

# **Epilepsy and epileptic seizures II.**

## **Classification and treatment**



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**Epileptic seizure  $\neq$  Epilepsy**

# Classification of epileptic seizures

## ■ Partial

- Simplex
  - Motor (cloni, version)
  - Sensory (auras: somato-sensory, temporal auras)
- Complex (loss of consciousness with oral and manual automatisms)

## ■ Generalized

- Absence
- Myoclonic
- Generalized tonic-clonic (grand mal seizure)

# Classification of epileptic seizures

- **Partial** (origin: circumscribed brain region)
  - **Simplex**
    - Motor (cloni: Jackson march)
    - Sensory
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# Sensory auras

- Simplex:
  - Somatosensory
  - Visual
  - Olfactory
  - Auditory (very rare)
  - Vertigo (very rare)
- Complex (usually temporal)
  - Gastric aura
  - Affective (positive or negative)
  - Dysmnestic (de ja vu, jamais vu)

# Complex partial seizures /psychomotor seizures/

- Occurs in temporal lobe epilepsy
- Preceded by „temporal” aura (gastric sensation, de ja vu., Olfactory auras)
- Consists of:
  - Loss of consciousness
  - Oral automatisms (swallowing, biting, munching)
  - Manual automatisms

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- ```
graph TD; subgraph Partial; P1[Simplex]; P2[Complex]; end; subgraph Generalized; G1[Absence]; G2[Myoclonic]; G3[Generalized tonic-clonic]; end; P1 --> G3; P2 --> G3; G1 --> G3; G2 --> G3;
```



## **Generalized tonic-clonic seizure:**

- it may be primer,
- But can be preceded by any other seizure types. („secondarily generalized seizure”: final common pathway)
- sequence: vocalisation, tonic phase (with apnea), clonic phase,
- postictal confusion or postictal sleep
- Postictal examination (not obligate signs):
  - tongue biting,
  - enuresis,
  - muscle pain
  - trauma,
  - Todd paresis: transient paresis due to exhaustion (only after some focal onset grand mal seizures)

# Epilepsy syndrome = a chronic condition with a typical symptoms

- Typical seizure type
- Typical etiology (genetics vs. Acquired)
- Typical localisation of the seizure focus in the brain (circumscribed or non-localisable)
- Typical age at onset

# Classification of epilepsy syndromes

Two main axis:

- Genetical (=idiopathic) or acquired
- Focal (localisation-related) or generalized

## **Focal epilepsy**

1. Benign centro-temporal epilepsy

## **Idiopathic (genetical) epilepsies**

## **Non-idiopathic epilepsies (symptomatic, cryptogenic)**

1. Temporal lobe epilepsy
2. Frontal lobe epilepsy
3. Parietal lobe epilepsy
4. Occipital lobe epilepsy

## **Generalized epilepsy**

1. Absence epilepsy
2. Juvenile myoclonic epilepsy
3. Grand mal epilepsy on awaking

1. West syndrome
