

# North American Society for Pediatric Gastroenterology, Hepatology and Nutrition and the Society for Pediatric Radiology Joint Position Paper on Noninvasive Imaging of Pediatric Pancreatitis: Literature Summary and Recommendations

<sup>\*†‡</sup>Andrew T. Trout, <sup>§</sup>Sudha A. Anupindi, <sup>||</sup>A. Jay Freeman, <sup>¶</sup>Jorge Alberto Macias-Flores, <sup>#</sup>J. Andres Martinez, <sup>\*\*</sup>Kalyan R. Parashette, <sup>††</sup>Uzma Shah, <sup>‡‡</sup>Judy H. Squires, <sup>§§</sup>Veronique D. Morinville, <sup>||||</sup>Sohail Z. Husain, and <sup>‡</sup>Maisam Abu-El-Hajja

## ABSTRACT

The reported incidence of pediatric pancreatitis is increasing. Noninvasive imaging, including ultrasound computed tomography (CT), and magnetic resonance imaging (MRI), play important roles in the diagnosis, staging, follow-up, and management of pancreatitis in children. In this position paper, generated by members of the Pancreas Committee of the North American Society for Pediatric Gastroenterology, Hepatology and Nutrition (NASPGHAN) and the Abdominal Imaging Committee of The Society for Pediatric Radiology (SPR), we review the roles of noninvasive imaging in pediatric acute, acute recurrent, and chronic pancreatitis. We discuss available evidence related to noninvasive imaging, highlighting evidence specific to pediatric populations, and we make joint recommendations for use of noninvasive imaging. Further, we highlight the need for research to define the performance and role of noninvasive imaging in pediatric pancreatitis.

**Key Words:** computed tomography, magnetic resonance imaging, radiography, ultrasound

An infographic is available for this article at: <http://links.bww.com/MPG/C9>.

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Recognition and the reported incidence of pancreatitis in children are increasing, with incidence currently estimated to be 1 in 10,000 for acute pancreatitis (AP) and 2 in 100,000 for chronic pancreatitis (CP) (1–3). Given that data related to imaging of pediatric pancreatitis are sparse, pediatric recommendations for imaging are largely based on adult data. The purposes of this document, which is jointly endorsed by the North American Society

## What Is Known

- The reported incidence of pancreatitis in children is increasing, currently estimated to be 1 in 10,000 for acute pancreatitis and 2 in 100,000 for chronic pancreatitis.
- The roles of imaging in acute pancreatitis are to: identify findings of acute pancreatitis at diagnosis; assess for local complications; identify potential etiologies of acute pancreatitis; monitor the evolution of local complications; and plan and guide interventions.
- The roles of imaging in chronic pancreatitis are to: contribute to/establish the initial diagnosis of chronic pancreatitis; stage and monitor disease, including complications; assess for superimposed acute pancreatitis; identify potential etiologies of chronic pancreatitis; characterize secretory (exocrine) function; and plan for surgical intervention.

## What Is New

- Little information is available regarding the optimal imaging strategy for pediatric pancreatitis.
- Current methods to prognosticate and predict pancreatitis severity and disease progression are inadequate.
- It is currently not possible to identify minimal change or early chronic pancreatitis in pediatric patients.

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From the <sup>\*</sup>Department of Radiology, Cincinnati Children's Hospital Medical Center, the <sup>†</sup>Department of Radiology, the <sup>‡</sup>Department of Pediatrics, University of Cincinnati College of Medicine, Cincinnati, OH, the <sup>§</sup>Department of Radiology, The Children's Hospital of Philadelphia, University of Pennsylvania Perelman School of Medicine, Philadelphia, PA, the <sup>||</sup>Department of Pediatrics, Emory University, Children's Healthcare of Atlanta, Atlanta, GA, the <sup>¶</sup>División de Gastroenterología Pediátrica, Hospital Infantil de Especialidades de Chihuahua, Mexico, the <sup>#</sup>Division of Pediatric Gastroenterology, Hepatology and Nutrition, Vanderbilt University Medical Center, Nashville, TN, the <sup>\*\*</sup>Department of Pediatrics, Loma Linda University School of Medicine, Loma Linda, CA,

the <sup>††</sup>Pediatric Gastroenterology, Hepatology and Nutrition, Massachusetts General Hospital for Children, Harvard Medical School, Boston, MA, the <sup>‡‡</sup>Department of Radiology, University of Pittsburgh Medical Center, Department of Radiology, UPMC Children's Hospital of Pittsburgh, Pittsburgh, PA, the <sup>§§</sup>Department of Pediatrics, Montreal Children's Hospital, McGill University, Montreal, Quebec, Canada, the <sup>||||</sup>Division of Gastroenterology, Hepatology, and Nutrition, Department of Pediatrics, Stanford School of Medicine, Stanford, CA, and the <sup>¶¶</sup>Division of Gastroenterology, Hepatology and Nutrition, Cincinnati Children's Hospital Medical Center, Cincinnati, OH.

for Pediatric Gastroenterology, Hepatology and Nutrition (NASPGHAN) and The Society for Pediatric Radiology (SPR), are to: summarize existing literature and experience regarding imaging of pediatric AP and CP; provide recommendations for the role of imaging in the diagnosis and management of pediatric pancreatitis; and identify knowledge gaps and areas for future study. This document focuses on noninvasive imaging and will not emphasize endoscopic ultrasound (EUS) or endoscopic retrograde cholangiopancreatography (ERCP), which also contribute to the diagnosis and management of these diseases (4–6).

## METHODS

This document was generated through collaboration between members of the NASPGHAN Pancreas Committee and The SPR Abdominal Imaging Committee with concept approval by the boards of both organizations before generation of the document. Members of each committee volunteered to participate with 8 gastroenterologists and 3 radiologists contributing to drafting the document, grading available evidence, and generating and voting on recommendations. One radiologist (S.A.A., J.H.S., or A.T.T.) and 2 gastroenterologists (A.J.F., J.A.M-F., J.A.M., V.D.M., K.R.P., or U.S.) were primarily responsible for each section of the text. One radiologist (A.T.T.) and 1 gastroenterologist (M.A-E-H) led the project, providing global oversight and structure. Although a systematic literature review was not performed, contributing authors reviewed pertinent literature through April 2019 for their respective section(s). Project leads confirmed inclusion of relevant pediatric articles by performing a PubMed search in July 2019 for the following MeSH terms: “Pancreatitis/diagnostic imaging,” “Tomography, X-Ray Computed/methods,” “Ultrasonography/methods,” “Magnetic Resonance Imaging/methods,” limited by the PubMed “Child: birth-18 years” filter.

On the basis of a complete draft of the document, project leads generated or extracted specific recommendations from the text. Project leads also assigned classifications for the recommendations based on a modified version of the GRADE system, applying the criteria for studies on diagnostic accuracy (7). Grades incorporated a score of recommendation strength (1 = Strong, 2 = Weak) and evidence quality (A = high quality, B = moderate quality, C = low quality). The criteria for studies of diagnostic accuracy consider cross sectional or cohort studies with comparison to an appropriate

reference standard to reflect high-quality evidence in lieu of randomized controlled trials. Of note, the process of applying the modified GRADE system for this document did not utilize independent evaluators to review the recommendations and supporting evidence, and a formal GRADE report of the literature was not created.

Grades assigned by project leads were preliminarily affirmed by the authors of each manuscript section. The full draft manuscript was then reviewed and approved by all members of the project team who provided suggested edits and commented on the proposed recommendations and GRADE classifications. All members of the project team had reviewed the modified GRADE methodology before affirming and commenting on GRADE classifications. After final edits, all members of the project team voted on the recommendations via a survey built in REDCap, assigning a 5-point Likert score (5, strongly agree; 4, agree; 3, neutral; 2, disagree; 1, strongly disagree) to each recommendation (8,9). Voting results were submitted to a research coordinator at Cincinnati Children’s Hospital who was not involved in generation of this document or the recommendations. A priori, a minimum 75% frequency of “strongly agree” or “agree” ratings was defined as the threshold required to be considered consensus. Other references that have used this system include a position paper on Nutritional Considerations in Pediatric Pancreatitis by Abu-El-Haija et al (10).

Although high-quality literature to support and direct the use of specific imaging modalities in pediatric pancreatitis is limited, recommendations are based on expert opinion informed by adult literature and the pediatric literature that exists. Comments are included wherever needed to explain recommendations.

Subsequent to study team affirmation of recommendations, a final version of the document was submitted to committees of NASPGHAN and The SPR for review and comment. Comments provided by the reviewing organizations were reviewed by the project leads and incorporated as appropriate in the final document, which was approved by the NASPGHAN Council and The SPR Board.

## BACKGROUND

### Acute Pancreatitis

The diagnosis of AP in children has been defined as the presence of at least 2 of the following: abdominal symptoms

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Address correspondence and reprint requests to Andrew T. Trout, MD, Associate Professor of Radiology and Pediatrics, Department of Radiology, Cincinnati Children’s Hospital Medical Center, MLC 5031, 3333 Burnet Avenue, Cincinnati, OH 45229-3026 (e-mail: andrew.trout@cchmc.org).

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TABLE 1. Categorical etiologies of pediatric pancreatitis

Category	Examples*
Obstructive	Biliary stone(s) Pancreatic duct anomalies (eg, complete divisum, annular pancreas) Choledochal cyst Tumor
Genetic	Cationic trypsinogen (PRSS1) Serine protease inhibitor Kazal type 1 (SPINK1) Cystic fibrosis transmembrane regulator (CFTR) Chymotrypsin C (CTRC) Calcium-sensing receptor (CASR) Carboxypeptidase 1 (CPA1) Carboxyl ester lipase (CEL)
Medication related	Anti-epileptics Asparaginase
Trauma	
Systemic illness	Infections (eg, mumps, herpes virus) Inflammatory disease (eg, hemolytic uremic syndrome, systemic lupus erythematosus)
Metabolic	Hypertriglyceridemia Hypercalcemia Kidney disease
Autoimmune	
Substance/toxic	Alcohol Smoking

Any of the listed etiologies can contribute to a single episode of acute pancreatitis (AP). Genetic and obstructive causes become leading etiologies in ARP and CP. Modified from Uc and Husain (14).

\*Examples are not meant to be exhaustive lists.

consistent with AP; serum amylase or lipase values  $\geq 3$  times the upper normal level; and imaging findings consistent with AP (11). Biliary causes, anatomic causes, and genetic pancreatitis represent the most common etiologies of AP in children, but up to 20% of cases remain idiopathic (Table 1) (12–14). Severity staging of AP has only recently been defined for pediatrics (Table 2), and is structured to classify which children are most at risk of complicated courses (15,16). Mild AP is defined as AP without organ failure or local or systemic complication, and usually resolves within 1 week. Moderately severe AP is defined as either the presence of transient ( $\leq 48$  hours) organ failure, the presence of local complications, or the exacerbation of comorbid disease. Severe AP (SAP) is defined by organ failure lasting longer than 48 hours. Moderately severe or

TABLE 2. North American Society for Pediatric Gastroenterology, Hepatology and Nutrition acute pancreatitis working group classification of pediatric acute pancreatitis severity

Severity	Findings
Mild	No organ failure/dysfunction* No local or systemic complication(s) <sup>†,‡</sup>
Moderately severe	EITHER Development of transient ( $\leq 48$ hours) organ failure/dysfunction* OR Local or systemic complications <sup>†,‡</sup>
Severe	Organ failure/dysfunction <sup>†</sup> lasting $> 48$ hours

Modified from Abu-El-Haija et al (15).

\*Organ failure/dysfunction = defined according to the International Pediatric Sepsis Consensus (15).

<sup>†</sup>Local complications = pancreatic or peripancreatic necrosis and/or fluid collections.

<sup>‡</sup>Systemic complications = exacerbation of co-morbid disease.

SAP have been reported to occur in approximately 13% to 30% of children with AP (15,17). To date, pediatric-specific risk factors for SAP remain unclear.

