Diffuse interstitial pulmonary fibrosis in patients with a positive sheep cell agglutination test

T. B. STRETTON AND J. T. LEEMING

From the Manchester Royal Infirmary

In 1944 Hamman and Rich described an acute form of diffuse interstitial pulmonary fibrosis in four patients who died within six months of the onset of symptoms. Many cases with a similar syndrome have since been reported. The majority of these, however, have run a more chronic course and the outcome has not always been fatal. The disorder has been fully reviewed by Rubin and Lubliner (1957) and Scadding (1960); its aetiology remains unknown. A number of patients with rheumatoid disease have had diffuse pulmonary fibrosis indistinguishable from the idiopathic variety (Ellman and Ball, 1948; Rubin and Lubliner, 1957; Smith and Rothermich, 1957; Scadding, 1960; Ognibene, 1960; Cudkowicz, Madoff, and Abelmam, 1961). These patients have all had overt arthritis, except for one of the patients described by Ognibene (1960); in this instance there was diffuse pulmonary fibrosis and the sheep cell agglutination test (S.C.A.T.) was positive but there was no gross arthritis, a diagnosis of anarthritic rheumatoid disease (Bagratuni, 1956) being made. We report here two further similar cases.

CASE REPORTS

CASE 1 A woman aged 45 was admitted in May 1960. For seven years she had suffered from increasingly severe breathlessness on exertion and for five years she had had a dry cough which recently had become productive of white, mucoid sputum. This illness was accompanied by general tiredness and lassitude. During the two years before her admission she had been receiving digitalis and occasional chlorothiazide from her general practitioner for the treatment of ankle swelling; the swelling improved but there was no change in the increasing breathlessness.

On examination there was gross finger clubbing and slight central cyanosis. She was in sinus rhythm, the blood pressure was 115/80 mm. Hg and there was no cardiac abnormality; the venous pressure was normal. Coarse crepitations were audible over all areas of the lungs. The liver edge was palpable just below the right costal margin. No abnormalities were found in the other systems.

Investigations The chest radiograph (Fig. 1) showed changes in the lungs typical of gross, widespread fibrosis with slight honeycombing in the lower zones. Serial radiographs had been taken at a chest clinic during the previous seven years and these were available for comparison. They revealed that the fibrosis had steadily increased in extent and severity over the years.

The patient’s vital capacity was 0·95 litre, 27% of her predicted vital capacity of 3·55 litres. The one second forced expiratory volume (F.E.V.1) was 0·9 litre, giving a normal F.E.V.1 to vital capacity ratio (95%).

The sputum contained normal upper respiratory tract flora and a beta-haemolytic streptococcus.

The S.C.A.T. was positive at a titre of 1/2048.

The serum albumin was 2·5 g./100 ml. and the globulin 6·8 g./100 ml. Electrophoresis revealed a marked increase in the gamma-globulin, and there were no abnormal protein bands. The Sia test was positive; there was no cryoglobulin. Serum bilirubin was 0·3 mg./100 ml.; thymol turbidity 25 units, zinc sulphate turbidity 44 units, and the alkaline phosphatase 63 King-Armstrong units/100 ml. Serum glutamic-oxaloacetic transaminase was 52 units/100 ml. and the serum glutamic-pyruvate transaminase was 66 units/100 ml. The prothrombin activity was 80% of normal but rose to 100% after the administration of vitamin K1. A bromsulphthalein test revealed 19% retention at 45 minutes.

A liver biopsy revealed granulomatous foci in the portal tracts; these foci consisted of endothelioid cells, lymphocytes, occasional plasma cells, and polymorph leucocytes.

The haemoglobin (Hb) was 11·8 g./100 ml. (80%). The total white cell count (W.B.C.) was 9,000/c.mm.; of these 60% were polymorphs, 33% lymphocytes, 3% mononuclears, and 4% eosinophils. Lupus erythematosus (L.E.) cells were not found. The erythrocyte sedimentation rate (E.S.R.) was 52 mm. in one hour (Wintrobe). The Mantoux test was positive 1/100. A Kveim test was negative.

Serum sodium was 140, potassium 5·4, chloride 92, and plasma bicarbonate 27 mEq/l. The blood urea was 26 mg./100 ml. Serum calcium was 9·3 and phosphorus 3·6 mg./100 ml.

1 Present address: Bolton District General Hospital, Lancashire
FIG. 1. Case 1. (a) Chest radiograph; (b) enlarged view of the left lower zone of the chest radiograph to show in detail the pulmonary abnormality.
An electrocardiogram was normal apart from digitalis effect.

