CASE STUDY

Metastatic pulmonary calcification after renal transplantation


ABSTRACT: Metastatic pulmonary calcifications, unlike dystrophic calcifications, occur in the normal healthy lung. The radiological pattern is quite specific. The disease is commonly described in chronic renal failure with calcium disorders. The prognosis is totally unpredictable.

In 1992, a 50 yr old man underwent a successful renal transplantation during the final stage of chronic renal failure. He subsequently developed asymptomatic diffuse nodular opacities, that were discovered in 1995. An open lung biopsy confirmed the diagnosis of metastatic pulmonary calcification. There was no calcium disorder in this patient.

In contrast to the benign course of pulmonary calcification in most patients, some fulminant pulmonary calcifications complicating renal transplantation or hypercalcaemia have been described. Radiographic identification of such entities is important to permit correction of calcium disorders. Otherwise, the condition is a potentially progressive and fatal cause of respiratory failure.


*Service de Pneumologie-Allergologie, and +Service de Néphrologie, Hôpital de Rangueil, Toulouse, France. **Service d’Anatomie Pathologique, and ++Service de Radiologie, Hôpital de Purpan, Toulouse, France.

Correspondence: M. Murris-Espin
Service de Pneumologie-Allergologie
Hôpital de Rangueil
31054 Toulouse
France

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Soft tissue calcification usually occurs when tissue is damaged or in the presence of calcium disorders. We report the case of a man who, following kidney transplantation, developed diffuse confluent nodular opacities, visible on chest radiography, without any clinical symptoms. The computed tomography (CT) scan revealed the presence of calcifications.

Case report

A 50 yr old man developed progressive renal failure as a result of nephropathic interstitial disease. In May 1992, he underwent a thyroparathyroidectomy for secondary hyperparathyroidism. The surgical specimen demonstrated features consistent with intracapsular adenocarcinoma. He received a cadaveric renal transplant in November 1992, after 12 yrs of chronic haemodialysis. In the early post-transplant period, he showed slightly elevated parathormone levels, i.e. 135 ng·mL⁻¹ in May 1993, 118 ng·mL⁻¹ in November 1993, and 67 ng·mL⁻¹ in September 1994. Neither calcium nor phosphorus serum abnormalities occurred in the early or later post-transplant phases. Chest radiographs were normal. The patient was maintained on daily doses of cyclosporin (200 mg), prednisolone (15 mg), isradipine (10 mg), atenolol and L-thyroxine. In March 1995, the chest radiograph revealed numerous diffuse confluent nodular opacities, 5–12 mm in diameter, with poorly-defined margins, predominantly in the upper lung zone (fig. 1). The patient was asymptomatic and the clinical examination was normal. He had no fever, pain, coughing or dyspnoea.

On the CT scan, the nodules appeared fluffy. Calcifications were observed in some of the nodules (fig. 2). The chest radiograph had been normal prior to transplantation.

The following serum levels were recorded (normal values in parentheses): calcium 9.2 mg·dL⁻¹ (9.0–10.2
Metastatic pulmonary calcification is characterized by diffuse calcium salt deposition, whether localized in apical zones or throughout the lungs. It may be associated with myocardial calcification in chronic renal disease.

Although calcium is merely interstitial in location, the opacities observed radiographically may simulate airspace disease, such as pulmonary oedema or infarction (parenchymal opacification, poorly-defined infiltrates). Numerous nodular opacities measuring 2–12 mm in diameter are usually observed. Calcification is rarely identified on a conventional chest radiograph, and this low detectability may be related to the small size of the calcium deposits and the high kilovoltage radiographic technique [1]. The pulmonary nodules are more clearly defined on CT scan. They may be unilateral or diffuse but predominate in the upper zones. Calcifications of vessels of the chest wall are better seen on the mediastinal windows. Other findings on the CT scan include ground-glass attenuation and patchy consolidation [2]. Sometimes, only high resolution sections of CT scan disclose the calcified nature of the nodules [2]. A roentgenographic technique known as dual energy digital chest radiography may be useful in detecting lung calcifications [3]. Visible calcifications may remain stable for many years or may progress rapidly.

Metastatic calcifications of vital parenchyma are related to chronically elevated calcium levels as in chronic renal failure, primary hyperparathyroidism, D hypervitaminosis, milk alkali syndrome [2], or diffuse myelomatosis. Calcifications are usually composed of calcium, phosphate and magnesium. The cause of deposition of calcium salt in the body tissue is not fully understood.

Metastatic calcifications occur in disorders of calcium and phosphorus metabolism, and are most likely to develop when the calcium-phosphorus product exceeds 70, but may occur with normal levels [4]. In the case of chronic renal failure with secondary hyperparathyroidism, a significant association has been found between lung calcium content, elevated phosphate level and the calcium-phosphate product. No correlation has been observed with levels of parathormone, calcium, creatinine, protein, alkaline phosphatase or albumin, or with a history of parathyroidectomy. Moreover, no correlation was found with the duration of haemodialysis or with the type of dialysate [3].

Occasionally, metastatic lung calcification has been observed in the absence of supersaturation of extracellular fluid with calcium and phosphate ions [5]. Metastatic pulmonary calcifications have been shown to be present in 60–80% of autopsied haemodialysis patients, although rarely recognized during life [6], their extent is usually reflected in the survival of the patient. Although many patients remain asymptomatic, some develop severe respiratory symptoms which are related to the extent of calcification. Restrictive lung diseases with progressive decrease in TL,CO, VC and hypoxaemia may be associated with rising levels of pulmonary calcium levels [3].

The predilection of metastatic calcification for the upper lung area may be due to the high ventilation/perfusion ratio in this region, creating high oxygen and low carbon dioxide levels. Consequently relative alkalinity....
favours deposition of calcium salt. Normally, pH at the apex of the lung reaches roughly 7.51, whilst at the base it is 7.39 [7, 8].

Pathologically, lung metastatic calcification is preferentially located in the alveolar septa. Nevertheless, it can occur in pulmonary arteries and bronchial walls. In mild cases, calcium deposits may be present along the alveolar epithelial basement membrane and in the alveolar capillary walls without significant desmoplasia or septal thickening. Sometimes a foreign body giant cell reaction to the calcium can be found. Severe calcification tends to be associated with interstitial fibrosis [6].

The main differential diagnosis is lung dystrophic calcification. Dystrophic calcification consists of calcium deposition in dead cells or tissue damaged by infections (tuberculosis, fungal infections, chickenpox pneumonitis), silicosis, sarcoidosis, metastatic tumour, rheumatic mitral stenosis, microlithiasis and broncholithiasis or extensive bone malignancy [9]. The roentgenographic images are usually specific: and calcispherites of alveolar microlithiasis; popcorn ball of hamartoma, etc.

The present case report is of interest for several reasons. To our knowledge, this is the second report of a progressive pulmonary calcification occurring in a transplant recipient with a normally functioning renal transplant [1]. The precise mechanism of pulmonary calcification remains unknown. As in the first description, this patient showed no increase in calcaemia, or phosphoraemia. The case confirms that chest radiography is not a sensitive tool for detecting pulmonary calcifications. In contrast to the benign course of pulmonary calcification in most patients, some fulminant pulmonary calcifications complicating renal transplantation or hypercalcaemia have been described. It is important to recognize such entities so as to correctly identify this potentially progressive and fatal cause of respiratory failure [10, 11].

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References