In general terms, the roles of imaging in AP are to: identify findings of AP at diagnosis; assess for local complications; identify potential etiologies of AP; monitor the evolution of local complications, and plan and guide interventions. Largely on the basis of the lack of ionizing radiation, transabdominal ultrasound is favored as the initial imaging modality for the diagnosis of AP in children, whereas computed tomography (CT) and/or magnetic resonance imaging (MRI) are reserved for more complicated cases or to answer specific clinical questions (4).

## Acute Recurrent Pancreatitis

Acute recurrent pancreatitis (ARP) has been defined as 2 distinct attacks of AP with more than 1 month pain-free interval between attacks, or with normalization of pancreatic enzymes and complete resolution of pain regardless of interval between episodes (11). ARP is believed to develop in 15% to 35% of pediatric patients who suffer from an initial event of AP (3,17,18). In 1 study, the majority of patients who developed ARP had a second attack within 5 months after their initial episode (19). Genetic mutations represent the most common risk factor for the development of ARP with almost 50% of patients carrying a mutation in CFTR, PRSS1, CTRC or SPINK1 in 1 series. Additionally, approximately 1/3 had a pancreatic duct obstructive risk factor, such as pancreas divisum (3,18). It should be noted, however, that pancreas divisum alone does not necessarily cause pancreatitis.

In general terms, the roles of imaging in ARP are to: confirm attacks of AP; assess for local complications; identify potential etiologies of ARP; monitor the evolution of complications; plan and guide interventions; and assess for imaging findings suggestive of progression to CP. There are no robust data to define an optimal imaging modality or strategy for ARP though most favor MRI because of its ability to optimally assess both parenchyma and duct (11).

## Chronic Pancreatitis

CP results from progressive inflammation that results in fibrotic replacement of pancreatic parenchyma, and eventually, exocrine and endocrine dysfunction. CP in children has been defined as imaging findings of CP combined with at least 1 of the following: abdominal pain consistent with pancreatic origin; exocrine pancreatic insufficiency (EPI); or pancreatic endocrine insufficiency (11). Less frequently, surgical or biopsy specimens consistent with CP are obtained. In children, genetic factors (seen in up to 73% of patients) are the most prevalent etiology of CP followed by obstructive causes, similar to those seen in ARP (3,18).

In general terms, the roles of imaging in CP are to: contribute to/establish the initial diagnosis of CP; stage and monitor disease, including complications; assess for superimposed AP; identify potential etiologies of CP; identify findings that might herald endocrine or exocrine dysfunction; characterize secretory (exocrine) function; and plan for intervention. Although findings of CP may be identified on ultrasound or CT, MRI/magnetic resonance cholangiopancreatography (MRI/MRCP) is favored for the diagnosis and characterization of CP given its superiority in visualizing parenchymal and duct changes (20).

## IMAGING TECHNIQUES AND GENERALITIES

### Transabdominal Ultrasound

Current consensus recommendations favor transabdominal ultrasound as the initial imaging examination to evaluate suspected

AP as it is widely available, the examination can be performed with minimal patient preparation, and the examination does not use sedation, contrast material, or ionizing radiation (4,21–23). The major advantages of ultrasound over other imaging modalities are availability and portability. The latter allows bedside imaging of critically ill and difficult-to-transport patients. Ultrasound can, however, be limited in patients with large body habitus, or with excessive bowel gas. In addition, ultrasound may underestimate or poorly delineate the extent of extrapancreatic sequelae of AP (24).

A right upper quadrant ultrasound examination typically does not provide full imaging of the pancreas and will not evaluate other areas of the abdomen for potential pancreatitis complications. A complete abdominal ultrasound allows evaluation of both upper quadrants and the pancreas, and may include assessment of the lower quadrants for fluid.

Ultrasound examinations are ideally performed fasting ( $\geq 4$  hours) to reduce bowel gas that can obscure the pancreas and to distend the gallbladder. When the patient is able, drinking water immediately prior to the examination to distend the stomach with fluid and displace gastric air may provide an improved acoustic window for imaging the pancreas. A complete ultrasound examination should assess pancreatic size, contour, echogenicity, pancreatic duct diameter (and for duct filling defects), and should assess for peripancreatic edema and pancreatic or peripancreatic fluid collections (25). In addition, the gallbladder should be assessed for calculi (an etiology of pancreatitis), and the biliary tree should be assessed for dilation and calculi (4,22,26). Color Doppler along with gray-scale imaging can evaluate the peripancreatic vascular structures for complications, such as splenic vein or portal vein thrombosis and can assess vascularity of the pancreas.

Contrast-enhanced ultrasound (CEUS), which involves the intravenous administration of contrast material consisting of microbubbles, has been explored for assessment of both AP and CP but is not yet accepted as standard of care, so its role in pediatric pancreatitis remains to be defined (27,28).

## Computed Tomography

The major advantage of CT versus other noninvasive imaging modalities is that the examinations are short and can generally be achieved without sedation or anesthesia. For this reason, CT is the modality of choice for acute assessment of traumatic injury of the pancreas. Body habitus and air-filled bowel loops are not limiting factors for CT (compared with ultrasound).

CT for pediatric pancreatitis should utilize intravenous (IV) contrast material, which optimizes assessment of the solid organs and vasculature. CT with IV contrast material can be performed as a single (arterial or portal venous phase) or multiphase examination. When performed as a single-phase examination for pancreatic indications, portal venous phase imaging is most common. Intravenous iodinated contrast material carries a very low risk of allergic-like reactions. Risk of exacerbation of renal dysfunction in children with estimated glomerular filtration rate less than  $60 \text{ mL/min/1.73 m}^2$  should be balanced with the potential benefit of the examination (29).

Use of oral contrast material for CT in pancreatitis is inconsistent; specific recommendations do not exist for children. In adults, oral water is recommended for imaging of pancreatitis (20). A potential benefit of positive oral contrast, which is high in attenuation, is distinguishing fluid collections from bowel loops but oral contrast material can be difficult for pediatric or acutely ill patients to consume and prolongs the preparatory phase of the CT examination.

Limiting CT to the abdomen only is discouraged. CT limited to the abdomen allows assessment of the pancreas, adjacent vessels, and surrounding structures but does not allow assessment for extension of complications (eg, fluid collections) into the lower

abdomen and pelvis. CT of the abdomen and pelvis allows assessment of the pancreas and the full extent of associated complications.

## Magnetic Resonance Imaging and Magnetic Resonance Cholangiopancreatography

MRI has the best soft tissue contrast of available cross-sectional imaging modalities. This soft tissue contrast optimizes parenchymal characterization and visualization and characterization of ducts and fluid collections. MRCP, which is sometimes arbitrarily distinguished from other MRI examinations, simply reflects a type of MRI sequence that utilizes heavy T2-weighting to accentuate fluid-filled structures including the pancreatic and biliary ducts. Like CT, body habitus and air-filled bowel loops are not limitations for MRI. Potential need for sedation or general anesthesia in the pediatric population to accomplish relatively long examinations is the primary disadvantage of MRI.

Given its superior soft tissue contrast, MRI is the preferred imaging modality for ARP, CP, and autoimmune pancreatitis where characterization of both the pancreatic parenchyma and duct is important (30). Protocol guidelines exist for adults but have not been formalized for children (20). An MRCP sequence should be included in most pancreatic MRI examinations. IV gadolinium-based contrast material can be useful for characterization of the vasculature and for diagnosis of autoimmune pancreatitis but is not required for routine assessment of CP. IV contrast material is not required to acquire an MRCP sequence and hepatobiliary contrast material can compromise MRCP sequences.

Use of intravenous secretin as an adjunctive medication may improve visualization of the pancreatic duct and allows assessment of exocrine function (31–39). According to adult studies, secretin improves visualization of the pancreatic duct by MRCP with a higher diagnostic accuracy of detecting pancreas divisum (34–37). The reported sensitivity and specificity of diagnosing pancreas divisum with secretin-MRCP falls in the range of 73% to 100% and 97% to 100%, respectively (40).

## Timing of Imaging

Optimal timing of imaging, particularly relative to an attack of AP, depends on the unique patient situation. Few studies have published pediatric data specific to this question, but adult data suggest that imaging within the first 48 hours of an attack of AP infrequently alters management and poorly predicts the severity of organ failure (41). However, if imaging is necessary to make a diagnosis or to manage a patient, it should not be deferred. In patients with suspected or known CP where imaging confirmation of findings of CP is needed, imaging when the patient does not have superimposed AP is preferred to prevent obscuration of findings by acute inflammation.

## General Imaging Technique Summary Statements and Recommendations

1. CT should be performed with intravenous contrast material as a single portal venous phase examination unless specific arterial detail is needed (Table 3).  
GRADE: 1C, agreement 100% (11/11; 6, strongly agree; 5, agree; average score = 4.5)
2. When imaging with MRI, intravenous contrast material is not always needed but contributes to the diagnosis and definition of necrosis, assessment of the vasculature, and the diagnosis of autoimmune pancreatitis.  
GRADE: 2C, agreement 100% (11/11; 3, strongly agree; 8, agree; average score = 4.3)

TABLE 3. Summary statements and recommendations

Statement number	Statement/recommendation	Grade (7)	Agreement	Average score
<b>General imaging</b>				
1	CT should be performed with intravenous contrast material as a single portal venous phase examination unless specific arterial detail is needed	1C	100% (11/11)	4.5
2	When imaging with MRI, intravenous contrast material is not always needed but contributes to the diagnosis and definition of necrosis, assessment of the vasculature and the diagnosis of autoimmune pancreatitis	2C	100% (11/11)	4.3
<b>Acute pancreatitis</b>				
3	Transabdominal ultrasound is recommended as a first-line noninvasive imaging modality for suspected AP	1B	91% (10/11)	4.7
4	If ultrasound is negative for AP and an imaging diagnosis of AP is needed, either CT or MRI is recommended	1B	100% (11/11)	4.6
5	CT or MRI is recommended for identification and assessment of known or suspected complications of AP	1C	91% (10/11)	4.5
6	Ultrasound can be used to follow known AP fluid collections for resolution or progression (changes in size)	2C	82% (9/11)	4.3
7	CT or MRI should be used to characterize the degree of organization of collections before intervention	1C	100% (11/11)	4.5
<b>Acute recurrent pancreatitis</b>				
8	MRI is recommended to identify structural or obstructive causes for ARP	1B	100% (11/11)	4.8
9	When clinically indicated, MRI is recommended to follow children with ARP and to assess for progression to CP	1C	100% (11/11)	4.6
10	In a child who requires sedation for imaging, it is reasonable to alternate MRI with ultrasound or CT for serial monitoring of ARP	2C	82% (9/11)	4
<b>Chronic pancreatitis</b>				
11	MRI is the recommended modality for imaging of suspected CP	1C	91% (10/11)	4.6
12	When imaging is needed to assess a suspected or known episode of AP in a child with CP, transabdominal ultrasound is the preferred first-line imaging modality	1B	91% (10/11)	4.5
13	If ultrasound is negative for AP in a child with CP and an imaging diagnosis of AP is needed, either CT or MRI are recommended	1B	100% (11/11)	4.5
14	CT or MRI are recommended for planning of endoscopic or surgical interventions in a patient with known CP	2C	100% (11/11)	4.5
15	MRI is recommended for clinically indicated serial imaging of CP	1B	100% (11/11)	4.8

AP = acute pancreatitis; CP = chronic pancreatitis; CT = computed tomography; MRI = magnetic resonance imaging.

