

NEUROLOGY REFERENCE CARD

WHO TO CALL FOR CONSULTS:

Patient service	Consultant
7S, 7N, 8S, 8E, 11S, BI NICU, BWH NICU, BWH Nursery	Neurology ICU resident
ED	Neurology ED resident
Floor (except 8E), ICP*	Neurology consult

*For daytime floor consults: if patient is followed by Epilepsy (see clinic notes), page Epilepsy Consult Fellow

Information to prepare for consults:

1. Acuity: Stroke STAT (call 52170)? Currently seizing? Impending herniation?
2. Consult question
3. Relevant Neurologic history
4. Seizure type/frequency (describe)
5. Current neuro meds (AEDs, tone meds, rescue meds). Calculate doses in mg/kg/d, times given.
6. Pertinent findings on YOUR neurologic exam (for headache, please do a fundoscopic exam)

MANDATORY CONSULTS:

1. Status epilepticus
2. Therapeutic hypothermia (in NICUs)
3. All ECMO patients
4. Cardiac arrest (most)

If a patient does not need a consult, but would benefit from urgent follow-up (<1-2 weeks), please have your Attending page the NOW Attending (Neurologist of the week).

IF PATIENT IS DUE FOR AN AED DOSE, PLEASE ADMINISTER ON TIME REGARDLESS OF WHETHER OR NOT OUR CONSULT IS DONE UNLESS OTHERWISE SPECIFIED. Consider trough levels.

THE NEUROLOGIC EXAM

Please try to do as much as possible. The more you practice, the better you'll get!

- MENTAL STATUS (describe interactions):
1. Awake, comfortable, fussy, distracted, somnolent, obtunded
 2. Oriented to person, place, day, month, year.
 3. Follows directions
 4. Maintains attention (months of the year or days of the week backwards)
 5. Fund of knowledge appropriate for age
 6. Memory (3 word recall at 1, 5 minutes)
 7. Language: Speaking fluently, coherent, paraphrased errors, neologisms, naming, repetition.

CRANIAL NERVES:

CN II: Visual acuity, visual fields, PERLA, fundoscopic examination (disc margins at least)

CN III, IV, VI: Fixing and following, smooth eye movements or nystagmus.

CN V: Facial sensation to light touch

CN VII: Facial movements (smile, grimace, cheek puff)

CN VIII: Do they hear finger rub bilaterally?

CN IX-XII: Swallow function, any changes in articulation or voice quality, palate elevation, tongue midline or deviated. Test strength of shoulder shrug, neck rotation.

MOTOR: Describe tone (axial and appendicular), especially in newborns. Strength testing can be tricky with kids <3, but try to push and pull extremities and see how much they reciprocate. Describe abnormal movements (speed, quality, stereotyped?, suppressible?)

REFLEXES: Check especially for clonus and any asymmetry in reflexes. DTRs should be checked at brachioradialis, biceps, triceps, patellae and achilles tendons. Toes up or down with plantar reflex?

SENSATION: Check light touch at least. If there is question of a sensory deficit, please also do temp/pinprick and vibration/proprioception.

CEREBELLAR/COORDINATION: Finger-nose-finger (or describe if little kids reach for toys smoothly). Finger tapping, rapid alternating movements. Any sway on Romberg?

GAIT: Test normal gait. Do heel, tip toe and tandem if possible.

HEADACHES

Type	Characteristics
Migraine	Throbbing, pulsating pain. Unilateral in 60-70%, but often bilateral in younger patients. Worse with exertion, better with hydration, rest, darkness.
Tension	Bilateral pressure that waxes and wanes.
High pressure headache (e.g.: idiopathic intracranial hypertension)	Variable HA. Gradual. Some constant, some throbbing, variable location, though often retrobulbar. Worse when supine, Valsalva.
Trigeminal autonomic cephalgia (e.g.: cluster HA)	Unilateral, around the eye or temple. Rapid onset (minutes), pain is continuous, excruciating.
Medication overuse headache	Characteristics vary, but usually preceded by another HA disorder.

