

Clinical Case Studies in Epilepsy

INTRODUCTION

A *seizure* is a temporary alteration in brain function due to excessive and/or hypersynchronous neuronal activity. Epilepsy is the tendency to have recurrent unprovoked seizures. Approximately 10% of individuals will have 1 seizure in his or her lifetime, and up to 3% will suffer from epilepsy.¹ However, the prevalence of active epilepsy is only 0.8%. In the United States, approximately 2 million people have epilepsy, with 100,000 new cases diagnosed per year. People of every background and age are affected. Epilepsy is associated with impaired quality of life, and its treatment can have serious consequences. Unfortunately, 20% to 40% of patients with epilepsy continue to experience occasional seizures despite treatment.²⁻⁴

Epilepsy is not a single condition but a symptom of various disorders and reflects underlying brain dysfunction. The International League Against Epilepsy (ILAE) currently has 2 classification schemes for epilepsy that are designed to be used together. The first divides seizures into 3 types, with subtypes of each: partial (focal seizures involving only part of the brain), generalized (seizures involving both hemispheres of the brain), and unclassifiable (Table 1).⁵ This system allows for simple classification that may determine diagnostic evaluations, choice of medication, and prognosis. A supplement to this system, the ILAE Classification of Epilepsies and Epileptic Syndromes,⁷ divides epilepsies into 4 groups: localization-related (involves 1 or more focal areas of the brain), generalized (involves both hemispheres of the brain at the same time), undetermined, and special syndromes. The localized and generalized groups further divide into idiopathic (no identifiable cause and no associated neurologic abnormalities, although underlying genetic mutations are increasingly being discovered), symptomatic (cause is identified), or cryptogenic (presumed symptomatic, but the cause is unknown) epilepsy syndromes. Epilepsy syndromes are defined by the specific seizure type, clinical findings (including age of onset), and type of EEG abnormality. Epilepsy syndromes include the catastrophic epilepsy syndromes of infancy and childhood (eg, West syndrome, Lennox-

Gastaut syndrome, myoclonic epilepsies), idiopathic partial epilepsy syndromes (eg, benign epilepsy with centrotemporal spikes), idiopathic generalized syndromes (eg, benign neonatal convulsions, juvenile myoclonic epilepsy [JME]), and special syndromes such as febrile seizures. A specific epilepsy syndrome may require specific anticonvulsant drug treatment and is frequently associated with a predictable prognosis. The ILAE is currently developing a third diagnostic scheme aimed at providing a standardized description of individual patients rather than a fixed classification.⁶ This scheme uses 5 axes: ictal phenomenology (axis 1); seizure type (axis 2); syndrome, when known (axis 3); genetic defect or specific pathologic substrate for symptomatic focal epilepsies (axis 4); and impairment classification (axis 5).

This review presents 4 cases that evolve over the course of the discussion to encompass evaluation of a first seizure, diagnosis and management of epilepsy in the elderly, management of epilepsy during pregnancy, and evaluation and management of acute mental status change in a patient with epilepsy.

EVALUATION OF A FIRST SEIZURE

CASE 1 PRESENTATION

A 10-year-old woman is referred to a neurologist for evaluation after experiencing a convulsion while jogging. The patient is otherwise healthy and takes no medications besides oral contraceptives. Records from the emergency department (ED) visit document that a physician who was jogging behind the patient witnessed the event and did not notice any prodrome or focal findings. Evaluation in the ED showed a postictal period of confusion for 20 minutes followed by a normal examination. Electrolytes, renal function, complete blood count, and a computed tomography (CT) scan of the patient's head were normal.

DIAGNOSING EPILEPSY

- What is the differential diagnosis for a seizure?

Epilepsy In Clinical Practice A Case Study Approach

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Table of Contents Epilepsy In Clinical Practice A Case Study Approach

1. Understanding the eBook Epilepsy In Clinical Practice A Case Study Approach
 - The Rise of Digital Reading Epilepsy In Clinical Practice A Case Study Approach
 - Advantages of eBooks Over Traditional Books
2. Identifying Epilepsy In Clinical Practice A Case Study Approach
 - Exploring Different Genres
 - Considering Fiction vs. Non-Fiction
 - Determining Your Reading Goals
3. Choosing the Right eBook Platform
 - Popular eBook Platforms
 - Features to Look for in an Epilepsy In Clinical Practice A Case Study Approach
 - User-Friendly Interface
4. Exploring eBook Recommendations from Epilepsy In Clinical Practice A Case Study Approach
 - Personalized Recommendations
 - Epilepsy In Clinical Practice A Case Study Approach User Reviews and Ratings
 - Epilepsy In Clinical Practice A Case Study Approach and Bestseller Lists
5. Accessing Epilepsy In Clinical Practice A Case Study Approach Free and Paid eBooks
 - Epilepsy In Clinical Practice A Case Study Approach Public Domain eBooks
 - Epilepsy In Clinical Practice A Case Study Approach eBook Subscription Services
 - Epilepsy In Clinical Practice A Case Study Approach Budget-Friendly Options
6. Navigating Epilepsy In Clinical Practice A Case Study Approach eBook Formats

- ePub, PDF, MOBI, and More
- Epilepsy In Clinical Practice A Case Study Approach Compatibility with Devices
- Epilepsy In Clinical Practice A Case Study Approach Enhanced eBook Features
- 7. Enhancing Your Reading Experience
 - Adjustable Fonts and Text Sizes of Epilepsy In Clinical Practice A Case Study Approach
 - Highlighting and Note-Taking Epilepsy In Clinical Practice A Case Study Approach
 - Interactive Elements Epilepsy In Clinical Practice A Case Study Approach
- 8. Staying Engaged with Epilepsy In Clinical Practice A Case Study Approach
 - Joining Online Reading Communities
 - Participating in Virtual Book Clubs
 - Following Authors and Publishers Epilepsy In Clinical Practice A Case Study Approach
- 9. Balancing eBooks and Physical Books Epilepsy In Clinical Practice A Case Study Approach
 - Benefits of a Digital Library
 - Creating a Diverse Reading Collection Epilepsy In Clinical Practice A Case Study Approach
- 10. Overcoming Reading Challenges
 - Dealing with Digital Eye Strain
 - Minimizing Distractions
 - Managing Screen Time
- 11. Cultivating a Reading Routine Epilepsy In Clinical Practice A Case Study Approach
 - Setting Reading Goals Epilepsy In Clinical Practice A Case Study Approach
 - Carving Out Dedicated Reading Time
- 12. Sourcing Reliable Information of Epilepsy In Clinical Practice A Case Study Approach
 - Fact-Checking eBook Content of Epilepsy In Clinical Practice A Case Study Approach
 - Distinguishing Credible Sources
- 13. Promoting Lifelong Learning
 - Utilizing eBooks for Skill Development
 - Exploring Educational eBooks
- 14. Embracing eBook Trends
 - Integration of Multimedia Elements
 - Interactive and Gamified eBooks

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