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Familial pneumothoraces and bullae

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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 (Bolande and Tucker, 1964) and α_1 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 α_1 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 several 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 (DL_{CO}) 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 instillation

Table 1 Details of patients

Patient	Height (cm)	Present age (years)	Age at onset (years)	No. of pneumothoraces	Smoking history	Evidence of bullae	α_1 Antitrypsin (g/100 ml) (normal 1.8-3.0)
1	151	57	28	2R 1L	-	+	2.6
2	152	55	32	2L 1R	+	+	2.3
3	161	43	37	1L	+	-	2.7

R = right; L = left.

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

Table 2 Pulmonary function

Patient	FEV ₁		VC		FEV ₁ /VC %	RV ¹		TLC ¹		DLco ²		Kco ²	
	Litres	% Predicted	Litres	% Predicted		Litres	% Predicted	Litres	% Predicted	mmol min ⁻¹ kPa ⁻¹	% Predicted	mmol min ⁻¹ kPa ⁻¹ l ⁻¹	% Predicted
1	0.9	45	1.3	54	69	2.7	180	4.0	100	5.1	74	1.5	83
2	1.5	75	2.2	96	66	1.5	100	3.7	95	6.8	98	1.8	105
3	3.0	115	3.8	115	79	2.0	121	5.8	117	6.6	79	1.2	67

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 α_1 antitrypsin deficiency and Marfan's syndrome are usually part of generalised emphysema but the relative preservation of carbon monoxide transfer factor and

Fig. 2 Patient 2. Tomogram of left lower zone showing large bullae. A rim of pneumothorax is present.

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.

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