

National Jewish Health[®] Breathing Science is Life[®]

NTM Lecture Series for Providers

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Challenging cases

NTM Provider Course

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Disclosures

- Insmed: speaker, advisory board, investigator
- AN2: advisory board
- Paratek: speaker

Objectives

- Recognize underlying host vulnerability to bronchiectasis
- Appreciate diagnostic tools for NTM-LD
- Differentiate clinically relevant infection from non-pathogenic infection



Case One

- 40 year old female
- Healthy child
- Pneumonia at 17, 20 years old
- Pulmonary MAC diagnosed and treated on 3 separate occasions
 - First incidence at age 25
 - No culture data while on treatment
- No history of sinusitis, otitis media or other recurrent infections
- No history of infertility
- No family history of pulmonary disease



Case One

- Current symptoms of chronic cough, sputum production and fatigue
- 3/3 sputum cultures are smear -, culture + M. avium







What additional diagnostic testing is most likely to reveal her underlying susceptibility to NTM lung disease?

- 1. Sweat testing
- 2. Alpha 1 antitrypsin level and phenotype
- 3. Quantitative immunoglobulins
- 4. Interferon gamma auto-antibody testing



- Sweat chloride: 21 mmol/L (within normal)
- Quantitative immunoglobulins: normal
- Alpha 1 level: undetectable
- Alpha 1 phenotype: ZZ





NEJM., 2020, Vol.382(15), p.1443-1455

Follow up 5 years later

- Patient is receiving Alpha 1 augmentation (Prolastin-C) therapy
- She completed 18 months of MAC treatment
 - With 12 months of negative cultures
 - No evidence of recurrence
- Airway clearance: vest, aerobika, hypertonic saline twice daily
- Chronic lower respiratory infection with MRSA



Case Two

- 38-year-old female presented to NJ for an evaluation of severe asthma
- Childhood asthma, recurrent pneumonia, chronic sinusitis
- Hemoptysis at age 36, hospitalized with pneumonia
- First CT imaging at age 38







Diagnostic evaluation

- Smear +, culture + for MAC (3/3)
- Bacterial cultures + pseudomonas aeruginosa
- FEV1 57% predicted (ratio 60)
- IgE >5000



What is the most likely diagnosis

- Asthma
- APBA
- Cystic fibrosis
- All of the above



Follow up

- Sweat chloride 26, 39 mmol/L (above normal range)
- CFTR: deltaF508, R117H mutations
- She completed 12 months of negative cultures for MAC
- While on treatment for MAC, she began isolating *M. abscessus*
- She completed an additional 12 months of *M. abscessus* therapy
- 2018 she started modulator therapy





Follow up CT scan 6 years later-2019





Case three

- 23 year old female
- In Thailand teaching English
- 18 months (7/16) into stay: 3 weeks of dry cough without fever/chills/NS/wt loss



Initial CXR







Initial evaluation (Thailand)

- Sputum x 1: afb smear negative
- 9/2016 BAL:
 - smear negative for afb
 - Routine cultures negative
 - Fungal cultures and serology negative
 - Cytology negative for malignancy
 - Meloidosis serology negative
- Started on empiric TB therapy Rifampin/Isoniazid/PZA/Ethambutol
- 2 month follow up: symptoms unchanged, AFB cultures negative
- 11/16 she elected to return to the US



Initial CT 11/2016







NJ Evaluation 10/2017

- Continued cough, productive
- CBC WBC 11.1, Hgb 11.7, Plt 338
- No eosinophilia
- CMP normal
- HIV negative
- Immunology evaluation normal
- Fungal serologies negative
- QFT negative





As an NTM suspect-What additional testing would be indicated?

- 1. Second bronchoscopy for repeat AFB cultures
- 2. Universal PCR on BAL fluid
- 3. Sputum induction x 3 for AFB smear and culture
- Serum IgA antibodies against mycobacterial glycopeptidolipid (GPL) core antigen



NJ evaluation 10/2017

- BAL cell count: 94% neutrophils, 3% macrophage, 3% Lymph, 0 eosinophils
- BAL culture negative (bact/fungal/afb)
- BAL galactomannan negative
- 3 induced sputum: + *M. abscessus*





CT scan consistent with congenital abnormality: bronchial atresia



Medicine and Infectious Diseases

Volume 28, Issue 12, December 1998, Pages 937-939



Effect of lidocaine (Xylocaine $\ensuremath{^{\ensuremath{\mathbb{R}}}}$) on culture of tubercle bacilli in liquid medium

F. Evreux **, C. Lemort **, A. Hauchecorne **, C. Lacroix ***, A. Morel **



M. abscessus, subspecies abscessus: C28 sequevar



11/2017

1/2018



M. abscessus, subspecies abscessus: C28 sequevar



11/2017

Follow up



- Aggressive IV + PO therapy
- Right upper lobectomy
- Completion of 12 months of negative cultures
- 3.5 years later, no evidence of recurrence



Case Four

- 64 year old female with chronic cough of 4 years
- Repeated courses of azithromycin for bronchitis
- 2014 AFB cultures +M. intracellulare, macrolide resistant
- Treatment at NJH with ethambutol/rifampin/moxifloxacin + IV amikacin
- RML, lingulectomy



Case Four

- 1/2016, she achieves 12 months of negative cultures and d/c therapy
- Intermittent cough, mild and stable
- Follow up surveillance cultures form outside hospital



With these follow up cultures would you

- 1. Restart prior 4 drug NTM therapy
- 2. Start Azithromycin, ethambutol, rifampin (A/E/R) while waiting for susceptibility results
- 3. Order repeat imaging and start A/E/R therapy if her CT scan shows progression





KEY POINTS

- Flares in symptoms over the years were attributed to pseudomonas aeruginosa infections and symptoms abate with antipseudomonal therapy.
- CT scan has remained stable now 5 years off therapy despite intermittent growth of different NTM species.
- Underlying bronchiectasis increases her risk of NTM-LD in the future
- Counseling includes: continued airway clearance, avoidance of immunosuppressive agents including ICS, environmental avoidance

