



100 CARDINAL PED PRESENTATIONS

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CASES 21-30



What is this lecture about?

- Pediatric – exclusive to or commonly seen in kids
- Not a Zebra (ie something I've either seen during my career or know has been seen in our PED)
- Not a horse either – ie something you may make it through training without seeing
- Emergency practitioner can make the diagnosis or at least suspect it
- Emergency practitioner *should* make the diagnosis or at least suspect it, and can make a difference by either getting diagnostic studies, appropriate consultations, and starting initial management or by not doing unnecessary work-up

Quick info in 3 slides

- Classic case – build on illness scripts to reinforce when you should suspect this entity
- What should make you suspect this diagnosis
- Basics of diagnosis and ED management
- You can look it up for more detail, but you can't look it up until you at least suspect it
- FOAM resources for additional readings

How to use this lecture

- After the initial case presentation, think about the differential diagnosis
- Helpful framework: SPIT
 - *What is the most Serious diagnosis?*
 - *What is the most Probable diagnosis?*
 - *What is the most Interesting diagnosis?*
 - *What is the most Treatable diagnosis (ie what diagnosis should the EP do something about ASAP)?*
- Write down what you think is the diagnosis – commit!
 - *At the end, see how many you got right*

4 month old with abnormal movements

- 4 month old ex-full term infant born without any complications
- Brought in by parents because has been having repetitive brief abnormal movements as shown in their video
 - *Movements occur in clusters of 10-30 at a time, several times a day*
 - *Occur mostly during awake time*
 - *Not yet rolling over*
- VS normal, weight 6.1kg, neuro exam unremarkable, not hypotonic



Video can be found at

<https://www.youtube.com/watch?v=aVoJtslvqOU>

4 month old brought in for poor feeding and constipation

- 4 month old ex-full term infant brought in by parents because has been taking longer to feed and is constipated
- Pre parents, baby's cry also sounds different, less loud
- No fever, vomiting, cyanosis, ill contacts
- VS temp 37.6, HR 130, RR 30, BP 74/38, O2 sat 97%. Baby is hypotonic and appears to have mild ptosis
- You observe the baby feeding from a bottle and the baby has a weak suck



5 month old with rash on face and in diaper area

- 5 month old boy, ex full term NSVD to 28yo G1P1, no complications
- Brought in for 1 month of face and diaper rash, not improving despite trying antifungal, barrier cream, mupirocin ointment
- Also having some diarrhea x 1 month, no fever
- Formula feeding 6-8 oz four times a day + baby food, weaned from breast milk 1 ½ months ago
- Temp 37.5, HR 140, RR 32, BP 76/34
- Rash consists of erythematous, well demarcated plaques on cheeks and buttocks



5 month old with fever, rash, diarrhea

- 5 month old with 6 days of fever and 3 days of rash and “diarrhea”
 - *“Diarrhea” is 6 loose stools per day instead of usual 2, no blood*
- Previously healthy, born NSVD, no complications, no medications
- Saw PMD on day 2 of fever, and was diagnosed with otitis media and prescribed amoxicillin
- PE: mildly toxic appearing, no nuchal rigidity, HEENT normal (including no otitis), lungs clear, heart RRR no murmur, abdomen benign, no lymphadenopathy, exam otherwise normal
- CBC WBC 14.5, 70% N, Hct 27, Platelets 350, ESR 60, ALT 80, AST 40, Urine wbc 15, gram stain negative, viral respiratory panel negative

6 month old inconsolable and blue

- Just came to LA from Mexico
- Child woke up from a nap and started crying, then turning blue, then bluer and bluer
- History of “sopla” and told will need “cirugia” of ”corazon”
- Patient is crying non-stop, hyperpneic, cyanotic with pulse ox 60% and not improving with blow by O2 by mask
- He is alert, well hydrated, and afebrile. Lung, cardiac, and abdominal exam appear normal (but are difficult to do because of the child’s crying), and you do not hear a murmur.

