

SHOCK



Stabilization Essentials in Pediatrics (StEP) is an interdisciplinary two-day course with components of didactic lectures, high fidelity simulations and hands-on workshops, prepared and delivered by PICU faculty. The target audiences are MDs, RNs, and RTs who care for critically ill children over the short term, usually while they await transport. These practitioners may be part of different departments depending on local workflows (ie. ED4, Adult ICU or High Acuity Pediatric Units).

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LEARNING OBJECTIVES

 To review a case of a neonate presenting in shock
 To review the approach to initial stabilization and management of shock
 To describe differences in management according to

suspected etiology of shock



CASE 1



6-day old term, 4 kg female presents with 2-day history of poor feeding and 1-day history of lethargy and increased work of breathing

Heart Rate	194 bpm sinus
Blood Pressure	55/38 mmHg
Respiratory Rate	80 breaths/min
SpO2	91% in Room Air
Temperature	36 degrees Celsius

Examination: Lethargic, CRT 5 secs, peripheral and central pulses weak, mottled

Signs of shock



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Signs of Poor Organ perfusion:

- Brain altered level of consciousness
- Skin Mottled, cool, or warm, Flash or prolonged capillary refill time
 - Kidneys: decreased urine output (< 1 mL/kg/h)</p>



Shock in Children

- Hypotension is a LATE feature of shock in children
- Cardiac output in small infants is heart rate dependent (limited ability to increase their stroke volume)
 - Vascular access may be difficult



Causes of shock



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DISTRIBUTIVE

- Anaphylaxis
- Septic Shock

HYPOVOLEMIC

 Urine or gastrointestinal losses, inadequate oral intake

• Hemorrhage

CARDIOGENIC

- Congenital Heart Disease
- Arrhythmia
- Myocarditis

OBSTRUCTIVE

- Pericardial Tamponade
- Tension
 Pneumothorax

Initial Management

B

D





ABCDE approach independent of suspected cause

- Assess airway patency, suction secretions as needed
- Support breathing by providing supplemental FiO2 1.0 via pediatric non-rebreather mask
- Insert peripheral IV access x 2, consider IO after 5 minutes or 2 failed attempts at IV access
- Initiate fluid resuscitation 10-20 mL/kg normal saline over 10-20 minutes*
- Verify blood glucose and correct as needed if glucose <3.3mmol/L
- 5 mL/kg bolus D10W or D10NS IV
 - Expose the patient and identify any visible rashes, measure temperature

CASE 1



6-day old term, 4 kg female presents with 2-day history of poor feeding and

1-day history of lethargy and increased work of breathing

Heart Rate	194 bpm sinus
Blood Pressure	55/38 mmHg
Respiratory Rate	80 breaths/min
SpO2	91% in Room Air
Temperature	39.6 degrees Celsius

precipitous delivery in the context of +ve GBS status, not treated with antibiotics, + maternal fever intra-partum

Examination: Lethargic, CRT 5 secs, peripheral and central pulses weak, mottled

CHBC Provincial Pediatric Sepsis Screening Tool

Sepsis is a MEDICAL EMERGENCY; Early Recognition and Treatment is Imperative for Survival





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https://www.childhealthbc.ca/clinicianresources/pediatric-sepsis-0

Sepsis is a **MEDICAL EMERGENCY: Early Recognition and Treatment is Imperative for** Survival

- Immediately notify Most Responsible Practitoner
- Proceed to CHBC Provincial Pediatric Sepsis Clinical Care Algorithm
- Consult local Pediatrician on-call: or CHARLiE via Zoom or phone; or higher level of care center via PTN

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https://www.childhealthbc.ca/clinician-resources/pediatric-sepsis-0



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ARE THERE SIGNS OF SUSPECTED SEPSIS? (patient has 2 or more of the following in the context of infection)

- Looks sick or toxic (feels cold to touch, blotchy, blue or pale skin)
- Parental/caregiver concern
- Critical heart rate
- Temperature greater than 38° Celsius or less than 36° Celsius
- Altered mental state (drowsy, difficult to wake, irritable or confused)
- Tachypnea, increased work of breathing, cough, grunting, chest pain
- Decreased feeding
- Reduced urine output or other signs of dehydration
- Abdominal pain, distension, vomiting, diarrhea
- Joint pain or swelling, rash or other signs of skin infection

IS THE CHILD AT HIGH RISK OF COMPLICATIONS FROM SEPSIS?

