

# Evidence for protein oedema, neutrophil influx, and enhanced collagen production in lungs of patients with systemic sclerosis

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## Abstract

**Bronchoalveolar lavage fluid from patients with systemic sclerosis was analysed for evidence of pulmonary vascular leakage, inflammatory cell influx, and enhanced type III collagen synthesis. Eighteen patients with systemic sclerosis and computed tomographic evidence of fibrosing alveolitis were compared with 16 patients with a normal scan. The albumin concentration in lavage fluid was higher in all patients than in normal volunteers. Patients with an abnormal computed tomogram as a group had increased proportions of all inflammatory cell types, whereas those with a normal scan had increased neutrophils only. Increased lavage type III procollagen peptides were found in all patients with an abnormal computed tomogram and eight of those with a normal scan. These results suggest that pulmonary vascular leakage and neutrophil influx may be early pathological features of lung disease in systemic sclerosis and frequently associated with enhanced collagen production. Thus lavage of patients with systemic sclerosis may identify lung inflammation and altered collagen metabolism early in the evolution of fibrosing alveolitis.**

Systemic sclerosis is characterised by a vascular disorder and excessive deposition of collagen and other matrix proteins in the skin and internal organs.<sup>1</sup> Symptoms and signs of lung disease are rare when patients first present, yet up to 80% have abnormal lungs at necropsy.<sup>2,3</sup> The histological appearances are identical to those of cryptogenic fibrosing alveolitis.<sup>4</sup> The investigation of patients with systemic sclerosis offers an opportunity therefore to study fibrosing alveolitis from the earliest stages of its evolution.

Symptomless patients with collagen vascular disorders, including a few with systemic sclerosis, have evidence of subclinical alveolitis.<sup>5</sup> There are, however, currently no data to indicate whether this alveolitis is associated with other pathological processes, such as pulmonary vascular leakage of plasma proteins or altered lung collagen metabolism.

The aim of this study was to examine the lower respiratory tract of patients with systemic sclerosis for evidence of pulmonary vas-

cular leakage, inflammatory cell infiltration, and enhanced lung collagen metabolism.

## Methods

### PATIENTS

We investigated 34 patients fulfilling the American Rheumatism Association's preliminary criteria for the diagnosis of systemic sclerosis.<sup>6</sup> All underwent clinical evaluation, including chest radiography, lung function tests, thin (3 mm) section thoracic computed tomography,<sup>7</sup> and liver and renal function tests. The clinical data are summarised in table 1. Patients were divided into those with and those without computed tomographic evidence of fibrosing alveolitis.<sup>8</sup> Each scan was assessed by a radiologist who was unaware of the patient's clinical condition.

### BRONCHOALVEOLAR LAVAGE

All patients underwent fibreoptic bronchoscopy and bronchoalveolar lavage.<sup>9</sup> Lavage fluid was collected in a siliconised container on ice, and all subsequent manipulations were performed at 4°C to minimise protein degradation during processing. Lavage was also performed on 10 healthy, non-smoking volunteers who acted as controls. The median age of the control group was 29 (range 21-36) years.

Lavage fluid was centrifuged at 300 *g* for five minutes, and the supernatant stored at -40°C. The cell pellet was resuspended and the total and differential cell count determined immediately.<sup>9</sup> Blood was collected from all subjects at the time of bronchoscopy and serum stored at -40°C. Albumin concentrations were measured calorimetrically in 20 µl aliquots of serum and 1 ml aliquots of unconcentrated lavage fluid, the Bromocresol Green binding reaction being used (Sigma Chemical Company, Poole, Dorset).

