CrackCast Episode 10 – Pediatric Resuscitation

Episode Overview:

1) Describe the CPR technique for an infant
2) Describe the PALS cardiac arrest, bradycardia and tachycardia algorithms
3) Describe the PALS septic shock algorithm
4) List the dose and primary indication for core pediatric resuscitation medications
5) What are 8 risk factors for Sudden Infant Death Syndrome (SIDS)?
6) List 8 strategies for preventing SIDS
7) Describe the management of a SIDS death
8) Define Apparent Life Threatening Events (ALTE) and list 10 causes

Wisecracks

1) Epidemiology and pathophysiology of pediatric cardiac arrest
2) What does Rosen’s say about atropine and succinylcholine use in pediatric resuscitation?
3) What are the pertinent anatomic features of children as it relates to laryngoscopy and intubation?

1) Describe the CPR technique for an infant

Rosen’s advocates for the “encircling” hands technique in infant CPR, which according to animal models gives better hemodynamics than using the standard two finger technique.

Remember the high quality CPR mantra: "push hard, push fast at a rate of at least 100 bpm, minimize interruptions, allow full chest recoil, do not over-ventilate"

2) Describe the PALS cardiac arrest, bradycardia, and tachycardia algorithms

PALS Cardiopulmonary Arrest

- high quality CPR (at least 100bpm)
- attach patient to monitor, attempt IV access
- ventilations at a rate of 10-12 breaths per minute
- shock shockable rhythm at 2-4 J/kg
- epinephrine 0.01mg/kg every 3-5 minutes
- pulse/rhythm checks every 2 minutes
- think through reversible causes (Hs and Ts)

**Chest Compression to Ventilation Ratio**
- no ideal ratios are known; recommendations made based on educational theory and physiology
- less O2 is needed during arrest
- in adults the ratio is 30:2
- in kids (infants and children):
  - 1 rescuer CPR: 30:2
  - 2 rescuer CPR: 15:2
- neonates: 3:1
- once an advanced airway is in place the goal is 8-10 breaths per minute.

**Pediatric VF or VT**
- much less common than PEA or asystole
- common etiologies:
  - infectious cardiomyopathies, congenital heart disease, WPW, channelopathies, electrolyte abnormalities
  - another specific pediatric entity: "asphyxia-associated VF" in foreign body aspirations, or drowning cases
- outcomes after initial VF/VT rhythm are good (~25%)
- 10% decrease in survival for each minute that VF/VT is not defibrillated

**PALS Bradycardia**

Treat bradycardia with poor perfusion, altered mental status, hypotension or shock

1) supplemental O2 and assist respiration if needed
2) start CPR if heart rate less than 60 if **signs of hypotension** (kids can’t change their contractility as much as adults, thus are rate dependent)
3) give epinephrine 0.01mg/kg IV/IO Q3-5min
4) atropine 0.02mg/kg IV/IO
   - min dose 0.1 mg
   - max dose 0.5 mg
   - may repeat x 1
6) consider trans-cutaneous or trans-venous pacing

**PALS Tachycardia**

1) Are they stable or unstable?
2) QRS Wide or Narrow?
   2a) QRS > 90ms (2.25 small boxes) - possible VT
   2b) QRS < 90ms - probably SVT if history of abrupt rate change, absent/abnormal p-waves, non-variable HR, HR > 220 in infants or 180 in children
   2c) probably sinus tach if history of known cause, normal p-wave morphology, variable R-R, constant P-R interval, HR < 220 in infants, or 180 in children
3) **Unstable** = shock 0.5-1 J/kg, increase to 2 J/kg if ineffective
4) stable wide complex tachycardia - consider adenosine if regular rhythm and monomorphic (do not delay defibrillation though… if unsure use electricity)
   - adenosine 0.1 mg/kg rapid IV push, max 6mg,
   - second dose 0.2 mg/kg rapid IV push, max 12mg
   4b) amiodarone 5mg/kg IV/IO over 20-50 minutes OR procainamide 15mg/kg IV/IO over 30-60 minutes
5) narrow complex - probable SVT
   5a) consider vagal maneuvers - ice bath, carotid sinus massage, or any Valsalva (REVERT trial?)
   5b) adenosine as above or synchronized DCCV if no vascular access
6) If probable sinus tachycardia: search for and treat the cause NOT the HR
3) Describe the PALS septic shock algorithm