## IMAGING OF ACUTE PANCREATITIS

### Purpose/ Indication/Rationale for Imaging in Acute Pancreatitis

Imaging in the context of suspected or known AP serves the multiple purposes previously described. At diagnosis, it is particularly important to identify gallstones or biliary obstruction (usually because of choledocholithiasis) as etiologies of AP. These entities can be urgently addressed with endoscopic and/or surgical intervention (4,42–44). Identification of local complications of AP including necrosis, acute fluid collections, venous stenosis/thrombosis or arterial aneurysms, and hemorrhage has relevance for clinical staging of attack severity (15,45). In adults, severity scoring can help prognosticate and triage appropriate management but there is currently no consensus imaging severity staging/scoring system in children. In a clinical report, the Pancreas Committee of NASPGHAN has proposed stratifying AP in children as mild, moderately severe, or severe utilizing a combination of clinical and imaging criteria (Table 2) (15).

The most common complication of AP is the development of acute peripancreatic fluid collections and pseudocysts (13%–15%) (46,47). The frequency of these fluid collections secondary to AP has been reported in pediatric studies to be between 8% and 41% (48). Pancreatic and peripancreatic necrosis (sterile or infected) occur less commonly (47,49). As an attack of AP progresses, imaging serves to assess the evolution and maturity of fluid collections (necrotic or simple) to help define the timing of interventions. Fluid collections generally need to have a well-defined

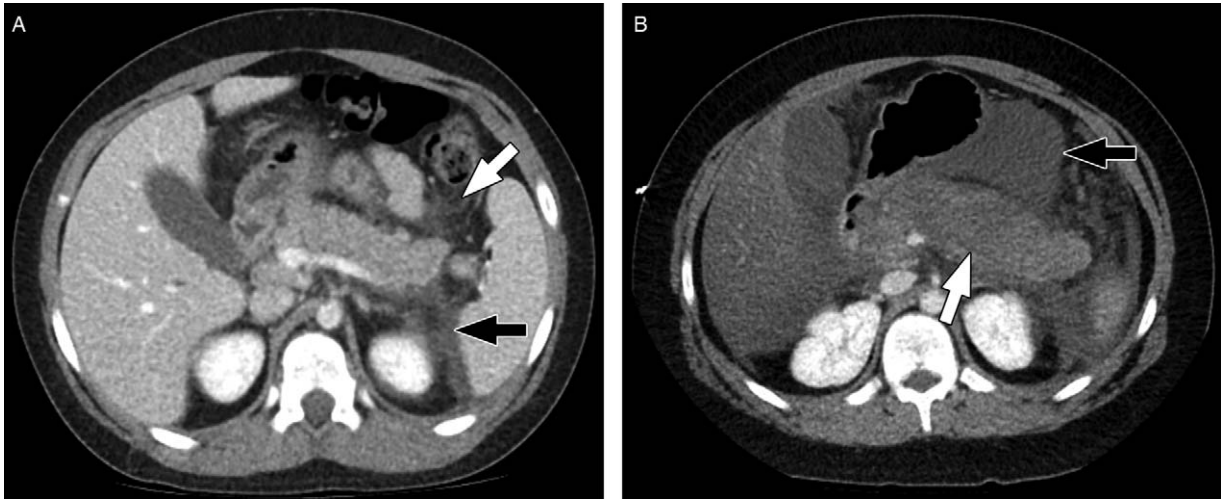
wall to be amenable to intervention, particularly endoscopic intervention.

### Imaging Findings of Acute Pancreatitis

Two forms of AP are distinguishable by imaging: interstitial edematous and necrotizing pancreatitis, with the former more common (Fig. 1). The imaging features of acute interstitial pancreatitis are similar on ultrasound, CT, and MRI (Table 4). In the very early stages of AP, ultrasound and CT may show no abnormal findings, as laboratory abnormalities often precede imaging findings of AP (22). This does not, however, seem to apply to MRI, which has higher soft tissue contrast resolution (50).

Findings of necrotizing pancreatitis depend on the stage of necrosis. Early in the course of necrotizing pancreatitis, there is decreased or absent vascularity/perfusion of the gland with hypoenhancement following contrast administration. As necrosis evolves, the gland and peripancreatic tissues may be replaced by necrotic collections. By ultrasound and MRI, these collections will contain debris. By CT, the collections may deceptively appear simpler. Superinfection of these collections may occur, with air in the collection(s) being a specific, but not sensitive, finding (50).

Fluid collections associated with AP have specific definitions, which were updated in the 2012 Revised Atlanta Classification (Table 5) (41). The definitions apply to adults but have been extrapolated to children. One important nuance of the Revised Atlanta Criteria is that if acute necrotic collections organize, these are by definition walled off necrosis (not pseudocysts) regardless of



**FIGURE 1.** Examples of interstitial edematous acute pancreatitis and necrotizing acute pancreatitis in 2 different patients. (A) Axial image from a CT performed with intravenous contrast material in a 10-year-old boy with interstitial edematous pancreas shows a swollen but homogeneously enhancing pancreas with peripancreatic stranding (white arrow) and with an acute peripancreatic fluid collection. (B) Axial image from a CT performed with intravenous contrast material in a 10-year-old boy with necrotizing pancreatitis shows a swollen pancreas with a large area of absent enhancement (white arrow) indicative of necrosis. There is more normal enhancement of the pancreatic tail. An acute necrotic collection is also present in the lesser sac (black arrow). CT = computed tomography.

how simple they appear. Pseudocysts occur only in the context of acute interstitial edematous pancreatitis, or, rarely, in the case of disconnected duct, due to prior necrosis or intervention.

**Diagnostic Performance of Imaging Modalities in Acute Pancreatitis**

Little data are available on the diagnostic performance of ultrasound, CT, and MRI for the assessment of AP in children.

**Transabdominal Ultrasound**

Pediatric-specific data regarding the ability of transabdominal ultrasound to detect gallstones as an etiology for AP are not available. Adult data have shown ultrasound to be approximately 99% sensitive for gallstones in the gallbladder (51). The sensitivity of abdominal ultrasound in detecting AP, based on adult data, is reported to be as high as 79% (52). The sensitivity of ultrasound in diagnosing AP in children has not been well-defined. In a study of 112 children with AP, 75% (n = 84) had

**TABLE 4.** Imaging findings of acute pancreatitis

Imaging modality	Findings in interstitial edematous pancreatitis	Findings in necrotizing pancreatitis
Ultrasound	Normal (early stage) Enlarged pancreas, focal or diffuse Hypo- or hyperechoic parenchyma Ill-defined borders Dilated pancreatic duct Thickened, echogenic peripancreatic fat Peripancreatic fluid	Avascular areas of parenchyma (Doppler or CEUS) Fluid collections replacing parenchyma
CT	Hypoattenuating areas in parenchyma Enlarged pancreas, focal or diffuse Ill-defined borders Peripancreatic edema Peripancreatic fluid Fluid elsewhere in abdomen and pelvis	Absent enhancement of parenchyma Intra and extrapancreatic collections (+/- debris)
MRI/MRCP	Decreased T1W signal Increased T2W signal Hypoenhancing parenchyma Enlarged pancreas, focal or diffuse Dilated pancreatic duct Peripancreatic edema Peripancreatic fluid Fluid elsewhere in the abdomen and pelvis	Absent enhancement of parenchyma Intra and extrapancreatic fluid collections containing debris High T1W signal in pancreas or collections (hemorrhage)

CEUS = contrast-enhanced ultrasound; CT = computed tomography; MRCP = magnetic resonance cholangiopancreatography; MRI = magnetic resonance imaging.

TABLE 5. Definitions of pancreatic and peripancreatic collections based on the Revised Atlanta Classification

Fluid collection	Morphologic features
Acute peripancreatic fluid collection <i>Develops in setting of interstitial edematous pancreatitis</i>	Peripancreatic fluid associated with interstitial edematous pancreatitis No associated necrosis Applies only to fluid seen within the first 4 weeks after onset of interstitial edematous pancreatitis and without features of a pseudocyst Contrast-enhanced computed tomography criteria: Homogeneous collection with fluid density Defined by normal fascial planes No definable wall encapsulating the collection Adjacent to pancreas (no intrapancreatic extension)
Acute necrotic collection <i>Develops in setting of necrotizing pancreatitis</i>	Collection containing variable amounts of both fluid and necrotic debris Associated with necrotizing pancreatitis/peripancreatitis Contrast-enhanced computed tomography criteria: Heterogeneous and/or nonliquid density (some appear homogeneous early in their course) No definable wall encapsulating the collection Location—intrapaneatic and/or extrapancreatic
Pancreatic pseudocyst	Encapsulated collection of simple fluid with a well-defined inflammatory wall Usually occurs >4 weeks after onset of interstitial edematous pancreatitis (though best defined by maturity of wall rather than time course) Following necrosectomy, a completely debrided necrotic collection can be considered a pseudocyst Contrast-enhanced computed tomography criteria: Well circumscribed, usually round or oval Homogeneous fluid density No nonliquid component Well defined wall (ie, completely encapsulated)
Walled-off necrosis (WON)	Encapsulated collection of pancreatic and/or peripancreatic necrosis that has developed a well-defined inflammatory wall Usually occurs >4 weeks after onset of necrotizing pancreatitis (though best defined by maturity of wall rather than time course) Contrast-enhanced computed tomography criteria: Heterogeneous with liquid and nonliquid material (some may appear homogeneous) Well defined wall (ie, completely encapsulated) Location—intrapaneatic and/or extrapancreatic

Adapted from Revised Atlanta Classification (41).

an ultrasound performed and, ultrasound was only 52% (95% confidence interval: 41%–63%) sensitive for AP diagnosed based on symptoms and serum enzymes (53). Prior studies have shown widely variable performance of ultrasound for diagnosis of AP. Benifla and Weizman (46) reported a diagnosis of AP by ultrasound in 81% of 589 children, but Coffey et al (54) reported ultrasound findings of AP in only 23% of 77 patients with AP diagnosed by elevated enzymes.

Chao et al and Siegel et al reported the most useful indicator of AP to be a dilated pancreatic duct. The sensitivity and specificity of a dilated pancreatic duct in children on ultrasound range between 78% to 83% and 87% to 92%, respectively, with positive-predictive value (PPV) of 86–91%, and negative-predictive value (NPV) of 75% to 84% (55,56).

### Computed Tomography

On the basis of adult literature, CT with IV contrast material is considered the imaging reference standard for AP (26,57,58). IV contrast material allows evaluation for necrosis, based on absent parenchymal enhancement, and optimizes identification and assessment of intra- or extra-pancreatic fluid collections. IV contrast material also allows evaluation of the peripancreatic vasculature to ensure patency and assess for pseudoaneurysm formation (22).

Other than for specific assessment of the arteries, adult data suggest that a single-phase portal venous phase examination is sufficient for assessment of AP (59).

Adult data suggest CT is more sensitive than ultrasound for AP, particularly for severe pancreatitis and acute necrosis (22). Diagnostic performance has not been specifically defined for children; however, Coffey et al (54) reported CT to show findings of AP in 62% of 42 patients with AP based on positive enzymes (vs 23% for ultrasound).

Compared with ultrasound, CT with intravenous contrast material provides improved characterization of the location and extent of fluid collections and abscesses, integrity of the splenic vein and portal system, and presence of parenchymal necrosis (43,52,60,61). A small surgical series (n=13) in adult patients showed CT to have 100% per patient sensitivity for necrosis (62). Of note, however, on a per-segment basis, CT was only approximately 64% sensitive, missing additional sites of necrosis in several patients (62). When infected necrosis is present, gas pockets are more readily visible on CT (Fig. 2) than on ultrasound. CT may, however, underestimate the complexity of fluid collections relative to ultrasound or MRI (Fig. 3) (63,64). For serial imaging of children with AP, the ionizing radiation associated with CT should be considered when selecting an imaging modality for follow-up.



**FIGURE 2.** A 5-year-old girl with infected pancreatic necrosis. Axial image from a CT performed with intravenous contrast material shows gas locules in the nonenhancing pancreas (arrows). No normal pancreas is visible and acute fluid is present in the abdomen. The patient also had acute renal cortical necrosis accounting for absent enhancement of the renal cortex. CT = computed tomography.

