Open Repair of Large Hepatic Artery Pseudoaneurysm Without Collateral Circulation: A Case Report

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Hepatic artery pseudoaneurysm is a rare arterial disease. This case report describes a patient with hepatic artery pseudoaneurysm who presented with recurrent epigastric pain over a 4-month period. Computed tomography angiography (CTA) showed aneurysmal enlargement of the hepatic artery measuring 55 mm × 46 mm. The angiographic information is as follows: (1) the common hepatic artery originated from the superior mesenteric artery; (2) the proper hepatic artery originated from the common hepatic artery; (3) the proper hepatic aneurysmal disease had no collateral circulation. After careful consideration, the patient underwent an open surgical repair (OSR). The patient recovered well without any associated complications. The 1-year follow-up of patients did not reveal any relevant complications. The treatment choice, puzzles, and reflections of this case are all discussed in this article.

Keywords: hepatic artery pseudoaneurysm, open surgery repair, endovascular approach, reconstruction, good prognosis

INTRODUCTION

Hepatic artery aneurysm (HAA) is a rare aneurysm with an overall prevalence of 0.002–0.4%, accounting for about 20% of visceral aneurysms and a rupture rate of 44% (1, 2). Arterial pseudoaneurysms account for 25–80% of reported cases and usually occur after medically induced injury or penetrating or blunt liver trauma, leading to symptomatic presentation of these aneurysms (3–5). HAA is the second most common visceral aneurysm after splenic artery aneurysm (6–10). Most common cases of HAA were observed during the sixth decade of life, with a 3:2 male predominance (1, 4). Majority of HAA cases are extrahepatic (75–80%) (1, 11, 12). A solitary aneurysm with multiple HAAs was reported in only 8% of cases (1). The risk factors of HAA include fibromuscular dysplasia, bacterial endocarditis, vasculitis, systemic lupus erythematosus or polyarteritis nodosa, Takayasu arteritis, Wegener granulomatosis, and congenital causes such as Marfan syndrome, Ehlers–Danlos syndrome, and Osler–Weber–Rendu syndrome (11–17). Abbas et al. (1) reported that patients with fibromuscular dysplasia and polyarteritis nodosa are at significant risk of HAA rupture, accounting for 50% of ruptured HAA. Hepatic artery pseudoaneurysm often occurs after iatrogenic injury or penetrating or blunt liver trauma, leading to symptomatic presentation of these aneurysms (3–5). The onset, diagnosis, treatment options, and advantages and disadvantages of various options for this patient, prognosis of this disease, and reflections of this case are discussed in this article.
CASE REPORT

A 69-year-old man was admitted to our hospital with complaints of intermittent epigastric pain for 4 months. Physical examination showed slight pressure pain in the upper abdomen but no rebound pain, and he had blood pressure of 136/84 mmHg, pulse of 96/min, and temperature of 36.3°C. Laboratory values on admission were as follows: 11.4 μmol/L direct bilirubin (DBIL), 237 IU/L alanine aminotransferase (ALT), 120 IU/L aspartate aminotransferase (AST), 106 g/L hemoglobin, 819 × 10^9/L white blood cells, and negative results of other blood tests. CTA revealed a 46 mm × 55 mm proper hepatic artery aneurysm arising from the common hepatic artery with multiple calcifications and local mural thrombosis (Figures 1A,B). Considering the risk of rupture, angiography was performed and endovascular treatment was prepared if needed. A 45 mm × 50 mm aneurysmal disease was observed about 30 mm from the initial segment of the proper hepatic artery without collateral vessels (Figure 1C). Deploying the stent was difficult considering the tortuosity of the delivery route. Otherwise, embolization was also excluded for the following reasons: first, the patient's hepatic artery pseudoaneurysm had no collateral vessels, which may lead to liver necrosis after embolization; second, the HAA was so large that it may compress the biliary tract and duodenum, causing jaundice, gastrointestinal obstruction, and even duodenal fistula. OSR was performed by right subcostal incision, approaching the proper hepatic artery pseudoaneurysm through the right gastrocolic ligament. The size of the pseudoaneurysm was ∼50 mm × 60 mm and closely adhesive to the omental sac and surrounding tissues. It is located between the medial duodenum, the head of the pancreas and the superficial part of the bile duct. Given the obvious peripheral adhesions, we did not explore the periphery of the aneurysm to reduce the possibility of tissue damage. The proximal and distal parts of the proper hepatic artery were mobilized, and the sac of the pseudoaneurysm was directly opened (Figure 2A). After removed the amount of thrombus from the sac, direct end-to-end anastomosis was performed with 6-0 Prolene, because the inflow and outflow vessels were adjacent (Figure 2B). The operation went smoothly. Hepatic artery clamp time was 25 min. Post-operatively, the patient had no special discomfort. Three days later, the patient was transferred to another hospital to continue his recovery without any complication. The patient's 1-year follow-up did not reveal any late complications (Figure 3). The results of blood reexamination showed 3 μmol/L DBIL, 30 IU/L ALT, and 34 IU/L AST.

