

Respiratory infections

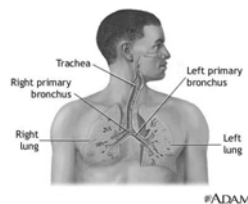
Community acquired pneumonia: a review of common pathogens

Natalie Neu, MD
MID 2008

CAP: general principles

- Presentation
- Etiology
- Specific organisms and pneumonia
 - *Streptococcus pneumoniae*
 - Mycoplasma
 - Chlamydia
 - Legionella
 - Pertussis

Respiratory tract anatomy



Pneumonia: presentation and working up the etiology

- Common complaints
 - Dyspnea, fever, cough (productive or not), chills, chest pain, myalgia, headache
- History
 - Age, co-morbidities, sick contacts, unusual exposures, social situation/support
- Physical exam findings
 - Oxygen saturation
 - Rales, tactile fremitus, decreased breath sounds, rhonchi
- Radiology
 - Confirming the diagnosis; may or may not help narrow the diagnosis e.g. *S. pneumoniae*: lobar; *S. aureus*: multilobar/abscess; Mycoplasma- diffuse interstitial

Community acquired pneumonia: CAP

- 5.6 million cases annually
- #1 cause of death due to infectious diseases in the U.S.
- \$9.7 billion dollars annually
- 3 groups for patient management
 - Outpatient, inpatient (non- ICU), ICU
- Etiology:
 - *Streptococcus pneumoniae* (#1)
 - “Atypical organisms”
 - Viral (e.g. RSV, influenza, adenovirus)
 - Gram negative
 - Other

Bartlett, J. G. et al. N Engl J Med 1995;333:1618-1624

Table 2. Microbiologic Pathogens in Community-Acquired Pneumonia.

Microbial Agent or Cause	Prevalence (%)	
	North American Studies*	British Thoracic Society†
Bacteria		
<i>Streptococcus pneumoniae</i>	20–60	60–75
<i>Haemophilus influenzae</i>	3–10	4–5
<i>Staphylococcus aureus</i>	3–5	1–5
Gram-negative bacilli	3–10	Rare
Miscellaneous‡	3–5	—
Atypical agents	10–20	—
Legionella	2–8	2–5
<i>Mycoplasma pneumoniae</i>	1–6	3–18
<i>Chlamydia pneumoniae</i>	4–6	—
Viruses	2–15	8–16
Aspiration	6–10	—

*Based on 15 published reports from North America.¹⁻¹⁵ None of these studies used techniques adequate to detect anaerobes in respiratory secretions; these organisms account for 20 to 30 percent of cases in some reports.¹⁶⁻¹⁷ *P. carinii* is excluded but may account for up to 15 percent in recent reports from urban centers.¹⁸

†Based on an analysis of 453 adults in a prospective study of community-acquired pneumonia in 25 British hospitals.^{19,20} Dashes indicate that the pathogen was not included in the study.

‡Includes *Moraxella catarrhalis*, group A streptococcus, and *Moraxella meningitidis* (each accounting for 1 to 2 percent of cases).

Table 6. Most common etiologies of community-acquired pneumonia.

Patient type	Etiology
Outpatient	<i>Streptococcus pneumoniae</i> <i>Mycoplasma pneumoniae</i> <i>Haemophilus influenzae</i> <i>Chlamydia pneumoniae</i> Respiratory viruses ^a
Inpatient (non-ICU)	<i>S. pneumoniae</i> <i>M. pneumoniae</i> <i>C. pneumoniae</i> <i>H. influenzae</i> <i>Legionella</i> species Aspiration Respiratory viruses ^a
Inpatient (ICU)	<i>S. pneumoniae</i> <i>Staphylococcus aureus</i> <i>Legionella</i> species Gram-negative bacilli <i>H. influenzae</i>

NOTE. Based on collective data from recent studies [171]. ICU, intensive care unit.
^a Influenza A and B; adenovirus, respiratory syncytial virus, and parainfluenza.

