



IMAGING MODALITIES FOR PANCREATIC DISEASE

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Outline

- Imaging generalities based on disease category
- Modality advantages / disadvantages
- Protocol considerations / Tips
- Overview of findings of pancreatitis
- Future directions

Pancreatitis



TA Ultrasound CT MRI Pancreatic Tumors

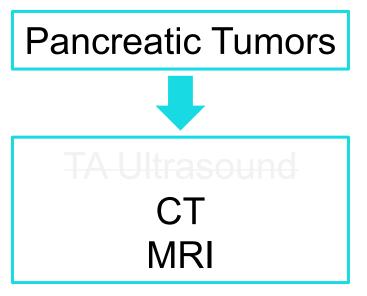


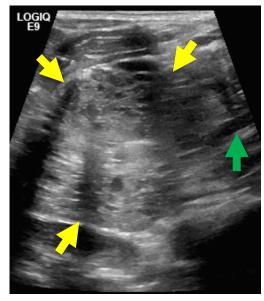
TA Ultrasound

MRI

Modality	Advantages	Disadvantages
Transabdominal Ultrasound	No radiation*AvailabilityPortabilityNo IV contrast	 Limited by bowel gas / habitus Limited FOV Suboptimal vessel assessment Lower sens / spec
СТ	Optimal for Ca⁺⁺Wide FOVSpeed	Radiation*Suboptimal duct assessmentIodinated contrast
MRI	 No radiation* Best soft tissue contrast (duct) Wide FOV Possible non-contrast exam 	 Poor assessment of Ca⁺⁺ Long exams[*] Small space +/- gadolinium contrast
Endoscopic Ultrasound	ProximityPotential to biopsy	InvasiveLimited FOV
ERCP	Potential to intervene	Invasive

- +/- tumor
- Status of vessels
- Invasion of local structures
- +/- metastatic disease







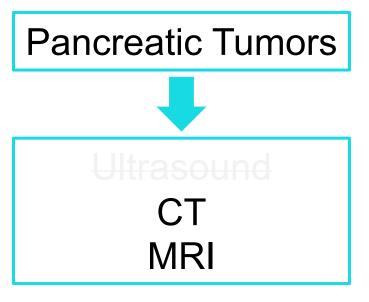
Pancreatic Tumors



CT MRI

- +/- tumor
- Status of vessels
- Invasion of local structures
- +/- metastatic disease

IV Contrast

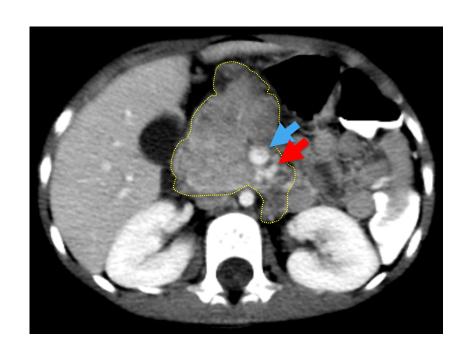


Protocol Considerations / Tips

Pancreatic Tumors

- IV contrast is a must
 - Multiphase post-contrast imaging can help
- Oral contrast can help
- MRI vs. CT is a preference

Pancreatic Tumors





- +/- tumor
- Status of vessels
- Invasion of local structures
- +/- metastatic disease

Pancreatic Tumors



CT MRI FDG PET EUS/ERCP

Protocol Considerations / Tips

Pancreatitis

- IV contrast
 - A must for CT
 - Single phase (portal venous) sufficient*
 - Can help for MRI (autoimmune, necrosis, vessels)
- Oral contrast can help (CT)
- Build redundancy into MRCPs

Protocol Considerations / Tips

Radiology

Reporting Standards for Chronic Pancreatitis by Using CT, MRI, and MR Cholangiopancreatography: The Consortium for the Study of Chronic Pancreatitis, Diabetes, and Pancreatic Cancer

Temel Tirkes, MD • Zarine K. Shah, MD • Naoki Tahahashi, MD • Joseph R. Grajo, MD • Stephanie T. Chang, MD • Sudhakar K. Venkatesh, MD • Darwin L. Conwell, MD • Evan L. Fogel, MD • Walter Park, MD • Mark Topazian, MD • Dirniy Yadan, MD, MPH • Anil K. Dasyam, MD • For the Consortium for the Study of Chronic Puncrentitis, Diabetes, and Puncrentic Cancer

Tirkes T, et al. *Radiology*. 2019 Jan;290(1):207-215

Table 2: Guidelines for Imaging of CP by Using MRI, MR Cholangiopancreatography, and CT				
MR Protocol Guidelines	CT Protocol Guidelines			
Axial T1-weighted 3D two-point Dixon gradient-echo sequence generating water-only and fat-only images in addition to in-phase and opposed-phase images	All imaging must be performed with a multidetector helical CT scanner.			
T2-weighted images (axial with fat suppression)	Scanning settings including kV, mAs, and pitch can be per individual institutional preference.			
T2-weighted images: fast spin-echo or a variant of fast spin-echo (axial and coronal without fat suppression)	Imaging should be performed per pancreas protocol, which should include unenhanced and contrast material—enhanced phases (specifically, pancreatic parenchymal and portal venous phases).			
MR cholangiopancreatography images	Water should be administered as oral contrast agent.			
2D MR cholangiopancreatography: 40-mm thick, eight paracoronal projections to best show the pancreatic duct from different angles	Scanning should be performed from hepatic dome to iliac crest in all phases. Scanning in portal venous phase can extend from hepatic dome to pubic symphysis.			
3D MR cholangiopancreatography: 2–3 mm, respiratory synchronized 3D fast spin-echo sequence. Create maximum intensity projection from source images.	Axial images should be reconstructed preferably at thickness of \leq 2.5 mm.			
Secretin-stimulated 2D MR cholangiopancreatography (thick slab): Secretin dose for adults is 16 µg administered via slow intravenous injection over 1 min and flushed with 20 mI of saline. After infusion, ductal system is imaged with a coronal single-shot fast spin-echo sequence, repeated every 30 sec up to 10 min.	Contrast dose should preferably be 100–150 mL but can be altered at the discretion of the radiologist.			
Postcontrast T1 weighted images: 3D gradient-echo with fat suppression (precontrast, arterial, portal venous, 5-min delayed venous phases)	Contrast injection rate should be greater than or equal to 2 mL/sec, preferably 4–5 mL/sec.			
Note.— CP = chronic pancreatitis, $3D$ = three-dimensional, $2D$ = two-dimensional.				

