“Brain on Fire”: Hyperperfusion as a Hallmark of Hyperammonemic Encephalopathy

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Abstract

We describe a very rare case of acute fulminant hepatic failure as a complication of acute viral hepatitis caused by hepatitis A virus, complicated by hyperammonemic encephalopathy. The brain magnetic resonance imaging (MRI) findings were suggestive of cytotoxic edema involving bilateral cerebral hemispheres. The novel findings of hyperperfusion on arterial spin labeling perfusion MRI and hyperemic hypoxia on susceptibility weighted imaging are discussed. The patient had a rapid progression of cerebral edema and succumbed to the illness despite supportive care. Characteristic neuroimaging findings may help in the diagnosis of acute hyperammonemic encephalopathy of brain MRI, which may be useful in leading to appropriate clinical workup and diagnosis of the underlying cause of hyperammonemia. In our case, hyperammonemic encephalopathy was precipitated by fulminant hepatic failure caused by hepatitis A virus, which is a rare occurrence.

Keywords

► hepatic encephalopathy
► hyperammonemia
► hepatitis

Introduction

Fulminant acute hepatic failure (FAHF) is a rare clinical syndrome characterized by rapidly progressive hepatic dysfunction that has a very high case fatality rate of the order of 50 to 70%.1,2 Drug overdose (such as acetaminophen), viral hepatitis, alcohol abuse, and toxin exposure account for most of the cases of fulminant hepatic failure reported in the published literature. FAHF is an extremely rare complication of viral hepatitis caused by hepatitis A virus (HAV).3 HAV, a picornavirus, is spread by the fecal–oral route and usually leads to a self-limiting acute hepatitis, which confers a lifelong immunity to the infected person. FAHF has been reported in less than 1% of patients infected with HAV. Patients with FAHF develop acute hepatic encephalopathy, markedly raised serum bilirubin levels, coagulopathy, and transaminitis (of the order of >1,000 U/L). Development of acute hepatic encephalopathy has been postulated to be due to accumulation of toxins within the blood, which are normally handled by metabolism in the liver.3 Hyperammonemia has been reported to be the most implicated cause for acute hepatic encephalopathy. Hyperammonemic encephalopathy has a characteristic neuroimaging phenotype, which is distinct from hypoxic and other toxic encephalopathies and, in relevant clinical setting, may be pathognomonic of acute hyperammonemic encephalopathy.4 Herein we report of case of acute encephalopathy in which neuroimaging findings led to a diagnosis of acute hyperammonemia, which was subsequently found to be due to HAV-associated FAHF. Our case is unique as we report the findings of arterial spin labeling (ASL) magnetic resonance imaging (MRI) perfusion.

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in acute hyperammonemic encephalopathy due to HAV-associated FAHF. To the best of our knowledge, such findings have not been reported in the published literature.

**Case Description**

A 17-year-old adolescent girl was brought to the emergency department of our institution after she was found unconscious after a brief febrile illness of 2 days’ duration. There was no history suggestive of any previous comorbid illness. The bystanders did not report any seizure or trauma. There was no fecal or urinary incontinence and no tongue bite. On examination, the patient was found to be in altered sensorium with a Glasgow Coma Scale (GCS) score of 9/15. The clinical examination was remarkable for icterus. The abdominal examination was within normal limits. Review of systems did not reveal any significant abnormality. There was no history suggestive of drug overdose or toxin intake. The following vitals were recorded at presentation: heart rate of 110 beats per minute (regular), respiratory rate of 22/min (irregular), SPO2 of 93% on room air, and blood pressure of 140/90 mm Hg.

Considering the patient was in acute encephalopathy with reduced GCS, MRI of the brain was done, which revealed diffuse symmetric edema involving the cerebral cortex in both the cerebral hemispheres, manifesting as a cortical swelling and hyperintense signal on T2-weighted (T2W) image (Fig. 1a) and fluid attenuated inversion recovery (FLAIR) images (Fig. 1b). Effacement of the sulcal spaces and basal cisterns was noted (Fig. 1c). Bilateral uncal herniation was evident (Fig. 1d). Diffusion weighted images revealed cortical diffusion restriction, characterized by involvement of the bilateral insula and cingulate gyri with relative sparing of bilateral peri-Rolandic cortex and parieto-occipital regions (Fig. 2a). On apparent diffusion coefficient (ADC) maps, symmetric hypointense signal involving the cortex and juxtacortical white matter was noted corresponding to the areas of diffusion restriction (Fig. 2b). These findings suggested diffuse cytotoxic edema. Symmetric cytotoxic edema was also noted in bilateral thalami (Fig. 2c). Sparing of the cerebellum, brainstem, and basal ganglia was noted (Fig. 2d). On susceptibility weighted images (SWI), relative hyperintense signal was noted in cerebral arteries as well as cortical veins (Fig. 2e), indicating reduced cerebral oxygen extraction due to cytotoxic edema and subsequent increased concentration of oxyhemoglobin in the cortical veins. No hemorrhages were noted. ASL MRI perfusion revealed diffuse increases in perfusion in the bilateral cerebral cortices, corresponding to the areas involved on T2W, FLAIR, and DWI sequences (Fig. 3). Considering the typical neuroimaging findings and the pattern of involvement, a diagnosis of acute hyperammonemonic encephalopathy was considered. The patient was intubated and managed with intravenous (IV) mannitol and lactulose administered through a nasogastric tube.

