Seizures and Epilepsy Biomedical Engineering - URJC

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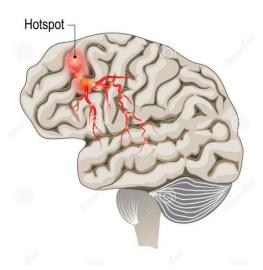


Overview

Overview of seizures

- **Definition:** Uncontrolled, abnormal electrical activity in the brain.
- Manifestations: Varying from tonic-clonic to absence seizures.
- Duration: Typically less than two minutes.

Characteristics



Introduction

Definition of seizures

- **Seizure:** Abnormal, excessive, synchronized **electrical discharge**.
- **Epilepsy:** Defined by **two or more** unprovoked seizures.
- Status Epilepticus: Enduring epileptic condition, a medical emergency.

Terms (I)

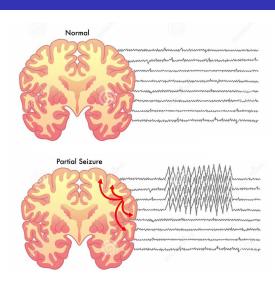
Seizures:

- They are only one manifestation of neurologic or metabolic diseases
- Paroxysmal phenomenons caused by electric hyper-excitability of a group of neurons
- Many causes, including a genetic predisposition, head trauma, stroke, brain tumors, alcohol or drug withdrawal, hypoglycemia.

■ Epilepsy:

- Medical disorder marked by recurrent (more than two), unprovoked seizures
- 2 Chronic condition

Characteristics



Terms (II)

Status Epilepticus

- Single generalized convulsion lasting more than five minutes or a series of generalized seizures without full return of consciousnes.
- Enduring epileptic condition.
- There are many types of status epilepticus.
- Generalized convulsive status epilepticus is a medical emergency.

Etiology

- **Provoked Seizures:** Result from various causes like electrolyte disorders, head injury, toxins, etc.
- Unprovoked Seizures: It's the definition of epilepsy, occur without an apparent cause.
- **Incidence:** 25-30 % provoked, 70-75 % idiopathic (unknown origin).

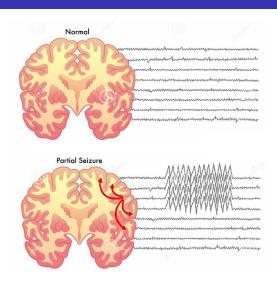
Epidemiology

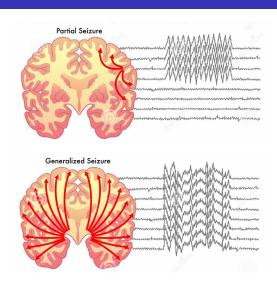
Epidemiology

- Epilepsy incidence is highest in **younger** and **older** age groups and increases steadily after 50 years of age.
- The most common cause of seizures and epilepsy in **older** people is **cerebrovascular disease**.

- **Seizure Threshold:** Everyone has **some propensity** to have seizures. Influenced by medications, genetics, electrolyte imbalances, etc.
- On a cellular Level: Excitation of cerebral neurons, involving neurotransmitters like glutamate and GABA, that can trigger epileptiform activity.
- Disbalance of neurotransmitters, for any cause: electric discharge.
- Status Epilepticus: If prolongued over time: can cause systemic changes, damaging to the brain.

- Seizures occur due to an epileptogenic lesion, and then they can propagate or not.
- Some focal onset seizures can be preceded by an aura, which refers to symptoms and signs that occur before the onset of seizure activity. These symptoms may include vision changes, dyspepsia, déjà vu, paresthesias, hearing disturbances, and sensation of abnormal taste or smell.





Type of Seizures

Type of movements

- Motor Onset: Focal and generalized seizures with motor manifestations.
- Non-Motor Onset: Focal and generalized seizures with non-motor manifestations.

Level of awareness

- 1 Conscious: it's called SIMPLE.
- Unconscious: it's called COMPLEX.

The term complex means that the patient loses consciousness. "Simple means that the patient remains aware (i.e., conscious).

Type of Seizures



Seizures that begin here may cause a person to experience an odd smell, odd taste, buzzing or ringing in the ears, fear or panic, dėjà vu, or abdominal discomfort.