Clinical scenario 1

- Francisco is a 2 year old, previously well
- Presented with URI symptoms and fever to PMD in July
- Respiratory symptoms worsened, cxr revealed right sided pneumonia, WBC 24K with 80% PMN and 3% bands
- Initially treated with IV therapy without resolution in 4 days
- CT scan showed large right sided effusion

Table 8. Epidemiologic conditions and/or risk factors related to specific pathogens in community-acquired pneumonia.

Condition	Commonly encountered pathogen(s)
Alcoholism	<i>Streptococcus pneumoniae</i> , oral anaerobes, <i>Klebsiella pneumoniae</i> , <i>Acinetobacter</i> species, <i>Mycobacterium tuberculosis</i>
COPD and/or smoking	<i>Haemophilus influenzae</i> , <i>Pseudomonas aeruginosa</i> , <i>Legionella</i> species, <i>S. pneumoniae</i> , <i>Moraxella carolinensis</i> , <i>Chlamydia pneumoniae</i>
Aspiration	Gram-negative enteric pathogens, oral anaerobes
Lung abscess	CA-MRSA, oral anaerobes, endemic fungal pneumonia, <i>M. tuberculosis</i> , atypical mycobacteria
Exposure to bat or bird droppings	<i>Histoplasma capsulatum</i>
Exposure to birds	<i>Chlamydia psittaci</i> (if poultry; avian influenza)
Exposure to rabbits	<i>Francisella tularensis</i>
Exposure to farm animals or petting zoo	<i>Coccidia</i> (if fever)
HIV infection (early)	<i>S. pneumoniae</i> , <i>H. influenzae</i> , <i>M. tuberculosis</i>
HIV infection (late)	The pathogens listed for early infection plus <i>Pneumocystis jirovecii</i> , <i>Cryptococcus</i> , <i>Histoplasma</i> , <i>Aspergillus</i> , atypical mycobacteria (especially <i>Mycobacterium kansasii</i>), <i>P. aeruginosa</i> , <i>H. influenzae</i>
Hotel or cruise ship stay in previous 2 weeks	<i>Legionella</i> species
Travel to or residence in southwestern United States	<i>Coccidioides</i> species, <i>Hantavirus</i>
Travel to or residence in Southeast and East Asia	<i>Burkholderia pseudomallei</i> , avian influenza, SARS
Influenza active in community	Influenza, <i>S. pneumoniae</i> , <i>Staphylococcus aureus</i> , <i>H. influenzae</i>
Cough >2 weeks with whoop or posttussive vomiting	<i>Bordetella pertussis</i>
Structural lung disease (e.g., bronchiectasis)	<i>Pseudomonas aeruginosa</i> , <i>Burkholderia cepacia</i> , <i>S. aureus</i>
Injection drug use	<i>S. aureus</i> , anaerobes, <i>M. tuberculosis</i> , <i>S. pneumoniae</i>
Endobronchial obstruction	Anaerobes, <i>S. pneumoniae</i> , <i>H. influenzae</i> , <i>S. aureus</i>
In context of bioterrorism	<i>Bacillus anthracis</i> (anthrax), <i>Yersinia pestis</i> (plague), <i>Francisella tularensis</i> (tularemia)

NOTE. CA-MRSA, community-acquired methicillin-resistant *Staphylococcus aureus*; COPD, chronic obstructive pulmonary disease; SARS, severe acute respiratory syndrome.

IDSA/ATS Guidelines for CAP in Adults • CID 2007;44 (Suppl 2)



Microbial causes of CAP in childhood

Age	Organisms
Birth to 3 weeks	Group B Streptococcal, Gram negative enteric bacilli, Cytomegalovirus, <i>Listeria monocytogenes</i> , HSV
3 weeks- 3 months	<i>Chlamydia trachomatis</i> , Respiratory syncytial virus (RSV), Parainfluenza virus type 3 (PIV), <i>Streptococcus pneumoniae</i> , <i>Bordetella pertussis</i> , <i>Staphylococcus aureus</i>
3 months - 5 years	RSV, PIV, influenza, adenovirus, rhinovirus, <i>Streptococcus pneumoniae</i> , <i>Haemophilus influenzae</i> , <i>Mycoplasma pneumoniae</i> , <i>Mycobacterium tuberculosis</i>
5-15 years	<i>Mycoplasma pneumoniae</i> , <i>Chlamydia pneumoniae</i> , <i>Streptococcus pneumoniae</i> , <i>Mycobacterium tuberculosis</i>

