Familial pneumothoraces and bullae

G. J. GIBSON
From the Department of Medicine, Hammersmith Hospital, Du Cane Road, London W12 0HS

Gibson, G. J. (1977). Thorax, 32, 88–90. Familial pneumothoraces and bullae. The cases of three sisters who presented with spontaneous pneumothoraces are described. In two of the patients large bullae were clearly demonstrable. No recognisable associations of bullae or pneumothorax were present and there was no evidence of generalised emphysema. The cases suggest a familial predisposition to the development of bullae in otherwise apparently healthy lungs.

The classification of bullae usually depends on whether the non-bullous lung is healthy or emphysematous (Ogilvie and Catterall, 1959). Familial emphysema, such as occurs with Marfan’s syndrome (Boland and Tucker, 1964) and α1 antitrypsin deficiency (Eriksson, 1965) may be accompanied by bullae, but bullae occurring in otherwise normal lungs are thought to be acquired and degenerative (Belcher and Siddons, 1954), probably following infection (Almeyda, 1949; Ogilvie and Catterall, 1959).

I here report the cases of three sisters who presented with spontaneous pneumothorax; in two of the patients multiple bullae were present but no predisposing factors were recognisable and there was no functional evidence of widespread emphysema.

Case reports

The patients presented to the Chest Clinic at Hammersmith Hospital between 1947 and 1970 with spontaneous pneumothoraces (Table 1). None had any features of Marfan’s syndrome and blood levels of α1 antitrypsin were normal. The patients’ father had also suffered two pneumothoraces but no further details are available.

Patient 1, a non-smoker, first presented at the age of 28 with a right pneumothorax. Because of slow resolution, thoracoscopy was performed and showed a large bulla, 4 cm in diameter, arising from the upper lobe. Camphorated oil was instilled and full expansion followed; a subsequent chest radiograph was described as normal. A left-sided pneumothorax developed in 1960 and, because of failure of re-expansion after seven months, a further instillation of camphorated oil was performed with full resolution. In 1971, the patient again noticed dyspnoea and became aware of a clicking sensation in the chest. The radiograph (Fig. 1) showed probable localised pneumothorax at the right base with evidence of further bullae in the same area and also in the right upper zone and probably the left lower zone. The radiographic appearance has not changed over the subsequent five years. Pulmonary function (Table 2) shows a small vital capacity (VC) and a large residual volume (RV) (measured plethysmographically) with a slightly reduced forced expiratory ratio (FEV1/VC). The carbon monoxide transfer factor (DLCO) was well preserved in relation to the volume of functioning lung, ie, the transfer of CO per litre alveolar volume (Kco) was normal.

Patient 2 first presented with a left pneumothorax at the age of 32 when thoracoscopy showed no cysts or bullae. Resolution followed the instil-

### Table 1  Details of patients

<table>
<thead>
<tr>
<th>Patient</th>
<th>Height (cm)</th>
<th>Present age (years)</th>
<th>Age at onset (years)</th>
<th>No. of pneumothoraces</th>
<th>Smoking history</th>
<th>Evidence of bullae</th>
<th>α1 Antitrypsin (g/l) (normal 1.8–3.0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>151</td>
<td>57</td>
<td>28</td>
<td>2R 1L</td>
<td>–</td>
<td>+</td>
<td>2.6</td>
</tr>
<tr>
<td>2</td>
<td>152</td>
<td>55</td>
<td>32</td>
<td>2L 1R</td>
<td>+</td>
<td>+</td>
<td>2.3</td>
</tr>
<tr>
<td>3</td>
<td>161</td>
<td>43</td>
<td>37</td>
<td>1L</td>
<td>–</td>
<td>–</td>
<td>2.7</td>
</tr>
</tbody>
</table>

R = right; L = left.
Familial pneumothoraces and bullae

Fig. 1 Patient 1. Chest radiograph showing extensive bullae; there is probably a pneumothorax at the right base.