4) List the dose and primary indication for core pediatric resuscitation medications

a. Adenosine
   0.1 mg/kg – rapid IV push - for probable SVT, max first dose 6 mg
   0.2 mg/kg – second dose, max second dose 12 mg

b. Amiodarone
   5 mg/kg - for stable wide complex tachycardia; for persistent pulseless VF/VT

c. Atropine
   0.02 mg/kg - for persistent unstable bradycardia

d. Calcium
   CaCl - 20mg/kg - for hypocalcemia, hyperkalemia

e. Dextrose
   0.5-1g/kg IV/IO - for hypoglycemia (adults D50; kids generally D25)

f. Dobutamine
   2-20 mcg/kg/min - for cardiogenic shock

f. Dopamine
   2-20 mcg/kg/min - shock and hypotension

g. Epinephrine
   0.01 mg/kg - for persistent unstable bradycardia; or PEA, pulseless VF/VT
despite defibrillation, or shock
   0.01-1 mcg/kg/min - for shock

h. Furosemide
   1 mg/kg - for pulmonary edema

i. Lidocaine
   1 mg/kg - as a pre-treatment agent for intubation in increased ICP, infants

j. Magnesium sulfate
   25-50 mg/kg - for Torsades or refractory VF/VT

k. Naloxone
   0.01 - 0.1 mg/kg - for opiate overdose

l. Norepinephrine
   0.05 - 2 mcg/kg/min - for shock (septic particularly)

m. Procainamide
   15 mg/kg for stable wide complex tachycardia

n. Sodium bicarbonate
   1-2 mEq/kg - for profound metabolic acidosis or Na channel blocker overdose (e.g. TCA)

Two more medications that Rosen’s lists:

Alprostadil (PGE1) - Prostaglandin
   used for ductal dependent congenital heart disease
   infusion: start at 0.05 - 0.1 mcg/kg/min
5) What are 8 risk factors for Sudden Infant Death Syndrome (SIDS) and describe its epidemiology

Risk factors for SIDS:

1) between 1-12 months of age
2) low socioeconomic status
3) multiple gestation pregnancy
4) male
5) prone sleeping
6) exposure to second-hand smoke

Epidemiology:

- 3rd leading cause of death in infants
- most common cause of death in children < 1 yr
- can occur anytime before age 2
  - rare < 1 month or > 1 yr.
  - peak 2-4 months of age
  - no association between Apparent Life Threatening Events (ALTE) and SIDS
- In the ED:
  1. rarely benefit to CPR or resuscitation (baby is already dead)
  2. primary role is to support the family
  3. blood and urine samples but rarely helpful
- most SIDS due to genetic variants in ion-channels/proteins
- autopsy usually done
- psychological considerations:
  1. ER physician should be clear, pediatrician backup may be helpful
  2. parents feel intense guilt and burden
  3. many have increased rates of infertility, miscarriage, and divorce post SIDS
- parental support is key
- for more see the AAP guideline: “death of a child in the ED”

6) List 8 strategies for preventing SIDS

1) avoid smoking, alcohol, or illicit drugs in the home
2) regular prenatal care
3) supine sleep position  
4) empty crib, on a flat surface, and no sleep positioners (i.e. wedges)  
5) firm sleeping surface  
6) room-sharing but no bed-sharing  
7) avoid overheating  
8) pacifier use decreases risk

7) Describe the management of a SIDS death

See the AAP guideline: “Death of a child in the ED”:

http://pediatrics.aappublications.org/content/134/1/e313

Standard protocol includes an autopsy, death scene investigation, clinical and family history.

8) Define and list 10 causes of Apparent Life Threatening Emergencies (ALTE)

- ALTE = “a description of a clinical presentation”  
- not clearly defined  
  - very challenging for ER docs because the events range from minor to life threatening  
  - up to 50% of cases have no definitive diagnosis

10 Causes of ALTE

1. infection  
2. GERD  
3. congenital pulmonary or vascular malformations  
4. seizure disorders  
5. cardiac dysrhythmias  
6. metabolic issues (i.e. hypoglycemia)  
7. head injury / non-accidental trauma  
8. apnea NYD  
9. breath holding  
10. reactive airways

Etiology (more detailed):  
  - Infection  
    - sepsis
RSV / other resp. viruses - 9%
- febrile convulsion - 12%
- pertussis - 6%
- UTI
  - GERD - 18%
    - with or without obstructive apnea
  - congenital malformations
    - tracheomalacia, vascular rings, pulmonary slings
  - seizure disorder - 25%
  - cardiac dysrhythmias
  - congenital cardiac malformations - 2%
  - metabolic
    - hypoglycemia
  - child abuse/ non-accidental trauma (NAT)
    - NAT don’t ever forget!!!!
  - apnea - 9%
  - asthma / head injury / feeding difficulties / breath holding

- ALTE child may look totally normal at the time of ED presentation (50% of the time) but any non-hospital person who witnessed choking or cyanosis should be taken seriously
- ED evaluation depends on hx and physical exam
- investigations +/-:
  - CBC, glucose, lytes, urine/blood cultures
  - tox screen, ECG
  - +/-CT, CXR,
- children < 60 days should undergo thorough screening for infection with//without pediatrics assessment
  - EMRAP Claudioius and Keens suggests kids <1 month be admitted
- outcome:
  - variable depending on etiology
  - one retrospective review of 196 pts. showed no deaths: mostly asthma, GERD, and seizures
  - most well appearing infants can be discharged with close follow-up
- safe discharge criteria: according to Fu and Moon
  - brief, non-severe, and resolving episode
  - non progressive cause - GERD
  - no comorbidities
  - no previous ALTEs in 24 hrs
Wisecracks:

