

# Diagnostic Imaging Pathways - Pancreatitis (Acute)

## Population Covered By The Guidance

This pathway provides guidance on the investigation of adult patients with suspected acute pancreatitis and its complications.

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## Quick User Guide

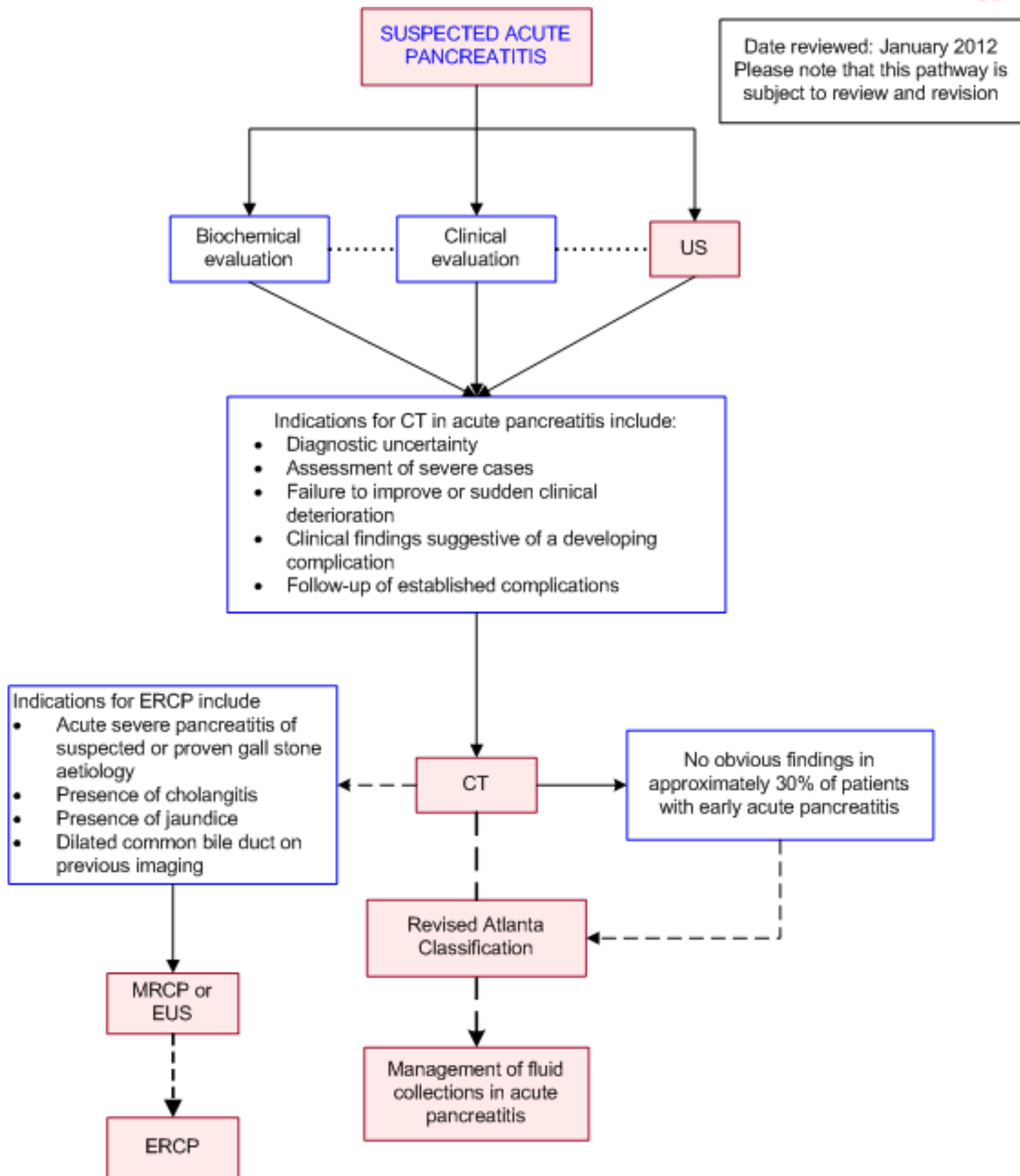
Move the mouse cursor over the **PINK** text boxes inside the flow chart to bring up a pop up box with salient points.

