Microbial Diseases of the Respiratory System

Chapter 24
Microbial Respiratory Infections

• INTRODUCTION
  – Infections of the upper respiratory system are the most common type of human infection.
  – Pathogens that enter the respiratory system may infect other parts of the body by hematogenous spread (Ex: septicemia, meningitis, distant focal infection).
I. Structure and Function of the Respiratory System

• The upper respiratory system consists of the nose, pharynx, and associated structures, such as the middle ear and auditory tubes.
• Coarse hairs in the nose filter large particles from air entering the respiratory tract.
• The ciliated mucous membranes of the nose and upper respiratory system trap airborne particles and remove them from the body.
• Lymphoid tissue, tonsils, and adenoids provide immunity to certain infections.
I. Structure and Function of the Respiratory System

- The ciliary escalator of the lower respiratory system helps prevent microorganisms from reaching the lungs.
- The lower respiratory system consists of the larynx, trachea, bronchial tubes, and alveoli.
- Microbes in the lungs can be phagocytized by alveolar macrophages.
- Respiratory mucus contains IgA antibodies.
Anatomy of Upper and Lower Respiratory Track System
Expectorated sputum, smear, Gram stain, light microscopy, low power view (LPV). Purulence none. Contaminating bacteria and epithelial cells heavy. No pathogens seen. Please submit carefully collected sample of lower respiratory tree material. The sample is saliva, not sputum. There could be several reasons for submission of this sample to the laboratory. The patient could have been poorly directed and simply "spit" into the collection container, or the patient's cough may not be productive of sputum.
B&S 21-4
Gram stain of sputum specimen. A. **This specimen contains numerous polymorphonuclear leukocytes and no visible squamous epithelial cells, indicating that the specimen is acceptable for routine bacteriologic culture.**
II. Normal Microbiota of the Respiratory System

• The normal microbiota of the nasal cavity and throat can include pathogenic microorganisms in a carrier status.

• Don’t cause disease because of competition with predominant microorganisms.

• The lower respiratory system is usually sterile because of the action of the ciliary escalator.
III. Microbial Diseases of the Upper Respiratory System

• Specific areas of the upper respiratory system can become infected to produce pharyngitis, laryngitis, tonsillitis, sinusitis, and epiglottitis.
  – Pharyngitis – sore throat
  – Laryngitis – infected larynx
  – Tonsillitis – inflamed tonsils
  – Sinusitis – infected sinus
  – Epiglottitis – inflammation of the flap like structures of cartilage that prevents swallowed material from entering the larynx – possible life-threatening when inflamed and occludes airway. H. influenzae type b can cause epiglottitis.

• These infections may be caused by several bacteria and viruses, often in combination.

• Most respiratory tract infections are self-limiting.
III. Microbial Diseases of the Upper Respiratory System

• A. Streptococcal Pharyngitis (Strep Throat)
  – This infection is caused by group A β-hemolytic streptococci, the group that consists of the species *Streptococcus pyogenes*.
  – Symptoms of this infection are inflammation of the mucous membrane and fever, tonsillitis, and otitis media may also occur. At least half of pharyngitis cases are caused by viruses.
  – Preliminary rapid clinic diagnosis is made by indirect agglutination tests or next day culture in the micro lab. Definitive diagnosis is based on a rise in IgM antibodies.
  – Penicillin is used to treat streptococcal pharyngitis.
  – Immunity to streptococcal infections is type-specific.
  – Strep throat is usually transmitted by droplets but at one time was commonly associated with unpasteurized milk.
Streptococcal pharyngitis
III. Microbial Diseases of the Upper Respiratory System

