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Neuro Nugget: Seizures



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AMERICAN PHYSICIAN INSTITUTE FOR ADVANCED PROFESSIONAL STUDIES

Neuro Nugget: Seizures

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Seizure: Definitions & Background

- Seizures are defined as disorders characterized by excessive or oversynchronized discharges of cerebral neurons. Note the concept of "cerebral" neurons.
- Seizures Disorders are Classified as follows:
 - Generalized seizures
 - Tonic-Clonic Seizures (previously called Grand Mal Seizures)
 - Absence (or Petit Mal) Seizures
 - Tonic Seizures (stiffening seizures)
 - Clonic Seizures (shaking seizures)
 - Myoclonic Seizures
 - Atonic Seizures (drop attack seizures)
 - Partial Seizures
 - Simple Partial Seizures
 - Complex Partial Seizures (Temporal or Psychomotor)
- Etiologies include
 - o Primary Neurologic Disorders:
 - Benign Febrile Convulsions: Childhood seizures, occurring in 2-4% of children 3 months to 5 years of age. Occur with fever and in the absence of CNS infection. Last for 15 minutes or less and lack local features. 90% of children have recurrences. Probability of developing a chronic seizure disorder is 2-6%.
 - Idiopathic Epilepsy: idiopathic means that no specific etiology can be established. Account for up to 75% of all seizures. Usually begins between 5-25 years of age. 30-70% of patients will have recurrences.
 - Head Trauma: common when due to a depressed skull fracture or intracerebral or subdural hemorrhages.
 - Stroke: Strokes that effect the cerebral cortex lead to seizures in 5-15% of patients. Usually, post-stroke seizures do not indicate development of a seizure disorder and long-term anti-convulsant therapy often is not given.
 - Meningitis or Encephalitis: bacterial, fungal, tubercular, and parasitic infections can cause seizures.
 - Tumor and other mass lesions: glioblastomas, astrocytomas, meningiomas can cause seizures.
 - Systemic Disorders
 - Hypglycemia
 - Hypnatremia
 - Hyperosmolar states
 - Hvpocalemia
 - Uremia
 - Hepatic encephalopathy
 - Porphyria
 - Drug overdose or withdrawal
 - Global cerebral ischemia from cardiac arrest

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- Hypertensive encephalopathy
- Eclampsia

Seizure Assessment

- Main Point of Assessment: Must decide if generalized or partial. If partial, work up to find focal lesion
- Family History
 - History of childhood febrile convulsions
 - o History of seizures in family members.
 - History of neurologic disorders such as mental retardation and cerebral palsy. Ask about and *look* for neurocutaneous syndromes.
- Social History
 - o Travel-consider cystercicosis.
 - o Occupation-consider chemical exposure, head trauma.
 - o EtOH, IVDA (when in doubt get urine tox!)
- Witness descriptions of seizure
 - o Ictal cry? (tonic phase or seizure)
 - o Any tonic-clonic movements? Are they in all 4 extremities? (generalized)
 - o Did the movements happen all at once or march across the body (partial simple with Jacksonian march).
 - Were there any eye deviations or head deviation? (partial)
 - o Consciousness maintained? (simple partial seizures)
 - Consciousness clouded but partial (partial complex)
 - o Automatisms? (partial complex seizures)
- Physical Exam
 - o Dysmorphism of facial features.
 - o Signs of injury or trauma suggestive of brain injury.
 - o Mental status-evaluate for aphasia and short term memory loss.
 - o Field cut-for temporal lobe seizures.
 - o Fundi: papilledema, cherry-red spot
 - Eye movements: Look for nystagmus. Can be seen in certain epilepsy syndromes/toxicity of seizure meds.
 - o Nuchal rigidity: Kernig's/Brudzinski's for possible meningitis
 - o Cardiovascular system: r/o orthostatic hypotension
 - o Skin: any port-wine nevus? (Sturge Weber Syndrome)
 - o Cafe-au-lait spots? (Neurofibromatosis)
 - Hypopigmented macules? (tuberous sclerosis)
 - o Gingival hypertrophy? (dilantin use)
 - o Alopecia (depakote) or hirsutism (dilantin)?
 - o Ext: limb-length discrepancy- check thumb nail side by side
- Labs:
- o Accu check!
- o CBC (WBC), BMP (Na/Ca/Mg/Cr), CK/Trp, LFT, urine tox
- o AED levels
- o If infant/child, needs serum aminoacids, urine organic acids, baseline serum lactate/pyruvate.

- EKG: to rule out any cardiac arrhythmias or if concern for cardiogenic syncope.
- Chest x-ray: if any concern for aspiration during seizure.
- LP if has altered mental status, fever, nuchal rigidity. (Get head CT first!)
- Brain MRI pre and post infusion
- Why EEG
 - Seizures
 - New onset seizure in a patient.
 - Can assist in choosing anti-epileptic medication.
 - Increased seizure frequency or change in character of seizures in a known epileptic patient.
 - Consideration of anti-epileptic medication (AED) discontinuation.
 - Patients with history of well controlled seizures who wish to taper off medications.
 - Altered Mental Status
 - Can detect and grade encephalopathies from metabolic, toxic, infectious and iatrogenic etiologies.

