

#### **COUGH to determine a gram-negative pneumonia**

#### Adriana Cecchini, MSN, RN, CIC

*Clinical Documentation Specialist* Brigham and Women's Faulkner Hospital Boston, MA





#### **Presented By**



- Adriana Cecchini, MSN, RN, CIC
- Adriana is a CDI specialist at Brigham and Women's Faulkner Hospital in Boston, Massachusetts. Adriana has been a nurse for more than 15 years. Adriana's prior clinical experience includes acute care, infectious disease, and public health. She holds a certification in infection control and epidemiology. She has been working in CDI since 2019.



#### **No disclosures**



www.shutterstock.com · 1029115462



### **Learning Objectives**

- At the completion of this educational activity, the learner will be able to:
  - Apply the COUGH acronym when evaluating a patient admitted with pneumonia
  - Describe the rationale for recommendations on selected diagnostic and treatment strategies for adult patients with gram-negative (GN) pneumonia (PNA)
  - Evaluate a query opportunity in the Cystic Fibrosis (CF) patient with PNA or pulmonary exacerbation
  - Identify opportunities for collaboration at their own facility to increase capture of patients with gram-negative pneumonia



## Why query to specify a gram-negative pneumonia as the Principle Diagnosis?

Scenario One	Scenario Two
<ul><li>81 year old female with a history of CAD, HTN and GERD.</li><li>Now admitted to the hospital with pneumonia being treated with Ceftriaxone and Azithromycin.</li></ul>	81 year old female with a history of CAD, HTN, GERD and COPD with recent exacerbation requiring intubation 2 weeks ago. Now admitted to the hospital with a gram-negative pneumonia being treated with Piperacillin / Tazobactam (Pip/Tazo).
DRG: 195: Simple Pneumonia without	DRG 179: Respiratory Infections and
CC/MCC	Inflammation without CC/MCC
Weight: 0.6658	Weight: 0.8727
ALOS: 2.9	ALOS: 3.66
GMLOS: 2.5	GMLOS: 3.07
Severity of Illness: 1	Severity of Illness: 1
Risk of Mortality: 1	Risk of Mortality: 1



#### What is one of the main symptoms of Pneumonia?



#### What is one of the main symptoms of Pneumonia?

Cough





www.shutterstock.com · 1689695362



When you are reviewing a possible gram-negative pneumonia case, work through the COUGH...

C-comorbidities O-organisms U-use of antibiotics G-guidelines for treatment H-hospitalizations



#### **C-Comorbidities**

- Chronic heart, lung, liver, or renal disease; diabetes mellitus; malignancy; or asplenia
  - Ex. COPD, Cystic Fibrosis, ESRD on HD
- According to the Clinical Practice Guideline of the American Thoracic Society and Infectious Diseases Society of America (IDSA):

"Patients with comorbidities should receive broader-spectrum treatment for two reasons. First, such patients are likely more vulnerable to poor outcomes if the initial empiric antibiotic regimen is inadequate. Second, many such patients have risk factors for antibiotic resistance by virtue of previous contact with the healthcare system and/or prior antibiotic exposure (see Recommendation 10) and are therefore recommended to receive broader-spectrum therapy to ensure adequate coverage. In addition to H. influenzae and M. catarrhalis (both of which frequently produce β-lactamase), S. aureus and gram-negative bacilli are more common causes of CAP in patients with comorbidities, such as COPD."

https://www.idsociety.org/practice-guideline/community-acquired-pneumonia-cap-in-adults/

Gram negative Pneumonia is more common in HAP/VAP
 https://www.idsociety.org/practice-guideline/hap\_vap/



#### **O-Organisms**

- Look at respiratory cultures: BAL, endotracheal aspirate, induced sputum etc.
  - Current cultures
  - Past cultures (including possible colonization)
  - Viral respiratory panels
  - Urine antigen testing
- IMPORTANT: Respiratory cultures <u>without</u> growth does NOT mean a patient does not have a GN PNA. Respiratory cultures with growth help guide treatment when available but are not required for diagnosis.
- Challenges with respiratory cultures:
  - Difficult to obtain good culture
  - Need to obtain before starting antibiotics



