First two cases of literature: Caustic sclerosing cholangitis due to percutaneous treatment of hydatid liver disease causing liver transplantation

Short title: Sclerosing cholangitis due to PAIR

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Abstract
Echinococcus granulosus is predominantly found in the liver and can be effectively managed through antihelminthic therapy, surgical intervention, or interventional radiology. Percutaneous treatment (PT) has gained widespread popularity due to its minimally invasive nature. An integral step in surgical and PT procedures involves the utilization of protoscolicidal agents to eliminate the parasites. However, the administration of protoscolicidal agents carries the risk of inducing caustic sclerosing cholangitis (SC) if there exists a communication between the cyst and the biliary tract. In this pioneering study, we present two cases of caustic SC that occurred subsequent to PT for hydatid liver, necessitating further progression of the disease and ultimately leading to liver transplantation.

**Keywords:** hydatid disease; hydatid cyst; PAIR; cholangitis; liver transplantation

**Introduction**

Hydatid disease, attributed to the parasite Echinococcus granulosus, primarily affects the liver and lungs, although it can manifest in various tissues and organ systems. Treatment options for liver hydatid cysts encompass medical, surgical, or percutaneous interventions. Percutaneous interventions involve the guidance of ultrasound or computed tomography to insert a catheter into the cyst cavity. A scolicidal agent is then injected and subsequently aspirated through this catheter. Suggested scolicidal agents include 20-30% hypertonic saline, 95% ethanol, or 0.5% cetrimide [1]. However, the use of these agents carries the risk of inducing severe toxicity to the biliary tract in the presence of a cystobiliary connection. This study presents two cases of secondary sclerosing cholangitis (SC) that arose following the administration of scolicidal agents during percutaneous therapy (PT) and necessitated liver transplantation. To the best of our knowledge, the requirement for liver transplantation as a consequence of SC subsequent to PT for hepatic hydatid cysts has not been reported previously.

**Patients and Methods**
Case 1

A 50-year-old male patient was initially admitted to another medical center with complaints of right upper quadrant pain. Upon evaluation, a type 1 hydatid cyst located in segments 6-7 of the liver was detected. The patient underwent medical therapy and subsequently underwent a procedure known as puncture-aspiration-injection reaspiration (PAIR), using a 20% saline solution. Following the procedure, the patient was discharged on the first day and reported no symptoms except for intermittent pain, which persisted for eight months. During outpatient follow-up visits, the attending physician confirmed the eradication of the cyst.

After eight months, the patient was readmitted to the hospital due to jaundice and right upper quadrant pain. Clinical, laboratory, radiological, and endoscopic retrograde cholangiopancreatography (ERCP) findings confirmed the diagnosis of SC (see Figure 1). The patient underwent regular follow-up, receiving endoscopic biliary stenting for five years to manage recurrent cholangitis attacks. Eventually, the patient was referred to our clinic for the evaluation of secondary biliary cirrhosis. Considering the patient's moderately impaired liver function (Model for End-Stage Liver Disease score [MELD Na]: 13, Child Class B), liver transplantation was not initially considered. However, due to recurring episodes of cholangitis and the development of liver abscesses, multiple percutaneous drainage procedures were required.

In the sixth year following the PAIR procedure, the patient was admitted to our institute with advanced cirrhosis (MELD Na: 26, Child Class C), leading to the decision to perform a living donor right lobe liver transplantation. Intraoperative findings indicated cholestasis and fibrosis in the liver, with the common bile duct displaying sclerotic changes and thickened walls resulting from recurrent ERCP procedures and SC episodes. The postoperative period was complicated by the occurrence of biliary fistula and hepatic artery thrombosis, necessitating a
relaparotomy. Fortunately, the patient had an uneventful clinical course following the relaparotomy and was successfully discharged.

Pathological evaluation of the explanted liver confirmed the presence of cirrhosis and obliteration of the biliary tree. Tragically, the patient succumbed to sepsis caused by secondary infection of a bilioma resulting from a biliary leak attributed to late hepatic artery thrombosis in the fourth month following the liver transplantation.

Case 2

A 26-year-old female patient underwent PAIR procedure in another medical center for the management of a type 1 hydatid cyst located in segment 7 of the liver. Ethanol was employed as the scolicidal agent during the intervention. Immediately after the procedure, the patient developed persistent abdominal pain, fever, and hyperbilirubinemia. Subsequent imaging with computerized tomography and endoscopic retrograde cholangiopancreatography (ERCP) (see Figure 2) confirmed the diagnosis. After 26 days of hospitalization, the patient was referred to our institute with a secondary diagnosis of SC accompanied by liver failure.

