

# **Pediatric Pneumonia and Bronchiolitis Diagnosis and Treatment: An Evidence-Based Approach**



Show me the  
**EVIDENCE!**

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# Disclosures: Nothing to Declare!



# Today's Road Map

- Pneumonia topics
- Bronchiolitis topics
- Case Study
- Summary & Questions
- After this talk you will:
  - Understand that most pediatric CAP is viral
  - Know why narrow spectrum antibiotics are best AND know best amox CAP dosing
  - Reserve labs/imaging only for potentially hospitalized CAP pt's
  - Know that evidence supports minimal bronchiolitis tx (and what the tx's are)



# Pneumonia- Epidemiology

- US: affects 2.6% of children under 17
- PNA hospitalization decreases with age:
  - <2 y/o: 62 per 10,000
  - 10-17 y/o: 4 per 10,000
- PNA is #1 cause of pediatric hospitalizations: \$1 billion annually



# Pneumonia- Definitions

- PNA- LRTI usually associated with fever, resp sx's, evidence of parenchymal involvement (exam or infiltrates)
- Community acquired pneumonia (CAP)- acute lower respiratory tract infection in previously healthy pt





# Pneumonia- Side Note on Antibiotic Stewardship/Overuse

Increased abx resistance

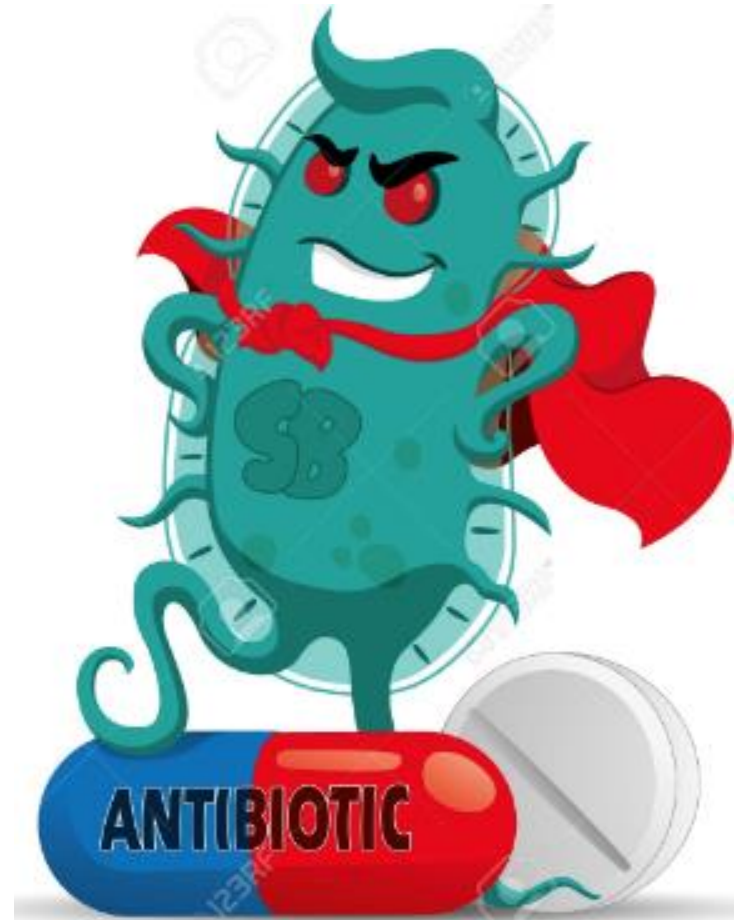
- 2 million annual abx-resistant infections
- 23,000 resultant deaths
- 50% of pediatric abx are not needed or sub-optimally prescribed

Increased *c. difficile*

Increased atopy/asthma

Broad spectrum abx in AOM, strep throat, sinusitis:

