

# Respiratory tract infection

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## Respiratory tract infection in children

- To know how common this problem in pediatric medicine.
- How to differentiate between upper respiratory tract infection and lower respiratory tract infection.
- To know epiglottitis in details (History, physical examination, etiology, differential diagnosis, management).
- To know the pneumonia (bacterial vs viral)

# MANAGEMENT OF COMMUNITY ACQUIRED PNEUMONIA IN CHILDREN

- Clinical features (How do children with CAP present?)
- Etiology Causes of CAP (virus, bacterial, atypical organism) does the etiology alter by age.
- Investigations.
- Severity assessment
- Managements
- Complications of CAP pneumonia (pneumatocele necrotizing pneumonia)

# Pulmonary TB



- Local Epidemiology vs. inherited epidemiology
- Diagnosis, Intervention, managements.
- How to approach children with PPD (children and family)

## INTRODUCTION

- Children have about 6 respiratory infection per year
- They are the the greatest of all causes of medical morbidity in pediatrics.
- Majority of acute respiratory infections are URTI but infection of lower respiratory tract are sufficiently frequent to pose almost daily problems for clinician caring for children
- Large number of different miroorganisms are capable of infecting the lower respiratory tract produces several respiratory syndromes and illnesses.

# **Etiology**

- Viral: Infleunza,
  Parainfleunza,RSV,Rhinovirus,Entero,
   Corona, Measles, Varicella, Adeno,
   EBV,CMV, Herpes
- Mycoplasma: M. pneumoniae
- Rickettsia: Coxiella burnetii (Q fever)
- Chlymedia

# Etiology (CONt.)

- Bacterial: staph, H flu
   Pneumococcus
- Fungi: Candida, Histoplasma,
   Aspergillus
- Parasites: Pneumocystis carinii,
   Toxoplasmosis

#### TABLE 13.2. Common causes of upper airway obstruction in children.

#### Anatomic Altered level of consciousness (airway muscle laxity) Postextubation airway obstruction Tonsillar hypertrophy Subglottic stenosis (acquired or congenital) Macroglossia Vocal cord paralysis External or internal compression Tumor Hemangioma Hematoma Cyst Papilloma Vascular rings and slings Infectious Laryngotracheobronchitis (croup) Peritonsillar abscess Retropharyngeal abscess **Bacterial tracheitis** Supraglottitis (epiglottitis) Infectious mononucleosis Miscellaneous Postextubation airway obstruction Angioedema Foreign body aspiration Airway trauma

# Upper airway obstruction illneses

## INFECTIOUS CAUSES

**Epiglottis** 

Larynogotracheobronchitis(Croup)

**Bacterial tracheitis** 

Diphtheria

Retropharyngeal abscess

Peritonsillar abscess

# Upper airway obstruction illnesses

• NONINFECTIOUS CAUSES:

Foreign body

Trauma

Angioneurotic edema

Hypocalcemic tetany

Caustic burns

# Upper airway obstruction illnesses

## **Acute Epiglottitis**

- Life- threatening condition characterized by upper airway inflammation and obstruction.
- infection of epiglottis and supraglottic structures.
- High risk of death(7%)
- Most common in male (ration of 2.5 to 1).
- may occur at any age.
- Age incidence 2-7 Y
- Vulnerable population incluse infants less than 12 months elderly more than 85 years old

- Caused by almost always H.Influenza type B(HIB) 90%
- No seasonal predilection in incidence of epiglottitis.
- Risk factors:

absence of immunization against HIB immunocompromised state smoking

Pathology:

Direct invasion by HIB

Cellulitis with marked edema of the epiglottis, aryepiglottic folds, ventricular band and arytenoid.

Edema increase the epiglottis curls posteriorly and inferiorly

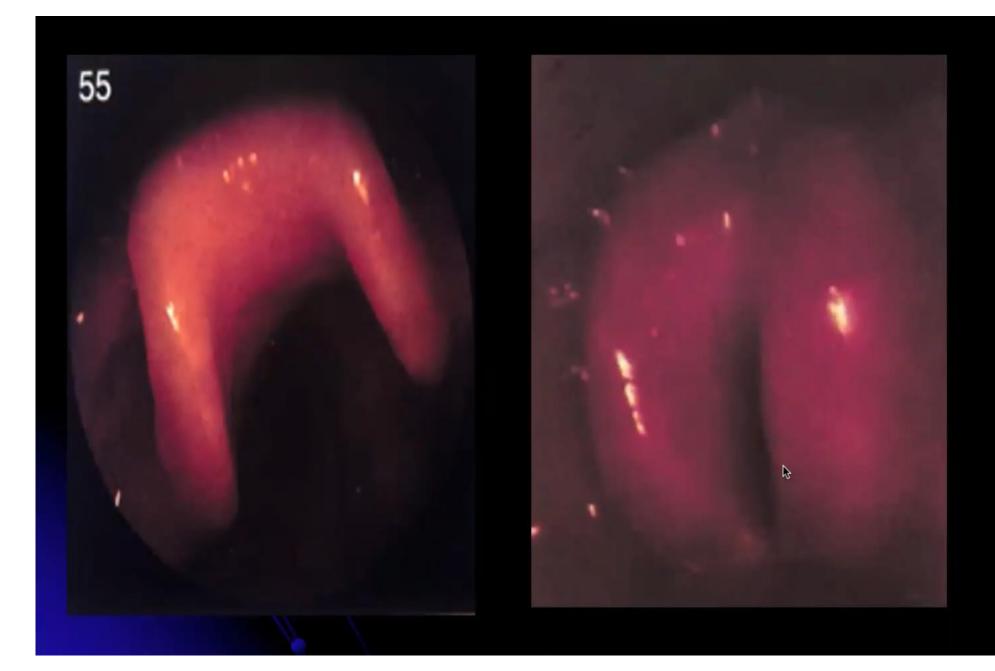
Airway obstruction

(Inspiration tend to draw the inflamed supraglottic ring into the laryngeal inlet)

#### Clinical features:

- Sore throat, followed by Odynophagia accompanied by drooling, retching and difficulty breathing
- Voice is not hoarse but speech is muffled
- Cough is not croupy
- Stidor
- Marked Fever(38.8-40C)
- Child assuming posture that maximize the increase in diameter of the obstructed airway( Sitting& leaning forward with hyperextension of the neck and protrusion of the chin)





- Lateral Neck: Thumb sign
- Treatment

Avoid stimulation of the child. This includes radiographs, drawing bloods, starting IVF until secure airway established

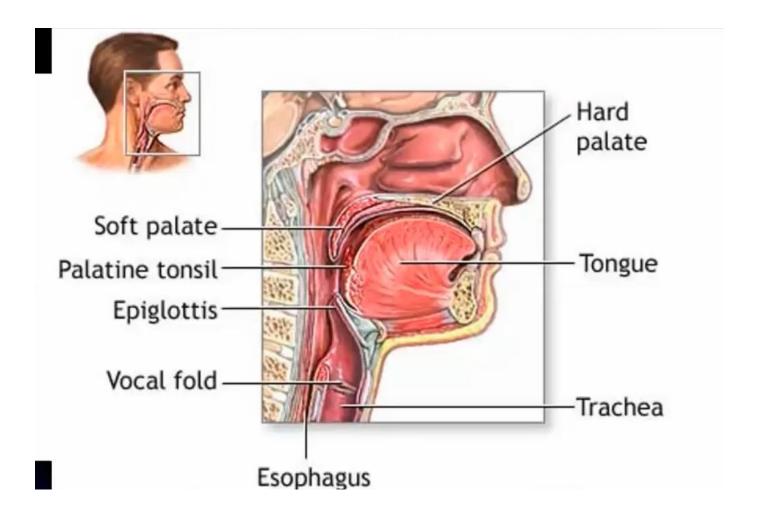
Emergency Intubation: Elective nasopharyngeal intubation (ETT 0.5mm smaller than that required is recommended)

