High probability ventilation-perfusion scan in primary pulmonary hypertension

SAT SHARMA MD FRCPC, WILLIAM D LESLIE MD FRCPC, MORLEY LERTZMAN MD FRCPC
Sections of Respiratory and Nuclear Medicine, Department of Medicine, University of Manitoba, Winnipeg, Manitoba


The perfusion lung scan is a valuable noninvasive tool in the evaluation of patients with pulmonary arterial hypertension of undetermined cause and for the exclusion of occult large-vessel pulmonary thromboembolism. Peripheral patchy defects have been reported in primary pulmonary hypertension (PPH) but there are no well documented reports of segmental or larger perfusion defects. A case of a 55-year-old male with severe pulmonary hypertension of unknown etiology who had persistent high probability perfusion scan patterns over a period of two years is reported. No evidence of thromboembolism was present on pulmonary angiography. A discussion of the case and a review of the literature on the role of lung scan in PPH are presented. Most patients with PPH have normal or low probability perfusion scans; high probability scans occur rarely.

Key Words: Perfusion lung scan, Primary pulmonary hypertension

Scintigraphie de perfusion et de ventilation à haute probabilité dans l'hypertension pulmonaire primitive

RÉSUMÉ : La scintigraphie pulmonaire de perfusion est une étude précieuse sans effraction tissulaire qui permet d'évaluer les patients souffrant d’hypertension artérielle pulmonaire d’origine indéterminée et d’exclure une thromboembolie occulte d’un gros vaisseau pulmonaire. Un déficit de perfusion parcellaire et périphérique a été observé dans l’hypertension pulmonaire primitive (HPP) mais il n’y a pas de rapports bien documentés sur un déficit de perfusion segmentaire ou plus étendu. On rapporte le cas d’un homme de 55 ans souffrant d’une hypertension pulmonaire grave d’origine inconnue chez qui pendant plus de deux ans les scintigraphies de perfusion se sont constamment révélées à haute probabilité pour une thromboembolie. L’angiographie pulmonaire n’avait révélé aucune thromboembolie. On présente une discussion de ce cas ainsi qu’une revue de la littérature sur le rôle de la scintigraphie pulmonaire dans l’HPP. La plupart des patients souffrant d’HPP présentent des scintigraphies de perfusion normales ou à faible probabilité; les scintigraphies à haute probabilité sont rares.

EXTENSIVE EXPERIENCE OF VARIOUS AUTHORS INDICATES that the perfusion lung scan is an excellent noninvasive tool for distinguishing between pulmonary hypertension due to large-vessel chronic thromboembolism and pulmonary hypertension of undetermined cause (1-4). The pattern of abnormality seen in pulmonary hypertension of undetermined cause due to thrombotic occlusion of microvasculature, or primary pulmonary hypertension (PPH), consists of diffuse patchy peripheral defects. Perfusion defects of a lobar or segmental nature have not been well documented in PPH. The present case is reported to demonstrate that multiple segmental defects typical of pulmonary emboli may be present in a patient with nonthromboembolic pulmonary hypertension.

Correspondence and reprints: Dr S Sharma, B-1009, 409 Taché Avenue, Winnipeg, Manitoba R2H 2A6. Telephone 204-237-2217, Fax 204-231-1927
CASE PRESENTATION

A 55-year-old Caucasian male with a 10 pack-year smoking history presented with gradually progressive exertional dyspnea and a single episode of exertional syncope. He had a past history of systemic hypertension and a long history of Raynaud’s phenomenon. Physical examination revealed elevated venous pressure, right ventricular lift, right ventricular fourth heart sound and a systolic murmur of tricuspid regurgitation over the right sternal border. Electrocardiography showed typical findings of right ventricular hypertrophy. A chest radiograph showed cardiomegaly and large pulmonary arteries (Figure 1). Transthoracic echocardiogram revealed normal left ventricle size and function, and a markedly dilated right ventricle with a severe reduction in right ventricular systolic function. Severe tricuspid regurgitation was present and paradoxic motion of interventricular septum was seen. Pulmonary artery systolic pressure was estimated by transvalvular flow velocity to be in excess of 90 mmHg. A six-view technetium-99m macroaggregated albumin perfusion lung scan showed multiple, bilateral, triangular wedge-shaped segmental and subsegmental defects (Figure 2). 133Xenon ventilation scan showed mild asymmetry but no segmental ventilatory abnormalities, and therefore established the presence of multiple areas of ventilation-perfusion mismatch (Figure 2, bottom). Subsequent ventilation perfusion scans done at an interval of one week and one month were unchanged from the first study. No evidence of deep venous thrombosis in the lower extremities was found on venography. Pulmonary function studies were consistent with a moderate restrictive pattern: forced vital capacity (FVC) 1.92 L (58% predicted); forced expiratory volume in 1 s (FEV1) 1.51 L/s (56% predicted); FEV1:FVC ratio 78%; total lung capacity 61% predicted; and carbon monoxide diffusing capacity of the lung 19.4 mL/min/mmHg (80% predicted). Arterial blood gases showed pH 7.42, PaCO2 32 mmHg, PaO2 71 mmHg and bicarbonate 21 mmol/L. No mediastinal or parenchymal abnormalities were identified on an uninfused computed tomography (CT) scan of the chest. Pulmonary angiography was not performed. The patient was initially treated with intravenous heparin followed by adequate long term anticoagulation with oral warfarin therapy.