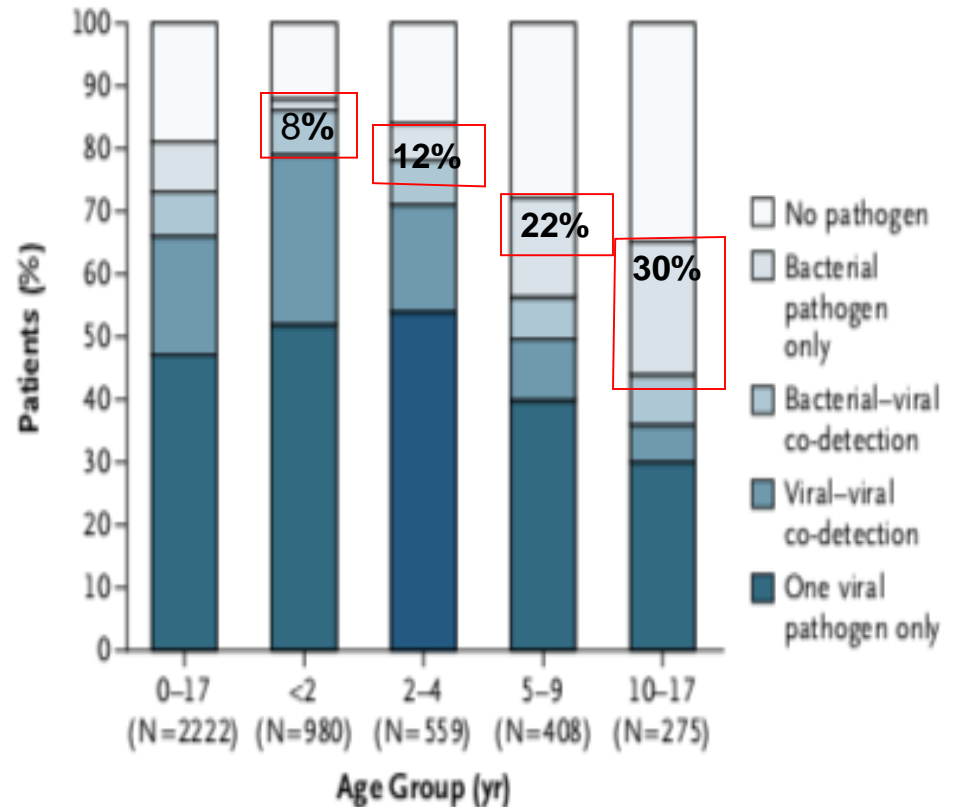
- not more effective
- increase risk of AE's
- lower QOL



# Pneumonia- Etiologies

- CAP 3 main bacterial suspects: *S. pneumoniae*, *S. aureus*, *S. pyogenes*
- EPIC 2015 Study:
  - >2300 pts 0-17 admitted with PNA
  - **Only 15% had bacteria** (% increases with age; largely *mycoplasma*-driven)

A Detection of Bacterial and Viral Pathogens

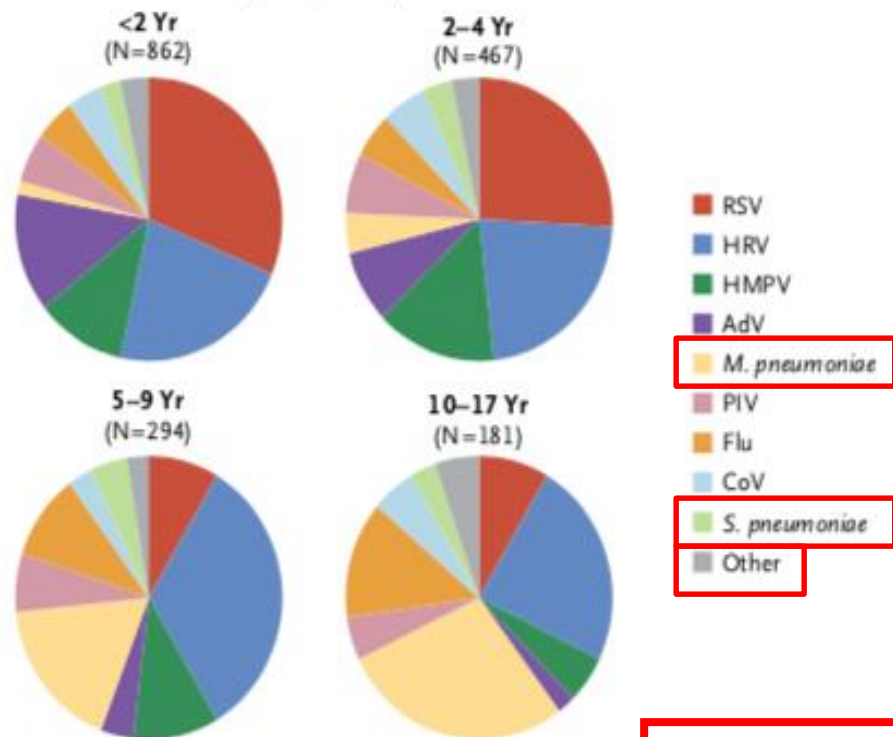


\*Any bacterial etiology percentage



# Pneumonia- Etiologies: EPIC Study Continued

C Detection According to Age Group



- Mycoplasma only exceeds 5% in 5-9 y/o group
- *S. pneumo* and other group increase slightly but stay in 5-10% range

