

Role of contrast enhanced computed tomography in evaluation of vascular complications of pancreatitis: A cross-sectional study

BOTCHA NUTHAN¹, G.S.KEJRIWAL²

¹(Department of Radiodiagnosis/Maharajahs institute of medical sciences/NTRUHS/India)

²(Department of Radiodiagnosis/Maharajahs institute of medical sciences/NTRUHS/India)

Abstract:

Background: pancreas is surrounded by various blood vessels any inflammation during pancreatitis causes exogenous release of the pancreatitis amylases and lipases causing direct action on the arterial vessel walls leading to pseudoaneurysm and hemorrhage. on the other hand the inflamed pancreatic tissue and collections may compress on the adjacent veins leading to thrombosis. contrast enhanced computed tomography helps in identifying these vascular complications.

Materials and Methods: this is a cross sectional study conducted in department of Radiodiagnosis in Maharajah's institute of medical sciences, Vizianagaram, India for a duration of 1 year from 1st June 2021 to 31st may 2022. the study population consist of patients with complaints of upper abdominal pain clinically diagnosed as pancreatitis based on raised serum amylases and lipases or those patients which are diagnosed as pancreatitis on ultrasound. These patients are subjected to CT both non-enhanced and enhanced using Iohexol(omnipaque) as contrast medium. After assessing the pancreatitis and local complications, the vessels around the pancreases are examined to look for any thrombosis or aneurysmal dilatations.

Results: Of the 40 patients included, Splenic Vein Thrombosis was seen in 26(65%), followed by Portal vein thrombosis in 11 (27.5%), Superior Mesenteric Vein thrombosis in 9 (22.5%), all the three vein thrombosis was seen in 2 (5%) and arterial pseudoaneurysms was seen in 2 (5%).

Conclusion: Vascular complications in cases of pancreatitis are not uncommon and should be look in each and every case of pancreatitis. Splenic vein thrombosis is the most frequently found vascular complications in cases of pancreatitis. Timely diagnosis and management will reduce the associated morbidity and mortality.

Key Word: complication, pseudoaneurysm, venous thrombosis

Date of Submission: 06-08-2022

Date of Acceptance: 21-08-2022

I. Introduction

Pancreatitis is one of the common disease characterised by inflammation of the pancreatic tissue leading to exogenous release of the proteolytic enzymes, causing both local and systemic complications [1]. Acute pancreatitis causes both local and systemic complications leading to large changes in the clinical features and prognosis [1]. Chronic pancreatitis is an irreversible inflammatory condition of pancreas where there is gradual parenchymal thinning and fibrosis Patients with chronic pancreatitis develop an inflammatory pseudo mass, resulting infocal enlargement with fibrosis. [2]. local complications ranging from acute fluid collection to pseudocyst, when compared with the other complications the vascular complication are relatively rare but they all need to be evaluated due to potential risk. one of the most dangerous complication may be a ruptured pseudoaneurysm[3]. since majority of these complications are lifeendangering these vascular complication are need to be evaluated as early identification can be lifesaving.

As contrast enhanced computed tomography helps in identification of the both pancreatic parenchymal, extrapancreatic and vascular complications. The study is done to describe the CECT findings of vascular complications associated with pancreatitis.

II. Material And Methods

This is a cross sectional study conducted in department of Radiodiagnosis in Maharajah's institute of medical sciences, Vizianagaram, India for a duration of 1 year from 1st June 2021 to 31st may 2022. the study population consist of patients with complaints of upper abdominal pain clinically diagnosed as pancreatitis based on raised serum amylases and lipases or those patients which are diagnosed as pancreatitis on ultrasound. These patients are subjected to CT both non-enhanced and enhanced using Iohexol(omnipaque) as contrast

medium. After assessing the pancreatitis and local complications, the vessels around the pancreases are examined to look for any thrombosis or aneurysmal dilatations.

Study Design:cross sectional study

Study Location: This was a tertiary care teaching hospital based study done in Department of Radiodiagnosis, at Maharajah's institute of medical sciences, Vizianagaram, Andhra Pradesh, India.

Study Duration: 1st June 2021 to 31st May 2022.

