

PEDIATRIC POST CARDIAC ARREST

ALGORITHM

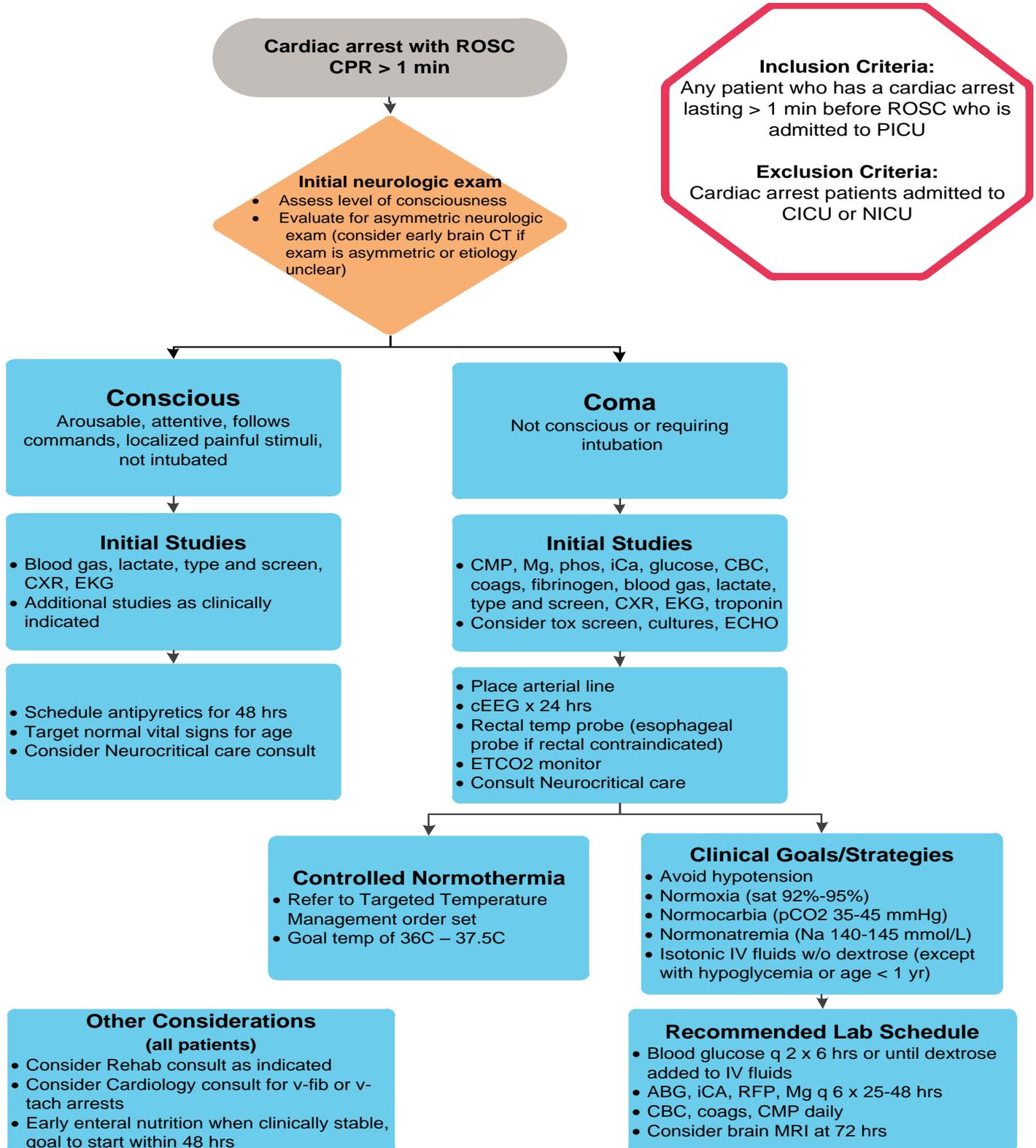


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TARGET POPULATION

Inclusion Criteria

- Sustaining cardiac arrest who regains spontaneous circulation after CPR lasting greater than or equal to 1 minute, OR
- Who undergo eCPR (CPR with cannulation to ECMO), AND
- Admitted to the Pediatric Intensive Care Unit (PICU)