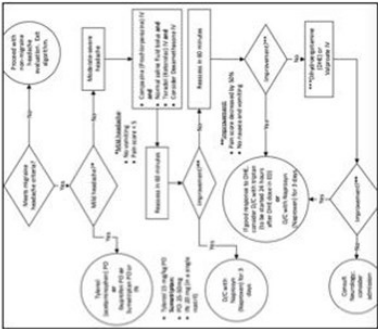
In the ED, see **Migraine EBG** (right):

Inclusion: Age 7+, low suspicion for other etiologies, HCG testing if of child-bearing age

HEADACHE RED FLAGS:

- acute onset
- atypical headache for patient
- neck stiffness
- worse when supine or with Valsalva
- waking from sleep
- vomiting w/o nausea or diarrhea
- focal neurologic symptoms
- altered mental status
- blurry/double vision

Associated symptoms	Risk factors
Associated with N/V, photo/phonophobia. Auras: usually visual, but can involve speech, sensory or motor deficits as well.	family history, female, R-L shunt
Associated with stress.	female, weight, drugs
Often associated vomiting. Transient visual obscurations in over half. Some with photopsia (flashes of light). Some with diplopia. Can have CN VI palsy. Associated with lacrimation (ipsilateral), injection, congestion, sweating. +sensitivity to alcohol.	obesity, prior history
Associated with use of opioids, NSAIDs, Tylenol, Fioricet for >=2 days/week x 3 mo.	