Sample size: 40 patients.

Subjects & selection method: the study population consists of patients with complaints of upper abdominal pain clinically diagnosed as pancreatitis based on raised serum amylases and lipases or those patients which are diagnosed as pancreatitis on ultrasound. These patients are subjected to contrast enhanced computed tomography and the vessels around the pancreas are evaluated for vascular complication. Those patients with positive vascular complication are included in the study.

Inclusion criteria: those patients with pancreatitis who are having vascular complications like splenic vein thrombosis, portal vein thrombosis, superior mesenteric vein thrombosis and pseudoaneurysms of adjacent arteries with normal renal function tests.

Exclusion criteria: patients who are allergic to iodinated contrast, altered renal function test and uncooperative individuals.

Procedure methodology

As per the inclusion and exclusion 40 patients are included in the study population informed written consent was obtained from all the patients of study population, contrast enhanced computed tomography of abdomen and pelvis is performed using GE 16 SLICE CT machine, initially plain computed tomography was performed followed by contrast study. Patients are asked to stay nil by mouth prior to study for a duration of 6 hours and get their renal function tests done. After completion of the plain computed tomography contrast enhanced computed tomography is done with iodinated contrast agent iohexol given at 2ml/kg body weight. The scan field is included from domes of diaphragm to pubis. Patient lies in supine position with hands placed over the head the parameters used in image acquisition are 120kV, 300mA with 0.9sec of tube rotation time, FOV:350mm images are acquired with slice thickness of 5mm. These images obtained are reconstructed with 1mm thickness in sagittal and coronal planes. Images are obtained during arterial, venous and delayed phases.

Statistical analysis

The results are evaluated using Microsoft excel 2016, and depicted in tables and figures

III. Result

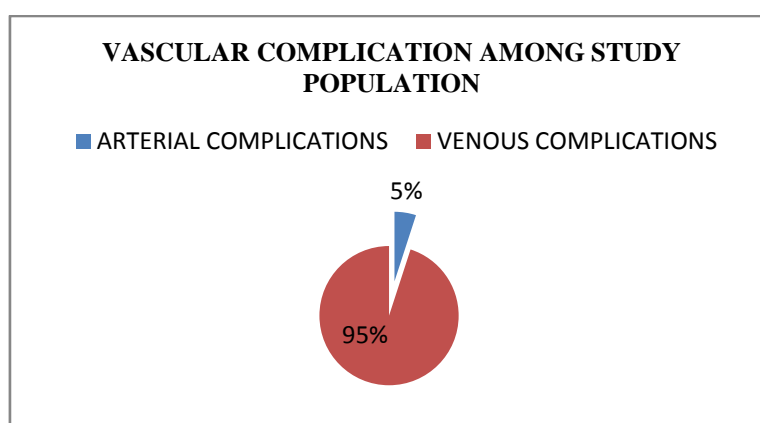
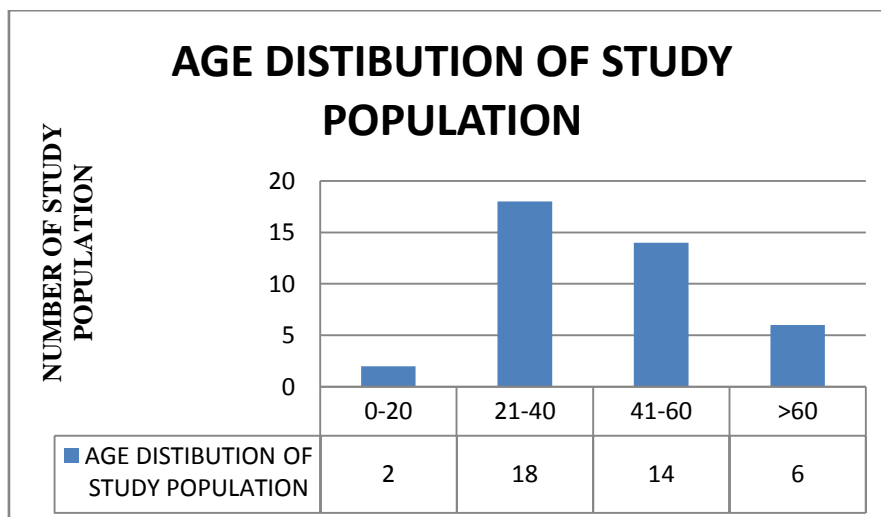
In present study we have 40 patients of pancreatitis in those vascular complications have been identified using CECT. The patients' age are ranging from 3 to 72 years. Most of the study population effected are of between 20 to 40 years of age (45%). The least effected age group is of less than 20 years (5%), among the study population the male individuals are 34 and female individuals are 6.