Clicking on the **PINK** text box will bring up the full text.

The relative radiation level (RRL) of each imaging investigation is displayed in the pop up box.

SYMBOL	RRL	EFFECTIVE DOSE RANGE
	None	0
	Minimal	< 1 millisieverts
	Low	1-5 mSv
	Medium	5-10 mSv
	High	>10 mSv

## Pathway Diagram



## Teaching Points

### Role of Imaging in acute pancreatitis



- Exclude an underlying cause (e.g. gallstones)
- Assess severity
- Detect complications
- Guide treatment of complications (e.g. fluid collection drainage)

**CT SCAN** - routine CT scan is not indicated. Indications include

- Where diagnosis is in doubt
- Clinically severe cases to assess degree of pancreatic necrosis
- Failure to improve or sudden deterioration
- Imaging complications of pancreatitis

**US Scan**

- To help determine aetiology of pancreatitis
- Assess for gallstone-induced pancreatitis
- Assess bile duct if abnormal liver function

**ERCP (Endoscopic Retrograde Cholangiopancreatography)** - indications include

- Severe pancreatitis of proven or suspected gallstone aetiology
- Presence of cholangitis
- Presence of jaundice

## Acute Pancreatitis

- The diagnosis of pancreatitis is usually made clinically and biochemically [1](#)
- In suspected acute pancreatitis, imaging is used to [1](#)
  - Exclude an underlying cause (e.g. gallstones)
  - Assess severity
  - Detect complications
- Clinical definition of acute pancreatitis (whether or not chronic pancreatitis is present) requires at least 2 out of 3 of the following [2](#)
  - Abdominal pain strongly suggestive of acute pancreatitis
  - Serum amylase / lipase levels of ? 3 times normal level
  - Characteristic imaging findings on imaging (US / CT / MRI)

## Revised Atlanta Classification

- In 2008, acute pancreatitis Classification Working group revised the 1992 Atlanta classification to clarify previous areas of confusion, improve clinical assessment & management, standardise the description of patients for reporting clinical studies and to offer a standardised means of data collection for future studies to allow objective evaluation of new therapies [2](#)
- **Summary of Revised Atlanta Classification** [17,18](#)

Morphologic Type	Associated Collections
<4 weeks	
Interstitial edematous pancreatitis	Sterile / Infected Acute Peri-pancreatic fluid collection



	(APFCs)
Necrotizing pancreatitis	Sterile / Infected Acute parenchymal necrotic collections or Sterile / Infected Acute Peri-pancreatic necrotic collection or Sterile / Infected combined pancreatic and peri-pancreatic necrotic collection (ANCs)
?4 weeks	
Interstitial edematous pancreatitis	Sterile / Infected pseudocyst
Necrotizing pancreatitis	Sterile / Infected walled-off necrosis (WON)

- Clinical severity and organ failure is calculated using scoring systems like Marshall system, SOFA, APACHE-II or Ranson scoring systems [19,20,21,22](#)