6 month old with hypoglycemic seizure

- 6 month old ex-full term infant noted to be having a seizure on parent's video monitor at 6am
- EMS called, blood glucose 20 in the field, 5cc/kg D10 given with resolution of seizure
- Parents had decided to let baby "cry it out" and baby slept through the night for first time; they are feeling guilty
 - *Especially because +FH of SIDS death in sibling at age 4 months*
- VS: temp 37.7, HR 130, RR 32, BP 76/34, O2 sat 100%
- Groggy (post-ictal?), anterior fontanelle soft and flat, heart no murmur, lungs clear, abdomen soft, hepatomegaly with liver down 4cm, no rash
- Labs: glucose now 100, anion gap acidosis with bicarb 14, urine no ketones, ammonia 90

7 month old with yellow skin

- 7 month old female, ex 37 wk born NSVD to 35yo G2P2 mom, no complications
- Brought in for gradual yellowing of skin noted over last ½-1 week
- No fever, vomiting, diarrhea, rash
- Was jaundiced in first week of life which gradually resolved without phototherapy
- Eating commercial baby food – fruits and vegetables, started solids around age 5 months
- VS normal, yellow skin but anicteric, remainder of exam normal, no hepatomegaly



8 month old with vomiting and lethargy

- 8 month old ex-26 week premie, has been home from NICU x 4 months
- 2 days of vomiting 3-4 x/day, non-bloody, non-bilious, and per parent seems lethargic
- Parent is brand new foster parent, has had baby for only 3 days, and is unable to give additional history on NICU problems or course
- VS temp 37.7, HR 140, RR 28, BP 75/40
- Baby is awake but quiet – per foster parent sleeps a lot, no nuchal rigidity, lungs clear, heart RRR, no murmur, abdomen benign, no rash, nonfocal neuro
- What do you check the head for?



<https://oliversmiracle.files.wordpress.com>

9 month old with altered mental status

- 9 month old ex-full term, brought in for 1 day of lethargy
- Patient was well until a few hours ago when he became unarousable
 - *It was not noted to be his nap time*
 - *Parents tried verbal and gentle physical stimulation*
 - *After about 10 minutes, he awoke again, although remained less active than usual*
 - *20 minutes later he had another episode of unresponsiveness*
- On exam, he is sleepy but not comatose, temp 37.6, HR 120, RR 28, BP 76/34, O2 sat 100%. Anterior fontanelle flat, lungs clear, heart RRR no murmur, perfusion adequate, abdomen soft, nondistended, nontender, no rash
- Labs unremarkable, head CT negative, CXR/KUB shown



<http://oftankonyv.reak.bme.hu/tiki-index.php?page=Pediatric+radiology>



10 month old with rash after fever

- 10 mo old had 3 days of high fever
- Immunizations up to date
- Seen in ED on day 2, UA negative, RSV/Flu negative, discharged home
- Fever resolved, then broke out in rash
 - *Blanching pink macules and papules on face, trunk, extremities*
- Has been well appearing and nontoxic throughout illness, feeding well



Did you write down what you thought the answers were? Answers on following slides

4 month old with abnormal movements

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- Brought in by parents because has been having repetitive brief abnormal movements as shown in their video
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Infantile spasms (West syndrome) – suspect it

- Presents at < 1 year old, peak between 3 and 7 months old
- Brief spasms occurring in clusters of 2-125 spasms at a rate of up to 13 per minute
 - *Flexion spasms (“salaam seizures”) that appear similar to bowing at the waist, flexion of extremities*
 - *Extension spasms with flexion of upper body and extension of legs*
- Typically occur when infant awake or just woke up
- Neurodevelopmental regression frequently

Infantile spasms (West syndrome) – diagnose and manage it

- Once suspect diagnosis clinically, refer to pediatric neurologist for confirmation
 - *Neurologist may work up as outpatient or admit to expedite work-up*
- EEG shows characteristic pattern known as hypsarrhythmia
- 50-70% will have an associated underlying etiology, eg CNS malformation, chromosomal abnormality, inborn error, TORCH infection
 - *MRI and genetic/metabolic testing are indicated (although not in ED)*
- Main treatment is ACTH, mechanism of action unknown
 - *Anti-convulsants other than Vigabatrin are not effective*
- 85% have poor neurodevelopmental outcome, even with treatment
- For more information: <https://www.youtube.com/watch?v=3Nx975d2J7U>

4 month old brought in for poor feeding and constipation

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- Pre parents, baby's **cry also sounds different**, less loud
- No fever, vomiting, cyanosis, ill contacts
- VS temp 37.6, HR 130, RR 30, BP 74/38, O2 sat 97%. Baby is **hypotonic** and appears to have mild **ptosis**
- You observe the baby feeding from a bottle and the baby has a **weak suck**



Infant botulism – suspect it

- From intestinal colonization of *Clostridium botulinum* with toxin release
 - *Associated with ingesting honey and corn syrup (never give to baby < 1 year old), home canned foods, soil contamination*
- Affects infants < 12 months old, peak 2-8 months (median 3-4 months)
 - *Infant bowel flora and decreased gastric acid predisposes*
- Commonly present with constipation, poor feeding
 - *Weakness, hypotonia, weak cry, decreased gag and suck reflexes, ptosis, decreased extraocular movements*
 - *Main complication is respiratory failure from muscle weakness*

