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5-FU induced acute toxic leukoencephalopathy: early recognition and reversibility on DWI-MRI

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INTRODUCTION

Toxic leukoencephalopathy is a progressive structural damage of white matter tracts involved in higher mental function. It is also known as toxic spongiform leukoencephalopathy.1 Clinically, it is known to be reversible on withdrawal of toxic agent. Reversibility of imaging findings have been described in correlation with clinical improvement and categorically referred to as ‘acute toxic leukoencephalopathy’.2 Diffusion Weighted Imaging-Magnetic Resonance Imaging (MRI - DWI) play a pivotal role in picking up the abnormality in acute and subacute phase.

We describe this drug induced complication in a young patient with esophageal malignancy being treated with 5-FU.

CASE REPORT

A 29 years old young male, with advanced carcinoma of esophagus, was admitted for chemotherapy as consolidated treatment. On day 4th of 5-FU infusion, the patient developed discomfort and shortness of breath. ECG showed ventricular tachycardia run and rush call was generated. Loading dose of 1500 mg Magnesium was given intravenously (I/V) over 15 minutes with infusion rate of 125 mg/kg/hour. Calcium gluconate 1000 mg I/V stat was also injected. This reverted the rhythm to sinus rhythm. In view of 5-FU induced coronary spasm patient had non-ST elevation myocardial infarction and echocardiography showed ejection fraction of 15 - 20%. Due to severe left ventricular systolic dysfunction, patient also suffered from acute kidney injury as his serum creatinine level rose from 0.9 mg/dl to 1.7 mg/dl and the creatinine clearance level was 49 ml/minute. He was, therefore, put on diuretics. Within 24 hours, the patient developed aphasia and right hemianopia with no motor weakness. EEG showed mild theta wave slowing of encephalopathy. MRI revealed bilateral symmetrical periventricular and deep white matter signal changes with diffusion restriction (Figure 1a), in keeping with acute toxic leukoencephalopathy.

His remaining chemotherapy was withheld and was given I/V methylprednisolone and levetiracetum. Patient responded well to this treatment and recovered completely from neurological symptoms in 2 - 3 days. His creatinine levels also normalized to 1.2 mg/dl from

ABSTRACT

Acute toxic leukoencephalopathy (ATL) is a rare adverse effect of 5-Fluorouracil (5-FU) chemotherapeutic agent. It is imperative for the radiologist to confidently identify the white matter changes caused by this agent in case of toxicity. This will help in early detection and appropriate management of patient, as the condition is reversible both clinically and on imaging. We report a case of a 29 years old gentleman, known case of carcinoma of esophagus who suffered from acute toxic leukoencephalopathy secondary to leukotoxic therapeutic agent 5-FU, and illustrate the reversible imaging findings of this condition on withdrawal of the inciting agent.

Key Words: 5-FU. Diffusion weighted imaging (DWI). MRI. Toxic leukoencephalopathy.
maximum of 2.5 mg/dl. His hospital stay was about 10 days and he was discharged in stable condition. Repeat MRI performed after 37 days of the initial scan revealed almost complete resolution of the DWI hyperintense signals in periventricular deep white matter with residual hyperintensity in splenium of corpus callosum (Figure 1b).

**DISCUSSION**

Toxic leukoencephalopathy typically presents with wide range of neurobehavioural symptoms including seizures, coma and death. It appears as diffuse bilateral symmetrical areas of diffusion restriction in periventricular and deep white matter and corpus callosum. There is sparing of cortex and subcortical white matter as well as basal ganglia. Exposure to variable extrinsic agents, like chemotherapeutic agents, cranial irradiation, narcotics, and environmental toxins lead to this condition.

In this patient, the cause of toxic leukoencephalopathy was induction therapy by 5-FU, which was reversible on withdrawal of drug and the imaging findings on DWI-MRI also showed reversibility on follow-up scan. Similar cases have been reported in the literature. McKinney et al. emphasized on the clinical and radiological correlation and reversibility of acute toxic leukoencephalopathy.

Several chemotherapeutic drugs are responsible to induce leukoencephalopathy including methotrexate, vincristine, ifosfamide, fluorarabine, cytarabine, 5-fluorouracil, cisplatin and the interferons. Among them, 5-FU has been frequently reported as a causative agent of leukoencephalopathy. However, the reported incidence is less than 5% and the cause is multifactorial.

5-FU readily penetrates the blood brain barrier and is a fluorine-substituted analogue of pyrimidine uracil, which blocks DNA synthesis. Dyhydropyrimidine dehydrogenase (DPD) deficiency is a risk factor for 5-FU-induced leukoencephalopathy as it is responsible for major catabolism of this drug in vivo. The exact pathophysiology of drug induced leukoencephalopathy is unknown. Past studies in vitro and in vivo has postulated that in acute phase there is myelin destruction, vacuolization; myelin swelling and macrophage infiltration resulting in restricted movement of free water. This explains the high intensity signals in DWI-MRI secondary to this cytotoxic oedema.

The imaging mimickers of ATL are posterior reversible encephalopathy syndrome (PRES) and radiation induced angiopathy. PRES can be induced by similar drugs and chemotherapeutic agents, such as cyclosporin, tacrolimus, and interferon alfa. Posterior reversible encephalopathy typically involves cortex and subcortical white matter on FLAIR in early phase and subsequently involves periventricular white matter in severe cases. Diffusion restriction occurs in minority of cases.

On the other hand, in radiation injury, there is small vessel ischaemic demyelination which in acute phase, returns hyper intense signals on FLAIR in periventricular deep white matter which is asymmetrical and lacks diffusion restriction.

Other close differentials of ATL includes carbon monoxide poisoning and inhaled opiates. They are also potentially reversible and show diffusion restriction in acute phases, however, have subtle variations on imaging, therefore, clinical history plays a pivotal role. Acute carbon monoxide poisoning affects deep white matter in early stages with diffusion restriction and later involves the deep gray matter, which is a reversible finding. Heroin induced leukoencephalopathy typically shows symmetric subcortical and periventricular white matter hyperintensity on diffusion weighted images, specifically involving the cerebral and cerebellar white matter and deep gray matter.

In conclusion, it is imperative for the radiologist to rightly identify the distribution and pattern of the abnormality to give a prompt diagnosis and alert and expedite the withdrawal of causative agent. The role of primary physician in providing appropriate clinical history cannot be denied in the same instance, as many of the toxic agents have overlapping imaging features.

**REFERENCES**

