

CASE REPORT

PEER REVIEWED | OPEN ACCESS

Cholecystitis causing rupture of the cystic artery and massive hemoperitoneum

Rowan Klein Nulend, Andrew Tse, Timothy Pollitt

ABSTRACT

Introduction: Cholecystitis is a common surgical pathology; however, it is uncommonly associated with hemorrhage. Cases of hemoperitoneum secondary to cholecystitis are usually in the context of cystic artery pseudoaneurysms; however, no reports describe rupture of a non-aneurysmal cystic artery.

Case Report: We present a case of hemoperitoneum and hemorrhagic shock secondary to cholecystitis, with resultant rupture of a branch of the cystic artery, in the absence of a pseudoaneurysm. The patient was a middle-aged man who presented with severe right upper quadrant pain on a background of biliary colic. Imaging was consistent with cholecystitis, and significant volume free fluid in the right upper quadrant. The patient became shocked and was taken for laparoscopy, with conversion to laparotomy. 1.6 liters of hemoperitoneum was evacuated, and a cholecystectomy was performed. It appeared that a gallbladder perforation has caused rupture of a branch of the cystic artery, with resultant intraperitoneal hemorrhage. No pseudoaneurysm was identified. The patient made a complete recovery.

Conclusion: This was a previously unreported, yet life-threatening complication of cholecystitis. This report aims to present a novel case and promote awareness of this dangerous complication.

Keywords: Cholecystitis, Cystic artery rupture, Hemoperitoneum

How to cite this article

Klein Nulend R, Tse A, Pollitt T. Cholecystitis causing rupture of the cystic artery and massive hemoperitoneum. J Case Rep Images Surg 2024;10(2):1–4.

Article ID: 100139Z12RN2024

doi: 10.5348/100139Z12RN2024CR

INTRODUCTION

Despite cholecystitis being a common intra-abdominal pathology, hemorrhage secondary to cholecystitis is rare. The above case describes a case of cholecystitis complicated by massive hemoperitoneum secondary to cystic artery rupture, which is extremely rare. Cystic artery rupture most commonly occurs secondary to a cystic artery pseudoaneurysm (CAPA), which was not identified in this case.

CASE REPORT

A middle-aged male presented to the emergency department with a 9-hour history of severe right sided abdominal pain and nausea. This was on a background of recurrent episodes of biliary colic. He had a previous medical history of bipolar disorder and 30 pack year history of smoking. Examination revealed a tender right upper quadrant with peritonism. Blood tests are seen in Table 1.

Table 1: Preoperative blood results

Hemoglobin (g/L)	167
Bilirubin (umol/L)	19
ALP (U/L)	67
GGT (U/L)	446
ALT (U/L)	169
AST (U/L)	126

Abbreviations: ALP: Alkaline phosphatase; GGT: Gamma-glutamyl transferase; ALT: Alanine transaminase; AST: Aspartate transaminase.

Rowan Klein Nulend¹, Andrew Tse², Timothy Pollitt²

Affiliations: ¹Department of Surgery, Westmead Hospital, Westmead, NSW, Australia; ²Department of Surgery, Port Macquarie Base Hospital, Port Macquarie, NSW, Australia.

Corresponding Author: Rowan Klein Nulend, Westmead Hospital, Cnr Hawkesbury Rd and Darcy Rd, Westmead, NSW 2145, Australia; Email: rowankleinnulend@hotmail.com

Received: 26 May 2024

Accepted: 03 July 2024

Published: 14 August 2024

Bedside ultrasound showed a thickened gallbladder, with features suggestive of localized perforation. On computed tomography (CT) imaging (Figures 1 and 2) he was found to have extensive intraperitoneal free fluid, primarily surrounding the liver and subhepatic space. Hyperdense material was visualized within the gallbladder, which was consistent with hemocholecyst. Linear hyperdensity traversing the gallbladder and extending along the falciform ligament, and the liver surface implied active extravasation. There was no evidence of a cystic artery pseudoaneurysm on either imaging modality.

The patient was commenced on intravenous ceftriaxone and metronidazole. Shortly after CT, the patient became shocked with a blood pressure of 88/70 mmHg. He was urgently taken for laparoscopy, with conversion to laparotomy. Intraoperative findings were approximately 1.6 L of hemoperitoneum, and free gallstones in the right upper quadrant. The source of hemorrhage was at Hartmann's pouch, where a gallbladder perforation appeared to have ruptured a branch of the cystic artery. There was no evidence of a cystic artery pseudoaneurysm. The hemoperitoneum was evacuated, and a cholecystectomy was performed. Intraoperative cholangiography was not done at this stage. Gallbladder histopathology was consistent acute-on-chronic hemorrhagic cholecystitis.

