

Epilepsy in the Adolescent Population

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Epilepsy diagnosis

Basic approach

- Is the epilepsy focal or generalized?
 - Semiology
 - EEG features
- Diagnostic workup to identify etiology of seizures and epilepsy syndrome
 - Structural
 - Metabolic
 - Genetic

Epilepsy diagnosis: Relevance to adolescent population

- Current operational clinical definition of epilepsy: At least 2 unprovoked (or reflex) seizures occurring > 24 h apart
 - Unprovoked seizure definition
 - Provoked seizure definition
- Recurrent reflex seizures in response to photic stimulation
- Diagnosis of a specific epilepsy syndrome after a single seizure

Epilepsy classification

1. Localization-related epilepsy in adolescents
 - Non-lesional focal epilepsy: 20-30%
 - Lesional focal epilepsy: 20-30%
 - Pharmaco-resistant epilepsy: 20%
2. Idiopathic generalized epilepsy (IGE)
 - IGE constitutes 15-20% of all epilepsies in adults and children
 - Classification of the electroclinical syndrome based on:
 - Seizure types
 - Age of onset

- EEG characteristics
- Electrographic signature of IGE:
 - Interictal and ictal discharges
 - Normal background
 - Photoparoxysmal response
 - Atypical features
 - Fragments of generalized discharges

IGE subtypes in the adolescent population

A. Juvenile Absence Epilepsy

- Clinical features
 - Age at onset
 - Seizure types
- EEG features
 - Generalized spike wave with faster frequency
 - Polyspikes
 - Activation procedures
- Treatment
 - valproate
 - lamotrigine
 - ethosuximide may be effective for absences only
- Prognosis
 - Typically lifelong

B. Juvenile Myoclonic epilepsy

- Clinical features
 - Age at onset
 - Seizure types
- EEG features
 - Generalized 4-6 Hz polyspike and polyspike-wave discharges
 - Correlate with a myoclonic jerk
 - Photoparoxysmal response in up to half of cases
- Treatment
 - Valproate with high efficacy (use limited in women)
 - Lamotrigine is often first choice (may worsen myoclonic seizures)
 - Others: topiramate, levetiracetam, zonisamide
 - Carbamazepine, phenytoin, vigabatrin can worsen seizures

- Prognosis
 - Most cases are pharmaco-responsive
 - Lifelong therapy
 - Percentage achieving remission off medication
- C. Generalized tonic-clonic seizures upon awakening (GTCA)/Epilepsy with Grand Mal Seizures upon awakening (EGMA)
 - Clinical features
 - Age at onset
 - Seizure types
 - EEG features
 - Generalized spike-wave discharges
 - Polyspikes
 - Photoparoxysmal response
- D. Genetic Generalized Epilepsy NOS
 - IGE subgroups not recognized by the ILEA
 - All exhibit EEG features similar to classic IGE
 - Distinct sub-syndromes
 - Perioral myoclonia with absences (onset 2-13 yrs)
 - IGE with phantom absences (onset of first GTC in adulthood)
 - Jeavons syndrome (typically presents in childhood)
 - Adult onset IGE (onset at > 18-20 years)

Non-epileptic Paroxysmal Events in the Adolescent Population

- A. Syncope
 - Most common non-neurologic mimic of epilepsy
 - Vasovagal syncope most common
 - EEG findings during a syncopal event
- B. Parasomnias
 - REM sleep behavior disorder
 - Clinical features
 - Non-REM sleep parasomnias
 - Clinical features
 - EEG features
 - Comparison to nocturnal (epileptic) seizures

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- Clinical features
 - EEG features
 - C. Psychogenic Non-epileptic Spells
 - Diagnosis of psychogenic non-epileptic spells
 - Historical features
 - Clinical features differentiating them from epileptic seizures
 - Semiology of childhood psychogenic non-epileptic spells
 - EEG features that support the diagnosis of non-epileptic spells

Treatment adherence in the adolescent population

Factors affecting adherence:

- in adults
- versus in adolescents

Potential solutions to barriers to medication adherence

- Internet based programs
- Smartphone applications

Psychosocial considerations in adolescents with epilepsy

- Driving
- Sports
- Emerging independence
- Parental protection

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