Rare Case of Hypertensive Crisis Secondary to Diencephalic Epilepsy

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Introduction

Hypertensive crisis (HC) is a clinical situation characterized by the quick, inappropriate, intense and symptomatic elevation of blood pressure, with or without risk of target-organ deterioration.\(^1\) HC has shown a decrease in incidence, most likely due to the advancement of outpatient antihypertensive therapy,\(^2\) but it still represents a challenge in emergency clinical practice. In over 80% of cases, patients already have a previous diagnosis of arterial hypertension (AH).\(^3\) However, undiagnosed AH or secondary causes may be the etiology of HC.

Case report

Female patient, 49 years old, from Angola, went to the emergency unit of a secondary hospital. She presented with lethargy and mental confusion, profuse cold sweating, elevated blood pressure levels (238 x 146 mmHg) and tachycardia (102 bpm). Cardiac and pulmonary auscultation were normal and without focal neurological signs. It was characterized as hypertensive emergency. She reported similar episodes during adolescence, with profuse cold sweating, followed by erection of hairs, but with spontaneous resolution and short duration. The patient denied the use of illicit drugs, contraceptives and hormone replacement therapy.

Lab exams such as hemogram and serum blood glucose, sodium, potassium, CK-MB, and troponin 1 came back normal. ECG (Figure 1) showed sinus tachycardia. Head CT was normal. Diagnostic investigation was complemented with a transthoracic echocardiogram, echo-doppler of renal arteries, and aortography, as well as a lab research for pheochromocytoma, Cushing’s syndrome, acromegaly, hyperaldosteronism and dosage of thyroid hormones, with all exams within the parameters of normality.

The patient was started on treatment for hypertensive emergency, and the patient was transferred to the intensive care unit (ICU) on venous nitrate, reaching a value of 162 x 104 mmHg in the fourth hour after admission.

During her stay in the infirmary, antihypertensives were suspended since the patient showed intolerance to the medications, with hypotension maintained with minimal doses (enalapril maleate 5 mg, twice a day).

During outpatient follow-up, the patient remained asymptomatic, with physical examinations persistently normal even without the use of antihypertensives.

Approximately three months after the first episode, the patient manifested the same symptoms, with new admission to the ICU, where she displayed a similar clinical behavior; that is, notably elevated levels of blood pressure responsive only to intravenous nitrate, with posterior hypotension and intolerance to low doses of antihypertensives.

Based on the clinical history of the first admission, an electroencephalogram (EEG) was performed (Figure 2) during this second hypertensive crisis, showing positive spicules (14 and 16 per second); and so the diagnosis was of diencephalic epilepsy. Plasma catecholamines were measured at the time of the crisis and 24 hours after that. The first one was elevated, and the second was normal, which corroborated the diagnosis.\(^4\) Treatment with carbamazepine was initiated,\(^4\) which yielded good therapeutic response.

The patient had a third episode nine months after the first one; however, during her stay, she admitted to having suspended the treatment with carbamazepine. Since then, she has remained asymptomatic under regular outpatient follow-up and on continuous carbamazepine.

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Discussion

Even though most cases of HC occur in previously hypertensive patients, especially those on irregular treatment, and the main etiology is idiopathic, secondary causes must always be on the assisting doctor’s radar. With the patient in question, the crises were due to a diencephalic epilepsy. It is important to highlight the importance of epilepsy in developing countries of Africa, Latin America, and Asia, where the prevalence rate is 4 to 5 times higher than in industrialized countries.

The diencephalon is the central nervous system structure formed by the thalamus, hypothalamus, epithalamus and subthalamus. In the context of this clinical case, it is worth mentioning the role of the thalamus in the emotional behavior and cortical activation, and of the hypothalamus in the regulation of the autonomous nervous system and body temperature. Diencephalic epilepsy is a clinical entity, in which discharges from the hypothalamus or from the anterior portion of the thalamus may cause tremors, erection of hairs, sensation of fear, sweating, plenitude, nausea, and dyspnea, as well as tachycardia and blood pressure elevation, among other signs of sympathetic dysfunction. EEG, in addition to the clinical evaluation and response to anticonvulsants, is an important diagnostic instrument, because it demonstrates positive spicules (14 and 16 seconds). Anxiety disorders are included in its differential diagnosis. It is worth mentioning that in a HC derived from a status epilepticus and not related to hypertensive encephalopathy, the patient may present with hypotension after the first acute hypertensive event, just like the patient in question. It is also important to mention that most cases of diencephalic epilepsy associated to severe hypertension are reported in comatose patients, victims of serious traumatic brain injury; however, the patient in question did not present a history of brain injury. Treatment with carbamazepine yields good responses, with gabapentin as a second option.

In conclusion, despite being a rare cause for HC, diencephalic epilepsy must be considered when other differential diagnoses are ruled out and the patient presents with hypotension secondary to the use of maintenance antihypertensives.
Figure 2 – EEG: positive spicules (14 and 16 per second).

Author contributions

Writing of the manuscript: Ribeiro RNF, Martins WA, Ribeiro BNF. Critical revision of the manuscript for intellectual content: Martins WA, Ribeiro BNF.

Potential Conflict of Interest

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References


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Study Association

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