Background: Pulmonary hypertension (PH) is a rare complication of sarcoidosis, although it is not uncommon in advanced disease.

Methods: A retrospective series of 22 sarcoidosis patients (16 men) of mean (SD) age 46 (13) years with PH was divided into two groups depending on the absence (stage 0: n = 2, stage II: n = 4, stage III: n = 1) or presence (n = 15) of radiographic pulmonary fibrosis at the time of PH diagnosis.

Results: In both groups PH was moderate to severe and there was no response to acute vasodilator challenge. In non-fibrotic cases no other cause of PH was found, suggesting a specific sarcoidosis vasculopathy, although no histological specimens were available. In cases with fibrosis there was no correlation between haemodynamics and lung volumes or arterial oxygen tensions, suggesting other mechanisms for PH in addition to pulmonary destruction and hypoxaemia. These included extrinsic arterial compression by lymphadenopathies in three cases and histologically proven pulmonary veno-occlusive disease in the five patients who underwent lung transplantation. Ten patients received high doses of oral prednisone for PH (stage 0: n = 1, stage II: n = 4 and stage IV: n = 5); three patients without pulmonary fibrosis experienced a sustained haemodynamic response. Survival of the overall population was poor (59% at 5 years). Mortality was associated with NYHA functional class IV but not with haemodynamic parameters or with lung function.

Conclusion: Two very different phenotypes of sarcoidosis combined with PH are observed depending on the presence or absence of pulmonary fibrosis. PH is a severe complication of sarcoidosis.

Methods

Patients

This retrospective study was conducted in two centres, Avicenne and Antoine Béclère hospitals. Patients with a history of sarcoidosis and PH observed between January 1988 and December 2002 were evaluated. Complete information was obtained from hospital and referring physician medical records and reviewed by two of the authors (HN and DV). The diagnosis of sarcoidosis was established by pathological examination and compatible clinical and radiographic features. PH was defined as a mean resting pulmonary artery pressure (mPAP) of ≥25 mm Hg and normal pulmonary wedge pressure (PWP ≤12 mm Hg) during right heart catheterisation. The haemodynamic study was performed at the Antoine Béclère Hospital (MH, OS, GS) as described elsewhere. Other causes of precapillary PH including thromboembolic disease, appetite suppressant ingestion, portal hypertension, connective tissue disease, HIV infection, sickle cell disease, and congenital heart disease were excluded on the results of history, physical examination, ventilation-perfusion lung scan and/or pulmonary angiography, Doppler echocardiography, abdominal Doppler echography, HIV serology, and antinuclear antibody.

Twenty five consecutive patients with sarcoidosis and PH were reviewed. Three were excluded from the study: one patient with splenectomy had chronic thromboembolic disease, one had evidence of mixed connective tissue disease and had also taken fenfluramine, and one had portal

Abbreviations: CI, cardiac index; FEV₁, forced expiratory volume in 1 second; FVC, forced vital capacity; mRAP, mean right atrial pressure; KCO, carbon monoxide transfer coefficient; mPAP, mean pulmonary artery pressure; PaO₂, arterial oxygen tension at room air; PH, pulmonary hypertension; PVOD, pulmonary veno-occlusive disease; PVRI, pulmonary vascular resistance index; PWP, pulmonary wedge pressure; SACE, serum angiotensin converting enzyme; SvO₂, mixed venous oxygen saturation; TLC, total lung capacity; TlCO, carbon monoxide transfer factor
hypertension. The 22 studied patients underwent a standard evaluation during first right heart catheterisation including chest radiography, pulmonary function tests as recommended,20 contrast enhanced high resolution computed tomographic (HRCT) scanning, and measurement of serum angiotensin converting enzyme (SACE).

Controls
Patients with sarcoidosis and PH were compared with control patients with sarcoidosis without PH on Doppler echocardiography. Each patient was matched with two controls for sex, age, and radiographic stage. The pneumology department of Avicenne Hospital is highly specialised in sarcoidosis and has a unique and large cohort of patients with the condition. Each patient with sarcoidosis referred to the department is registered according to the date of admission, age, sex, and radiographic stage and is assigned a recruitment number according to the date of admission. Controls were selected by two steps: (1) for each patient with sarcoidosis and PH the two sarcoidosis patients whose number was immediately before and after and who had the same sex, age (± 5 years), and radiographic stage were reviewed; (2) the sarcoidosis patient was selected as a control if a Doppler echocardiogram was available in his chart and had no evidence of PH. If not, the following sarcoidosis patient according to his recruitment number was reviewed, and so forth.