Complicated pneumonia with empyema



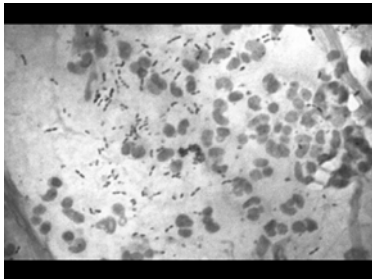
Streptococcus pneumoniae

- Gram-positive; oval or lancet-shaped, occur in pairs or short chains (diplococci)
- Capsular polysaccharide is most important virulence factor; approximately 85 capsular types
- Decreasing incidence but remains the **most commonly isolated pathogen** in patients with pneumonia
- Organism causes pneumonia, meningitis, otitis media, sinusitis, bacteremia, pericarditis, arthritis

S. Pneumoniae

Diagnosis, treatment and prevention

- Diagnosis:
 - Blood culture, urine antigen test, sputum culture
- Treatment: Beta-lactam antibiotics
 - Risk factors resistance in *Streptococcus pneumoniae*
 - Age >65, receipt of β -lactam therapy within 3 months, alcoholism, immune suppression, multiple medical co-morbidities, exposure to child in daycare
 - PCN Resistance classified by breakpoints
 - Sensitive MIC ≤ 0.6
 - Intermediate MIC 0.1-1 mcg/ml
 - Resistant MIC ≥ 2 mcg/ml
 - Cephalosporins, vancomycin, macrolides, linezolid
- Prevention: Vaccines
 - Conjugated pneumococcal vaccine (Prevna[®])
 - 23 valent pneumococcal vaccine (Pneumovax[®])



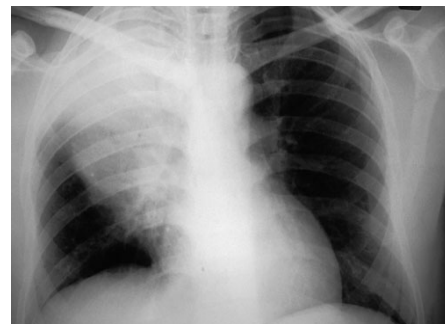
Clinical scenario 2

- Myra is a 21 year old medical student living in the dorm room studying for exams
- She goes to student health complaining of low grade fever, headache, non-productive cough, sore throat and general malaise
- Her exam reveals mild fine inspiratory rales- nothing impressive
- The Dr sends her for an xray that reveals bilateral infiltrates

Structure, Virulence Factors and Pathogenesis

- Capsular polysaccharide is most important virulence factor; approximately 85 capsular types
- Protein adhesins: allow binding to epithelial cells in the oropharynx
- Secretory **IgA protease** - inhibits function of secretory IgA which normally binds bacteria to mucin to facilitate clearance from the respiratory tract
- Pneumolysin - creates pores in and destroys ciliated epithelial cells
- Hydrogen peroxide - reactive O₂ intermediate causes tissue damage
- Teichoic acid, peptidoglycan and pneumolysin activate complement

Mycoplasma pneumonia



Mycoplasma

- Does not have a cell wall
- Cell membrane contains sterols not present in other bacteria
- Special enriched media needed for growth
- Laboratory cultures rarely done- diagnosis usually by serology (IgG)
- Bedside test- cold agglutinins

Erythema multiforme



Mycoplasma- pathogenesis and immunity

- P1- protein attachment factor- facilitates attachment to sialic acid receptors of respiratory epithelium and RBC surface
- Remains extracellular
- Causes local destruction of cilia, interferes with normal airway clearance which leads to mechanical irritation and persistent cough
- Acts as a super antigen stimulating PMS's and macrophages to release cytokines (TNF α , IL1, and IL 6)

Clinical scenario 3

- JM 10 week old infant born to a 16 year old mom
- Pregnancy history limited due to lack of prenatal care but baby born full term, no complications, left hospital 2 days
- Seen by pediatrician at 2 weeks old with eye discharge was given eye drops
- Returned to ER: RR 60, cough but no fever
- Xray done and bloods drawn