DISCUSSION

In spite of their rarity with a reported incidence rate of 0.01–0.2%, visceral artery aneurysms hold a very important clinical significance, especially if we consider their natural history characterized by their propensity to rupture (18). These aneurysms are usually asymptomatic and difficult to detect until they rupture and cause abdominal pain and hypovolemic shock. The mortality rate after rupture of a visceral artery aneurysm remains quite high (30% as reported over the past decade) (19). The total prevalence rate of HAAs is 0.002–0.4%, accounting for about 20% of visceral aneurysms, and HAAs have a rupture rate of 44% (2, 20). HAAs are often related to hypertension, followed by malignancy and peripheral vascular disease. Other risk factors include chronic pancreatitis, trauma, and immune system disease (4).

HAAs can be diagnosed by ultrasound scan, computed tomography angiography (CTA), and digital subtraction angiography (DSA). According to The Society for Vascular Surgery clinical practice guidelines on the management of visceral aneurysms, CTA is the recommended diagnostic tool for patients who are thought to have HAA (Grade 1B); meanwhile, mesenteric angiography for pre-operative planning is recommended for patients with HAA who are considered for intervention (Grade 1B) (4).

Pseudoaneurysms can be distinguished from true aneurysms by antecedent clinical events such as iatrogenic injury or penetrating or blunt liver trauma along with specific imaging, focal arterial disruption in the setting of otherwise normal arteries, and inflammatory changes around an irregular aneurysm sac (4, 21). In contrast to true aneurysms, most pseudoaneurysms present with symptoms and are accompanied by gastrointestinal bleeding or bile bleeding (22, 23).

![Figure 1](https://example.com/figure1.png)

**Figure 1** (A) Shows the pre-operative three-dimensional CT imaging of the hepatic pseudoaneurysm. (B) Displays the hepatic pseudoaneurysm shown by CT in the pre-operative patient, with a maximum cross-section of about 55 mm × 46 mm. (C) Shows the hepatic pseudoaneurysm shown by hepatic arteriography; the red arrow indicates the hepatic artery pseudoaneurysm; the green arrow indicates the proper hepatic artery; the black arrow points to the common hepatic artery; and the blue arrow points to the gastroduodenal artery.
However, despite recent advances in therapeutic techniques and diagnostic tools, management of visceral artery aneurysm remains clinically challenging. Rupture is the most emergent and life-threatening situation for HAAs (4).

According to the clinical practice guidelines of the Society for Vascular Surgery on the management of visceral aneurysms, all hepatic artery pseudoaneurysms, given the high propensity of rupture and significant antecedent mortality, regardless of cause, should be repaired as soon as the diagnosis is made (Grade 1A) (4). However, in true HAAs, repair is recommended in the following situation: (1) all symptomatic HAAs regardless of size (Grade 1A); (2) asymptomatic patients without significant comorbidity, if true HAA is > 2 cm (Grade 1A) or if aneurysm enlarges >0.5 cm/y (Grade 1C), and patients with significant comorbidities, if HAA is >5 cm (Grade 1B); (3) patients with vasculopathy or vasculitis, regardless of size (Grade 1C) and patients with HAA with positive blood cultures (Grade 1C) (4).

Treatment approaches mainly include the following: arterial embolization, endovascular stent–graft repair, or OSR (4). The ideal surgical procedure is to remove the aneurysm while maintaining hepatic circulation. An endovascular-first approach is recommended to all HAAs if it is anatomically feasible (i.e., if this approach maintains arterial circulation to the liver) (Grade 1A) (4). In patients with extrahepatic aneurysms, open and endovascular techniques are recommended to maintain liver circulation (Grade 1A) (4). Coil embolization of the affected artery is recommended for patients with intrahepatic aneurysms (Grade 1B). Meanwhile, resection of the involved lobe of the liver is recommended for patients with large infrahepatic AAA to avoid significant liver necrosis (Grade 1C) (4). OSR or endovascular repair of visceral artery aneurysms yields similar long-term results, but morbidity is significantly worse with open repair (24, 25). Therefore, endovascular techniques should be preferentially offered for anatomically suitable candidates. Overall, endovascular therapy has become the mainstream technique. However, open repair remains the therapeutic regimen with definite efficacy and is mostly chosen for HAA cases of ruptured, asymptomatic common hepatic artery (>2 cm), or asymptomatic common hepatic artery in patients with fibromuscular dysplasia or polyarteritis nodosa, and proper hepatic and proximal right or left hepatic branches (4, 5, 21, 24–26).

A summary of treatment recommendations for extrahepatic aneurysms is shown in Table 1 (4).

