Management of primary pulmonary hypertension

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When Paul Wood first described primary pulmonary hypertension he believed it to have a vasoconstrictive origin and showed that the pulmonary vascular resistance could be selectively lowered in some but not all of these patients by infusion of acetyl choline into the pulmonary artery in contrast to patients with the Eisenmenger syndrome, who were invariably unresponsive. The pathogenesis has since been argued, microthromboembolism and thrombosis in situ having some proponents, but the concept of an increased smooth muscle tone perhaps generated by endothelial damage and platelet activity has gained ground. Efforts to find an effective orally active specific pulmonary arteriolar dilator have continued over the years but with little success, though there has been considerable recent resurgence of interest.

Observations made in beef cattle grazing at high altitude and in humans living and exercising at altitude have shown an individual and species variability suggesting a genetically determined hyperreactivity. This pulmonary hypertension is reversible on return to sea level, and the incidence of primary pulmonary hypertension is seemingly higher in populations living at high altitudes. Childhood and familial cases are well known, and an association with Raynaud’s phenomenon was early noted.

In the European epidemic of primary pulmonary hypertension, which was linked with the slimming drug aminorex fumarate, only an estimated 0.2% of individuals taking the drug developed the disease and the condition was usually reversible. Primary pulmonary hypertension has also been linked with fenfluramine, an anorectic which is still available. A concept of idiosyncratic pulmonary hypertension of possible dietary origin grew from this experience and from observations of a malignant type of pulmonary hypertension in rats who had eaten seeds of Crotalaria spectabilis which contains the alkaloid monocrotaline.

Primary pulmonary hypertension is commoner in women of childbearing age than in any other group, symptoms often develop after puberty or postpartum, and progression of pulmonary vascular disease has been associated with the use of oral contraceptives. There is some experimental basis for an association with female hormones, and the use of a non-virilising antioestrogen such as tamoxifen might even find a place in treatment.

The prognosis of primary pulmonary hypertension is grave, though a few patients remain stable without specific treatment and many survive longer than the median two years of Wood’s first cases. A clinical diagnosis can often be made, but it is essential that every patient has full cardiac and pulmonary investigations to ensure that no treatable cause of secondary pulmonary hypertension is left undiscovered.

The place of lung biopsy is controversial. It has the academic attraction of aiding prediction of likely vasodilator responsiveness as well as the possibility of uncovering a specific underlying cause. Unfortunately, transbronchial biopsy produces too small a sample, and since haemorrhage or pneumothorax would be especially hazardous in these patients an open lung biopsy is safer but it has not yet been shown to justify itself. Pulmonary veno-occlusive disease can usually be recognised clinically, although lung biopsy provides confirmation of this quickly fatal disorder. Medial hypertrophy in primary pulmonary hypertension appears to precede intimal proliferation, obliteration fibrosis, fibrinoid necrosis, and the “plexiform arteriopathy” of Wagenvoort, but the therapeutic dilemma is how to get hold of patients at a stage of medial hypertrophy before these irreversible changes occur. Symptoms are absent until the disorder is far advanced, and the appearances at necropsy are usually identical to those in the Eisenmenger syndrome. Fanciful wishful thinking suggests that if increased vasomotor tone and reactivity generate more muscle then this muscle will carry more receptors and at an earlier stage might allow a beneficial selective pulmonary vasodilator response to normally non-selective agents. A recent report describes a rare case of a...
In man in whom sublingual isoprenaline repeatedly induced a transient fall to normal of a moderately increased pulmonary artery pressure and achieved this in four separate studies over a four year period during six years of clinical benefit, a seemingly clear case of reversible vasoconstrictive pulmonary hypertension. Another reported case showed a dramatic response to acetyl choline, but three years later the pulmonary artery pressure was higher and no longer fell with acetyl choline.20 These cases, however, differ greatly from most patients with primary pulmonary hypertension, most of whom have higher pulmonary artery pressures when first seen and show at best a much less dramatic response to any drug.21-24

Therapeutic endeavour is encouraged by a few patients who have shown spontaneous regression of primary pulmonary hypertension.25-26 Two reported patients had both developed symptoms in childhood25 or adolescence,26 and one had already advanced to a stage of right ventricular failure when death would have been no surprise.25 This shows that clinically advanced disease should not necessarily be equated with irreversibility. In another personal case regression to normal from pulmonary hypertension at systemic level occurred in a woman who had been treated with an early calcium antagonist, prenylamine, prescribed to her years earlier for angina presumed to be from the right ventricle.