## Computed Tomography (CT)

- Contrast enhanced CT (CE-CT) is the imaging modality of choice for evaluating pancreas and the surrounding tissues [3,7](#) and is often the first radiological investigation for suspected acute pancreatitis in many institutions
- Routine CT is not indicated in mild acute pancreatitis unless there are clinical or other signs of deterioration [1,7,8,9](#), and there is no advantage of performing early imaging to predict the clinical severity of acute pancreatitis more than a clinical evaluation [9](#)
- 14-28% of CT scans are normal in mild pancreatitis [7,8](#)
- Ideal time for CE-CT is 48 hours after onset of symptoms for better accuracy in detecting pancreatic necrosis but in practice, patients with undiagnosed abdominal pain CE-CT is often performed on admission or the diagnosis of AP would have been made on CE-CT
- Indications for CT scan include [1,3,8](#)
  - Diagnostic uncertainty
  - Assessment of severity and to detect complications
  - Failure to improve on treatment (>48 hrs)
  - Clinical findings suggesting a developing complication (e.g. fever, pain, hypotension, decreasing haematocrit)
  - Sudden deterioration in clinical status following an initial response to medical treatment
  - Follow-up and monitoring of established complications
  - Guidance of interventional procedures such as percutaneous fine needle aspiration and/or catheter drainage of fluid collections
- Combination of pre and post-contrast enhancement appearances permits the assessment of the degree of pancreatic necrosis and surrounding peri-pancreatic and intra-abdominal fluid collections. The severity of disease as demonstrated on CT (CT severity index) correlates with the risk of morbidity and mortality [10](#)
- Disadvantages - exposure to ionising radiation with repeat scanning

## Endoscopic Retrograde Cholangiopancreatography (ERCP)

- Mainly used to locate and remove gallstones in the common bile duct among patients with severe pancreatitis attributable to gallstones [1](#)
- Other indications for ERCP in the setting of acute pancreatitis include [1](#)
  - Presence of ascending cholangitis
  - Presence of jaundice
  - Dilated common bile duct on previous imaging
- Urgent ERCP and sphincterotomy is indicated in patients with severe gallstone pancreatitis who fail to respond to treatment within 48 hours [14,15](#)
- Similarly patients with gallstone acute pancreatitis who develop ascending cholangitis stand to benefit from early ERCP and endoscopic sphincterotomy [16](#)

## Fluid Collections In Acute Pancreatitis

- The RAC classifies fluid collections in acute pancreatitis based on both the morphologic classification they are associated with and the disease timeframe
- Collections can be sterile or infected at any time and can occur in all the collection types
- Fluid collections associated with IEP in the first 4 weeks of onset are called acute peri-pancreatic fluid collections (APFC). If these collections progress / persist for 4 weeks or more, they are termed pancreatic pseudocysts. Pseudocysts occur in 10-20 % of patients as a complication of acute pancreatitis [17](#)
- Fluid collections associated with necrotizing pancreatitis are called acute necrotic collections (ANC) if occurring within 4 weeks and walled-off necrosis (WON) after 4 weeks. ANCs can be further divided based on the morphological classification of the pancreatitis they are associated with. Parenchymal collections occurring within the first 4 weeks should also be classified as necrotic collections [17](#)
- The question of intervention (usually percutaneous aspiration/drainage) for relatively symptomatic pseudocysts/fluid collections is a balance between on the one hand, the risks of introducing infection into a sterile collection and draining an "immature" cyst and on the other hand, the complications of a large untreated, unresolved fluid collection
- Fluid collections in acute pancreatitis can be categorised into the following (general guidelines only)
  1. Acute peri-pancreatic fluid collections (APFCs)
    - Infection is extremely rare
    - Majority get reabsorbed with no complications
    - Fine needle aspiration (FNA) is only indicated if strong suspicion of infection. Otherwise no active invasive treatment is necessary
  2. Pancreatic pseudocysts [23](#)
    - A pancreatic pseudocyst consists of enzyme-rich fluid surrounded by a wall of granulation or fibrous tissue
    - May be localised to the pancreas or located remotely. Communication with the pancreatic ductal system is present in up to 80% of cases
    - Spontaneous regression occurs in 30-50% of cases and most pseudocysts less than 4cm in diameter resolve within 6 weeks
    - Infection can be noted by the presence of gas locules within pseudocyst. If no gas is visible on CE-CT, FNA can be done to rule out infection but risk of introducing infection by performing FNA should be taken into consideration
    - Drainage is indicated for pseudocysts larger than 5cm, that are growing, symptomatic or infected