Infant botulism – diagnose and manage it

- Must clinically suspect, consult neurology and ID, and admit to PICU (due to risk of respiratory failure)
- Diagnosis confirmed by stool sample for *C. botulinum* spores or botulinum toxin, but may take several days to result
- Presumptive diagnosis based on clinical presentation and EMG findings of short, low-amplitude motor potentials
- Treated empirically if highly suspect with Botulism immune globulin (BIG) obtained from California DHS www.infantbotulism.org
 - *Antibiotics not indicated and may worsen disease by increasing toxin release*
- May require respiratory support with mechanical ventilation until recovers
- For more information: <http://dontforgetthebubbles.com/infantile-botulism/> and <http://www.medbullets.com/step2-3-pediatrics/20607/infantile-botulism>

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<http://www.anatomybox.com/wp-content/uploads/2012/03/acrodermatitis-erteropathica.jpg>



<http://www.rheumatologynetwork.com>

Acrodermatitis enteropathica – suspect it, diagnose it, manage it

- Zinc deficiency – may be congenital or acquired
- Diaper dermatitis that is erythematous, well demarcated, and resistant to usual treatments for diaper rash
- Lesions also common on cheeks, periorificial and acral areas of body
- Alopecia and nail dystrophy common, appreciated mainly in older patients
- Congenital form presents a few weeks to a month after weaning from breast milk, or during breastfeeding if mom zinc deficient
- Diagnosis confirmed by serum zinc levels
- Treat with zinc supplementation, expect improvement within 5-10 days

Review: other common diaper rashes

<https://www.firstderm.com/diaper-rash/>



- **Irritant diaper dermatitis**
- Affects areas touching the diaper, folds often spared, can become erosive (Jacquet's)
- Treat with frequent diaper changes, thick layer barrier creams, hypoallergenic products (soap, wipes, diapers etc), mild corticosteroids topical BID if very inflamed

For more information:

<http://pedemmorsels.com/diaper-dermatitis/> and
<http://learnpediatrics.sites.olt.ubc.ca/files/2010/07/Diaper-rash.pdf>

<https://pedclerk.bsd.uchicago.edu/page/diaper-rash>



- **Candidal diaper dermatitis**
- Bright red, moist appearing, in skin folds, with satellite lesions
- Risk factors: recent antibiotics, check for oral thrush
- Treat with topical nystatin ointment or clotrimazole cream

PEARL

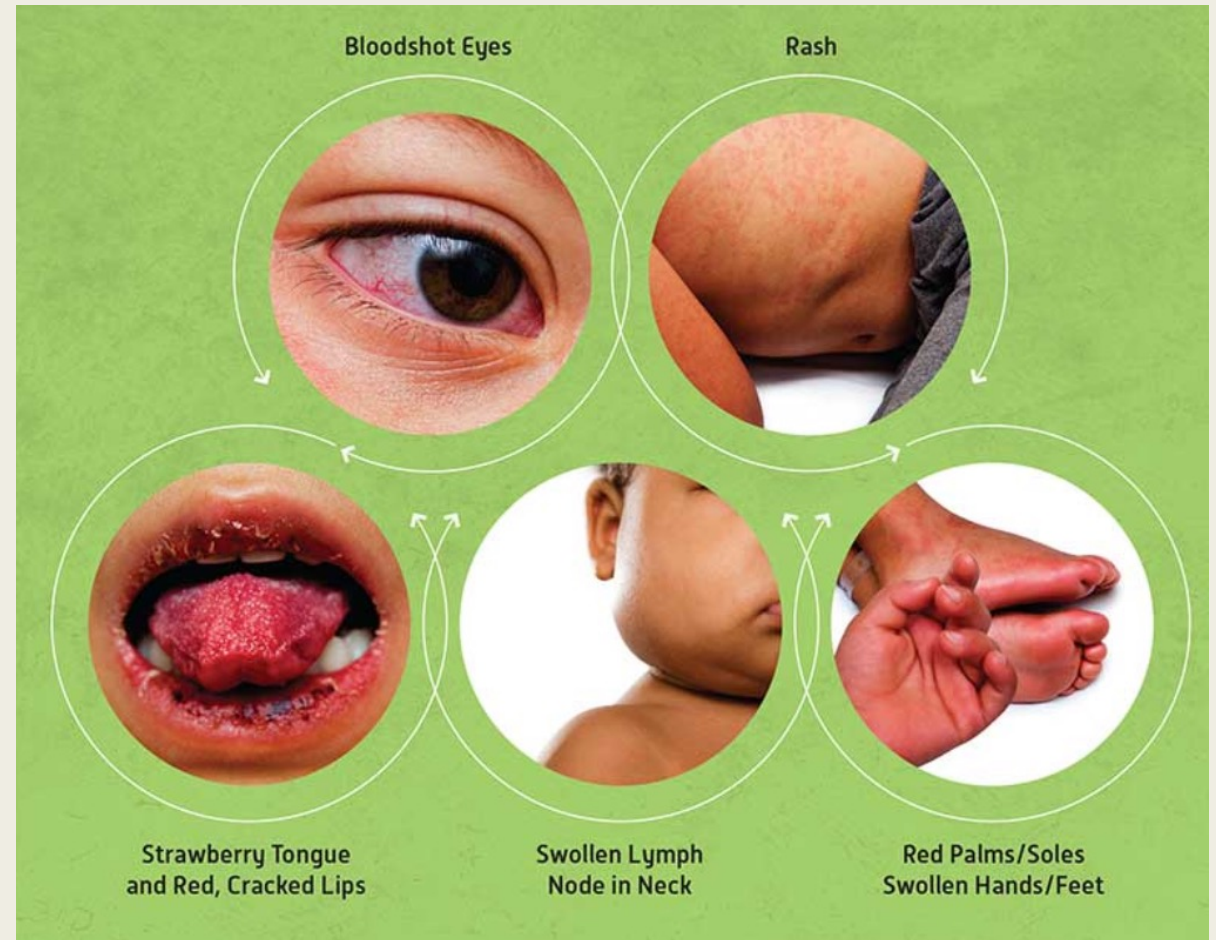
Do NOT use Mycolog or Lotrisone combined steroid + antifungal as the steroid component is too potent for baby skin