- Age less than 3 months or born premature
- Immunocompromised

AND/OR

- Cardiac, respiratory, or neuromuscular disease
- Significant developmental delay
- Indwelling vascular access or medical device
- Recent surgery or hospitalization
- Recent inpatient episode of sepsis (within 6-12 weeks)
- Intravenous recreational drug use



Clinical Care Algorithm

center via PTN

 Consult local Pediatrician on-call; or CHARLiE via Zoom or phone; or higher level of care

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Child Health BC Pediatric Sepsis Clinical Care Algorithm

Sepsis is a MEDICAL EMERGENCY; Early Recognition and Treatment is Imperative for Survival





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https://childhealthbc.ca/sepsis/algorithm/ clinical_care

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CHILD HEALTH BC PROVINCIAL PEDIATRIC SEPSIS RECOGNITION AND MANAGEMENT GUIDELINE

For Emergency Departments and Urgent Care Settings

MAY 2023

LEAD BENEFACTOR

saveonfoods

childhealthbc.ca



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https://childhealthbc.ca/sepsis/guideline/ recognition management

Definitions



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INVESTIGATION OF SEPTIC SHOCK



First Line Investigations



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First Line Lab Investigations in Pediatric Sepsis		
Blood culture	Prioritize over other blood tests. It is recommended to obtain blood cultures before initiating antimicrobial therapy in situations where this does not substantially delay antimicrobial administration. ¹⁵	
	Culture sensitivity increases with blood volume. Recommended volume to collect is 1mL/kg max 40mL. ^{26,27}	
Blood gas	Base deficit more than 5.0 mEq/L marker of possible sepsis ¹⁶	
Complete blood count	WBC can be normal, high or low in early sepsis. ¹⁶ Platelet count less than 100,000 uL in sepsis or disseminated intravascular coagulation (DIC). ¹⁶	
Lactate	Do NOT attribute increased lactate to difficult venipuncture. Lactate more than 2.0 mmol/L marker of possible sepsis. ¹⁶ Repeat lactate (q2h) if greater than 2mmol/L. Lactate greater than 4 mmol/L requires urgent action. ¹⁶	
Electrolytes, glucose, urea and creatinine	Include sodium, potassium, chloride, ionized <i>or</i> total calcium, magnesium, phosphate, glucose, urea and creatinine. C Reactive Protein: low value does not exclude early sepsis. ¹⁶	
Urinalysis, urine culture and sensitivity	Obtain via in and out/indwelling catheter.	

Second Line Investigations



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Second Line Investigations in Pediatric Sepsis		
Blood group and screen	If indicated.	
Liver function tests	Total bilirubin, AST, ALT. Increased bilirubin or alanine aminotransferase (ALT) in sepsis. ¹⁶	
Coagulation studies	PT (INR), PTT, fibrinogen if clinical evidence of bleeding or Disseminated Intravascular Coagulation (DIC). Altered values in the context of sepsis with thrombocytopenia indicative of DIC. ¹⁶	
Lumbar puncture & cerebrospinal fluid investigations	Recommended for neonates with suspected sepsis as long as clinically stable. For older children, lumbar puncture may be considered if meningitis is suspected and the child is clinically stable (i.e. no signs of increased ICP, coagulopathy or hemodynamic compromise). ^{16,19} Can do WBC and PCR for meningitis diagnosis on CSF from delayed LP ¹⁶ • cell count with differential • gram stain and culture • glucose • protein level Nucleic acid testing (NAT) for HSV and other potential pathogens.	
Cultures & Swabs	If diarrhea present, stool for culture and virology if available. Nasopharyngeal Flocked Swab; Respiratory Nucleic Acid Testing (NAT) panel. Culture & Sensitivity of wound that appears infected.	
Radiology	Consider Chest x-ray for respiratory distress or signs on examination. Other imaging as directed by the focus of infection e.g. septic joint. ¹⁶	
Cardiac	Electrocardiogram 12 Lead. Echocardiogram.	

*Consider sending ammonia level in neonate with suspected sepsis



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MANAGEMENT OF SEPTIC SHOCK



Management of Septic Shock

Initial Resuscitation



Surviving Sepsis

Campaign•

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Algorithm for Children Systematic Screening for Sepsis in Children SEPTIC SEPSIS SHOCK SUSPECTED Within 1 Within 3 hour of initial hours of initial recognition of suspicion of Expedited septic shock sepsis diagnostic Shock evaluation develops **Diagnostic evaluation supports** sepsis-associated organ dysfunction 3 5 2 4 6 Obtain Collect Start empiric Measure Administer fluid Start vasoactive agents if shock **IV/IO** blood broad-spectrum lactate. bolus(es) if shock antibiotics. access. culture. is present.* persists.*

FLUID RESUSCITATION



Intravenous Fluid 10-20 mL/kg crystalloids over 5-20 minutes

Titrate to markers of cardiac output: improved mentation, heart rate, capillary refill time, urine output

Reassess closely after each bolus markers of cardiac output and signs of fluid overload

Up to 40-60ml/kg in the first hour*

EMPIRIC ANTIMICROBIAL THERAPY



Should be started as soon as possible, ideally within 1 hour of sepsis recognition

