Case Report

Cerebral venous thrombosis in ulcerative colitis

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ABSTRACT

Ulcerative colitis has been reported to show hyper coagulation leading to peripheral and rarely central thrombosis. A 35-year-old female was admitted with chief complaints of increased frequency of bloody diarrhea, abdominal pain, and weight loss for 2 months. The patient was diagnosed to have ulcerative colitis after sigmoidoscopy and biopsy and she was started on treatment. Two days later, the patient developed headache and seizures. Magnetic resonance imaging of brain showed cerebral venous thrombosis with venous infarcts. A high index of clinical suspicion is needed to diagnose this uncommon condition so that appropriate treatment can be initiated.

Key words: Cerebral venous thrombosis, irritable bowel syndrome, ulcerative colitis

Introduction

Cerebral venous thrombosis (CVT) presents with protean manifestations which may mimic several other neurological disorders and hence is occasionally misdiagnosed. Ulcerative colitis is a type of chronic inflammatory bowel disease (IBD) that affects the lining of the large intestine (colon) and rectum. The etiology is unknown. Patients with active IBD are at an increased risk of thromboembolism which includes deep vein thrombosis, pulmonary embolism, cerebral venous thrombosis, and rarely arterial thrombosis.^[1] We report a patient who presented with gastrointestinal symptoms and during the acute phase of the illness developed cerebral venous thrombosis.

A 35-year-old female was admitted with chief complaints of increased frequency of bowels and abdominal pain for 2 months. The patient had a history of significant weight loss. The patient used to have frequency of 7-8 motions with blood and mucus. She had no history of fever, night sweats, and arthritis. The patient was not on any medication nor had any other major ailment in the past. On physical examination, the patient was moderately

built and pale. She was afebrile; pulse 90/minute and blood pressure was 110/60 mm Hg. There was no lymphadenopathy. The chest was clear. Her abdomen was flat, with normal bowel sounds. There was mild tenderness diffusely but no rebound tenderness or guarding. Rectal examination showed perianal ulcers. Sigmoidoscopy revealed perianal fistula and perianal ulcers suggestive of inflammatory bowel disease. Biopsies were taken from both the affected and normal-appearing areas. H and E stain of a colonic biopsy [Figure 1a showed a crypt abscess (arrow) and goblet cell loss (arrow head) confirming ulcerative colitis. The patient was started on steroids, 5-aminosalicylic acid, and sulfasalazine. Three days later she developed headache. Headache was insidious, progressive, holocranial that gradually became severe in the next 2 days. She developed two episodes of generalized seizures and became drowsy. Fundus showed early papilledema. She had no cranial or limb weakness but had extensor plantar.

Laboratory findings showed hemoglobin 8.4 g%; the white blood cell count was slightly elevated at 11,200/mm³. Erythrocyte sedimentation rate was 25 mm in the first hour, prothrombin time (PT) of 13.4 seconds (reference range: 12.0 seconds), with an INR at 1.5, and a normal PTT at 38.4 seconds (reference range: 32-40 seconds). Liver function, renal function, serum electrolytes were within normal limits.

Noncontrast-computed tomography scan of the head was normal. With suspicion of venous thrombosis,

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magnetic resonance venogram of the brain was done. Venogram showed loss of flow signal in superior sagittal sinus [Figure 1b], left transverse sinus, and sigmoid sinus [Figure 1c]. MRI brain showed left frontal infarct [Figure 1d]. The patient was started on low molecular heparin 0.4 ml subcutaneous 12 hourly, phenytoin, intravenous dexamethasone, and antiedema measures. Her symptoms subsided and she developed no further complications. She was discharged on aspirin, sulfasalazine, steroids, and phenytoin. Her prothrombotic workup such as factor V Leiden, protein C and S, factor VIII, and antithrombin III was normal after 6 months. Antinuclear antibodies and anticardiolipin antibodies were negative and homocysteine levels were normal. She is asymptomatic and on regular follow-up.

Discussion

Cerebral venous thrombosis (CVT) is a rare but potentially devastating complication of IBD. The diagnosis needs to be considered in any patient with IBD presenting with neurological symptoms. Ulcerative colitis is a type of chronic inflammatory bowel disease (IBD) that affects the lining of the large intestine (colon) and rectum. The etiology is unknown. Ulcerative colitis may affect any age group, although there are peaks at ages 15-30 and then again at ages 50-70. A total of 40%

of adult patients have extra-intestinal manifestations in skin, joints, bones, lungs, blood, eyes, kidneys, liver, and peripheral, and central nervous system.^[2] Incidence of thrombosis is 6.5% in patients with active IBD.^[3] CVT is more common in UC than in Crohn's disease.^[4]

Various mechanisms have been postulated for thrombosis in UC which include hypercoagulation (elevated FVIII, fibrinogen, decrease in antithrombin, protein S and protein C), hypofibrinolysis [elevated PAI-1 and lipoprotein (a)], platelet abnormalities, endothelial dysfunction (increased von Willebrand factor), and immunological abnormalities (antiphospholipid antibodies).^[5]

Our patient had a negative prothrombotic workup on follow-up 6 months later. Probably in the acute phase of the illness patient may have had a hypercoagulable state. Cerebral thrombosis is a matter of high concern as it can lead to high mortality. Hence a high index of suspicion should be there in the case of headache which is increasing in severity as was present in this case. Clinical manifestations of CVT can vary from headache to major neurological manifestations. The usual presenting complaint is headache in 70-90% of cases.^[6] Patients may present with idiopathic intracranial hypertension with headache, papilledema and visual disturbances, focal neurological deficit, and headache or seizures. Seizures are a common manifestation occurring in about 40% of cases.^[7] Focal neurological deficits are also present but were not seen in our patient though her imaging showed venous infarct. Treatment of the condition is by antiedema measures and heparin.^[8] Steroids reduce the intracerebral edema and are also indicated in the active cases of ulcerative colitis. Anticoagulant in the form of low molecular weight heparin is indicated in CVT, even in cases with intracerebral bleed to dissolve the thrombus and decreases the thrombus spread.^[9] Our patient improved on treatment and is asymptomatic since last 2 years on follow-up. CVT can be fatal in IBD if not promptly diagnosed. This case report signifies the importance of considering the diagnosis of CVT in a case of IBD.

Figure 1: (a) MR venogram of the brain showing loss of flow signal in superior sagittal sinus (b) MR venogram of the brain showing loss of flow signal in left transverse sinus and sigmoid sinus (c) MRI axial sections of the brain showing hyperintense signal in the left frontal region suggestive of infarct (d) H and E stain of a colonic biopsy showing a crypt abscess (arrow) and goblet cell loss (arrow head) confirming ulcerative colitis

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