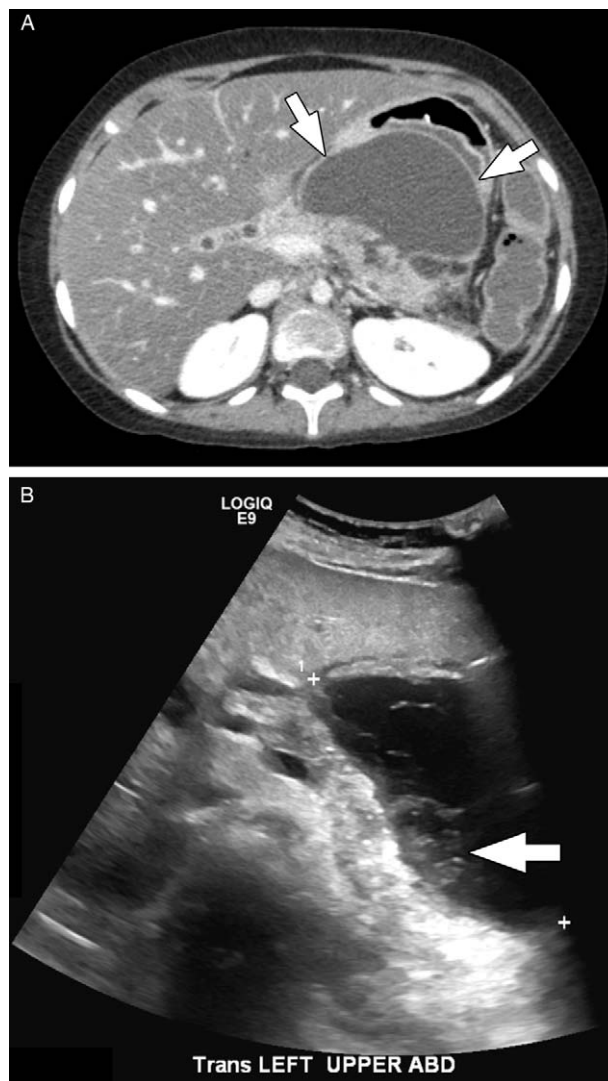
CT severity scoring indices (eg, CT severity index [CTSI], Balthazar score) were established in adult populations, but more recently have been applied to children (15,65). Similar to adult studies, the CTSI has been shown to be a better predictor of the severity of AP compared with clinical scores in children (65–67). A recent pediatric study applying the CTSI scores in 211 children with AP found the sensitivity and specificity of the CTSI in predicting a severe course of AP to be 81% and 76%, respectively, with positive-predictive and negative-predictive values of 62% and 90%, respectively (68). In this same pediatric cohort, the presence of necrosis in AP was associated with higher rate of major complications (68).

### **Magnetic Resonance Imaging/Magnetic Resonance Cholangiopancreatography**

Studies are lacking regarding the diagnostic performance of MRI in diagnosing AP in children, particularly compared with other imaging modalities. MRI can be used for assessment of AP, but because of the need for long periods of holding still may not be suitable for children, especially if critically ill. MRI may contribute to confirmation of an attack of AP or to identification/confirmation of acute duct obstruction (see below) but pancreatic edema because of an acute episode of pancreatitis can obscure pancreatic duct anomalies that may be relevant to the cause of pancreatitis.

The greater soft tissue contrast of MRI (vs CT) is advantageous when assessing the pancreatic parenchyma and biliary and pancreatic ducts and when characterizing fluid collections (22). Adult data suggest that MRI is more sensitive than CT for findings of AP including edema and hemorrhage with up to 15% to 30% of patients with a normal CT showing findings of AP on MRI (69–71). Adult data have also shown the diagnostic performance of MRI to be as good as CT for pancreatic necrosis (72).

MRI, particularly MRCP, has also been shown to be more sensitive than CT for biliary etiologies of pancreatitis (20). Specifically, in adults, MRCP has up to 100% sensitivity for pancreatic and biliary duct stones greater than 3 mm in size (73). MRCP can be

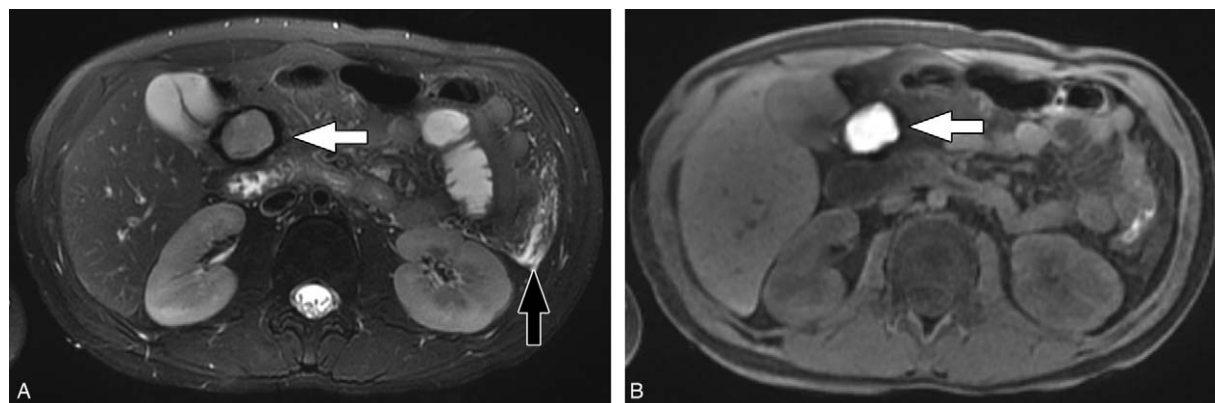


**FIGURE 3.** A 10-year-old girl with walled off pancreatic and peripancreatic necrosis. (A) Axial image from a CT performed with intravenous contrast material shows a walled off collection involving the body of the pancreas and the peripancreatic tissues in the lesser sac (arrows). Note how the content of the collection appears relatively simple by CT. (B) Transverse image from a transabdominal ultrasound performed the next day shows the same collection but with layering solid/semi-solid debris (arrow). CT = computed tomography.

particularly useful in the evaluation of choledocholithiasis when biliary duct dilatation is found on ultrasound without stone(s) visible in the duct(s) (74–78). Structural abnormalities of the pancreatic duct and parenchyma, such as pancreas divisum and an abnormal union of the pancreaticobiliary junction with a long common channel have also been associated with acute pancreatitis (79), and MRCP is the optimal noninvasive imaging modality for these entities.

MRI can help distinguish acute necrotic collections from acute peripancreatic fluid collections by identifying and characterizing the internal content of these collections (50). MRI is also superior to CT in detecting hemorrhage, which can be a complication of necrosis (80) (Fig. 4). In clinical practice, MRI is often used for assessment and monitoring of late complications of AP, such as





**FIGURE 4.** A 9-year-old girl with acute on chronic pancreatitis with a hemorrhagic peripancreatic collection. (A) Axial T2-weighted, fat-saturated MR image shows a fluid collection above the head of the pancreas and adjacent to the gallbladder (white arrow). The collection shows peripheral susceptibility artifact related to evolving blood products. Edema in the left hemiabdomen (black arrow) reflects acute pancreatitis. (B) Axial T1-weighted, fat-saturated MR image shows the content of the collection (white arrow) to be T1-weighted hyperintense compatible with blood products.

fluid collections, to time and guide therapeutic interventions (22,81). This, in part, not only capitalizes on the high soft tissue contrast but also on the fact that MRI does not involve exposure to ionizing radiation, and is thus, more acceptable for serial examinations.

## Acute Pancreatitis Summary Statements and Recommendations

### Initial Diagnosis

3. Transabdominal ultrasound is recommended as a first-line noninvasive imaging modality for suspected AP (Table 3).
  - a. This recommendation reflects the availability and portability of ultrasound and the role of ultrasound in identifying biliary causes of AP.
  - b. Note: A negative ultrasound does not exclude AP (low-to-moderate sensitivity).

GRADE 1B, agreement 91% (10/11; 9 strongly agree, 1 agree, 1 neutral, average score = 4.7)
4. If ultrasound is negative for AP and an imaging diagnosis of AP is needed, either CT or MRI is recommended.
  - a. This recommendation reflects the only moderate sensitivity of ultrasound and the greater sensitivity of CT and MRI.

GRADE 1B, agreement 100% (11/11, 7 strongly agree, 4 agree, average score = 4.6)

### Suspected Complications of Acute Pancreatitis

5. CT or MRI is recommended for identification and assessment of known or suspected complications of AP.
  - a. Note: CT has the potential to underestimate the complexity of fluid collections.

GRADE 1C, agreement 91% (10/11, 6 strongly agree, 4 agree, 1 neutral, average score = 4.5)

### Follow-up of Known Complications, With or Without Planning for Intervention

6. Ultrasound can be used to follow known AP fluid collections for resolution or progression (changes in size).

GRADE 2C, agreement 82% (9/11, 6 strongly agree, 3 agree, 1 neutral, 1 disagree, average score = 4.3)

7. CT or MRI should be used to characterize the degree of organization of collections before intervention.
  - a. Note: CT has the potential to underestimate the complexity of fluid collections

GRADE 1C, agreement 100% (11/11, 6 strongly agree, 5 agree, average score = 4.5)

## Acute Recurrent Pancreatitis Summary Statements and Recommendations

The recommendations for AP above also apply to assessment of repeated episodes of AP in the child with ARP.

8. MRI is recommended to identify structural or obstructive causes for ARP.
  - a. This recommendation reflects the high soft tissue contrast and ability to assess the pancreatic and biliary ducts afforded by MRI.

GRADE 1B, agreement 100% (11/11, 9 strongly agree, 2 agree, average score = 4.8)

## Serial Follow-up for Progression to Chronic Pancreatitis

9. When clinically indicated, MRI is recommended to follow children with ARP and to assess for progression to CP.
  - a. This recommendation reflects the strengths of MRI in monitoring changes in both parenchyma and duct. This also reflects the lack of ionizing radiation associated with MRI
  - b. Note: The need for, and frequency of, serial follow-up in children with ARP as a means for assessing for progression to CP has not been defined.

GRADE 1C, agreement 100% (11/11, 7 strongly agree, 4 agree, average score = 4.6)
10. In a child who requires sedation for imaging, it is reasonable to alternate MRI with ultrasound or CT for serial monitoring of ARP.
 