2. Lennox-Gastaut syndrome

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# Temporal lobe epilepsy

- Start: 5-18 y
- Auras: usually gastric sensation but psychic (affective), dysmnestic (de ja vu) or smell auras are also common
- Auras can be isolated but usually followed by complex partial seizures (LOC, automatisms) Grand mal seizures are rare
- Memory disturbances are characteristics
- EEG: temporal lobe epileptic focus
- MRI: usually hippocampalis sclerosis, occasionally: tumor or dysgenesis
- 50% of patients had febrile convulsions in childhood

# Idiopathic generalized epilepsy

1. generalized seizure without focal features:
  - Absence: Childhood Absence epilepsy
  - Myoclonic and grand mal seizures: Juvenile myoclonic epilepsy
  - Pure Grand mal seizures: Grand mal on awaking epilepsy
2. Sleep deprivation and alcohol can provoke seizures
3. Often (not always) positive family history of seizures
4. Typical age at onset (Juvenile myoclonic epilepsy: 13-16 ys)
5. Normal intelligence
6. EEG: 3 Hz- spike and wave complex
7. Normal brain MRI
8. Seizures responds to valproic acid in  $> 90\%$  of patients

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## Focal epilepsy

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## Focal drugs

### Idiopathic (genetical) epilepsies

### Non-idiopathic epilepsies (symptomatic, cryptogenic)

## Focal drugs

1. Temporal lobe epilepsy
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## Surgery

## Generalized epilepsy

1. Absence epilepsy
2. Juvenile myoclonic epilepsy
3. Grand mal epilepsy on awaking

## Broad-spectrum

## drugs

1. West syndrome
2. Lennox-Gastaut syndrome

# Antiepileptics: 60-70% of epileptic patients become seizure-free on chronic drug treatment

## ■ Focal drugs:

- 1. Carbamazepine
- Oxcarbazepine
- Phenytoin
- Gabapentin

## ■ Broad spectrum drugs:

- 1. Valproic acid
- Lamotrigine
- Benzodiazepines
- Phenobarbital
- Levetiracetam

Almost all antiepileptic drugs should be  
started with low doses and titrated  
slowly



# Status epilepticus

- Definition:
    - If the seizures reoccur and the patient is not conscious between two seizures
  - OR
  - Seizures occurred in  $> 30$  minutes
- 
- Status epilepticus can be developed from any seizure types

# Status epilepticus is dangerous

- Life threatening conditions:
  - grand mal status epilepticus
  - status epilepticus accompanied by extended motor phenomena (cloni)
- without adequate treatment patients may die due to
  - Aspiration
  - Aspiration-related infections
  - Cardiopulmonal insufficiency

# Status epilepticus - treatment

- If an epileptic patient has a seizure, no additional treatment (no hospital, no acute medication) is needed if there is no injury