5 month old with fever, rash, diarrhea

- 5 month old with **6 days of fever** and 3 days of **rash** and “diarrhea”
 - *“Diarrhea” is 6 loose stools per day instead of usual 2, no blood*
- Previously healthy, born NSVD, no complications, no medications
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- CBC WBC 14.5, 70% N, **Hct 27**, Platelets 350, **ESR 60**, **ALT 80**, AST 40, **Urine wbc 15**, gram stain negative, viral respiratory panel negative

(Atypical / Incomplete) Kawasaki's – suspect it

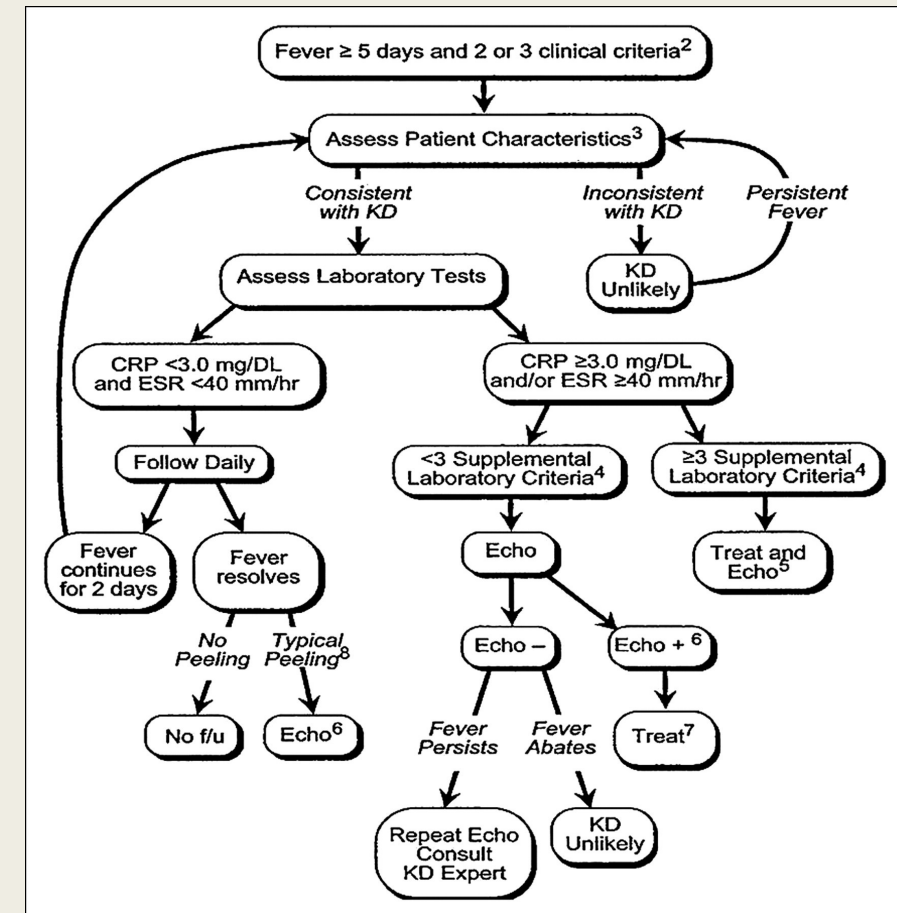
- Classic Kawasaki
 - 5 days of fever + 4/5 criteria
- Incomplete or Atypical Kawasaki
 - 5 days of fever + 2-3 criteria
 - < 6 months old and ≥ 7 days of fever, even if no other criteria
- Whenever you write "fever x 5 days" in your note, think Kawasaki's!