Walking pneumonia

- Lacks seasonal pattern, spread by droplet secretions
- Common in children and young adults
- Mild respiratory symptoms
- Complications: otitis media, erythema multiforme, hemolytic anemia, myocarditis, pericarditis, neurologic abnormalities
- Treatment: erythromycin

Chlamydia trachomatis xray



Chlamydial pneumonias: *trachomatis, pneumoniae, psittaci*

- Intracellular parasites- use host high energy phosphate compounds
- Trilaminar outer membrane which contains LPS
- Two phase life cycle- Elementary body (infectious) and reticulate body (divides by binary fission in the host)

C. pneumoniae

- Single strain- TWAR
- Prolong incubation period
- Common in school age children
- Indolent course-sore throat, chronic cough, no fever
- Chest xray variable (lobar, diffuse, bilateral)
- Diagnosis: PCR and serology
- Treatment: macrolide, doxycycline, levofloxacin

Chlamydial pneumonias

- Infect non-ciliated columnar cells
- Multiply in alveolar macrophages
- Perivascular and peribronchiolar infiltrates
- Clinical symptoms due to host immune response
- Immunity not long-lasting
- Diagnosis by serology- four fold rise in titer

C. psittaci

- History: Parrot exposure
- Mild clinical respiratory symptoms, fever, rash
- Concomitant symptoms: cns- headache, confusion, cranial nerve palsy, seizures; hepatitis; pericarditis
- Xray-consolidation, reticular nodular pattern, adenopathy
- Titers: > 1:64 diagnostic
- Treatment: doxy, tetracycline, erythromycin

C. trachomatis pneumonia

- Neonatal infection presents at 1-3 months of age
- Staccato-like cough, rapid respiratory rate
- NO FEVER
- Evaluation: minimal chest findings, xray hyperinflation and diffuse infiltrates, peripheral eosinophilia
- Associations: atherosclerotic heart disease
- Treatment: erythromycin
- Prevention: maternal screening

Psittacosis

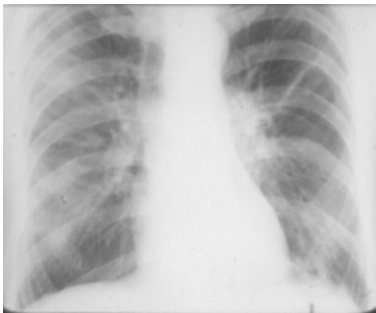


Clinical scenario 4

- Charlie is a 68 year old retired plumber who recently underwent a renal transplantation
- Felt great and was tinkering around his house updating his bathroom fixtures
- Came for follow up visit complaining of high fever, cough, chills and his wife said that he was acting confused at times
- Laboratory studies reveal WBC 35,000 with left shift, LDH >1000
- Chest xray reveals multilobar process

The 1976 Legionnaire's Convention, Philadelphia, PA

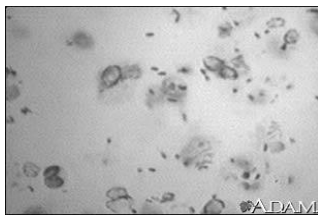
- 29/180 patients died due to pneumonia
- Identification of a gram negative bacilli
- Epidemiologic link to being in the lobby of Hotel A; historical link to 1966 outbreak in a psychiatric hospital
- National panic- worries about biologic and chemical warfare- media frenzy
- 6 months to identify the organism



Legionella pneumophila and micdadei

- 2-6% community acquired pneumonias
- Risk: immunocompromised, hospitalized, and outbreak situations
- Gram negative bacilli- don't stain with common reagents
- Fastidious and grow on supplemented media
- Organisms contaminate water sources: air conditioning systems and water tanks

Legionella species



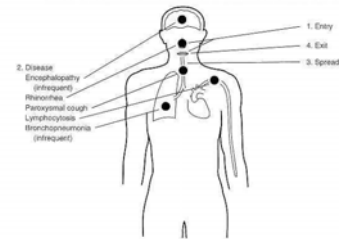
Legionella: pathogenesis and immunity

- Intracellular pathogen- multiply in macrophages and monocytes
- Proteolytic enzymes kill the infected respiratory cells leading to formation of microabscesses
- Immunity- Cell mediated immunity (T cells) needed for immune response