Although the endovascular-first approach to all HAAs is proposed if it is anatomically feasible (4), Young Erben et al. (27) reported that among HAA cases requiring intervention, 81% are treated by OSR (66.7% open reconstruction, 4.8% endoaneurysmorrhaphy alone, 4.8% patch and 4.8% ligation), and that 19% are treated with endovascular techniques (coil embolization performed 9.5% in the common and 9.5% in the right hepatic arteries). Furthermore, overall mortality was 14% (6% after elective OT, 40% for emergency OT, 0% for ET). Some retrospective case series have shown that the
outcome for visceral artery aneurysms after OSR or endovascular repair yielded similar long-term results, but that morbidity is significantly worse with open repair than with the endovascular approach (25). Empirically, pseudoaneurysms tended to have higher mortality and reintervention rates (2). The main complications of endovascular treatment are hepatic ischemia, abscess, cholecystitis, possible recanalization, rebleeding, and internal fistula (27). In this case, we did not choose endovascular approaches mainly based on the following reasons: first, the patient's hepatic artery pseudoaneurysm with a huge size of about 45 mm × 50 mm had no collateral vessels. If we performed embolization, it is likely to compress the pancreas, duodenum, and anastomosis because of the short distance of the proximal to the distal part of the HAA.

CONCLUSION

Although endovascular therapy is the first choice of treatment in most cases, open surgery still has its unique role. We should not only strictly grasp the indications of various surgical procedures but also make clinical decisions according to specific conditions of patients. What is more, timely and correct diagnosis is the first step in treatment; although an HAA is a rare disease, it can be potentially lethal especially if left untreated or becomes late diagnosed.

DATA AVAILABILITY STATEMENT

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

ETHICS STATEMENT

Written informed consent was obtained from the individual(s) for the publication of any potentially identifiable images or data included in this article.

AUTHOR CONTRIBUTIONS

XW and XC were contributed equally to this article and mainly responsible for the collection of data and article writing. JZ and BH were mainly responsible for treatment plan selection, surgical design, and implementation. QG, XL, and XD were mainly responsible for data collection and surgical assistance. All authors contributed to the article and approved the submitted version.

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REFERENCES

1. Abbas MA, Fowl RJ, Stone WM, Panneton JM, Oldenburg WA, Bower TC, et al. Hepatic artery aneurysm: factors that predict complications. J Vasc Surg. (2003) 38:41–5. doi: 10.1016/s0741-5214(03)00090-9
2. Barrionuevo P, Malas MB, Nejim B, Haddad A, Morrow A, Ponce O, et al. A systematic review and meta-analysis of the management of visceral artery aneurysms. J Vasc Surg. (2019) 70:1694–9. doi: 10.1016/j.jvs.2019.02.024
3. Scott AB. Hepatic and splenic artery aneurysms. Semin Vasc Surg. (2005) 18:196–201. doi: 10.1053/j.semvasc.2005.09.005
4. Chaer RA, Abularrage CJ, Coleman DM, Eslami MH, Kashyap VS, Rockman C, et al. The Society for Vascular Surgery clinical practice guidelines on the management of visceral aneurysms. J Vasc Surg. (2020) 72:35–39. doi: 10.1016/j.jvs.2020.01.039
5. Fankhauser GT, Stone WM, Naidu SG, Oderich GS, Ricotta JJ, Bjarnason H, et al. The minimally invasive management of visceral artery aneurysms and pseudoaneurysms. J Vasc Surg. (2011) 53:966–70. doi: 10.1016/j.jvs.2010.10.071
6. Gjoreski A, Risteski F, Damjanoski G. Successful endovascular treatment of a giant hepatic artery aneurysm with dual layer stents placement as flow-diverting option: case report. Open Access Maced J Med Sci. (2019) 7:403–6. doi: 10.3889/oamjms.2019.120
7. Hossain A, Reis ED, Dave SP, Kersten MD, Hollis LH. Visceral artery aneurysms: experience in a tertiary-care center. Am Surg. (2001) 67:432–7.
8. Messina LM, Shanley CJ. Visceral artery aneurysms. Surg Clin North Am. (1997) 77:425–42. doi: 10.1016/s0039-6109(97)07055-4
9. Pulli R, Dorigo W, Trosi N, Pratesi G, Innocenti AA, Pratesi C. Surgical treatment of visceral artery aneurysms: a 25-year experience. J Vasc Surg. (2008) 48:334–42. doi: 10.1016/j.jvs.2008.03.043
10. Zhakhayev M, Maruya Y, Takatsuki M, Baimakhanov Z, Soyama A, Hidaka M, et al. Stent treatment for huge aneurysm of the common hepatic artery: a case report. Radial Case Rep. (2019) 14:44–7. doi: 10.1016/j.radcr.2018.09.008
11. Erskine JM. Hepatic artery aneurysm. Vasc Surg. (1973) 7:106–25. doi: 10.1177/15385747300700205
12. Lal RB, Strohl JA, Piazza S, Aslam M, Ball D, Patel K. Hepatic artery aneurysm. J Cardiovasc Surg. (1989) 30:509–513.
13. Caputo AE, Roberts WN, Yee YS, Posner MP. Hepatic artery aneurysm in corticosteroid-treated, adult Kawasaki's disease. Ann Vasc Surg. (1991) 5:533–7. doi: 10.1016/bf02015278
14. Chan RJ, Goodman TA, Aretz TH, Lie JT. Segmental mediolycytic arterioapathy of the splenic and hepatic arteries mimicking systemic necrotizing vasculitis. Arthritis Rheumat. (1998) 41:935–8. doi: 10.1002/1529-0131(199805)41:5<935::aID-art22>3.0.Co;2-n
15. den Bakker MA, Tangkau PL, Steffens TW, Tjiarm SL, van der Loo EM. Rupture of a hepatic artery aneurysm caused by Wegener's granulomatosis. Pathol Res Pract. (1997) 193:61–6. doi: 10.1016/s0343-0338(97)80096-9
16. Hasen-Khodja R, Declemy S, Batt M, Castanet J, Perrin C, Ortonne JP, et al. Visceral artery aneurysms in Von Recklinghausen's neurofibromatosis. J Vasc Surg. (1997) 25:572–5. doi: 10.1016/s0741-5214(97)0270-2
17. Koyama M, Tanaka M, Shimizu M, Nomura S, Kako N, Suzuki S, et al. Surgical treatment of mesenteric infarction, thoracoabdominal aortic aneurysm, and proper hepatic aneurysm in a middle-aged woman with Takayasu's arteritis. J Cardiovasc Surg. (1995) 36:33–41.
18. Chen XY, Ge JT, Zhao JC, Yuan D, Yang Y, Huang B. Duodenal necrosis associated with a threatened ruptured gastroduodenal artery pseudoaneurysm complicated by chronic pancreatitis: case report. Ann Vasc Surg. (2020) 68:571.e9–571.e13. doi: 10.1016/j.avsg.2020.04.050
19. Zhao JC. Massive upper gastrointestinal bleeding due to a ruptured superior mesenteric artery aneurysm duodenum fistula. *J Vasc Surg.* (2008) 48:735–7. doi: 10.1016/j.jvs.2008.04.015

