



## General principles:

- Hydration
- Oxygenation
- Promt & adequate analgesia
- Early treatment of infection
- Identification and treatment of complications



- AIM: prevent dehydration - Start hyper-hydration at 150%
  - maintenance (can be given PO/NG/IV)
  - Strict input-output monitoring

### Oxygen

- AIM: sats >95%
- Administer high flow oxygen if sats< 95%

#### RED FLAGS

- · Chest pain
- · Sats <95%
- Toxygen requirement
- · Fever >38°C
- T work of breathing

## Analgesia

- Start Within 30 minutes of arrival
- · If severe give IV/IN morphine
- Consider PCA/NCA to control pain
- · Remember paracetamol + NSAIDs
- Don't forget antiemetics & laxatives

NUH guidelines: Sickle cell disease (acute management) NICE guideline: Sickle cell disease: managing acute painful episodes in hospital

# **Emed Sickle Cell Crisis**

**Monica K. Wattana, Molly Estes** 

#### **Emed Sickle Cell Crisis:**

Hematologic and Oncologic Emergencies, An Issue of Emergency Medicine Clinics of North America Monica K.

Wattana, Molly Estes, 2025-08-01 In this issue of Emergency Medicine Clinics guest editors Drs Monica K Wattana and Molly Estes bring their considerable expertise to the topic of Hematologic and Oncologic Emergencies Top experts discuss approach considerations for the oncologic patient leukopenia and febrile neutropenia tumor lysis hyperviscosity syndrome mechanical obstruction and more Contains 14 relevant practice oriented topics including tumor hemorrhage chemotherapy complications immunotherapy complications acute leukemia multiple myeloma and hypercalcemia sickle cell and more Provides in depth clinical reviews on hematologic and oncologic emergencies offering actionable insights for clinical practice Presents the latest information on this timely focused topic under the leadership of experienced editors in the field Authors synthesize and distill the latest research and practice guidelines to create clinically significant topic based reviews

Kelley and Firestein's Textbook of Rheumatology Gary S. Firestein, Ralph C. Budd, Sherine E Gabriel, Iain B McInnes, James R. O'Dell, 2016-06-21 Consult the definitive resource in rheumatology for an in depth understanding of scientific advances as they apply to clinical practice Masterfully edited by Drs Gary S Firestein Ralph C Budd Sherine E Gabriel Iain B McInnes and James R O Dell and authored by internationally renowned scientists and clinicians in the field Kelley and Firestein's Textbook of Rheumatology 10th Edition delivers the knowledge you need for accurate diagnoses and effective patient care From basic science immunology anatomy and physiology to diagnostic tests procedures and specific disease processes this state of the art reference provides a global authoritative perspective on the manifestations diagnosis

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paramedics and emergency medical technicians who are treating the patients FEATURES AND BENEFITS General suggestions for working with people with disabilities such as person first language wheelchair etiquette and interacting with service animals Suggestions for assessment and treatment are specific to EMS The disabilities discussed in this text include Autism spectrum disorders Causes and capacities of intellectual disabilities such as pain and Down syndrome Hearing vision and speech impairments Physical disabilities such as arthritis cerebral palsy and spinal cord injury Mental health disorders such as ADHD anxiety and schizophrenia Traumatic brain injury Alcohol use disorder Cystic fibrosis PANDAS PANS Rett Hematology/Oncology Emergencies, An Issue of syndrome Sickle cell disease Systemic lupus erythematosus **Emergency Medicine Clinics of North America** John C. Perkins, 2014-08-28 This issue of Emergency Medicine Clinics edited by Drs John Perkins and and Jonathan Davis focuses on emergencies related to Hematology Oncology and covers topics such as Oncologic Mechanical Emergencies Neutropenic Fever Oncologic Metabolic Emergencies Acute Leukemias Pediatric Oncologic Emergencies Chemotherapeutic Medications and their Emergent Complications Anemia Thrombotic Microangiopathies TTP HUS HELLP Congenital Bleeding Disorders Acquired Bleeding Disorders and Antithrombotic agents Sickle Cell Disease and more Gastrointestinal Emergencies, An Issue of Emergency Medicine Clinics Angela Mills, Anthony Dean, 2011-05-28 This issue of Emergency Medicine Clinics guest edited by Drs Angela Mills and Anthony Dean will include articles on the following topics Approach to acute abdominal pain Evaluation of abdominal pain in older adults Evaluation of abdominal pain in the pediatric population Imaging and laboratory testing in acute abdominal pain Esophageal and gastric emergencies and Anorectal emergencies and foreign bodies in the gastrointestinal tract Observation Medicine, An Issue of Emergency Medicine Clinics of North America R. Gentry Wilkerson, Christopher Baugh, 2017-08-01 This issue of Emergency Medicine Clinics focuses on Observation Medicine in the ED Editors R Gentry Wilkseron and Christopher Baugh have assembled an expert team of authors on topics such as History and Principles of Observation Medicine Observation Unit Design and Establishing an Observation Unit Care of the Patient with Chest Pain in an Observation Unit Care of Cardiovascular Conditions in an Observation Unit Care of Gastrointestinal Conditions in an Observation Unit Care of Metabolic Endocrine Conditions in an Observation Unit Care of Neurologic Conditions in an Observation Unit Care of Respiratory Conditions in an Observation Unit Care of Infectious Conditions in an Observation Unit Care of Traumatic Conditions in an Observation Unit Care of Extremes of Age in an Observation Unit Pediatric Emergency Medicine, An <u>Issue of Emergency Medicine Clinics of North America, E-Book Mimi Lu, Ilene Claudius, Christopher S. Amato, 2021-07-06</u> This issue of Emergency Medicine Clinics guest edited by Drs Mimi Lu Ilene Claudius and Chris Amato focuses on Pediatric Emergency Medicine This is one of four issues each year selected by the series consulting editor Dr Amal Mattu Articles in this issue include but are not limited to High risk medicolegal issues in PEM meningitis appendicitis fractures Respiratory Emergencies intubation non invasive trach emergencies supraglottic airways Neck Trauma cervical spine seatbelt signs