Vasodilator drugs can improve cardiovascular function only in patients with primary pulmonary hypertension who have a high level of reversible vasomotor tone in the pulmonary bed and, conversely, they must have a detrimental effect in patients with irreversible anatomical changes.22,27 Selective systemic vasodilatation will cause hypotension and may be fatal unless left ventricular filling increases. The right ventricular output is characteristically very low in primary pulmonary hypertension, and it may paradoxically fall further after the chosen vasodilator if the drug either reduces venous return and right ventricular filling pressure or if it has negative inotropic properties and reduces right ventricular contractile force without relieving its afterload. Thus nifedipine has been reported to cause a reduction in cardiac output associated either with a fall in right atrial pressure28 from systemic vasodilatation or with an increase in right atrial pressure associated with evidence of right ventricular failure.29 These adverse effects are not apparent when nifedipine unloads the right ventricle by pulmonary vasodilatation and the cardiac output rises.30,31 In just the same way net benefit from nifedipine has been reported in left ventricular failure.

Patients with a patent foramen ovale were early observed by Wood to have a better prognosis; they can keep the left ventricle filled and are less likely to become hypotensive or to die suddenly. In such patients a favourable response to a drug may be seen by an increase in systemic arterial oxygen content while a fall signals selective systemic vasodilatation with an increase in right to left shunting. Unlike in mitral stenosis successful reduction in pulmonary vascular resistance in primary pulmonary hypertension is not normally associated with a fall in systemic arterial saturation, caused by perfusion of underventilated lung, provided the cardiac output rises.27 Recruitment of previously unperfused vessels rather than a general relaxation of vasomotor tone may possibly account for some observed rises in cardiac output associated with a fall in both systemic and pulmonary artery pressures. This phenomenon may explain refractory hypotension and death reported with incremental dosage of a drug which had seemed to be well tolerated27 or there may be rapid attenuation of an initially favourable response.32 In some patients given a vasodilating inotropic agent such as isoprenaline the pulmonary artery pressure has fallen33,34; in others it has caused a rise in right ventricular output with a deleterious rise in pulmonary artery pressure35 associated with failure to dilate the lung vessels; calculated resistance may stay the same or actually fall, tachycardia is usual, and tremor uncomfortable.

Conflicting responses (many favourable but some adverse) have been reported from acute studies of many different vasodilators either infused into the pulmonary artery or given intravenously: acetylcholine,4 tolazoline,36 phen tolamine,37-39 isoprenaline,33,34 and diazoxide,40-43 but long term oral treatment has been disappointing. Prostacyclin suggested itself as a suitable drug for acute assessment,44-46 being a powerful relaxant of vascular smooth muscle, and it may be particularly relevant if abnormal platelet behaviour is important in the pathogenesis of primary pulmonary hypertension. Although non-selective, prostacyclin is easily titrated and can be used safely for assessing drug responsiveness. Since the vasodilator effect of hydralazine is thought to be mediated by endogenous prostaglandins its use in primary pulmonary hypertension was attractive47-49 but as with nifedipine both beneficial and serious adverse reactions have been reported. Recently, the use of diltiazem50,51 and captopril52,53 have been reported. Diltiazem has the least myocardial depressant effect of the calcium antagonists so is the most attractive for use in primary pulmonary hypertension. Enalapril with its longer action may prove preferable to captopril. New synthetic beta adrenergic agonists such as pirbuterol,54 with a high selectivity for pulmonary beta2 receptors, will no doubt be "better" than isoprenaline.

Sadly, structural obliteration of the pulmonary cir-
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culation is probably the major factor by the time most patients are seen, but even if only 5% or 10% of patients are capable of pulmonary vasodilatation every patient with primary pulmonary hypertension should be given the benefit of an inpatient, invasive, acute haemodynamic assessment. There is yet no guide to the agent of choice. Positive responders probably respond to each of the drugs, and those few more fortunate patients stand to gain a great deal from long-term treatment. There is no role for digitalis in patients with sinus rhythm, and diuretic dosage should be minimal. Oral anticoagulants should usually be given to prevent secondary thromboembolism as in other low output states. Oxygen inhalation at home aided by an oxygen condenser may be beneficial in some. Portable oxygen is helpful for patients with arterial desaturation. Patients with primary pulmonary hypertension should be screened for evidence of collagen vascular disease, but it is even more important for patients who present with symptoms of a connective tissue disorder to be investigated for pulmonary hypertension at a time when vasodilator treatment may be most rewarding; they are patients with systemic lupus erythematosus or scleroderma, particularly in the CREST syndrome or mixed connective tissue disorder and rheumatoid disease. Patients with systemic lupus erythematosus who are found to have the lupus anticoagulant should be early suspects for pulmonary hypertension; it is of interest that nifedipine is the usual drug of choice for the treatment of Raynaud's phenomenon and might be preventing the development or progression of associated pulmonary hypertension in some of these patients.

Heart and lung transplantation is the final radical treatment for pulmonary vascular disease in primary pulmonary hypertension and the Eisenmenger syndrome and may offer the only realistic hope for palliation of advanced disease.

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