Dear Editor,

We read with great interest the recent letter to the editor by Cadiñanos Loidi et al. which described a 45-year-old woman who presented with endobronchial tuberculosis and cutaneous and lymph node involvement. Although tuberculosis is prevalent worldwide, certain unusual presentations can make the etiologic diagnosis challenging. We would like to highlight the imaging findings of an uncommon presentation of thoracic tuberculosis in a case we recently encountered.

A 67-year-old man presented with a subcutaneous nodule in the sternal region with no inflammatory signs and progressive growth noted 2 months previously. One month later, the lesion exuded an odorless white secretion, and similar lesions in the axillary and clavicular regions also showed signs of fistulization. The patient reported daily intermittent fever during the evening, asthma and weight loss. Physical examination showed bilateral cervical, supraclavicular and axillary lymphadenopathy with fistulization. Laboratory test results were normal and an HIV test was negative. A tuberculin skin test was positive, with 8 mm induration.

Chest CT showed fluid collection in the soft tissues of the right anterior chest wall, adjacent to the ipsilateral chondrosternal joint, with cortical thickening of the sternal body (Fig. 1A–C). The lungs were normal. Sagittal T2-weighted MRI with fat saturation showed bone marrow edema in the sternal (Fig. 1D). A biopsy of the sternal lesion showed chronic, granulomatous, supplicative inflammation with caseous necrosis. Sternal fluid secretion and biopsy material were positive for *Mycobacterium tuberculosis*. The bacteria were sensitive to anti-tuberculosis drugs. The patient was given a four-drug (isoniazid, pyrazinamide, ethambutol and rifampin) anti-tuberculosis treatment for 2 months (intensive care phase) and isoniazid and rifampin for 4 months (maintenance phase), and achieved complete clinical recovery. One year later the patient was asymptomatic.

Mycobacterial infections of the sternum are very rare and usually develop secondary to sternotomy, Bacillus Calmette-Guérin vaccination, or immunosuppression. Complications of tuberculosis sternal osteomyelitis include secondary infection, fistula formation, spontaneous sternal fracture, tracheal compression, and rupture of tuberculous abscess into the mediastinum, pleural cavity, or subcutaneous tissues.

On standard X-rays and CT images, sternal changes may be minimal and unassociated with significant bone destruction. In this situation, it may be difficult to differentiate between osteomyelitis and inflammatory soft tissue lesions. Cortical thickening may merely be a periosteal reaction secondary to the inflammation of surrounding soft tissue. In these cases, MRI may provide valuable diagnostic information by showing inflammatory changes in sternal bone marrow characterized by low and variable signal intensity on T1- and T2-weighted images, respectively. Indeed, some associated features suggestive of tuberculosis, including fistulae, heterogeneous soft-tissue masses and sternoclavicular joint abnormalities, are valuable for diagnosis. However, no radiographic (CT or MRI) finding is pathognomonic for sternal tuberculosis. Thus, definitive distinction between tuberculous sternal osteomyelitis and other causes of sternal infection rests largely on histopathological and/or microbiological examination.

The majority of authors have argued that standard antituberculous chemotherapy is sufficient to treat sternal tuberculosis, although some believe that surgical debridement should be utilized. No consensus guideline regarding treatment regimen and duration has been developed. Aspiration and anti-tuberculosis chemotherapy are the treatments of choice. Close follow-up is essential to detect complications that may necessitate surgery. Surgical treatment is advised when removal of a large sequestrum is necessary, when the diagnosis is doubtful, and in non-responding cases. In conclusion, MRI may be useful for the diagnosis of sternal tuberculosis.

**Conflicts of Interest**

The authors declare that they have no conflicts of interest.

**References**

Sandblasting in the Naval Industry: Another Life-Threatening Activity Related to Silicosis

Pulido con arena en la industria naval: otra actividad potencialmente letal relacionada con la silicosis

Dear Editor,

We read with great interest the case reported by Bueno Palomino,¹ which alerted pulmonologists to the occupational risks faced by sandblasting.

We would like to highlight the risks of a similar activity: sandblasting used in heavy industry, especially in shipbuilding to clean boilers and ship hulls. Abrasive blasting involves the projection of abrasive particles against a surface (often glass or metal) using compressed air. In 1949, the United Kingdom adopted regulations for blasting activities that severely restricted the use of abrasive products containing free silica. In 1992, the National Institute for Occupational Safety and Health recommended that the use of sand in abrasive blasting should be banned in the United States.

Silicosis in sandblasting workers in the shipbuilding industry represents the most aggressive form of the disease, with evidence of progressive massive fibrosis in about 40% of cases.² Cases of silicoproteinosis have been reported in this group of patients.

Silicoproteinosis may develop after a relatively short period of heavy exposure to fine particulate silica (e.g., in sandblasting activities). The disease often progresses rapidly, manifesting within months or a few years after initial exposure. Symptoms include progressive shortness of breath that invariably leads to acute respiratory failure. Prognosis is very poor and most reported cases die within months. The pathological features of silicoproteinosis differ from those of chronic silicosis and resemble those of primary alveolar proteinosis. No effective treatment has been described and management consists only of supportive care.²–⁴

The high-resolution CT findings of silicoproteinosis consist of bilateral air-space disease manifesting as consolidation, ground-glass opacities, and centrilobular nodules. Punctate calcifications superimposed in areas of consolidation and calcified lymph nodes are commonly seen.³,⁴ (Fig. 1).

Although it is difficult to envision the continued occurrence of silicoproteinosis in modern society, sporadic cases continue to be reported, especially in underdeveloped countries. Radiologists and pulmonologists should be aware of the clinical presentation and imaging patterns of this serious and progressive disease, which can rapidly lead to fatal outcomes after the onset of symptoms.

Conflicts of Interest

The authors declare that they have no conflicts of interest.

References


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