

Incidence and Clinical Characteristics of Healthcare Events

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Abstract

Adverse neurological events, including stroke-like syndrome, are critical concerns in healthcare due to their potential impact on patient outcomes and quality of life. Understanding the incidence and clinical characteristics of these events is crucial for early recognition, accurate diagnosis and appropriate management. This article aims to provide a comprehensive review of the incidence rates and clinical features of adverse neurological events and stroke-like syndrome, shedding light on their presentation, underlying causes and implications for patient care. Adverse neurological events encompass a broad spectrum of conditions that can result in neurological dysfunction, ranging from transient symptoms to severe neurological deficits.

Keywords: Neurological deficits • Metabolic disturbances • Cognitive impairment • Hyperglycemia

Introduction

These events can be caused by various factors, including vascular abnormalities, medication-related adverse effects, metabolic disturbances, infections, autoimmune disorders and neoplastic processes. The incidence of adverse neurological events varies depending on the specific etiology and population under study. Stroke-like syndrome refers to a constellation of clinical symptoms and signs that mimic an acute ischemic stroke but without evidence of vascular occlusion on neuroimaging. These events can be caused by diverse etiologies, including mitochondrial disorders, autoimmune encephalitis, epilepsy, vasculitis, and metabolic disturbances. Stroke-like syndrome can present with focal neurological deficits, altered consciousness, seizures, and other neurological abnormalities. Prompt recognition and differentiation from true strokes are crucial to guide appropriate management and prevent unnecessary interventions. The incidence rates of adverse neurological events vary depending on the underlying cause and population studied. For example, medication-related adverse neurological events can range from mild cognitive impairment and tremors to more severe manifestations such as extrapyramidal symptoms and drug-induced movement disorders. The incidence of these events is influenced by factors such as medication type, dosage, duration of use, and individual susceptibility. Neurological complications can also occur following surgical procedures, with varying incidence rates depending on the type of surgery and patient characteristics [1].

Literature Review

The clinical characteristics of adverse neurological events can vary widely depending on the underlying etiology. For medication-related events, symptoms may include cognitive impairments, movement disorders, tremors, mood changes, and peripheral neuropathy. In contrast, adverse neurological events associated with metabolic disturbances, such as hyponatremia or hyperglycemia, may manifest as altered mental status, seizures, or focal neurological deficits. Autoimmune encephalitis and vasculitis can present

with a wide range of neurological symptoms, including cognitive impairment, seizures, psychosis, and stroke-like episodes. Stroke-like syndrome typically presents with focal neurological deficits that mimic ischemic stroke, such as hemiparesis, aphasia, sensory deficits, and visual disturbances. However, the absence of vascular occlusion on neuroimaging distinguishes stroke-like syndrome from true ischemic strokes. The duration and progression of symptoms in stroke-like syndrome can vary, with some cases exhibiting acute-onset symptoms, while others may have subacute or fluctuating presentations. Seizures and altered mental status may also occur in association with stroke-like syndrome [2].

Discussion

Diagnosing adverse neurological events and stroke-like syndrome can be challenging due to their diverse etiologies and overlapping clinical features. Comprehensive evaluation, including detailed medical history, physical examination, neuroimaging, laboratory investigations and sometimes specialized tests, is necessary to identify the underlying cause accurately. Collaborative multidisciplinary approaches involving neurologists, radiologists, immunologists, and other specialists are often required to guide diagnosis and appropriate management. Recognizing and understanding the incidence and clinical characteristics of adverse neurological events and stroke-like syndrome is critical for providing optimal patient care. Early recognition and accurate diagnosis enable appropriate interventions, including the discontinuation or adjustment of offending medications, initiation of specific treatments for underlying causes and supportive care to address symptoms and complications. Multidisciplinary care, patient education, and long-term follow-up are essential to address the physical, cognitive, and psychosocial aspects of these conditions and optimize patient outcomes [3].

