SEIZURES AND EPILEPSY: PATHOPHYSIOLOGY AND PRINCIPLES OF DIAGNOSIS

Thomas Pt. Henry, MD1

(Scroll down for answers and explanations, which appear after question 5.1

CALLESTIONS

- 1. A full-term male recease has repeated tionsic metaures and the
 - electroencephalogram (EEG) alvows a generalized burst suppression pattern interiorally. What is the Shely discoverie?
 - A. Benign familial reconstal epilepsy
 - B. Early myoclonic encephalopathy
 - C. Charlesiania prendinomia
 - Cl. West syndrome
 - E. Myoclonic epilepsy in intency.
- 2. A flyth-team female infant at 10 months has parmel development, single or clustered focal or massive jerks, and a normal interioted BBG. What is the likely disconnection?
 - A. Epilepsy of intancy with migrating focal Market Street or Street
 - **B.** West syndrome
 - C. Myoclonic epilepsy in intency
 - Ch. Disavest symphomes
- E. Progressive myocionus epilepey 3. A 5-year-old boy has new onset of rational temperation of medianes as well a on selectors with empairs and unresponsiveness. The interioral EEG shows generalized 3-per-second spikewaye bursts and bilateral occipital apilities. What is the likely discovers?
 - A. Lennou-Gastaut syndrome
 - R. Childhood absence epileosy
 - C. Benign epilepsy with centrotemporal Secretarion of
 - D. Panaylotopoulos syndrome
 - E. Epilepsy with myocionic absences.
- 4. A 25-year-old graduate student has a first nevalized tomic-clionic seizure after staying up all night to study, has a negative wrine taxicology screen in the ergency department, and has a normal cranial computed tomography (CT) scan. Later in the neurology clinic, on further questioning the reports that for the last year she has had multifocal jeries on recomings after staying up late. The brainmagnetic resonance imaging (MRII) study

atheres auditte bitmontal autonoriinal white matter signal changes. The interiotal EEG is normal except for a single brief burst of diffuse, bifrontally predominant spikewave complexes repeating at about 4 Hz. What is the likely diagnosis?

- A. Juvenile absence epitepsy
- Juvenile myodlonic epilepsy.
- C. Progressive myodionus epileasy
- D. Autosomal-dominant necturnal frontal PORTOR RECORDS INCOME.
- E. Plasmussen syndrome
- 5. A 28-year-old cafeteria worker has a history of febrile commutations in early childhood and poor school performance. Non-sentences become property of four property years off entirelaure medications, until a eneralized tomic-clonic seleure is observed at work. In retrospect, the patient has had intense "daydreams" at work or home for the last year, and others often have had to repeat his name to get. him to answer. The interictal EEG shows frequent delta-theta transfents over the right temporal lobe and occusional right temporal agriko wawe complexes. On brain MRI the right hippocampus is smaller and brighter than the left hippocampus. What is the likely diagnosis?
 - A. Epitegsy with generalized tonic clonic section atoms
 - B. Autosomal dominant epilepsy with Intelligence Specificantelli-
 - C. Lectron Cinetaut syndrome
 - D. Gelastic selbures with hypothalamic Programme Services
 - E. Mesial temporal lobe opliepsy with Microscopimical actionosis.

AMESIANTERS:

- The correct answer is (C), Ohtshara.
 - syndrome. Only the syndromes in answers A. B. and C-benign femilial reconstal. epitecey (ISFNII), early myoclonic encephalopathy, and Ohtahara syndromepopular in meanwheel. Of theses, only DENE and Ohtahara syndrome involve tonic seisures Ealthough West syndrome involves tonic solzures as well). Finally, only the Ontahura

- supplierance background background-supplierations are the HEREON.
- The correct enswer is ICs. Myoclonic epillopsy in intancy. Only the syndromes in answers C and E-myoclonic epilepsy in intency and progressive myoclonus epitepsy -involve myodonus without other seizures. and of the listed sundrames, only myoclanic epitiopsy in intency features normat development. In addition, mycclonic epilepsy in Infancy, along with Dravet syndrome, often has normal interletal EDGs in infancy."
- 3. The correct answer is (D),
 - Panayiotopoulos syndrome Monoporcyalative selections with lotal emesis are come exponent in the Paragologicanousless. syndrome. Interiotal generalized 3-persecond spike wave bursts and bilateral occipital spikes can occur in benign epilepsy with controllersposal spikes (BECTS) and Panishotopoulos syndrome, but the absence of centrolemporal spikes excludes BECTS.11
- 4. The correct answer is (8), Juvenille myoclonic policety. Juvenile myoclonic epilepsy (JME) most often features. generalized tonic-clonic seizures and myoclomus on wakening; each of the other Sinther's exercises remaining at leasest mean additional type of seizure that this patient does not have. Generalized fast solkeways and polyspike-waves discharges are most characteristic of JME, and not of the other syndromes. Onset of myocionus in JME occasionally occurs in early adulthood, although enset of myodishus usually is descriptions.
- 5. The correct answer is (E), Mesial temporal lobe epilepsy with hippocampal aclerosis. The brain MRII is described as showing the fundament of histogramman actions is, and these together with the ignitateral interictal. temporal EEG spikes are competible only partition properties the management tenture continuous of TO ETA with hippocampal scienosis. Many patients with messal TLE can provide a more definite. description of auras and other lotal and posticial phenomena, but subjective symptoms of the partial seigures in medial TLE also can be quite vague."

Epilepsy Board Review Manual

Peipei Pang



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