• B. Scarlet Fever
  – Strep throat, caused by an erythrogenic toxin-producing *S. pyogenes*, results in scarlet fever.
  – *S. pyogenes* produces erythrogenic toxin when lysogenized by a phage.
    • *Means Strep A has to have a bacterial phage carrying the toxin gene.*
  – Symptoms include a red rash, high fever, and a red, enlarged tongue, peeled skin. Death is a possible outcome.
Scarlet Fever - Caused by *Strep pyogenes*. The rash begins as a facial erythema sparing the area around the mouth, and spreads to the trunk and limbs. The classical appearance is described as a punctate erythema, and is followed by extensive peeling, which may continue for two or three weeks. The tongue, furred at first, later looks raw with prominent papillae.
Scarlet Fever - The throat is generally red and the tonsils swollen and dark red spots of exudate. If the organism is a producer of the erythrogenic toxin, the local signs are accompanied by the punctate erythematous rash of scarlet fever.
Scarlet Fever - In the early stages there is a dense white coating. Later this peels off leaving a raw red surface, with prominent follicles, the “strawberry tongue”.
Scarlet fever rash showing a ‘strawberry tongue’.
Streptococcus pyogenes on blood agar giving small colonies surrounded by a clear zone where the blood cells have been lysed (beta hemolysis).
Fig 1
Gram Stain of *Streptococcus pyogenes*
III. Microbial Diseases of the Upper Respiratory System

• C. Diphtheria - *Corynebacterium diphtheriae*
  
  – Diphtheria is caused by exotoxin-producing *Corynebacterium diphtheriae*.
  
  – *Gram positive non-spore forming pleomorphic rod. Dividing cells often fold into V and Y shapes.*
  
  – Exotoxin is produced when the bacteria are lysogenized by a phage.
  
  – *Many well people are symptomless carriers.*
  
  – A membrane, containing fibrin and dead human and bacterial cells, forms in the throat and can block the passage of air. "diphtheria" means leather
  
  – The exotoxin inhibits protein synthesis, and heart, kidney, or nerve damage may result.
III. Microbial Diseases of the Upper Respiratory System

• C. Diphtheria - *Corynebacterium diphtheriae* (cont.)
  – Laboratory diagnosis is based on isolation of the bacteria and the appearance of growth on differential media.
  – Antitoxin must be administered to neutralize the toxin, and antibiotics can stop growth of the bacteria.
  – Routine immunization in the U.S. includes diphtheria toxoid in the DTaP vaccine. Prior to this diphtheria was the leading killer of children.
  – Slow-healing skin ulcerations are characteristic of cutaneous diphtheria.
    • *Cutaneous diphtheria characterized by skin lesions is fairly common in tropical countries. In US affects mainly lower socio-economic groups.*
    • *There is minimal dissemination of the exotoxin in the bloodstream.*
A diphtheria leathery membrane caused by the diphtheria toxin.
Diphtheria infection by *Corynebacterium diphtheriae* is still common in some developing countries. Clinical manifestations vary between carrier state and life-threatening illness. This photograph shows severe diphtheria with gross swelling and congestion of the whole pharynx and tonsillar area. Dirty white exudate covers both tonsils and is spreading to the posterior pharyngeal wall.
A *Corynebacterium diphtheria* gram stain showing club shaped morphology. Dividing cells can show V or Y shapes.
III. Microbial Diseases of the Upper Respiratory System

D. Otitis Media: an uncomfortable infections of the middle ear.

– Earache, or otitis media, can occur as a complication of nose and throat infections.

– Pus accumulation causes pressure on the eardrum. 8 million cases/yr.

– Bacterial causes include *Streptococcus pneumoniae, Hemophilus influenzae, Moraxella (Branhamella) catarrhalis, Streptococcus pyogenes*, and *Staphylococcus aureus*. 
The ear anatomy
Otitis media. In the early stages the redness is most prominent in the region of the malleus. The most common bacterial causes are *Streptococcus pneumoniae*, and *Hemophilus influenza* with Strep pyogenes and Staph aureus less common. A small portion are caused by *Moraxella catarrhalis*. 
Acute otitis media. Advanced stage with bulging drum. These appearances are seen just before the drum perforates.
Acute otitis media with bulging eardrum.
Direct smear of otitis media specimen illustrating intracellular gram-negative diplococci. The organism was identified biochemically as *M. catarrhalis* from cultures.
IV. Viral Disease of the Upper Respiratory System

