Pulmonary alveolar microlithiasis

A 37-year-old patient was admitted to the hospital with severe dyspnoea and cough. Baseline investigations revealed haemoglobin of 16.2 g/dl, haematocrit of 45.8% and a total leucocyte count of 8800/mm³. Comprehensive metabolic profile and serum and urine protein electrophoresis findings were normal. Arterial blood gas analysis showed type 1 respiratory failure. Spirometry showed restrictive ventilatory disturbances (vital capacity 53%, forced expiratory volume in 1 s 91%). A posterior—anterior chest radiograph showed bilateral diffuse, nodular (‘sandstorm-like’) calcifications in both lungs (figure 1A). A high-resolution CT scan demonstrated multiple bilateral micronodulation of calcific densities throughout both lungs (figure 1B). On transbronchial biopsy, the diagnosis of massive parenchymal and alveolar calcification was confirmed. Echocardiography showed valve calcification with moderate mitral-aortic regurgitation and severe pulmonary hypertension. Abdominal ultrasonography revealed cholelithiasis and medullary nephrocalcinosis. The diagnosis of pulmonary alveolar microlithiasis (PAM) with concomitant valve calcification, cholelithiasis and idiopathic medullary nephrocalcinosis was established.

Francesco Giallauria,1 Gabriele Giallauria2
1Department of Clinical Medicine, Cardiovascular and Immunological Sciences, Cardiac Rehabilitation Unit, University of Naples “Federico II”, Naples, Italy; 2Radiodiagnostic Unit, “Martini del Villa Malta” Hospital, Sarno, Italy.

Correspondence to Dr Francesco Giallauria, Department of Clinical Medicine, Cardiovascular and Immunological Sciences, Cardiac Rehabilitation Unit, University of Naples “Federico II”, Naples, Italy; giallauria@libero.it

Competing interests None.

Patient consent Obtained.

Ethics approval This study was conducted with the approval of the ‘Martini del Villa Malta’ Hospital.

Provenance and peer review Not commissioned; externally peer reviewed.

Received 9 January 2011
Accepted 31 March 2011


Learning points

- PAM is a rare idiopathic slowly progressive disease; usually with familial association leading to alveolar deposition of calcium/phosphate microliths.1 The underlying disturbances in calcium and phosphate homeostasis have not yet been identified. A discrepancy between the paucity of symptoms and the degree of pulmonary involvement is usually reported. PAM is found worldwide, but particularly predominates in Japan, India, Turkey and Italy.2

- PAM is prevalent among family units with a high rate of consanguinity among the parents of affected individuals. It has been recently found that mutations in the SLC34A2 gene (the type Ib2 sodium phosphate co-transporter gene) are associated with PAM.3 4

- The characteristic picture of PAM on the chest radiograph shows the infiltrates as fine sand-like calcific micronodules (‘sandstorm lung’) diffusely involving both lungs, usually most marked in the middle and lower zones. CT reveals relatively symmetrical abnormalities in all patients and that calcifications were most prominent in the peripheral, mediastinal and fissural subpleural regions, and each lobe appeared surrounded by a fine dense outline, giving the overall appearance of a ‘stony lung’. The use of high-resolution CT enables the identification of parenchymal abnormalities and provides additional information about the distribution of the disease at the level of the secondary pulmonary lobe.

REFERENCES


Figure 1 Chest radiograph showing bilateral diffuse, nodular (‘sandstorm-like’) calcifications in both lungs (figure 1A). A high-resolution CT scan demonstrating multiple bilateral micronodulation of calcific densities throughout both lungs (figure 1B).