Seminar in Epileptology

Epileptic Disord 2020; 22 (4): 399-420

How to diagnose and classify idiopathic (genetic) generalized epilepsies

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Received April 06, 2020; Accepted May 24, 2020



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Idiopathic (Genetic) Generalized Epilepsies

- Electroclinically well-defined group that accounts for almost one third of all people with epilepsy
- Consist of four well-established syndromes and some other rarer phenotypes

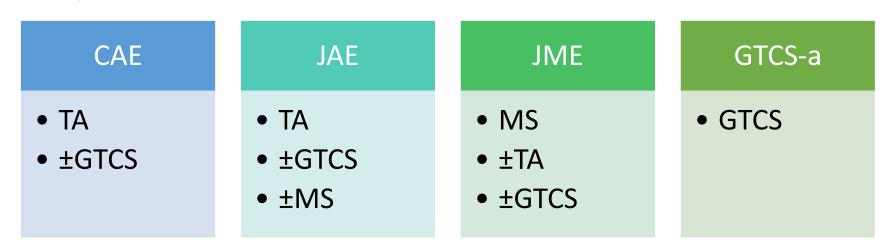
childhood absence epilepsy juvenile absence epilepsy juvenile myoclonic epilepsy IGE with generalized tonic clonic seizures alone



older

Seizure types

- Three main seizure types
- Occur either alone or in any combination
 - generalized tonic-clonic seizures
 - typical absences
 - myoclonic seizures





Diagnosing IGE

Thorough medical history

- -tructured interviews may aid young clinicians
- Asking too many open-ended questions may distract the patient
- Asking only yes/no questions may create a suggestion bias
- Specific focus on seizure types, their age at onset, timing and triggers
- Co-existance of multiple seizure types
- Comorbidities and family history should be questioned comprehensively
- Obtain collateral history from witnesses when possible

Neurological and physical examination

• Additional neurological or cognitive signs and symptoms are red flags

EEG is indispensable

Imaging is not essential in typical cases



EEG in IGE

- «Normal» routine EEG is not unexpected
- Methods to increase sensitivity
 - Sleep deprivation and sleep can enhance generalized spike-wave discharges (GSWD)
 - Hyperventilation is the foremost method triggering GSWDs and TA
 - **Photic stimulation** can reveal photosensitivity
 - *Historical triggers* can be tested under controlled and safe conditions
 - Scheduling the *EEG* appointment to *after awakening* may disclose GSWDs and MS which could have been otherwise missed in a JME patient
 - 24-hour EEG or longer telemetry may allow registration of GSWD when others do not
- Clinicians should be precise about the objective of the EEG and know what they are looking for in order to find it

