A STUDY ON AETIOLOGICAL CAUSES AND CLINICAL MANIFESTATIONS OF PORTAL VEIN THROMBOSIS
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ABSTRACT

BACKGROUND
Portal Vein Thrombosis (PVT) has become an increasingly recognisable disorder during evaluation of cases of abdominal pain with usage of widespread imaging techniques. VT can result due to various clinical conditions like chronic liver disease, infections, malignancies and hypercoagulable states.

The aim of the study is to observe the clinical presentation and to do the aetiological workup of cases of PVT in a tertiary care centre.

MATERIAL AND METHODS
The study is a cross-sectional observational study done on patients having PVT who presented to Institute of Medical Gastroenterology, MMC and RGGGH, during the period of January 2016-July 2017 were taken up for the study. The clinical presentation of the above patients was observed and their aetiological workup were done.

Results
Totally, 45 cases were taken into study. 27 were males and 18 were females. Clinical presentation- The main symptoms were abdominal distension (18 patients, 51%), abdominal pain (10 patients, 27%), pain associated with diarrhoea and vomiting (5 patients, 14%), pain with nausea and anorexia (3 patients, 8%). Aetiological workup showed chronic liver disease (24 patients, 54%), prothrombotic states (9 patients, 20%), local factors, prothrombotic risks and idiopathic causes (12 patients, 26%). Detection of PVT were done mostly by portal vein Doppler (32 patients, 72%) and computed tomography (13 patients, 27%).

CONCLUSION
Higher incidence of PVT were seen among patients with chronic liver disease. Prothrombotic states like myeloproliferative disorders and coagulation defects were the next common causes detected. PVT presenting as plain abdominal pain, pain associated with nausea, vomiting and diarrhoea were seen in patients as well thereby suggesting that PVT is an important differential diagnosis in patients presenting as abdominal pain with a negative workup for common causes. With the help of widespread and improved imaging techniques, earlier diagnosis of PVT can be achieved and early intervention can greatly reduce the morbidity of patients.

KEYWORDS
Hypercoagulable State, Myeloproliferative Disorder, Portal Vein Thrombosis.


BACKGROUND
Portal Vein Thrombosis (PVT) is defined as a state of obstruction of blood flow in portal vein, which can be either partial or complete and associated with thrombus inside the vascular lumen.1 First case of portal vein thrombosis was reported in a patient with ascites, splenomegaly and varices by Balfour and Steward in 1868.2 PVT in general is a rare clinical state with a low incidence of 0.7% and with a prevalence of 3.7% among general population.3 Higher incidence and prevalence rates are seen among cirrhotic
patients with about 10% of PVT cases. Acute Portal Vein Thrombosis (PVT) is diagnosed by the presence of thrombus in PV. Chronic PVT is made out by the presence of collaterals resulting due to longstanding PVT. Patients with acute PVT present with nonspecific symptom making the diagnosis of PVT difficult. Diagnosis of PVT is done mostly using USG or CT imaging. Chronic liver disease-cirrhosis accounts for nearly 11.2%–25.2% of PVT patients. In patients with no chronic liver disease, hypercoagulable states and myeloproliferative disorders accounts for 40%–70%. Local factors like intra-abdominal surgeries, cancers and inflammatory conditions accounts for 10%–50% of PVT cases. Mortality due to PVT accounts for 2–3% in cases with no underlying liver disease or cancers. Previous few studies explains us about the clinical features and outcomes of PVT. To increase our knowledge and awareness about PVT, we hereby report our study on 45 cases of PVT who presented during January 2016-July 2017 in our centre, Institute of Medical Gastroenterology, Madras Medical College, Chennai, India.

Aim and Objectives
1. To observe the clinical presentation of the portal vein thrombosis cases presented to the centre.
2. To do the aetiological workup of cases of PVT.

MATERIAL AND METHODS
Patients who presented to our centre with various symptomatology and diagnosed to have PVT by USG-portal vein Doppler and computed tomography were taken into study. A total of 45 patients with PVT were identified, their clinical profile and aetiological workup done. The clinical presentation includes patient’s age, sex, thrombus staging acute or chronic. Acute cases identified as having thrombus in portal vein and chronic cases identified as having established collaterals, patients presenting symptoms. PVT aetiological workup and diagnostic procedures includes routine blood haemogram (leucocyte count, platelet count) blood chemistry like C-reactive protein, LDH, aspartate and alanine aminotransferase, alkaline phosphatase, serum amylase, procoagulant workup and imaging techniques.