Upon evaluation, it was determined that the patient had end-stage liver failure, with a Model for End-Stage Liver Disease score (MELD Na) of 31.5 and Child Class C11. To address her condition, a living donor liver transplantation was performed. The patient's postoperative course was uneventful, and she was discharged on the 21st day after the surgery. Pathological examination of the explanted liver revealed the presence of a segmental hydatid cyst that had partially ruptured into the major bile ducts. Furthermore, mucosal ulceration was observed in the extrahepatic biliary tract, accompanied by increased connective tissue thickening of the duct. During the 31st-month follow-up after the transplantation, the patient's condition remained stable and favorable.

Discussion
While vague abdominal pain is a common symptom associated with hydatid liver cysts, these cysts often remain asymptomatic and are typically discovered incidentally during patient examinations. Over time, the cysts gradually grow and become symptomatic when they rupture into the biliary tract, resulting in obstructive jaundice. In some cases, the cysts may rupture spontaneously into the peritoneal cavity [2]. Laboratory tests of patients with uncomplicated cysts often yield normal results. However, in cases where there is communication between the cyst and the biliary system, liver function tests and cholestasis enzymes may show elevated levels. Due to the non-specific nature of the symptoms and findings from history and physical examination, radiological imaging, such as ultrasound and computed tomography, becomes essential for accurate diagnosis. Radiological classification of hepatic hydatid cysts is based on the World Health Organization Informal Working Group on Echinococcosis and Gharbi classification system [3].

The choice of treatment for hydatid cysts depends on the type of cyst. Percutaneous treatment (PT) is commonly employed in patients who are not suitable candidates for surgery, those who prefer non-surgical options, cases of recurrence, and individuals who do not respond to benzimidazole treatment. PT modalities are also used for World Health Organization cyst types CE1 and CE3a, as well as for cysts smaller than 5 cm, pregnant women, and children under 3 years old [4]. However, PT is generally not recommended for WHO cyst types CE2, CE3b, CE4, and CE5, as well as for cysts located in the lungs and those that communicate with the biliary tree [4]. For cysts larger than 10 cm, the preferred approach is often the placement of a drainage catheter into the cystic cavity, which is then removed when the drainage volume falls below 10 ml [5].

Although percutaneous intervention has relatively low reported rates of mortality and morbidity, it is not without specific and serious complications. In a study, the overall rate of various complications associated with PT, including anaphylaxis, allergic skin rash,
pneumothorax, cavity infection, and cystobiliary fistula, was reported to be approximately 9% [6]. The most concerning complications are the risk of anaphylactic shock (0.38%) and secondary intra-abdominal hydatidosis (1.3%) resulting from the dissemination of cyst content into the abdomen during the intervention. Additionally, there is a risk of scolicidal agents causing cholangitis in cases where there is communication between the cyst and the biliary tract. However, such a complication has not been reported thus far [7].

Caustic SC is a term coined by Belghiti et al. in 1986 [8] to describe SC caused by scolicidal agents. It is believed to arise from a combination of four key factors: injection of scolicidal agents into the cyst cavity, presence of cystobiliary communication, prolonged exposure of the biliary tree to the scolicidal solution, and individual sensitivity to the scolicidal agent [9]. In our presented cases, caustic SC was diagnosed, which distinguishes them from primary SC based on two distinctive characteristics.

Firstly, the disease in both patients exhibited a rapid progression compared to the natural course of primary SC. In patients with primary SC, the time interval between the initial diagnosis and liver transplantation typically spans between 10 to 14 years. However, in our cases, the disease progressed rapidly over a period ranging from months to several years.

Secondly, unlike primary SC where fibrotic and stenotic biliary lesions affect the entire biliary tree, caustic SC observed in our cases was localized only to the portion exposed to the scolicidal agent. This finding aligns with the observations made in our presented patients.

Bakdik et al. reported a cystobiliary communication rate of 5.4% in patients who underwent PAIR treatment [9]. Determining whether there is a cystobiliary connection before PAIR is the most important point in preventing caustic SC formation. For this purpose, it is necessary to clarify the relationship between the cyst and the biliary tree before using the scolicidal agent during PAIR. However, it is important to note that cystobiliary connection alone is insufficient
for the development of cholangitis. The presence of a certain degree of biliary obstruction, which can occur in hydatid cysts that rupture into the biliary tract, as well as individual sensitivity to the agent, are additional contributing factors [10].

**Conclusion**

While there have been reports in the literature regarding caustic cholangitis resulting from the use of scolicidal agents during surgery, there have been no documented cases of caustic cholangitis following PAIR. Our presented cases demonstrate that percutaneous treatment of liver hydatid cysts has the potential to induce caustic SC in patients with abnormal cystobiliary communication. This highlights the importance of considering this potential complication in patients undergoing percutaneous interventions for hydatid cysts, particularly those with cystobiliary communication.

**References**


**Figure legends:**

**Figure 1.** Strictures biliary areas.

**Figure 2.** Hydatid cyst with PAIR catheter
Figure 2.