Splenic vein thrombosis is noted 26 patients that denotes a percentage of 65%. Portal vein thrombosis is seen in 11 patients that is about 27.5%. Superior mesenteric vein thrombosis is seen in 9 patients that is about 22.5%, all three veins thrombosis is seen in 2 patients that is about 5%. Among all the following the spleen vein thrombosis is the most commoner followed by the portal vein thrombosis.

Only 2 cases of the pseudo aneurysms have been noticed in the study one is arising from the splenic artery whereas the other is arising from the gastroduodenal artery

AGE (YEARS)	NUMBER	PERCENTAGE
0-20	2	5
21-40	18	45
41-60	14	35
>60	6	15
TOTAL	40	100

Table 1 showing age wise distribution of study population



Vascular complications	Complication findings	Number of cases	Percentage
Arterial complications	Pseudoaneurysms	2	5
	Splenic artery pseudoaneurysm	1	2.5
	Gastroduodenal artery pseudoaneurysms	1	2.5
Venous complications	Splenic vein thrombosis	26	65
	Portal vein thrombosis	11	27.5
	Superior mesenteric vein thrombosis	9	22.5
	Splenic vein +portal vein thrombosis	2	5
	Splenic vein +superior mesenteric vein thrombosis	3	7.5
	Portal vein +superior mesenteric vein thrombosis	1	2.5
	Splenic + portal vein +superior mesenteric vein thrombosis	2	5

TABLE2 showing vascular complications

IV. Discussion

In the present study of 40 cases of pancreatitis with vascular complications among the vascular complications the venous complications(95%) are most commoner than arterial complications (5%). The most commonly effected age group in the present study is 21-40 years(45%).the most common complication is splenic vein thrombosis which is seen in 26 patients (65%) followed by portal vein thrombosis which is seen in 11 patients (27.5%).superior mesenteric vein thrombosis is seen in 9 patients(22.5%).only two cases of pseudoaneurysm is seen one in the splenic artery and the other in the gastroduodenal artery. The present study

findings are similar to that of study conducted by Sanjay D and Magdum, in their study population of 47 cases of pancreatitis, the vascular complications are seen in 17 cases, out of the 17 cases 11 cases are of splenic vein thrombosis (64.7%) and 2 cases are of portal vein thrombosis. (6)

Vascular complications are not so common in cases of pancreatitis but these are around seen in 25% cases of pancreatitis.

One of the most common causes of vessel injury is severe pancreatic inflammation and necrosis, which results in the local release of exocrine proteolytic and lipolytic enzymes. These cause erosion of the vessel wall, leading to formation of a pseudoaneurysm, a hematoma (when the pseudoaneurysm is thrombosed or there is no active extravasation), or intraperitoneal hemorrhage due to rupture of pseudoaneurysm (13).

As the pancreas is in close relation to splenic vein, portal vein and superior mesenteric vein any inflammation around the pancreas like pseudocyst/acute fluid collection lead to adjacent compression of these vessels further leading to thrombosis. Splenic vein thrombosis is quite common in those cases of pancreatitis with pseudocyst and collections [13]. Portal vein rupture and pancreaticopotal fistula are very rare. [14]

Pseudoaneurysms are very rare about 3.5-10% in patients with pancreatitis (18) splenic artery being the most common artery involved in pseudoaneurysm (40%) followed by gastroduodenal (30%), pancreaticoduodenal (20%), gastric (5%), hepatic (2%), and other rarely involved are superior mesenteric, jejunal, ileocaecal, aorta which constituted for 1-3% [9,19-23]

