Complex Partial Seizure as a Manifestation of Non-Ketotic Hyperglycemia: The Needle Recovered From Haystack?

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Abstract

We present a case of a 75-year-old gentleman with undiagnosed type 2 diabetes mellitus presenting with acute onset expressive dysphasia and right hemi-paresis with no prior history of seizure. He developed clusters of stereotypical complex partial seizures which were refractory to anti-epileptic agents. He was not known to have diabetes and his brain MRI was normal. His random blood sugar measurement on admission to hospital was 30 mmol/L with HbA1c measurement of 14.8%. His seizures terminated completely when his hyperglycemia was corrected with insulin and rehydration therapy.

Keywords: Non-ketotic hyperglycemia; Seizures; MRI

Introduction

First onset seizure in a patient aged over 50 years would normally suggest an intra-cranial lesion as an initial hypothesis. However, it could also be related to an underlying metabolic disturbance such as non-ketotic hyperglycemia (NKH) seen in diabetes (generally type 2). Rapid diagnosis is essential for seizure management, as they are usually refractory to anti-epileptic agents, stopping spontaneously when the hyperglycemia is corrected [1-3]. The global prevalence of diabetes has increased significantly. For instance, diabetes mellitus (DM) affects about 8.3% of the total world population and represents a challenge for all health authorities across the globe. It is estimated that 382 million people worldwide have diabetes, and this will increase to 592 million by 2035. Importantly, it is estimated that as many as 175 million people with DM worldwide, or close to half of all people with DM, are unaware of their disease [4-6]. Therefore, in the era of diabetes epidemic, we believe it is crucial that acute and emergency medicine physicians should familiarize themselves with all diabetes complications, including neurological manifestations of NKH.

Case Report

A 75-year-old gentleman presented to the accident and emergency department (A&E) following an episode of fall at home and was found by his wife on a bathroom floor. Initial neurological examination revealed global aphasia with right-sided hemiparesis. He was noted to be incontinent of urine.

His medical history included atrial fibrillation, hypertension, benign prostatic hypertrophy and chronic obstructive pulmonary disease. His medications included warfarin, tamsulosin, seretide, felodipine and candesartan. He has no physical disability, independent prior to hospital admission. On presentation, he was hemodynamically stable with a GCS of 11/15 (E4/V1/M6). Initial blood parameters including inflammatory markers were within normal limits. Each episode of seizures lasted less than a minute. They were all self-terminating. He was commenced on phenytoin infusion. CT brain revealed an old lacunar infarct in the left basal ganglia. He was transferred to the stroke ward for further observation and treatment. It was noted he had been developing further seizure clusters despite IV phenytoin. Each seizure episode lasted between 3 and 5
min. Nursing documentation revealed at least 20 stereotypical movements within the first 24 h.

On the second day of his admission, his CRP had risen and he was commenced treatment for aspiration pneumonia. MRI brain was unremarkable with no evidence of temporal or meningeval enhancement (Fig. 1). Lumbar puncture was delayed in view of his abnormal INR. His random blood sugar measurement was recorded at 30 mmol/L with HbA1c measurement of 14.8%. He was commenced on IV insulin with subsequent reduction of seizure frequency. He was seizure free within 24 h and discharged uneventfully with oral metformin. He was reportedly still seizure free on annual review without any anti-epileptic medication.

**Discussion**

NKH is a clinical syndrome consisting of severe hyperglycemia, hyper-osmolarity and intra-cellular dehydration without keto-acidosis. Maccario et al first described focal seizures as a manifestation of hyperglycemia without keto-acidosis, in their report of seven cases, published in 1965 [1]. In the intervening years, several case series and retrospective reviews have described such a neuro-endocrine entity [2, 3]. In majority of cases, seizures reveal previously undiagnosed diabetes and they may be preceded by osmotic symptoms [4].

Neurological deficits associated with NKH vary and can include an alteration in consciousness level. Motor seizures are most often partial and may be followed by a post-ictal motor deficit. According to the literature, they often begin as partial seizures that generalize secondarily [7]. Other presentations include visual hallucinations, aphasia, visual field defect and hemisensory loss [8, 9]. States of “petit mal” are seen in the form of epilepsy partialis continua [10].

Pathophysiology for neurological manifestation of NKH remains unclear. A multi-factorial etiology has been suggested, with hyperglycemia as one of the factors likely to promote the onset of seizures. In 1973, Sing and Strobos suggested that hyperglycemia may create a hyper-osmolar gradient between the intra- and extra-cellular neuronal environments, thereby inducing intra-cellular dehydration, which induces seizures [11]. According to Ozer et al, hyperglycemia increases GABA metabolism and thereby diminishes the seizure threshold [12]. Dong Wook et al suggest reversible blood brain barrier (BBB) disruption as a cause of hyperglycemia-induced seizures in NKH patients [13]. However, Maccario concluded that serum hyper-osmolality, leading to intra-cellular dehydration and disturbance of electrolyte equilibrium of the intra- and extra-cellular compartments, is the probable etiologic factor [14].

Correction of hyperglycemia, hyper-osmolarity or electrolyte disorders is vital. Initiation of insulin therapy and rehydration are required as correction of hyperglycemia stops seizures effectively. Anti-epileptic treatment is likely to aggravate seizures. Indeed, phenytoin inhibits insulin secretion and consequently is harmful.

This case report highlights the importance of a thorough clinical evaluation in patients presenting with stroke mimics. All patients presenting to the emergency department with stroke-like symptoms should have their blood sugar measured on arrival to avoid an unnecessary delay in initiating appropriate treatment.

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