Pulmonary Infection Due to Nocardia nova

To the editor: The genus Nocardia includes various species that have been implicated in human infection. *N. asteroides* is the species most frequently isolated in patients with infection by Nocardia. During a recent taxonomical review, 2 new species were added to this genus: *N. farcinica* and *N. nova*. Below we describe a case of pulmonary disease due to *N. nova* which contributes to the small body of literature on the role of this microorganism in pulmonary infections.

The patient was a 78-year-old man with a personal history of rheumatic polymyalgia undergoing treatment with 10 mg/day of oral corticosteroids for more than 9 months. He also met clinical criteria for chronic bronchitis, with grade II/IV dyspnea in stable condition. He came to our hospital complaining of dyspnea with minimal effort which had steadily increased over the 4 weeks prior to admission. He also showed signs of orthopnea and paroxysmal nighttime dyspnea. Up to 90 mg/day of oral prednisone was prescribed to treat the increase in dyspnea and the rheumatic polymyalgia. The patient reported having cough and purulent sputum as well as a fever of around 38 ºC on the days prior to admission. On physical examination the patient was alert and oriented; his blood pressure was 160/90 mm Hg and temperature was 38 ºC. Dependent edema with pitting was noted along the full length of the lower extremities (including the groin). Basal crackles on both sides, best heard over the lower third of the right hemithorax, were detected on pulmonary auscultation. Respiratory rate was 24 breaths/min. Arterial blood gas analysis showed an arterial oxygen pressure of 35 mm Hg, an arterial carbon dioxide pressure of 30 mm Hg, a pH of 7.44, and oxygen saturation at 89%. A chest x-ray showed satisfactory improvement. He was afebrile and chest x-rays revealed satisfactory improvement.

*Figures.* Profile of bands obtained by polymerase chain reaction/restriction fragment length polymorphism of gene hsp65 using Msp1 and Hinf1 restriction enzymes. Lane 1: molecular weight marker VIII (Roche); lane 2: restriction fragment using Hinf1 enzyme; lane 3: restriction fragment using Msp1 enzyme.

*LETTERS TO THE EDITOR*


M.J. Unzaga, A. Gaafar, and R. Cisterna

Departamento de Microbiología, Hospital de Basurto, Bilbao, Spain.