### 3. Necrotic collections (ANCs and WONs)

- FNA is useful to distinguish between infected and sterile necrosis, with a sensitivity of 88-96% and specificity of 90-96% [24,25](#)
  - Indications for FNA include: failure to improve within 48-72 hours of commencing medical therapy, persistent symptoms for more than 7 days with greater than 30% necrosis or clinical suspicion of sepsis with less than 30% necrosis
  - Sterile ANCs may be drained based on patient's clinical condition. Percutaneous drainage is preferable though surgery and endoscopic procedures may be done rarely
  - Infected ANCs are drained with percutaneous drainage but surgery / endoscopic procedure may be needed later if recurs / inadequate
  - Sterile WON are drained based on clinical circumstances and percutaneous drainage is preferred but surgical drainage / endoscopic drainage may be needed for a cure
  - Infected WON are drained with percutaneous drainage as an interim with surgery to follow
- Indications for aspiration/drainage include [26,27,28,29](#)

1. Diagnosis of possible infection/abscess. If aspiration confirms infection, possible therapeutic options are dependent on the morphology of the collection and the clinical status of the patient. They include
  - Percutaneous catheter drainage either as a definitive procedure or as a "holding" measure pending surgery
  - Surgical drainage/debridement as a first-line treatment
  - Endoscopic drainage via the stomach or duodenum
2. Continuing symptoms considered due to the mass effect of the fluid collection
3. Cyst enlarging on serial follow-up imaging. In this situation ERCP may be useful. If communication between the pancreatic duct and the fluid collection is demonstrated, the need for prolonged drainage is likely and surgery may be a better option
4. Some authorities suggest size alone as a criterion for drainage (usually around 5 cm)

## Ultrasound

- Recommended to help determine the aetiology in all patients with suspected acute pancreatitis [1,2,3](#)
- Primarily used to assess the biliary tree for gallstones, duct dilatation/obstruction and to exclude other pathology [1,2,3](#)
- Helps distinguish fluid collections from solid inflammatory masses
- Useful for follow-up of pancreatic fluid collections if seen well on initial ultrasound [4](#)
- Limitations
  - Visualisation of the pancreas is usually sub-optimal due to overlying bowel gas from a coexistent ileus [5,6](#)
  - Detection of intra-parenchymal and retroperitoneal fluid collections correlates poorly with pancreatic necrosis [3](#)
  - Often underestimates the presence, extent and complexity of fluid collections

## Magnetic Resonance Cholangiopancreatography (MRCP) and Endoscopic

## Ultrasonography (EUS)

- In many centres MRCP and EUS are performed following CT scanning if gall stone pancreatitis is being suspected prior to patients undergoing invasive ERCP if needed
- MRCP is reported to have a high negative predictive value of 100% for CBD stones [11,12](#)
- MRCP is non-invasive and has no ionising radiation risk compared to CT. It is reported to have a sensitivity of around 62% and specificity of around 98% for CBD stones [12](#)
- EUS is an invasive imaging method but is reported to have a higher diagnostic yield (51% vs 20%) compared to MRCP in a prospective study looking for causes of idiopathic pancreatitis following traditional cross-sectional imaging [11](#)
- Some studies report higher diagnostic yield for EUS and MRCP compared to ERCP in idiopathic pancreatitis [13](#)

## Endoscopic Ultrasonography (EUS)

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- Some studies report higher diagnostic yield for EUS and MRCP compared to ERCP in idiopathic pancreatitis [13](#)

## References

References are graded from Level I to V according to the Oxford Centre for Evidence-Based Medicine, Levels of Evidence. [Download the document](#)