Kawasaki Disease Foundation

(Atypical / Incomplete) Kawasaki's – diagnose and manage it

- Send CRP or ESR, CBC, Albumin, ALT, UA
- Supplementary lab criteria
 - *Albumin ≤ 3 , anemia for age, ALT > 50 , Platelets > 450 after 7 days, WBC > 15 , pyuria > 10*
- Echocardiogram if CRP ≥ 3 or ESR ≥ 40
 - *Treat empirically if ≥ 3 lab criteria or echo positive*
- Admit, ID consult, high-dose aspirin, IVIG
- Algorithm <http://pemsources.org/wp-content/uploads/2016/06/Kawasaki.pdf>
- More info: <http://pedemmorsels.com/tag/kawasaki-disease/>



Fimbres AM, Shulman ST: Kawasaki Disease. Pediatrics in Review 2008;29(9):308-315

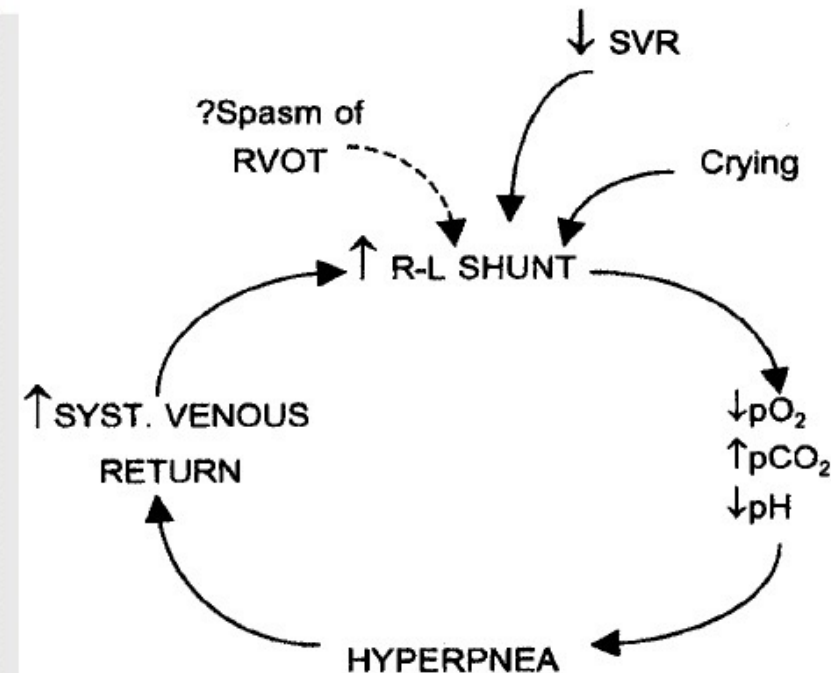
6 month old inconsolable and blue

- Just came to LA from Mexico
- Child woke up from a nap and started crying, then turning blue, then bluer and bluer
- History of “sopla” and told will need “cirugia” of ”**corazon**”
- Patient is crying non-stop, **hyperpneic**, cyanotic with pulse ox 60% and **not improving with blow by O2** by mask
- He is alert, well hydrated, and afebrile. Lung, cardiac, and abdominal exam appear normal (but are difficult to do because of the child’s crying), and you **do not hear a murmur**.