Rupture of these pseudoaneurysms is a fatal complication. The pseudoaneurysm may rupture into the pseudocyst, GI tract, peritoneum, or pancreatic parenchyma. [24,25] There is an entity called "Haemosuccus pancreatitis" where the pseudoaneurysm ruptures into the pancreatic duct and bleeds from ampulla of Vater which is seen in ERCP. Another entity called as "Pseudoaneurysmatic pseudocyst" where an artery is involved within the pseudocyst. [28]. A rare case of rupture of gastroduodenal artery pseudoaneurysm into superior mesenteric vein was reported by Suzuki T et al. [29]. Rupture of the pseudoaneurysm is not determined by the size. [30] although pseudoaneurysms are rare those with potential rupture need to be treated immediately as undetected rupture can be life threatening. Endovascular embolisation is the best first-line treatment, and if that fails, emergency surgery should be performed as soon as possible [31,32].

Angiography is considered the gold standard, but Computerised Tomography (CT) scans are usually used for these patients as angiography is not always available.

The presence of contrast enhancement within or adjacent to a suspected pseudocyst on a CT scan is suggestive of a pseudoaneurysm.

On plain scan, increased attenuation within a fluid collection may indicate a recent hemorrhage. Contrast enhanced CT plays an important role in detecting these vascular complications of pancreatitis [33] and helps in identifying the cause of pancreatitis [34].

A more recent study of 127 patients over two years by Gonzelez HJ et al. found 20 patients with splenic vein thrombosis. They proposed that venous thrombosis in patients of pancreatitis is the common finding and due to overlap of peripancreatic collections and splenic vein thrombosis strongly advocated the mechanism of thrombosis to be compression and perivascular inflammation [37].

LIMITATIONS: This study did not aim to determine the intervention and management of the condition, and there was no follow-up to determine the consequences of these complications. Furthermore, because the study was conducted in a single institute, it was limited to a smaller number of patients.



Figure 1 Contrast enhanced CT showing thrombosis in splenic vein extending upto spleno-portal confluence

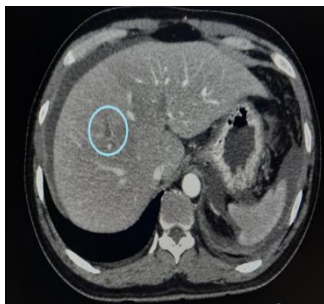


Figure 2 Contrast enhanced CT showing thrombosis in portal vein division

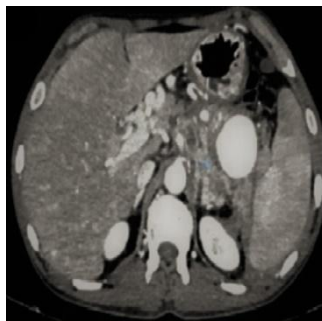


Figure 3 Contrast enhanced CT showing pseudoaneurysm of splenic artery in case of chronic pancreatitis

V. Conclusion

Vascular complications in cases of pancreatitis are not uncommon and should be look in each and every case of pancreatitis. Splenic vein thrombosis is the most frequently found vascular complications in cases of pancreatitis. Timely diagnosis and management will reduce the associated morbidity and mortality.