\*Any bacteria





# Pneumonia- Diagnosis

Combo of H&P, imaging, labs

## Exam-

- Fever, cough, increased WOB, hypoxia, tachypnea
- Tachypnea is most sensitive; decreases w/ age
- Lower lobe: possible ab pain
- Younger pt's: nonspecific signs/symptoms
- Consolidation: crackles, fremitus ("ninety-nine"), egophany ("E" sound), bronchophony, dullness to percussion

## CXR-

- **Don't need for mild disease with consistent H&P**
- Uses:
  - Severe illness/admitted pt's
  - Rule out other entities (foreign body, PTX, etc) *when suspected*
  - Inconclusive H&P
  - Treatment failure/complications
  - Febrile infant with leukocytosis
  - Recurrent PNA or suspected anatomic PNA set-up
- Returns to normal in 2 months in 90% of pts

## US/CT-

- US for empyema/effusion
- CT for abscess/fistula suspicion



# Pneumonia- Diagnosis: Labs

**Reserve for severe dz or hospitalization candidates**

## Bloodwork-

- Blood culture- if admitted, tx failure, complicated dz
- Inflammatory markers-
  - One piece of puzzle; not sole determinant
  - **Procalcitonin**-
    - **<0.1 ng/ml: zero pt's with typical bacteria**
    - **<0.25 ng/ml: 96% NPV for typical bacteria**
  - ESR > 35 mm/h PPV = 38%
  - CRP > 6.0 mg/dL PPV = 43%
  - ESR/CRP PPV increases when combined
  - **CBC not useful to determine viral vs. bacterial**
  - Aids in disease monitoring, esp complicated/severe cases

## PCR's and viral studies-

- Viral PCR's- only if results will change mgmt.
- Upper-airway samples can help diagnose: mycoplasma pneumoniae, Chlamydia pneumoniae, Bordetella pertussis, Bordetella parapertussis
- Mycoplasma PCR- more sensitive when done alone compared to viral panel array

## Other specimen studies-

- Pleural fluid analysis- only if complicated dz
- Consider sputum sample in older, mod/severe hospitalized patients



# Pneumonia- Diagnosis: Bacterial, Atypical, or Viral?- THIS IS CHALLENGING!

## Hints at bacterial:

- Abrupt onset, possibly following URI sx's
- Ill appearance
- Complicated pneumonia
- Truly focal exam (note: viruses cause focal pneumonia too)
- Elevated inflammatory markers (esp if both CRP and procalcitonin)

## Viral hints:

- Insidious onset
- Non ill appearance
- Coincident URI symptoms
- Diffuse (non-focal) auscultation
- Proven virus (e.g. respiratory infection panel)
- Wheezy, hypoxic kid more likely viral
- Diffuse x-ray pattern

Hints at atypical (*c. pneumoniae* or *mycoplasma*): See next slide



# Pneumonia- Diagnosis: Hints at Atypical Bacteria

## Labs-

- CRP, WBC, % PMN's lower in atypical vs. *S. pneumo* (although std deviations overlap)

**CXR- no studies have shown differences bw typical and atypical PNA via CXR**

**H&P- Mycoplasma:** URI and LRTI sx's; also hemolytic anemia, rash/SJS, polyarthritits, GI (pancreatitis, hepatitis), carditis

## Clinical-

- Pt age: mycoplasma rates <5% in pt's < 5 y/o**
- Milder, longer course (~3-5 days)
- Low-grade fever
- Wheezing
- Diffuse x-ray exam/findings
- Constitutional findings
- Mycoplasma can also cause effusion (usually small)
- Mycoplasma may start w sore throat & myalgia**