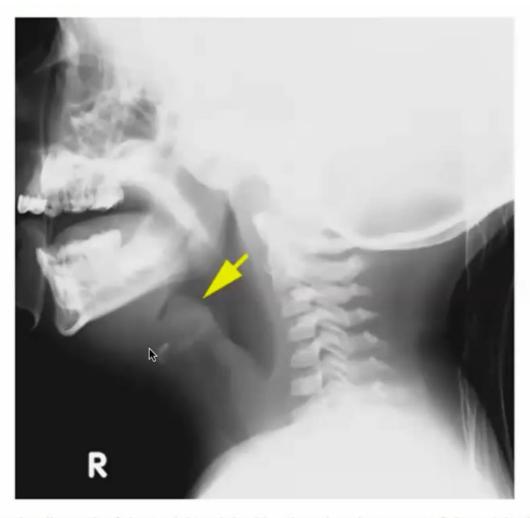
I.V Ceferuxime or cefotaxime for 7 days

NO steroid

Criteria for extubation: Aferbril

swallowing comfortable





Lateral radiograph of the neck in epiglottitis: there is enlargement of the epiglottis (arrow) giving the "thumb sign. (Courtesy of B Poss)

# **CROUP**

- Age: 6months 3 years
- More in fall winter and spring
- URI (Rhinorrhea, Mild to moderate fever
- Progress to inspiratory stridor, hoarseness and croupy cough
- Rib cage and abdominal asynchrony occurs as the condition deteriorates

Am review Resp dis,1990;142,540-544

# Upper airway obstruction illnesses CROUP

Etiology: Parainfluenza 1(most common), 2(less frquent),3(less common) Influenza **RSV** Adeno, rhino, entero, herpes M Peneumonia

# **CROUP**

- Most children have mild disease
- Last 7-14 days
- Among children admitted with croup less than 1% require intubation
- Lateral neck X-ray: Narrowing of subglottic area



Fig 2. AP radiograph of the neck in patient with LTB showing narrowing in the subglottis ("steeple sign"). (Photo courtesy of the Department of Radiology, University of Texas Medical Branch at Galveston.)

# **CROUP**

## Treatment:

Moist air

Oxygen

IV fluid

Steroid therapy: IM,PO, Inhalation (suppression of the local inflammatory reaction, shrinking of lymphoid swelling and reduction in capillary permeability)

#### 0.6mg/kg IM

Aerosols: Recemic epi.; increase the airway diameter within 30 mints, however the effect is short lived lasting for 2 hours

TABLE 13.3. Infectious causes of upper airway obstruction.

	Croup	Epiglottitis	Bacterial tracheitis	Retropharyngeal abscess
Onset	Gradual Viral prodrome 1–7 days	Rapid onset 6–12 hr	Viral prodrome followed by rapid deterioration	Viral prodrome followed by rapid deterioration
Typical age at onset	6 months to 4 years	2–8 years	6 months to 8 years	<5 years
Seasonal occurrence	Late fall to winter	Throughout the year	Fall to winter	Throughout the year
Causative agents	Parainfluenza, respiratory syncytial virus, influenza A	Haemophilus influenzae type b (classically), Streptococcus pneumoniae, GABHS	Staphylococcus aureus (classically), GABHS, Streptococcus pneumoniae	Anaerobic bacteria, GABHS, Staphylococcus aureus
Pathology	Subglottic edema	Inflammatory edema of supraglottis	Thick, mucopurulent, membranou stracheal secretions	Abscess formation in the deep cervical fascia
Fever	Low-grade	High fever	High fever	High fever
Cough	"Barking" or "seal-like"	None	Usually absent	Usually absent
Sore throat	None	Severe	None	Severe
Drooling	None	Frequent	None	Frequent
Posture	Any position	Sitting forward, mouth open, neck extended ("tripod position")	Any position	Sitting forward, mouth open, neck extended ("tripod position")
Voice	Normal or hoarse	Muffled	Normal or hoarse	Muffled
Appearance	Nontoxic	Toxic	Toxic	Toxic

Note: GABHS, group A  $\beta$ -hemolytic Streptococcus.

## **PNEUMONIA**



## Factor relating to the etiology:

- 1) Host normal or compromised.
- 2) **Age**
- 3) Season
- 4) Environment: animate (human and animals) and inanimate.

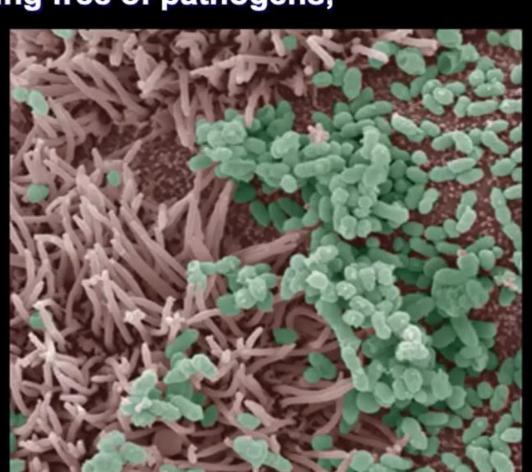
 Innate immunity in the lung is truly amazing. The lung has ~100 square meters of surface area (roughly the size of a tennis court) and is directly exposed to the outside environment with every breath we take. Despite this, the lower airway is normally sterile. There are many levels of innate immunity that keep the lung free of pathogens,

including

filtering in the upper airways,

mucociliary clearance,

antimicrobial factors



#### **PNEUMONIA**

## **Etiology:**

Bacterial: S. pneumonia, H flu, Staph, GAS, TB.

Viral: RSV, parainfl (1,2,3), Inf, adeno, rhino, entero.

Immuno-compromised: Broader spectrum of etiological agents: fungi, gram negative bacili, pneumocystis carini. anaerobes, CMV.

#### Most Common Causes of Pneumonia in Immunocompetent & Immunocompromised Children over 1 month of Age

	IMMUNOCOMPETENT	IMMUNOCOMPROMISED
• Bacterial	Streptococcus pneumonia Haemophilus influenzae Staphylococcus aureus Group A Streprococci Bordetella pertussis Moraxella catarrhalis Yersinia pestis Pasteurella multocida Brucella spp. Francisella tularensis Neisseria meningitidis Salmonella spp.	Pseudomonas spp. Enterobacteriaceae Legionella pneumophilia Nocardia spp. Rhodococcus equi Actinomyces spp. Anaerobic bacteria Enterococcus spp.
Bacteria –     Like Agents	Mycoplasma pneumonia Chlamydia pneumonia Chlamydia trachomatis Chlamydia psittaci Coxiella burnetii	

## VIRAL PNEUMONIA

Most common cause of LRTI, RSV, parainfl (1,2,3), inf, adeno, rhino, entero.

#### Treatment:

Difficult to distinguish from bacterial pneumonia.

Oxygenation and ventilatory assistance in severe cases.