Obtain blood cultures before initiating antimicrobials, if does not delay their administration

Empiric broad-spectrum therapy to cover all likely pathogens Narrow empiric anti-microbials once the pathogen and sensitivities are identified

If no pathogen is identified, antimicrobials should be narrowed or stopped according to clinical presentation, site of infection, patient risk factors, clinical improvement

EMPIRIC ANTIMICROBIAL THERAPY

Antibiotic Recommendations*		
Age	Unknown Source CNS Infection <i>Not</i> Suspected	Suspected CNS Infection
Term neonate less than 29 days of life	All infants under 28 days presenting with suspected sepsis should be treated for suspected CNS infection until proven otherwise	ampicillin AND cefotaxime AND acyclovir
29 days of life to 17 years of age less 1 day	cefTRIAXone** PLUS OR MINUS vancomycin***	cefTRIAXone ** AND vancomycin*** PLUS OR MINUS acyclovir

*Consult infectious diseases for antibiotic recommendations for children with allergies or contraindications to recommended antimicrobial therapies

**If ceftriaxone unavailable substitute with cefotaxime

***Consult pharmacy for therapeutic drug monitoring recommendations should duration of vancomycin be greater than 48 hours (sooner in patients with renal impairment)



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VASOACTIVE MEDICATIONS



Consider after 40-60mL/kg IV fluids • Avoid excessive fluid resuscitation • Additional fluid resuscitation may be concurrent

1st line: Epinephrine or Norepinephrine
Should be based on individual patient physiology
May both be administered via PIV/IO*



VENTILATION IN SEPTIC SHOCK



Positive pressure ventilation may be indicated in patients with fluid- and catecholaminerefractory shock to **decrease metabolic demand and support cardiac output**

Lower doses of induction drugs should be used given risk of hypotension/cardiac arrest

Non-Invasive Ventilation may be appropriate in patients responding to initial therapy who have respiratory distress due to fluid overload, evolving ARDS

ADDITIONAL THERAPIES



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STEROIDS

✓ If known corticosteroid deficiency
 ✓ Consider in fluid and catecholamine refractory shock

GLUCOSE CONTROL

Tight glycemic control is NOT recommended
 Lack of benefit and risk of hypoglycemia





CALCIUM

✓ Measure and replace calcium as needed

INTRAVENOUS IMMUNOGLOBULIN (IVIg)

✓IVIG may be of benefit in Toxic Shock Syndrome and suspected MIS-C (not routinely indicated in sepsis)

Resuscitation Targets?



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"Normalize" global perfusion HR to normal range for age BP to low normal range (5th to 50th percentile) Warm extremities Cap refill ≤ 2-3 seconds



"Normalize" end organ perfusion Urine output >1mL/kg/hr Normal mental status Lactate < 2

CASE 2



6-day old term, 4 kg female presents with 2-day history of poor feeding and

1-day history of lethargy and increased work of breathing

Heart Rate	194 bpm sinus
Blood Pressure	55/38 mmHg
Respiratory Rate	80 breaths/min
SpO2	91% in Room Air
Temperature	36 degrees Celsius

Physical Exam: CRT 5 seconds, peripheral and central pulses weak, mottled, Lower limb blood pressure not measurable, femoral pulses not palpable

CXR: Cardiomegaly





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Management of

Suspected Cardiogenic Shock



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Prostaglandin E1: 0.05 to 0.1 mcg/kg/min* Epinephrine infusion initiated at: 0.05 mcg/kg/min

Fluid resuscitation: 5-10 mL/kg over 20 minutes

Positive pressure ventilation (PPV) to minimize metabolic demand and support cardiac output Non-invasive PPV while preparing for intubation

Reduced doses of induction medications given potential for cardiac arrest e.g., Ketamine 0.5mg/kg/dose, Rocuronium 1mg/kg/dose

*Consultation with Pediatric Cardiology and Intensive Care

CASE 3



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6-day old female presents with 2-day history of poor feeding,

profuse watery diarrhea since birth, and 1-day history of lethargy

Heart Rate	194 bpm sinus
Blood Pressure	55/38 mmHg
Respiratory Rate	80 breaths/min
SpO2	91% in Room Air
Temperature	36 degrees Celsius

Physical Exam: CRT 5 seconds, peripheral and central pulses weak, mottled with cool extremities Decreased skin turgor

Management of Suspected Hypovolemic Shock



In the presence of a clinical history suggestive of <u>hypovolemia</u>, fluid resuscitation should be undertaken rapidly and titrated to clinical response (improved heart rate, urine output, perfusion, and mentation)

Glucose should be checked and corrected

Electrolytes (e.g., K+) should be measured and corrected as needed

References



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