



### Always Evaluate:

- Cardiopulmonary S/Sx
- Hemoglobin Level
- Reticulocyte Count

# Foundations Frameworks

## Approach to the Sickle Cell Crisis

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### Vaso-Occlusive Crisis

- Caused by anemia, dehydration
- These patients typically have higher resting Hgb
- Management:
  - Fluids: PO vs D5 1/2 NS at maintenance rate vs NS bolus if hypotensive
  - Opioid
  - Acetaminophen
  - Avoid NSAIDs or O<sub>2</sub> (unless hypoxic)
- Rule out: osteomyelitis, avascular necrosis, gallstones, septic arthritis, acute chest syndrome
- Cardiopulmonary Symptoms → Evaluate for **Acute Chest Syndrome**
  - Leading cause of death
  - Diagnosis: Infiltrate on XR chest plus...
    - Hypoxia
    - Fever
    - Chest pain
    - Respiratory signs/symptoms
  - Management: antibiotics, fluid resuscitation as needed, exchange transfusion, hematology consult, consider ICU admission

### Anemic Crisis

- Presents with less pain
- Patients may have pulmonary HTN, leg ulcers, increased risk of sudden death
- Obtain a hemoglobin, reticulocyte count, perform rectal exam

### Causes:

- Transient Red Cell Aplasia
  - Low reticulocyte count
  - Management: isolation and call hematologist
- Splenic Sequestration
  - Typically seen pediatric patients
  - Large spleen, LUQ pain, rapid drop in hemoglobin, high reticulocyte count
  - Management: transfusion, volume resuscitation
- Hemolytic Crisis
  - Elevated LDH, elevated reticulocyte count, elevated LFTs/bilirubin

### Special Circumstances

- Hyphema, priapism, CVA (even in peds), gallstones, osteomyelitis, sepsis/fever