GRADE 2C, agreement 82% (9/11, 3 strongly agree, 6 agree, 1 neutral, 1 disagree, average score = 4)

## IMAGING OF CHRONIC PANCREATITIS

### Purpose/Indication/Rationale for Imaging in Chronic Pancreatitis

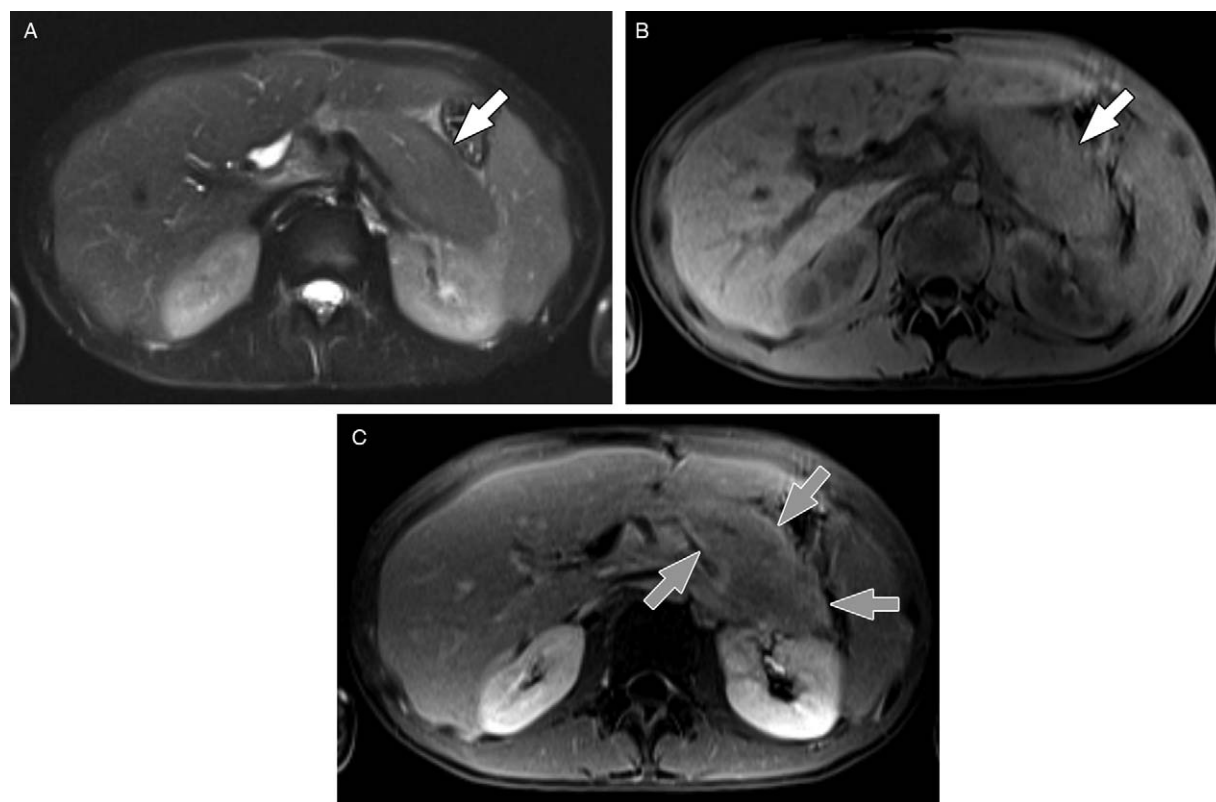
Imaging in CP serves multiple purposes. The dominant role of imaging at the time of diagnosis is to identify findings of CP that can be leveraged in combination with other criteria (clinical and histologic whenever available) to make the diagnosis of CP (11). Accurate diagnosis of CP will lead to altered management as it is now understood that children diagnosed with CP require specific clinical and laboratory follow-up as well as nutritional management (10).

In addition to diagnosing CP (11), there are specific imaging features that are of interest to the clinician in the child with known or suspected CP. Signs of acute inflammation or complications that may require therapeutic intervention/drainage, and, certainly, suspected pancreatic masses are relevant to diagnosis and management. One relatively rare but increasingly recognized and important differential diagnosis of a pancreatic “mass” is autoimmune pancreatitis (AIP) (82,83). AIP is highly responsive to steroid therapy, and hence its diagnosis will lead to a medical therapeutic intervention. Focal, segmental, or global enlargement of the pancreas with loss of the normal contour can be evidence of AIP, especially with delayed enhancement and the presence of a capsule-like rim, which is uncommon but very specific (Fig. 5) (84,85). Focal pancreatic enlargement should be differentiated from tumors as management is completely different (82).

The presence of pancreatic atrophy is helpful in interpretation of biochemical markers of CP and estimating the clinical risk for pancreatic exocrine and endocrine dysfunction. Studies correlating imaging findings and function are, however, lacking (86). The presence of parenchymal calcifications is a major feature of most criteria (generally adult-focused) for CP, though anecdotally, calcification appears to be less common in pediatrics.

Characterization of biliary and pancreatic duct anatomy is also important, particularly with regard to diagnostic and therapeutic decisions. Congenital anomalies, such as pancreaticobiliary maljunction and pancreas divisum can contribute to the underlying pancreatitis (87,88). Duct filling defects/calcifications can be both diagnostic criteria and therapeutic targets. An irregular narrow main pancreatic duct without marked upstream dilation and with smooth tapering of the common bile duct can be suggestive of AIP in the appropriate clinical context (89,90).

Imaging plays a critical role in surgical planning for patients with CP. Decompressing surgeries, such as lateral pancreaticojejunostomies, can be utilized in cases of very dilated pancreatic ducts, with or without the presence of intraductal stones (91). Pancreaticoduodenectomy-type procedures are considered particularly with pancreatic head pathologies (92). Total pancreatectomy-islet cell autotransplantation (TP-IAT) surgery is considered not only based on symptoms and the underlying etiology of CP but also based on the overall imaging appearance of pancreas, including perceived capacity to retrieve a critical mass of islet cells (93). As such, characterization of duct abnormalities and the extent of parenchymal change are important.



**FIGURE 5.** A 15-year-old boy with autoimmune pancreatitis. (A) Axial T2-weighted, fat-saturated MR image shows a diffusely enlarged, mildly T2-weighted hyperintense pancreas (arrow). (B) Axial T1-weighted, fat-saturated MR image shows a diffusely enlarged, T1-weighted hypointense pancreas (arrow). (C) Axial T1-weighted, fat-saturated MR image obtained 5 minutes after administration of intravenous contrast material shows a thin rim of enhancement surrounding the enlarged pancreas (grey arrows). CT = computed tomography; MR = magnetic resonance.

TABLE 6. Glossary of imaging terms/findings for chronic pancreatitis in adults

Location	Feature	Definition
Duct	MPD dilatation	>3.5 mm in body >1.5 mm in tail Lack of tapering of MPD from body to tail
	Side branch dilatation	≥3 tubular structures extending from the MPD
	Stricture	Focal narrowing of the MPD with or without upstream dilation
	Irregular contour of MPD or side branches	Qualitative
	Intraductal calculus	Filling defect (at EUS, must be ≥2 mm echogenic shadowing focus)
	Obstruction	No consensus definition Suggested definition: duct completely occluded because of calculus or stricture in the absence of malignancy
	Duct/periductal fibrosis	Histopathologic finding extrapolated to EUS and MRCP EUS finding of hyperechoic duct wall involving greater than 50% of body and tail of pancreas Qualitative MRCP finding where MPD does not dilate after secretin administration
Parenchymal	Generalized or focal atrophy	Gland thickness < norms defined for pediatrics (see Table 6) Specifically, in adults: pancreatic body thickness measured at level of left margin of vertebral body
		Mild 21 mm Moderate 14 mm Severe 7 mm
		Focal atrophy involves <30% of pancreatic parenchyma Diffuse atrophy involves ≥70% of pancreatic parenchyma
	Irregular contour of gland or accentuated lobular pattern	EUS finding of ≥3 lobules, each measuring ≥5 mm in body or tail
	Honeycombing lobularity	EUS finding of ≥3 continuous lobules
	Coarse calcifications	Best seen at CT, may not be apparent by MRI At EUS, echogenic shadowing focus must be larger than 3 mm At CT, >7 considered severe CP At CT >50 considered innumerable
	Cavities	EUS finding of pancreatic or peripancreatic collections that fill with contrast at ERCP Large defined as >10 mm diameter
	Decreased T1 signal	Qualitative Or signal intensity ratio compared with spleen, paraspinal muscle, and/or liver Or based on T1 relaxometry
	Pancreatic parenchymal enhancement ratio	Signal intensity during arterial phase divided by signal intensity during portal venous phase. ≤1 considered abnormal
Exocrine function	Qualitative assessment of duodenal filling after secretin administration	Matos criteria defines filling beyond the genu inferius as normal
	Quantitative assessment of fluid secretion after secretin administration	Adult and pediatric norms have been defined but cut-offs for EPI have not

Applicability to pediatrics has yet to be defined. Data from (11,20,38,106,116–120). These features have been extrapolated to pediatrics but very few have been validated. CP = chronic pancreatitis; CT = computed tomography; EPI = exocrine pancreatic insufficiency; EUS = endoscopic ultrasound; MPD = main pancreatic duct; MRCP = magnetic resonance cholangiopancreatography.

## Imaging Findings of Chronic Pancreatitis

Imaging features of advanced CP have been well characterized and described in the adult literature (Table 6) (21). Imaging findings of advanced CP specific to pediatrics have not yet been defined, and findings of early or probable CP, when intervention might be attempted to slow disease progression, have not been well-defined for any population (94). Features that may herald “early” CP include a few ectatic duct side branches, parenchymal volume loss, or mild T1 signal changes but these remain to be validated (95).

Currently, no standard imaging classification system exists for CP in children. The Cambridge classification, based on pancreatic duct findings on ERCP in adult patients (Table 7), has been adapted to MRCP in adults but this classification does not incorporate parenchymal changes of CP, and this classification has not been validated in children (96,97). The Cambridge classification defines CP based on the number of abnormal side branches, cavities, filling defects, or obstruction visualized (20). The M-ANNHEIM

Classification for adult CP published in 2007 relied partially on the Cambridge classification as well as other imaging findings at transabdominal ultrasound, CT, EUS, and/or MRI but has not been validated in children (Table 8) (98). Recently, the Consortium for

TABLE 7. Cambridge classification of chronic pancreatitis in adults by endoscopic retrograde cholangiopancreatography

Grade	MPD	Number abnormal side branches
0. Normal	Normal	None
1. Equivocal	Normal	<3
2. Mild CP	Normal	≥3
3. Moderate CP	Abnormal	≥3
4. Severe CP	Abnormal	≥1 large cavity, obstruction, filling defect, severe dilatation or contour irregularity

Adapted from (20). CP = chronic pancreatitis; MPD = main pancreatic duct.

TABLE 8. M-ANNHEIM diagnostic criteria for chronic pancreatitis

<i>Definite</i> chronic pancreatitis (one or more of the following criteria)	Pancreatic calcifications Moderate or severe duct findings (see Table 7) Marked and persistent exocrine insufficiency (pancreatic steatorrhea markedly reduced by enzyme supplementation) Typical histology (with adequate histologic specimen)
<i>Probable</i> chronic pancreatitis (one or more of the following criteria)	Mild duct findings (see Table 7) Recurrent or persistent pseudocysts Pathologic test of pancreatic exocrine function (such as fecal elastase-1, secretin test, secretin-pancreozymin test) Endocrine insufficiency (ie, abnormal glucose tolerance test)
<i>Borderline</i> chronic pancreatitis	Typical clinical history but without additional above criteria

Adapted from (98).

the Study of Chronic Pancreatitis, Diabetes, and Pancreatic Cancer (CPDPC) proposed new reporting standards for CT and MRCP in adults with CP (20). These remain to be validated in adults, and their applicability to pediatrics is unknown.

Given the expected growth and maturation of the pancreas during childhood, knowledge of normal anatomy and gland and duct size based on age is critical to define abnormalities (eg, atrophy and duct dilation) suggestive of CP. Normative values for gland thickness exist for ultrasound and CT (56,99). Based on no significant difference between CT measurements and measurements previously reported at ultrasound, Trout et al (99) suggested that thickness values could likely be extrapolated to MRI, though this remains to be confirmed. Normative values for pancreatic duct diameter also exist for ultrasound and MRI (Table 9) (56,99). Although normal values exist, cut-off values for diagnosis of CP have not been defined.

## Diagnostic Performance of Imaging Modalities in Chronic Pancreatitis

Data specific to the diagnostic performance of specific imaging modalities for pediatric CP are not available. A meta-analysis of adult literature concluded ultrasound, MRI, and CT all have high diagnostic sensitivity and specificity for CP without significant differences (100). Reference standards varied across the included studies. Specifically, in 3460 adults, estimated sensitivities were 67% (95% CI: 53%–78%) for ultrasound, 78% (95% CI: 69%–85%) for

MRI, and 75% (95% CI: 66%–83%) for CT. Estimates of specificity were 98% (95% CI: 89%–100%) for ultrasound, 96% (95% CI: 90%–98%) for MRI, and 91% (95% CI: 81%–96%) for CT.

## Transabdominal Ultrasound

When planning for surgical procedures, ultrasound provides insufficient anatomic assessment, particularly with regard to vascular variants relevant to surgical approach (eg, right hepatic artery or accessory right hepatic artery arising from the superior mesenteric artery).