FRONTAL LOBE

Seizures that begin here may cause loss of motor control, a change in behavior, or change in language expression.

OCCIPITAL LOBE

Seizures that begin here may cause a person to see multicolored shapes, such as circles and flashes, or experience temporary loss of vision.

PARIETAL LOBE

Seizures that begin here may cause a person to feel numbness or tingling, or feel burning or cold sensations.

Partial or focal seizure

A focal onset seizure refers to abnormal neural activity in only one brain area within one brain hemisphere with a fixed focal or localized onset. They can be divided in:

- **1 Simple Partial Seizures**. The patient loses consciousness.
- Complex Partial Seizures. the patient remains aware (conscious).

Motor symptoms

Motor manifestations of focal motor onset seizures include abnormal movements, like the following:

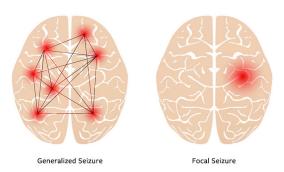
- Tonic movements.
- Clonic movements are characterized by repeated, short contractions.
- Atonic movements are characterized by loss of tone in a limb.
- Myoclonic movements are characterized by irregular, nonrhythmic jerking of the limbs.

Non-motor symptoms

The clinical manifestations of focal non-motor seizure include autonomic, behavioral arrest, cognitive, emotional, or sensory symptoms:

- Changes in blood pressure, heart rate, sweating, or digestive upset.
- Behavioral seizures: weird behaviors.
- Cognitive seizures: abnormal language or thinking, jamais vu, déjà vu, hallucinations.
- Emotional changes such as fear, dread, anxiety, or pleasure.
- Sensory seizures: abnormal sensations of vision, paresthesias, hearing, smell, or pain.

A focal seizure can rapidly spread and evolve into a bilateral tonic-clonic seizure, that is, becomes a **secondary generalized seizure**.



Generalized seizures

Generalized seizures

- A generalized tonic-clonic seizure, also known as grand mal seizure, is defined as a seizure that has a tonic phase followed by clonic muscle contractions.
- Among patients, families, and observers, they are most feared of seizure types.
- They are usually associated with impaired awareness or complete loss of consciousness.

Generalized seizures



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Generalized seizures

Generalized seizures

- They result from diffuse cortical activation at seizure onset (primarily generalized)...
- or generalization of partial seizure activity (secondary generalized).

Absence seizures consists of generalized, **non-convulsive** seizures characterized by impairment of awareness and intermittently has other manifestations such as automatisms or subtle myoclonic, tonic, atonic, or autonomic phenomena.

Special Types of Seizures

- Absence Seizures: Brief, unresponsive seizures, common in children.
- **Febrile Seizures:** Associated with fever, typically in young children.
- Juvenile Myoclonic Epilepsy: Involves absence, myoclonic, and tonic-clonic seizures.
- Lennox-Gastaut Syndrome: Rare, severe childhood epilepsy with multiple seizure types.

Evaluation and management

Evaluation and management

- Clinical Assessments: History, physical examination, laboratory work, neuroimaging, EEG.
- **EEG:** Biomarker for epilepsy, helps in risk stratification.
- **Treatment:** Address reversible causes, medication options for chronic seizure disorders.

Diagnosis

- **Testing:** Electrolytes, serum glucose, sodium.
- Consider Lumbar Puncture: For fever, immunosuppression, or signs of CNS infection.
- Neuroimaging: Suspicion of acute intracranial process, focal findings on neurologic examination, head trauma, malignancy history, etc.

Electroencephalography (EEG)

- **Biomarker for Epilepsy:** Identifies focal or generalized epileptiform discharges.
- Risk Stratification: Assists in determining the likelihood of future seizures.
- Follow-up: Abnormal EEG may prompt MRI to locate structural abnormalities.

Treatment Options

- Addressing Reversible Causes: Correcting abnormalities like hypoglycemia, alcohol withdrawal. NO NEED OF MEDICATION.
- **Medication if indicated:** Carbamazepine, benzodiazepines, topiramate, valproic acid, phenytoin, etc.