# Cutaneous Mycetoma due to Nocardia brasiliensis With Delayed Pulmonary Involvement in an Immunocompetent Patient



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### INTRODUCTION

- Nocardia species are aerobic, variably acidfast, Gram-positive filamentous bacteria found in soil
- Cutaneous nocardiosis is rare and often mimics hidradenitis suppurativa (HS), acne conglobata, cellulitis, or sporotrichosis, leading to delayed diagnosis
- N. brasiliensis most commonly causes mycetoma, defined by triad of swelling, draining tracts, and bacterial granules.
- Early recognition and culture are key to prevent dissemination and morbidity

## **CASE DESCRIPTION**

- 27-year-old uninsured male construction worker, immigrant from Mexico, with recurrent draining nodules on upper back previously labeled HS and acne.
- Multiple ED visits and empiric antibiotics, brief isotretinoin trial, over 7 years
- Serial chest CTs in 2025 reveal enlarging 1 cm LLL subpleural nodule and bilateral ground-glass opacities
- **Neck CT**: sinus tracts and soft-tissue infiltration.
- Labs: WBC 22.7  $\times$  10 $^{9}$ /L, Hgb 10 g/dL, lactate 2.3 mmol/L.
- **Histopathology:** suppurative granulomatous inflammation with **sulfur granules (GMS+)**
- Culture grew N. brasiliensis
- Treated with IV amikacin + imipenem → oral
   TMP-SMX + azithromycin
- Marked improvement by **7 weeks**; 4-month CT showed **resolution of pulmonary nodules**
- No immunodeficiency identified.

# DISCUSSION

- 1. Epidemiology & Pathogenesis:
- Nocardia brasiliensis is a soil-borne, filamentous, variably acid-fast bacterium causing most mycetoma in Mexico and Latin America.
   Infection follows minor trauma or inoculation. Though often linked to immunosuppression, up to 45% occur in healthy hosts, typically laborers with soil exposure.
- 2. Clinical Presentation & Differential Diagnosis:
  - Cutaneous nocardiosis mimics HS, acne conglobata, and cellulitis.
     Chronic draining tracts and yellow granules are classic. Our patient's HS-like back nodules underscore the challenge of delayed recognition due to the clinical overlap between nocardiosis and chronic inflammatory skin disorders
  - Pulmonary involvement by N. brasiliensis is extremely rare;
     concurrent cutaneous + pulmonary disease in an
     immunocompetent host is exceptionally rare
- 3. Diagnostic Work-up:
- Biopsy with GMS/AFB stains plus culture is diagnostic. Imaging (CT/MRI) helps detect dissemination to (lungs/central nervous system). N. brasiliensis growth confirmed both cutaneous and pulmonary disease in an immunocompetent patient.
- 4. Treatment & Clinical Course:
- First-line TMP-SMX for 1–3 months (longer if disseminated).
- $\circ$  Severe disease: **combination IV therapy** (typically amikacin  $\pm$  imipenem) followed by transition to oral agents.
- 5. Clinical & Social Implications:

Figures 1A-C: A. Adult male with

dissemination five years prior to

presentation to ED C. After six

cutaneous nocardiosis and

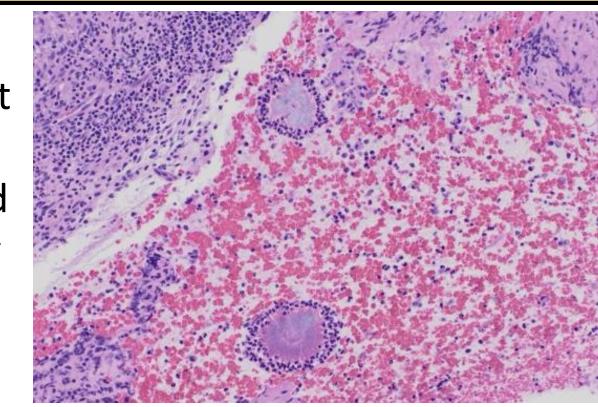
presentation at ED B. At

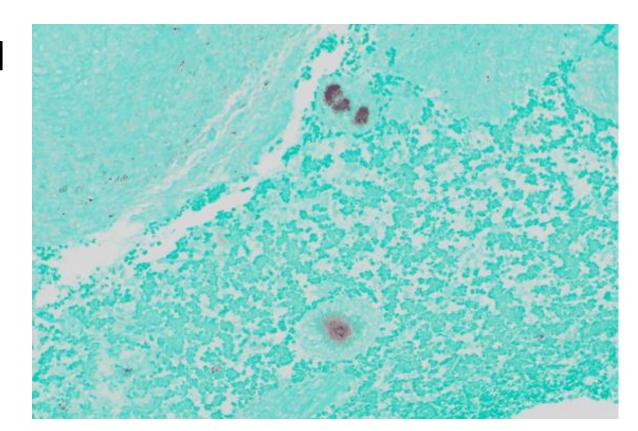
weeks after diagnosis and

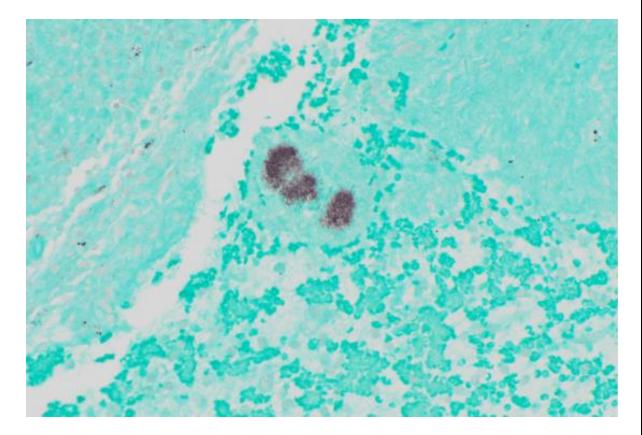
initiation of targeted therapy

delayed pulmonary

- Mortality varies 7–64% depending on immune status and dissemination.
- Fragmented, ED-based care and lack of insurance can delay evaluation of persistent skin infections. Diagnosis and follow-up can transform prolonged morbidity into curable infection.







Figures 2A-C. A. Hematoxylin & Eosin stain at 20x magnification B. Grocott Methenamine Silver (GMS) special stain at 20x magnification C. GMS stain at 40x magnification

#### **LEARNING POINTS & CONCLUSION**

- Evaluation of treatment-resistant HS or acne should include biopsy with bacterial, fungal, and AFB stains plus culture or molecular identification, especially in immigrants from endemic regions/with soil contact
- Mycetoma-type nocardiosis can occur in immunocompetent hosts and may rarely spread to lungs
- Early, prolonged, targeted therapy prevents morbidity and scarring
- Social determinants of health such as access barriers and fragmented care may delay diagnosis.

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