Parameter	Infection due to			Undiagnosed cases (n = 86)
	<i>Streptococcus pneumoniae</i> (n = 48)	Atypical bacteria (n = 46)	Mixed <i>S. pneumoniae</i> and atypical bacteria (n = 16)	
WBC count, mean cells/ $\mu$ L $\pm$ SD	16,669 $\pm$ 8831 <sup>a,b,c</sup>	12,554 $\pm$ 5404	13,141 $\pm$ 4540	12,960 $\pm$ 5670
Neutrophils	69 $\pm$ 17 <sup>a,b,c</sup>	59 $\pm$ 18	63 $\pm$ 16	62 $\pm$ 16
Lymphocytes	22 $\pm$ 15	28 $\pm$ 17	25 $\pm$ 16	27 $\pm$ 17
Monocytes	7 $\pm$ 3	8 $\pm$ 3	7 $\pm$ 3	8 $\pm$ 3
Eosinophils	1 $\pm$ 2	1 $\pm$ 1	1 $\pm$ 2	1 $\pm$ 2
Basophils	0.3 $\pm$ 0.6	0.4 $\pm$ 0.7	0.3 $\pm$ 0.4	0.3 $\pm$ 0.6
CRP level, mean $\mu$ g/dL $\pm$ SD	109 $\pm$ 110 <sup>a,b</sup>	59 $\pm$ 88	77 $\pm$ 79	69 $\pm$ 82
ESR, mean mm/h $\pm$ SD	57 $\pm$ 28	47 $\pm$ 27	52 $\pm$ 44	49 $\pm$ 39

**NOTE.** Data are mean  $\pm$  SD of WBCs  $\pm$  SD, unless otherwise indicated. Unless indicated, differences were not significant. CRP, C-reactive protein; ESR, erythrocyte sedimentation rate.

<sup>a</sup>  $P < .05$  compared with atypical bacterial infection.

<sup>b</sup>  $P < .05$  compared with mixed *S. pneumoniae*-atypical bacterial infection.

<sup>c</sup>  $P < .05$  compared with undiagnosed cases.



# Pneumonia- Diagnosis: Bacterial, Atypical, or Viral?

Clinical and radiographic clues to the etiology of pneumonia in children\*

Etiology	Clinical features	Radiographic features
Bacteria (most commonly <i>Streptococcus pneumoniae</i> )	<ul style="list-style-type: none"> <li>Children of all ages</li> <li>Abrupt onset</li> <li>Ill-appearance</li> <li>Chills</li> <li>Moderate to severe respiratory distress</li> <li>Focal auscultatory findings</li> <li>Localized chest pain</li> <li>WBC count &gt;15,000/microL (if obtained)</li> <li>Elevated acute phase reactants (if obtained)</li> </ul>	<ul style="list-style-type: none"> <li>Alveolar infiltrates</li> <li>Segmental consolidation</li> <li>Lobar consolidation</li> <li>"Round" pneumonia</li> </ul> <p>Complications:</p> <ul style="list-style-type: none"> <li>Pleural effusion/empyema</li> <li>Lung abscess</li> <li>Necrotizing pneumonia</li> <li>Pneumatocele</li> </ul>
Atypical bacterial ( <i>Mycoplasma pneumoniae</i> , <i>Chlamydia pneumoniae</i> )	<ul style="list-style-type: none"> <li>Children of all ages (most common in children &gt;5 years)</li> <li>Abrupt onset with constitutional findings (malaise, myalgia, headache, rash, conjunctivitis, photophobia, sore throat, headache)</li> <li>Gradually worsening nonproductive cough</li> <li>Wheezing</li> <li>Extrapulmonary manifestations or complications (eg, Stevens-Johnson syndrome, hemolytic anemia, hepatitis, etc)</li> </ul>	Interstitial infiltrates
Viral	<ul style="list-style-type: none"> <li>Usually children &lt;5 years</li> <li>Gradual onset</li> <li>Preceding upper airway symptoms</li> <li>Nontoxic appearing</li> <li>Diffuse, bilateral auscultatory findings</li> <li>Wheezing</li> <li>May have associated rash (eg, measles, varicella)</li> </ul>	Interstitial infiltrates
Afebrile pneumonia of infancy (most commonly <i>Chlamydia trachomatis</i> )	<ul style="list-style-type: none"> <li>Usually in infants 2 weeks to 4 months</li> <li>Insidious onset</li> <li>Rhinorrhea</li> <li>Staccato cough pattern</li> <li>Peripheral eosinophilia (if CBC obtained)</li> </ul>	Hyperinflation with interstitial process



