Introduction

Yellow nail syndrome (YNS) has been considered a rare condition; however, at least in part, it seems to be under detected or underreported. YNS was first described by Samman and White in 1964. They reported this entity in a study on 13 patients, and most of them presented ankle edema and nails with slow growth. Although the classical triad of YNS includes yellowish discolored nails, lymphedema and pleural effusion, two of these changes are enough to establish the diagnosis. The etiology of YNS is not well clear, and the pathogenesis involves changes in lymphatic drainage. Lymphedema is the initial manifestation of YNS in approximately one-third of cases. The purpose of this study is to describe the syndrome in a woman of the oldest-old age group with renal failure.

Case Report

A 94-year-old woman, with antecedent of chronic bronchitis, bronchiectasis, recurrent pneumonitis, arterial hypertension and chronic renal failure was admitted to control an episode of cardiac and respiratory insufficiency. Yellow nail changes and a tendency to pincer nails developed in her hand and toe fingers, preceded by longstanding course of respiratory diseases with pleural involvement. Laboratory tests detected moderate anemia and mildly elevated levels of urea and creatinine, thyroid function was normal. This case study is about yellow nail syndrome in the absence of ankle lymphedema, and affecting a woman of the oldest-old age group with renal failure.
1.15 ng/dL, and negative blood cultures. Arterial gasometry determinations showed pH: 7.48, HCO₃: 22.6 mmol/L, base excess: –0.5, PaCO₂: 30.3 mmHg, PaO₂: 35.2 mmHg, SaO₂: 68.4%, and anion gap: 4.3 mmol/L. Chest CT scan showed septal thickening, fibrosis, bronchiectasis, honeycombing, and small pulmonary nodules, in addition to basal areas with pleural thickening (Figure 2). Diagnoses of YNS and renal failure were based on clinical and complementary data. The patient was clinically managed with antibiotics to control respiratory infection. After hospital discharge, the patient was referred to Pneumology outpatient follow-up.

Discussion

The complete triad of YNS is reported in about one-third of patients; yellow nails occur in 89% of cases, lymphedema in 80%, and pulmonary or pleural changes in 63%. More frequently, pleural effusions are the last component of the syndrome to appear. Nevertheless, chronic rhinosinusitis, bronchitis, bronchiectasis, repeated pneumonitis, restrictive or obstructive pulmonary diseases, and pleural sequelae are often reported. Letheulle et al., reported five males with YNS, aged from 52 to 71 years, and median age of 56 years. Chronic sinusitis occurred in all patients; three had the triad and two did not have ankle lymphedema, but facial edema was observed. Nordkild et al., reviewed 97 patients with YNS, most of them in early middle age, and with male/female ratio of 1/1.6; yellow nails occurred in 99%, lymphedema in 80%, and respiratory changes in 63% of cases. Valdés et al., reviewed studies of 150 patients with diagnosis of YNS in a period of approximately 50 years, with male/female ratio of 1.2/1, median age of 60 years and from birth to 88 years; 78% of patients were 41 to 80 years old. YNS and pleural effusions occurred mainly between the fifth to eighth decades of life, often associated with lymphedema. Although lymphedema was found in all the individuals with yellow nails plus pleural effusions, only 85.6% of the total group of patients presented with yellow nails. Bilateral pleural effusions occurred in 68.3% of the cases; moreover, the fluid was serous in 75.3%, milky in 22.3% and purulent in 3.5%. Pericardial effusions have been also described in YNS, and drainage may be needed. Maldonado et al., analyzed 41 patients with YNS, 20 males and 21 females, with median age of 61 years and range from 18 to 82 years. Forty patients had chronic respiratory changes - pleural effusions (46%), bronchiectasis (44%), chronic sinusitis (41%), and recurrent pneumonitis (22%). Interestingly, 15 patients (37%) had no lymphedema.

In the patient herein described, non-pitting edema was observed exclusively in the upper extremities, involving all the hands fingers. This finding is not common in YNS, although some authors have described

Figure 1. Swollen hands with indicative features of lymphedema, in association with nail changes typically found in cases of yellow nail syndrome - inspissations, longitudinal ridging, hyper curved nail plates and yellowish discoloration in toe fingers; deficient lunula is more evident in the hand fingers. Worthy of note is the moderate edema involving the hand fingers, and the absence of distal edema in the lower limbs.

Figure 2. Computed tomography scan showing septal thickening, areas of fibrosis and multiple bronchiectasis, sparse foci of honeycombing at the periphery of the lungs, and small pulmonary nodules, in addition to basal areas with pleural thickening.
lymphedema uniquely affecting the face. Chronic rheumatoid arthritis has been reported in association with the occurrence of YNS. Worthy of note, this patient had significant sequels of osteoarthritis in the upper extremities, which could have played a role in the origin of the well localized edema.

Conclusions

Incomplete triad of YNS may pose diagnostic challenges; however, higher index of suspicion about the main features of the syndrome lead to more accurate evaluation. The concomitance between hyper curved nails (pincer nails) and the yellowish discolored nails of the syndrome may follow either undiagnosed or scarcely reported.

References