Analysis of HRCT scans
The scans of the patients and controls were randomly reviewed by one chest radiologist who was blind to the diagnoses (MB). They were analysed for the presence or absence of pulmonary elementary lesions as previously described.20 21

Statistical analysis
Analysis was performed using the Statview version 5.0 statistical package (SAS Institute, Cary, NC, USA). Results are expressed as mean (SD). Groups were compared by unpaired Student’s t test and χ² test as appropriate. Simple regression analysis was used for correlations. The probability of survival of patients and controls was estimated by the Kaplan-Meier method and compared using the log rank test. Survival was calculated from PH diagnosis for patients and from normal Doppler echocardiography for controls until the end of the follow up period. Transplanted patients were considered as censored at the date of transplantation. Univariate analysis was based on the proportional hazards model. The results are expressed as hazard ratios with 95% confidence intervals.

RESULTS
Clinical findings
The study population included 22 patients (16 men and 6 women) of mean (SD) age 46 (13) years. Seventeen patients were referred because of detection of PH by Doppler echocardiography while they had increased dyspnoea; five were referred for lung transplantation and systematic right heart catheterisation evidence of PH. The patients were divided into two groups according to the absence (group A) or presence (group B) of pulmonary fibrosis on the chest radiograph. Group A consisted of seven patients (two with stage 0 disease, four with stage II, and one with stage III). Group B comprised 15 patients (all with stage IV disease).

The demographic and clinical characteristics of the patients are shown in table 1. Sarcoidosis involved at least one extrapulmonary organ in 11 of the 22 patients (skin (n = 6), liver (n = 4), peripheral adenopathy (n = 3), spleen (n = 1), articular (n = 1), eye (n = 1), and stomach (n = 1)) and one patient had hypercalcaemia. Sarcoidosis and PH were diagnosed simultaneously in one patient in group A with radiographic stage II disease. One patient in group A had symptoms of Raynaud’s disease.

Two patients had radiographic stage 0 disease at the time of PH diagnosis. The first was a 55 year old woman with known sarcoidosis affecting the skin and the stomach and hypercalcaemia. We had no information about her chest radiograph until she was clinically asymptomatic. She was placed on corticosteroids for near syncope, n (%) 0 3 (20)

Right heart failure, n (%) 1 (14.3) 4 (26.7)

Near syncope, n (%) 0 3 (20)

Mean (SD) baseline haemodynamics

<table>
<thead>
<tr>
<th></th>
<th>Group A (n = 7)</th>
<th>Group B (n = 15)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>48 (13)</td>
<td>45 (13)</td>
</tr>
<tr>
<td>M/F</td>
<td>3/4</td>
<td>13/2</td>
</tr>
<tr>
<td>White/black</td>
<td>4/3</td>
<td>13/2</td>
</tr>
<tr>
<td>Non/ex/current smokers</td>
<td>4/3/0</td>
<td>8/7/2</td>
</tr>
<tr>
<td>Mean (SD) time interval between sarcoidosis and PH (years)</td>
<td>3.7 (3.1)</td>
<td>12.5 (6.2)</td>
</tr>
<tr>
<td>NYHA functional class, n (%)</td>
<td>I 1 (14.3)</td>
<td>II 4 (57.1)</td>
</tr>
<tr>
<td>Right heart failure, n (%)</td>
<td>1 (14.3)</td>
<td>4 (26.7)</td>
</tr>
<tr>
<td>Near syncope, n (%)</td>
<td>0</td>
<td>3 (20)</td>
</tr>
<tr>
<td>Mean (SD) baseline haemodynamics</td>
<td>mRAP (mm Hg) 7.3 (4.2)</td>
<td>6.4 (6.7)</td>
</tr>
<tr>
<td>mPAP (mm Hg)</td>
<td>51.7 (16.0)</td>
<td>40.1 (11.6)</td>
</tr>
<tr>
<td>PWP (mm Hg)</td>
<td>8.8 (2.3)</td>
<td>8.1 (3.6)</td>
</tr>
<tr>
<td>Cl (l/min/m²)</td>
<td>2.45 (0.69)</td>
<td>3.26 (0.09)</td>
</tr>
<tr>
<td>PVRI (BU/m²)</td>
<td>23.0 (10.4)</td>
<td>14.6 (8.9)</td>
</tr>
<tr>
<td>SvO₂ (%)</td>
<td>63 (16)</td>
<td>68 (19)</td>
</tr>
<tr>
<td>Acute vasodilator response, n (%)</td>
<td>0 (0)</td>
<td>0 (0)</td>
</tr>
</tbody>
</table>