## What are some examples of common GN organisms that can cause pneumonia?

#### **Community-Acquired Pneumonia**

- Klebsiella pneumoniae
- Haemophilus influenzae
- Moraxella catarrhalis
- Atypical (GN) bacteria
  - Mycoplasma pneumoniae
  - Chlamydia pneumoniae
  - Legionella pneumophila
  - Chlamydia psittaci
- Rarely:
  - Pseudomonas aeruginosa
  - Escherichia coli
  - Acinetobacter baumannii

#### Hospital-acquired & Ventilatorassociated Pneumonia

- Pseudomonas aeruginosa
- Gram-negative enteric bacilli:
  - Klebsiella pneumoniae
  - Enterobacter spp
  - Escherichia coli
- Acinetobacter baumannii
- Stenotrophomonas maltophilia



#### **Other common organisms that can cause PNA**

- Gram positive organisms: Streptococcus pneumoniae (most common cause of CAP) and Staphylococcus aureus, Group A streptococci
- Viruses: Influenza, SARS-CoV-2 (COVID-19), Rhinoviruses, Adenovirus, Parainfluenza Virus, Respiratory Syncytial Virus
- Fungi: Pneumocystis jirovecii, Aspergillus species (especially A. fumigatus), and Cryptococcus neoformans



#### **U-Use of antibiotics**

- Currently prescribed antibiotics
  - If indicated do they provide GN coverage?
- Recent antibiotic use for any condition/infection
- Recent antibiotic treatment failure for pneumonia



#### **G-Guidelines /CMS Conditions of Participation**

- Hospitals are required to comply with the Federal requirements set by the Medicare Conditions of Participation (CoP) in order to receive Medicare/Medicaid payment.
  - §482.42 Condition of Participation: Infection Prevention and Control and Antibiotic Stewardship Programs: The hospital must have active hospital-wide programs for the surveillance, prevention, and control of HAIs and other infectious diseases, and for the optimization of antibiotic use through stewardship. The programs must demonstrate adherence to nationally recognized infection prevention and control guidelines, as well as to best practices for improving antibiotic use where applicable, and for reducing the development and transmission of HAIs and antibiotic resistant organisms. Infection prevention and control problems and antibiotic use issues identified in the programs must be addressed in collaboration with the hospital-wide quality assessment and performance improvement (QAPI) program.



- Your institution may already have empiric antibiotics guidelines that you can use to help identify query opportunity or avoid unnecessary queries
- Identify the CMS antimicrobial stewardship chapter author for guidance and collaboration with your CDI department
- Learn about restricted use antibiotics at your organization. There may be an opportunity to query if a restricted antibiotic is ordered



Pulmonary	1	1	NOTES:	
1. Community-Acquired Pneumonia (CAP) – Low risk for antibiotic resistant pathogens	Streptococcus pneumoniae (40% resistant to azithromycin). Haemophilus Influenzae (20% resistant to amoxicillin), Atypical pathogens (Mycoplasma pneumoniae, Chiamydophila pneumoniae, Legionella species)	(Ceftriaxone 1-2gm IV Q24hr <u>PLUS</u> Azithromycin 500mg PO Q24hr) OR Levofloxacin 750mg PO Q24hr Patients admitted to the ICU may require broader treatment (see CAP High Risk section below).	<ol> <li>Appropriate diagnostics for CAP include: sputum gram stain and culture, urine Legionella and urine Pneumococcal antigen testing. Consider testing for respiratory viruses as well as viruses may account for ≥50% of community-acquired pneumonias.</li> <li>Azithromycin and levofloxacin both treat atypical pathogens. Atypical pathogens. Atypical pathogens. however, account for &lt;5% of CAP cases in hospitalized patients. May consider withholding in patients with mild disease.</li> <li>Doxycycline 100mg PO BID is an alternative to azithromycin for co- administration with ceftriaxone.</li> <li>Oral step-down therapy for ceftriaxone: amoxicillin/clavulanate or levofloxacin.</li> </ol>	5 days
<ol> <li>Community Acquired Pneumonia (CAP) – High risk for antibiotic resistant pathogens</li> <li>(septic shock, need for mechanical ventilation, IV antibiotics in last 90 days, cystic fibrosis, bronchiectasis, known colonization with Pseudomonas or MDR pathogen)</li> </ol>	Above pathogens and Pseudomonas, MDR GNRs	Cefepime 2gm IV Q8hr OR Piperacillin/tazobactam 4.5gm IV Q6hr PLUS Levofloxacin 750 mg PO/IV OR azithromycin 500 mg PO/IV * If septic shock, known MRSA colonization, necrotizing pneumonia, or new need for mechanical ventilation:	NOTES: 1. Appropriate diagnostics for CAP include: sputum gram stain and culture, urine Legionella and urine Pneumococcal antigen. Consider testing for respiratory viruses as well. Viruses may account for 250% of community- acquired pneumonias. 2. Narrow therapy once organism and	5 days



