A Case of Mycobacterium Abscessus in an Immunocompetent Patient

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1. Abstract

1.1. Presentation

A 71-year-old lady from the USA, presented with a four-week history of cough and dyspnoea with a cavitatory lesion on her chest x-ray.

1.2. Diagnosis

Subsequent CT Thorax and bronchoscopy with bronchoalveolar lavage (BAL) and transbronchial needle aspirate (TBNA) biopsies confirmed a diagnosis of mycobacterium abscessus.

1.3. Treatment

Having initially been treated with anti-tuberculous therapy, this was ceased, and she completed a course of oral clarithromycin before being lost to follow-up on her return to the USA.

1.4. Discussion

She had no identifiable risk factors or underlying lung disease and work-up for immunosuppression was negative.

It is very rare for abscessus to cause lung pathology in immunocompetent patients.

2. Case Report

A 71-year-old American tourist from Florida on holiday in Ireland, presented to hospital with a 6 week history of persistent cough productive of green purulent sputum without haemoptysis.

Her background history was unremarkable; she had no underlying lung disease and was a non-smoker. She had never been admitted to hospital acutely before and had no recent courses of antibiotics or steroids. She was never ill as a child and had no significant family history. She is a retired schoolteacher and had no history of tuberculosis (TB) or TB exposure. A chest radiograph was performed showing a 3 cm thin-walled cavity in the right upper lobe.

Sputum cultures were sent, and she was admitted to the ward in an isolation room and put on an IV penicillin-based antibiotic. She was referred urgently to the respiratory service and underwent a HRCT and bronchoscopy. The CT showed an abnormal right upper lobe with a 6 cm slightly thick-walled cavity associated with local peribronchial thickening and multiple soft nodular opacities in the right upper lobe. A transbronchial biopsy (TBNA) and bronchoalveolar lavage (BAL) were taken and sent for TB culture and microscopy. She was commenced on quadruple TB therapy at that point and discharged from hospital. She was advised to refrain from flying. She returned to the outpatient department 2 weeks later. Her FBC was normal, negative HIV screen and her immunoglobulins, complement levels, HbA1c and autoimmune screen all returned as normal. The results of the sputum and bronchial washings returned a diagnosis of mycobacterium abscessus. It was resistant to ciprofloxacin, cotrimoxazole and doxycycline and sensitive to amikacin, clarithromycin, and linezolid. Her TB therapy was stopped, and she was commenced on oral clarithromycin. She was lost to follow-up when she flew back home to the USA.

3. Discussion Mycobacterium Abscessus

Mycobacterium abscessus is a rapidly growing mycobacterium (RGM) distantly related to TB and Leprosy [1]. M. abscessus can cause a variety of clinical manifestations, mostly skin and soft tissue infections. Granuloma formation is the hallmark of its disease process. Rarely, in individuals with underlying respiratory conditions or impaired immunity, M. Abscessus can cause lung infection. Host risk factors for disease include increased age, smoking/vaping, underlying cystic fibrosis or bronchiectasis, immunosuppression (HIV/AIDS, mutations in interferon gamma pathway or...
immunosuppressive drug therapy with TNF-alpha inhibitors) and broad-spectrum antibiotic use [2]. It is even rarer form. Abscessus to cause lung pathology in immunocompetent patients. This case of mycobacterium abscessus infection occurred in a previously healthy lady without underlying lung disease or immunosuppressive disease. The secondary work up for underlying immunological deficiency was negative with a negative HIV screen, normal immunoglobulins and complement and she was never on immunosuppressive therapy. She had a negative sweat test and no evidence of underlying lung damage on CT scan or PFTs. Is there a geographical component to this case? In a 2012 study by Adjemian et al. [3]. Identified Florida as a particular high-risk area for pulmonary NTM. A 2018 study of hospitalised patients with NTM based in Miami, Florida showed a similar pattern of resistance as in our case.

Could this lady have an as-yet undiagnosed subclinical immunosuppressive disease? This is possible as new advances in throughput technology means an increasing recognition of subtle genetic mutations causing immune dysfunction. There is no family history whatsoever however this does not exclude de novo mutations. The mystery continues.

References
2. https://www.sciencedirect.com/topics/medicine-and-dentistry/mycobacterium-abscessus#:~:text=Predisposing%20conditions%20that%20are%20commonly,usually%20younger%20than%2040%20years