#### Clinical case

#### January...... 1 months......

- An older brother with an upper respiratory tract infection
- 3 days of rhinorrhoea and cough accompanied by low grade fever.
- Admitted to the Emergency Department for an episode of apnoea with mild respiratory distress with retractions and reduced oral intake of fluid (<50%)in the last 12 hrs</li>

#### **CLINICAL DEFINITION**

The diagnosis of bronchiolitis is a clinical one based on typical history and findings on physical examination. Clinicians in different countries use different criteria to diagnose acute bronchiolitis.

A consensus guidelines panel reported a 90% consensus on the definition of bronchiolitis as a seasonal viral illness characterized by fever, nasal discharge and dry, wheezy cough.

On examination there are fine respiratory crackles and/on high pitched expiratory wheeze.



#### Bronchiolitis: Definition



Age <24 months

Signs of viral infection + Wheeze

American Academy of Pediatrics. 2006; 118(4):1774-1793



Age <12 months

Signs of viral infection

+
widespread crackles
+/wheeze

Everard, Ped Clin N Am 2009;56:119

## **BRONCHIOLITIS**

Lower respiratory tract infection in children < 24 months of age.

It involves large and small airways tracheobronchitis, bronchiolitis and alveolar and interstitial lung involvement (pneumonia).

#### Etiology:

Viral: RSV; adenovirus (3,7,21), influenza; parainfluenza (3); rhinovirus; mumps.

Others: Mycoplasma pneumoniae.

## RSV season?

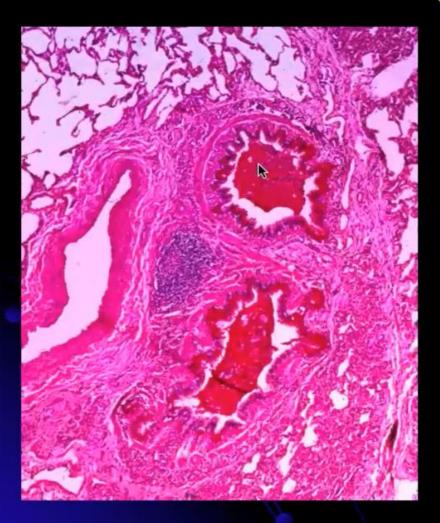
- Ubiquitous throughout the world
- Seasonal outbreaks
  - Temperate Northern hemisphere: November to April, peak January or February
  - Temperate Southern hemisphere: May to September, peak May, June or July
  - Tropical Climates: rainy season
  - In Saudi Arabia RSV appears in November and the seasonal peak occurs during Jan. &Feb.



#### Bronchiolitis: clinical manifestations

- Exposure to children or adults with a respiratory viral infection.
- The initial symptoms are rhinorrhoea, cough and sometimes low grade fever. In 18% of cases the first clinical symptom could be episodes of apnoea.
- With the relief of fever the child manifests tachypnoea, retractions, nasal flaring, rales and hypoxemia
- Dehydration and metabolic acidosis.
- Syndrome of inappropriate secretion of antidiuretic hormone is common with severe respiratory distress.
- It is a dynamic disease and its clinical characteristics can quickly change

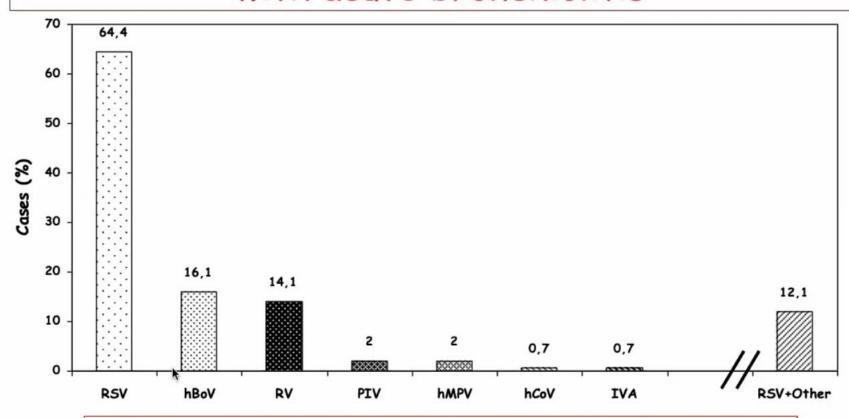
## Pathophysiology



- Upper → lower airways
  - Peribronchiolar mononuclear infiltration
  - Epithelial necrosis
  - Airway plugging
  - Hyperinflation, atelectasis,
     V/Q mismatching
  - Hypoxemia, work of breathing



## Distribution of identified virus in 182 infants with acute bronchiolitis



Midulla et al. Arch Dis Child 2010;95(1):35-41.

### **BRONCHIOLITIS** (cont.)

Pathology: Inflammation of small bronchi and bronchioles, sloughing of resp. ciliated epi.

The bronchioles are plugged with fibrin and mucus – bronchiolar obst. – increase work of breathing + V/Q mismatching – hypoxemia.

CO<sub>2</sub> retention is uncommon, but if present it may lead to assisted ventilation.



#### Pathogenesis

- Mechanical occlusion of terminal and respiratory bronchiole with mucus, fibrin, epithelial cells and inflammatory cells.
- Effects of the immunological reaction and of inflammatory mediators.



### **BRONCHIOLITIS** (cont.)

#### Complications:

Atelectasis, apnea and respiratory failure. Bronch obliterans (esp. with adeno) &.myocarditis

Majority: Mild to moderate disease lasting 3-10 days. 2% require hospitalization; of those 3-7% develop respiratory failure and 1% die.

B

#### High risk:

Children with Cardiopulmonary disease (e.g. BPD, CF, VSD), immune deficiency and neonates.

## RISK FACTORS OF SEVERITY

- Prematurity
- Low birth weight
- Age less than 6-12 weeks
- Chronic pulmonary disease
- Hemodynamically significant cardiac disease
- Immunodeficiency
- Neurologic disease
- Anatomical defects of the airways

# ENVIRONMENTAL RISK FACTORS

- Older siblings
- Concurrent birth siblings
- Native American heritage
- Passive smoke exposure
- Household crowding
- Child care attendance
- High altitude



#### Bronchiolitis: "Indications for hospitalization"

- Prematurity
- Age <3 months</li>
- Apnoea
- Severe underlying conditions
- Poor feeding (less than 50%)
- Respiratory distress (RR >60/min, nasal flaring, retractions) and cyanosis
- Oxygen saturation <92%</li>

Phase of illness should be considered in the decision for timing of review or admission to hospital

### BRONCHIOLITIS (cont.)

#### Treatment:

Admit: if sig respiratory distress, dehydration, underlying disease.

O2 sat, CXR, NP aspirate.

Oxygen, IPPV (apnea, fatigue)

IV fluid

BD: 30% respond to salbutamol.

Steroids: not recommended.

Ribvirin: for RSV, Inf A & B to high risk group, given nebulization 12-18 hr/day for 3-7 days

Recemic epi: .

Martin Chalumeau Laurence Foix-l'Helias Pierre Scheinmann Pierre Zuani Dominique Gendrel Hubert Ducou-le-Pointe

## Rib fractures after chest physiotherapy for bronchiolitis or pneumonia in infants



Table 1. Summary of medical histories and radiological signs

Patient	Age (months) at the time of chest physiotherapy	Chest physiotherapy indication	Age (months) at the time of rib-fracture diagnosis	Fractured tibs
ı	1	Bronchiolitis	2	Lateral left 3rd to 7th
2	1	Pneumonia	2	Posterior right 6th, 7th Posterior left 7th, 8th
3	2	Bronchiolitis	3	Lateral left 7th
4	2	Bronchiolitis	3	Lateral right 3rd, 4th
5	6	Bronchiolitis	7	Lateral right 3rd, 6th

#### TABLE 1

#### Prophylaxis with Palivizumab or RSV-IG\*— Categories of Risk

Infants and children less than two years of age with known CLD who required medical therapy for CLD within six months of the anticipated start of the RSV season.

