

Related Concepts

Alteration in bleeding
Acute Pain
Impaired gas exchange
Ineffective tissue perfusion

Related Exemplars/Diseases

Sickle Cell Disease
Sickle Cell Crisis

Reading/Resources - Clinical Judgment

Cause - inheritance of sickle hemoglobin genes - lack of healthy red blood cells to carry O₂
Sickle-shaped red blood cells become rigid & sticky affecting O₂ transport

Most common symptoms

- Anemia
- Acute pain
- Swelling of hands & feet
- Frequent infections

Class/Lab/Clinical - Clinical Judgment

Monitor for pain → prompt administration of analgesics
Monitor O₂ saturation
Monitor for changes in cardiovascular status/efficiency
Monitor changes in skin integrity related to decrease in tissue perfusion

Priority Assessments or Cues

- 1 Pain
- 2 SPO₂/oxygen level
- 3 Signs of infection

Priority Labs & Diagnostics

- 1 RBC's
- 2 H/H
- 3 Sickling tests

Priority Nursing Interventions

- 1 Prompt administration of pain meds
- 2 Monitor fluid volume - hydration
- 3 Pace activities (energy conservation)

Priority Medications

- 1 Analgesics
- 2 Antibiotics
- 3 Hydroxyurea

Priority Potential & Actual Complications

- 1 Sickle cell crisis
- 2 Thrombotic crisis leading to blockage of small blood vessels - stroke, vascular occlusion
- 3 pain
Splenic sequestration (most often seen in infants & toddlers)

Priority Collaborative Goals

- 1 Dietary teaching - Na restriction & need for protein
- 2 Pain Management
- 3 Genetic testing



NurseThink® Quick

SICKLE <u>S</u>trokes/swelling of hands & feet / spleen problems <u>I</u>nfections / Infarctions <u>C</u>risis (painful, sequestration, aplastic) chronic hemolysis, cardiac problems <u>K</u>idney disease <u>L</u>iver disease <u>E</u>rection (priapism), <u>E</u>ye problems - retinopathy		
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NEXT GEN LEARNING – NCLEX® TEST PLAN

Safe and Effective Care: Management of Care, Coordinated Care, Safety and Infection Control Monitor for progression of symptoms, monitor for signs of occlusive issues & thrombotic crisis - hypoxia, change in oxygenation, circulation to extremities Health Promotion and Maintenance Monitor for progression of symptoms that can lead to complications Teach patient & family re care & treatment of condition Psychosocial Integrity Assess level of coping with illness, provide emotional support to patient & family, encourage patient/family to ask questions re care/treatment, diet, activity & follow up care Physiological Integrity: Basic Care and Comfort, Pharmacological and Parenteral Therapies, Reduction of Risk Potential, and Physiological Adaptation Promote bedrest, hydration & exchange transfusions when in sickle cell crisis, monitoring of vital signs, O₂ saturation level, pain level, peripheral circulatory status.
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QUALITY AND SAFETY COMPETENCIES

Patient-Centered Care Respect of patient values & preferences, involve patient & family in care decisions, provide education, emotional support Teamwork and Collaboration Collaboration with geneticist, dietary & other system specific specialists or needed Evidence-Based Practice Pain management protocols, hydroxyurea (antiproliferative agent) Quality Improvement Prevent hospital-acquired infections & complications, reinforcement of teaching Safety Monitor patient for adverse blood transfusion reactions, medication reactions Informatics Record & trend vital signs & lab results in EHR.

Peer Review: _____ Faculty Review: _____

Grade Tracker

67	77	87	92											
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