Legionnaires disease

- Incubation period up to 10 days
- Clinical- influenza like illness or severe manifestation= pneumonia
- Fever (105), rigors, cough, headache
- Multilobular infiltrates and microabscesses
- Extrapulmonary manifestations: CNS, diarrhea, abdominal pain, nausea
- High white counts, abnormal liver, renal panel
- High mortality-15-20% depending on host

Bordetella pertussis



Legionella: Diagnosis, prevention and treatment

- Urine antigen detection assays- EIA for *L.pneumophila* only
- Serology >1:128 positive however late development of antibodies
- Culture on special media
- Treatment: macrolide or levofloxacin
- Prevention: hyperchlorination, super heating, continuous copper-silver ionization

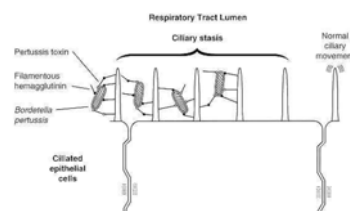
Bordetella pertussis “Whooping cough”

- Fastidious, gram negative coccobacilli
- *Pertussis, parapertussis, and bronchiseptica*
- Spread by respiratory droplets
- Rapid multiplication in mucus membrane
- No bacteremia
- Toxins cause local tissue damage

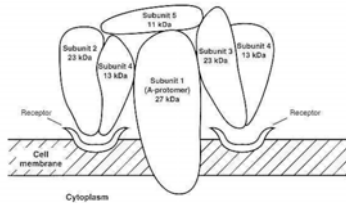
Clinical scenario 5 (Loyola Univ Medical Center)

- Jerry, a 7 month old child, comes to clinic with a running nose, sneezing and slightly irritable
- Diagnosed with URI
- Returns 2 weeks later because he is turning blue with coughing spells. Spells are worse at night, seems to have spasms and then he “whoops” for air.
- Examination reveals mildly dehydrated, not distressed, clear lung exam
- WBC reveals leucocytosis with lymphocytosis

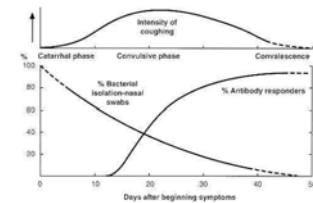
Binding and uptake by phagocytic cells



Pertussis toxin



Pertussis clinical symptoms



G protein and ADP ribosylation

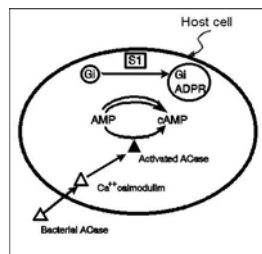
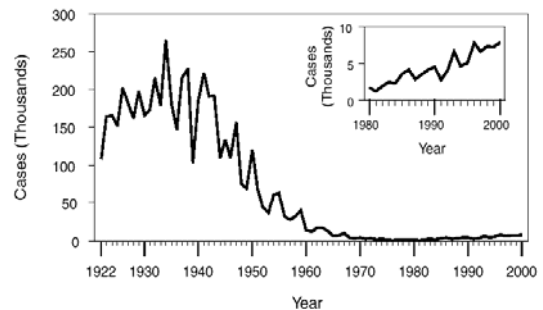
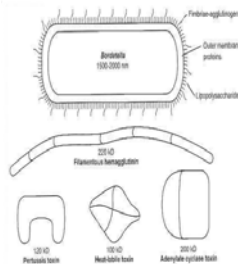


FIGURE 1. Number of reported pertussis cases, by year — United States, 1922–2000



Toxin production and pathophysiology

Pertussis toxin- ↑ cAMP	↑ secretions (paroxysmal stage)
Adenylate cyclase and hemolysin toxin	Inhibit WBC chemotaxis, phagocytosis, and killing
Heat-labile toxin	Local tissue destruction
Tracheal cytotoxin	Destroys ciliated cells, IL-1 (fever), NO (kills epithelial cells)
Lipid A and Lipid X	Activate alternative complement, cytokine release