Hypercyanotic spell – suspect it

- Not just Tetralogy of Fallot
 - *Mixing lesions and pulmonary stenosis*
- Most common 6 wks – 6 mos
- Triggers: fever, defecation, feeding, awakening, dehydration, crying
- Inconsolable crying, worsening cyanosis, hyperpnea, quieter or no murmur, may have seizure

Pathophysiology of hypoxic spell



Hypercyanotic spell – diagnose and manage it

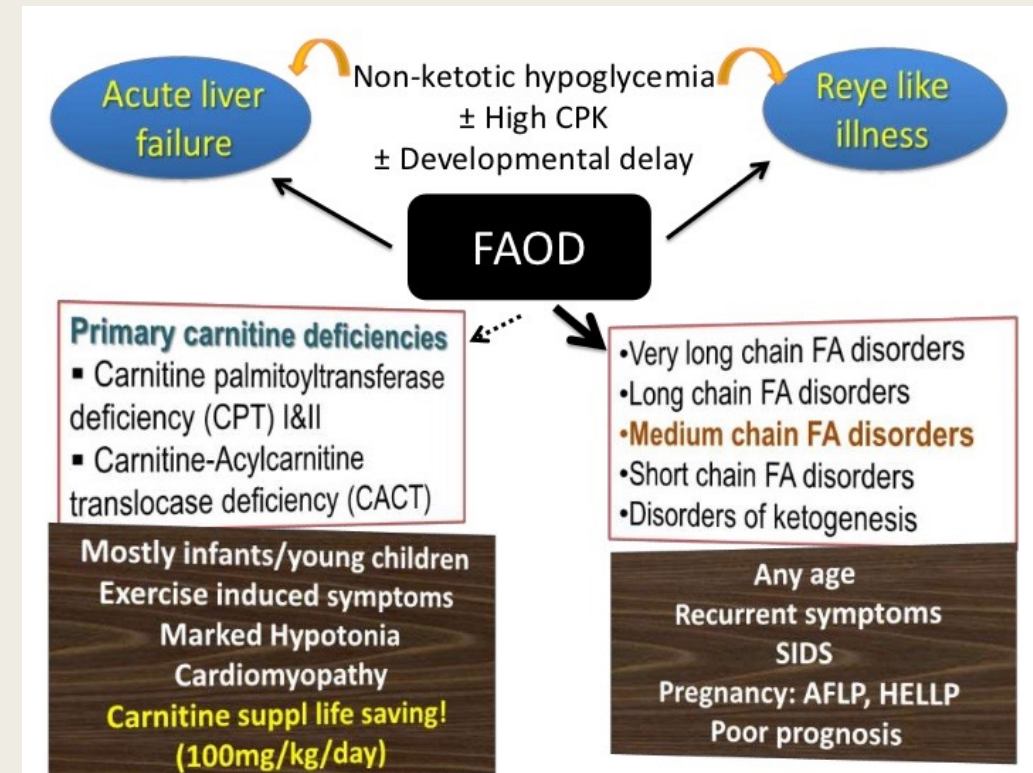
- Clinical diagnosis based on h/o congenital heart disease, classic presentation of cyanosis / low O₂ sat not improved by O₂, hyperpnea, somewhat abrupt onset of symptoms
- Knee-chest position (increase SVR), calm & comfort child, oxygen (decrease PVR), morphine (intranasal fentanyl if no IV), 10 cc/kg NS bolus (increase SVR), ketamine, propranolol or metoprolol or esmolol, phenylephrine, bicarb if significant metabolic acidosis
- Don't mess around with getting CXR, EKG, Echo if strongly suspicious
- Do consult pediatric cardiology
- For more info: <http://pedemmorsels.com/hypercyanotic-spells/> and for drug doses http://cardioiap.org/hyper_cyanotic_spell.aspx

6 month old with hypoglycemic seizure

- 6 month old ex-full term infant noted to be having a seizure on parent's video monitor at 6am
- EMS called, blood glucose 20 in the field, 5cc/kg D10 given with resolution of seizure
- Parents had decided to let baby "cry it out" and baby **slept through the night for first time**; they are feeling guilty
 - *Especially because **+FH of SIDS death** in sibling at age 4 months*
- VS: temp 37.7, HR 130, RR 32, BP 76/34, O2 sat 100%
- **Groggy** (post-ictal?), anterior fontanelle soft and flat, heart no murmur, lungs clear, abdomen soft, **hepatomegaly** with liver down 4cm, no rash
- Labs: glucose now 100, anion gap acidosis with bicarb 14, **urine no ketones**, ammonia 90

Fatty acid oxidation disorder – suspect it

- Typically presents in < 2 year old, but may present throughout life, even adulthood if mild
 - Often after period of fasting (as infant gives up night feeds)
 - Sometimes associated with stress eg viral illness leading to decreased po intake
 - May have had previous unexplained episodes
 - May have family history of SIDS, sudden death, seizures
- Vomiting, lethargy/coma, non-ketotic hypoglycemia, hepatomegaly, anion gap acidosis, hyperammonemia (but mild-mod, less than urea cycle disorder)