References

- [1]. Lenhart DK, Balthazar EJ. MDCT of acute mild (necrotizing) pancreatitis: abdominal complications and fate of fluid collections. *American Journal of Roentgenology*. 2008 Mar;190(3):643-9.
- [2]. Balthazar EJ. Acute pancreatitis: assessment of severity with clinical and CT evaluation. *Radiology*. 2002 Jun;223(3):603-13.
- [3]. Sharma PK, Madan K, Garg PK. Hemorrhage in acute pancreatitis: should gastrointestinal bleeding be considered an organ failure?. *Pancreas*. 2008 Mar 1;36(2):141-5.
- [4]. Masud R, Waheed A, Aurangzeb, Malik FI, Mian A. Role of radiology in pancreatic disorders, Pak. *Armed Forces Med J*. 2009;59(1):103-08.
- [5]. Quinlan JD. Acute pancreatitis. *Am Fam Physician*. 2014;90(9):632-39.
- [6]. Sanjay D, Magdum A. MDCT evaluation of acute pancreatitis. *JMSCR*. 2017;05(08):26894-97.
- [7]. Mallick IH, Winslet MC. Vascular complications of pancreatitis. *JOP*. 2004;5:328-37.
- [8]. Bernades P, Baetz A, Levy P, Belghiti J, Menu Y, Fekete F. Splenic and portal venous obstruction in chronic pancreatitis. A prospective longitudinal study of a medical-surgical series of 266 patients. *Dig Dis Sci*. 1992;37(3):340-46.
- [9]. Boudghene F, L'Hermine C, Bigot JM. Arterial complications of pancreatitis: Diagnostic and therapeutic aspects in 104 cases. *J VascIntervRadiol*. 1993;4:551-58. doi: 10.1007/BF01307725.
- [10]. De Rosa A, Gomez D, Pollock JG, Bungay P, De Nunzio M, Hall RI, et al. The radiological management of pseudoaneurysms complicating pancreatitis. *JOP*. 2012;13(6):660-66.
- [11]. Vogelzang RL, Gore RM, Anschuetz SL, Blei AT. Thrombosis of the splanchnic veins: CT diagnosis. *AJR Am J Roentgenol*. 1988;150(1):93-96.
- [12]. Hourani S Al, Al-Bdour MN, Rashaideh MA, Al-Nawayseh KR, Alasmara AA, Alkawasbeh MA, et al. Pseudo aneurysm complicates pancreatic pseudo cyst: Importance of early detection and management. *J Surg Pak (International)*. 2008;13: PD09-10: doi: 10.7860/JCDR/2017/25314.9651.
- [13]. Flati G, Andr n-Sandberg A, La Pinta M, Porowska B, Carboni M. Potentially fatal bleeding in acute pancreatitis: Pathophysiology, prevention, and treatment. *Pancreas*. 2003;26(1):08-14.
- [14]. Charvat F, Maskova J, Belina F, Buric I, Lacman J, Fuksa Z, et al. Portal vein erosion: A rare hemorrhagic complication of acute pancreatitis treated by percutaneous stent-graft placement. *J VascIntervRadiol*. 2010;21(3):411-12.
- [15]. Rattner DW, Warshaw AL. Venous, biliary, and duodenal obstruction in chronic pancreatitis. *Hepatogastroenterology*. 1990;37(3):301-06.
- [16]. Belli AM, Jennings CM, Nakielny RA. Splenic and portal venous thrombosis: A vascular complication of pancreatic disease demonstrated on computed tomography. *ClinRadiol*. 1990;41(1):13-16.
- [17]. Mam CS, Edgar KA, Francis IR. CT diagnosis of splenic vein occlusion: Imaging features, etiology and clinical manifestations. *Abdom Imaging*. 1995;20:78-81. [PMID 7894307]
- [18]. White AF, Baum S, Buranasiri S. Aneurysms secondary to pancreatitis. *AJR Am J Roentgenol*. 1976;127(3):393-96.
- [19]. Woods MS, Traverso LW, Kozarek RA, Brandabur J, Hauptmann E. Successful treatment of bleeding pseudoaneurysms of chronic pancreatitis. *Pancreas*. 1995;10:22-30. [PMID 7899456].
- [20]. Suzuki T, Ishida H, Komatsuda T, Oyake J, Miyauchi T, Heianna J, et al. Pseudoaneurysm of the gastroduodenal artery ruptured into the superior mesenteric vein in a patient with chronic pancreatitis. *J Clin Ultrasound*. 2003;31:278-82. [PMID 12767023].