The patient spent one night in intensive care unit (ICU) for postoperative monitoring before being transferred to the general surgical ward.

Subsequently, the patient improved clinically with normalizing biochemical and hematological markers. The patient was discharged on the fifth postoperative day. The patient was followed up and made an expectant full recovery.

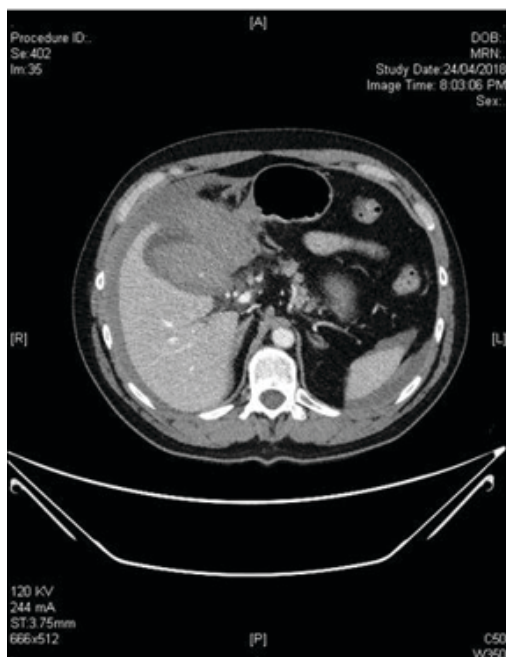


Figure 1: Axial preoperative CT abdomen.



Figure 2: Coronal preoperative CT abdomen.

DISCUSSION

Cystic artery pseudoaneurysms are extremely rare, most commonly occurring secondary to biliary instrumentation [1]. Although cholecystitis is a common surgical pathology, CAPAs have very rarely been described in cholecystitis. Clinical features of CAPA rupture remain unclear; however, it is hypothesized that in these instances, chronic cholecystitis can cause a CAPA, and that a superimposed acute inflammatory episode causes this pseudoaneurysm to rupture [2]. Cystic artery pseudoaneurysms have been associated with hemorrhagic cholecystitis resulting in hemobilia and upper gastrointestinal bleeds (UGIB), with some cases occurring with (supra)therapeutic antiplatelet/anticoagulation [3–14]. Four cases of gastrointestinal bleed secondary to CAPA have been described with concurrent cholecysto-intestinal fistulas [4, 15–17]. An even smaller number of CAPAs leading to hemoperitoneum have been described, again with some in the context of therapeutic antiplatelet/anticoagulation therapy [2, 18–21].

We found one previous case of massive hemoperitoneum secondary to cystic artery rupture in the absence of any aneurysm, secondary to acute cholecystitis in a patient with ulcerative colitis and primary sclerosing cholangitis [22]. This case was diagnosed through extravasation of contrast during a CT abdomen with contrast and was treated with angio-embolization followed by urgent laparotomy and cholecystectomy.

Two cases of cystic artery rupture and hemoperitoneum have been described without cholecystitis in patients with significant vascular risk factors. One case of hemoperitoneum due to spontaneous cystic artery rupture in the absence of cholecystitis or underlying arterial

pathology has been described in a patient with severe cardiopulmonary disease [23]. One case of spontaneous fatal cystic artery rupture has been described in a patient with vascular Ehlers–Danlos syndrome and multiple previous vascular anomalies [24].

Transhepatic gallbladder perforation has also been identified as a cause of hemoperitoneum secondary to acute cholecystitis, with subsequent hemorrhage from the liver bed [25–28]. Further cases of hemoperitoneum secondary to cholecystitis have been described with a range of related underlying pathologies, including coagulopathy (and therapeutic anticoagulation), cirrhosis, chronic primary sclerosing cholangitis, and rupture of a hemorrhagic cholecystitis [29, 30].