# Pneumonia- When to Admit for CAP?

**TABLE 1. Criteria to Consider Hospitalization for Pediatric Pneumonia**

• Hypoxemia (oxygen saturations <90% to 92% at sea level)
• Infants <3 to 6 months of age with suspected bacterial community-acquired pneumonia
• Tachypnea:
◦ Infants <12 months of age: respiratory rate >70 breaths per min
◦ Children: respiratory rate >50 breaths per min
• Respiratory distress: apnea, grunting, difficulty breathing, and poor feeding
• Signs of dehydration or inability to maintain hydration or oral intake
• Capillary refill time >2 s
• Infants and children with toxic appearance
◦ Suspected or confirmed to have infection with a virulent organism (community-acquired methicillin-resistant <i>Staphylococcus aureus</i> or group A <i>Streptococcus</i> )
• Underlying conditions/comorbidities that:
◦ May predispose patients to a more serious course (eg, cardiopulmonary disease, genetic syndromes, neurocognitive disorders, neuromuscular disorders)
◦ May be worsened by pneumonia (eg, metabolic disorder)
◦ May adversely affect response to treatment (eg, immunocompromised host, sickle cell disease)
• Complications (eg, effusion and/or empyema)
• Failure of outpatient therapy (48–72 h with no clinical response)
• Caretaker unable to provide appropriate observation or to comply with prescribed home therapy
Indications for intensive care unit admission include:
• Severe respiratory distress or impending respiratory failure that requires:
◦ Intubation and mechanical ventilation
◦ Positive pressure ventilation
• Recurrent apnea or slow irregular respirations
• Cardiopulmonary monitoring due to cardiovascular compromise secondary to:
◦ Sustained tachycardia
◦ Inadequate blood pressure
◦ Requirement of pharmacological support for blood pressure or perfusion
◦ Altered mental status due to hypercarbia or hypoxemia
◦ Pulse oximetry measurement of <92% on fractional inspired oxygen concentration of >0.50
• Pediatric Early Warning Score >6

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# Pneumonia- Outpatient Mgmt

Tx is empiric

**Abx not routinely recommended in preschool age**  
(unless strong bacterial suspicion)

**Oral cephalosporins are inferior to amox**

Beta-lactam allergy-

- Mild: amox w/ observation, PO cephalosporin (e.g. cefpodoxime, cefprozil, cefuroxime)
- Severe: levoflox, linezolid, macrolide (resistance often high)

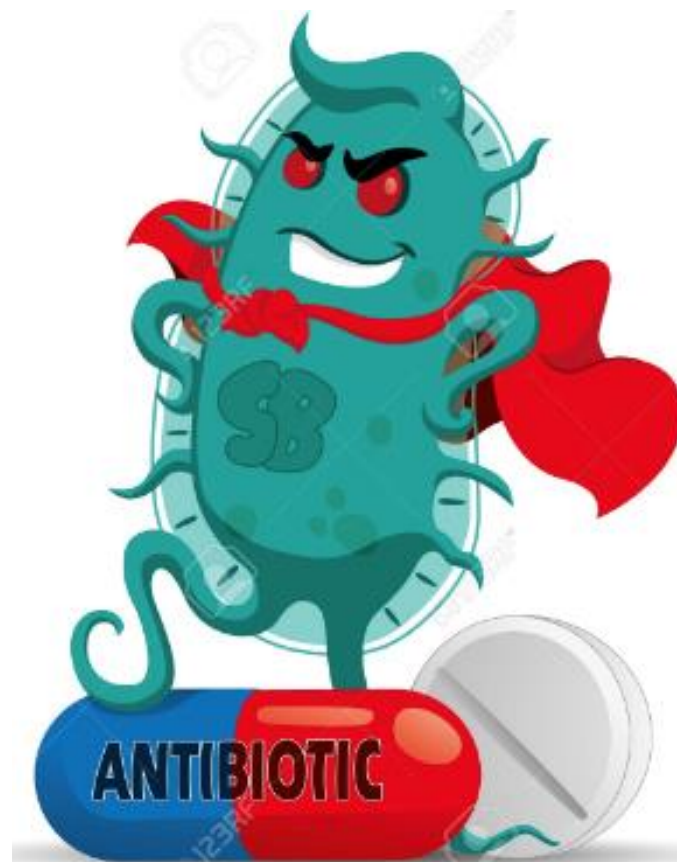
**Amox dosing: 33 mg/kg TID (MDD 4 g)**

Duration- 5-7 days

Amox-clavulanate- adds HiB and *M. catarrhalis* coverage (not needed in fully immunized, normal child)