- [21]. Toscano RL, Ruiz OR, Gerace CA Jr. Rupture of splenic artery pseudoaneurysm. *Am Surg.* 1995;61:940-42. [PMID 7486420].
- [22]. Yeh TS, Jan YY, Jeng LB, Hwang TL, Wang CS, Chen MF. Massive extraenteric gastrointestinal hemorrhage secondary to splanchnic artery aneurysms. *Hepatogastroenterology.* 1997;44:1152-56. [PMID 9261616].
- [23]. Giles RA, Pevec WC. Aortic pseudoaneurysm secondary to pancreatitis. *J Vasc Surg.* 2000;31:1056-59. [PMID 10805901].
- [24]. Dinu F, Deviere J, Van Gossum A, Golzarian J, Dussaussois L, Delhaye M, et al. The wirsungorrhagies: Causes and management in 14 patients. *Endoscopy.* 1998;30:595-600. [PMID 9826136].
- [25]. Kuganeswaran E, Smith OJ, Goldman ML, Clarkston WK. Hemosuccuspancreaticus: Rare complication of chronic pancreatitis. *GastrointestEndosc.* 2000;51:464-65. [PMID 10744821].
- [26]. Wagner WH, Cossman DV, Treiman RL, Foran RF, Levin PM, Cohen JL. Hemosuccuspancreaticus from intraductal rupture of a primary splenic artery aneurysm. *J Vasc Surg.* 1994;19:158-64.
- [27]. Igari K, Ochiai T, Aihara A, Kumagai Y, Iida M, Yamazaki S. Hemosuccuspancreaticus caused by a primary splenic artery aneurysm as a rare cause of gastrointestinal bleeding: Report of a case. *Int Surg.* 2010;5:325-28.
- [28]. Masatsugu T, Yamaguchi K, Yokohata K, Mizumoto K, Chijiwa K, Tanaka M. Hemorrhagicpseudocyst and pseudocyst with pseudoaneurysm successfully treated by pancreatectomy: Report of three cases. *J HepatobiliaryPancreat Surg.* 2000;7(4):432-37.
- [29]. Suzuki T, Ishida H, Komatsuda T, Oyake J, Miyauchi T, Heianna J, et al. Pseudoaneurysm of the gastroduodenal artery ruptured into the superior mesenteric vein in a patient with chronic pancreatitis. *J Clin Ultrasound.* 2003;31:278-82. [PMID 12767023].
- [30]. Tessier DJ, Stone WM, Fowl RJ, Abbas MA, Andrews JC, Bower TC, et al. Clinical features and management of splenic artery pseudoaneurysm: Case series and cumulative review of literature. *J Vasc Surg.* 2003;38:969-74. [PMID 14603202].
- [31]. De Rosa A, Gomez D, Pollock JG, Bungay P, De Nunzio M, Hall RI, et al. The radiological management of pseudoaneurysms complicating pancreatitis. *JOP.* 2012;13(6):660-66.
- [32]. Varshney P, Songra B, Mathur S, Gothwal S, Malik P, Rathi M, et al. Splenic artery pseudoaneurysm presenting as massive hematemesis: A diagnostic dilemma case. *Case Rep Surg.* 2014;2014:501937.
- [33]. Balthazar EJ, Fisher LA. Hemorrhagic complications of pancreatitis: Radiologic evaluation with emphasis on CT imaging. *Pancreatology.* 2001;1:306-13. [PMID 12120209].
- [34]. Chandna P, Yadav RK. Importance of MDCT in Evaluation of Pancreatitis. *IOSR Journal of Dental and Medical Sciences. (IOSR-JDMS).* 2019;18(4):74-83.
- [35]. Di Cesare E, Di Sibio A, Gennarelli A, Felli V, Vellucci V, Casazza I, et al. An unusual case of ascending pancreatitis with mediastinal involvement: A case report with CT and MRI findings. *Case Rep Radiol.* 2014;2014:925105.
- [36]. Heider TR, Azeem S, Galanko JA, Behrns KE. The natural history of pancreatitis-induced splenic vein thrombosis. *Ann Surg.* 2004;239(6):876-72.
- [37]. Gonzelez HJ, Sahay SJ, Samadi B, Davidson BR, Rahman SH. Splanchnic vein thrombosis in severe acute pancreatitis: A 2-year, single-institution experience. *HPB (Oxford).* 2011;13:860-64.

BOTCHA NUTHAN, et. al. "Role of contrast enhanced computed tomography in evaluation of vascular complications of pancreatitis: A cross-sectional study." *IOSR Journal of Dental and Medical Sciences (IOSR-JDMS)*, 21(08), 2022, pp. 11-16.