### TYPE III PROCOLLAGEN N PEPTIDE ASSAY

Bronchoalveolar lavage fluid was concentrated by dialysing it extensively against an aqueous solution of 0.02 M ammonium bicarbonate at 4°C with a molecular weight cut off of 2000 (Sigma Chemical Company). Known volumes (1.25-2.5 ml) were then lyophilised in 3.5 ml polystyrene test tubes (Sarstedt, West Germany) and redissolved in phosphate buffered saline containing 0.4 g/l Tween 20 to give 50-100 fold concentrates. Duplicate samples of concentrated lavage fluids and 25 µl

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Table 1 Clinical, radiographic, and physiological characteristics of 34 patients with systemic sclerosis (medians and ranges unless otherwise specified)

Group	Age (y)	Duration (y) of systemic sclerosis	Smoking habit (NS:S)	Radiograph (N:Abn)	Lung function tests		
					FVC (% pred)	TLCO (% pred)	Exercise A-aDO <sub>2</sub> (kPa)
CT abnormal	47 (24-62)	7 (1-16)	18:0	4:14	75 (47-106)	52 (37- 80)	3.6 (1.9-8.5)
CT normal	49.5 (31-64)	3* (1-16)	13:3	16: 0	99** (81-112)	76** (54-104)	2.4 (1.2-3.6)

\*p < 0.05; \*\*p < 0.01.  
CT—computed tomogram; NS:S—non-smokers:smokers; N:Abn—normal:abnormal; FVC—forced vital capacity; TLCO—carbon monoxide transfer factor; A-aDO<sub>2</sub>—alveolar-arterial oxygen tension difference.

Table 2 Characteristics of lavage fluid from patients with scleroderma

Group	Lavage fluid (% recovered)	Albumin (mg/ml)	BALF:serum albumin ratio (× 10 <sup>3</sup> )	Total cells (× 10 <sup>6</sup> )
Controls (n = 10)	63 (32-79)	0.04 (0.027-0.09)	1.45 (0.6-1.37)	12.2 (4.4-21.7)
Patients:				
CT abnormal (n = 18)	42 (19-56)	0.23* (0.1-0.36)	7.3* (2.9-11.6)	17.7 (6.2-48)
CT normal (n = 16)	40 (25-52)	0.19* (0.1-0.32)	5.75* (2.8-9.1)	11.5 (3.0-39)

\*p < 0.01 in the comparison with controls.  
CT—computed tomogram; BALF—bronchoalveolar lavage fluid.

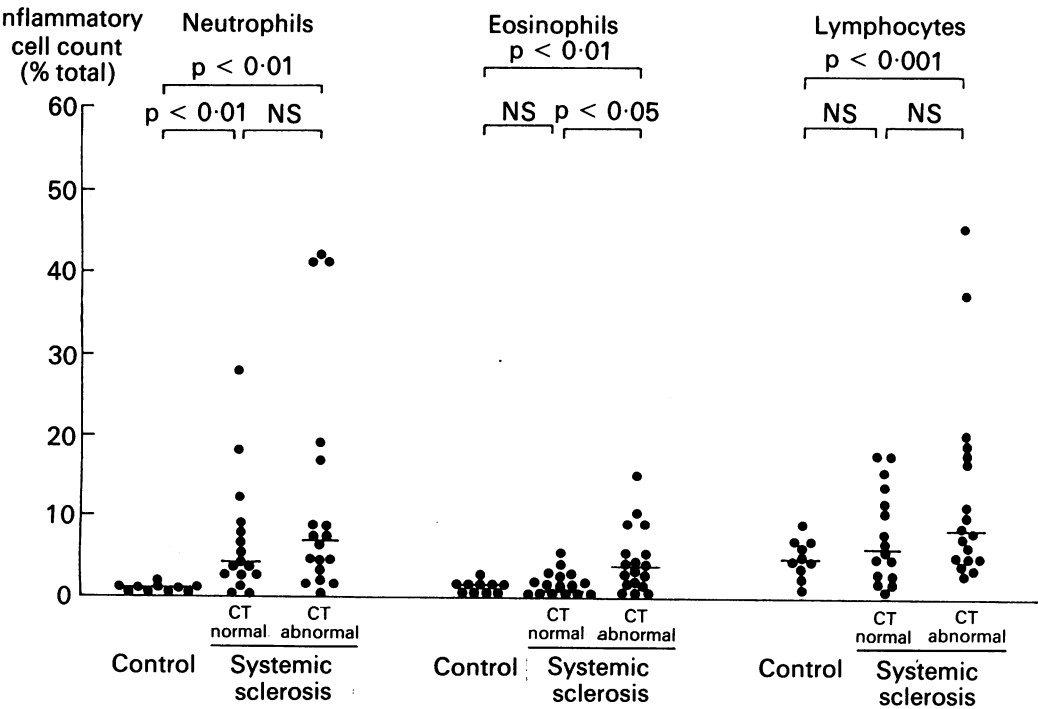
aliquots of serum were assayed for antigens related to type III procollagen N terminal peptide by a commercially available radio-immunoassay (Hoechst, West Germany), which uses Fab fragments of antibody to the peptide.<sup>10</sup> Serial dilutions of an unlabelled procollagen N peptide standard were assayed in triplicate and used to construct an inhibition curve. The Fab fragments have equal affinity for the different antigenic forms of the peptide that occur in biological fluids, and therefore inhibition curves generated by serial dilutions of such fluids parallel those generated by dilutions of the standard.<sup>10</sup>