Preterm infants born at 28 weeks of estimated gestational age or earlier may benefit from prophylaxis during their first RSV season, whenever that occurs during the first year of life, even if CLD is not present.

Preterm infants born at 29 to 32 weeks of estimated gestational age or earlier may benefit from prophylaxis during their first RSV season, whenever that occurs during the first six months of life, even if CLD is not present.

Infants born at 32 to 35 weeks of estimated gestational age must have two of the following risk factors to be candidates for prophylaxis: attendance at a child-care center, school-aged siblings, exposure to environmental pollution, abnormalities of the airways, or severe neuromuscular problems.

RSV-IG = intravenous RSV immune globulin (RespiGam); CLD = chronic lung disease; RSV = respiratory syncytial virus.

<sup>\*—</sup>RSV-IG is contraindicated in children with cyanotic congenital



#### Bronchiolitis: "Indications for hospitalization"

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## RISK FACTORS OF SEVERITY

- Prematurity
- Low birth weight
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- Chronic pulmonary disease
- Hemodynamically significant cardiac disease
- Immunodeficiency
- Neurologic disease
- Anatomical defects of the airways

## Prevention?

- Prevention of spread
  - In hospital: Meticulous infection control
    - Isaacs D. Arch Dis Child 66,p226;1991
  - At home?
- Immunization?
  - Formalin inactivated → worse disease
- Heat inactivated?
- Passive immunity?
  - **RSV-IG**
  - **Palivizimub**



## PREVALENCE OF BACTERIAL INFECTION IN CHILDREN WITH A DOCUMENTED RESPIRATORY SYNCYTIAL VIRUS LOWER RESPIRATORY TRACT INFECTION

#### **METHODS**

In this retrospective study\* the authors reviewed the medical records of children aged <6.5 yrs discharged between July 1, 2000 and June 30, 2002 with a documented diagnosis of respiratory syncytial virus (RSV) infections. The presence and degree of fever at the time or before admission, complete and differential blood cell count and culture results (cerebrospinal fluid, blood culture, suprapubic aspiration urine, catheterised urine) obtained on the first day of hospitalisation were assessed.

#### RESULTS

A total of 912 patients with a median age of 135 days (range 6–2,398 days) were selected. Fever was present at or before admission in 628 (68.9%) of the 912 patients. None of the 63 (95.5%) patients aged <30 days, and none of the 135 (89.4%) patients aged 30–90 days had positive blood cultures. Urine cultures were positive in six of 45 patients aged <30 days, 14 of 94 patients aged 30–90 days and eight of 95 patients aged >90 days. Cerebrospinal fluid cultures obtained in 57.6% of febrile patients aged <30 days, 29.8% of patients aged 30–90 days and 4.4% of patients

#### CONCLUSIONS

Concurrent serious bacterial infections appear to be rare in children with a documented RSV lower respiratory tract infection. Blood and cerebrospinal fluid cultures may not be necessary in nontoxic-appearing infants and young children with fever and a documented RSV lower respiratory tract infection, even in those aged <3 months,



#### Bronchiolitis: discharge criteria

- Oxygen saturation stably remains >90-94%
- Absence of respiratory distress
- Adequately oral intake to prevent (>75% of usual intake) to prevent dehydration
- Adequate parental care and family education

American Accademy of Pediatrics DOI:10.1542/peds.2014-2742

### **MYCOPLASMA PNEUMONIA**

- Peak incidence 5-15 year (account for 75% of pneumonia in this age group)
- C/P: Insidous onset of fever, headache and sore throat followed by dry cough that can last for months.
- Other organs: Meningoencephalitis, carditis, migratory arthralgia and arthritis, hemolytic anemia, +ve coomb's and cold agglutinins.

### MYCOPLASMA PNEUMONIA (cont.)

#### Investigations:

- CXR: Not specific, unilateral or bilateral disease, 20% has pleural effusion.
- CBC: WBC is usually normal.
- Cold agglutinin > 1:64.
- Serology: 4 fold increase in CFT.
- Treatment: Erythromycin, may not alter the duration or sequela (may decrease the duration of cough).



#### **BACTERIAL PNEUMONIA**

#### **Etiology:**

Neonatal period: GBS, listeria monocytogenes and gram –ve bacilli.

After neonatal period: S pneumoniae, H. flu type B, staph. Aureus, GAS.

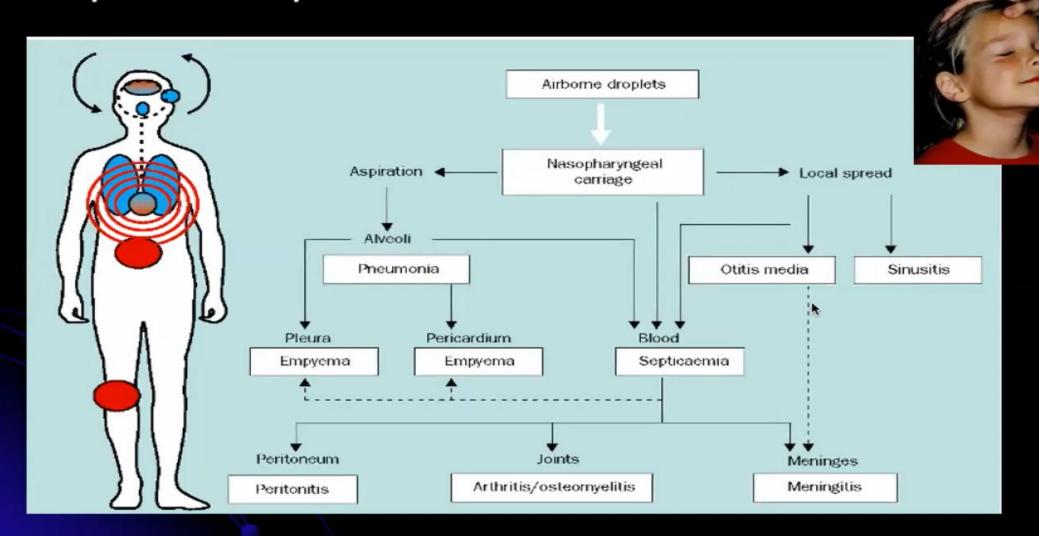
Mycoplasma is quite common >5y.

After age of 4 – 5 years: S. pneumoniae and mycoplasma responsible for the majority of cases.

#### **Bacterial adhesion and invasion:**

قعمات الملك سعود Kmg Saud University

- Streptococcus pneumoniae -



### **BACTERIAL PNEUMONIA (cont.)**

#### Pathology:

Normally the resp. tract is **sterile** below the vocal cords. Pneumonia result from asp. Of pathogen to lower resp. tract. Concurrent viral infection aid this process (present in 30-50% of cases) esp. RSV, measles and influenza.