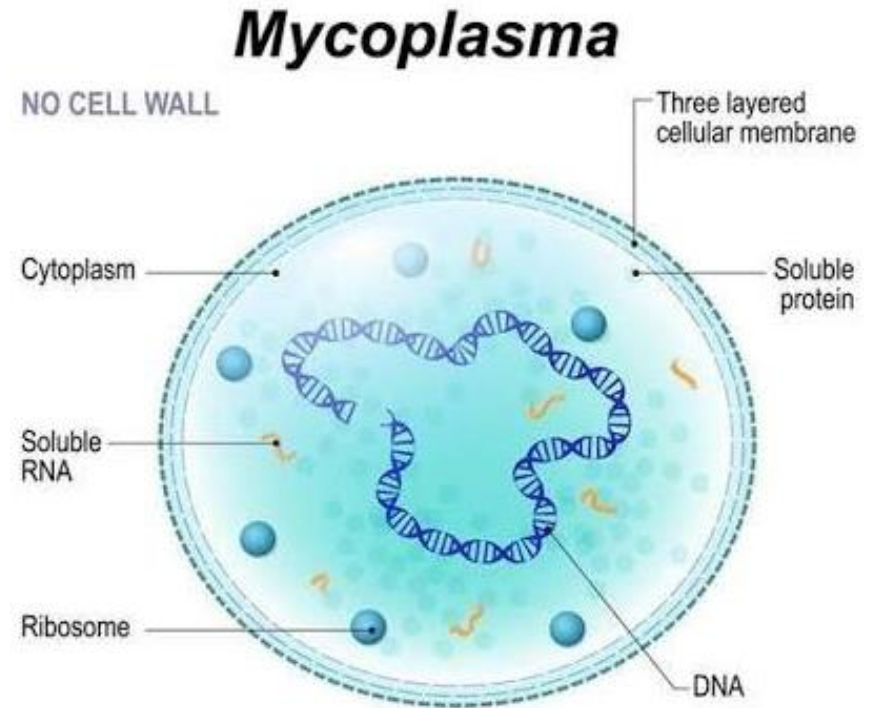
Mycoplasma- azithro for 5 days

Remember your local antibiogram



# Pneumonia- Outpt Mgmt: Should I Test and Treat Mycoplasma?

- Azithro will not adequately treat most CAP bugs
- Testing- consider if unclear H&P; results are not immediate
- Treatment- when high suspicion based on H&P/testing



# Pneumonia- Outpt Mgmt: Tx Failure

- Definition- clinical worsening despite *48 hours properly chosen/dosed abx*
- Consider repeat CXR/hospitalization
- If hospitalized, expand abx if more virulent bugs (*S. aureus*, *S. pyogenes*)



# Pneumonia- Inpt Mgmt Basics

1st line (generally)- ampicillin

Broaden your microbial ddx-

- *S. pneumo* still most common
- Flu with superimposed bacterial PNA: suspect *S. aureus*
- Rapidly progressive or sepsis/shock: consider *S. aureus* or *S. pyogenes*
- Likely do not need to cover for HiB or *M. catarrhalis* (unless under-immunized)
- Complicated dz: *consider* ceftriaxone first line plus staph coverage. Amp OK for simple effusion.

PCN allergy- ceftriaxone

- Ceftriaxone other cases: under-immunized, high local PCN-R strains, severe/complicated disease

Beta-lactam allergy- levofloxacin

- Confirmed atypical- \*azithro
- MRSA coverage- Vanc. Add clinda for toxin-mediated (complicated) process.
- Duration-
  - Uncomplicated: 7-10d
  - Complicated: longer
- Steroids- Consider in asthma pt when signs of reversible obstruction
- Chest physiotherapy- no benefits in outcomes or LOS
- No improvement or worsening 48-72 hours of appropriate abx- assess for other cause, consider resistance, look for complications

\*no change in outcomes when azithro used



# Pneumonia- Inpt Mgmt:

## Complications

- Will save most of this for another talk!
- Complicated CAP- effusion, empyema, abscess, fistula, necrotizing PNA.
- Few pointers-
  - Small uncomplicated effusion <10 mm on lateral CXR or <1/4 opacification of hemithorax: can still tx empirically only w amp
  - Broaden abx coverage beyond 3 main suspects: *HiB*, *mycoplasma*, *Legionella/Aspergillus* (necrotizing), anaerobes/*Klebsiesla* (abscess)