The controls were well matched with regard to their demographic characteristics (32 men and 12 women, mean (SD) age 46 (13) years, 75% white and 25% Afro-Caribbean) and extrapulmonary manifestations of sarcoidosis.

Haemodynamic findings
The results are shown in table 1. PH tended to be more severe in group A patients (mPAP 51.7 (16.0) mm Hg vs 40.1 (11.6) mm Hg, p = 0.07, and pulmonary vascular resistance index (PVRI) 23.0 (10.4) BU/m² vs 14.6 (8.9) BU/m², p = 0.06).
PH was moderate (mPAP <35 mm Hg) in one patient in group A (14.3%) and in seven patients in group B (46.7%), p = NS.

Functional ventilatory findings

The results are shown in table 2. Patients in group A had nearly normal spirometric parameters or only a mild restrictive defect but a severe decrease in carbon monoxide transfer factor (TLCO) and carbon monoxide transfer coefficient (KCO). Two patients (28.6%) had significant hypoxaemia of ≤55 mm Hg, one of whom had right-to-left intracardiac shunting through a patent foramen ovale. No significant difference in lung volumes and flows were found between patients in group A and controls, but TLCO, KCO, and PaO2 were significantly lower in patients with sarcoidosis and PH (table 2).

Pulmonary function was markedly changed in patients in group B (table 1). PaO2 was ≤7.3 kPa in six patients (40.0%). All parameters were significantly more reduced in patients in group B than in controls, except for total lung capacity (TLC) and FEV1/FVC which were similar (table 2).

HRCT findings

An HRCT scan was unavailable in one patient in group B. The occurrence of ground glass attenuation in patients in group A was significantly higher than in controls (85.7% v 14.3%, p<0.01). Patients in group B had a significantly higher frequency of septal lines than controls (78.6% v 46.4%, p = 0.047), particularly when polygonal reticulations were drawn (71.4% v 25%, p = 0.004; fig 1). Extrinsic compression of large pulmonary arteries by adenopathies, defined as a reduction in the vascular lumen by over 50% after injection of contrast material, did not occur in group A and was present in three patients in group B (21.4%) at the mediastinal or hilar level. In these patients extrinsic arterial compression was established by pulmonary angiography. There was no evidence of pulmonary venous compression on the HRCT scans.

Serum angiotensin converting enzyme

SACE was measured in 18 patients at the time of PH diagnosis; it was increased in three of five cases in group A and in three of 13 cases in group B.

Correlations between haemodynamic and functional ventilatory findings

In each group we attempted to correlate mPAP and PVRI with the following pulmonary function parameters (% predicted): FEV1, FVC, TLC, FEV1/FVC, TLCO, KCO, and PaO2 on room air. In group A mPAP was only correlated significantly with TLCO (r = 0.847, p = 0.03) and tended to correlate with KCO without reaching the level of significance (r = 0.762, p = 0.08). PVRI was only significantly correlated with PaO2 (r = 0.775, p = 0.04). In group B mPAP was only significantly correlated with TLCO (r = 0.684, p = 0.01) and there was a trend for MPAP to correlate with KCO (r = 0.521, p = 0.07). No relation was found between PVRI and any pulmonary function parameters.

Treatment

Fourteen patients (63.6%) were treated with long term oxygen and six received warfarin. No patient was given long term systemic vasodilator agents. One patient with radiographic stage IV disease received nebulised iloprost without any clinical benefit and died within 4 months while on the lung transplant waiting list.