### **BACTERIAL PNEUMONIA (cont.)**

C/P: Fever, chills, cough, chest and abdominal Pain. Younger infants less specific symptom and signs

#### Diagnosis:

CBC – dif, cold agglutinin

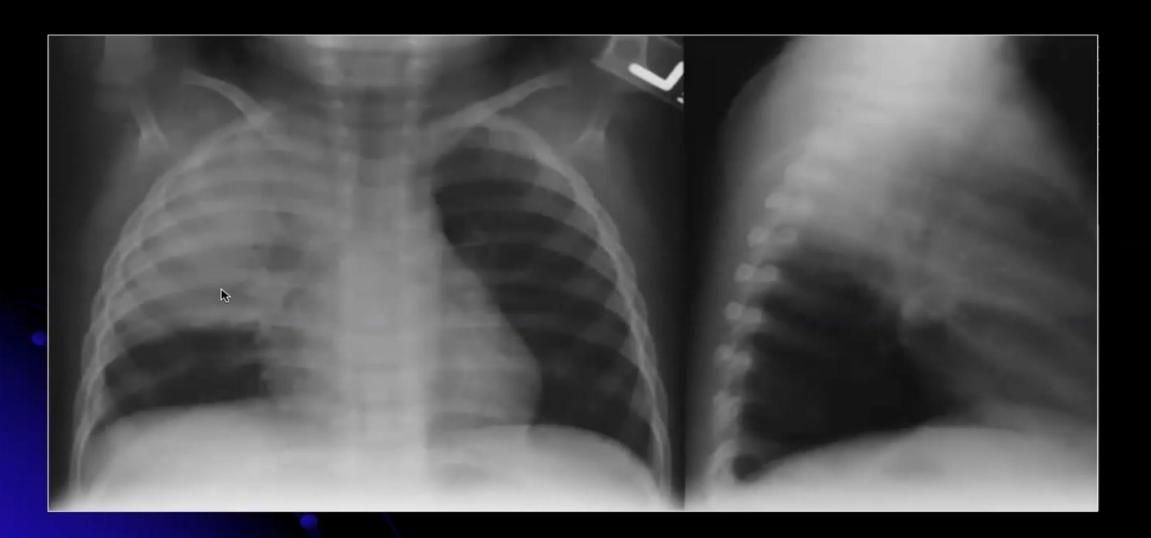
CXR

Blood culture, sputum in older children

#### Treatment:

- 1) Adequate oxygenation.
- Depends on severity and age. Ampi or amoxacillin (10-30% of H flu are resistance) cefuroxime 75-100mg/kg/day
- Older child: Pen or macrolides =>erythro (clarythromycine or zithromax)

## RUL





## What are the indications for transfer to intensive care?

There are two main scenarios when a child is likely to need admission to an intensive care unit:

- (1)When the pneumonia is so severe that the child is developing severe respiratory failure requiring assisted ventilation;
- (2) A pneumonia complicated by septicaemia.

Key features that suggest a child requires transfer include:

- √Failure to maintain oxygen saturation >92% in fractional inspired oxygen of >0.6
- √Shock
- ✓Rising respiratory and pulse rate with clinical evidence of severe respiratory distress and exhaustion, with or without a raised arterial carbon dioxide tension
- ✓ recurrent apnoea or slow irregular breathing.

## Features of severe disease in an older child include:

- ➤Oxygen saturation <92%, cyanosis
- ➤ Respiratory rate >50 breaths/min
- ➤ Significant tachycardia for level of fever (values to define Tachycardia vary with age and with temperature
- ➤ Prolonged central capillary refill time > 2 s
- ➤ Difficulty in breathing; grunting
- ➤ Signs of dehydration
- chronic conditions (eg, congenital heart disease, chronic lung disease of prematurity, chronic respiratory conditions leading to infection such as cystic fibrosis, bronchiectasis, immune deficiency).



### BACTERIAL PNEUMONIA (cont.)

#### Complications:

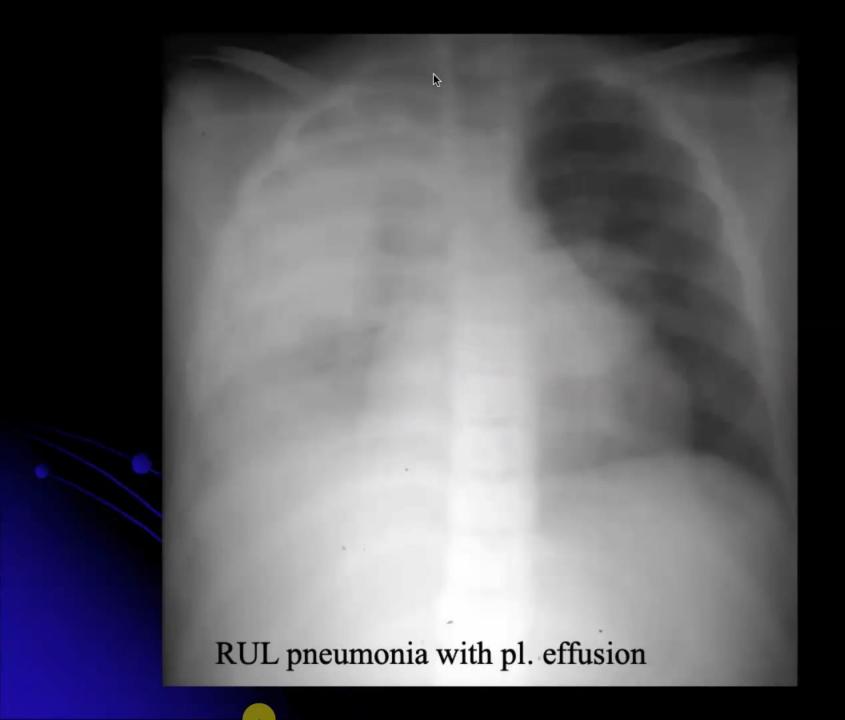
#### 1) Parapneumonic effusion:

Esp. with S. aureus, H flu, S pneumoniae. Can be thin transudate or thick exudates (empyema)

Send pl fluid for cell count, glucose, protein, pH, LDH and culture.

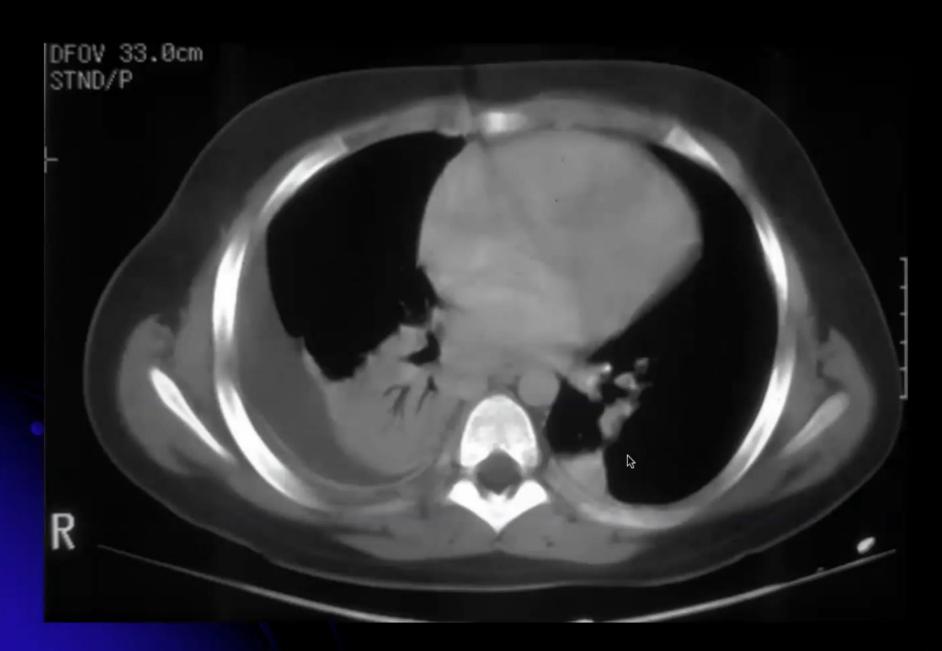
Empyema WBC > 15,000/mm3, protein >3 g/dl, pH <7.2

Management: ABX + drainage, recovery is slow, fever continue for 1 – 2 weeks.