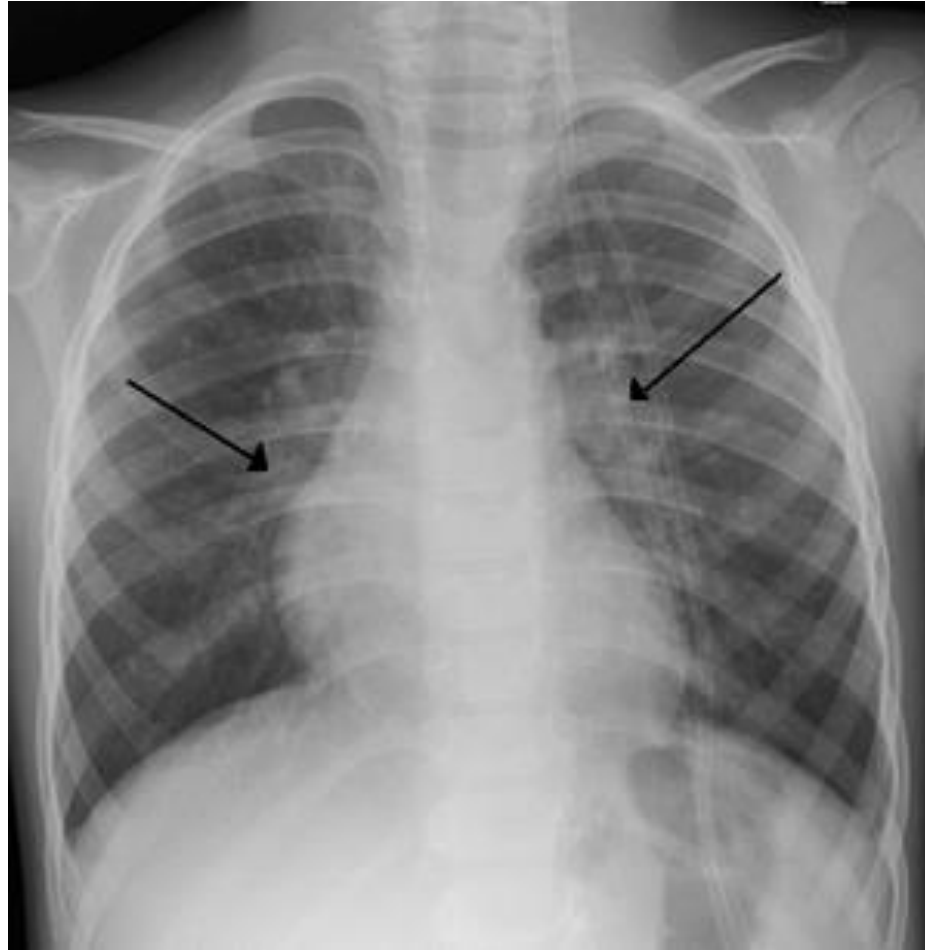
# Pneumonia- Summary

- Reserve labs/imaging for potentially hospitalized pt
- Bacteria only involved in 15% of hospitalized PNA pt's; increases with age
- Hints at etiology-
  - Age-
    - mycoplasma only 5% in <5 y/o; increases substantially age 5
    - Bacterial is 8-12% of cases age 0-4
  - Signs & Symptoms-
    - Hints at viral/atypical-
      - Preceding/coincident URI/constitutional symptoms
      - Wheezing
      - Diffuse exam
      - Mild illness/appearance
- Treatment-
  - Abx not indicated in preschool age (unless strong evidence for bacteria)
  - Outpt: amox 33 mg/kg TID (MDD 4 g) for 5-7 days
  - Inpatient- amp first line. Broaden your ddx/pathogens. *Consider* azithro only for *proven mycoplasma*





# Bronchiolitis



# Bronchiolitis- Epidemiology

- 90% of kids infected with RSV in first 2y;  
40% of these will have LRTI
- Most common cause of infant hospitalization:  
100,000 hospitalizations annually in US =  
\$1.73 billion
- Preterm infants have higher hospitalization  
rates



# Bronchiolitis- Definition, Pathophysiology, Clinical

- Common LRTI in infants, characterized by acute inflammation, edema, epithelial necrosis, increased mucus production in small airways (bronchioles)
- Almost always virally-mediated: RSV >> rhinovirus > flu > hMPV > paraflu/others
- Usually URI prodrome followed by respiratory distress (retractions, grunting, tachypnea, nasal flaring), wheezing, hypoxia, extrapulmonary signs/symptoms (AMS, decreased PO)