20. Haghighatkah H, Taheri MS, Kharazi SM, Zamini M, Khorasgani SR, Zarkani ZI. Hepatic artery aneurysms as a rare but important cause of abdominal pain; a case series. *Arch. Acad Emerg Med.* (2019) 7:e25.

21. Nirman T, Kashyap VS, Greenberg RK, Sarac TP, Clair DG, Pierce G, et al. The endovascular management of visceral artery aneurysms and pseudoaneurysms. *J Vasc Surg.* (2007) 45:276–83. Discussion 283. doi: 10.1016/j.jvs.2006.10.049

22. Graff J, Schälte G, Jovanovic V. Rupture of a renal artery aneurysm during delivery. *Aktuelle Urol.* (2003) 34:350–3. doi: 10.1055/s-2003-42004

23. Pierre-François Plouin PF, Perdu J, Batide-Alanore AL, Boutouyrie P, Gimenez-Roqueplo AP, Jeunemaitre X. Fibromuscular dysplasia. *Orphanet J Rare Dis.* (2007) 2:28. doi: 10.1186/1750-1172-2-28

24. Cochennec F, Riga CV, Allaire E, Cheshire NJW, Hamady M, Jenkins MP, et al. Contemporary management of splanchnic and renal artery aneurysms: results of endovascular compared with open surgery from two European vascular centers. *Eur J Vasc Endovasc Surg.* (2011) 42:340–6. doi: 10.1016/j.ejvs.2011.04.033

25. Sachdev U, Baril DT, Elozy SH, Lookstein RA, Silverberg D, Jacobs TS, et al. Management of aneurysms involving branches of the celiac and superior mesenteric arteries: a comparison of surgical and endovascular therapy. *J Vasc Surg.* (2006) 44:718–24. doi: 10.1016/j.jvs.2006.06.027

26. Cordova AC, Sumpio BE. Visceral Artery aneurysms and pseudoaneurysms—should they all be managed by endovascular techniques? *Ann Vasc Dis.* (2013) 6:687–93. doi: 10.3400/avd.ra.13-00045

27. Erben Y, Martino RRD, Bjarnason H, Duncan AA, Kalra M, Oderich GS, et al. Operative management of hepatic artery aneurysms. *J Vasc Surg.* (2015) 62:610–5. doi: 10.1016/j.jvs.2015.03.077

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