1. UK Working Party on Acute Pancreatitis. **United kingdom guidelines for the management of acute pancreatitis**. Gut. 2005;54(suppl III):iii1-iii9 (Guidelines). [View the reference](#)
2. Sarr MG, Banks PA, Bollen TL et al. **Revision of the Atlanta Classification of acute pancreatitis**. Acute Pancreatitis Classification Workgroup, 2008. April. Accessed 25/05/2012. (Guidelines). [View the reference](#)
3. Balthazar EJ. **Acute pancreatitis: assessment of severity with clinical and CT evaluation**. Radiology. 2002;223:603-13. (Review article)
4. Dalzell DP, Scharling ES, Ott DJ, et al. **Acute pancreatitis: The role of diagnostic imaging**. Crit Rev Diagn Imaging. 1998;39(5):339-363. (Review article)
5. Silverstein W, Isikoff M, Hill M, et al. **Diagnostic imaging of acute pancreatitis: prospective study using CT and sonography**. AJR Am J Roentgenol. 1981;137:497-502. (Level III evidence)
6. McKay A, Imrie C, O'Neill J, et al. **Is an early ultrasound scan of value in acute pancreatitis?** Br J Surg. 1982;69:369-72. (Level III evidence)
7. Jacobs JE, Birnbaum BA. **Computed tomography evaluation of acute pancreatitis**. Semin Roentgenol. 2001;36(2):92-8. (Review article)
8. Balthazar EJ, Freeny PC, vanSonnenberg E, et al. **Imaging and intervention in acute pancreatitis**. Radiology. 1994;193:297-306. (Review article)
9. Morteale KJ, Ip IK, Wu BU, Conwell DL, Banks PA, Khorasani R. **Acute Pancreatitis: Imaging utilization practices in an urban teaching hospital - analysis of trends with assessment of independent predictors in correlation with patient outcomes**. Radiology. 2011;258:174-81.



(Level III evidence)

10. Balthazar EJ, Robinson DL, Megibow AJ, et al. **Acute pancreatitis: value of CT in establishing prognosis.** Radiology. 1990;174:331-6. (Level II/III evidence)
11. Ortega AR, Gomez-Rodriguez R, Romero M, Fernandez-Zapardiel S, Cespedes Mdel M, Carrobes JM. **Prospective comparison of endoscopic ultrasonography and magnetic resonance cholangiopancreatography in the etiological diagnosis of "idiopathic" acute pancreatitis.** Pancreas. 2011;40(2):289-94. (Level II evidence)
12. Srinivasa S, Sammour T, McEntee B, Davis N, Hill AG. **Selective use of magnetic resonance cholangiopancreatography in clinical practice may miss choledocholithiasis in gallstone pancreatitis.** Can J Surg. 2012;53(6):403-7. (Level III evidence)
13. Mariani A, Arcidiacono PG, Curioni S, Giussani A, Testoni PA. **Diagnostic yield of ERCP and secretin-enhanced MRCP and EUS in patients with acute recurrent pancreatitis of unknown aetiology.** Dig Liver Dis. 2009;41(10):753-8. (Level III evidence)
14. Neoptolemos JP, Carr-Locke DL, London NJ, et al. **Controlled trial of urgent retrograde cholangiopancreatography and endoscopic sphincterotomy versus conservative treatment for acute pancreatitis due to gallstones.** Lancet. 1988;2:979-83. (Level I/II evidence). [View the reference](#)
15. Fan ST, Lai ECS, Mok FPT, et al. **Early treatment of acute biliary pancreatitis by endoscopic papillotomy.** N Engl J Med. 1993;328:228-32. (Level I/II evidence). [View the reference](#)
16. Neoptolemos J, Carr-Locke D, Leese T, et al. **Acute cholangitis in association with acute pancreatitis: incidence, clinical features and outcome in relation to ERCP and endoscopic sphincterotomy.** Br J Surg. 1987;74:1103-6. (Level III evidence)
17. Thoeni RF. **The revised Atlanta classification of acute pancreatitis: its importance for the radiologist and its effect on treatment.** Radiology. 2012;262(3):751-64. (Review article)
18. Bharwani N, Patel S, Prabhudesai S, Fotheringham T, Power N. **Acute pancreatitis: the role of imaging in diagnosis and management.** Clin Radiol. 2011;66(2):164-75. (Review article)
19. Marshall JC, Cook DJ, Christou NV, et al. **Multiple organ dysfunction score: a reliable descriptor of a complex clinical outcome.** Crit Care Med. 1995;23(10):1638-52. (Level I/II evidence)
20. Vincent JL, Moreno R, Takala J, et al. **The SOFA (sepsis-related organ failure assessment) score to describe organ dysfunction/failure. On behalf of the working group on sepsis-related problems of the European society of intensive care medicine.** Intensive Care Med. 1996;22(7):707-10. (Level II evidence)
21. De Campos T, Cerqueira C, Kuryura L, et al. **Morbimortality indicators in severe acute pancreatitis.** JOP. 2008;9(6):690-7. (Level III evidence)
22. Ranson JH, Rifkind KM, Roses DF, et al. **Prognostic signs and the role of operative management in acute pancreatitis.** Surg Gynecol Obstet. 1974;139(1):69-81. (Level III evidence)
23. Rosso E, Alexakis N, Ghaneh P, et al. **Pancreatic Pseudocyst in Chronic Pancreatitis: Endoscopic and Surgical Treatment.** Dig Surg. 2003;20:397-406. (Review article)
24. Buchler MW, Gloor B, Muller CA, Friess H, Seiler CA, Uhl W. **Acute necrotizing pancreatitis: treatment strategy according to the status of infection.** Ann Surg. 2000;232:619-26. (Level II evidence)
25. Uhl W, Warshaw A, Imrie C, et al. **IAP guidelines for the surgical management of acute pancreatitis.** Pancreatolgy. 2002;2:565-73. (Evidence-based guidelines). [View the reference](#)
26. Mithofer K, Mueller PK, Warshaw AI. **Interventional and surgical treatment of pancreatic abscess.** World J Surg. 1997;21:162. (Review article)
27. VanSonnenberg E, Wittich GR, Casola G, et al. **Percutaneous drainage of infected and non infected pancreatic pseudocysts: experience in 101 cases.** Radiology. 1989;170:757. (Level III evidence)
28. VanSonnenberg E, Wittich GR, Chon KS, et al. **Percutaneous radiologic drainage of**