Patients treated with corticosteroids

Specific treatment for sarcoidosis was instituted because of PH in 10 patients (stage 0 (n = 1), stage II (n = 4), stage IV (n = 5), all of whom were given high doses of oral prednisone ranging from 0.5 to 1 mg/kg/day combined with methotrexate 15 mg weekly in one case and with 3 monthly boluses of cyclophosphamide in another case. At the start of

<table>
<thead>
<tr>
<th>Table 2</th>
<th>Functional ventilatory characteristics of patients at the time of PH diagnosis</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Group A (n = 7)</td>
</tr>
<tr>
<td>PaO2 (kPa)</td>
<td>8.4 (2.3)</td>
</tr>
<tr>
<td>FEV1 (ml)</td>
<td>2141 (481)</td>
</tr>
<tr>
<td>FEV1 (% pred)</td>
<td>75 (20)</td>
</tr>
<tr>
<td>FEV1/FVC (%)</td>
<td>75 (8)</td>
</tr>
<tr>
<td>FVC (ml)</td>
<td>2851 (649)</td>
</tr>
<tr>
<td>FVC (% pred)</td>
<td>82 (18)</td>
</tr>
<tr>
<td>TLC (ml)</td>
<td>4688 (1022)</td>
</tr>
<tr>
<td>TLC (% pred)</td>
<td>84 (13)</td>
</tr>
<tr>
<td>TLCO (% pred)</td>
<td>39 (28)</td>
</tr>
<tr>
<td>KCO (% pred)</td>
<td>41 (29)</td>
</tr>
</tbody>
</table>

Group A, patients without pulmonary fibrosis on the chest radiograph; group B, patients with pulmonary fibrosis on the chest radiograph; PaO2, arterial oxygen tension at room air; FEV1, forced expiratory volume in 1 second; FVC, forced vital capacity; TLC, total lung capacity; TLCO, carbon monoxide transfer factor; KCO, carbon monoxide transfer coefficient.

Comparison between patients in group B and controls.

*p Comparison between patients in group A and controls.

+C Comparison between patients in group B and controls.

†TLCO and KCO were not available in one patient in group A and two patients in group B.
treatment the patients underwent right heart catheterisation and concurrent Doppler echocardiography. Systolic PAP was estimated as previously described. The echocardiographic measures of systolic PAP were in good agreement with those obtained invasively ($r = 0.967$, $p < 0.0001$). The therapeutic response was evaluated after 3 or 6 months by Doppler echocardiography performed by the same cardiologist in all cases and also by right heart catheterisation in three cases. The results are shown in table 3. Systolic PAP remained stable or increased in seven patients (two with stage II disease and five with stage IV) and decreased more than 20% from baseline in the three remaining cases (one stage 0 and two stage II). In these three patients, corticosteroids were maintained at low doses and Doppler echocardiography totally normalised at 12, 14, and 36 months, respectively.

Interestingly, before being referred to our centre, one of these three patients already had a history of sarcoidosis with PH that was reversed with corticosteroids but relapsed after the treatment was stopped (see above).

### Mortality and survival

Patients were observed for 3.7 (3.5) years. At the end of the follow up period 10 patients were alive, seven had died, and five were transplanted (five stage IV). The causes of mortality were right ventricular failure in three cases (two stage II, one stage IV), sudden death in one (stage IV), end stage respiratory insufficiency in one (stage IV), pulmonary cancer in one (stage IV), and the cause was unknown in the remaining case (stage IV). Controls were observed for 7.5 (4.6) years. Two of them died (one of pulmonary cancer and one from a surgery complication) and none was transplanted. The probability of survival of patients was significantly poorer than that of controls at 1, 2, and 3 years (84.8% vs 100%, 73.5% vs 96.4%, and 59.0% vs 96.4%, respectively; $p = 0.003$).

Univariate analysis was used to examine the relationship between survival and selected variables measured at initial right heart catheterisation. Mortality was not associated with patient age, sex, radiographic stage IV, any pulmonary function test including FEV1 % predicted, FEV1/FVC%, FVC % predicted, TLC % predicted, TLCO % predicted, and PaO2, 6 minute walk test, and any haemodynamic parameters including mRAP, mPAP, CI, and PVRI. Only NYHA functional class IV was significantly related to an increased risk of death (15.15 (95% CI 2.77 to 83.33), $p = 0.002$).