The case we present features a hemoperitoneum due to the rupture of a branch of the cystic artery secondary to acute perforated cholecystitis, in an otherwise healthy patient. We identified only one similar case, in a patient with complex autoimmune gastrointestinal/biliary pathology. Other cases of cystic artery hemorrhage are typically associated with bleeding from (pseudo) aneurysms. A clinical picture of acute cholecystitis and hypovolemic shock, alongside imaging findings consistent with hemoperitoneum were suggestive of such hemorrhage, which was confirmed intraoperatively. Due to appropriate investigations and timely intervention, the patient was able to make a complete recovery.

CONCLUSION

A clinical picture of acute cholecystitis and hypovolemic shock, alongside imaging findings consistent with hemoperitoneum was suggestive of such hemorrhage, which was confirmed intraoperatively. Due to appropriate investigations and timely intervention, the patient was able to make a complete recovery.

REFERENCES

1. Loizides S, Ali A, Newton R, Singh KK. Laparoscopic management of a cystic artery pseudoaneurysm in a patient with calculus cholecystitis. *Int J Surg Case Rep* 2015;14:182–5.
2. Fujimoto Y, Tomimaru Y, Hatano H, et al. Ruptured cystic artery pseudoaneurysm successfully treated with urgent cholecystectomy: A case report and literature review. *Am J Case Rep* 2018;19:187–93.
3. Gapp J, Mukherjee T, Gross J, Sirineni G, Abdalla A. Ruptured cystic artery pseudoaneurysm as a cause of cholestasis: 1344. *American Journal of Gastroenterology* 2018;113.
4. Carey F, Rault M, Crawford M, Lewis M, Tan K. Case report: Cystic artery pseudoaneurysm presenting as a massive per rectum bleed treated with percutaneous coil embolization. *CVIR Endovasc* 2020;3(1):8.
5. Nakajima M, Hoshino H, Hayashi E, et al. Pseudoaneurysm of the cystic artery associated with

- upper gastrointestinal bleeding. *J Gastroenterol* 1996;31(5):750–4.
6. Saluja SS, Ray S, Gulati MS, Pal S, Sahni P, Chattopadhyay TK. Acute cholecystitis with massive upper gastrointestinal bleed: A case report and review of the literature. *BMC Gastroenterol* 2007;7:12.
7. Tanaka T, Takakura K, Maruyama Y, et al. Hemobilia derived from cystic artery pseudoaneurysm. *Case Rep Gastroenterol* 2019;13(1):89–94.
8. Delgadillo X, Berney T, de Perrot M, Didier D, Morel P. Successful treatment of a pseudoaneurysm of the cystic artery with microcoil embolization. *J Vasc Interv Radiol* 1999;10(6):789–92.
9. Parathithasan N, Croagh D. Cystic artery pseudoaneurysm: Over warfarinisation and management. *Journal of Case Reports* 2013;3:308–12.
10. Rossini M, Bonati E, Cozzani F, Marcato C, Del Rio P. Hemobilia due to cystic artery pseudoaneurysm following cholecystectomy: Diagnosis and management, a case report. *Acta Biomed* 2019;90(4):595–8.
11. Arata R, Yanagawa S, Miyata Y, Ishitobi T, Kodama S, Sumimoto K. Hemobilia after laparoscopic cholecystectomy that was successfully treated conservatively: Case report. *Int J Surg Case Rep* 2020;77:307–10.
12. Anand U, Thakur SK, Kumar S, Jha A, Prakash V. Idiopathic cystic artery aneurysm complicated with hemobilia. *Ann Gastroenterol* 2011;24(2):134–6.
13. Al'aref SJ, Abdel-Rahman H, Hussain N. Idiopathic cystic artery aneurysm complicated with hemobilia and acute pancreatitis. *Hepatobiliary Pancreat Dis Int* 2008;7(5):547–50.
14. Totolici BD, Neamțu C, Bodog FD, et al. Hemobilia by idiopathic aneurysm of cystic artery, fistulized in the biliary ways – Clinical case. *Rom J Morphol Embryol* 2017;58(1):267–70.
15. Kim D, Kim T, Kim C, et al. A Ruptured cystic artery pseudoaneurysm with concurrent cholecystoduodenal fistula: A case report and literature review. *The Korean Journal of Helicobacter and Upper Gastrointestinal Research* 2018;18:135.
16. Aggarwal A, Goyal S, Choudhary J, Mishra H, Baberwal MC. Cystic artery pseudoaneurysm with concurrent cholecysto-enteric fistula: A rare case report. *Journal of Evolution of Medical and Dental Sciences* 2015;4(2):273–7.
17. Glaysher MA, Cruttenden-Wood D, Szentpali K. A rare cause of upper gastrointestinal haemorrhage: Ruptured cystic artery pseudoaneurysm with concurrent cholecystojejunal fistula. *Int J Surg Case Rep* 2014;5(1):1–4.
18. Nguyen D, Goodwin JS, Bhowmik N, Boiteau G, Potts J. Acute hemorrhagic cholecystitis with large hemoperitoneum: Treatment with microcoil embolization and subsequent cholecystectomy. *J Radiol Case Rep* 2021;15(2):25–34.
19. Liang X, Lü JM, Meng N, Jin RA, Cai XJ. Hemorrhagic shock caused by rupture of cystic artery pseudoaneurysm secondary to calculous cholecystitis. *Chin Med J (Engl)* 2013;126(23):4590–1.