<https://www.slideshare.net/childrenliverindia/interpret-tests-for-metabolic-diseases-talk-sk-yachha>

Fatty acid oxidation disorder – diagnose and manage it

- Reverse hypoglycemia with D10W 2 cc/kg for neonates, 5 cc/kg for older, then start D10 1/2NS maintenance fluids
 - *Recheck bedside glucose often, may need 12-15 mg/kg/min glucose*
- Suspect inborn error? Order CBC, Chem panel (iStat if ill appearing or seizure), LFTs, VBG, ammonia, lactate, pyruvate, bedside glucose, urine/serum ketones
 - *Send urine organic acids, urine and serum amino acids, free and total carnitine levels, urine acylglycine specifically for suspected fatty acid oxidation disorder*
- Consult genetics specialist immediately
- Consider IV carnitine 25-100 mg/kg/day in 3 divided doses
- Avoid: salicylates, fat-binding/producing meds eg propofol, steroids, intralipids
- Admit for monitoring for recurrent hypoglycemia and confirmation of diagnosis
- Long-term will require low-fat, high-carbohydrate diet with frequent meals/snacks

7 month old with yellow skin

- 7 month old female, ex 37 wk born NSVD to 35yo G2P2 mom, no complications
- Brought in for **gradual yellowing of skin** noted over last ½-1 week
- No fever, vomiting, diarrhea, rash
- Was jaundiced in first week of life which gradually resolved without phototherapy
- **Eating commercial baby food** – fruits and vegetables, started solids around age 5 months
- VS normal, **yellow skin but anicteric**, remainder of exam normal, no hepatomegaly



Carotenemia – suspect it

- Yellow skin from eating carotenoids (esp beta-carotene), in yellow/orange fruit and vegetables such as: carrots, sweet potatoes, pumpkin mangoes, apricots, melons
- No icterus, normal bilirubin level, normal LFTs
- Commonly present in second half of first year of life, after introduction of solid foods and beginning to wean from milk diet
- Can also get secondary carotenemia at any age from eg hyperlipidemia (binds carotenoids preventing excretion), liver disease / diabetes / hypothyroidism (prevents conversion of carotenoids to retinol), nephrotic syndrome (prevents excretion of carotenoids)



Source of this and previous picture:
<http://elkhuntersjournal.com/photorfu/carrots-turn-skin-orange>

Carotenemia – diagnose and manage it

- Clinical diagnosis based on dietary history and physical exam
 - *Serum beta-carotene level elevated, but not routinely sent*
- Do not need to send serum bilirubin for confirmation if not icteric
 - *Mucous membranes will also not be yellow*
- Advise parents to decrease intake of high carotene foods and have follow up with primary physician for improvement
 - *Some green vegetables are also high in carotene*
- For more information: <http://www.dermnetnz.org/topics/carotenaemia/> and <https://www.ncbi.nlm.nih.gov/pmc/articles/PMC2280429/pdf/canfamphys00155-0083.pdf>

8 month old with vomiting and lethargy

- 8 month old **ex-26 week premie**, has been home from NICU x 4 months
- 2 days of **vomiting** 3-4 x/day, non-bloody, non-bilious, and per parent seems **lethargic**
- Parent is brand new foster parent, has had baby for only 3 days, and is unable to give additional history on NICU problems or course
- VS temp 37.7, HR 140, RR 28, BP 75/40
- Baby is awake but quiet – per foster parent sleeps a lot, no nuchal rigidity, lungs clear, heart RRR, no murmur, abdomen benign, no rash, nonfocal neuro
- What do you check the head for?
 - *Palpable ventriculoperitoneal shunt, bulging anterior fontanelle, measure the head circumference*



<https://oliversmiracle.files.wordpress.com>

Hydrocephalus and/or VP shunt malfunction – suspect it

- Very low birth weight premature infants often have intraventricular hemorrhage and get VP shunts placed
 - *Parents may not give the history – palpate the head to feel the shunt*
- This infant is showing “setting sun sign” = rim of white sclera seen above irises due to increased ICP affecting cranial nerves
- Other signs of increased ICP: bulging anterior fontanelle, increased head circumference, vomiting, lethargy, headache, papilledema
- VP shunt + fever = consider shunt infection, especially if shunt placed within last 6 months

Hydrocephalus and/or VP shunt malfunction – diagnose and manage it

- Noncontrast CT head to diagnose increased ventricular size or hydrocephalus increased ICP
 - *For shunt malfunction, helpful to compare to baseline CT or MRI*
 - *Newer rapid MRI protocols to decrease number of CTs patients with VP shunts get over lifetime*
- Plain radiograph “shunt series” to check for continuity of shunt
 - *Some studies show 4-8% of patients with malfunction have normal CT but abnormal shunt series*
- Acutely manage any critically high ICP, consult Neurosurgeon ASAP
- For more info: <http://dontforgetthebubbles.com/ventriculoperitoneal-shunts/>
- For critical increase in ICP: <https://first10em.com/2015/04/14/vp-shunt-malfunction/>

