CASE REPORT

Acute Encephalopathy in a Child with Coronavirus Disease-2019 Infection

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ABSTRACT

Coronavirus disease-2019 (COVID-19) is a global health crisis. Children account for 1–5% of diagnosed COVID-19 cases with relatively mild presentation and less mortality compared to adults.1 While patients typically present with fever, shortness of breath, and cough, neurologic manifestations have been reported, mainly in adults. We report a case of COVID-19-associated encephalopathy in an 11-year-old child who presented with altered sensorium and seizures.

Our patient was an 11-year old boy who premorbidly had intellectual disability and a single episode of generalized seizures at the age of 4 years, but was never on antiepileptics. Details of his past history or investigations for intellectual disability were not available as the child stayed with his aunt after death of both parents (when child was 3 years old). He presented with a 2-day history of fever and loose motions 2 days prior to admission. The child developed altered mental status and had multiple episodes of generalized tonic clonic (GTC) seizures 12 hours before admission to our hospital (after being denied admission by two hospitals).

On admission to our hospital, he had severe tachypnea and active GTC seizures. The hospital protocol for pediatric status epilepticus was followed and his seizures were controlled after about 15 minutes (lorazepam, fosphenytoin, and levetiracetam were used). The child’s Glasgow coma scale on admission was 6/15 and he was immediately intubated and ventilated. He had bradycardia (heart rate 60/minute), hypertension (130/90 mm Hg), tachypnea (respiratory rate 52/minute), and bilateral mid-dilated pupils unresponsive to light. Chest auscultation did not reveal any foreign sounds. The pulse oximeter showed oxygen saturation (SpO2) of 92%. The ECG showed sinus bradycardia.

Initial laboratory workup revealed severe metabolic acidosis (pH 6.8, PaO2 68 mm Hg, PaCO2 15 mm Hg with bicarbonate of 3 mEq/L) with deranged renal functions (BUN: 136 mg/dL, creatinine 14 mg/dL) and liver function (ALT: 240 U/L, AST: 340 U/L, PT INR 2.1, bilirubin 1.4 mg/dL). There was hyperglycemia (RBS 453 mg/dL) with grossly elevated serum amylase (979 U/L); HbA1c was 4.8% indicating acute hyperglycemia probably due to pancreatitis. There was hyperonatremia (serum sodium 127 mEq/L); Hemogram revealed hemoglobin of 9.1 g/dL, platelet count of 90,000 with white cell count of 17.6 x 109, and the neutrophil lymphocyte ratio of 8.8 (neutrophil 71%, lymphocytes 9%). C reactive protein (CRP) was positive (3.6 mg/dL); however, procalcitonin was not elevated (0.4 ng/mL). Serum ferritin was high (2400 ng/mL) and D-dimers were elevated (3.5 mg/dL). Chest X-ray showed mild right parahilar and lower-zone infiltration. Creatine kinase-myocardial band (CK-MB) was elevated (99 IU/L) suggestive of myocardial injury.

SARS-CoV-2 real-time reverse transcriptase (rRT)-PCR was positive. Respiratory pathogen polymerase chain reaction (PCR) panel was negative. The cerebrospinal fluid (CSF) analysis did not show any cells. Tests for SARS-CoV-2 and standard meningitis/encephalitis PCR panel on CSF were negative. Blood and urine cultures were sterile. We could not obtain neuroimaging due to the critical condition of the patient and bedside EEG could not be done. NS-1 and rapid malaria test were negative. The HIV test was nonreactive. Ultrasound kidney ureter bladder (KUB) showed bilateral raised renarcheogenic suggestivity of acute medical renal disease.

Patient was given IV boluses of normal saline, sodium bicarbonate infusion, insulin drip (in view of hyperglycemia), antibiotic (meropenem), inotropes, steroids, enaxaparin, oseltamivir, and acyclovir. Hemodialysis was done, which reduced the creatinine to 11.5 mg/dL. However, despite all measures, child succumbed within 24 hours of admission to COVID-19 PICU.

We considered a differential diagnosis of acute necrotizing encephalopathy (ANE) due to SARS-CoV-2 infection, uremic encephalopathy, and septic encephalopathy. Additionally, possibility of status epilepticus due to an abnormal brain was...
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considered (in view of premorbid history intellectual disability and single episode of seizure in past). Acute necrotizing encephalopathy is a rare complication of influenza and other viral infections and has been related to intracranial cytokine storms, which result in blood–brain barrier breakdown, but without direct viral invasion or parainfectious demyelination. Accumulating evidence suggests that a subgroup of patients with severe COVID-19 might have a cytokine storm syndrome which our patient had. Acute necrotizing encephalopathy has been predominantly described in the pediatric population. The most characteristic imaging feature includes symmetric, multifocal lesions with invariable thalamic involvement. Unfortunately, neuroimaging could not be obtained in our patient and CSF was negative for SARS-CoV2; hence, we could not prove the diagnosis of COVID encephalopathy. However, there was a strong temporal association of encephalopathy and COVID-19 infection in our patient.

The mechanism by which COVID-19 affects the brain is not known. Direct brain infection and/or an autoimmune process may also be possible. Because the virus binds the surface spike protein to the human angiotensin-converting enzyme 2 receptor (ACE-2), and the fact that ACE-2 is present in the brain vascular endothelium, a vascular process with clotting and infarction may also be possible and this not mechanism may be possible in our patient.

Our patient had multiorgan dysfunction (encephalopathy/pancreatitis/acute kidney injury/coagulopathy/liver dysfunction/myocarditis) with evidence of cytokine storm (high ferritin, D dimer, CRP, CPK MB) and fulfilled WHO criteria of hyperinflammatory syndrome. There is one pediatric case report from India of hyperinflammatory syndrome treated successfully with tocilizumab and IVIG. In our case, delayed presentation of the child to our hospital probably contributed to the fatal outcome.

Atypical presentations of COVID-19 in children include COVID toes and hyperinflammatory syndrome. Neurological presentation with seizures is rare. This is the first reported case of COVID-19-associated encephalopathy from India in pediatric population. As the number of patients with COVID-19 increases worldwide, pediatricians should be aware of atypical presentation with altered mental status and seizures and presentation as hyperinflammatory shock with multiorgan dysfunction among children. It may be necessary to screen children presenting with encephalopathy or encephalitis for COVID-19 as infected patients require special precautions to prevent further spread.

**REFERENCES**