		Add Vancomycin 30-45 mg/kg/day IV divided Q8- 12hr (goal trough 15-20 mcg/mL) OR Linezolid 600mg IV/PO Q12h	susceptibility results are known. 3.if vancomycin or linezolid is started, stop at 48hrs if no MRSA cultured.	
3. Aspiration Pneumonia	Gram-negative enteric pathogens, oral anaerobes	Ceftriaxone 1-2gm IV Q24hr OR Levofloxacin 750mg PO Q24hr Only add Metronidazole 500mg PO Q8hr if abscess	Clindamycin 600mg IV Q8hr or 450mg PO Q6hr	If rapid clinical improvement, consider aspiration pneumonitis and stop antibiotics early. If slow to improve, then 5 days
4. Hospital-Acquired Pneumonia [HAP] (risk factors for MDRO: IV antibiotics within the past 90 days)	Overlap with CAP in terms of common pathogens for patients with early onset HAP (<5 days from admission). Aerobic gram- negative bacilli including <i>Pseudomonas aeruginosa</i> more common with prolonged hospitalization/ventilation. <i>Staphylococcus aureus</i> more common in patients with known MRSA colonization, necrotizing pneumonia, empyema, septic shock, or recent influenza.	Gram-negative: Cefepime 2gm IV Q8hr OR Piperacillin/tazobactam 4.5gm IV Q6hr (give over 3 hours) OR Meropenen 1gm IV Q8h * If septic shock, known MRSA colonization, necrotizing pneumonia, or new need for mechanical ventilation add MRSA treatment: Vancomycin 30-45 mg/kg/day IV divided Q8-12hr (goal trough 15-20 mcg/mL) OR Linezolid 600mg IV/PO Q12h AND If septic shock add second Gram-negative agent: Levofloxacin 750 mg IV or gentamicin or tobramycin 7 mg/kg (re- dose per pharmacy)	NOTES: 1. Send respiratory sample for Gram stain and culture for all patients. 2. Consider urine legionella antigen testing. 3. Deescalate therapy once organism and susceptibility results are known. 4. If vancomycin or linezolid is started, stop at 48hrs if no MRSA cultured.	7 days



- Other resources and guidelines:
- CAP guidelines: <u>https://www.idsociety.org/practice-guideline/community-acquired-pneumonia-cap-in-adults/</u>
- HAP/VAP guidelines: <u>https://www.idsociety.org/practice-guideline/hap\_vap/</u>
- Johns Hopkins Antibiotic Guide
   https://www.unboundmedicine.com/ucentral/index/Johns\_Hopkins\_ABX\_Guide/All\_Topics/A



#### **IDSA/ATS CAP Treatment Guidelines**

Table 4. Initial Treatment Strategies for Inpatients with Community-acquired Pneumonia by Level of Severity and Risk for Drug Resistance