A. The Common Cold - rhinoviruses (50%), corona viruses (15-20%)

- Any one of approximately 200 different viruses can cause the common cold; rhinoviruses cause about 50% of all colds.
  - Symptoms include sneezing, nasal secretions, and congestion.
  - Sinus infections, lower respiratory tract infections, laryngitis, and otitis media can occur as complications of a cold. Usually no fever in uncomplicated cases.
  - Colds are most often transmitted by indirect contact.
  - Rhinoviruses prefer temperatures slightly lower than body temperature.
  - The incidence of colds increases during cold weather, possibly because of increased interpersonal indoor contact or physiological changes.
  - Antibodies are produced against the specific viruses.
V. Microbial Diseases of the Lower Respiratory System

• Anatomy & Collection of Specimen from Lower Respiratory System

• Lower respiratory tract starts at the larynx.
Tracheal secretions received in the laboratory in a Lukens trap. Tracheostomy - Do get colonization- Not significant, but if get into lungs and can cause pneumonia.
Overview of obtaining a protected catheter bronchial brush during a bronchoscopy examination.  **Top left,** The bronchoscope is introduced into the nose and advanced through the nasopharyngeal passage into the trachea. The bronchoscope is then inserted into the lung area of interest.  **Top right,** A small brush that holds 0.01 to 0.001 ml of secretions is placed within a double cannula. The end of the outermost tube or cannula is closed with a displaceable plug made of absorbable gel. The cannula is inserted to the proper area.  **Lower right,** The brush is then extended beyond the inner cannula and the specimen is collected by brushing the area involved. The brush is withdrawn into the inner cannula, which is withdrawn into the outer cannula to prevent contamination by upper airway organisms as it is removed.
A perinasal swab - A thin malleable wire with a small cotton/wool wisp. The swab is gently passed through the anterior nares, quickly twisted and withdrawn and transported immediately to the laboratory in transport medium containing charcoal.
VI. Bacterial Diseases of the Lower Respiratory System

• Many of the same microorganisms that infect the upper respiratory system also infect the lower respiratory system.

• Diseases of the lower respiratory system include bronchitis and pneumonia.

• *Pneumonia is a term applied to severe complications of bronchitis with the alveoli involved.*
VI. Bacterial Diseases of the Lower Respiratory System

• A. Pertussis (Whooping Cough) - *Bordetella pertussis*
  – Pertussis is caused by *Bordetella pertussis*, a Gram negative coccal bacillus
  – The initial stage of pertussis resembles a cold and is called the catarrhal stage.
  – The accumulation of mucus in the trachea and bronchi causes deep coughs characteristic of the paroxysmal (second) stage.
  – Trying to clear mucous leads to violent coughing with gasping (whooping) for breaths. Ciliary action is blocked by accumulation of dense masses of these bacteria in the trachea and bronchi.
  – The convalescence (third) stage can last for months.
  – Laboratory diagnosis is based on isolation of the bacteria on enrichment and selective media (or PCR), followed by serological tests.
  – Have to collect specimen with thin wire with a special swab. See slide below.
  – Regular immunization for children has decreased the incidence of pertussis. The acellular vaccine has minimal side effects vs. the whole cell.
Ciliated cells of the respiratory system infected with *Bordetella pertussis*. Cilia will eventually be destroyed.
VI. Bacterial Diseases of the Lower Respiratory System