Adverse neurological events, including stroke-like syndrome, encompass a wide range of conditions with varying incidence rates and clinical presentations. Understanding the incidence and clinical characteristics of these events is crucial for timely recognition, accurate diagnosis and appropriate management. The diverse etiologies and overlapping features of adverse neurological events and stroke-like syndrome require a comprehensive evaluation involving multiple specialties. Collaborative multidisciplinary approaches, patient-centered care, and long-term follow-up are vital to optimize patient outcomes and improve quality of life. Continued research and vigilance in monitoring adverse neurological events will contribute to enhancing our understanding of these conditions and refining their management strategies. Adverse neurological events encompass a range of conditions and symptoms that can occur as a result of various factors, including medication use, underlying medical conditions, and other environmental or genetic factors. Among these events, stroke-like syndrome represents a distinct subset characterized by transient or permanent neurological deficits resembling a stroke. Understanding the

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incidence, clinical characteristics, and underlying mechanisms of these adverse neurological events is crucial for accurate diagnosis, appropriate management, and improved patient outcomes. This article aims to review the incidence and clinical characteristics of adverse neurological events, with a specific focus on stroke-like syndrome [4].

Adverse neurological events encompass a wide range of conditions and symptoms that affect the central nervous system. These events can be categorized into various subtypes, including but not limited to cognitive impairment, seizures, movement disorders, sensory disturbances, and stroke-like syndrome. Adverse neurological events can occur as a result of medication side effects, toxic exposures, metabolic disturbances, autoimmune disorders, or underlying neurological conditions. The incidence and prevalence of adverse neurological events vary depending on the underlying causes and the population being studied. Some events, such as seizures, have a relatively high incidence in the general population, while others, like stroke-like syndrome, may be rarer and occur in specific clinical contexts. Large-scale epidemiological studies and population-based registries are important sources of information to estimate the incidence and prevalence of these events accurately. The clinical presentation of adverse neurological events can be diverse and depend on the specific condition or syndrome involved. Some events, such as cognitive impairment or movement disorders, may develop gradually over time, while others, such as seizures or stroke-like syndrome, may have an acute onset. Common clinical characteristics observed in adverse neurological events include focal neurological deficits, altered consciousness, and changes in sensory perception, motor abnormalities, speech disturbances, and cognitive impairments. The specific clinical features will vary based on the underlying cause and the affected region of the nervous system [5].

Stroke-like syndrome refers to a clinical presentation characterized by transient or permanent neurological deficits that mimic a stroke but are not caused by a vascular occlusion. Stroke-like syndrome can occur in various contexts, including mitochondrial disorders, autoimmune diseases and certain toxic or metabolic conditions. The clinical presentation of stroke-like syndrome can vary widely, with symptoms such as hemiparesis, aphasia, visual field defects, and altered mental status. The duration and reversibility of these deficits may differ depending on the underlying cause and the promptness of intervention. The underlying mechanisms of adverse neurological events and stroke-like syndrome are multifactorial and often depend on the specific condition or syndrome involved. For example, mitochondrial dysfunction and impaired energy metabolism play a crucial role in the pathogenesis of stroke-like syndrome associated with mitochondrial disorders. Autoimmune-mediated processes, including inflammation and microvascular dysfunction, contribute to stroke-like syndrome observed in autoimmune disorders. Genetic mutations can disrupt various cellular processes, leading to neurological dysfunction and stroke-like symptoms. Toxic exposures, metabolic disturbances, and medication side effects can also result in adverse neurological events through various mechanisms, such as oxidative stress, excitotoxicity, or disruption of neurotransmitter systems [6].

Conclusion

Accurate diagnosis of adverse neurological events and stroke-like syndrome requires a thorough clinical evaluation, including a detailed medical history, neurological examination, neuroimaging studies and laboratory investigations. In certain cases, specialized diagnostic tests such as genetic testing, cerebrospinal fluid analysis, or neurophysiological studies may be necessary. Treatment and management strategies vary depending on the underlying cause and clinical context. In some cases, addressing the underlying cause or discontinuing the offending medication may lead to symptom resolution. Other treatment approaches may include supportive

care, symptomatic management, immunosuppressive therapy, or metabolic interventions tailored to the specific condition. Adverse neurological events, including stroke-like syndrome, represent a diverse group of conditions that can significantly impact patient health and well-being. Understanding the incidence, clinical characteristics, and underlying mechanisms of these events is crucial for accurate diagnosis and appropriate management. Further research and epidemiological studies are needed to determine the true incidence and prevalence of adverse neurological events, including stroke-like syndrome, in various populations. By increasing awareness, improving diagnostic approaches, and tailoring management strategies to the underlying cause, healthcare professionals can optimize patient care and outcomes for those affected by adverse neurological events.

Acknowledgement

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Conflict of Interest

None.

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