Chronic eosinophilic pneumonia secondary to long-term use of nitrofurantoin: high-resolution computed tomography findings*

Pneumonia eosinofílica crônica secundária ao uso prolongado de nitrofurantoína: achados da tomografia computadorizada de alta resolução do tórax

Rosane Rodrigues Martins1, Edson Marchiori2, Sérgio Lopes Viana3, Luiz Sérgio Pereira Grillo Júnior4, Vera Luiza Capelozzi5, Laércio Moreira Valença6

Abstract
The authors report the case of a female patient who developed chronic eosinophilic pneumonia secondary to long-term use of nitrofurantoin for prophylaxis of recurrent urinary tract infections due to urethral stenosis. On high-resolution computed tomography scans, the pulmonary reaction to nitrofurantoin most commonly manifests as an interstitial-alveolar pattern in both lung bases. However, in this case, the alterations were most pronounced in the periphery of the upper lobes. In itself, this tomographic profile is strongly indicative of chronic eosinophilic pneumonia. The patient had previously been submitted to an open lung biopsy. The diagnosis of chronic eosinophilic pneumonia was confirmed through a review of the biopsy.

Keywords: Pneumonia; Pulmonary eosinophilia; Nitrofurantoin/adverse effects; Tomography, X-ray computed.

Introduction
Nitrofurantoin has been widely used as an efficient and relatively safe medication in the treatment of genitourinary infections, and complications due to its use are rare. However, pulmonary parenchymal lesions have occasionally been associated with long-term nitrofurantoin therapy. The authors report the case of a patient with chronic eosinophilic pneumonia secondary to long-term nitrofurantoin use, with emphasis on the findings obtained through high-resolution computed tomography (HRCT) scans of the chest.

Case report
In September of 2003, a 74-year-old female patient sought treatment at our facility, reporting a long history (years) of chronic cough with minimal clearance of secretion. The patient reported having undergone a chest X-ray, which had shown ‘pulmonary fibrosis’, twenty months prior. Pulmonary function tests revealed a pattern of mild restriction (79% total lung capacity) and decreased diffusing capacity of the lung for carbon monoxide (63% of predicted). The initial treatment with oral and inhaled corti-
corticosteroid therapy previously performed in another facility was unsuccessful, and an open-lung biopsy was then performed, resulting in a histopathological diagnosis of diffuse interstitial fibrosis.

In the pathological history, the report of uterine cancer at the age of 35, which was treated with radiotherapy and chemotherapy, was worthy of note. As a complication of that condition, the patient presented urethral stenosis, with recurrent episodes of urinary tract infection. During the preceding three years, the management of the urinary tract infection had included self-catheterization every 4 h and prophylaxis with nitrofurantoin. The physical examination revealed only a scar from a left thoracotomy, decreased chest expansion, and crackling rales in the lung bases.

The review of the lung biopsy by a pathologist resulted in a histological diagnosis of chronic eosinophilic pneumonia, possibly attenuated by previous use of steroids, accompanied by alterations resulting from bronchial obstruction. There was infiltration by eosinophils and mononuclear cells throughout the alveolar septa and alveolar spaces (Figure 1).

Since the anatomopathological findings might be associated with a reaction to nitrofurantoin, this medication was discontinued, and oral corticosteroid therapy was initiated.

A chest X-ray performed four months after the initial medical visit showed areas of subpleural interstitial thickening in the upper lobes. An HRCT scan, performed concomitantly, revealed focal subpleural thickening of the intralobular and interlobular interstitium accompanied by faint ground-glass opacities, these alterations being present only in the upper lobes, and alveolar opacity being observed in the anterior segment of the left upper lobe (Figure 2). There were no mediastinal alterations. A control HRCT scan performed four months later showed no significant changes, except for an improvement in the area of alveolar opacity in the left upper lobe (Figure 3).

Dyspnea and rales progressively resolved over the course of the treatment, and the dose of oral corticosteroid was progressively reduced until the medication was completely discontinued, after fifteen months of treatment. After the corticosteroid therapy had been discontinued, there was no recurrence of symptoms.

**Discussion**

Nitrofurantoin pulmonary toxicity was first reported in 1957 by Fisk. Since then, there have been various reports of nitrofurantoin-induced pulmonary reactions, with polymeric profiles that include pulmonary fibrosis, alveolar hemorrhage, bronchiolitis obliterans organizing pneumonia (BOOP), lupus-like syndrome, desquamative, usual, and nonspecific interstitial pneumonias, allergic reactions, diffuse alveolar damage, and eosinophilic pneumonia.

Nitrofurantoin-induced pulmonary disease has two principal forms of presentation: the acute form, which is more common, develops hours to days after the use of the medication, and is probably related to a hypersensitivity reaction or to an immunolog-
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In the case presented here, the histopathological diagnosis was chronic eosinophilic pneumonia, since there have been no specific reports of this disease being diagnosed (based on HRCT findings) as an adverse reaction to nitrofurantoin. The principal chronic alterations associated with nitrofurantoin are usually found in the lung bases, although sparse peribronchovascular alterations have been described in patients presenting BOOP as a reaction to nitrofurantoin administration. There can be an overlap between the clinical, radiological, and pathological findings of chronic eosinophilic pneumonia and those of BOOP. However, in many cases, HRCT helps to differentiate between these two diseases.

The acute form is clinically characterized by tachypnea, fever, nonproductive cough, tachycardia, chest pain, skin rash, and arthralgia. Chest X-rays show a reticular or alveolar pattern in both lung bases, with septal lines and pleural effusion, mimicking pulmonary edema. Peripheral eosinophilia and reduced diffusing capacity of the lung for carbon monoxide are typical, and pulmonary function test results can include hypoxemia and a restrictive pattern. The differential diagnosis should primarily include pulmonary embolism, infection, myocardial infarction, and heart failure. In addition, the temporal relationship between the onset of the symptoms and the initiation of the medication, as well as the fact that there is significant improvement after discontinuation of the medication, are fundamental.

The chronic form has a more insidious onset, with progressive dyspnea, dry cough, tachypnea, and no fever. The most common finding on chest X-rays is interstitial thickening in the lung bases. Pulmonary function tests reveal a restrictive pattern, with hypoxemia and reduced diffusing capacity of the lung for carbon monoxide. The differential diagnosis includes idiopathic pulmonary fibrosis and other causes of interstitial thickening in the lung bases. Within a few months after discontinuation of the drug, symptoms typically improve, with or without regression of the interstitial thickening.

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Figure 2 - High-resolution computed tomography scan of the chest performed in January 2004. Slices slightly above and at the level of the aortic arch: a) ground-glass opacities accompanied by subpleural interstitial thickening in the upper lobes; and b) small area of subpleural alveolar opacity in the anterior segment of the left upper lobe.
nitrofurantoin for prolonged periods be submitted to X-rays every six months as a tool for the early diagnosis of pulmonary alterations.

References