**Treatment and subsequent investigations** In view of the steady deterioration over the years, both symptomatically and radiologically, corticosteroid therapy was begun: prednisone was given in a dose of 10 mg. twice daily and has been continued up to the present time. Digitalis was continued and she had occasional courses of tetracycline for intercurrent pulmonary infections. No change has occurred in the patient's condition during the two years of prednisone therapy although the amount of pulmonary fibrosis appears radiologically to have increased slightly.

In December 1961 the results of further pulmonary function tests were as follows: the vital capacity was 1·08 litres (30% of the predicted value) and the F.E.V.1 was 0·9 litre (83% of the measured vital capacity). The arterial Pco2 (determined by the method of Campbell and Howell, 1960) was 40 mm. Hg.

Cardiac catheterization was performed in view of the suggestion from the history that the patient had previously been in congestive cardiac failure. (If this were so it was felt that there must be pulmonary hypertension although this was not manifest clinically. Because of the prognostic importance direct estimation of the pulmonary arterial pressure was thought to be desirable.) By the steady state Fick method the cardiac output was 3·59 l./min. at rest and 5·32 l./min. on exercise with an oxygen consumption of 244 ml./min. and 343 ml./min. respectively. The arterial and mixed venous oxygen saturations were 81% and 40% at rest and 81% and 41% on exercise. The mean pulmonary artery pressure at rest was 14 mm. Hg and on exercise it was 16 mm. Hg.

The liver function tests have been repeated on numerous occasions. The pattern of abnormality has not changed: the serum albumin has varied from 2·3 to 3·6 g./100 ml.; globulin 5·8 to 6·8 g./100 ml.; thymol turbidity 15 to 25 units; zinc sulphate turbidity 44 to 63 units; alkaline phosphatase 58 to 86 units/100 ml.; the serum bilirubin has been consistently normal.

**Case 2** A woman aged 61 was admitted in November 1960. She had been perfectly well until 10 months previously when she became increasingly breathless, at first on exertion but later also at rest. For four months she had had a dry, irritating cough, productive of a little white, frothy sputum. During this time she noticed that the fingers were becoming clubbed. On admission she was desperately breathless with a respiratory frequency of about 40/minute. There was central cyanosis. The slightest exertion in bed, such as sitting forwards unaided whilst her back was examined, produced the most intense dyspnoea and increased the degree of cyanosis. These symptoms disappeared when she was given oxygen to breathe. Clubbing of the fingers and toes was marked. She had a sinus tachycardia of 100/minute; the blood pressure was 110/70 mm. Hg. The cardiac impulse was normal; there was a systolic ejection click and a mid-systolic murmur in the pulmonary area. The jugular venous pressure was 4 cm. above the sternal angle and there was a trace of sacral oedema. Showers of crepitations were heard over all areas of the lungs. No other abnormalities were found.

**Investigations** Chest radiographs showed widespread changes in both lungs consistent with diffuse interstitial fibrosis together with some honeycombing. (The appearances were almost identical with those in Case 1.)

The sputum yielded a mixed growth of organisms with a predominant growth of *Staphylococcus saprophyticus*.

The S.C.A.T. was positive at a titre of 1/256.

The serum albumin was 3·3 g./100 ml., the globulin was 4·4 g./100 ml., bilirubin 0·8 mg./100 ml., thymol turbidity 6-2 units, and the alkaline phosphatase was 60 King-Armstrong units/100 ml. The blood urea was 55 mg./100 ml. Serum sodium was 143, potassium 4-8, chloride 92, and the plasma bicarbonate 34 mEq/litre.

The Hb was 16·6 g./100 ml. (114%). The total W.B.C. was 13,400/c.mm. (77% polymorphs; 16% lymphocytes; 7% monocytes). No L.E. cells were found. The E.S.R. was 15 mm. in one hour (Wintrobe).

An electrocardiogram demonstrated the presence of left bundle branch block.

**Treatment and subsequent investigations** Continuous oxygen therapy was given. Prednisone was commenced at a total daily dose of 60 mg. Antibiotics were administered. She was also given digitalis and diuretics. Her condition slowly improved so that in three weeks she was comfortable without the added oxygen. Serial radiographs, however, showed no change in the pulmonary fibrosis. Left bundle branch block persisted. The blood urea fell to 30 mg./100 ml. and the plasma bicarbonate fell to 30 mEq/l. The serum globulin fell to 3-2 g./100 ml. and the thymol turbidity to 2 units.

The vital capacity three weeks after admission was 1·21 litres, 37% of her predicted vital capacity (3·3 litres). The F.E.V.1 was 0·86 litre, giving a ratio of F.E.V.1 to vital capacity of 71%. At this time the estimated arterial Pco2 was 42 mm. Hg. Five weeks later the vital capacity had increased to 1·39 litres (42% of the predicted value), and the F.E.V.1 was 1·24 litres (giving an 89% F.E.V.1 to vital capacity ratio).