	Standard Regimen	Prior Respiratory Isolation of MRSA	Prior Respiratory Isolation of Pseudomonas aeruginosa	Recent Hospitalization and Parenteral Antibiotics and Locally Validated Risk Factors for MRSA	Recent Hospitalization and Parenteral Antibiotics and Locally Validated Risk Factors for <i>P. aeruginosa</i>
Nonsevere inpatient pneumonia*	β-Lactam + macrolide <sup>†</sup> or respiratory fluroquinolone <sup>‡</sup>	Add MRSA coverage <sup>§</sup> and obtain cultures/nasal PCR to allow deescalation or confirmation of need for continued therapy	Add coverage for <i>P. aeruginosa</i> <sup>  </sup> and obtain cultures to allow deescalation or confirmation of need for continued therapy	Obtain cultures but withhold MRSA coverage unless culture results are positive. If rapid nasal PCR is available, withhold additional empiric therapy against MRSA if rapid testing is negative or add coverage if PCR is positive and obtain cultures	Obtain cultures but initiate coverage for <i>P. aeruginosa</i> only if culture results are positive
Severe inpatient pneumonia*	$\beta$ -Lactam + macrolide <sup>†</sup> or $\beta$ -lactam + fluroquinolone <sup>‡</sup>	Add MRSA coverage <sup>§</sup> and obtain cultures/nasal PCR to allow deescalation or confirmation of need for continued therapy	Add coverage for <i>P. aeruginosa</i> <sup>  </sup> and obtain cultures to allow deescalation or confirmation of need for continued therapy	Add MRSA coverage <sup>§</sup> and obtain nasal PCR and cultures to allow deescalation or confirmation of need for continued therapy	Add coverage for <i>P. aeruginosa</i> <sup>  </sup> and obtain cultures to allow deescalation or confirmation of need for continued therapy

https://www.idsociety.org/practice-guideline/community-acquired-pneumonia-cap-in-adults/



#### **IDSA/ATS HAP/VAP Treatment Guidelines**

#### Table 4. Recommended Initial Empiric Antibiotic Therapy for Hospital-Acquired Pneumonia (Non-Ventilator-Associated Pneumonia)

Not at High Risk of Mortality <sup>a</sup> and no Factors Increasing the Likelihood of MRSA <sup>b,c</sup>	Not at High Risk of Mortality <sup>a</sup> but With Factors Increasing the Likelihood of MRSA <sup>b,c</sup>	High Risk of Mortality or Receipt of Intravenous Antibiotics During the Prior 90 d <sup>a,c</sup>
One of the following:	One of the following:	Two of the following, avoid 2 β-lactams:
Piperacillin-tazobactam <sup>d</sup> 4.5 g IV q6h	Piperacillin-tazobactam <sup>d</sup> 4.5 g IV q6h	Piperacillin-tazobactam <sup>d</sup> 4.5 g IV q6h
OR	OR	OR
Cefepime <sup>d</sup> 2 g IV q8h	Cefepime <sup>d</sup> or ceftazidime <sup>d</sup> 2 g IV q8h	Cefepime <sup>d</sup> or ceftazidime <sup>d</sup> 2 g IV q8h
OR	OR	OR
Levofloxacin 750 mg IV daily	Levofloxacin 750 mg IV daily	Levofloxacin 750 mg IV daily
	Ciprofloxacin 400 mg IV q8h	Ciprofloxacin 400 mg IV q8h
	OR	OR
Imipenem <sup>d</sup> 500 mg IV q6h	Imipenem <sup>d</sup> 500 mg IV q6h	Imipenem <sup>d</sup> 500 mg IV q6h
Meropenem <sup>d</sup> 1 g IV q8h	Meropenem <sup>d</sup> 1 g IV q8h	Meropenem <sup>d</sup> 1 g IV q8h
	OR	OR
	Aztreonam 2 g IV q8h	Amikacin 15–20 mg/kg IV daily
		Gentamicin 5–7 mg/kg IV daily
		Tobramycin 5–7 mg/kg IV daily
		OR
		Aztreonam <sup>e</sup> 2 g IV q8h
	Plus: Vancomycin 15 mg/kg IV q8–12h with goal to target 15–20 mg/mL trough level (consider a loading dose of 25–30 mg/kg × 1 for severe illness)	Plus: Vancomycin 15 mg/kg IV q8–12h with goal to target 15–20 mg/mL trough level (consider a loading dose of 25–30 mg/kg IV × 1 for severe illness)
	OR	OR
	Linezolid 600 mg IV q12h	Linezolid 600 mg IV q12h
		If MRSA coverage is not going to be used, include coverage for MSSA Options include: Piperacillin-tazobactam, cefepime, levofloxacin, imipenem, meropenem. Oxacillin, nafcillin, and cefazolin are preferred for the treatment of proven MSSA, but would ordinarily not be used in ar empiric regimen for HAP.
		llin allergy and aztreonam is going to be used ased antibiotic, include coverage for MSSA.