1) Epidemiology and pathophysiology of pediatric cardiac arrest

Epidemiology:
- rare compared with adults
- most common causes:
  - SIDS – 25%
  - traumatic – 20%
  - drowning – 15%
- outcome of pediatric cardiac arrest
  - in hospital – ROSC 50 -60%
    - survival to hospital discharge 9-27%
  - out of hospital – ROSC 18-50%
    - survival to hospital discharge 6-9%
- much of the survival depends on patient specific factors (reason for arrest) but also is affected by system factors (number of trained personnel on hand)

Pathophysiology

Three main pathways:
- **asphyxia** - # 1 cause
  - acute hypoxia/hypercarbia
- **ischemic**
  - insufficient myocardial blood flow
    - hypovolemic shock, sepsis, MI
- **arrythmogenic** <10%
  - VF/VT

Clinical Features

- “pulselessness, apnea, and unresponsiveness”
- new ILCOR and AHA guidelines advise against pulse checks in kids given the frequent occurrence of bradycardia and hypovolemia
  - ***start CPR if 2/3 features:*** (e.g. apnea and unresponsive)
  - C-A-B
  - no pulse check or rescue breaths for lay rescuers
    - pulse checks: only 85% sensitive ; only 64% specific (36% of the time CPR wasn’t needed)
    - based on ECMO studies
    - health care workers may empirically start CPR without a pulse check
Four phases of cardiac arrest:

○ 1) pre arrest
  ■ avoid the insult that caused asphyxia or ischemia
    ● gates around swimming pools, seat belts
    ● recognize and treat shock early

○ 2) no flow
  ■ untreated cardiac arrest:
    ● recognize ASAP

○ 3) low flow (aka CPR phase)
  ■ do CPR

○ 4) post-resuscitation: complex number of factors involved
  ■ 1. post arrest brain injury
    ● coma, seizures, myoclonus,
    ● spectrum: full function--> brain death
    ● probably best to keep temp <36 degrees (based on adult studies in post VF arrests) and prevent fever
  ■ 2. post arrest myocardial dysfunction
    ● consider dobutamine/milrinone/levosimendan
    ● epinephrine
    ● fluids
  ■ 3. systemic ischemia-reperfusion response
  ■ 4. unresolved pathologic process that caused the cardiac arrest
    ● treat the underlying cause if known

2) What does Rosen’s say about atropine and succinylcholine use in pediatric resuscitation?

**Atropine**

- ACEP recommendations:
  ○ “pre-treatment of children < 1 year and those undergoing intubation with succinylcholine
- Thought to:
  ○ minimize bradycardia during intubation
  ○ very little to no evidence supporting this recommendation

**Succinylcholine**

- FDA as of 1992
  - routine use of SC in children should be avoided except in special circumstances:
    - laryngospasm, full stomach, or when no IV access is available (so IM is needed)
• rocuronium is preferred
  ○ however many places still use SCh for RSI…
  ○ kids need a higher dose: 2 mg/kg

3) What are the pertinent anatomic features of children as it relates to laryngoscopy and intubation?

<table>
<thead>
<tr>
<th>Feature</th>
<th>Description</th>
<th>Implication</th>
</tr>
</thead>
<tbody>
<tr>
<td>size</td>
<td>smaller, narrow field of vision</td>
<td>difficult for visually impaired providers</td>
</tr>
<tr>
<td>adenoidal hypertrophy</td>
<td>large tonsils</td>
<td>risk for obstruction, bleeding, etc.</td>
</tr>
<tr>
<td>developing teeth</td>
<td>friable alveolar ridge in edentulous kids</td>
<td>risk for disruption, bleeding</td>
</tr>
<tr>
<td>primary teeth</td>
<td>weak in young children</td>
<td>easily avulsed or aspirated</td>
</tr>
<tr>
<td>tongue</td>
<td>large relative to oropharynx</td>
<td>obstructs</td>
</tr>
<tr>
<td>anterior, superior larynx</td>
<td>opening is higher at C3</td>
<td>more acute angle of the laryngeal opening = difficult to visualize</td>
</tr>
<tr>
<td>weak hyoepiglottic ligament</td>
<td>base of tongue connected to epiglottis loosely</td>
<td>blade in vallecula doesn't elevate the epiglottis as much</td>
</tr>
<tr>
<td>narrow, angled epiglottis</td>
<td>may be more difficult to mobilize</td>
<td>covers the tracheal opening,</td>
</tr>
<tr>
<td>narrowest point</td>
<td>at the cricoid cartilage instead of glottis</td>
<td>tough to pass tube into the cords</td>
</tr>
</tbody>
</table>