Cardiac catheterization was performed two months after admission, when there was clinical improvement, for the same reasons as in the first patient. The mean pulmonary arterial pressure was 18 mm. Hg. The arterial blood was 83% saturated and the mixed venous blood was 56%. The oxygen uptake at rest was 220 ml./min. and the cardiac output was 4·67 litres/min. Exercise studies were not feasible.

The initial improvement on treatment was maintained apart from an attack of supraventricular tachycardia, and the patient was discharged home five months after her admission. At this time she was able to sit out of bed and could walk short distances.
without gross dyspnoea. Two weeks later, however, her condition abruptly deteriorated and she died suddenly at home.

A necropsy was performed by Dr. G. Greville Williams. The main abnormalities were limited to the lungs. The central parts of these appeared normal to the naked eye but the peripheral halves were composed of a mass of smooth-walled cavities 1 to 1.5 mm. in diameter. Histologically, in the less affected areas there was patchy fibrosis of the alveolar septa. The cysts (Fig. 2) mostly had no epithelial lining and were surrounded by a mixture of collagen and smooth muscle. There were a few lymphocytes and rather more plasma cells in the interstitial tissue with occasional cholesterol or fat-containing giant cells. A superimposed mild acute inflammation was seen. The overall picture was considered to be one of honeycomb lung, as described by Heppleston (1956). Skin and subcutaneous tissue from the elbows and hee's and the metacarp-phalangeal joint of the right index finger were taken for histological study by Dr. J. Ball. There were no gross macroscopical changes in these regions but microscopically there was evidence of tissue damage. There was some fibrous thickening of the olecranon bursa of one elbow. At all sites there was hyaline fibrosis of the subcutaneous nerves with loss of myelin and axons; this was thought to be related to the obliterator endarterial fibrosis which was a feature of many of the small arteries in the subcutaneous tissues. No active arthritis was found though there was evidence of an old, healed, minor erosive arthropathy.

No significant abnormalities were found in other organs.

**COMMENT** Both these patients had restrictive pulmonary disease with normal alveolar ventilation ($P_{\text{CO}_2}$ of 40 and 42 mm. Hg) and arterial unsaturation (81% and 83% respectively) breathing room air at rest. In the first patient the arterial saturation did not fall further on exercise, suggesting that this was due not to a limitation in diffusion but to a venous to arterial shunt (of some 30% of the cardiac output). Exercise studies could not be done in the second patient but clinical examination showed that the slightest exertion whilst breathing room air caused further brisk arterial unsaturation and intense dyspnoea; these observations suggested that the patient had a critical defect of gaseous diffusion. A surprising finding in both patients in the face of such extensive pulmonary fibrosis was the normal pulmonary arterial pressure and vascular resistance.

**DISCUSSION**

Oswald and Parkinson (1949) were the first authors to emphasize that honeycomb lungs occur in various diseases in which there is diffuse interstitial pulmonary infiltration. In six of their patients there was an associated disorder such as tuberous sclerosis, xanthomatosis, biliary cirrhosis or pituitary disease. In the other 10 cases no such recognizable disease was noted and, where histological examination was carried out, extensive interstitial pulmonary fibrosis was found. Heppleston (1956), in an extensive study of the pathology of honeycomb lung, considered that the essential abnormality was an obliteration of the respiratory or non-respiratory bronchioles. He suggested that neighbouring unaffected bronchioles then underwent compensatory enlargement to form the cystic spaces. In the majority of his cases the obliterator process was a non-specific, interstitial fibrosis though in some a recognizable granulomatosis was seen. Heppleston concluded that honeycomb lung was, in most instances, the result of patchy, interstitial fibrosis. The distinction between the two conditions is probably, therefore, somewhat artificial and, as shown by our case 2, radiological differentiation may be quite impossible.
The aetiology of diffuse interstitial fibrosis of the lungs is often obscure (Scadding, 1960) but it has been found in association with a variety of diseases, including the connective tissue disorders (Rubin and Lubliner, 1957). It has been found in a number of patients with rheumatoid disease (Ellman and Ball, 1948; Rubin and Lubliner, 1957; Smith and Rothermich, 1957; Scadding, 1960; Ognibene, 1960; Cudkowicz et al., 1961). Honeycomb lung (sic) has also been found in rheumatoid patients (Hart and Mackenzie, 1955; Dixon and Ball, 1957; Brinkman and Chaikof, 1959). Other pulmonary disorders occur in rheumatoid disease: these include pleural effusions (Emerson, 1956; Mason and Steinberg, 1958; Horler and Thompson, 1959; Ward, 1961), rheumatoid nodules in the pleura or lung parenchyma (Christie, 1954; Gresham and Kellaway, 1958), Caplan’s syndrome (Caplan, 1953), and a pulmonary arteritis (Price and Skelton, 1956; Gardner, Duthie, Macleod, and Allen, 1957). Of considerable interest is the occurrence of certain of these pulmonary disorders before the onset of the arthritis; this has been described in pleural effusions (Horler and Thompson, 1959) and in Caplan’s syndrome where the development of pulmonary nodules has been known to antedate the onset of arthritis by up to 10 years (Caplan, 1959).