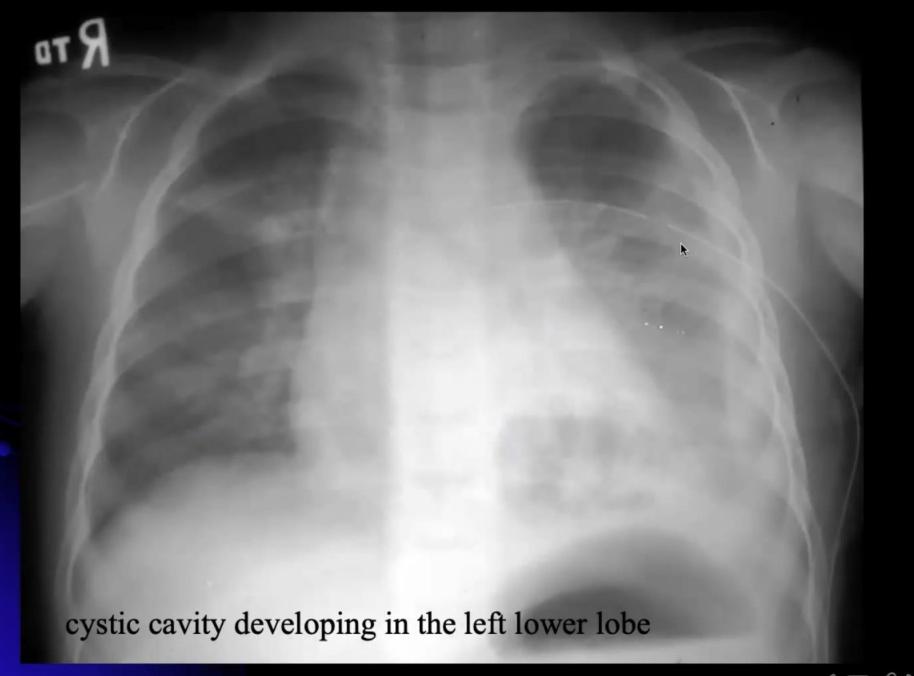








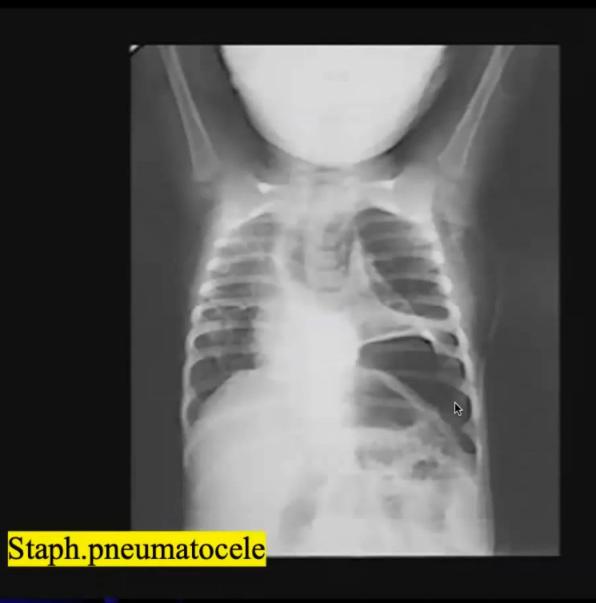




- 2) Pneumatoceles: Thin wall cavity, complicate 40% of staph pneumonia, unusual with other types. Usually asymptomatic unless rupture pneumothorax or pyothorax. Resolve spontaneously within 3 months
- 2) Lung abscess: Esp in aspiration pneumonia in mentally retarded children. Esp. in the dependent portion of the lung.

Growth: mixed anaerobic bacteria

Treatment: Pen G, clinda or flagyl.





#### The predisposing factors to necrotizing pneumonia include

- > Congenital cysts
- **≻**Sequestrations
- > bronchiectasis,
- >neurological disorders
- **≻Immunodeficiency**
- certain serotypes of pneumococcal disease are more likely to lead to necrotizing pneumonia and abscess formation than others
- S aureus with Pantone Valentine leukocidin toxin can lead to severe lung necrosis with a high risk of mortality

#### 3. Septicaemia and metastatic infection

Children can present with symptoms and signs of pneumonia but also have features of systemic infection. Children with septicaemia and pneumonia are likely to require high dependency or intensive care management. Metastatic infection can rarely occur as a result of the septicaemia associated with pneumonia.

Osteomyelitis or septic arthritis should be considered, particularly with S aureus infections.

#### 4. Haemolytic uraemic syndrome

S pneumoniae is a rare cause of haemolytic uraemic syndrome.

A recent case series found that, of 43 cases of pneumococcal haemolytic uraemic syndrome, 35 presented with pneumonia and 23 presented with empyema. Although a rare complication.

in cases with pallor, profound anaemia and anuria, this should be considered.



## TUBERCULOSIS



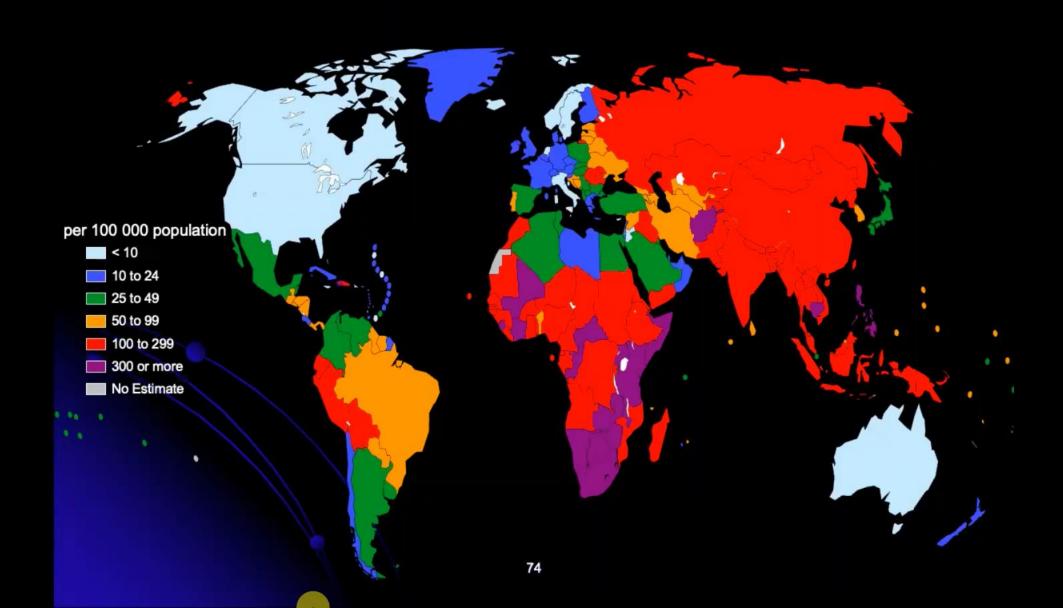
#### **TUBERCULOSIS**

✓ The vast majority of childhood TB occur in children < 4 y usually after exposure to an infected adults.
</p>

(i.e. children infected with TB always have an adult with active TB in their environment).