Laboratory investigations showed total serum bilirubin level of 10.8 mg/dL, alkaline phosphatase level of 320 U/L, alanine aminotransferase (ALT) level of 1,400 U/L, and aspartate aminotransferase level of 1,700 U/L. Prothrombin time was noted to
the astrocytes. Ions lead to osmotic derangements resulting in swelling of astrocytes, the elevated levels of ammonia and ammonium active transport of quaternary ammonium ions and diffusion (normal range of 80 µmol/L). Thus, a hyperperfusion was followed by hyperperfusion. In one of the studies reporting ASL findings in hyperammonemic encephalopathy due to urea cycle disorders, initial hyperperfusion in bilateral corona radiata has also been described in cases of hyperammonemic encephalopathy in the index case, few publications have shown elevated CBF, particularly in type B hepatic encephalopathy, which results from portosystemic shunting. In our case, these findings are corroborated by the findings on SWI, which shows an increase in intracranial venous signal due to reduced utilization of oxygen by brain and thus increase in oxyhemoglobin concentration in the venous blood, a state that may be referred to as hyperemic hypoxia.

Although the neuroimaging findings were characteristic of hyperammonemic encephalopathy in the index case, few differential diagnoses need to be considered in such cases presenting with encephalopathy. Status epilepticus may result in similar findings of diffuse cytotoxic edema and hyperperfusion with hyperemic hypoxia. However, the clinical scenario excluded convulsive seizure. Still, nonconvulsive status epilepticus needs to be kept in the differential diagnosis. Hypoxic encephalopathy in adults could result in diffuse cortical cytotoxic edema; however, peri-Rolandic involvement and cerebellar and basal ganglia involvement are usually seen in hypoxic-ischemic encephalopathy. Somatoline demyelination may show changes of cytotoxic edema in pontine and extrapontine distribution; however, basal ganglia involvement is characteristic in extrapontine forms of osmotic demyelination. Also, clinical scenario of rapid correction of hyponatremia is suggestive of the diagnosis in such cases. Hypoglycemic encephalopathy in adults is suggested by cytotoxic edema involving the basal ganglia and cerebral cortex, in particular, parieto-occipital and insular involvement. In adults with hypoglycemic encephalopathy, the cerebellum and brainstem are spared. Diffusion restriction in bilateral corona radiata has also been described in hypoglycemic encephalopathy. On the other hand, hyperglycemic encephalopathy may be encountered in diabetic ketoacidosis and is characterized by diffuse cytotoxic cerebral edema. However, the diagnosis is usually evident by clinical and laboratory parameters. Posterior reversible encephalopathy syndrome (PRES) is characterized by bilaterally symmetrical edema involving the parieto-occipital
region and may involve the cerebellum and brainstem. In typical cases, no diffusion restriction is seen. Pattern of hypoperfusion is commonly encountered in PRES, although some cases of hyperperfusion have also been described.\textsuperscript{18}

**Conclusion**

We have described an exceedingly rare complication of HAV infection wherein the patient developed rapidly progressive hyperammonemic encephalopathy following FAHF. Characteristic neuroimaging findings have been described. We emphasize on diffuse hypoperfusion in the cerebral cortex in the setting of cytotoxic edema, exquisitely depicted by ASL perfusion. Also, the findings of hyperemic hypoxia on SWI reinforce the hypothesis of reduced cerebral oxygen utilization in the setting of cytotoxic edema in this case. To the best of our knowledge, such findings are novel and have not been described previously in a case of HAV-associated FAHF and hyperammonemic encephalopathy.

**Ethics Statement**

The authors declare that the manuscript conforms to the Declarations of Helsinki. Informed written consent was taken from the guardians of the patient for data acquisition and publication.

**Authors’ Contribution**

S.P. contributed to drafting of the manuscript and data acquisition. P.S. contributed to data acquisition, and approval of the final draft of the manuscript. R.G. contributed to drafting of the manuscript and approval of the final draft of the manuscript. P.B. contributed to data collection and approval of the final draft of the manuscript.

**Conflict of Interest**

None declared.

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