

Introduction

Lemmel's syndrome is defined as obstructive jaundice caused by a periampullary duodenal diverticulum (PAD), leading to bile duct compression and dilation.

Most obstruction is at the ampulla of Vater, with an incidence as high as 27%. Most PAD are found incidentally and are suspected to be found in up to 22% of the population.

Some patients can have life threatening pathology, necessitating quick intervention.

Case Report

A 77-year-old Caucasian female with a history of cholecystectomy 20 years ago presented to the emergency department with a 1-day history of epigastric abdominal pain, mild fever, jaundice, and non-bloodly diarrhea.

Laboratory evaluation was notable for a lipase of 3326 U/L and transaminitis with ALP 908 IU/L, AST 234 U/L, ALT 153 U/L, and total bilirubin 7.2 mg/dL. CT abdomen with IV contrast revealed moderate-severe pancreatitis and intra/extrahepatic biliary ductal dilation up to 19 mm in diameter with additional dilation of the pancreatic duct.

A subsequent MRCP revealed a large periampullary duodenal diverticulum with obstruction of the common bile duct, consistent with Lemmel's syndrome, and possible duodenal diverticulitis [A, B].

Piperacillin-tazobactam was initiated due to concern for ascending cholangitis and diverticulitis. ERCP was performed the following morning, which revealed a large duodenal diverticulum in the second portion of the duodenum with fecalized material impacted within the diverticulum [C].

The entire biliary ampulla was obscured by obstructing material and surrounding ulceration was noted. Several attempts to remove the impaction were unsuccessful.

A temporary PTC drain was placed with interval improvement in her pain and lab abnormalities. The patient was transferred to an outside facility for advanced endoscopy, where she had successful removal of obstruction and intra/extrahepatic stenting.

Figures

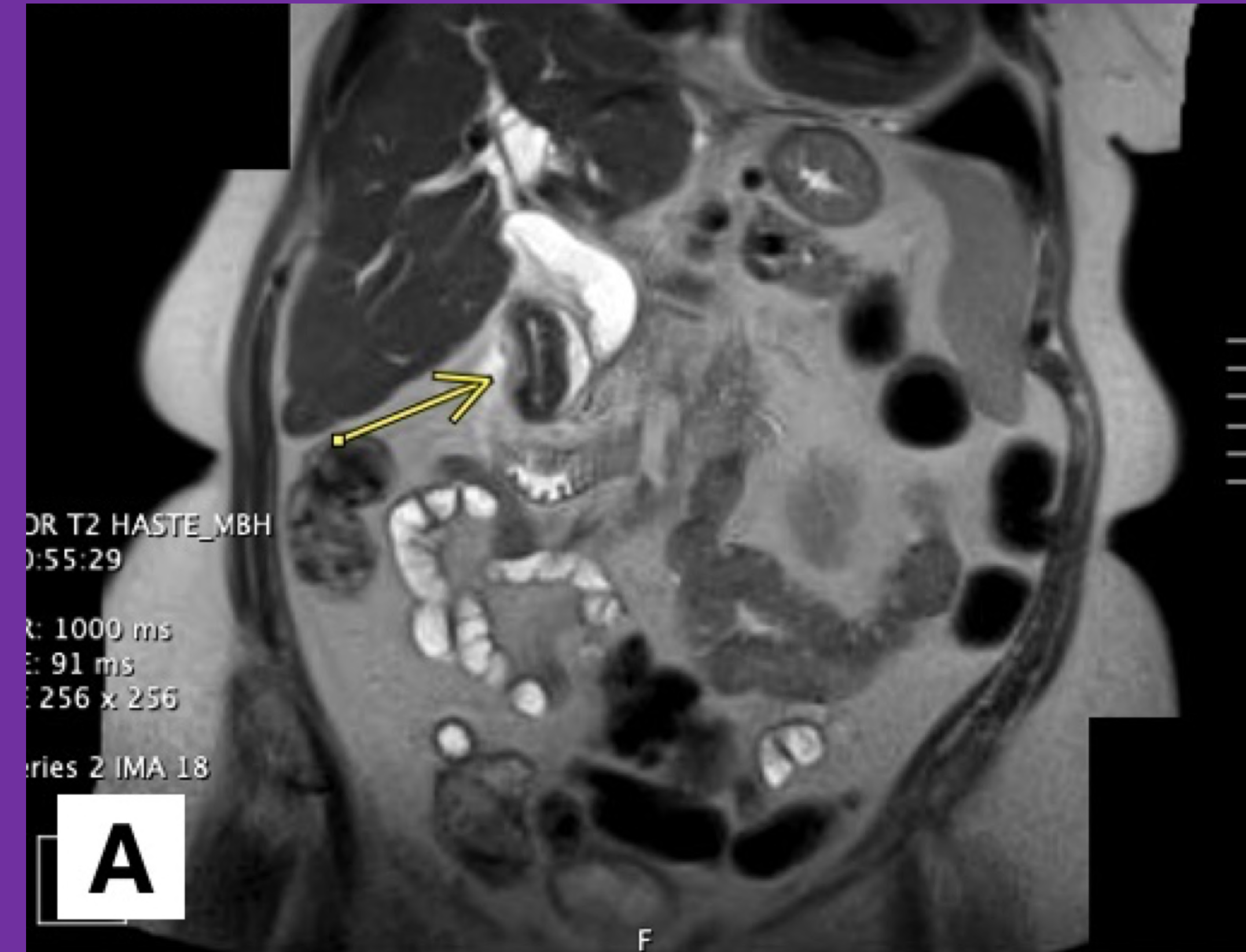


Figure A: Coronal slice from MRI abdomen and pelvis demonstrating a large periampullary duodenal diverticulum with obstruction of the common bile duct.