Table 2 Pulmonary function

<table>
<thead>
<tr>
<th>Patient</th>
<th>FEV₁</th>
<th>% Predicted</th>
<th>VC</th>
<th>% Predicted</th>
<th>FEV₁/VC</th>
<th>%</th>
<th>RV¹</th>
<th>% Predicted</th>
<th>TLC¹</th>
<th>% Predicted</th>
<th>DlCO¹</th>
<th>mmol min⁻¹ kPa⁻¹</th>
<th>% Predicted</th>
<th>Kco¹</th>
<th>mmol min⁻¹ kPa⁻¹</th>
<th>% Predicted</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>0.9</td>
<td>45</td>
<td>1.3</td>
<td>54</td>
<td>0.6</td>
<td>69</td>
<td>2.7</td>
<td>180</td>
<td>4.0</td>
<td>100</td>
<td>5.1</td>
<td>74</td>
<td>1.5</td>
<td>83</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>1.5</td>
<td>75</td>
<td>2.2</td>
<td>96</td>
<td>1.8</td>
<td>66</td>
<td>1.5</td>
<td>100</td>
<td>3.7</td>
<td>95</td>
<td>6.8</td>
<td>98</td>
<td>1.8</td>
<td>105</td>
<td></td>
<td></td>
</tr>
<tr>
<td>3</td>
<td>3.0</td>
<td>115</td>
<td>3.8</td>
<td>115</td>
<td>2.0</td>
<td>79</td>
<td>2.1</td>
<td>121</td>
<td>5.8</td>
<td>117</td>
<td>6.6</td>
<td>79</td>
<td>1.2</td>
<td>67</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

1By whole-body plethysmography (DuBois et al., 1956).
2By single-breath method (Ogilvie et al., 1957).

lation of camphorated oil. Four years later a pneumothorax occurred on the opposite side and thoracoscopy showed several thin-walled bullae arising from the middle lobe, the largest 3 cm in diameter. Resolution followed application of silver nitrate. During the next two years bullae at the left base became visible in the radiograph and a further left pneumothorax followed. A tomographic cut of the left lower zone showed extensive bullae (Fig. 2). Because of slow resolution talc was instilled into the pleural cavity and no subsequent pneumothoraces occurred. Fifteen years later the patient remains well and a bulla is still visible at the left base. Pulmonary function (Table 2) shows only mild airways obstruction.

Patient 3 presented with a left pneumothorax at the age of 37; gradual re-expansion occurred spontaneously over several months and no bullae were visible. Pulmonary function showed a mild reduction of Kco.

Discussion

The patients were unlike the typical patient with spontaneous pneumothorax in that all were female, their height was not above average, the age of onset was in the third or fourth decade, and re-expansion of the lung was usually slow. Although bullae were visualised in only two of the patients, it seems likely that each event was due to rupture of a bulla. The patients had none of the recognised familial associations of bullae or pneumothorax. Bullae with α₁ antitrypsin deficiency and Marfan’s syndrome are usually part of generalised emphysema but the relative preservation of carbon monoxide transfer factor and
absence of hyperinflation in these patients suggests virtually normal function of the non-bullous lung. The minor abnormalities of pulmonary function in patients 2 and 3 might be due to cigarette smoking; patient 1 was, however, a lifelong non-smoker and her high residual volume and slightly reduced forced expiratory ratio are probably a direct consequence of the bullae and resulting distortion of the normal anatomy.

The familial association described here suggests a predisposition to the development of bullae, and hereditary factors may be involved in their pathogenesis even in the absence of generalised emphysema.

I am grateful to Drs. P. Stradling and N. B. Pride for helpful discussion and permission to publish details of the patients.

References


Requests for reprints to: Dr. G. J. Gibson, Department of Medicine, Hammersmith Hospital, Du Cane Road, London W12 0HS.
Familial pneumothoraces and bullae.

G J Gibson

Thorax 1977 32: 88-90
doi: 10.1136/thx.32.1.88

Updated information and services can be found at:
http://thorax.bmj.com/content/32/1/88

Email alerting service

These include:
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

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/