- pancreatic abscesses.** AJR Am J Roentgenol. 1997;168:979. (Level III evidence)
29. Balthazar EJ. **Complications of acute pancreatitis: clinical and CT evaluation.** Radiol Clin North Am. 2002;40:1211-27. (Review article)
  30. Trout AT, Elsayes KM, Ellis JH, Francis IR. **Imaging of acute pancreatitis: prognostic value of computed tomographic findings.** J Comput Assist Tomogr. 2010;34(4):485-95. (Review article)
  31. Chang JH. Lee IS. Lim YS. Jung SH. Paik CN. Kim HK. Kim TH. Kim CW. Han SW. Choi MG. Jung IS. **Role of magnetic resonance cholangiopancreatography for choledocholithiasis: analysis of patients with negative MRCP.** Scand J Gastroenterol. 2012;47(2):217-24. (Level III evidence)
  32. Rifkind KM, Lawrence LR, Ranson JHC. **Initial roentenographic sign in acute pancreatitis: a study of findings in 73 patients.** N Y State J Med. 1976;76:1968. (Level II /III evidence)
  33. Siegelman SS, Copeland BE, Saba GP, Cameron JL, Sanders RC, Zerhouni EA. **CT of fluid collections associated with pancreatitis.** AJR Am J Roentgenol. 1980;134:1121-32. (Level III evidence)

## Further Reading

1. Piironen A. **Severe acute pancreatitis: contrast-enhanced CT and MRI features.** Abdom Imaging. 2001;26:225-33. (Review article)
2. Steinberg W, Tenner S. **Acute pancreatitis.** N Engl J Med. 1994;330:1198-1210. (Review article)

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