Central Pontine Myelinolysis (CPM): An Atypical Presentation of HIV

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Abstract

HIV affects the Nervous System at different stages of disease ranging from central nervous system to peripheral nervous system. Nervous system manifestation of HIV is estimated to affect approximately 25-90\% of the patients. In spite of wide range of the manifestations of neurological complications Central Pontine Myelinolysis (CPM) remains an uncommon presentation of HIV. A sixty year male presented with history of sub acute onset of fever, quadriplegia and dyssarhria of two month duration. This case discusses the atypical neurological presentation in HIV patient.

Keywords: HIV; Atypical presentation; Neurological manifestation

Introduction

A wide range of nervous system involvement occurs in HIV [1]. Central Pontine Myelinolysis is an uncommon condition characterized by the clinical Syndrome of quadriplegia, pseudo bulbar palsy and symmetrical lesions in the center of basis pontis. The risk factors associated with CPM includes presence of hypoxemia, preexisting hypokalemia, malnutrition and chronic alcoholism. It is frequently associated with rapid correction of hyponatremia [2]. The clinical presentation varies between rapidly evolving spastic paraparesis with pseudo bulbar palsy, and changes in sensorium with confusion or coma. Some cases are clinically ‘silent’ and are only discovered at autopsy. CPM in patients with HIV infection is usually not recognized in life. There are few reports of CPM occurring in HIV patients with identified risk factors for developing CPM [3]. Diagnosing a case presenting with signs and symptoms of CPM in HIV patient with no identified risk factor is difficult.

Case Report

A 60 years old male presented with history of low grade fever of 2 month duration along with sub acute onset of quadriplegia and slurring of speech. He had altered sensorium of 1week duration. The patient was admitted in neurology ICU. On Physical examination he was average built. His vitals were stable pulse rate 86/min regular, BP: 130/70 mmHg, R/R 16/min, Afebrile. Cardiovascular, respiratory system and per abdomen examination was not significant. On Neurological examination: Patient was in altered sensorium, obeying commands with gestures and by making incomprehensible sounds. Cranial nerves - Fundus - Normal. He had bilateral lateral gaze paresis. Pupils were normal size reacting to light. Gag reflex was brisk .Rest other cranial nerves were normal. The muscle power was 3/5 on MRC grading in all the limbs. All the Deep tendon reflexes were exaggerated. Planters were bilaterally withdrawal.

Laboratory investigations

Hb: 10 g\%, TLC: 5700 cumm, Differential counts was normal. Random Blood -153 mg/dl, renal function test were normal. ESR: 77 mm 1st hr. Serum electrolytes- S. Na\textsuperscript{+}: 138 meq/L, Serum Sodium remained normal throughout hospital.
stay. Liver function test-Se Bilirubin 0.8 mg/dL, SGOT 24 U/L, SGPT 28 U/L, Serum cholesterol -182 mg/dL. Serum Proteins - 7.1 g/dL, Albumin 3.6 g/dL, Serum Osmolality 285 mmol/dL, CSF Examination - Cytology - Acellular, Sugar: 54 mg/dl, Protein 79 mg/dL, Chloride 113 mmol/L. All the biochemical parameters were within normal limits. CD4 count was 250/mL. HIV I and II antibodies were detected by using three different methods e.g. Elisa/ Rapid and simple test in patients' serum.

He was started on antiretroviral therapy and symptomatic treatment.

MRI Brain

Bilateral Symmetrical Hyper intense signals are seen in pons consistent with central pontine Myelinolysis (Fig. 1, 2).

Discussion

Central pontine Myelinolysis (CPM) was first described in 1959 in four patients, three of whom were alcoholic and one was severely malnourished. This clinicopathological syndrome is characterized by quadriplegia, pseudo bulbar palsy and symmetric lesion in centre of basis pontis. Extra-pontine Myelinolysis also occurs and patient can have varied presentation according to the site involved. The unusual clinical presentations include ataxia, cataplexy, neurobehavioral Syndromes and extra pyramidal involvement. CPM is usually found in sick patients with malnutrition, malignancy, burn injury, chronic alcoholism, hypoxemia and pre-existing hypoaluminaemia. HIV associated CPM had been documented in Autopsy by various authors [4]. There are few reports of CPM occurring in AIDS patients with potential risk factors for CPM development [5]. There has been no previous report of CPM presenting in as neurological manifestation in HIV patient. Our case is unique as CPM is documented without any risk or precipitating factor. There was no evidence of electrolytic imbalance in our patient. His albumin levels were normal. CPM has been described in liver transplant recipients and in severe burns cases, as well as in association with alcoholism and malnutrition. It has been thought that the rapid correction, or over correction, of hyponatremia was the crucial factor in causing CPM in most studied cases. Our patient did not have an episode of serum hyperosmolality or hypoalbuminaemia. The Diagnosis was made on clinical examination which was consistent with CPM and the neuroimaging confirmed the involvement of pons (Fig. 1). In absence of the risk factors or precipitating factors the possibility is that CPM may be thought as a primary effect of HIV itself, representing another neurological manifestation of this virus which is not well known and needs to be studied. The p24 HIV protein was detected immunocytochemically in rare perivascular macrophages in the brain stem and beyond the margin of the area involved with CPM in neuropathological studies. Further studies are required to understand the effect of HIV on the demyelinization of pons and other brain structures.

Conclusion

CPM is demyelination syndrome of the pons. In HIV patients it is usually not recognized during life and diagnosed on autopsy. Clinical presentation may vary, ranging from mutism and dysarthria to spastic quadriplegia with a pseudo bulbar palsy. Our case illustrates that CPM should be considered as the cause for neurological manifestation in patients with AIDS in the appropriate clinical setting, regardless of
presence or absence of the risk factors responsible for causing the fatal condition.

References