Exclusion Criteria

- Admitted to the CICU or NICU

DEFINITIONS

- Cardiac Arrest: The cessation of cardiac mechanical activity as confirmed by the absence of signs of circulation
- Respiratory Arrest: The cessation of spontaneous respiratory effort such that there is ineffective ventilation and oxygenation
- Return of Spontaneous Circulation (ROSC): The restoration of a spontaneous perfusing rhythm that results in more than spontaneous gasp, fleeting palpable pulse or arterial waveform
- Conscious: arousable, attentive, follows commands, localizes painful stimuli, and not intubated
- Coma: not conscious and/or requiring intubation

INITIAL EVALUATION

Initial Evaluation in the PICU

- Comprehensive physical exam including comprehensive neurologic exam
- Specific attention to certain aspects of the neurologic exam:
 - Assess cranial nerves, Glasgow Coma Scale, level of consciousness (careful consideration given to pre-arrest neurologic baseline, as patients who have had short in-hospital cardiac arrest with no change in neurologic status from baseline proceed down the algorithm differently than those with change in neurologic functioning)

- For patients who are conscious with minimal new neurologic impairment (arousable, attentive, follow commands, localize painful stimuli, not intubated)
 - Arterial or Venous Blood Gas
 - Lactate
 - Type and Screen
 - Chest radiograph
 - EKG
 - Other labs to be obtained at the discretion of the attending based on clinical scenario: CMP, Magnesium, Phosphorus, ionized Calcium, CBC, Coagulation panel, fibrinogen, troponin
- For patients who are comatose (unconscious and/or requiring intubation)
 - Complete Metabolic Panel
 - Magnesium
 - Phosphorus
 - Ionized calcium
 - Complete Blood Count
 - Coagulation Panel
 - Fibrinogen
 - Lactate
 - Troponin
 - Arterial Blood Gas
 - Type and Screen
 - Chest radiograph
 - EKG
- Additional studies to be obtained for selected patients at the discretion of the attending physician:
 - Echocardiogram
 - Urine toxicology screen
 - Co-oximetry
 - Blood culture
 - Urine cultures
 - Cortisol
 - Non-contrast head CT to evaluate for acute pathology

CLINICAL MANAGEMENT

For patients who are conscious with minimal new neurologic impairment (arousable, attentive, follow commands, localize painful stimuli, not intubated)

Monitoring

- Routine PICU monitoring

Clinical Goals and Strategies

- Avoidance of fever: schedule antipyretics for 48 hours
- Target normal vital signs for age
- Consider Neurocritical care team consultation if abnormal neurologic findings or change from baseline exam

For patients who are comatose (unconscious and/or requiring intubation)

Monitoring

- Routine PICU monitoring
- Arterial catheter
- Continuous EEG for 24 hours
 - Prolonged monitoring may be indicated based on clinical scenario
- Rectal temperature probe, esophageal probe if rectal contraindicated (refer to normothermia order set)
- Continuous end-tidal CO₂ monitor while intubated

Clinical Goals and Strategies

- Controlled normothermia: use Targeted Temperature Management Order Set
- Normotension: hypotension is associated with worsened outcomes following pediatric cardiac arrest
- Normoxia (sat 92%-95%)
- Normocarbia (pCO₂ 35-45 mmHg; can target normal pH if the patient has evidence of chronic CO₂ retention)
- Normoglycemia (80-180 mg/dL)
- Normal serum sodium (140-145 mmol/L)
- IV fluids: use isotonic fluids without dextrose initially (recommend dextrose containing fluids in cases of documented hypoglycemia or age < 1 year)
 - Target euvolemia/even fluid balance once hemodynamically stable (defined as no fluid boluses and/or no escalation of vasoactive medications for 6 hours)
 - Add dextrose to IV fluids at 24 hours post-resuscitation or if serum glucose falls below 80 mg/dl
- Early enteral nutrition: recommend placing nasogastric tube when the patient is clinically stable with a goal of initiating enteral feeding within 48 hours of admission
- Recommended lab schedule:
 - Arterial Blood Gas, iCa, Renal Function Panel, Magnesium every 6 hours for 24-48 hours
 - CBC, coags, CMP daily
- Consult Neurocritical care team at admission

Other Considerations (all patients)