• B. Tuberculosis - Mycobacterium tuberculosis.
  – Tuberculosis is caused by *Mycobacterium tuberculosis*, a slow growing acid-fast bacillus
  – Large amounts of lipids in the cell wall account for the bacterium's acid-fast characteristic as well as its resistance to drying and disinfectants.
  – *M. tuberculosis* may be ingested by alveolar macrophages. If not killed, the bacteria reproduce in the macrophages. Resistant to phagocytosis.
  – Lesions formed by *M. tuberculosis* are called tubercles; dead macrophages and bacteria form the caseous lesion that might calcify and appear in an X ray as a Ghon complex. Ghon complex means calcified caseous lesions. *Caused by hypersensitivity reaction.*
  – Liquefaction of the caseous lesion results in a tuberculous cavity in which *M. tuberculosis* can grow.
  – New foci of infection can develop when a caseous lesion ruptures and releases bacteria into blood or lymph vessels; this is called miliary tuberculosis.
  – Miliary (characterized by lesions resembling millet seeds) tuberculosis is characterized by weight loss, coughing, and loss of vigor.
VI. Bacterial Diseases of the Lower Respiratory System

• B. Tuberculosis - Mycobacterium tuberculosis (cont.).
  – Chemotherapy usually involves two drugs taken for 1-2 years; multidrug-resistant *M. tuberculosis* is becoming prevalent. DOT therapy.
  – Drugs are Streptomycin, rifampin, isoniazid (INH) and ethanbutol.
  – *Intracellular growth shields TB from antibiotics.*
  – *Stress and genetic differences contribute to susceptibility.*
  – A positive tuberculin skin test can indicate either an active case of TB, or prior infection, or vaccination and immunity to the disease.
  – Sensitized T-cells are present at site of infection or skin test.
  – Laboratory diagnosis is based on the presence of acid-fast bacilli and isolation of the bacteria, which requires incubation of up to 8 weeks.
  – *Mycobacterium bovis* causes bovine tuberculosis and can be transmitted to humans by unpasteurized milk.
  – *M. bovis infections usually affect the bones or lymphatic system.*
  – *BCG vaccine for tuberculosis consists of a live, avirulent culture of M. bovis. Useful if given early in childhood. Mainly outside of US.*
  – *M. avium-intracellulare* complex infects patients in the late stages of HIV infection.
Mycobacterium tuberculosis acid fast stain, showing filamentous red-stained funguslike growth. Smear from lung tissue.
Expectorated sputum, concentrated smear, fluorochrome acid-fast stain, fluorescent microscopy, MPV. Typical acid-fast bacteria, numerous. Impression: Tuberculosis. The patient had been placed in respiratory isolation following physical examination and history and was immediately begun on antituberculosis therapy following receipt of the direct examination report. *Mycobacterium tuberculosis* was identified from culture.
Mycobacterium tuberculosis growing on Lowenstein-Jensen (LJ) medium. The medium contains egg, mineral salts and malachite (green). Growth appears after about 4 weeks incubation as granular buff-colored colonies.
Progression of a *Mycobacteria tuberculosis* infection of the lung.
Tuberculosis - This radiograph is typical of advanced chronic pulmonary disease. There is extensive nodular shadowing in both lungs, especially in the upper zones. Both upper lobes are contracted and there is cavitation in the right upper lobe.
Tuberculosis - Bones and joints may be affected by tuberculosis at any time in the natural history of the infection. There may be adjacent caseation and bone disease may present as a cold abscess, as in this patient with an indolent swelling of the chest wall. Radiographs showed tuberculosis of the underlying ribs.
A positive tuberculin skin test. Note the induration diameter of $\geq 10$mm.
VI. Bacterial Diseases of the Lower Respiratory System

• C. Bacterial Pneumonias-
  – 1. *Streptococcus pneumoniae*
    • *Pneumococcal pneumonia is caused by encapsulated* *Streptococcus pneumoniae*.
    • *Symptoms are fever, breathing difficulty, chest pain, and rust-colored sputum.*
    • *The bacteria can be identified by the production of* *alpha-hemolysins, inhibition by optochin, bile solubility, and through serological tests.*
    • *A vaccine consists of purified capsular material from 23 serotypes of S. pneumoniae.*
Encapsulated Streptococcus pneumoniae
Expectorated sputum, smear, Gram stain, light microscopy, MPV. Purulence light. Amorphous debris moderate. Gram-positive diplococci, encapsulated, extracellular. Impression: Pneumococcal disease. This is a typical smear presentation for early pneumococcal pneumonia. The pneumococci have proliferated to high numbers, and the lung is responding with increased mucus and fluid release.
C. Bacterial Pneumonias-

- 2. *Hemophilus influenzae*
  
  - Alcoholism, poor nutrition, cancer, and diabetes are predisposing factors for *H. influenzae* pneumonia.
  