penetrating palate injuries Cardiac Emergencies Fever and Tox New Designer Drugs Pediatric Emergency Medicine, An <u>Issue of Emergency Medicine Clinics</u> Mimi Lu, Dale P. Woolridge, Ann Dietrich, 2013-08-28 This issue focuses on Pediatric Emergency Medicine in the topic areas of Seizure Pain and Sedation Trauma Cardiac Emergencies Shock Asthma Infant Fever Head Injuries and Concussions and more Thrombosis and Haemostasis, 2007 Iron Disorders, An Issue of Hematology/Oncology Clinics Matthew M. Heeney, 2014-08-28 This issue of Hematology Oncology Clinics guest edited by Drs Matthew Heeney and Alan Cohen is devoted to Iron Disorders Articles in this issue include Hereditary Hemochromatosis HFE and Non HFE Iron Refractory Iron Deficiency Anemia IRIDA Sideroblastic Anemia Anemia of Chronic Disease Inflammation Pathophysiology of Transfusional Iron Overload Transfusional Iron Overload and Iron Chelation Therapy Iron Overload and its Management in Non Transfusion Dependent Thalassemia Treatment of Iron Deficiency Anemia and Iron Overload Assessment Data Base Directory ,1991 Index Medicus, 2002 Vols for 1963 include as pt 2 of the Jan issue Sickle Cell Disease in Clinical Practice Jo Howard, Paul Telfer, 2015-02-12 Sickle Cell Medical subject headings Disease is the most common genetic disease world wide and in the UK It has marked geographical variation in its distribution in the UK with a concentration in London and other major conurbations Birmingham and Manchester In these areas specialist centres have become established offering expert up to date care for both inpatients and out patients with Sickle Cell Disease Although patient numbers are increasing outside these areas the expertise of health professionals can be patchy This book aims to provide a user friendly accessible resource for areas with smaller numbers of patients to allow them to provide equitable care with the larger well established centres Sickle Cell Disease can be associated with acute life threatening complications when clear easily available advice is needed and with chronic long term complications which may need liaison with other health professionals Clear treatment protocols for all the common complications of sickle cell disease are outlined here with summaries of key evidence and references Handling Sickle Cell Disease Yvette LaPierre, 2021-08-01 This book explores sickle cell disease and how people with the disease manage it It examines how sickle cell affects daily life work and school and it explains the latest treatments available Features include a glossary web resources source notes and an index Aligned to Common Core Standards and correlated to state standards Essential Library is an imprint of Abdo Publishing a division of ABDO Sickle Cell Disease Graham R. Serjeant, Beryl E. Serjeant, 2001 Sickle cell disease is one of the grand diseases in the terminology of Victorian pathology because its widespread and multitudinous manifestations may affect any organ system in the body It has served as a model of molecular disease being one of the first genetic disorders to be explained at the molecular level Since publication of the Second Edition however further advances have been made Much has been learnt about the causes of variability in the natural history of the disease and of the reasons why there are such marked variations in the patterns of morbidity and mortality As Director of The Jamaican Sickle Cell Unit Graham Serjeant and his colleagues have shown the disease to be extremely variable ranging from a severe clinical course

with death in early childhood to mildly affected patients living to the age of 80 years This extreme variability within a single genotype implies that other factors both genetic and environmental must influence expression of the disease This new edition contains many new references and incorporates the latest thinking on the biology of the disease and the best practice in its management *Management and Therapy of Sickle Cell Disease* Clarice D. Reid, Samuel Charache, Bertram H.

Lubin, 1997-02 Serves as a guide for the health care worker involved in the management of patients with sickle cell disease Represents a collective summary of experiences with therapeutic regimens rather than the by product of controlled clinical trials Referred to as the Bible or cookbook Covers child adolescent and adult health care maint patient care coord psychosocial mgmt newborn screening infection painful events lung stroke transfusion eye contraception and pregnancy prenatal diagnosis gallbladder leg ulcers bones and joints etc

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