## Computed Tomography

Use of IV contrast material is recommended when performing CT for CP. IV contrast material allows optimal assessment of the pancreatic parenchyma and allows evaluation of the peripancreatic vessels for patency. For adults, multiphase protocols that include an unenhanced phase, parenchymal/arterial phase, and portal venous phase have been recommended (20). No such recommendations exist for pediatrics but given the relative infrequency of calcifications in pediatric pancreatitis, an unenhanced phase is likely unnecessary. As with AP, a parenchymal/arterial phase can be useful if clinical questions relate to the arteries but a single portal venous phase examination is generally sufficient to characterize CP in children.

CT with IV contrast material provides excellent assessment of the pancreatic parenchyma, allowing identification of features of ARP and CP, particularly calcifications and pancreatic atrophy. CT can also identify congenital anomalies, such as annular pancreas and can assess for superimposed acute pancreatitis and complications of pancreatitis, including established vascular collaterals because of chronic/established thrombosis. CT (and MRI) outperform ultrasound to define vascular anatomy relevant to surgical planning. CT is, however, limited by suboptimal visualization of the pancreatic and biliary ducts.

## Magnetic Resonance Imaging/Magnetic Resonance Cholangiopancreatography

MRI and MRCP have the benefits of providing information on both parenchymal and duct changes of CP but are limited in their ability to visualize calcifications. Adult and pediatric data suggest that the sensitivity of MRCP to detect pancreatic duct abnormalities may be improved by the administration of secretin (38). Theoretically secretin distends the pancreatic duct and may allow for earlier detection of side branch-ectasias and provide information on exocrine function by quantifying duodenal filling (101–103).

Although MRI is the favored noninvasive imaging modality for assessment of the pancreatic duct and does well in this capacity, ERCP remains the only modality that allows the pancreatic and

TABLE 9. Reference values for normal pancreatic duct diameter at ultrasound and magnetic resonance imaging in children

Age	Main pancreatic duct diameter (mm ± SD)		
Ultrasound			
1–3 y		1.13 ± 0.15	
4–6 y		1.35 ± 0.15	
7–9 y		1.67 ± 0.17	
10–12 y		1.78 ± 0.17	
13–15 y		1.92 ± 0.18	
16–18 y		2.05 ± 0.15	
	Head	Body	Tail
MRI			
0–12 mo	0.8 ± 0.2	0.7 ± 0.2	0.7 ± 0.2
1–23 mo	1.0 ± 0.3	0.9 ± 0.3	0.8 ± 0.2
24–59 mo	1.1 ± 0.3	1.1 ± 0.3	1.1 ± 0.3
60–95 mo	1.4 ± 0.3	1.3 ± 0.2	1.3 ± 0.2
96–120 mo	1.4 ± 0.3	1.4 ± 0.3	1.4 ± 0.3

Data from (56,121). MRI = magnetic resonance imaging; SD = standard deviation.

biliary ducts to be imaged distended under pressure, maximizing characterization of duct anomalies and abnormalities (104).

Imaging techniques are becoming increasingly quantitative. Adult data have shown that pancreatic exocrine function can be noninvasively assessed with ultrasound or MRI with reasonable agreement with direct stimulation tests with collection of intraduodenal fluid (105). In children, measurement of the volume of fluid secreted by the pancreas in response to secretin has been shown to be highly accurate by MRI (<10% error in measured volume) with high inter-reader reproducibility (106). Further, normative data for secreted fluid volume measured by MRI in children now exist but diagnostic cut-offs for EPI remain to be defined (107).

Elastography allows the measurement of tissue stiffness by either ultrasound or MRI, and in other organs, increased stiffness is associated with tissue fibrosis. Although correlation between pancreatic stiffness and fibrosis related to CP remains to be defined, studies have revealed differences in stiffness in patients with CP compared with controls. In an adult study, transabdominal strain ultrasound elastography, when combined with greyscale ultrasound, was shown to correctly diagnose presence of CP in approximately 95% of those confirmed to have CP (108). Pancreatic stiffness as measured by MR elastography has also been shown in adults to be significantly different between controls and patients with either mild CP or moderate/severe CP ( $1.21 \pm 0.13$  vs  $1.50 \pm 0.15$  vs  $1.90 \pm 0.16$  kPa;  $P < 0.001$ ) (109). A recent publication including 49 controls and 14 children with pancreatitis (acute or chronic) showed measured stiffness to be lower in those with pancreatitis (normal controls:  $1.7 \pm 0.3$  kPa for both of 2 readers; AP or CP:  $0.9 \pm 0.2$  and  $1.1 \pm 0.3$  kPa) (110).

T1 relaxation time, which quantifies the finding of T1 signal loss associated with CP, has been shown in adults to have 77% sensitivity and 83% specificity for CP (109). Further, Wang et al (109) showed that T1 relaxation times were significantly different between controls and patients with either mild CP or moderate/severe CP ( $865 \pm 220$  vs  $1075 \pm 22$  vs  $1350 \pm 139$  milliseconds;  $P < 0.001$ ) though AUCs were lower than for pancreatic stiffness. Notably, multiple regression analysis showed that T1 relaxation time and stiffness were independent predictors of mild CP. Pediatric data are just being generated for pancreatic T1 relaxation time; for example, Gilligan et al (111) reported values in a small cohort of healthy children at 1.5 and 3 T.

## Chronic Pancreatitis Summary Statements and Recommendations

### Initial Diagnosis

11. MRI is the recommended modality for imaging of suspected CP (Table 3).
  - a. This recommendation reflects inadequate characterization of findings (particularly duct findings) of CP by transabdominal ultrasound. This also reflects the superior soft tissue contrast of MRI, which allows characterization of both parenchyma and duct findings of CP.
 GRADE 1C, agreement 91% (10/11, 9 strongly agree, 1 agree, 1 disagree, average score = 4.6)

### Assessment for Superimposed Acute Pancreatitis

12. When imaging is needed to assess a suspected or known episode of AP in a child with CP, transabdominal ultrasound is the preferred first-line imaging modality.
  - a. This recommendation reflects the availability and portability of ultrasound and the role of ultrasound in identifying biliary causes of AP.

- b. Note: A negative ultrasound does not exclude AP (low to moderate sensitivity).

GRADE 1B, agreement 91% (10/11, 6 strongly agree, 4 agree, 1 neutral, average score = 4.5)

13. If ultrasound is negative for AP in a child with CP and an imaging diagnosis of AP is needed, either CT or MRI are recommended.
  - a. This recommendation reflects the only moderate sensitivity of ultrasound and the greater sensitivity of CT and MRI.
 GRADE 1B, agreement 100% (11/11, 6 strongly agree, 5 agree, average score = 4.5)

## Intervention Planning

14. CT or MRI are recommended for planning of endoscopic or surgical interventions in a patient with known CP.
  - a. This recommendation is based on the large field of view afforded by CT and MRI, optimal characterization of the degree of organization of fluid collections, and optimal characterization of the peripancreatic vasculature.
  - b. Note: When the intervention will target the duct, MRI is favored over CT.
 GRADE 2C, agreement 100% (11/11, 6 strongly agree, 5 agree, average score = 4.5)

## Serial Monitoring

15. MRI is recommended for clinically indicated serial imaging of CP.
  - a. This recommendation reflects the optimal soft tissue contrast of MRI, no need for intravenous contrast material (in most cases), and the lack of associated ionizing radiation.
  - b. Note: In the child who requires sedation for MRI, risks of sedation must be balanced with the need for serial imaging.
 GRADE 1B, agreement 100% (11/11, 9 strongly agree, 2 agree, average score = 4.8)

## FUTURE DIRECTIONS/GAPS IN KNOWLEDGE/NECESSARY RESEARCH

### Basic Imaging Strategies

A paucity of rigorous research exists on the diagnostic performance of transabdominal ultrasound (including CEUS), CT, and MRI for pediatric pancreatitis. These basic studies are needed to define the optimal imaging modality(ies) for AP, ARP, and CP in children and to inform future diagnostic guidelines. Such research should include exploration of the optimal timing, relative to attack(s) of AP, of imaging aimed at identifying a pancreatic duct cause of pancreatitis. Further, research is needed to define the optimal imaging strategies for pancreatitis that balance diagnostic performance with cost, and the risks of sedation, intravenous contrast material and radiation exposure.

### Prognostication and Prediction

Research into imaging (and other diagnostic) methods is needed to prognosticate and predict disease severity in the context of AP and disease progression in the context of AP and ARP. Currently, it is not possible to predict which patients will progress

from AP to ARP and from ARP to CP. However, preliminary data are emerging. A recent study in adults showed a decrease in pancreas volume after 3 episodes of AP, with volume loss possibly reflecting an early finding in the transition to CP (112). Understanding and predicting the progression through various stages of pancreatitis would allow for more effective counseling and optimization of the timing of interventions. Further, research is also needed into the role of imaging in prognostication based on genetic etiologies of pancreatitis.

## Identification of Minimal Change Chronic Pancreatitis

Currently the identification of CP is often delayed, with imaging findings only apparent when disease is well established. CP, and attendant pancreatic dysfunction, are sources of significant morbidity, particularly in children, and thus early identification and intervention are critical. As such, research is needed to facilitate identification of minimal change or early CP. Early studies in adult populations suggest quantitative MRI methods, such as T1 mapping and MR elastography may be able to identify early CP (109,113).

## Noninvasive Pancreatic Function Assessment

Diagnosis of exocrine and, to some degree, endocrine insufficiency remains invasive. Research is needed to identify noninvasive, or minimally invasive techniques to diagnose insufficiency, and more importantly to predict development of insufficiency to allow early intervention. Some data suggest that imaging can noninvasively assess exocrine and endocrine pancreatic function, but these techniques require further study (114,115).

## CONCLUSIONS

Pancreatitis in the pediatric population, both acute and chronic, is increasingly being recognized. As with adults, imaging plays a role in the diagnosis, staging, and follow-up of both acute and chronic pancreatitis. Pediatric-specific literature informing the use of imaging in pancreatitis is, however, sparse. For this reason, much of what we know and recommend regarding imaging of pediatric pancreatitis is extrapolated from the adult literature. This document provides summaries and recommendations of the literature regarding imaging of the child with pancreatitis that can be used to inform clinical decision-making. Many of the recommendations are largely based on expert opinion. Going forward, dedicated pediatric studies of imaging in pancreatitis are clearly needed. These studies should address not only optimal basic imaging strategies but should also address the more complex problems of identification of early stage disease and prognostication of disease course to enable generation of pediatric-specific evidence-based guidelines.