  - *H. influenzae* is a small gram-negative coccobacillus.
VI. Bacterial Diseases of the Lower Respiratory System

• C. Bacterial Pneumonias-

  — 3. *Mycoplasmal Pneumonia* - *Mycoplasma pneumoniae*

  • *Mycoplasmal pneumonia* is common in children and young adults; as many as 20% of cases. Lowgrade fever, cough of long 2-3 weeks duration.

  • *Organism has no cell wall* so is difficult to grow so presents like a viral infection.

  • Described as “atypical “ or “walking pneumonia”.

  • *M. pneumoniae* produces small fried-egg colonies after 2 weeks' incubation on enriched media containing horse serum and yeast extract.

  • A complement-fixation test, used to diagnose the disease, is based on the rising of antibody titer.

  • Treated with tetracyclines.
Figure 21-12.
Electron micrograph of effect of *M. pneumonias* on ciliated tracheal cells. *Left,* infected animal model; *right,* uninfected.
Typical chest radiograph of a patient with a 3-week course of “atypical” pneumonia. Note nonspecific interstitial pneumonia, patch infiltrate delineated by “feathery outline”.

Figure 21-14.
Colonies of *Mycoplasma pneuomoniae* showing the typical ‘fried egg appearance’.
VI. Bacterial Diseases of the Lower Respiratory System

• D. Legionellosis - *Legionella pneumophila*
  
  – The disease is caused by the aerobic gram-negative rod *Legionella pneumophila*.
  
  – Legionnaires Disease due to 1976 outbreak of the American Legion meeting.
  
  – The bacterium can grow in water, such as air-conditioning cooling towers, slow plumbing, and then be disseminated in the air.
  
  – This pneumonia does not appear to be transmitted from person to person. High fever, cough, general symptoms of pneumonia.
  
  – Bacterial culture, FA tests, and DNA probes are used for laboratory diagnosis.
  
  – Treated with erythromycin and rifampin
VI. Bacterial Diseases of the Lower Respiratory System

• E. Psittacosis (Ornithosis) *Chlamydia psittaci*
  
  – *Chlamydia psittaci* is transmitted by contact with contaminated droppings and exudates of fowl.
  
  – Psittacine birds such as parakeets and parrots but other birds are infected too.
  
  – Are obligate intracellular parasites but have an elementary body stage that allows the bacteria to survive outside a host.
  
  – Commercial bird handlers are most susceptible to this disease. Example: chicken and turkey farms
  
  – The bacteria are isolated in embryonated eggs, mice, or cell culture; identification is based on FA staining.
  
  – Treated with tetracycline.
*Chlamydia psittaci* detection from direct sputum smear using fluorescent labeled monoclonal antibody, highlighting cytoplasmic inclusions (magnification x700)
VI. Bacterial Diseases of the Lower Respiratory System

