymphocytosis is more frequent in patients having anal Crohn's disease (six of eight patients) than in those without (six of 14 patients) (p<0.05). There was no correlation between alveolar lymphocytosis and pulmonary function tests.
abnormalities, pulmonary scanning results, blood test data, or superoxide anion production by alveolar macrophages. There was a significant correlation between reduced carbon monoxide diffusing capacity and the distance delta between both centre of gravity of lung perfusion and ventilation (r=0.68 p<0.05). Four patients with an obstructive syndrome had a longer Crohn's disease history (66±18 months) than 15 without (30±8 months) (p<0.001).

Discussion

The main finding of our study is a high proportion (54%) of increased alveolar lymphocytosis among patients with Crohn's disease, free of clinical pulmonary symptoms and with normal chest roentgenograms. Alveolar lymphocytosis is persistent as it was present at the same level six months later in four tested patients.

The significance of increased percentage of bronchoalveolar lavage lymphocyte in Crohn's disease is not clear. An increased percentage of bronchoalveolar lavage lymphocytes reflects a shift in the relative proportion of alveolar cells which is classically associated with an alveolitis – that is, an accumulation of immune and inflammatory cells within the alveolar structures. It is well recognised that such alveolar lymphocyte alveolitis, is a characteristic feature of all classic lung granulomatous disorder, such as, sarcoidosis, hypersensitivity pneumonitis, and berylliosis. These diseases were excluded from our study.

An iatrogenic mechanism could be discussed. Indeed most patients with Crohn's disease are treated with sulphasalazine at some stage of the disease and could develop a latent sensitisation to the drug. Previous studies of pulmonary function in inflammatory bowel disease have shown, however, that variations in lung function, especially reduced carbon monoxide diffusing capacity are not a consequence of drug therapy. Moreover 18 of our patients had not received sulphasalazine during the three months before the study. Finally sulphasalazine is known as inducing hypersensitivity pneumonitis. This would have resulted in an increased number of eosinophil polymorphonuclear cells in the bronchoalveolar lavage cell count which we did not observe.

Our study shows the capacity of alveolar macrophages from patients with Crohn's disease to produce superoxide anion. The increased production of superoxide anion is likely to be caused by alveolar macrophages as there were neither neutrophils nor eosinophils polymorphonuclear cells in bronchoalveolar lavage specimens from controls or patients, and as lymphocytes do not produce superoxide anion. The release of superoxide anion by macrophages was not associated with increased alveolar lymphocytosis in patients with Crohn's disease in contrast with previous observations in patients with sarcoidosis. The production of superoxide anion may reflect a spontaneous activation of alveolar macrophages similar to that of circulating monocytes observed in Crohn's disease. Whether or not release of reactive material by macrophages may play a role in tran-
sient alteration of pulmonary function or in the occurrence of alveolar lymphocytosis remains to be established.

It is currently accepted that part of the immune system specific to the gastrointestinal tract is common to all mucosal surfaces.\textsuperscript{40} Therefore the association of Crohn's disease and lymphocyte alveolitis and/or superoxide anion production by alveolar macrophages may be the result of an immunological mechanism: the lymphocytes and/or the macrophages sensitised to antigen at one mucosal site could circulate and inhabit the mucosal surfaces of the lung.

Abnormal pulmonary function tests were noted in 15 of 21 patients and were similar to findings previously described in inflammatory bowel disease.\textsuperscript{22-26} Abnormal pulmonary function tests do not seem to be related to alveolar lymphocytosis which is therefore latent (Table 2).

Our patients have an increased physiologic dead space in the upper part of their lung related to any hypoperfusion of this lung portion. This may explain in part reduced carbon monoxide diffusing capacity in several Crohn's disease patients. Different perfusion rates in lung upper and lower part in patients with Crohn's disease is unexplained. Release of vasoactive substances inducing vasodilation in pulmonary circulation may be one hypothesis. Alveolar distension resulting in an increase of alveolar pressure is another hypothesis suggested by the existence of an obstructive ventilatory disorder in some patients. Crohn's disease related distensibility abnormalities of pulmonary vessels in lung bases under hydrostatic pressure could be another explanation.\textsuperscript{41,42}

Pulmonary abnormalities in Crohn's disease are identical to some reported in pulmonary sarcoidosis: alveolar lymphocytosis and activation of alveolar macrophage. Some differences exist between Crohn's disease and sarcoidosis, however: (a) Crohn's disease patients had dramatically low serum angiotensin converting enzyme; (b) no mediastino pulmonary \textsuperscript{67}Ga uptake was observed in Crohn's disease patients; (c) OKT\textsubscript{4} lymphocytes are not always raised in Crohn's disease patients.\textsuperscript{36} Whatever the significance of our data, persistent abnormalities of bronchoalveolar lavage and pulmonary function tests suggest a latent involvement of the lung in Crohn's disease which merits further investigation.

References


7 Collins WJ, Bending DW, Taylor WF. Pulmonary vasculitis complicating childhood ulcerative colitis. \textit{Gastroenterology} 1979; 77: 1051-3.


23 Healtry RV, Thomas P, Prokipchuk EJ, Gauldie J.
Latent pulmonary involvement in Crohn's disease


29 Best WR, Becktel JM, Singleton JW. Redevized values of the eight coefficients of the Crohn's Disease Activity Index (CDAI). *Gastroenterology* 1979; 77: 843–7.


32 Quanjer PH. *Standardized lung function testing report working party 'Standardization of lung function tests'*. Luxembourg: European Community for Coal and Steel, 1981.


P Bonniere, B Wallaert, A Cortot, X Marchandise, Y Riou, A B Tonnel, J F Colombel, C Voisin and J C Paris

_Gut_ 1986 27: 919-925
doi: 10.1136/gut.27.8.919

Updated information and services can be found at:
_http://gut.bmj.com/content/27/8/919_

_Email alerting service_
Receive free email alerts when new articles cite this article.
Sign up in the box at the top right corner of the online article.

_Topic Collections_
Articles on similar topics can be found in the following collections
_Crohn's disease_ (932)
_Gastrointestinal hormones_ (848)

_Notes_

To request permissions go to:
_http://group.bmj.com/group/rights-licensing/permissions_

To order reprints go to:
_http://journals.bmj.com/cgi/reprintform_

To subscribe to BMJ go to:
_http://group.bmj.com/subscribe_/