#### **H-Hospitalizations**

- Look for hospitalizations or admissions to a healthcare center (rehab, nursing home etc.) within the past 3 months
- Also think about....H-Hoses= respiratory equipment ex. recent intubation, CPAP, tracheostomy, bronchoscopy etc.



www.shutterstock.com · 1593004267



#### **Cystic Fibrosis and Pneumonia**





### **Cystic Fibrosis (CF)**

- Pulmonary disease remains the leading cause of morbidity and mortality in patients with CF
- Chronic airway obstruction caused by viscous secretions, leads to progressive pulmonary colonization with pathogenic bacteria
- Patients with CF are particularly prone to chronic infection with *P. aeruginosa*
- Hypoxia causes changes in *P. aeruginosa* (and some other GN bacteria), including loss of motility and causes alginate production
  - Alginate, or alginic acid, is a component of the biofilm
  - Biofilm bacteria share nutrients and are protected from harmful factors in the environment, Ex: antibiotics; the host body's immune system
  - Biofilm and *P. aeruginosa* have a high resistance to antibiotics
  - Once biofilm production, aka development of "bacterial macrocolonies" occurs, eradication of the organism/infection is nearly impossible



#### **Cystic Fibrosis (CF) continued**

- Treatment of exacerbations with systemic antibiotics is a mainstay of CF care and is recommended in virtually all consensus guidelines
- Patients with persistent *P. aeruginosa* infection, are typically treated with chronic inhaled Tobramycin
- Most patients with CF have chronic bacterial infection of the airways with one or more of these organisms:
  - Pseudomonas aeruginosa
  - Staphylococcus aureus (methicillin-sensitive or methicillin-resistant species)
  - Burkholderia cepacia complex
  - •Nontypeable Haemophilus influenzae
  - Stenotrophomonas maltophilia
  - Achromobacter species
  - •Nontuberculous mycobacteria



#### **Cystic Fibrosis (CF) continued**

- So when do you query for GN PNA in a patient with CF?
- ANYTIME the patient is admitted with for possible pneumonia or a CF exacerbation!



#### **General Reminders**





#### Reminder...

#### Check antibiotic order for indications

Components			Order Questions		
Component cefepime 2 gram Solr	Order Dose 2,000 mg	Admin Dose 2,000 mg	Question Indication	Answer Empiric	Comment
sodium chloride 0.9% Pgbk		100 mL	Infection Source	Pneumonia	
Compon	ent Details				



#### Reminder...

#### Check antibiotic order for indications

Components			Order Questions	5	
	Order	Admin	Question	Answer	Comment
Component	Dose	Dose	Indication	Definitive	
piperacillin-tazobactam 4.5 gram Solr	4.5 g	4.5 g		(documented infection)	
sodium chloride 0.9% Pgbk	100 mL	100 mL	Infection Source	Pneumonia	



#### **Remember to check allergies**

• Documented allergies may be the reason a patient is not on the standard antibiotics. Check notes/documentation to assure query is appropriate.





## When should you consider querying for Gram Negative Pneumonia?

- Patient with a positive respiratory culture growing Gram Negative organism (GN)
   OR
- Patient with a co-morbidities ex. COPD, CF, asplenia and/or recent treatment with a vent....**also being treated with:** 
  - Cefepime 2gm IV Q8hr **OR** Piperacillin/tazobactam 4.5gm IV Q6hr

#### PLUS

• Levofloxacin 750 mg PO/IV OR azithromycin 500 mg PO/IV

OR treatment with...

• Meropenem 1gm IV Q8h or Imipenem/cilastatin

#### Or

Ceftazidime or aztreonam

#### Or

• Gentamicin or tobramycin or ciprofloxacin



#### When not to query a GN pneumonia?

#### • Patient on standard CAP treatment

#### **Example treatment:**

- Ceftriaxone 1-2gm IV Q24hr PLUS Azithromycin 500mg PO Q24hr
   OR
  - Levofloxacin 750mg PO Q24hr
- (Or moxifloxacin 400 mg PO/IV q24hr depending on your organization)

# Steps for Attendees to Answer/View POLLING QUESTIONS



# Steps for Attendees to Answer/View POLLING QUESTIONS



- 1. Navigate to the Schedule in the main menu.
- 2. Tap the name of the current session to view the session details page.
- 3. Scroll down the page to Live Polls.
- 4. Tap the name of the poll.
- Tap your answer choice(s) and then tap Submit.

Polling Question 1 Using Event App

Question 1: Should you query for GN PNA on this patient?