• F. Other Bacterial Pneumonias
  – Gram-positive bacteria that cause pneumonia include *Staph aureus* and *Strep pyogenes*.
  – Gram-negative bacteria that cause pneumonia include *Moraxella (B.) catarrhalis, Klebsiella pneumoniae*, and *Pseudomonas species*.
Bronchoaveolar lavage, cytocentrifuge preparation, Gram stain, light microscopy, HPV. Purulence moderate. Gram-positive cocci, pairs, short chains, groups, intracellular. Impression: Staphylococcal disease. The background supports infection, the specimen is directly from the alveolar spaces of the lung, and the organism morphology is typical for staphylococci. Routine bacterial culture yielded a pure growth of methicillin-resistant Staphylococcus aureus (MRSA
Aspirated sputum, smear, Gram stain, light microscopy, MPV. Purulence light. Amorphous debris heavy. Gram-positive cocci, pairs, encapsulated, extracellular. Initial antibiotic therapy can be directed toward streptococci and staphylococci (Stomatococcus). Routine bacterial culture isolated a pure growth of an encapsulated strain of Streptococcus pyogenes. The heavy amorphous background is protein-rich edema fluid from the capillary bed damaged by S. pyogenes toxins. The patient subsequently died from the infection despite correct antibiotic therapy and a correct diagnosis.
Gram stain of bronchial specimen positive for a nonfermentative gram-negative bacillus, e.g., *Pseudomonas aeruginosa*. 
VII. Viral Diseases of the Lower Respiratory System

• A. Viral Pneumonia
  – A number of viruses can cause pneumonia as a complication of infections such as with viral influenza.
  – The etiologies viral pneumonia are not usually identified in a clinical laboratory because of the difficulty in isolating and identifying viruses.
    • If no other cause is found, for example negative bacteria and fungus cultures and mycoplasmal pneumonia has been ruled out, a viral etiology may be assumed.
VII. Viral Diseases of the Lower Respiratory System

• B. **Respiratory Syncytial Virus (RSV)**
  - RSV is the most common cause of pneumonia in infants with 110K hospitalizations and 4500 deaths in US/year. Epidemics appear in winter/early spring.
    - Cough and wheezing for > 1 week. Fever if bacteria become involved.
    - Rapid testing using serology. Nearly 100% show evidence of infection by age 2 with 1% requiring hospitalization.
    - Ribavirin may be given in aerosol to diminish symptoms.
CPE of RSV on Hep2.
VII. Viral Diseases of the Lower Respiratory System

• C. **Influenza (Flu) Influenza virus**
  
  – Influenza is caused by influenza virus and is characterized by chills, fever, headache, and general muscular aches.
  
  – Hemagglutinin (H) and neuraminidase (N) spikes project from the outer lipid bilayer of the virus.
  
  – Viral strains are identified by antigenic differences in the H and N spikes; they are also divided by antigenic differences in their protein coats (A, B, and C).
  
  – Viral isolates are identified by hemagglutination-inhibition tests and immunofluorescence testing with monoclonal antibodies.
  
  – **Antigenic shifts** that alter the antigenic nature of the H and N spikes make natural immunity and vaccination of questionable value and may lead to pandemics. Minor antigenic changes are caused by antigenic drift and are usually covered by yearly vaccines.
  
  – Deaths during an influenza epidemic are usually from secondary bacterial infections.
  
  – Multivalent vaccines are available for the elderly and other high-risk groups that are 70-90% effective for no more than 3 years for that strain.
  
  – Amantadine and rimantadine given early, reduce symptoms against A-type Influenzavirus. Also zanamivir Relenza) and oseltamivir (Tamiflu).
Detailed diagram of the influenza virus. Note the H and N spikes used to identify viral strains. If the H and N spikes change, this leads to an antigenic shift with possible pandemic implications.
Influenza virus types (protein coats) and their antigenic subtypes (H and N spikes).

**TABLE 24.2**  
**Human Influenza Viruses***

<table>
<thead>
<tr>
<th>Type</th>
<th>Antigenic Subtype</th>
<th>Year</th>
<th>Disease Severity</th>
</tr>
</thead>
<tbody>
<tr>
<td>A</td>
<td>H3N2 (the first “modern” pandemic; originated in southern China)</td>
<td>1889</td>
<td>Moderate</td>
</tr>
<tr>
<td></td>
<td>H1N1 (Spanish)</td>
<td>1918</td>
<td>Severe</td>
</tr>
<tr>
<td></td>
<td>H2N2 (Asian)</td>
<td>1957</td>
<td>Severe</td>
</tr>
<tr>
<td></td>
<td>H3N2 (Hong Kong)</td>
<td>1968</td>
<td>Moderate</td>
</tr>
<tr>
<td></td>
<td>H1N1 (Russian)†</td>
<td>1977</td>
<td>Low</td>
</tr>
<tr>
<td>B</td>
<td>None</td>
<td>1940</td>
<td>Moderate</td>
</tr>
<tr>
<td>C</td>
<td>None</td>
<td>1947</td>
<td>Very mild</td>
</tr>
</tbody>
</table>