Generalized Seizures

- Tonic phase lasts for about 10-30 s unconsciousness and extension of limbs and body arching occurs. Ventilation stops and cyanosis can manifest. Beginning of clonus can lead to a vocalization as air is exhaled from the contracting thoracic cavity. Usually, no aura is present (which would suggest secondarily generalized partial seizures).
- Clonic phase lasts for the length of the seizures alternating contraction and relaxation of limbs, neck, and torso. Ventilation returns in this phase.
- Recovery may last many minutes or even longer as the patient's level of consciousness returns to normal
- In the recovery phase, a neuro exam should be conducted to search for unilateral weakness. This would suggest a Partial seizure disorder with secondary generalization and the presence of a focal brain lesion.
- Status Epilepticus is defined as seizures that fail to abort spontaneously or that recur so frequently that full consciousness is not restored. It is a medical emergency because permanent brain damage may occur from anoxia or neuronal excitotoxic damage.
- EEG: spike and wave
- Treatment:
 - Start with Valproic Acid
 - o Adjunctive Meds: lamotrigine, topiramate, zonisamide, levetiracetam
 - o Consider also phenytoin, phenobarbitol, mysoline (older meds)

Absence Seizures

- Genetically transmitted.
- Always begin in childhood and usually end by age 20.
- Brief spells of 5-10 s with loss of consciousness but with retained postural tone. Recur hundreds of times a day. Leads to impaired school performance and social interactions (peers may find the child to be odd).
- Sometimes motor findings are present blinking, head turning, or automatisms
- EEG: 3 hz generalized spike and wave
- Treatment
 - Start Ethosuximide (and only for Absence Seizures0

Note that Carbamazepine may worsen Absence Seizures

Tonic, Clonic, and Myoclonic Seizures

- Tonic Seizures have the tonic phase without clonic phase. The tonic phase may last long enough to lead to cyanosis and anoxia.
- Clonic Seizures have the clonic phase without a preceding tonic phase.
- Myoclonic Seizures are brief, jerk-like contractions effecting the entire body or only certain muscle groups. This seizure type is related to several rare hereditary neurodegenerative disorders.
- Treat like Tonic-Clonic Seizures

Simple Partial Seizures

- In Simple Partial Seizures **consciousness is preserved** (despite presence of neuropsych symptoms as described below). Note that the seizure may secondarily spread to other cortical areas and at that point present with altered consciousness.
- Simple Partial Seizures can begin with motor, sensory or autonomic changes. The cortical area of discharge can spread and thus the seizure symptoms can spread across the body Jacksonian March
- Autonomic changes include sweating, flushing, piloerection, pupillary dilatation, and urinary incontinence.
- Neuropsychiatric symptoms may occur, such as feeling of déjà vu, distorted memory, visual illusions, hallucinations, affective lability, panic attack, slowed or forced cognition.
- Focal neurological deficits may be present in the post-ictal state, such as Todd's paralysis.

Partial Complex Seizures

- These seizures result in **impaired consciousness** without total LOC.
- Symptoms include
 - o Somatosensory: epigastric sensations
 - o Neuropsychiatric: fear, olfactory hallucinations, déjà vu
 - o Motor: automatisms such as sitting down, standing up, orobuccal movements, other facial or neck movements
- Seizures discharge from temporal lobe or medial frontal lobe.
- Seizures differ among patients but stereotyped for each individual.
- Last on average 1-3 min but up to 30 min.
- May secondarily generalize.
- Treatment of Simple and Complex Partial Seizures and Secondarily Generalized Seizures
 - Start with Carbamazepine
 - o Adjunctive Meds: Lamotrigine, Levetiracetam, Topiramate, Zonisamide
- In Partial Complex Seizure Disorder Surgery is superior to med response

Status Epilepticus

- Two Treatment Goals
 - o Aggressive Supportive Care
 - ABCs
 - Rapid glucose determination and correction
 - Hospitalize
 - Treat complications of status: hyperthermia, acidosis, hypotension, respiratory failure, rhabdomyolosis, or aspiration

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- o Termination of Seizure
 - Give AED if seizure continuing past 4-5 min
 - Give Lorazepam to abort
 - Or Alternatives: diazepam, midazolam
 - If refractory
 - phenytoin
 - general anesthetic: propofol
 - barbiturate: phenobarbitol

Other Treatment Tips

- Do not start anticonvulsant therapy after single new onset seizure: only 33% of second seizure in 5 yrs
- Rational AED choice
 - o Type of seizure
 - Safety profile
 - o Monotherapy when possible
 - Age of patient
 - o Drug-drug interactions
- Start monotherapy and increase until seizure-free or as can tolerate. If still with seizures, switch to another med as monotherapy with cross-titration or add adjunctive med to the first.
- Note that many of the new AED's are broad spectrum for both generalized and partial seizures
 - o Lamotrigine
 - Levetiracetam
 - Topiramate
 - o Zonisamide
- Renally excreted
 - Gabapentin
 - Levetiracetam
- Real Simple Seizure Treatment Tips
 - o Generalized: Valproic Acid
 - o Partial: Carbamazepine
 - o Absence: Ethosuximide
 - New AEDs: good for both generalized and partial