- For patients with documented or suspected ventricular fibrillation or ventricular tachycardia as initial arrest rhythm, consider Cardiology consultation to rule out arrhythmia syndrome
- Rehabilitation Medicine: consider consultation for assistance with tone, prognostication and/or transition of care out of the PICU

IMAGING

- Consider brain MRI as an aid for prognostication if clinical neurologic recovery is concerning
- If MRI brain is desired, do not obtain earlier than 72 hours after admission (wait an additional 48-72 hours if hypothermic arrest and/or if patient has undergone therapeutic hypothermia)

REFERENCES

1. Kleinman, ME, Chameides, L, Schexnayder, SM, et al. Part 14: pediatric advanced life support: 2010 American Heart Association Guidelines for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. *Circulation*. 2010; 122: S876-908.
2. Trzeciak, S, Jones, AE, Kilgannon, JH, et al. Significance of arterial hypotension after resuscitation from cardiac arrest. *Critical Care Medicine*. 2009; 37: 2895-2903.
3. Neumar, et al. Post-cardiac arrest syndrome: epidemiology, pathophysiology, treatment and prognostication. *Circulation*. 2008; 118: 2452-2483.
4. Kilgannon JH et al. Association between arterial hyperoxia Following Resuscitation from Cardiac Arrest and In-hospital Mortality. *Journal of the American Medical Association*. 2010; 303(21): 2165-2171.
5. Kuisma M et al. Comparison of 30 and 100% inspired Oxygen Concentrations during early post-resuscitation period: a Randomised Controlled Pilot Study. *Resuscitation*. 2006; 69: 199-206.
6. Herman ST et al. Consensus Statement on Continuous EEG in Critically ill Adults and Children, Part 1: Indications. *Journal of Clinical Neurophysiology*. 2015; 32: 87-95.
7. Neumar RW et al. Post-cardiac Arrest Syndrome: Epidemiology, Pathophysiology, Treatment and Prognostication: A consensus statement from the International Liaison Committee on Resuscitation. *Circulation*. 2008; 118: 2452-2483.
8. Kim YJ et al. Neuroprotective effects of L-Carnitine against Oxygen-Glucose deprivation in rat Primary Cortical Neurons. *Korean Journal of Pediatrics*. 2012; 55(7): 238-248.
9. Rittenberger JC et al. Postcardiac arrest Management. *Emergency Medicine Clinics of North America*. 2015; 33: 691-712.
10. Calaway et al. Part 8: Post-Cardiac Arrest Care. 2015 American Heart Association Guidelines update for Cardiopulmonary Resuscitation and Emergency Cardiovascular Care. *Circulation*. 2015; 132: S465-S482.
11. Rittenberger JC et al. Emergency Neurologic Life Support: Resuscitation following cardiac arrest. *Neurocritical Care*; 2015: S119-128.
12. Elmer J, et al. The association between hyperoxia and patient outcome after cardiac arrest: analysis of a high resolution database. *Intensive Care Medicine*. 2015; 41: 49-57.
13. Eastwood et al. Conservative oxygen therapy in mechanically ventilated patients following cardiac arrest a retrospective nested cohort study. *Resuscitation*. 2015.
14. Lee et al. Factors influencing outcome in patients with cardiac arrest in the ICU. *Acta Anesthesiologica Scandinavica*. 2013; 57: 784-792.
15. Naples et al. Cranial CT in the resuscitated patients with cardiac arrest. *American Journal of Emergency Medicine*. 2009; 27: 63-67.
16. Cocchi et al. The Role of cranial CT in the immediate post cardiac arrest period. *International Journal of Emergency Medicine*. 2010.
17. Lopez-Herce et al. Post return of spontaneous circulation associated with mortality in pediatric in hospital cardiac arrest a prospective multicenter observational study. *Critical Care*. 2014; 18: 607.
18. Moler et al. Therapeutic Hypothermia after out-of-hospital cardiac arrest in children. *The New England Journal of Medicine*. 2015; 372; 2197-2206.
19. Nielsen et al. Targeted temperature management at 33C versus 36C after cardiac arrest. *The New England Journal of Medicine*. 2013; 369; 2197-2206.

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- Neurocritical Care Committee – October 11, 2016
- Pharmacy & Therapeutics Committee – November 18, 2016
- Clinical Care Guideline and Measures Review Committee – November 29, 2016
- Medication Safety Committee – not applicable
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Scheduled for full review on November 29, 2020

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