20. Ghooz A, Kheir E, Kotru A, et al. Hemoperitoneum secondary to rupture of cystic artery pseudoaneurysm. *Hepatobiliary Pancreat Dis Int* 2007;6(3):321–3.
21. Praveen Kumar Sunkara PRV, Shah PK, Rakshit K, Choudhary SR, Bohidar NP, Dubey SK. Rupture of cystic artery pseudoaneurysm: A rare complication of acute cholecystitis. *Indian J Surg* 2018;80(1):87–9.
22. Aljiffry MM, Almulhim AN, Jamal MH, Hassanain MM. Acute cholecystitis presenting with massive intra-abdominal haemorrhage. *J Surg Case Rep* 2014;2014(4):rju019.
23. Ouazzani A, Bataille D, Boutkhil A, Guérin E, Lefebvre JC, Vaneukem P. Spontaneous cystic artery rupture: A rare cause of haemoperitoneum. *Acta Chir Belg* 2009;109(1):106–8.
24. Febres-Aldana CA, Castellano-Sanchez AA, Alexis J. Spontaneous perforation of small intestine followed by rupture of the cystic artery: The natural history of Vascular Ehlers-Danlos Syndrome. *Autops Case Rep* 2019;9(1):e2018054.
25. Nural MS, Bakan S, Bayrak IK, Baydin A, Danaci M. A rare complication of acute cholecystitis: Transhepatic perforation associated with massive intraperitoneal hemorrhage. *Emerg Radiol* 2007;14(6):439–41.
26. Kolder D, Geiger T, Tharakan AK, Kessel JW, Awad ZT. Massive hemoperitoneum from transhepatic perforation of the gallbladder. *Mt Sinai J Med* 2006;73(8):1135–6.
27. Syme RG, Thomas EJ. Massive hemoperitoneum from transhepatic perforation of the gallbladder: A rare complication of cholelithiasis. *Surgery* 1989;105(4):556–9.
28. Refsum SE, Wilson RH, Blake G. Haemoperitoneum following gallbladder necrosis. *Ulster Med J* 1992;61(2):207–8.
29. Chiapponi C, Wirth S, Siebeck M. Acute gallbladder perforation with gallstones spillage in a cirrhotic patient. *World J Emerg Surg* 2010;5:11.
30. Mechera R, Graf L, Oertli D, Viehl CT. Gallbladder perforation and massive intra-abdominal haemorrhage complicating acute cholecystitis in a patient with haemophilia A. *BMJ Case Rep* 2015;2015:bcr2014205971.

Author Contributions

Rowan Klein Nulend – Conception of the work, Design of the work, Acquisition of data, Analysis of data, Interpretation of data, Drafting the work, Revising the

work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Andrew Tse – Design of the work, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Timothy Pollitt – Conception of the work, Acquisition of data, Revising the work critically for important intellectual content, Final approval of the version to be published, Agree to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved

Guarantor of Submission

The corresponding author is the guarantor of submission.

Source of Support

None.

Consent Statement

Written informed consent was obtained from the patient for publication of this article.

Conflict of Interest

Authors declare no conflict of interest.

Data Availability

All relevant data are within the paper and its Supporting Information files.

Copyright

© 2024 Rowan Klein Nulend et al. This article is distributed under the terms of Creative Commons Attribution License which permits unrestricted use, distribution and reproduction in any medium provided the original author(s) and original publisher are properly credited. Please see the copyright policy on the journal website for more information.

Access full text article on
other devices



Access PDF of article on
other devices





Submit your manuscripts at
www.edoriumjournals.com