The presence of the positive S.C.A.T. in the two patients described here raised the possibility that each had a diffuse connective tissue disorder, the chief manifestation of which was the pulmonary abnormality. This possibility was strengthened in the first patient by the discovery of liver lesions although the exact nature of these was obscure. Similar lesions have been seen previously by one of us in a patient referred to by Scadding (1960) but in this patient the S.C.A.T. was negative (Scadding, personal communication, 1961). Furthermore, each patient was seen by Professor J. H. Kellgren who found evidence of minor abnormalities in the locomotor systems though at no time were there any symptoms referable to the joints. Finally, in the second patient, histological examination of tissue from the elbows and heels revealed evidence of tissue damage with obliterative endarterial fibrosis in many of the small subcutaneous arteries. Although there was no active arthritis, suggestive evidence of an old, minor, erosive arthropathy was found.

One similar case has been reported by Ognibene (1960): the patient was a 67-year-old woman who developed histologically proven diffuse interstitial fibrosis of the lungs and had a positive S.C.A.T. without arthritis; she had, however, had arthralgia and muscle pains for a year before the onset of her pulmonary symptoms.

Kellgren and Ball (1959) have shown that the S.C.A.T. is most constantly positive in rheumatoid disease though it is sometimes positive in other connective tissue diseases: in systemic lupus erythematosus and scleroderma positive titres were found when articular or peripheral vascular lesions (or both) were prominent; in polyarteritis nodosa Kellgren and Ball found that the S.C.A.T. was only positive when the polyarteritis developed as a complication of severe rheumatoid arthritis. In population studies Ball and Lawrence (1961) found that 1-6 to 5-4% of random samples of rural and urban communities had a positive S.C.A.T.; of these, only 20% had evidence of rheumatoid arthritis according to the criteria of the American Rheumatism Association (1959). But the incidence of rheumatoid arthritis rose sharply in subjects with S.C.A.T. titres greater than 1 in 16: in those with titres of 1 in 256 or more the incidence was 70%. Furthermore, in a five-year follow-up of a selected urban group, of seven subjects who originally had a positive S.C.A.T. but no joint disease, two had in the meantime developed rheumatoid arthritis. The S.C.A.T. was studied by Ball (1955) in a group of South Wales coal-miners (with rheumatoid arthritis) investigated by Miall (1955). He found that the incidence of a positive S.C.A.T. and the titre of the tests were higher in subjects with progressive massive fibrosis or Caplan’s syndrome than in those with simple pneumoconiosis or a normal chest radiograph. This difference could not be explained by differences in the severity or duration of the arthritis and was clearly related in some unknown way to the lung lesion. It seems quite possible, therefore, that the positive S.C.A.T.s in our patients were indicative of the presence of rheumatoid disease despite the absence of joint symptoms.

The present cases, together with that reported by Ognibene (1960), support the hypothesis that diffuse interstitial pulmonary fibrosis may occasionally be due to rheumatoid disease despite the absence of arthritis. We suggest that all such patients are screened by means of the S.C.A.T. to determine the frequency with which rheumatoid disease occurs in what may otherwise appear to be an idiopathic disorder.

**SUMMARY**

Two patients with severe diffuse interstitial pulmonary fibrosis are described. Each had a positive S.C.A.T. but neither had any symptoms
referable to the joints. Evidence is presented that these patients had a connective tissue disorder, possibly rheumatoid disease, despite the absence of arthritis.

We wish to thank Dr. C. S. D. Don and Dr. A. Morgan Jones for permission to publish details of these patients. We are also indebted to Professor J. H. Kellgren and Dr. E. G. Wade for advice, and to Dr. J. Ball and Dr. G. Greville Williams for the pathological reports.

REFERENCES


---

**ADDENDUM**

Since this paper was prepared, Dr. F. I. Lee and Dr. A. T. Brain (*Lancet*, 1962, 2, 693) have described three patients who had chronic interstitial fibrosis of the lungs and later developed rheumatoid arthritis. They concluded that diffuse pulmonary fibrosis may sometimes be a manifestation of rheumatoid disease even in the absence of overt arthritis.
Diffuse Interstitial Pulmonary Fibrosis in Patients with a Positive Sheep Cell Agglutination Test
T. B. Stretton and J. T. Leeming

Thorax 1964 19: 79-84
doi: 10.1136/thx.19.1.79