The patient returned two years later with increasing dyspnea and worsening right-sided heart failure. Medications included lisinopril, furosemide, ranitidine and warfarin.
The perfusion scan was repeated two years after the first examination. Projections are anterior (upper left), posterior (upper right), right posterior oblique (lower left) and right posterior oblique (lower right). Previously described defects are more extensive and there are new defects in the right posterior lung base. Perfusion abnormalities are peripheral and wedge-shaped, and they appear to conform to segmental and lobar boundaries. Simultaneous assessment of regional ventilation on a technetium-99m diethylenetriamine pentaacetic acid (DTPA) radioaerosol scan was normal (not shown).

Complete blood cell count, electrolytes, blood urea nitrogen, creatinine and creatine kinase were within normal ranges. Chest radiograph, electrocardiograph, pulmonary function studies and echocardiographic findings were unchanged. A perfusion scan showed multiple segmental perfusion defects in both lungs, with further deterioration in the right lung compared with the previous scans (Figure 3). The ventilation scan, performed with technetium-99m diethylenetriamine pentaacetic acid (DTPA) radioaerosol, was normal. Pulmonary artery angiography and hemodynamic measurement showed mean pulmonary artery pressure of 50 mmHg. Review of the study by two independent radiologists failed to reveal any evidence of proximal or distal vessel thromboembolic disease (Figure 4). No evidence of organized old thrombi such as arterial wall irregularity, web formation or narrowing of vessels was noted. Additional studies performed at this time showed positive serum antinuclear antibodies (ANA) in a titre of more than 1:10,000 dilutions. Erythrocyte sedimentation rate, DNA, extractable nuclear antigens and anticientromere antibodies were all negative. Positive ANA may indicate a disease process such as progressive systemic sclerosis or CREST (calcinosis cutis, Raynaud’s phenomenon, esophageal motility disorder, sclerodactyly and telangiectasia) syndrome, but no clinical or laboratory evidence of an underlying collagen vascular disease was found. A repeat CT scan of the chest failed to show any mediastinal or parenchymal abnormalities.

**DISCUSSION**

This is a well documented case of a false positive perfusion lung scan in a patient with severe pulmonary artery hypertension of undetermined origin. Our patient has either PPH or pulmonary arterial hypertension associated with collagen vascular disease limited to the pulmonary vasculature. The clinical picture and pulmonary angiogram are consistent with PPH. Perfusion lung scans are used as a guide to further evaluation of patients with pulmonary hypertension who may have chronic large vessel thromboembolism amenable to thromboendarterectomy. The value of the perfusion scan as a screening investigation before angiography has been firmly
established. D’Alonzo et al (1) reported a series of 25 patients with chronic pulmonary hypertension. All eight patients in whom thromboembolic disease was established on subsequent angiography had high probability scans, whereas scans were normal or low probability in patients with PPH. Powe et al (2) reviewed their experience with 58 patients who were evaluated for pulmonary arterial hypertension. Thirty-four patients had a definitive diagnosis made by open lung biopsy, angiography and/or autopsy. All eight patients with thromboembolism had high probability scans. None of the 13 patients with PPH had a high probability scan. Three of the 13 patients who had nonembolic secondary pulmonary hypertension had high probability scans, but they also had significant underlying thoracic disease known to alter pulmonary perfusion. Ryan et al (3) reviewed the perfusion scans in 25 consecutive patients with chronic thromboembolic pulmonary hypertension diagnosed on pulmonary angiography. All of the perfusion scans were abnormal, although they were noted to underestimate the severity of pulmonary arterial hypertension. Moser et al (4) published the largest series on the role of perfusion lung scanning as a guide to angiography in pulmonary arterial hypertension. This group reported that angiography should be performed in patients with pulmonary hypertension who have a segmental or larger perfusion defect. Their series included 64 patients with PPH and 46 patients with chronic thromboembolic pulmonary hypertension. On evaluating the perfusion lung scans of these 110 patients, no segmental defects were noted on the scans of 64 patients with PPH, whereas all 46 patients with thromboembolic hypertension had one or more defects that were segmental or larger in size.

Recently one other case of a scan-angiographic disparity was reported in which a false positive perfusion scan occurred in a patient with PPH. Worsley et al (5) retrospectively studied the records of 35 patients with PPH. One of the patients had high probability ventilation-perfusion scan but no thromboembolism on pulmonary angiography. However, this report does not include any clinical details of this case and there is no information on how the angiography was reviewed or whether the perfusion defects were segmental or persistent.

Our patient meets the criteria for PPH, although this is essentially a diagnosis of exclusion. There are no specific histopathological lesions of PPH and the diagnosis of this disease requires the close integration of clinical symptoms, laboratory investigations and histopathological findings (6,7). A full range of pulmonary hypertensive lesions including plexogenic arteriopathy, the hallmark of PPH, is present in patients with long standing pulmonary hypertension secondary to congenital and acquired cardiac conditions, collagen vascular disorders and chronic major vessel thromboembolic pulmonary hypertension (8,9). The incidence of positive ANA has been reported in 14 to 29% of cases in various registries (10,11). The persistently positive perfusion scan in our patient shows that, although most patients with PPH have normal or low probability perfusion scans, there will be rare patients who have high probability scans.

REFERENCES