Figure B: Transverse slice from MRI abdomen and pelvis demonstrating a large periampullary duodenal diverticulum with obstruction of the common bile duct

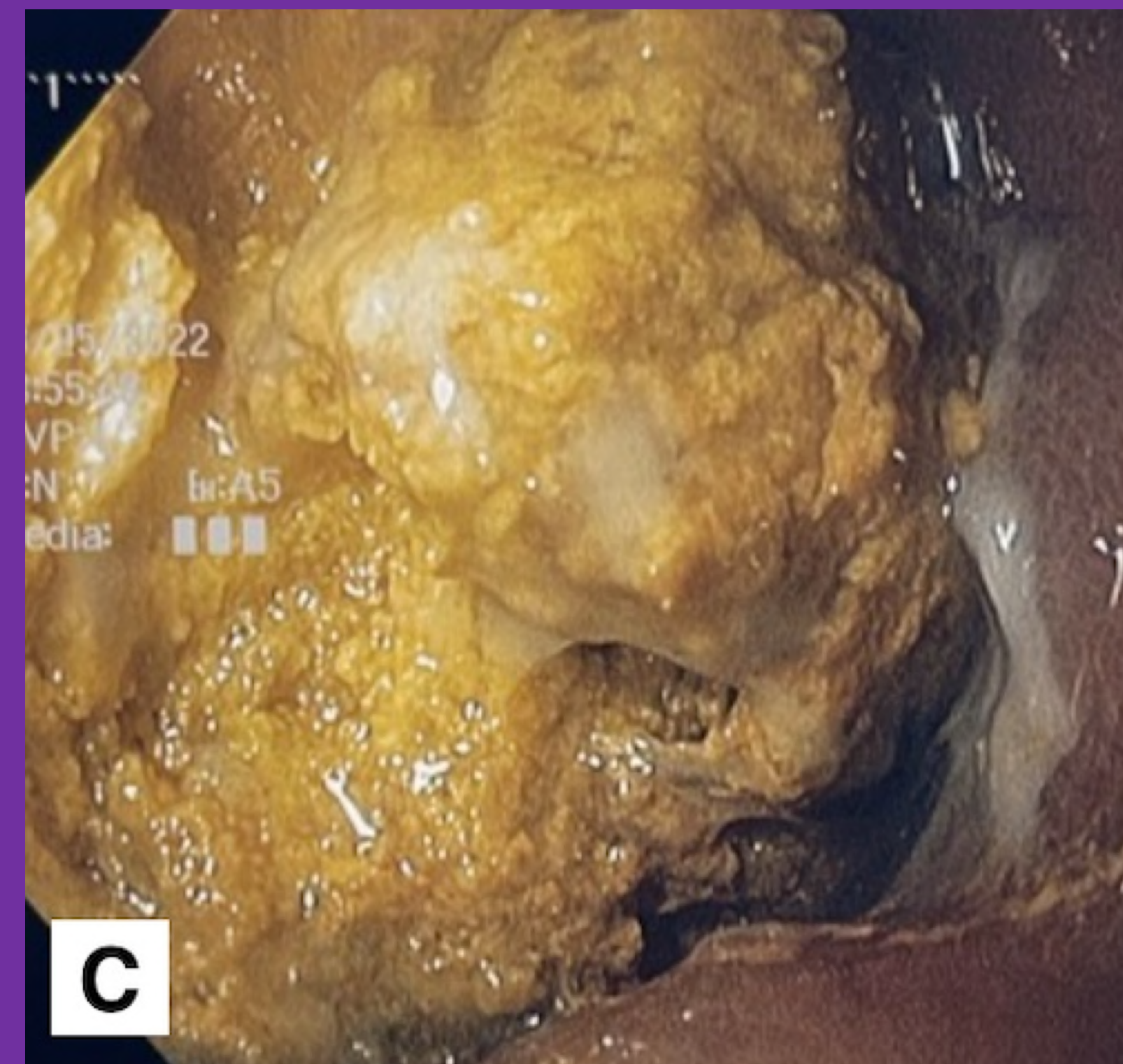


Image C: Image obtained during ERCP revealing impacted fecalized material within a large duodenal diverticulum in the second portion of the duodenum with surrounding ulceration.

Discussion

Most patients with periampullary duodenal diverticulum will remain asymptomatic and PAD will be found incidentally in most cases.

Lemmel's syndrome is classified as obstructive jaundice in the absence of choledocholithiasis or pancreato-biliary tumors.

Therapeutic options vary, but ERCP is considered the initial treatment of choice because it allows for the placement of a biliary stent and a sphincterotomy.

If Lemmel's is due to chronic papillary fibrosis or sphincter of Oddi dysfunction, then an endoscopic sphincterotomy is the preferred therapy.

If the disease is severe or complicated, a diverticulectomy is recommended in patients who are at lower risk for adverse complications.

Conclusion

A high index of suspicion is needed for urgent diagnosis and therapeutic intervention of this rare, but potentially life threatening pathology.

References

Michelle Bernshteyn, Suman Rao, Anuj Sharma, Umair Masood, Divey Manocha. Lemmel's syndrome: usual presentation of an unusual diagnosis. *Cureus*. 2020 April; 12(4): e7698.

LM Gorozieta-Rosales, J Gomez-Farias, KD Lopez-Garcia, DO Davila-Rodriguez. Lemmel syndrome: an extraordinary cause of obstructive jaundice – a case report. *J Surg Case Rep*. 2022