- Patient admitted for: CAP Antibiotic Treatment: Ceftriaxone and doxycycline
- Allergies: Levaquin
- Risks: Recent hospitalization after a fall, but no mention of intubation
  - Chose one answer:
    - 1.) YES 2.) NO

## POLLING RESULTS Question 1

Patient admitted for:CAPAntibiotic Treatment:Ceftriaxone and doxycyclineAllergies:LevaquinRisks:Recent hospitalization after a fall, but no mention of intubation

Correct Answer: 2. No, query is not indicated. Patient does not have risk factors for a GN PNA.

**Polling Question 2 Using Event App** 

Question 2: Should you query for GN PNA on this patient?

- Patient admitted for: Pneumonia Antibiotic Treatment: Pip/Tazo
- Allergies: None
- **Risks:** Recent Hospitalization for COPD exacerbation
  - Chose one answer:
    - 1.) YES 2.) NO
Patient admitted for: Pneumonia Antibiotic Treatment: Pip/Tazo Allergies: None Risks: Recent Hospitalization for COPD exacerbation

**Correct answer: 1. Yes**, high potential for GN PNA. Pip/Tazo will provide sufficient empiric of GN and GP organisms

**Polling Question 3 Using Event App** 

Question 3: Should you query for GN PNA on this patient?

- Patient admitted for: Septic shock and PNA
   Antibiotic Treatment: Vanco, Cefepime, Azithromycin
- Allergies: None
- Risks: COPD (uses 2L oxygen at night)
  - Chose one answer:
     1.) YES
    - 2.) NO

Patient admitted for: Septic shock and PNA Antibiotic Treatment: Vanco, Cefepime, Azithromycin Allergies: None Risks: Mild COPD (uses 2L oxygen at night)

**Correct answer: 1. Yes**, high potential for GN PNA. Above meds provide broad empiric treatment for GN, GP, and atypical pathogens. Risk factors: COPD.

Polling Question 4 Using Event App

Question 4: Should you query for GN PNA on this patient?

- Patient admitted for: PNA
   Antibiotic Treatment: Meropenem and Vancomycin
- Allergies: None
- Risks: Stage III Severe COPD and recent treatment for CAP
  - Chose one answer:
    - 1.) YES 2.) NO

Patient admitted for: PNA Antibiotic Treatment: Meropenem and Vancomycin Allergies: None Risks: Stage III Severe COPD and recent treatment for CAP

**Correct answer: 1. Yes**, high potential for GN PNA. Meropenem provides coverage of GN and GP organisms, and vancomycin provides additional GP against MRSA. Risk factors include COPD and recent CAP.

**Polling Question 5 Using Event App** 

Question 5: Should you query for GN PNA on this patient?

- Patient admitted for: CF exacerbation and PNA Antibiotic Treatment: Ceftazidime and Meropenem Allergies: None
- Risks: PMH- Cystic Fibrosis and history of respiratory colonization with drug-resistant *Pseudomonas aeruginosa*

• Chose one answer:



Patient admitted for: CF exacerbation and PNA
Antibiotic Treatment: Ceftazidime, Meropenem, and inhaled tobramycin
Allergies: None
Risks: PMH- Cystic Fibrosis and history of respiratory colonization with drug-resistant
Pseudomonas aeruginosa

**Correct answer: 1. Yes**, high potential for GN PNA. Ceftazidime, Meropenem, and inhaled tobramycin provide broad empiric treatment against GN organisms. Including drug-resistant pseudomonas

**Polling Question 6 Using Event App** 

Question 6: Should you query for GN PNA on this patient?

- Patient admitted for: PNA Antibiotic Treatment: Initially treated with IV Ceftriaxone and Azithromycin, now being discharged home on PO Levofloxacin Allergies: None
- Risks: PMH- mild Asthma, no recent antibiotics
  - Chose one answer:
    - 1.) YES 2.) NO

Patient admitted for: CAP PNA Antibiotic Treatment: Initially treated with IV Ceftriaxone and Azithromycin, now being discharged home on PO Levofloxacin Allergies: None Risks: PMH- mild Asthma, no recent antibiotics

**Correct answer: 2. No**, although current antibiotics do provide <u>some</u> GN treatment, the most common bacterial pathogen in CAP is GP (Streptococcus pneumoniae) and the current regimen is highly active vs Streptococcus pneumoniae and atypical organisms - GN pathogens are less of a concern in this situation as patient. Patient lacks additional risk factors to support a query.