STATISTICS  
Group data of non-parametric variables are expressed as medians with ranges and com-

pared by the Mann-Whitney U test. Correlations were assessed with Kendall's rank correlation coefficient.

Results  
LAVAGE FLUID ALBUMIN AND CELLULARITY  
The albumin concentrations in both groups of patients with systemic sclerosis were significantly greater than those from control subjects, when values were expressed both as absolute amounts and as lavage fluid: serum ratios (table 2). The total number of cells recovered from each subject varied considerably and no differences between the groups were observed (table 2). The differential cell counts show that patients with an abnormal computed tomogram had higher proportions of

Figure 1 Inflammatory cell profiles (% total cell count) in patients with systemic sclerosis. CT—computed tomogram.



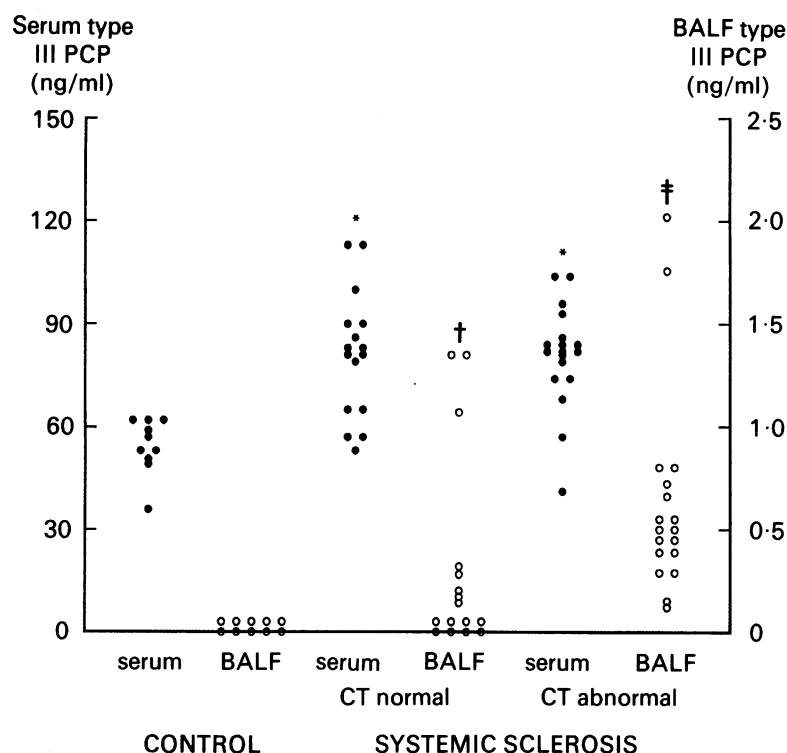


Figure 2 Type III procollagen N terminal peptides in serum (●) and bronchoalveolar lavage fluid (BALF: ○) from 34 patients with systemic sclerosis and 10 healthy volunteers. \* $p < 0.01$  controls; † $p < 0.02$ , both in the comparison with controls; ‡ $p < 0.02$  in the comparison with patients with normal computed tomograms. PCP—procollagen peptides; CT—computed tomogram.

neutrophils ( $p < 0.01$ ), eosinophils ( $p < 0.01$ ), and lymphocytes ( $p < 0.001$ ) than did normal volunteers (fig 1).

The patients with a normal scan had increased proportions of neutrophils ( $p < 0.01$ ), which remained significant ( $p < 0.02$ ) when the three smokers were excluded from the analysis. Eosinophils and lymphocytes were not significantly increased, though 11 of the 16 patients in this group had an increased proportion of at least one inflammatory cell type.

Lavage fluid from the seven patients with a normal scan and impaired carbon monoxide transfer factor (TLCO  $< 75\%$  predicted) did not differ significantly in albumin concentration or proportion of neutrophils from that of the remainder of the group.

#### TYPE III PROCOLLAGEN PEPTIDES IN SERUM AND LAVAGE FLUID

The serum concentrations of procollagen N peptide related antigenic material in patients were significantly greater than those seen in the normal volunteers, though there was considerable overlap (fig 2). The assay did not distinguish the patients with systemic sclerosis who had computed tomographic evidence of fibrosing alveolitis from those with a normal scan.