Transmission is by droplet nuclei

## Estimated TB incidence rate (2003)



## **TUBERCULOSIS** (Cont.)

- Its distribution is worldwide
- Multi-drug resistance has emerged as an important clinical problem
- Infection in patients with HIV infection initially lead to increases in the number of cases

## TUBERCULOSIS (Cont.)

• Adults with cavity harbor a great no. of bacilli for long time. They become noninfectious 2 weeks after therapy.

 Children with primary TB are rarely infectious, TB bacilli are sparse, but they are the long term reservoir of infection in the population.

## TUBERCULOSIS (Cont.)

#### **Etiology:**

- Mycobacterium tuberculosis & M. bovis
   Culture takes 4-6 w, sensitivity another 4 w.
- Radiometric methods, detection & sensitivity
   4-10 d.
- By DNA proves detection within 2 hrs.



# Problems in the diagnosis of TB

#### Active disease

- M. tuberculosis is difficult to isolate: even with good microbiological facilities, the bacillus is recovered in only 50-60% of cases

#### Latent infection

- M. tuberculosis cannot be cultured from latently infected individuals: no gold standard

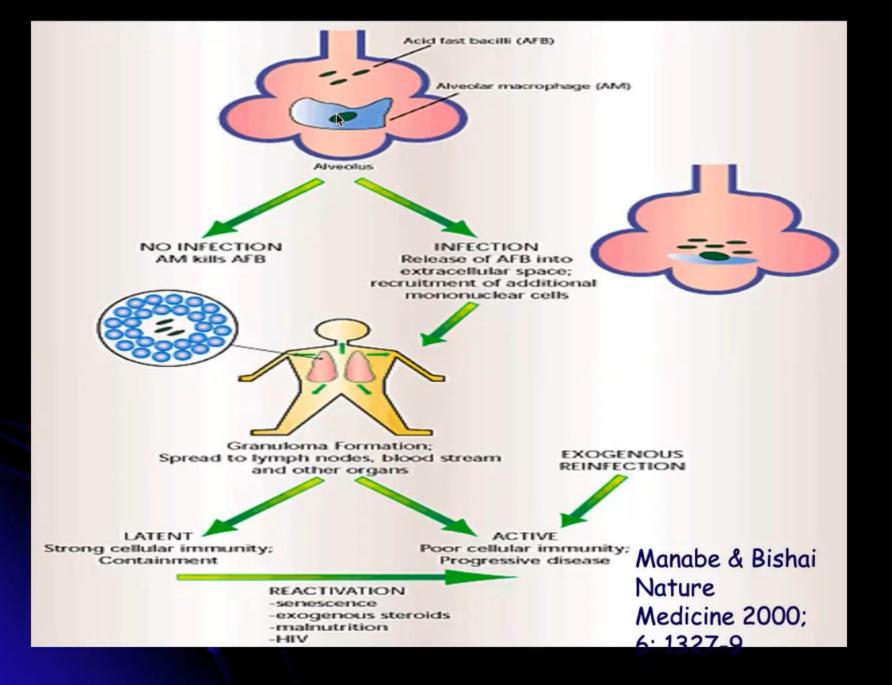


#### Clinical features

- insidious onset
- weight loss
- anorexia
- fever
- hepatosplenomegaly
- Headache almost always = meningitis
- abdominal pain and tenderness usually = peritonitis.
- Skin and eye tubercles (Tuberculous Uveitis)

PPD: - ve in 30%

CXR: may be characteristic.





#### **Bacteriologic confirmation**

Problematic in children

Pauci-bacillary disease, often poor yield

**NB! BUT** still do culture if possible

Which specimen to collect?

Gastric aspirate (fasting, early morning)

Induced sputum / Assisted sputum

Broncho-alveolar lavage

String test

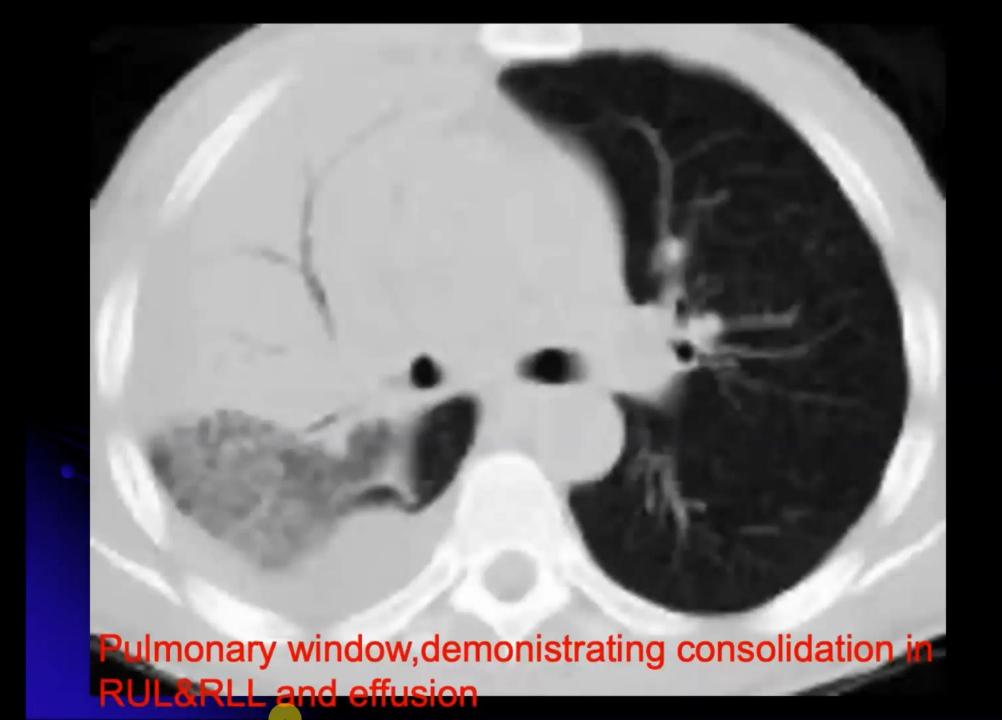
Fine needle aspiration or excision biopsy

**Not incision biopsy** 

## Timetable of disease after primary infection in children

- 3-8 weeks:
  - TST response
  - Hypersensitivity reactions
  - Erythema nodosum
- 1-3 months:
  - Hematogenous spread (meningitis and miliary in infants)
- 3-7 months:
  - Bronchial disease (< 5 years)</li>
  - Pleural effusions (>5 years)
- 1-3 years:
  - Osteo-articular disease
  - Calcifications
  - Adult-type disease
- > 3 years:
  - Reactivation





## Complications

## Most occur in the 1st year.

- Miliary TB & TB meningitis: not later than 3-6 mo. after initial infection.
- 2) Endobronchial TB: within 9 mo.
- 3) Bones or joints: within 1 y.
- 4) Renal: 5-25 y.
- 5) Secondary reactivation.