## REFERENCES

- Morinville VD, Barmada MM, Lowe ME. Increasing incidence of acute pancreatitis at an American pediatric tertiary care center: is greater awareness among physicians responsible? *Pancreas* 2010;39:5–8.
- Abu-El-Haija M, El-Dika S, Hinton A, et al. Acute pancreatitis admission trends: a national estimate through the Kids' Inpatient Database. *J Pediatr* 2018;194:147.e1–51.e1.
- Kumar S, Ooi CY, Werlin S, et al. Risk factors associated with pediatric acute recurrent and chronic pancreatitis: lessons from INSPPIRE. *JAMA Pediatr* 2016;170:562–9.
- Abu-El-Haija M, Kumar S, Quiros JA, et al. Management of acute pancreatitis in the pediatric population: a clinical report from the North American Society for Pediatric Gastroenterology, Hepatology and Nutrition Pancreas Committee. *J Pediatr Gastroenterol Nutr* 2018;66:159–76.
- Lin TK, Troendle DM, Wallihan DB, et al. Specialized imaging and procedures in pediatric pancreatology: a North American Society for Pediatric Gastroenterology, Hepatology, and Nutrition Clinical Report. *J Pediatr Gastroenterol Nutr* 2017;64:472–84.
- Liu QY, Gugig R, Troendle DM, et al. The roles of EUS and ERCP in the evaluation and treatment of chronic pancreatitis in children: a Position Paper from the NASPGHAN Pancreas Committee. *J Pediatr Gastroenterol Nutr* 2020;70:681–3.
- Shekelle P. Overview of clinical practice guidelines. <https://www.up-to-date.com/contents/overview-of-clinical-practice-guidelines>. Accessed 27 March, 2020.
- Harris PA, Taylor R, Minor BL, et al., REDCap Consortium. The REDCap consortium: building an international community of software platform partners. *J Biomed Inform* 2019;95:103208.
- Harris PA, Taylor R, Thielke R, et al. Research electronic data capture (REDCap)—a metadata-driven methodology and workflow process for providing translational research informatics support. *J Biomed Inform* 2009;42:377–81.
- Abu-El-Haija M, Uc A, Werlin SL, et al. Nutritional considerations in pediatric pancreatitis: a position paper from the NASPGHAN Pancreas Committee and ESPHAN Cystic Fibrosis/Pancreas Working Group. *J Pediatr Gastroenterol Nutr* 2018;67:131–43.
- Morinville VD, Husain SZ, Bai H, et al., REDCap Consortium. Definitions of pediatric pancreatitis and survey of present clinical practices. *J Pediatr Gastroenterol Nutr* 2012;55:261–5.
- Husain SZ, Srinath AI. What's unique about acute pancreatitis in children: risk factors, diagnosis and management. *Nat Rev Gastroenterol Hepatol* 2017;14:366–72.
- Abu-El-Haija M, Lowe ME. Pediatric pancreatitis-molecular mechanisms and management. *Gastroenterol Clin North Am* 2018;47:741–53.
- Uc A, Husain SZ. Pancreatitis in children. *Gastroenterology* 2019;156:1969–78.
- Abu-El-Haija M, Kumar S, Szabo F, et al., NASPGHAN Pancreas Committee. Classification of acute pancreatitis in the pediatric population: clinical report from the NASPGHAN Pancreas Committee. *J Pediatr Gastroenterol Nutr* 2017;64:984–90.
- Goldstein B, Giroir B, Randolph A, et al. International pediatric sepsis consensus conference: definitions for sepsis and organ dysfunction in pediatrics. *Pediatr Crit Care Med* 2005;6:2–8.
- Galai T, Cohen S, Yerushalmy-Feler A, et al. Young age predicts acute pancreatitis severity in children. *J Pediatr Gastroenterol Nutr* 2019;68:720–6.
- Garipey CE, Heyman MB, Lowe ME, et al. Causal evaluation of acute recurrent and chronic pancreatitis in children: consensus from the INSPPIRE Group. *J Pediatr Gastroenterol Nutr* 2017;64:95–103.
- Sweeny KF, Lin TK, Nathan JD, et al. Rapid progression of acute pancreatitis to acute recurrent pancreatitis in children. *J Pediatr Gastroenterol Nutr* 2019;68:104–9.
- Tirkes T, Shah ZK, Takahashi N, et al., Consortium for the Study of Chronic Pancreatitis, Diabetes, and Pancreatic Cancer. Reporting standards for chronic pancreatitis by Using CT, MRI, and MR cholangiopancreatography: the Consortium for the Study of Chronic Pancreatitis, Diabetes, and Pancreatic Cancer. *Radiology* 2019;290:207–15.
- Darge K, Anupindi S. Pancreatitis and the role of US, MRCP and ERCP. *Pediatr Radiol* 2009;39(Suppl 2):S153–7.
- Restrepo R, Hagerott HE, Kulkarni S, et al. Acute pancreatitis in pediatric patients: demographics, etiology, and diagnostic imaging. *AJR Am J Roentgenol* 2016;206:632–44.
- Parniczky A, Abu-El-Haija M, Husain S, et al. EPC/HPSG evidence-based guidelines for the management of pediatric pancreatitis. *Pancreatol* 2018;18:146–60.

24. Bollen TL. Imaging Assessment of Etiology and Severity of Acute Pancreatitis. <https://pancreapedia.org/reviews/imaging-assessment-of-etiology-and-severity-of-acute-pancreatitis>. Accessed 18 June, 2019.
25. Nievelstein RA, Robben SG, Blickman JG. Hepatobiliary and pancreatic imaging in children—techniques and an overview of non-neoplastic disease entities. *Pediatr Radiol* 2011;41:55–75.
26. Chang YJ, Chao HC, Kong MS, et al. Acute pancreatitis in children. *Acta Paediatr* 2011;100:740–4.
27. Golea A, Badea R, Socaciu M, et al. Quantitative analysis of tissue perfusion using contrast-enhanced transabdominal ultrasound (CEUS) in the evaluation of the severity of acute pancreatitis. *Med Ultrason* 2010;12:198–204.
28. Rickes S, Monkemüller K, Malfertheiner P. Acute severe pancreatitis: contrast-enhanced sonography. *Abdom Imaging* 2007;32:362–4.
29. Gilligan LA, Davenport MS, Trout AT, et al. Risk of acute kidney injury following contrast-enhanced CT in hospitalized pediatric patients: a propensity score analysis. *Radiology* 2020;294:548–56.
30. Sandrasegaran K, Menias CO. Imaging in autoimmune pancreatitis and immunoglobulin G4-related disease of the abdomen. *Gastroenterol Clin North Am* 2018;47:603–19.
31. Fitoz S, Erden A, Boruban S. Magnetic resonance cholangiopancreatography of biliary system abnormalities in children. *Clin Imaging* 2007;31:93–101.
32. Kim MJ, Han SJ, Yoon CS, et al. Using MR cholangiopancreatography to reveal anomalous pancreaticobiliary ductal union in infants and children with choledochal cysts. *AJR Am J Roentgenol* 2002;179:209–14.
33. Mortelet KJ, Rocha TC, Streeter JL, et al. Multimodality imaging of pancreatic and biliary congenital anomalies. *Radiographics* 2006;26:715–31.
34. Delaney L, Applegate KE, Karmazyn B, et al. MR cholangiopancreatography in children: feasibility, safety, and initial experience. *Pediatr Radiol* 2008;38:64–75.
35. Matos C, Metens T, Deviere J, et al. Pancreas divisum: evaluation with secretin-enhanced magnetic resonance cholangiopancreatography. *Gastrointest Endosc* 2001;53:728–33.
36. Rustagi T, Njei B. Magnetic resonance cholangiopancreatography in the diagnosis of pancreas divisum: a systematic review and meta-analysis. *Pancreas* 2014;43:823–8.
37. Sandrasegaran K, Tahir B, Barad U, et al. The value of secretin-enhanced MRCP in patients with recurrent acute pancreatitis. *AJR Am J Roentgenol* 2017;208:315–21.
38. Manfredi R, Lucidi V, Gui B, et al. Idiopathic chronic pancreatitis in children: MR cholangiopancreatography after secretin administration. *Radiology* 2002;224:675–82.
39. Sandrasegaran K, Cote GA, Tahir B, et al. The utility of secretin-enhanced MRCP in diagnosing congenital anomalies. *Abdom Imaging* 2014;39:979–87.
40. Mosler P, Akisik F, Sandrasegaran K, et al. Accuracy of magnetic resonance cholangiopancreatography in the diagnosis of pancreas divisum. *Dig Dis Sci* 2012;57:170–4.
41. Banks PA, Bollen TL, Dervenis C, et al., Acute Pancreatitis Classification Working Group. Classification of acute pancreatitis—2012: revision of the Atlanta classification and definitions by international consensus. *Gut* 2013;62:102–11.
42. Kandula L, Lowe ME. Etiology and outcome of acute pancreatitis in infants and toddlers. *J Pediatr* 2008;152:106.e1–110.e10.
43. Park A, Latif SU, Shah AU, et al. Changing referral trends of acute pancreatitis in children: a 12-year single-center analysis. *J Pediatr Gastroenterol Nutr* 2009;49:316–22.
44. Ma MH, Bai HX, Park AJ, et al. Risk factors associated with biliary pancreatitis in children. *J Pediatr Gastroenterol Nutr* 2012;54:651–6.
45. Izquierdo YE, Fonseca EV, Moreno LA, et al. Utility of CT classifications to predict unfavorable outcomes in children with acute pancreatitis. *Pediatr Radiol* 2018;48:954–61.
46. Benifla M, Weizman Z. Acute pancreatitis in childhood: analysis of literature data. *J Clin Gastroenterol* 2003;37:169–72.
47. Werlin SL, Kugathasan S, Frautschy BC. Pancreatitis in children. *J Pediatr Gastroenterol Nutr* 2003;37:591–5.
48. 9Bolia R, Srivastava A, Yachha SK, et al. Prevalence, natural history, and outcome of acute fluid collection and pseudocyst in children with acute pancreatitis. *J Pediatr Gastroenterol Nutr* 2015;61:451–5.
49. DeBanto JR, Goday PS, Pedrosa MR, et al., Midwest Multicenter Pancreatic Study Group. Acute pancreatitis in children. *Am J Gastroenterol* 2002;97:1726–31.
50. Manikkavasakar S, AlObaidy M, Busireddy KK, et al. Magnetic resonance imaging of pancreatitis: an update. *World J Gastroenterol* 2014;20:14760–77.
51. McIntosh DM, Penney HF. Gray-scale ultrasonography as a screening procedure in the detection of gallbladder disease. *Radiology* 1980;136:725–7.
52. Harvey RT, Miller WT Jr. Acute biliary disease: initial CT and follow-up US versus initial US and follow-up CT. *Radiology* 1999;213:831–6.
53. Orkin SH, Trout AT, Fei L, et al. Sensitivity of biochemical and imaging findings for the diagnosis of acute pancreatitis in children. *J Pediatr* 2019;213:143.e2–8.e2.
54. Coffey MJ, Nightingale S, Ooi CY. Diagnosing acute pancreatitis in children: what is the diagnostic yield and concordance for serum pancreatic enzymes and imaging within 96 h of presentation? *Pancreatol* 2014;14:251–6.
55. Chao HC, Lin SJ, Kong MS, et al. Sonographic evaluation of the pancreatic duct in normal children and children with pancreatitis. *J Ultrasound Med* 2000;19:757–63.
56. Siegel MJ, Martin KW, Worthington JL. Normal and abnormal pancreas in children: US studies. *Radiology* 1987;165:15–8.
57. Balthazar EJ, Ranson JH, Naidich DP, et al. Acute pancreatitis: prognostic value of CT. *Radiology* 1985;156:767–72.
58. Balthazar EJ, Robinson DL, Megibow AJ, et al. Acute pancreatitis: value of CT in establishing prognosis. *Radiology* 1990;174:331–6.
59. Avanesov M, Weinrich JM, Kraus T, et al. MDCT of acute pancreatitis: intraindividual comparison of single-phase versus dual-phase MDCT for initial assessment of acute pancreatitis using different CT scoring systems. *Eur J Radiol* 2016;85:2014–22.
60. Vaughn DD, Jabra AA, Fishman EK. Pancreatic disease in children and young adults: evaluation with CT. *Radiographics* 1998;18:1171–87.
61. Turkvatan A, Erden A, Turkoglu MA, et al. Imaging of acute pancreatitis and its complications. Part 2: complications of acute pancreatitis. *Diagn Interv Imaging* 2015;96:161–9.
62. Johnson CD, Stephens DH, Sarr MG. CT of acute pancreatitis: correlation between lack of contrast enhancement and pancreatic necrosis. *AJR Am J Roentgenol* 1991;156:93–5.
63. Kamal A, Singh VK, Akshintala VS, et al. CT and MRI assessment of symptomatic organized pancreatic fluid collections and pancreatic duct disruption: an interreader variability study using the revised Atlanta classification 2012. *Abdom Imaging* 2015;40:1608–16.
64. Morgan DE, Baron TH, Smith JK, et al. Pancreatic fluid collections prior to intervention: evaluation with MR imaging compared with CT and US. *Radiology* 1997;203:773–8.
65. Lautz TB, Turkel G, Radhakrishnan J, et al. Utility of the computed tomography severity index (Balthazar score) in children with acute pancreatitis. *J Pediatr Surg* 2012;47:1185–91.
66. Fabre A, Petit P, Gaudart J, et al. Severity scores in children with acute pancreatitis. *J Pediatr Gastroenterol Nutr* 2012;55:266–7.
67. Leung TK, Lee CM, Lin SY, et al. Balthazar computed tomography severity index is superior to Ranson criteria and APACHE II scoring system in predicting acute pancreatitis outcome. *World J Gastroenterol* 2005;11:6049–52.
68. Lautz TB, Chin AC, Radhakrishnan J. Acute pancreatitis in children: spectrum of disease and predictors of severity. *J Pediatr Surg* 2011;46:1144–9.
69. Busireddy KK, AlObaidy M, Ramalho M, et al. Pancreatitis-imaging approach. *World J Gastrointest Pathophysiol* 2014;5:252–70.
70. Stimac D, Miletic D, Radic M, et al. The role of nonenhanced magnetic resonance imaging in the early assessment of acute pancreatitis. *Am J Gastroenterol* 2007;102:997–1004.