### Histopathology

Native lungs were obtained from the five transplanted patients. Pulmonary explants were examined after endobronchial formalin fixation; 22–30 tissue blocks were processed. Paraffin sections were studied with haematoxylin/eosin, elastic and iron stains. The main histological findings are summarised in table 4 and fig 2. In all cases sarcoidosis granulomas were fibrotic and scattered, and were not observed on each block. They were present in the parenchyma in all cases and in the mediastinal lymph nodes in three of the five cases. The vascular location of the granulomas was predominantly the veins (four of the five cases), while arterial granulomas were seen in only two cases and neither venous nor arterial granulomas could be found in

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**Table 3** Patients treated with corticosteroids for sarcoidosis and PH*

<table>
<thead>
<tr>
<th>Sex/age</th>
<th>Chest radiographic stage</th>
<th>Associated treatment</th>
<th>Systolic PAP Baseline</th>
<th>3–6 months</th>
<th>Last evaluation</th>
</tr>
</thead>
<tbody>
<tr>
<td>F/55</td>
<td>0</td>
<td>Methotrexate</td>
<td>66</td>
<td>35</td>
<td>&lt;30 mm Hg at 12 months</td>
</tr>
<tr>
<td>M/61</td>
<td>II</td>
<td>Oxygen, warfarin</td>
<td>121</td>
<td>125†</td>
<td>Dead at 11 months</td>
</tr>
<tr>
<td>F/52</td>
<td>II</td>
<td></td>
<td>60</td>
<td>40</td>
<td>30 mm Hg at 14 months</td>
</tr>
<tr>
<td>M/28</td>
<td>II</td>
<td></td>
<td>77</td>
<td>60†</td>
<td>30 mm Hg at 36 months</td>
</tr>
<tr>
<td>M/63</td>
<td>II</td>
<td></td>
<td>80</td>
<td>82</td>
<td>Dead at 18 months</td>
</tr>
<tr>
<td>M/55</td>
<td>IV</td>
<td>–</td>
<td>50</td>
<td>55</td>
<td>Not re-evaluated</td>
</tr>
<tr>
<td>F/62</td>
<td>IV</td>
<td>–</td>
<td>45</td>
<td>45†</td>
<td>50 mm Hg at 18 months†</td>
</tr>
<tr>
<td>M/57</td>
<td>IV</td>
<td>Oxygen</td>
<td>80</td>
<td>85</td>
<td>Transplanted at 14 months†</td>
</tr>
<tr>
<td>M/47</td>
<td>IV</td>
<td>–</td>
<td>83</td>
<td>100</td>
<td>Transplanted at 39 months†</td>
</tr>
<tr>
<td>M/42</td>
<td>IV</td>
<td>Oxygen, Cyc</td>
<td>56</td>
<td>59</td>
<td>91 mm Hg at 48 months†</td>
</tr>
</tbody>
</table>

*Cyc, intravenous cyclophosphamide.

*Whatever the treatment at the time of PH diagnosis, each patient received at least high doses of oral prednisone (0.5–1 mg/kg).

†In these patients the measure obtained by Doppler echocardiography was confirmed by right heart catheterisation.

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Figure 2. Pulmonary explant, peripheral specimen, paraffin section. (A) Homogenous fibrosis of visceral pleura, interlobular septa (arrow) and perivascular spaces (magnification ×2.5; H&E stain). (B) Interlobular septal vein with occlusive intimal fibrosis (arrow) (magnification ×20; H&E stain). (C) Alveolar haemosiderosis (arrow 1) and iron deposits on elastic laminae (arrow 2) (magnification ×20, iron stain).
one patient. Besides granulomatous involvement, occlusive venopathy consisting of occlusive intimal fibrosis and recanalisation was found in all cases (fig 2), while arterial lesions were minor with no plexiform or thrombotic lesions. All cases had evidence of chronic haemosiderosis and iron deposits on elastic laminae. Parenchymal changes were present in all cases and were homogenous within all blocks: fibrosis occurred in the subpleural space, interlobular septa and peribronchovascular sheets (fig 2).