# Bronchiolitis- Mgmt

- Assess severity and risk factors for severe dz
  - Severe dz RF's: age < 12 w/o, prematurity, immunodeficiency, underlying cardio-pulm dz
- **Diagnose clinically; labs/imaging not routinely recommended**
  - Viral testing for infants on palivizumab to assess breakthrough RSV
    - *Can help when dx is in question*
  - CXR- often abnormal, but clinical significance unclear
    - Again, may assist if diagnosis is in question
    - LRTI kids with CXR were more likely treated with abx without difference in clinical outcome
- Pulse ox- unclear significance of hypoxia (<90%) on outcomes or borderline low (90-95%) on progression or need for follow-up



# Bronchiolitis- What is Tachypnea Anyways?

WHO criteria:

- 0-2 months:  $> 60$
- 2-12 months:  $> 50$
- 1-5 years:  $> 40$
- $> 5$  years:  $> 20$



# Bronchiolitis- Treatment

- Repositioning- Yes
- Albuterol- Not recommended
  - May improve symptoms scores
  - No change in LOS, disease resolution, hospitalization rate
  - May take into account h/o wheezing or atopy
- Hypertonic (3%) saline aerosol- Not in the ED, OK when admitted
- Steroids- Universally contraindicated
- Chest physiotherapy- No
- Suction- Nasal (non-invasive) probably OK. Deep suction may have negative effect
- Antibiotics- no, unless strong suspicion for bacterial infection



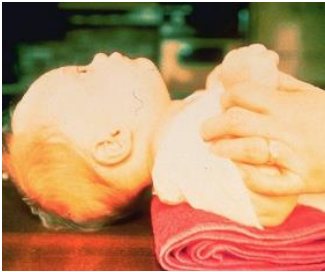


# Bronchiolitis- Treatment (continued)

- Oxygen-
  - Only for persistent sats  $< 90\%$
  - Pulse oximetry is optional
  - High-flow nasal cannula- early data suggest *modest* improvement



# Bronchiolitis Treatment Summary: Always Sometimes, \*Never



Reposition



Oxygen



Hypersaline



Deep suction



Physio-tx



Steroids



Albuterol\*



Antibiotics\*



# Bronchiolitis- Summary

- Clinical diagnosis; labs/imaging only if strong suspicion for something else
  - CXR abnormal in ~25% of pt's without difference in etiology or outcome
- Tx:
  - No routine abx, no steroids ever, no physiotherapy
  - Albuterol: not recommended, but can take into account atopic history
  - Supportive: positioning, suction
  - 3% hypersaline aerosol when admitted
  - O2 only if pulse ox < 90%



# Case Study- Ariel

- HPI: 3 y/o healthy female presents with 1 day h/o respiratory distress after 3-4 days rhinorrhea, congestion, sore throat, cough.
- Came from UC: pulse ox 88%, CXR obtained
- Vitals: T 100.9, RR 36, BP 96/60, P 120, P.O. 92-95%
- Exam: Non-toxic, grumpy, + clear rhinorrhea, b/l lateral cervical nodes, mild oropharyngeal erythema, faint diffuse wheezing, no increased WOB.



# Case Study (continued)

- CXR- bilateral lower lobe opacity (infiltrate vs. atelectasis)



# Case Study (continued)- Admit or Not?

- Any admission criteria?
  - No!

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# Case Study (continued)- Workup?

- Blood work?
  - No since not sick enough for admission
- Viral testing?
  - No since viral identification not going to change management
- Mycoplasma testing?
  - Probably not based on pt's age and classic viral findings/symptoms
- Was it wrong to get the CXR?
  - Probably OK since hypoxic at initial presentation



# Case Study (continued)- Treatment

- Consider observation period in ED for hypoxia redevelopment.
- Could consider bronchodilator if family h/o atopy
- Discharge with instructions for supportive care.



# 4 Points You MUST Remember!

1. Most pediatric CAP is viral, esp in the young
2. Narrow abx are best (superbugs, efficacy) for routine CAP
  - Amox CAP dosing: 33 mg/kg TID (MDD 4 g)
  - Evidence for adding azithro is *tenuous*
3. Reserve labs/imaging for potentially hospitalized patients
4. Bronchiolitis tx is minimal:
  - Yes: reposition, O2 (<90%)
  - Maybe: O2 (<90%)
  - Probably not: Deep suction, hypersaline, abx, albuterol
  - NO: steroids, chest physiotherapy



# With your help, we can:

- Keep our kids safe (duh!)
- Fight the good fight against abx resistance
- Reduce medical costs
- Reduce radiation exposure and other overdiagnosis



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