Procoagulation screen includes the following tests-Protein C and protein S levels, antithrombin levels, factor V Leiden mutation analysis, antiphospholipid antibodies and homocysteine levels. Patients on OCPs were taken into account as well because of thrombotic property of OCPs. Bone marrow biopsy done for detecting myeloproliferative disorders.

RESULTS
The study was carried out on 45 patients of portal vein thrombosis. The following results were observed. Median age was found to be 45 (ranging from 18-72), 23 (52%) were males. 36 patients had acute portal vein thrombosis and 9 patients had chronic presentation. In patients with acute presentation, the main presenting symptom was abdominal distension in 18 (51%) abdominal pain in 10 (27%) pain with vomiting and diarrhoea in 5 (14%) pain with nausea and anorexia in 3 (8%). One patient presented with fever, 1 with hypotension and 2 with haematemesis. In patients with chronic presentation, 2 presented with mild abdominal pain, rest 7 were asymptomatic.

Diagnostic procedure observations- Blood examination results showed elevated leucocyte count in 20 (55%), increased C-reactive protein in 28 (77%), increased LDH levels in 14 (22 tested, 64%) of the 36 acute cases tested. D-dimer levels were detected in all 18 patients (100%) tested. Patients with normal serum amylase levels were ruled out of acute pancreatitis being a cause. Thrombocytosis were seen in patients with myeloproliferative disorder.

Imaging study results- Portal vein Doppler aided in detecting 32 (72%) cases of PVT. Computed tomography were used to diagnose in 12 (26%). PVT was accidentally discovered in 1 (2%) patient who underwent MRI abdomen.

Aetiological workup results- Our study on 45 patients with PVT showed the following aetiological workup observations. 24 (54%) patients had chronic liver disease, some with cirrhosis, 9 (20%) patients had prothrombotic conditions, out of which, 6 (13%) patients had myeloproliferative disorders diagnosed using bone marrow biopsy and 3 (7%) patients had procoagulant states, 2-Antiphospholipid Antibodies (APLA) and 1-protein C and protein S deficiency. Prothrombotic risks were seen in 4 (9%) patients out of which, 2 had increased homocysteine levels and 2 patients were taking OCPs. The cause for PVT could not be identified in 2 (4%) cases. Local factors were leading to PVT in 6 (13%) of cases due to factors like postsurgical complication (post-subtotal gastrectomy and splenectomy), septic cholangitis, chronic pancreatitis and Crohn's disease. In 2 (4%) cases, cause could not be identified.
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**DISCUSSION**

Our analysis on 45 patients with PVT thrombosis who presented in our centre showed that the median age of the PVT patients was 45 ranging from 18 to 72.

Clinical presentation of PVT in acute state were abdominal distension (ascites), abdominal pain, which could not be explained by clinical examination or laboratory findings. Some patients with abdominal pain had associated vomiting, diarrhoea and anorexia symptoms as well. The blood investigation results were nonspecific, still some patients had elevated leucocyte count, C-reactive protein and LDH levels. Patients who underwent D-dimer testing showed elevated levels thereby aiding in suspecting PVT in cases of abdominal pain and elevated D-dimer levels, though it is sensitive, but not specific.

Imaging studies mostly used to establish PVT was USG-portal vein Doppler and computed tomography. Patients diagnosed to have PVT by portal vein Doppler were confirmed again by CT scan. CT scan used to picture thrombus in portal and mesenteric circulation also provided a clear picture of abdominal organ examination and aided in detecting local precipitating factors for PVT; e.g., pancreaticitis.

The aetiological workup of patients with PVT showed that more than one causative factor were seen in many cases, which showed that their occurrence were similar to some previous studies done to find the cause of PVT. The causes identified falls under three major categories, chronic liver disease, prothrombotic states and local factors with a prevalence of 54%, 29% and 13%, respectively, in our study. In two cases, cause could not be identified. Prothrombotic states includes myeloproliferative disorder, procoagulant states and patients with prothrombotic risks with a contribution of 13%, 7% and 9%, respectively. Prothrombotic state included patients on oral contraceptive pills and patients with elevated homocysteine levels. The local factors leading to PVT consists of inflammatory conditions like cholangitis, pancreatitis, inflammatory bowel disease, abdominal trauma, intra-abdominal surgeries, especially post splenectomy and carcinomas of stomach. Liver and pancreas encroaching the portal vein.

**REFERENCES**