Procollagen peptides were undetectable in lavage fluid from normal subjects but were detected in all patients with systemic sclerosis who had an abnormal scan and in eight of the patients with a normal scan. In those with a normal scan there was no statistical difference in procollagen N peptide concentrations in lavage fluid between those with impaired car-

bon monoxide transfer and those in whom it was normal.

When serum and lavage fluid concentrations of procollagen N peptide were corrected for albumin concentration they showed trends similar to those shown in figure 2, but lavage fluid concentrations in the patients often exceeded those in corresponding serum. The mean procollagen N peptide concentrations in lavage fluid in the patients with computed tomographic evidence of fibrosing alveolitis were two to three times greater than the concentrations in patients without evidence of lung disease ( $p < 0.001$ ). Concentrations of procollagen N peptide in lavage fluid did not correlate with the duration of systemic sclerosis, results of physiological tests of lung function, lavage fluid albumin concentration, or inflammatory cell counts in lavage fluid.

#### Discussion

Raised concentrations of albumin were found in the lavage fluid from all patients, irrespective of whether there was clinical evidence of pulmonary fibrosis. This suggests that increased pulmonary vascular leakage may be one of the earliest abnormalities of lung disease in systemic sclerosis and can be detected by bronchoalveolar lavage. This observation is consistent with findings from studies in experimental animals, where vascular leakage is detectable many days before pulmonary fibrosis is apparent biochemically or histologically.<sup>11-14</sup> The relevance of increased vascular permeability to the pathogenesis of pulmonary fibrosis is uncertain, but could indicate loss of endothelial integrity, allowing leakage of circulating blood proteins, inflammatory cells, or platelets from the circulation into the lung interstitium. These could provide a rich source of mediators capable of promoting lung collagen production by stimulating either fibroblast proliferation<sup>15-17</sup> or collagen production by individual fibroblasts.<sup>18</sup>

