Pulmonary infection with *Nocardia caviae* in a patient with diabetes mellitus and liver cirrhosis

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**Abstract** The first reported case of pulmonary infection with *N. caviae* in Britain occurred in a 67 year old woman, in whom diabetes mellitus and liver cirrhosis probably contributed to impaired cell mediated immunity. She was successfully treated with gentamicin 60 mg eight hourly and sulphadimidine 500 mg six hourly for six months and then with sulphadimidine 2 g/day for a further three months. When *Nocardia caviae* is isolated from the sputum of a patient with chronic respiratory infection and fever, effective treatment should be started promptly.

Nocardia organisms are Gram positive, partially acid fast, branching actinomycetes which uncommonly cause illness in man. Of a total of 17 cases of nocardiosis reported over the last three years in the United Kingdom (1984, 1985, and 1986), 12 were due to *N. asteroides* and the remainder due to untyped *Nocardia* species (Public Health Laboratory Service, unpublished). We report the first case in a patient with infection due to *N. caviae*, an organism first isolated from an infected ear of a guinea pig by Snijders in 1924.1

**Case report**

The patient is a 65 year old Pakistani woman with diabetes mellitus, who as a strict Muslim neither drinks alcohol nor smokes cigarettes; she has been resident in Britain for 12 years. She gave a two year history of intermittent fever, cough productive of persistently green sputum, malaise, and weight loss. Numerous sputum cultures carried out at the referring hospital had been negative for *Mycobacterium tuberculosis* and other mycobacteria, and the chest radiograph had shown patchy consolidation in both lung fields, with relative sparing of the apices (figure). She had had antituberculosis chemotherapy for almost one year and oral corticosteroids empirically for two to three months, without any clinical or radiological improvement. Examination after referral showed an obviously unwell, febrile, emaciated woman. She was breathless and had sinus tachycardia and cracks at the left base. She had abdominal swelling due to ascites but no apparent hepatosplenomegaly. Neurological examination at this stage showed nothing abnormal.

Investigations showed: haemoglobin 11·2 g/dl, white cells 14·8 × 10⁹/l (80% neutrophils), erythrocyte sedimentation rate 66 mm in one hour, and normal urea and electrolyte concentrations apart from sodium 127 mmol/l. No malarial parasites were seen on the blood film. Liver function was deranged: serum bilirubin 50 (normal 1–20) μmol/l, alkaline phosphatase 1050 (35–105) IU/l (almost all hepatic in origin), alanine aminotransferase 77 (0–35) IU/l, albumin 13 (33–48) g/l, total protein 63 (60–80) g/l. The prothrombin ratio was 1·8. No markers for hepatitis B infection were detected. Serum amylase activity was 115 (70–300) IU/l. Initial blood and stool cultures showed nothing remarkable. Abdominal ultrasound examination indicated that the liver was of abnormal texture and free fluid was present in the peritoneal cavity. An ascitic tap showed the fluid to be a transudate. A review of the histological appearances of the previous liver biopsy specimen showed well advanced micronodular cirrhosis.

The first indication that the patient had a nocardial infection was the finding of fine Gram positive branching filaments on a Gram stain of sputum. After five days of culture two colony types were identified, one white and the other off white, which were confirmed as *N. caviae* by the Mycological Reference Laboratory of the Central Public Health Laboratory. Antibiotic sensitivity testing was performed by a disc diffusion method, with DST agar (Oxoid Ltd) enriched with 5% lysed horse blood, the Oxford strain of *Staphylococcus aureus* being used as the control organism. The organism was sensitive to gentamicin and sulphadimidine but resistant to trimethoprim, rifampicin, and all the cephalosporins tested. The patient was therefore given intravenous gentamicin 60 mg eight hourly and oral sulphadimidine 500 mg six hourly. Within one week of her starting treatment the pyrexia had subsided, the white cell count had decreased to 9·2 × 10⁹/l and the patient felt somewhat better. Sputum cultures for *N. caviae* were repeatedly positive for two months, though the organism ceased to appear in the green stained sputum within three weeks of the start of effective treatment. During her recovery three main problems were encountered: firstly, unsteadiness of gait, which computed tomography showed to be due to a left cortical infarct, and from which she made a reasonable recovery; secondly, hypocalcaemic tetany (serum calcium 1·69 mmol/l), which was resistant to calcium supplements; and, thirdly, severe paraesthesia affecting the hands and feet. The serum magnesium concentration was 0·3 mmol/l and when it was restored to normal with magnesium gluconate supplements both the tetany and the paraesthesiae disappeared. Mild impairment of control of diabetes mellitus warranted temporary insulin treatment before she reverted to oral hypoglycaemic drugs.

The patient was treated with gentamicin and sulphadi-
midine for six months\(^4\) and then with sulphadimidine 2 g/day for a further three. Nocardia organisms were grown from her sputum two months after the start of treatment. The ascites resolved with treatment with spironolactone. A repeat chest radiograph six months later (figure, B) shows substantial resolution of the patchy consolidation.

Both the serum magnesium and the serum calcium concentrations have remained normal since the patient stopped gentamicin treatment.

**Discussion**

This is the first reported case of pulmonary infection with *Noccardia caviae* in Britain. As is often the case, it occurred in an immunocompromised setting, both diabetes mellitus and liver cirrhosis probably contributing to impaired cell mediated immunity.\(^3\) Disseminated infection due to this organism is invariably fatal.\(^3\)

The isolation of *Noccardia caviae* from the sputum in a clinical setting of chronic respiratory infection and pyrexia is highly significant, and should prompt effective treatment.

Occasionally, however, the organism occurs as an airborne contaminant, and occasionally it is isolated from sputum in the absence of any overt pulmonary disease.

Sulphonamides are the drugs of choice for treatment of all forms of nocardiosis. Trimethoprim is reported to act synergistically with these agents\(^5\) but this has not yet been clearly established. In this particular case antibiotic sensitivity testing indicated that the organism was resistant to trimethoprim but sensitive to gentamicin, so gentamicin was used in combination with sulphadimidine.

The metabolic complications seen in this patient are presumed to be secondary to gentamicin induced renal magnesium loss, previously reported with total doses exceeding 10 grams.\(^6\) Our patient had been given a total dose of 21 g by the time hypomagnesaemia and hypocalcaemia were observed. Impaired parathyroid hormone secretion is thought to provide a mechanism for this metabolic disturbance, but peripheral resistance to the action of this hormone also plays a part.\(^7\) Both serum magnesium and serum calcium concentrations have remained normal in this patient after she stopped the gentamicin.

Unfortunately there are no characteristic radiological changes associated with pulmonary nocardiosis, which can appear as consolidation, nodules (often cavitating), interstitial changes, pleural effusion, and lymphadenopathy.\(^8\)

In summary, this case illustrates that direct Gram staining of sputum may provide a clue to the aetiology of chronic pulmonary infection where tuberculosis appears to have been ruled out. Organisms such as actinomycetes are also likely to be missed by routine culture methods owing to their slow growth, as cultures are usually not continued for longer than 48 hours and anaerobic cultures are not usually performed unless there is a specific clinical indication.

**References**

1 Snijders EP. Verslag van het wetenschappelijk gedeelte der vergaderingen van der afdeling Sumatra's oostkust, *Genewest**