SEIZURES	STATUS EPILEPTICUS	CHECKLISTS FOR CONSULTS:	Seizures:																																											
<p>Seizures: Clinical manifestation of abnormal, excessive synchronous neuronal (cortical) discharges.</p> <p>Epilepsy: At least 2 unprovoked seizures occurring >24h apart</p> <p>Seizures are COMMON:</p> <ul style="list-style-type: none">• 3-5% of children <5yo have a febrile seizure• 1% of children <14yo have an afebrile seizure• 0.5-0.8% of children have epilepsy <p>Classification (ILAE 2017):</p> <p>Focal Onset (formerly "partial"): Originate in one hemisphere.</p> <ul style="list-style-type: none">• Can be Aware vs. Impaired Awareness• Can be Motor onset (automatisms, atonic, clonic, tonic, spasms; hyperkinetic, myoclonic) vs. Non-Motor onset (autonomic, behavior arrest, cognitive, emotional, sensory)• Can have focal to bilateral tonic-clonic (formerly "secondary generalization") <p>Generalized Onset: Bilaterally distributed origin.</p> <ul style="list-style-type: none">• Motor (tonic-clonic, clonic, tonic, myoclonic, atonic, spasms) vs. Non-Motor (absence) <p>Management (in general):</p> <p>Febrile seizures: no treatment, unless very recurrent, then consider benzo ppx with fever</p> <p>1st unprovoked seizure: no treatment, obtain outpatient routine EEG</p> <p>2nd unprovoked seizure: consider treatment, esp. if EEG abnormal</p> <p>Diazepam Dosing (consider script if prolonged seizure)</p> <table><tr><th>2-5 yr</th><th>6-11 yr</th><th>12+ yr</th></tr><tr><td>Weight (mg)</td><td>Weight (mg)</td><td>Weight (mg)</td></tr><tr><td>Dose (mg)</td><td>Dose (mg)</td><td>Dose (mg)</td></tr><tr><td>6-10</td><td>5</td><td>10-16</td></tr><tr><td>11-15</td><td>7.5</td><td>16-37</td></tr><tr><td>16-20</td><td>10</td><td>26-37</td></tr><tr><td>21-25</td><td>12.5</td><td>38-50</td></tr><tr><td>26-30</td><td>15</td><td>51-62</td></tr><tr><td>31-35</td><td>17.5</td><td>63-75</td></tr><tr><td>36-44</td><td>20</td><td>76-87</td></tr><tr><td></td><td></td><td>88-111</td></tr></table>	2-5 yr	6-11 yr	12+ yr	Weight (mg)	Weight (mg)	Weight (mg)	Dose (mg)	Dose (mg)	Dose (mg)	6-10	5	10-16	11-15	7.5	16-37	16-20	10	26-37	21-25	12.5	38-50	26-30	15	51-62	31-35	17.5	63-75	36-44	20	76-87			88-111	<p>Definition: failure of mechanisms responsive for seizure termination, leading to prolonged seizures with high risk of chronic consequences (neuronal death)</p> <p>Practical definition (for treatment): A seizure lasting longer than 5 minutes, or any ongoing seizures w/o return to baseline for 30 minutes.*</p> <p>*for convulsive seizures. Guidelines are not well-defined for non-convulsive seizures.</p> <p><i>Keep in mind, some of our Epilepsy patients have frequent and prolonged seizures every day that sometimes go beyond these criteria. It is often useful to ask the parents or consult clinic notes to get an idea of the severity of their Epilepsy.</i></p> <table><tr><th>Time</th><th>Agent</th></tr><tr><td>0-5 min</td><td>Lorazepam 0.1 mg/kg IV/IO/IM (max 4 mg)</td></tr><tr><td>5-15 min</td><td>Repeat lorazepam 0.1 mg/kg AND Fosphenytoin 20 mg/kg x1 IV/IO/IM (max 1,000 mg)</td></tr><tr><td>15-20 min</td><td>Phenobarbital 20 mg/kg x1 IV/IO OR Levetiracetam 30 mg/kg x1 IV/IO *if allergic, consider valproic acid 20 mg/kg IV/IO over 5 minutes</td></tr><tr><td>20-30 min</td><td>Repeat fosphenytoin 10 mg/kg OR Phenobarbital if LEV was used third, OR Levetiracetam if PHB was used third</td></tr></table>	Time	Agent	0-5 min	Lorazepam 0.1 mg/kg IV/IO/IM (max 4 mg)	5-15 min	Repeat lorazepam 0.1 mg/kg AND Fosphenytoin 20 mg/kg x1 IV/IO/IM (max 1,000 mg)	15-20 min	Phenobarbital 20 mg/kg x1 IV/IO OR Levetiracetam 30 mg/kg x1 IV/IO *if allergic, consider valproic acid 20 mg/kg IV/IO over 5 minutes	20-30 min	Repeat fosphenytoin 10 mg/kg OR Phenobarbital if LEV was used third, OR Levetiracetam if PHB was used third	<p>Headache:</p> <ul style="list-style-type: none"><input type="checkbox"/> Are you concerned for intracranial hemorrhage or impending herniation?<input type="checkbox"/> where is the pain (i.e. front, back, right, left)?<input type="checkbox"/> character (e.g. pounding, squeezing, sharp, etc.)<input type="checkbox"/> severity (1-10)<input type="checkbox"/> duration<input type="checkbox"/> frequency, change in frequency<input type="checkbox"/> time from onset to peak severity<input type="checkbox"/> are there associated symptoms (sensitivity to lights/noises, nausea/vomiting)<input type="checkbox"/> associated autonomic symptoms (e.g. eye tearing, eye redness, rhinorrhea, ptosis, change in facial color or temperature)<input type="checkbox"/> associated deficits (e.g. numbness, tingling, weakness, difficulty speaking or understanding others)<input type="checkbox"/> visual changes (double, blurry, flashes)<input type="checkbox"/> is the pain preceded by anything (scotoma, strange smell, feelings)<input type="checkbox"/> exacerbating factors (position, Valsalva, day/night, activity)<input type="checkbox"/> alleviating factors<input type="checkbox"/> do the headaches wake the patient from sleep and if so at what time?<input type="checkbox"/> family history<input type="checkbox"/> what is their neurologic examination?<input type="checkbox"/> what medications does the patient take to prevent headaches?<input type="checkbox"/> what medications has the patient taken to abort the headache?<input type="checkbox"/> what has he/she been given so far?	<p>Seizures:</p> <ul style="list-style-type: none"><input type="checkbox"/> actively seizing? Concern for herniation?<input type="checkbox"/> seizure history<input type="checkbox"/> baseline frequency, duration<input type="checkbox"/> semiology (not "GTC" – describe what happens)<input type="checkbox"/> history of status epilepticus?<input type="checkbox"/> clinic provider – Epilepsy or Neurology?<input type="checkbox"/> recent medication changes<input type="checkbox"/> current AEDs, dosing in mg/kg/d, dose timing<input type="checkbox"/> Missed/late AED doses?<input type="checkbox"/> When is next dose due? Get trough levels?<input type="checkbox"/> How is this seizure presentation different from their baseline/typical?<input type="checkbox"/> Current contributing factors (e.g. illness)?<input type="checkbox"/> Baseline developmental level (how much do they move/interact/see at baseline)? Are they now at their baseline?<input type="checkbox"/> Exam including VS (any O2 requirement), mental status (describe what they do spontaneously, and in response to a stimuli) <p>Stroke:</p> <ul style="list-style-type: none"><input type="checkbox"/> Stroke STAT (call 52170)? Acute/current neurologic deficits?<input type="checkbox"/> Last seen well time (if <5h, consider Stroke STAT; if >5h, call neuro consult)<input type="checkbox"/> acuity of onset?<input type="checkbox"/> deficits: speech (nonsensical, slurring, output), comprehension, vision (loss/double), vertigo, weakness, numbness, coordination, gait<input type="checkbox"/> Symptoms now (better, worse, same)<input type="checkbox"/> Risk factors: sickle cell, cardiac disease/shunt, personal or family history of stroke or clots (DVT/PE, miscarriage, stroke), hypercoagulable state
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