



Sepsis Clinical Pathway Synopsis

Objective of Clinical Pathway

The objective of the Sepsis Clinical Pathway is to provide care standards for infants or children with suspected sepsis, sepsis, or septic shock. The clinical pathway provides guidance on recommended evaluations, treatments, and reassessments to minimize variation in care.

Background

Sepsis, a life-threatening condition involving organ dysfunction caused by an exaggerated immune response to infection, is estimated to occur at a rate of 158.7 cases per 100,000 children in the United States.¹ More than 75,000 of these cases develop severe sepsis, including septic shock, each year, in which one or more organ systems fail.^{1,2} For up to 20% of the children diagnosed with sepsis, particularly if requiring admittance to the intensive care unit, mortality is the outcome.^{1,2}

Mortality for children experiencing sepsis is often associated with refractory shock, multiple organ dysfunction syndrome, or a combination of both.^{1,3} Early recognition, initial resuscitation, and ongoing management are critical to sepsis survivorship.³ The Sepsis Clinical Pathway aims to provide an evidence-based framework that guides early recognition, initial resuscitation, ongoing management, stabilization, and resolution, all of which are critical to optimizing outcomes.

Target Users

- Physicians (Emergency Department, Critical Care Medicine, Urgent Care, Neonatology, Hospital Medicine, Infectious Diseases, Hematology/Oncology, Fellows, Resident Physicians, Community Physicians)
- Nurse Practitioners
- Nurses
- Pharmacy

Target Population

Inclusion Criteria

- Infant or child with suspected sepsis, sepsis, or septic shock

Considerations

- For febrile ($\geq 38^{\circ}\text{C}$) full-term infants aged 8 to 60 days, consider referring to the [Febrile Infant Clinical Pathway](#)

AGREE II

The Surviving Sepsis Campaign International Guidelines for the Management of Septic Shock and Sepsis-Associated Organ Dysfunction in Children provided guidance to the Sepsis Clinical Pathway Committee.³ See Table 1 for AGREE II. Table 1

AGREE II Summary for the Surviving Sepsis Campaign Guidelines³

Domain	Percent Agreement	Percent Justification [^]
Scope and purpose	100%	The aim of the guideline, the clinical questions posed, and the target populations were identified.

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Stakeholder involvement	99%	The guideline <u>was developed</u> by the appropriate stakeholders and represents the views of its intended users.
Rigor of development	85%	The process used to gather and synthesize the evidence, the methods to formulate the recommendations, and to update the guidelines <u>were</u> explicitly stated.
Clarity and presentation	93%	The guideline recommendations <u>are</u> clear, unambiguous, and easily identified; in addition, different management options are presented.
Applicability	61%	The guideline <u>did not fully</u> address implementation barriers and facilitators, utilization strategies, or resource costs associated with implementation.
Editorial independence	98%	The recommendations <u>were not</u> biased by competing interests.
Overall guideline assessment	89%	
See Practice Recommendations		

Note: Four EBP Scholars completed the AGREE II on this guideline.

^Percentage justification is an interpretation based on the Children's Mercy EBP Department standards.

Practice Recommendations

Please refer to the Surviving Sepsis Campaign International Guidelines for the Management of Septic Shock and Sepsis-Associated Organ Dysfunction in Children for full practice recommendations, evaluation, and treatment recommendations.³

Additional Questions Posed by the Sepsis Clinical Pathway Committee

No clinical questions were posed for this review.

Recommendation Specific for Children's Mercy

Children's Mercy adopted the majority of the practice recommendations made by the Surviving Sepsis Campaign International Guidelines for the Management of Septic Shock and Sepsis-Associated Organ Dysfunction in Children.³ Variations/Additions include:

- Antibiotic recommendations were made in consideration of the guidance provided for the empiric antimicrobial treatment by the Children's Hospital Association's Improving Pediatric Sepsis Outcomes (IPSO) Collaborative Infectious Disease Workgroup and Antimicrobial Recommendations Working Group.^{4,5,6}

Measures

- Utilization of the Sepsis Clinical Pathway
- Utilization of the Sepsis-associated order sets
- Time to fluids
- Time to antibiotics
- Sepsis Huddle completion for the patient with sepsis alert

Value Implications

The following improvements may increase value by reducing healthcare costs and non-monetary costs (e.g., missed school/work, loss of wages, stress) for patients and families and reducing costs and resource utilization for healthcare facilities.

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- Decreased risk of overdiagnosis
- Decreased risk of overtreatment (i.e., treatment for meningitis when treatment for urinary tract infection is more appropriate)
- Decreased frequency of admission
- Decreased inpatient length of stay
- Decreased unwarranted variation in care
- Decreased morbidity and mortality

Potential Organizational Barriers and Facilitators

Potential Barriers

- Identification and diagnosis of sepsis
- Variability of an acceptable level of risk among providers

Potential Facilitators

- Collaborative engagement across care continuum settings during Sepsis Clinical Pathway development
- High rate of use of Sepsis Clinical Pathway
- Standardized order sets for Emergency Department, Hospital Medicine, and Pediatric Intensive Care

Diversity/Equity/Inclusion

Our aim is to provide equitable care. These issues were discussed with the Sepsis Clinical Pathway Committee, reviewed in the literature, and discussed before making any practice recommendations.