*The conventional wisdom is that H1, H2, and H3 are human-infecting strains; H4, H5, H6, and H7 primarily infect animals, especially swine and poultry. (Avian influenza strains H5N1 and H7N7 have caused human fatalities.)

†Probably escaped from a laboratory. At this time persons over age 20 were mostly immune from similar viruses circulating in the 1950s and earlier in the century.

VIII. Fungal Diseases of the Lower Respiratory System

• Fungal spores are easily inhaled; they may germinate in the lower respiratory tract.
• The incidence of fungal diseases has been increasing in recent years.
• The mycoses below can be treated with imidazole drugs (itraconazole) and amphotericin B (serious life threatening cases as this drug is very toxic).
VIII. Fungal Diseases of the Lower Respiratory System

A. Coccidioidomycosis: San Joaquin Valley Fever
   - Inhalation of the airborne arthrospores of *Coccidioides immitis* can result in coccidioidomycosis, typically a limited minor lung infection. Found in dry soils of SW US.
   - Most cases are subclinical with about 1% of cases progressing to a TB like course of illness. Predisposing factors such as fatigue and poor nutrition, older age, HIV may lead to disseminated disease.
   - See dimorphic forms in tissue: spherules with yeastlike endospores.
   - Same treatment as Histo using fluconazole (difulcan), ketoconazole, and itraconazole. Sometimes use Amphotericin B to treat serious cases but it is very toxic.
Coccidioides immitis – 2 wks growth at 28°C. Note its cottony, white appearance.
*Coccidioides immitis* – Room temperature culture: hyphae with barrel shaped – in alternating pattern.
Spherules of *Coccidioides immitis* in tissue (x300).
Coccidioides immitis – Spherule as seen from tissue, exudate, or sputum.
US endemic area for coccidiodomycosis. Note the high incidence that occurs in the San Joaquin Valley.
Life cycle of *Coccidiodes immitis*
VIII. Fungal Diseases of the Lower Respiratory System

• B. Pneumocystis Pneumonia
  – *Pneumocystis jiroveci (carinii)*
  – *Pneumocystis jiroveci*, currently classified as a fungus, is found in healthy human lungs. Taxonomy in question but probably related to yeasts.
  – *Pneumocystis carinii* causes disease in immunosuppressed patients. Transmitted by direct contact. Causes alveoli to become filled with a frothy exudate.
  – Common in AIDS patients or other immunosuppressed conditions (cancer, transplant pts). Untreated infections maybe lethal in these cases.
  – *Pneumocytis* pneumonia is currently being treated with trimethoprim or pentamidine.
Life cycle of *Pneumocystis jiroveci*

1. The mature cyst contains 8 intracystic bodies.
2. The cyst ruptures, releasing the bodies.
3. The bodies develop into trophozoites.
4. The trophozoites divide.
5. Each trophozoite develops into a mature cyst.
*Pnuemocystis jiroveci*. A) cysts of *P. jiroveci* from alveolar smear stained with silver stain. Cysts have a black outline. B) Trophozoite stage from chick embryo epithelial lung preparation. Note the extensions used to extract nutrients from the host cells.
Bronchoalveolar lavage, cytocentrifuge preparation, Gram stain, light microscopy, HPV. Purulence none. Local materials moderate. Alveolar cast composed of gram-negative matrix and intracystic bodies. Morphology consistent with *Pneumocystis carinii*