**DISCUSSION**

This study is the first to provide a comprehensive analysis of the underlying mechanisms of PH in a series of patients with sarcoidosis and PH. The results suggest that (1) sarcoidosis and PH can have two very different phenotypes depending on the presence or absence of pulmonary fibrosis; (2) specific mechanisms of PH can be multiple and vary from case to case, particularly in the presence of fibrotic disease; and (3) sarcoidosis and PH confers a severe vital prognosis.

Surprisingly, 31.8% of our patients developed PH in the absence of pulmonary fibrosis. In these patients the changes in lung function were mild, but mean TLCO was low and two patients had severe hypoxaemia (≤7.3 kPa). Mean PAP was significantly correlated with TLCO and PVRI with PaO2. However, PaO2 and TLCO were preserved or only slightly decreased in controls without PH, which suggests that hypoxaemia was the consequence rather than the cause of PH. In addition, no other cause of secondary PH was found. A fortuitous association between sarcoidosis and idiopathic PH cannot be definitively ruled out but seems unlikely because of the low incidence of each disease. Even if there is no histological proof, our findings strongly support the role of specific pulmonary vasculopathy, which presumably also induces the severe reduction in TLCO. Only a few similar cases have been reported in the literature,7 8 10–11 21 two of whom had striking evidence of a histological pattern of granulomatous PVOD.11 12 The two most typical HRCT features of PVOD are septal lines and ground glass attenuation.24 25 Although these signs can also be seen in pulmonary sarcoidosis, our patients with non-fibrotic sarcoidosis and PH differed significantly from controls with sarcoidosis without PH in the very high frequency of ground glass attenuation (85.7% v 14.3%, p<0.01), which may possibly reflect the presence of PVOD. Interestingly, two of our patients had radiographic stage 0 disease at the time of PH diagnosis. In one previously reported case, comparisons of subsequent specimens obtained before and after established PH showed a clear progression in pulmonary vascular involvement while the parenchymal lesions of sarcoidosis remained relatively stable with corticosteroid treatment.10 Similarly, in our two stage 0 patients we speculate that a pre-existing parenchymal component either responded to corticosteroid treatment or improved spontaneously while the vascular involvement worsened.

Most of the patients with sarcoidosis and PH had radiographic stage IV disease (68.2%). Pulmonary function, particularly PaO2, was significantly more changed in these patients than in controls without PH. A recent study by Shorr and colleagues1 in a large cohort of patients with advanced sarcoidosis listed for lung transplantation showed that patients with PH have similar pulmonary function but require more supplemental oxygen than patients without PH. The discrepancy between these findings may result from the different selection methods. We failed to find a correlation between mPAP or PVRI and pulmonary function or PaO2, except mPAP with TLCO. Furthermore, mPAP was over 35 mm Hg in 53.3% of our patients, which was certainly out of proportion with the changes in lung mechanics.20

Taken together, these findings suggest that—at least in some
stage IV cases—PH is not exclusively explained by the destruction of the vascular bed or hypoxaemia and that other mechanisms may also be involved. Extrinsic compres-
sion of large pulmonary arteries by mediastinal or hilar
adenopathies was seen in 21.4% of our patients with pul-
monary fibrosis and may be responsible for PH. With
regard to the HRCT findings, our patients with fibrotic
sarcoidosis and PH (group B) significantly differed from
controls without PH in the frequency of septal lines (78.6% v
46.4%, p = 0.047), particularly when they drew a polygonal
network (71.4% v 25.0%, p = 0.004), which raised the
possibility of PVOD.

Vascular involvement is very common in pulmonary
sarcoidosis, occurring in 69–100% of cases in pathological
studies, and consists of obliterative or destructive lesions
due to the invasion of vessel walls by granulomas or to the
perivascular fibrosis. These changes can be observed at all
levels from large branches of the pulmonary arteries to small
veins, but are predominantly on the venous side. Despite
frequent vascular involvement, however, PH is rare. In ourive transplanted patients with sarcoidosis and PH, examina-
tion of the lung revealed an original intrinsic venopathy with
marked lesions of intimal fibrosis (fig 2). The role of this
venopathy and its part in the development of PH is difficult
to confirm since all these patients had advanced pulmonary
fibrosis. Nevertheless, this hypothesis is supported by the fact
that the venous lesions were occlusive, as indicated by the
associated chronic haemosiderosis (fig 2C) in the absence of
another cause of venous hypertension including proximal
venous obstruction and mitral stenosis.