71. Heyn C, Sue-Chue-Lam D, Jhaveri K, et al. MRI of the pancreas: problem solving tool. *J Magn Reson Imaging* 2012;36:1037–51.
72. Hirota M, Kimura Y, Ishiko T, et al. Visualization of the heterogeneous internal structure of so-called “pancreatic necrosis” by magnetic resonance imaging in acute necrotizing pancreatitis. *Pancreas* 2002;25:63–7.
73. Nandalur KR, Hussain HK, Weadock WJ, et al. Possible biliary disease: diagnostic performance of high-spatial-resolution isotropic 3D T2-weighted MRCP. *Radiology* 2008;249:883–90.
74. Moon JH, Cho YD, Cha SW, et al. The detection of bile duct stones in suspected biliary pancreatitis: comparison of MRCP, ERCP, and intraductal US. *Am J Gastroenterol* 2005;100:1051–7.
75. Varghese JC, Liddell RP, Farrell MA, et al. Diagnostic accuracy of magnetic resonance cholangiopancreatography and ultrasound compared with direct cholangiography in the detection of choledocholithiasis. *Clin Radiol* 2000;55:25–35.
76. Singh A, Mann HS, Thukral CL, et al. Diagnostic accuracy of MRCP as compared to ultrasound/CT in patients with obstructive jaundice. *J Clin Diagn Res* 2014;8:103–7.
77. Freitas ML, Bell RL, Duffy AJ. Choledocholithiasis: evolving standards for diagnosis and management. *World J Gastroenterol* 2006;12:3162–7.
78. Tamir S, Braun M, Issachar A, et al. Yield of magnetic resonance cholangiopancreatography for the investigation of bile duct dilatation in asymptomatic patients. *United European Gastroenterol J* 2017;5:408–14.
79. Chavhan GB, Babyn PS, Manson D, et al. Pediatric MR cholangiopancreatography: principles, technique, and clinical applications. *Radiographics* 2008;28:1951–62.
80. Tang MY, Chen TW, Bollen TL, et al. MR imaging of hemorrhage associated with acute pancreatitis. *Pancreatol* 2018;18:363–9.
81. Barral M, Taouli B, Guin B, et al. Diffusion-weighted MR imaging of the pancreas: current status and recommendations. *Radiology* 2015;274:45–63.
82. Scheers I, Palermo JJ, Freedman S, et al. Autoimmune pancreatitis in children: characteristic features, diagnosis, and management. *Am J Gastroenterol* 2017;112:1604–11.
83. Scheers I, Palermo JJ, Freedman S, et al. Recommendations for diagnosis and management of autoimmune pancreatitis in childhood: consensus from INSPPIRE. *J Pediatr Gastroenterol Nutr* 2018;67:232–6.
84. Rehnitz C, Klauss M, Singer R, et al. Morphologic patterns of autoimmune pancreatitis in CT and MRI. *Pancreatol* 2011;11:240–51.
85. Shimosegawa T, Chari ST, Frulloni L, et al., International Association of Pancreatology. International consensus diagnostic criteria for autoimmune pancreatitis: guidelines of the International Association of Pancreatology. *Pancreas* 2011;40:352–8.
86. DeSouza SV, Singh RG, Yoon HD, et al. Pancreas volume in health and disease: a systematic review and meta-analysis. *Expert Rev Gastroenterol Hepatol* 2018;12:757–66.
87. Cavestro GM, Leandro G, Di Leo M, et al. A single-centre prospective, cohort study of the natural history of acute pancreatitis. *Dig Liver Dis* 2015;47:205–10.
88. Urushihara N, Hamada Y, Kamisawa T, et al. Classification of pancreaticobiliary maljunction and clinical features in children. *J Hepatobiliary Pancreat Sci* 2017;24:449–55.
89. Lee LK, Sahani DV. Autoimmune pancreatitis in the context of IgG4-related disease: review of imaging findings. *World J Gastroenterol* 2014;20:15177–89.
90. Matsubayashi H, Kakushima N, Takizawa K, et al. Diagnosis of autoimmune pancreatitis. *World J Gastroenterol* 2014;20:16559–6.
91. Hodgman E, Megison S, Murphy JT. Puestow procedure for the management of pediatric chronic pancreatitis. *Eur J Pediatr Surg* 2019;29:153–8.
92. Gurusamy KS, Lushuku C, Halkias C, et al. Duodenum-preserving pancreatic resection versus pancreaticoduodenectomy for chronic pancreatitis. *Cochrane Database Syst Rev* 2016;2:CD011521.
93. Bellin MD, Abu-El-Hajja M, Morgan K, et al., POST study consortium. A multicenter study of total pancreatectomy with islet auto-transplantation (TPIAT): POST (Prospective Observational Study of TPIAT). *Pancreatol* 2018;18:286–90.
94. Whitcomb DC, Shimosegawa T, Chari ST, et al., Working Group for the International (IAP – APA – JPS – EPC) Consensus Guidelines for Chronic Pancreatitis. International consensus statements on early chronic Pancreatitis. Recommendations from the working group for the international consensus guidelines for chronic pancreatitis in collaboration with The International Association of Pancreatology, American Pancreatic Association, Japan Pancreas Society, Pancreas-Fest Working Group and European Pancreatic Club. *Pancreatol* 2018;S1424-3903(18)30113-3. doi: 10.1016/j.pan.2018.05.008. [Epub ahead of print].
95. Kim DH, Pickhardt PJ. Radiologic assessment of acute and chronic pancreatitis. *Surg Clin North Am* 2007;87:1341–58viii.
96. Conwell DL, Lee LS, Yadav D, et al. American Pancreatic Association Practice Guidelines in chronic pancreatitis: evidence-based report on diagnostic guidelines. *Pancreas* 2014;43:1143–62.
97. Sarner M, Cotton PB. Classification of pancreatitis. *Gut* 1984;25:756–9.
98. Schneider A, Lohr JM, Singer MV. The M-ANNHEIM classification of chronic pancreatitis: introduction of a unifying classification system based on a review of previous classifications of the disease. *J Gastroenterol* 2007;42:101–19.
99. Trout AT, Preet-Singh K, Anton CG, et al. Normal pancreatic parenchymal thickness by CT in healthy children. *Pediatr Radiol* 2018;48:1600–5.
100. Issa Y, Kempeneers MA, van Santvoort HC, et al. Diagnostic performance of imaging modalities in chronic pancreatitis: a systematic review and meta-analysis. *Eur Radiol* 2017;27:3820–44.
101. Balci NC, Smith A, Momtahan AJ, et al. MRI and S-MRCP findings in patients with suspected chronic pancreatitis: correlation with endoscopic pancreatic function testing (ePFT). *J Magn Reson Imaging* 2010;31:601–6.
102. Boraschi P, Donati F, Cervelli R, et al. Secretin-stimulated MR cholangiopancreatography: spectrum of findings in pancreatic diseases. *Insights Imaging* 2016;7:819–29.
103. Tirkes T, Fogel EL, Sherman S, et al. Detection of exocrine dysfunction by MRI in patients with early chronic pancreatitis. *Abdom Radiol (NY)* 2017;42:544–51.
104. Tipnis NA, Dua KS, Werlin SL. A retrospective assessment of magnetic resonance cholangiopancreatography in children. *J Pediatr Gastroenterol Nutr* 2008;46:59–64.
105. Cappeliez O, Delhaye M, Deviere J, et al. Chronic pancreatitis: evaluation of pancreatic exocrine function with MR pancreatography after secretin stimulation. *Radiology* 2000;215:358–64.
106. Trout AT, Wallihan DB, Serai S, et al. Secretin-enhanced magnetic resonance cholangiopancreatography for assessing pancreatic secretory function in children. *J Pediatr* 2017;188:186–91.
107. Trout AT, Serai SD, Fei L, et al. Prospective assessment of normal pancreatic secretory function measured by MRI in a cohort of healthy children. *Am J Gastroenterol* 2018;113:1385.
108. Uchida H, Hirooka Y, Itoh A, et al. Feasibility of tissue elastography using transcutaneous ultrasonography for the diagnosis of pancreatic diseases. *Pancreas* 2009;38:17–22.
109. Wang M, Gao F, Wang X, et al. Magnetic resonance elastography and T1 mapping for early diagnosis and classification of chronic pancreatitis. *J Magn Reson Imaging* 2018. doi: 10.1002/jmri.26008. Epub ahead of print. PMID: 29537715; PMCID: PMC6138575.
110. Serai SD, Abu-El-Hajja M, Trout AT. 3D MR elastography of the pancreas in children. *Abdom Radiol (NY)* 2019;44:1834–40.
111. Gilligan LA, Dillman JR, Tkach JA, et al. Magnetic resonance imaging T1 relaxation times for the liver, pancreas and spleen in healthy children at 1.5 and 3 tesla. *Pediatr Radiol* 2019;49:1018–24.
112. DeSouza SV, Priya S, Cho J, et al. Pancreas shrinkage following recurrent acute pancreatitis: an MRI study. *Eur Radiol* 2019;29:3746–56.
113. Tirkes T, Lin C, Fogel EL, et al. T1 mapping for diagnosis of mild chronic pancreatitis. *J Magn Reson Imaging* 2017;45:1171–6.
114. Arifin DR, Bulte JW. Imaging of pancreatic islet cells. *Diabetes Metab Res Rev* 2011;27:761–6.



115. Serai SD, Dillman JR, Trout AT. Spin-echo Echo-planar imaging MR elastography versus gradient-echo MR elastography for assessment of liver stiffness in children and young adults suspected of having liver disease. *Radiology* 2017;282:761–70.
116. Uc A, Perito ER, Pohl JF, et al. INternational Study Group of Pediatric Pancreatitis: in search for a CuRE Cohort study: design and rationale for INSPPIRE 2 from the Consortium for the Study of Chronic Pancreatitis, Diabetes, and Pancreatic Cancer. *Pancreas* 2018;47:1222–8.
117. Catalano MF, Sahai A, Levy M, et al. EUS-based criteria for the diagnosis of chronic pancreatitis: the Rosemont classification. *Gastro-intest Endosc* 2009;69:1251–61.
118. Matos C, Metens T, Deviere J, et al. Pancreatic duct: morphologic and functional evaluation with dynamic MR pancreatography after secretin stimulation. *Radiology* 1997;203:435–41.
119. Axon AT, Classen M, Cotton PB, et al. Pancreatography in chronic pancreatitis: international definitions. *Gut* 1984;25:1107–12.
120. Balci NC, Alkaade S, Magas L, et al. Suspected chronic pancreatitis with normal MRCP: findings on MRI in correlation with secretin MRCP. *J Magn Reson Imaging* 2008;27:125–31.
121. Gwal K, Bedoya MA, Patel N, et al. Reference values of MRI measurements of the common bile duct and pancreatic duct in children. *Pediatr Radiol* 2015;45:1153–9.