Order Sets

- ED Sepsis EBP Pathway
- IP Sepsis EBP Pathway
- PICU Sepsis Pathway
- NICU Sepsis/NEC
 - The Sepsis Clinical Pathway does not address care provided in the ICN.

Education Materials

- The Sepsis Clinical Pathway has no associated educational materials

Clinical Pathway Preparation

This pathway was prepared by the EBP Department in collaboration with the Sepsis Clinical Pathway Committee, composed of content experts at Children's Mercy. If a conflict of interest is identified, the conflict will be disclosed next to the committee member's name.

Sepsis Clinical Pathway Committee Members and Representation

- Leslie Hueschen, MD | Hospital Sepsis Director & Emergency Department | Committee Chair
- Jay Rilinger, MD | Critical Care Medicine | Committee Member
- Josh Herigon, MD, MPH, MBI | Infectious Diseases | Committee Member
- Lauren Kirkpatrick, MD | Hospital Medicine | Committee Member
- Margaret Boyden, MD | Hematology/Oncology/BMT | Committee Member

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- Priya Tiwari, MD | Neonatology | Committee Member
- Grace Arends, MD | Pediatric Emergency Medicine | Committee Member
- Christopher Kaberline, MBA, RRT-NPS, CPHQ | Quality and Safety | Committee Member
- Jolene Palmer, MSN, RN, CPN | Quality and Safety | Committee Member

EBP Committee Members

- Todd Glenski, MD, MSHA, FASA | Anesthesiology, Evidence Based Practice
- Kelli Ott, OTD, OTR/L | Evidence Based Practice
- Kori Hess, PharmD | Evidence Based Practice

Clinical Pathway Development Funding

The development of this clinical pathway was underwritten by the following departments/divisions: Emergency Department, Critical Care Medicine, Infectious Diseases, Hospital Medicine, Hematology/Oncology/BMT, Neonatology, Pediatric Emergency Medicine, Quality and Safety, Evidence Based Practice.

Conflict of Interest

The contributors to the Sepsis Clinical Pathway have no conflicts of interest to disclose related to the subject matter or materials discussed in this care process.

Approval Process

- This product was reviewed and approved by the Sepsis Clinical Pathway Committee, Content Expert Departments/Divisions, and the EBP Department, and subsequently approved by the Medical Executive Committee.
- Products are reviewed and updated as necessary every 3 years within the EBP Department at Children's Mercy. Content expert teams are involved with every review and update.

Review Requested

Department/Unit	Date Obtained
Emergency Department	May 2023
Critical Care Medicine	May 2023
Infectious Diseases	May 2023
Hospital Medicine	May 2023
Hematology/Oncology/BMT	May 2023
Neonatology	May 2023
Pharmacy, Infectious Diseases	May 2023
Evidence Based Practice	May 2023

Version History

Date	Comments
April 2017	Version one - ED, Inpatient, PICU algorithms and power plans developed
June 2023	Version two - ED, Inpatient, PICU algorithms and power plans revised, Sepsis Synopsis developed
February 2024	Version three - ED, Inpatient, PICU algorithms revised to include LR and ceftriaxone incompatibility information

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March 2026	Version four – Clinical pathway algorithms and synopsis updated to include naming conventions of the associated Epic order sets
August 2026	Version five – ED, Inpatient algorithm revised to reflect age range changes when referencing the Febrile Infant Clinical Pathway
September 2026	Version six – Modified <i>Systemic Screening for Sepsis/Septic Shock</i> and <i>Sepsis Huddle</i> website pages to reflect updated proposed reference values for identifying age-based vital sign abnormalities, with separate values specified for Emergency Department and inpatient settings

Date for Next Review:

- June 2026

Implementation & Follow-Up

- Once approved, the clinical pathway was presented to the appropriate care teams and implemented. Care measurements will be assessed and shared with appropriate care teams to determine whether changes are needed.
- Order sets consistent with recommendations were created or updated for each care setting.
- Additional institution-wide announcements were made via email, hospital website, and relevant huddles.
- Metrics will be assessed and shared with appropriate care teams to determine if changes need to occur.

Disclaimer

When evidence is lacking or inconclusive, options in care are provided in the clinical pathway and the order set(s) and/or order panel(s) that accompany the clinical pathway.

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References

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2. Balamuth F, Weiss SL, Neuman MI, et al. Pediatric severe sepsis in U.S. children's hospitals. *Pediatr Crit Care Med*. 2014;15(9):798-805. doi: 10.1097/PCC.0000000000000225.
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