Abstract:

Introduction: Hyperhomocysteinemia is a risk factor for thrombosis and ischemia involving different organs of body, but its prevalence data from the Indian subcontinent is scarce. It usually presents with stroke or ischemic heart disease or portal and superior mesenteric vein thrombosis or deep vein thrombosis in lower limbs. Case report: We report a case of hyperhomocysteinemia in a 24 years old chronic alcoholic male, who presented with superior mesenteric thrombosis and stroke. The patient underwent segmental resection of small intestine and temporary ileostomy. He was treated with anticoagulants, vitamin supplementation and supportive care for stroke. He has recovered from hemiparesis, is doing well without any further thrombotic episodes during two years follow up. Conclusion: Young patients presenting with acute thrombotic episodes without any embolic source or atherosclerotic disease should be evaluated for hyperhomocysteinemia.

Key words: Hyperhomocysteinemia, Ileostomy, Mesenteric Veins, Stroke, Thrombosis.

Introduction

Hyperhomocysteinemia is a known risk factor for atherosclerosis, leading to venous and arterial thrombosis at an early age [1]. It can involve different organ systems and manifest as stroke or ischemic heart disease or peripheral vascular disease [2]. Hyperhomocysteinemia can be controlled by dietary supplements of vitamins B6 and B12 and folic acid, the cofactors in the metabolism of this amino acid [2]. Here we report a young patient of hyperhomocysteinemia, who presented with superior mesenteric thrombosis and stroke at the same time. Our case is the first case of hyperhomocysteinemia simultaneously involving 2 different systems.

Case Report

A 24 years old male presented with sudden onset of headache and left sided hemiparesis. He was a known alcoholic and nonsmoker. He had no history of diabetes mellitus, hypertension, hyperlipidemia or prior episode of vascular thrombosis. There was no known thrombotic disease in other family members. Magnetic resonance imaging of brain...
[Fig.1] showed subacute hemorrhagic infarct in the right temporo-parietal lobe causing mass effect and midline shift. Multifocal acute lacunar infarcts were seen in the right centrum semiovale. There was a minimal subarachnoid hemorrhage along the right fronto-parietal sulcal space with cortical vein thrombosis. Subsequently treatment was started in the line of management of stroke. After six hours of admission to hospital, he developed generalized abdominal distension with diffuse abdominal tenderness. Routine investigations showed elevated leucocyte count, neutrophilia, and mild elevated of serum creatinine. Contrast computerized tomography [Fig.2] of abdomen and pelvis showed complete thrombosis involving the distal segmental branch of superior mesenteric artery with signs of ischemic bowel of mid and terminal ileum and fluid in the abdomen. Patient underwent emergency laparotomy, which showed gangrene of entire mid and terminal ileum [Fig.3] with foul smelling hemorrhagic fluid in the abdomen. Resection of the gangrenous segment of the small intestine with proximal end ileostomy was performed [Fig.4].

A complete haemocoagulative profile comprising of protein C, protein S, antithrombin III, homocysteine, antiphospholipid antibodies and lupus anticoagulant, resistance to activated protein C, factors VII and VIII, mutation analysis of the prothrombin gene, and factor V Leiden was performed. All results were in the normal range except for the total homocysteine (HCY) level being high i.e. 174 μmol/L (normal value < 15 μmol/L). The levels of folic acid and cobalamin were low at 1 ng/mL (normal range 3-17 ng/mL) and 169 pg/mL (normal range 211-911 pg/mL), respectively. Lipid profile, ECG and cardiac echocardiography were normal. We did not check for methylene tetrahydrofolate reductase (MTHFR) gene mutations (both 677TT and 677CT) due to financial constraints. Post-operatively, the patient was put on anticoagulants, folate and vitamin B12 supplementation. He showed gradual improvement.
in motor and sensory functions in the left side of the body. On discharge his folate level improved and HCY levels decreased to 48 μmol/L. After 8 months, the side to side ileocolic anastomosis was performed to restore the bowel continuity. The patient is doing well after two years of follow up.

Discussion

Hyperhomocysteinemia is a metabolic disorder that results from abnormalities in the function of enzymes in the methionine metabolism pathway. It is a rare disorder with an incidence of the order of 1 in 2,00,000 to 10,00,000 population [3]. This can be due to congenital defect or acquired deficiency of vitamins such as folate, vitamin B6 or B12 [3]. Other less common causes of elevated HCY levels are advanced age, hypothyroidism, renal failure, lupus, leukemia, inflammatory bowel disease, liver disease, and certain drug such as oral contraceptives, theophylline, methotrexate, L-dopa, phenytoin, phenobarbital, and carbamazepine [4]. Another hypothesis is that MTHFR gene mutations (both 677TT and 677CT) raise plasma levels of HCY, which produce an increased thrombotic risk for cardiovascular and cerebral vessels. Vitamin B12 is required as a cofactor for the enzyme MTHFR, which remethylates homocysteine to form methionine [5]. Hence vitamin B12 deficiencies cause elevation HCY levels [5].

Young patients below the age of thirty with severe hyperhomocysteinemia are at a high risk of a vascular event like myocardial infarction and stroke [8]. Our case is first of its kind, where two different systems (gastrointestinal and neurological) are involved at the same time. All previous reports show either involvement of neuro or cardiac or other systems, involving only one system. This case highlights the importance of maintaining a high index.
of suspicion for any vascular thrombotic episodes in a young patient. In hyperhomocysteinemia, small doses of folic acid (1-5 mg/day) are generally efficacious. As additional therapy, low doses of pyridoxine, cyanocobalamin or choline usually normalize serum HCY levels in patients resistant to folate therapy [9]. Dietary supplementation of folic acid, vitamins B6 and B12 effectively reduces the plasma level of homocysteine and, hence, may play a role in the prevention and therapy of atherosclerotic vascular disease [10]. In our case chronic alcohol intake may also have contributed to vitamin deficiency. Abstinence from alcohol may be important to prevent nutritional deficiency in such cases.

**Conclusion**

Our case provides clinical evidence to support the association of hyperhomocysteinemia and vascular thrombosis. Young patients presenting with acute thrombosis of large vessels in the absence of an embolic source or atherosclerotic disease should be evaluated for abnormalities in HCY levels. A high index of suspicion and thorough evaluation are warranted to diagnose such rare cases.

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