PH is known to have a severe effect on outcome and to be a
predictor of mortality in patients with sarcoidosis listed for
lung transplantation. In our population survival was poor
with a rate of 59.0% at 5 years after the diagnosis of PH and
may have been overestimated since transplanted patients
were considered as censored at the date of transplantation.
Survival was significantly worse than in controls without PH.
PH was the cause of 57.1% of deaths, but haemodynamic
measures were not found to be associated with mortality.
NYHA functional class IV was the only predictor of mortality.
However, the statistical analysis was limited by the small
number of patients in the study.

These results support treating PH in patients with pulmonary
sarcoidosis, but the appropriate management of such a complica-
tion is not well defined. Little has been written about the benefits of corticosteroids in patients with sarcoidosis and PH, and the published data are somewhat discordant with some studies showing that PH can worsen despite corticosteroid treatment while others have reported a dramatic improvement. Gluskowski and colleagues studied the effects of corticosteroid treatment for 12 months on pulmonary haemodynamics in 24 patients with pulmonary sarcoidosis, three of whom had resting PH and 18 had an abnormal increase in mPAP on exercise. Regression of the radiological changes and an improvement in lung function was observed in almost all patients, but this was accompanied by an improvement in the pulmonary haemodynamics in only half of them. We found that corticosteroids had no effect in patients with pulmonary fibrosis but allowed a substantial and sustained improvement in PH in three of the five cases without pulmonary fibrosis, either alone or in association with methotrexate in one case. Interestingly, in one case with radiographic stage 0, PH recurred concomitantly with the reappearance of signs of sarcoidosis activity after corticosteroid treatment was stopped and was reversed when it was reintroduced. Pulmonary vasodilator therapy, including the use of new agents, has been successfully undertaken in patients with PH secondary to pulmonary fibrosis, but little is known about its usefulness in the context of sarcoidosis and PH.

In the small series studied by Preston and colleagues, five patients with sarcoidosis and PH received long term inhaled nitric oxide (in addition to intravenous epoprostenol in one case) and two patients received long term oral calcium channel blockers, but the results were not convincing.

Our study therefore indicates that there are two very
distinct phenotypes of sarcoidosis and PH depending on the
presence or absence of pulmonary fibrosis. Cases with
pulmonary fibrosis are not preceded by a prior history of
PH without pulmonary fibrosis. Specific vasculopathy seems
the sole mechanism of non-fibrotic sarcoidosis and PH but
gives rise to a higher level of mPAP and sometimes responds
to corticosteroid treatment. Combined with reports in the
literature, these findings suggest that vasculopathy may
differ between patients without or with pulmonary fibrosis.
On the one hand, PVOD may result from an active
granulomatous involvement of venous walls and, on the
other, from an authentic intrinsic venopathy. Interestingly,
pulmonary intrinsic venopathy has already been described in
another granulomatous disorder, pulmonary hirotiosis X.

The limitations of the study are that (1) it is a retrospective
analysis; (2) the two participant centres are highly specialised
in sarcoidosis (Avicenne Hospital) and pulmonary vascular
disease (Antoine Bécère Hospital) which creates a selection
bias and makes it impossible to evaluate the prevalence of PH
in the general sarcoidosis population; and (3) the patholog-
ical material is limited and was not available for patients
with non-fibrotic sarcoidosis and PH. However, we think it
would have been unethical and risky to perform a lung biopsy
in such cases.

We conclude that PH is a severe complication of sarcoidosis
which has two very different phenotypes. In the absence of
pulmonary fibrosis, treatment with an oral corticosteroid
should be considered to treat PH complicating sarcoidosis
since it may sometimes be efficacious. In patients with
fibrotic disease, corticosteroids seem to be inactive and
physicians may prefer to consider lung transplantation
sooner than they would have done solely on the basis of
lung function. Systemic vasodilator therapy should be used
with caution in patients with sarcoidosis and PH because of
its potential risk of precipitating pulmonary oedema in cases
of PVOD.

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