Pertussis

- Incidence declined due to vaccine
- Affects children under 1 and adults with waning immunity
- New recommendations for booster vaccine for children 11-18 years of age
- Incubation period 7-10 days
- Three stages of disease: catarrhal, paroxysmal, convalescent
- Diagnosis: special media- Bordet-Gengou- blood, charcoal, and starch. Nasopharyngeal culture
- Serologic testing: acute and convalescent titers

CDC 2003 National Hospital Discharge Survey: July 8, 2005

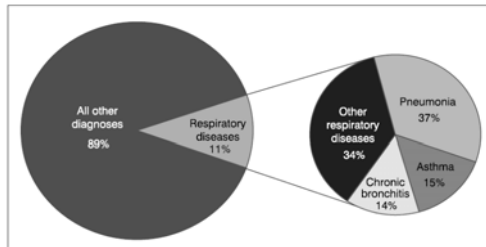


Figure 2. Percent distribution of patients hospitalized for respiratory diseases: United States, 2003

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THE BAD NEWS IS, THERE IS NO CURE FOR THE COMMON COLD. THE GOOD NEWS IS, I THINK YOU HAVE PNEUMONIA.

CAP guidelines: Implementing CAP guidelines locally

Table 2. Elements important for local community-acquired pneumonia guidelines.

All patients
Initiation of antibiotic therapy at site of diagnosis for hospitalized patients
Antibiotic selection
Empirical
Specific
Admission decision support
Assessment of oxygenation
Intensive care unit admission support
Smoking cessation
Influenza and pneumococcal vaccine administration
Follow-up evaluation
Inpatients only
Diagnostic studies
Timing
Types of studies
Prophylaxis against thromboembolic disease
Early mobilization
Thoracentesis for patients with significant parapneumonic effusions
Discharge decision support
Patient education

CID 2007::44

Treatment recommendations for CAP

Outpatient treatment

- Previously healthy and no use of antimicrobials within the previous 3 months
 - A macrolide (strong recommendation; level I evidence)
 - Doxycycline (weak recommendation; level III evidence)
- Presence of comorbidities such as chronic heart, lung, liver or renal disease, diabetes mellitus, alcoholism, malignancy, asplenia, immunosuppressing conditions or use of immunosuppressing drugs; or use of antimicrobials within the previous 3 months in which case an alternative from a different class should be selected
 - A respiratory fluoroquinolone (moxifloxacin, gemifloxacin, or levofloxacin 750 mg) (strong recommendation; level I evidence)
 - A β -lactam plus a macrolide (strong recommendation; level I evidence)
- In regions with a high rate ($\geq 25\%$) of infection with high-level (MIC ≥ 16 $\mu\text{g/mL}$) macrolide-resistant *Streptococcus pneumoniae*, consider use of alternative agents listed above in (2) for patients without comorbidities (moderate recommendation; level III evidence)

Inpatients, non-ICU treatment

- A respiratory fluoroquinolone (strong recommendation; level I evidence)
- A β -lactam plus a macrolide (strong recommendation; level I evidence)

Inpatients, ICU treatment

- A β -lactam (ceftriaxone, cefotaxime, or ampicillin-sulbactam) plus either azithromycin (level II evidence) or a respiratory fluoroquinolone (level I evidence) (strong recommendation) for penicillin-allergic patients; a respiratory fluoroquinolone and azithromycin are recommended

Special concerns

- If *Pseudomonas* is a consideration
 - An antipseudomonal β -lactam (piperacillin-tazobactam, ceftazidime, imipenem, or meropenem) plus either ciprofloxacin or levofloxacin (750 mg) or
 - The above β -lactam plus an aminoglycoside and azithromycin or
 - The above β -lactam plus an aminoglycoside and an antipseudomonal fluoroquinolone (for penicillin-allergic patients; substitute azithromycin for above β -lactam) (moderate recommendation; level III evidence)
- If CA-MRSA is a consideration, add vancomycin or linezolid (moderate recommendation; level III evidence)