- status epilepticus: should be treated immediately and the patient should be transported to neurological department, neurointensive units

# Status epilepticus – treatment 2

- ABC (free airways)
- FIRST STEP: iv benzodiazepine (20 mg diazepam vagy 2-3 mg clonazepam iv., or lorazepam)
- It can be repeated many times
- Benzodiazepine should be given very slowly because it can cause breathing stop
- Intramuscular benzodiazepine: no sense and should be avoided
- If there is no vein: 20-30 mg rectal diazepam

## Status epilepticus – treatment 3


- If the status epilepticus occurs in a patient without known epilepsy, then iv. glucose, vitamine B1 iv. should also be given immediately
- By adequate treatment the mortality of status epilepticus is low unless there is a serious underlying disorder (meningitis, encephalitis, stroke, SAH, tumor cerebri, CNS injury, delirium tremens)



# Refractory status epilepticus

- Repeat benzodiazepines
- Iv. phenytoin or valproic acid (start of action is late! After 15-60 min)
- Further seizures: full narcosis with short-acting barbiturates or midazolam or propofol

# Epilepsy surgery

- Localisation-related epilepsy *and* which is refracter to 2-3 antiepileptic drugs
- 
- Epilepsy surgery should be considered
    - Epilepsy surgery can be performed
      - if the epileptic focus can be localised (MRI, video-EEG, neurpsychological investigations)
      - and the focus can be removed without neurological, neuropsychological consequences
    - Generally **60-90%** of patients became seizure free after well-indicated surgery

# Treatment of Temporal lobe epilepsy

- Bad pharmacological prognosis:
  - In 75-90% pharmacoresistant
- Good surgical prognosis:
  - 75-90% of patients become long-term seizure-free
- Surgery: partial anterior temporal lobectomy

# EEG in epilepsy

- Routine, scalp-EEG with hyperventilation
- Sleep-EEG
- long-term EEG
  - „Holter-EEG”
  - Video-EEG monitoring

# Idiopathic generalized epilepsy

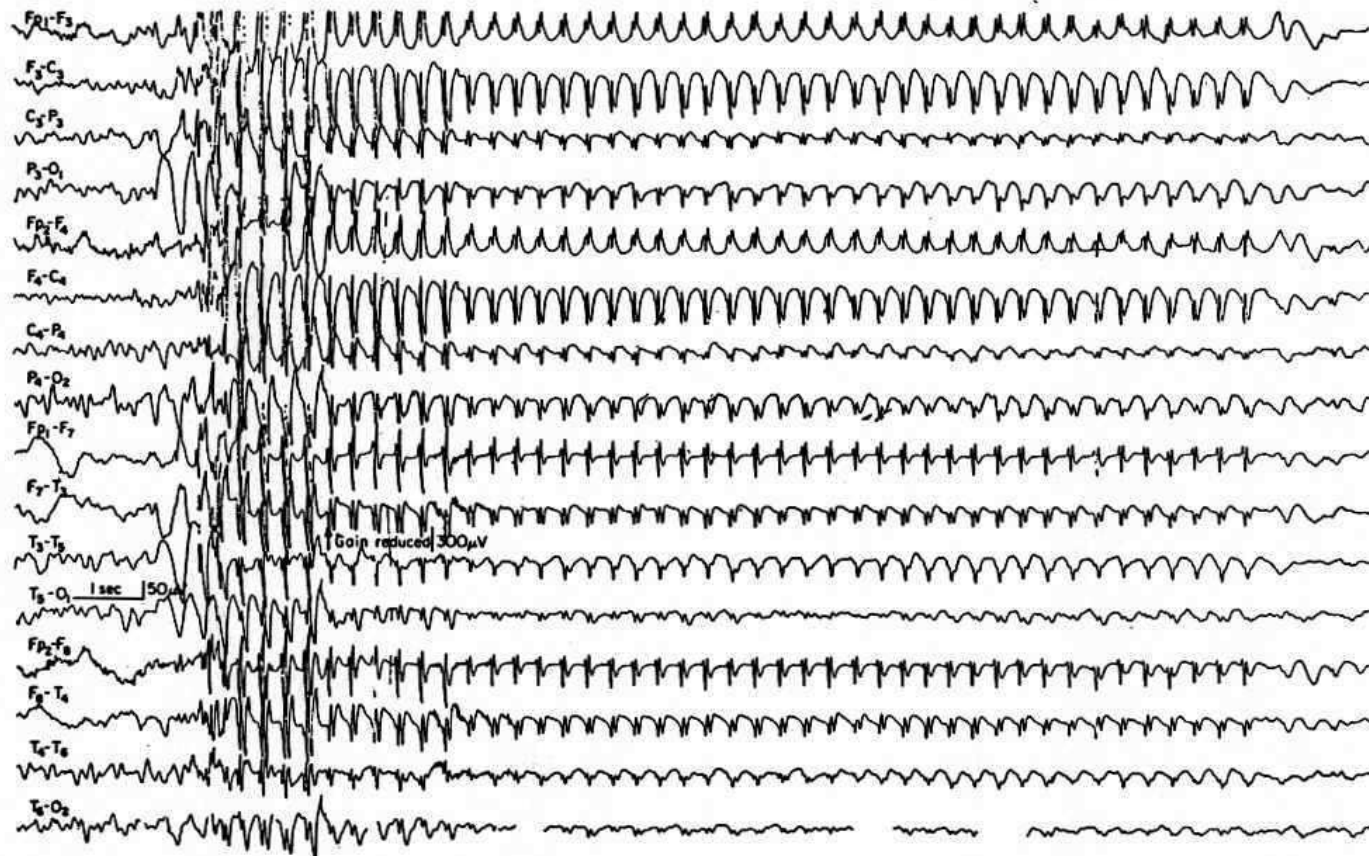


Fig. 23.3. Petit mal absence, age 8 yr. The spike wave burst is preceded by an aborted run of rhythmical posterior slow 3/sec waves; after two slow waves, the attack begins.

# Temporal lobe epilepsy

# Sensitivity of routine EEG:

- Absence 92%
- Other Idiopathic generalized epilepsy 58%
- Focal epilepsy: 30%



Normal EEG does not exclude and the  
abnormal EEG does not prove epilepsy:  
Epilepsy is a clinical diagnosis

# MRI

In all epilepsies beginning in adulthood  
MRI examination is obligatory

**Post-stroke**

brain tumor

Vascular malformations of the brain

**Post-encephalitis**

Malformations of cortical development (dysgeneses)      Posttraumatic

**Hippocampal sclerosis**