

# *Pneumonia*



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- ❑ The clinical presentation of childhood pneumonia varies depending upon the **responsible pathogen**, the **particular host**, and the **severity**. The presenting signs and symptoms are nonspecific; **no single symptom or sign** is pathognomonic for pneumonia in children
- ❑ Symptoms and signs of pneumonia may be **subtle**, particularly in **infants and young children**
- ❑ **Neonates and young infants** may present with **difficulty feeding**, **restlessness**, or **fussiness** rather than with cough and/or abnormal breath sounds
- ❑ **Older children** and adolescents may complain of **pleuritic chest pain** (pain with respiration), but this is an inconsistent finding. Occasionally, the predominant manifestation may be **abdominal pain** (because of referred pain from the lower lobes) or **nuchal rigidity** (because of referred pain from the upper lobes)

- Fever
- Cough
- Tachypnea
- Auscultation

## **Tachypnea: Good Predictor of PNEUMONIA**

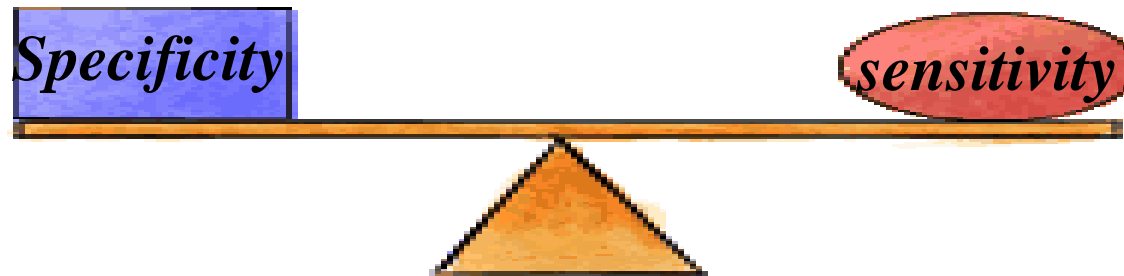
- ✓ Earliest sign
- ✓ Simplicity
- ✓ Reliability
- ✓ Ease in training

- The World Health Organization uses tachypnea (>60 breaths/min in infants 50 breaths/min in infants 2 to 12 months; >40 breaths/min in children 1 to 5 years; and >20 breaths/min in children  $\geq 5$  years) as the sole criterion to define pneumonia in children with cough or difficulty breathing
- In developed countries with a lower prevalence of pneumonia, multiple respiratory signs (eg, hypoxia, grunting, nasal flaring, retractions) are necessary to increase the certainty of pneumonia

## □ Reasonable Sensitivity & Specificity (around 80%)

Example: If in 100 cases of pneumonia, 80 cases were identified by “Fast Breathing”, then this would mean that:

- The sign has a mean sensitivity of **80 %**.
- 20 % would be missed “ *false negative* ”



- Low Sensitivity is a more serious problem than low specificity

- Use a timing device
- Count respirations for one full minute
- Best to count rate when the child is alert & quite.
- Fever can affect respiratory rate, but do not wait for fever to subside

- Auscultation:

- ✓ Babies and young children often cry during the physical examination
- ✓ Copious upper airway secretions. This creates another potential problem of transmission of upper airway sounds
- ✓ A subtle finding, particularly one at the pulmonary bases, can be missed due to shallow & rapid breathing
- ✓ Not all children with pneumonia have crackles

## HOSPITALIZATION:

- infants younger than **three to six months** of age
- **Hypoxemia** (peripheral capillary oxygen saturation [SpO<sub>2</sub>] < 90%)
- **Dehydration**, or inability to maintain hydration orally; inability to feed in an infant
- **Moderate to severe respiratory distress**: Respiratory rate >70 breaths/minute for infants 50 breaths per minute for older children; retractions; nasal flaring; difficulty breathing; apnea; grunting
- **Toxic appearance**
- **Underlying conditions** that may predispose to a more serious course of pneumonia (eg, cardiopulmonary disease, genetic syndromes, neurocognitive disorders), may be worsened by pneumonia (eg, metabolic disorder) or may adversely affect response to treatment (eg, immunocompromised host)
- **Complications** (eg, effusion/empyema, necrotizing process, abscess)
- **Failure of outpatient therapy** (worsening or no response in 48 to 72 hours)

## Extrapulmonary Manifestations of M pneumoniae Infections

1. A mucocutaneous rash is present in 10% of children with M pneumoniae and is fairly nonspecific. Several other skin manifestations have been associated with M pneumoniae infection, including erythema multiforme (EM), Stevens- Johnson syndrome (SJS), and the new clinical entity M pneumoniae–induced rash and mucositis (MIRM)

MIRM is characterized by more than 2 sites of mucosal involvement; cutaneous involvement itself may or may not be present, distinguishing it from EM and SJS. If cutaneous manifestations are seen, lesions are typically targetoid or vesiculobullous. Patients with MIRM have a more benign disease course than that of EM or SJS and typically make a full recovery, with recurrence in less than 10% of patients





2. M pneumoniae infection can cause **anemia(hemolytic anemia)**. Patients with underlying hematologic diagnoses such as sickle cell anemia are particularly at risk.

**3. Central nervous system manifestations** can occur in up to 7% of patients hospitalized with M pneumoniae infections.

A wide spectrum of central nervous system disease, including encephalitis, transverse myelitis, and cerebellar ataxia, have been reported

4. Cardiac manifestations of M pneumoniae are rare, occurring in less than 10%, but cases of myocarditis, pericarditis, complete heart block, and hemopericardium have all been reported.