**Polling Question 7 Using Event App** 

Question 7: Should you query for GN PNA on this patient?

- Patient admitted for: PNA Antibiotic Treatment: Ceftriaxone and doxycycline Allergies: azithromycin
- Risks: Positive respiratory culture for Streptococcus pneumoniae
- Chose one answer:
  - 1.) YES 2.) NO

Patient admitted for: PNA Antibiotic Treatment: PO amoxicillin-clavulanate Allergies: azithromycin Risks: Positive respiratory culture for *Streptococcus pneumoniae* 

**Correct answer: 2. No,** Streptococcus pneumoniae is a GP organism. Patient lacks risk factors for GN PNA.

**Polling Question 8 Using Event App** 

Question 8: Should you query for GN PNA on this patient?

- Patient admitted for: Multifocal PNA w/new O2 requirement
   Antibiotic Treatment: Pip/Tazo and Azithromycin Allergies: None
- Risks: Asthma, Bronchiectasis, Recurrent respiratory infection w/multidrug resistant Stenotrophomonas

Chose one answer:

1.) YES 2.) NO

Question 8: Should you query for GN PNA on this patient?

Patient admitted for: Multifocal PNA w/new O2 requirement Antibiotic Treatment: Pip/Tazo Allergies: Lisinopril and HCTZ Risks: Asthma, Bronchiectasis, Recurrent respiratory infection w/ multi-drug resistant Stenotrophomonas

**Correct answer: 2. Yes,** Patient has a history of recurrent infection with a GN organism (Stenotrophomonas) and Pip/Tazo will provide sufficient empiric of GN and GP organisms.



#### In closing

- Remember COUGH to determine if there is an opportunity to query for GN PNA
  - Comorbidities, Organism, Use of antibiotics, Guidelines (antimicrobial) and Hospitalization/"Hoses"
- Anytime a CF patient is being treated with IV antibiotics for PNA or CF exacerbation, assure coverage for GN PNA and send a query to clarify the GN PNA
- Collaborate with the ID pharmacist or ID physician at your organization that reviews antibiotic restrictions and prescribing guidelines



#### References

- Metlay, J. P., Waterer, G. W., Long, A. C., Anzueto, A., Brozek, J., Crothers, K., ... & Whitney, C. G. (2019). Diagnosis and treatment of adults with community-acquired pneumonia. An official clinical practice guideline of the American Thoracic Society and Infectious Diseases Society of America. *American journal of respiratory and critical care medicine*, 200(7), e45-e67.
- Kalil AC, Metersky ML, Klompas M, et al. Management of Adults with Hospital-acquired and ventilator associated Pneumonia: 2016 Clinical Practice Guidelines by the Infectious Diseases Society of America and the American Thoracic Society. Clin Infect Dis 2016; 63:e61
- Johns Hopkins Antibiotic Guide https://www.unboundmedicine.com/ucentral/index/Johns\_Hopkins\_ABX\_Guide/All\_Topics/A
- Asadi L, Sligl WI, Eurich DT, et al. Macrolide-based regimens and mortality in hospitalized patients with community-acquired pneumonia: a systematic review and meta-analysis. Clin Infect Dis 2012; 55: 371
- Cystic Fibrosis Foundation. 2019 Patient Registry: Annual Data Report. 2020. Available at: https://www.cff.org/Research/Researcher-Resources/Patient-Registry/2019-Patient-Registry-Annual-Data-Report.pdf (Accessed on January 7, 2022).
- Crull MR, Somayaji R, Ramos KJ et al. Changing Rate of Chronic Pseudomonas aeruginosa Infection in Cystic Fibrosis: A Population-Based Cohort Study. Clinical Infect Dis 2018; 76:1089



#### A SPECIAL THANK YOU...

- David W. Kubiak, PharmD, BCPS, BCIDP, FIDSA
- BWH/BWFH CDI Team







#### Thank you. Questions?

#### ACecchini@BWH.Harvard.edu

In order to receive your continuing education certificate(s) for this program, you must complete the online evaluation. The link can be found in the continuing education section of the program guide.