Lavage fluid cell counts in our study showed that patients with systemic sclerosis and evidence of fibrosing alveolitis as a group had higher proportions of neutrophils, eosinophils, and lymphocytes than control subjects, confirming earlier reports.<sup>19-21</sup> Patients without evidence of fibrosing alveolitis had increased proportions of neutrophils alone, even when the smokers were excluded. This agrees with an earlier report of increased neutrophils in the lavage fluid of six of 10 patients with systemic sclerosis and normal chest radiograph and normal lung function.<sup>5</sup> These observations suggest that neutrophil alveolitis is an early event in the pathogenesis of lung disease in systemic sclerosis, though analysis of lavage fluid may not reflect changes in the interstitium, where infiltration by other inflammatory cells, such as lymphocytes, may occur.<sup>4</sup>

There is currently no direct information on in vivo collagen production by the human lung, though several indirect techniques have been applied to the study of patients with fibrotic lung disorders. In general, these have been either measurements of post-translational



processing of procollagen molecules<sup>22</sup> or the enzymes concerned in this process.<sup>23</sup>

The serum procollagen *N* peptide concentrations we found are similar to those previously reported for both normal volunteers<sup>25</sup> and patients with systemic sclerosis.<sup>26</sup> Our results indicate that the measurement of serum procollagen *N* peptide does not distinguish patients with clinical evidence of pulmonary fibrosis from those without. By contrast, our results for bronchoalveolar lavage fluid suggest a progressive increase in type III collagen synthesis with the development of pulmonary fibrosis, though these data do not necessarily indicate that type III collagen is laid down as mature collagen fibrils in the lung interstitium. Nevertheless, our results complement those of previous morphometric<sup>27</sup> and biochemical<sup>28, 29</sup> studies, in which an increased lung collagen content was found in fibrosing alveolitis.

The lack of correlation between procollagen *N* peptide concentrations in lavage fluid and results of physiological tests of lung function requires explanation. Our data could mean that raised procollagen *N* peptide concentrations in lavage fluid, and by implication increased type III collagen synthesis by the lung, may occur very early in the pathogenesis of pulmonary fibrosis, at a stage undetectable by conventional techniques, including computed tomography of the thorax. It has been suggested that in areas of established pulmonary fibrosis type III collagen tends to be replaced by type I collagen<sup>27</sup>; in the patients with advanced lung disease therefore lavage fluid procollagen *N* peptide might be expected to return to more modest levels.

Our observations suggest increased pulmonary vascular permeability in patients with systemic sclerosis and concentrations of procollagen *N* peptide in lavage fluid might reflect leakage of these antigens from the circulation. There is evidence to refute this proposal, however. There was no correlation between albumin and procollagen *N* peptide in lavage fluid and an appreciable number of patients had a concentration of the peptide in their lavage fluid which, in relation to the albumin concentration, still greatly exceeded the level in their serum. This suggests enhanced type III collagen production by the lungs.

Increased concentrations of procollagen *N* peptide in bronchoalveolar lavage fluid from patients with other interstitial lung diseases have been reported,<sup>22, 30–32</sup> but this is the first study of pulmonary disease in systemic sclerosis. All previous studies were based on a radioimmunoassay using whole IgG anti-procollagen *N* peptide.<sup>33</sup> Although this assay is very sensitive, it preferentially measures the Col 1–3 antigen. Type III procollagen peptides in lavage fluid may occur in several different antigenic forms<sup>34</sup> and the assay using whole antibody may be less appropriate for use on lavage fluid, though it is the Col 1–3 peptide that probably reflects collagen synthesis most accurately. Despite differences in the assays, data from the present and previous studies suggest that procollagen peptides in lavage fluid reflect altered type III collagen metabolism of the lung interstitium in vivo.

This study suggests that increased pulmonary vascular leakage and neutrophil alveolitis may be early features of interstitial lung disease in systemic sclerosis and that these abnormalities are frequently associated with evidence of enhanced lung type III collagen synthesis. Possibly these inflammatory changes could resolve in some patients without the development of lung fibrosis. These findings are likely to be relevant to the early clinical diagnosis of pulmonary fibrosis and to investigators interested in understanding its pathogenesis.

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