9 month old with altered mental status

- 9 month old ex-full term, brought in for 1 day of **lethargy**
- Patient was well until a few hours ago when he became unarousable
 - *It was not noted to be his nap time*
 - *Parents tried verbal and gentle physical stimulation*
 - *After about 10 minutes, he awoke again, although remained less active than usual*
 - **20 minutes later** *he had another episode of unresponsiveness*
- On exam, he is sleepy but not comatose, temp 37.6, HR 120, RR 28, BP 76/34, O2 sat 100%. Anterior fontanelle flat, lungs clear, heart RRR no murmur, perfusion adequate, abdomen soft, nondistended, nontender, no rash
- Labs unremarkable, head CT negative, CXR/KUB shown



<http://oftankonyv.reak.bme.hu/tiki-index.php?page=Pediatric+radiology>



<http://www.emed.ie/Paediatrics/Gastrointestinal/Intussusception.php>

Intussusception – suspect it

- Majority 6-36 months old, 60% < 1 year old, 80-90% < 2 years old
- Classic presentation: colicky episodes of abdominal pain (cry & draw up legs toward belly), often 15-20 min apart, non-bilious vomiting, heme+ stool
 - *Kids can appear very normal in between episodes, don't discharge them too fast!*
 - *Classic "currant jelly" stool is a late finding*
- Initial presentation can be lethargy alone, with no abdominal pain, vomiting, or bloody stool initially
- Consider intussusception in the differential diagnosis of altered mental status in infants and rule it out before getting a CT head (if no other signs/symptoms worrisome for head trauma)
- Plain films may have target or crescent sign, paucity of gas in colon



Intussusception – diagnose and manage it

- Ultrasound is 97.9% sensitive for the diagnosis, and is preferred due to lack of ionizing radiation exposure
 - *POCUS ED bedside ultrasound has been performed with sensitivity of 85-90% with training*
- Most intussusceptions are ileocolic, and nonoperative reduction with air, saline, or contrast enema by a radiologist is 80-95% successful
 - *Pediatric surgeon should be available in case of perforation or lack of success*
- Children outside the 6-36 month age should be evaluated for a “lead point,” (Meckel’s, polyp, lymphoma, parasite, fecalith, hemangioma, etc)
- For more info: <http://brownemblog.com/blog-1/2016/7/7/keep-your-eye-on-the-target-pocus-for-intussusception>
- PED EM Morsels has many great morsels on intussusception: <http://pedemmorsels.com/intussusception/> and <http://pedemmorsels.com/intussusception-altered-mental-status/> and <http://pedemmorsels.com/intussusception-discharge/>
- SAEM Bedside ultrasound video lecture <https://vimeo.com/37877537>



<http://radiology-information.blogspot.com/2013/02/ultrasound-images-of-intussusception.html>

10 month old with rash after fever

- 10 mo old had 3 days of high fever
- Immunizations up to date
- Seen in ED on day 2, UA negative, RSV/Flu negative, discharged home
- Fever resolved, then broke out in rash
 - *Blanching pink macules and papules on face, trunk, extremities*
- Has been well appearing and nontoxic throughout illness, feeding well



Viral exanthem – suspect it

- This patient's course classic for **roseola infantum**
- Typically occurs in 6mo to 2yo
- A few days of high fever; as fever comes down, rash breaks out
- Blanching pink macules and papules
- Well appearing and nontoxic throughout illness despite high fevers
- Original classification
- First disease: measles (rubeola)
- Second disease: scarlet fever
- Third disease: rubella
- Fourth disease: “Duke’s disease” – no longer a distinct entity
- Fifth disease: erythema infectiosum (parvovirus B19 – slapped cheeks appearance)
- Sixth disease: roseola infantum

Viral exanthem – diagnose and manage it

- Clinical diagnosis initially
 - *Specific viruses with characteristic findings, eg measles, fifth disease*
 - *Many other viruses beyond the original classification cause a nonspecific viral exanthem (eg enterovirus)*
- For disease of public health concern, eg measles, consider serology
- Supportive care with rest, antipyretics, attention to fluid intake
- For more information: http://www.medscape.com/viewarticle/734882_1 and <https://www.derm101.com/therapeutic/viral-exanthems-2/>