#### Suggested criteria for diagnosis of TB in children

#### Suspected/probable

#### Any three of the following:

- 1. History of contact with an adult with suspected or proven
- Symptoms and signs of TB such as persistent fever, cough, ,weight loss, failure to thrive, anorexia, respiratory distress, decreased breath sounds, rales on chest examination, lymphadenopathy, etc.
- 3. Positive Mantoux or PPD more than 10mm of indurations
- 4. Chest radiographic Findings such as an infiltrate or Lymphadenopathy

#### **Confirmed**

- •A positive AFB smear or culture of gastric aspirate or other body Fluid or
- Histological Findings consistent with TB

### **Treatment**

First line

: INH, rifampin, pyrazinamide ethambutol and streptomycin

Second line : para-aminosalicylic acid, ethionamide, caperomycin, kanamycin and cylcoserine.

### **Treatment**



- INH + rifampin X 9 mos. will cure 98%.
- Shorter courses (6 mos.) using more drugs; INH, rifampin and pyrazinamide for 2 mos. followed by 4 mos. of INH and rifampin.
- The 9 mos. approach is the one recommended for children.



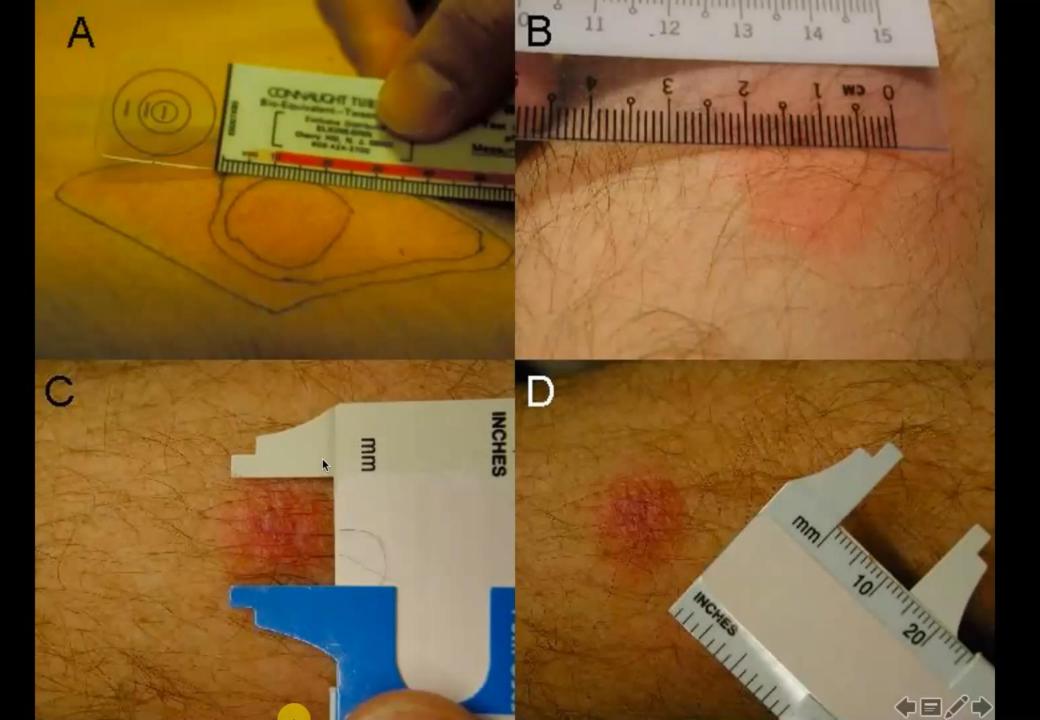
#### **Treatment**

#### **STEROIDS**

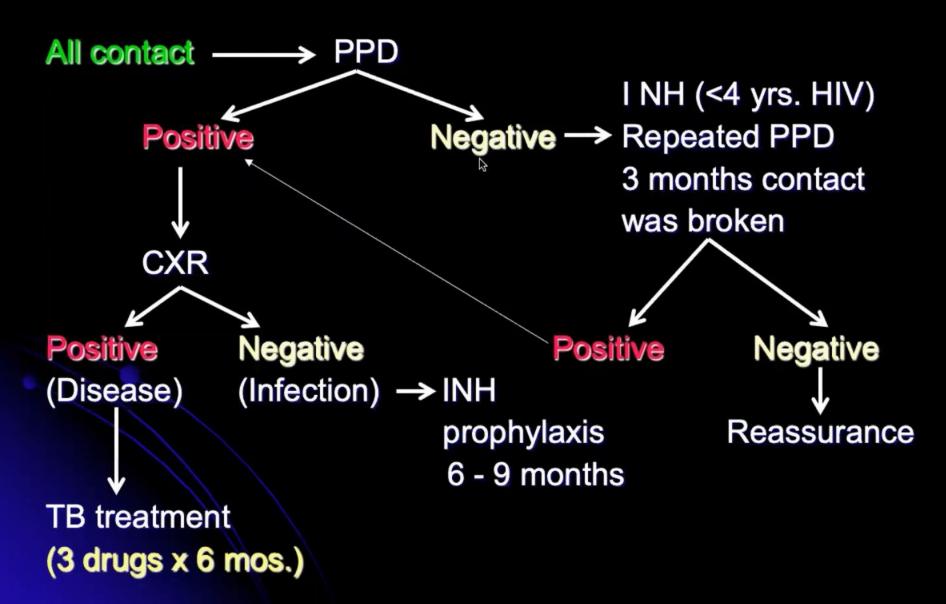
#### Use only with anti-TB med indicated in:

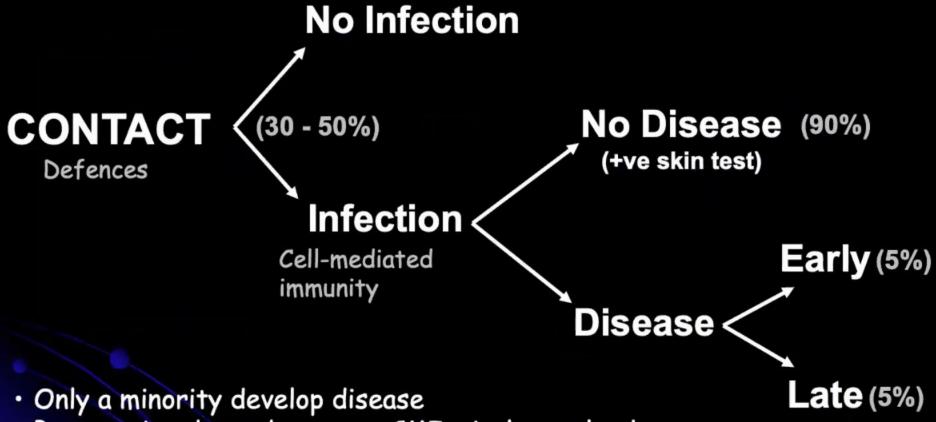
- 1) TB meningitis and increased ICP due to brainstem inflam and resultant HC.
- Endobronchial TB→ collapse or air trapping.
- Miliary TB with pericarditis, pleural effusion or peritonitis.





#### **Treatment of Contact**





- Progression depends on age, CMI, virulence, load
- If low prevalence area, new contact infrequent, early disease unlikely, most cases remote exposure
- Exogenous re-infection may occur esp. in immunocompromised or heavily exposed

#### Conclusion

- TB remains a permanent threat
- Importance of contact tracing and treatment adherence
- Future perspectives
  - Improved efficacy of TB vaccination
  - Improved diagnosis of TB infection (ELISPOT)
  - Shorter treatment for latent TB infection (and TB):
     new drugs