Findings of Pancreatitis

Acute Pancreatitis

- Swelling
- Edema
- Fluid collections
- Necrosis

Chronic Pancreatitis

Parenchyma

- Atrophy
- Loss of T1 signal
- Calcifications*

Duct

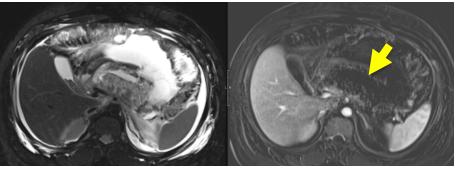
- Dilation
- Irregularity / Strictures
- Side branch dilation
- Filling defects

Findings of Pancreatitis

Acute Pancreatitis

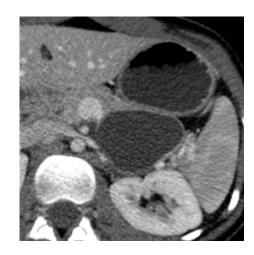
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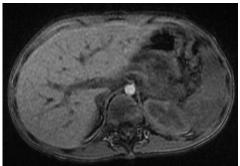
Findings of Pancreatitis – Fluid Collections

Fluid Collection	Description
Acute Peripancreatic Fluid Collection (APFC)	- Assoc. w/ interstitial edematous pancreatitis- No necrosis- No intrapancreatic extension- Simple fluid
Acute Necrotic Collection (ANC)	- Assoc. w/ necrotizing pancreatitis- Fluid + necrotic debris- Intra and/or extrapancreatic
Pseudocyst	- Develops from APFC - Encapsulated - Simple fluid
Walled off Necrosis (WON)	Develops from ANCEncapsulatedComplex



Findings of Pancreatitis







Chronic Pancreatitis

Parenchyma

- Atrophy
- Loss of T1 signal
- Calcifications*

Duct

- Dilation
- Irregularity / Strictures
- Side branch dilation
- Filling defects

Future Directions

- Faster MRI protocols
- Quantitative Imaging



Faster MRI Protocols

MR Protocol Guidelines

Axial T1-weighted 3D two-point Dixon gradient-echo sequence generating water-only and fat-only images in addition to in-phase and opposed-phase images T2-weighted images (axial with fat suppression)

T2-weighted images: fast spin-echo or a variant of fast spin-echo (axial and coronal without fat suppression)

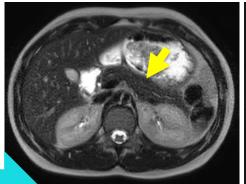
MR cholangiopancreatography images

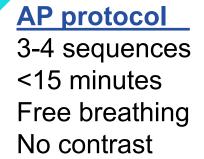
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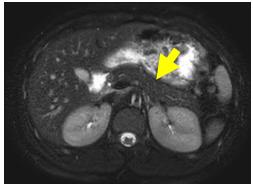
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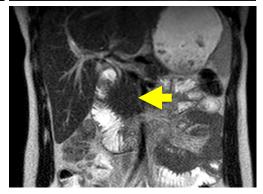
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Postcontrast T1 weighted images: 3D gradient-echo with fat suppression (precontrast, arterial, portal venous, 5-min delayed venous phases)







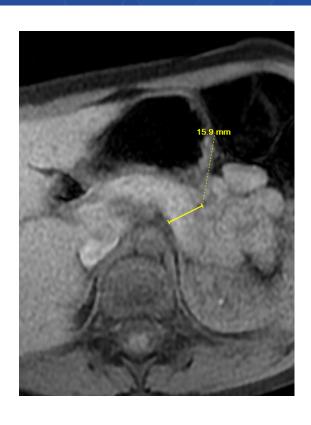


Quantitative Imaging

- Pancreatic measurements
- Elastography
- MR signal mapping
- Secretin for exocrine function



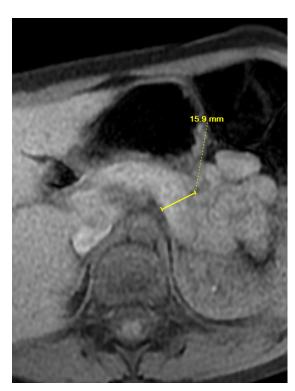
Quantitative Imaging – Panc. measurements



Norms exist

- Adult
 - Heuck A, et al. Gastrointest Radiol. 1987;12(1):18-22
- Pediatric
 - CT: Trout AT, et al. Pediatr Radiol.
 2018 Oct;48(11):1600-1605
 - Ultrasound: Siegel MJ, et al.
 Radiology. 1987 Oct;165(1):15-8

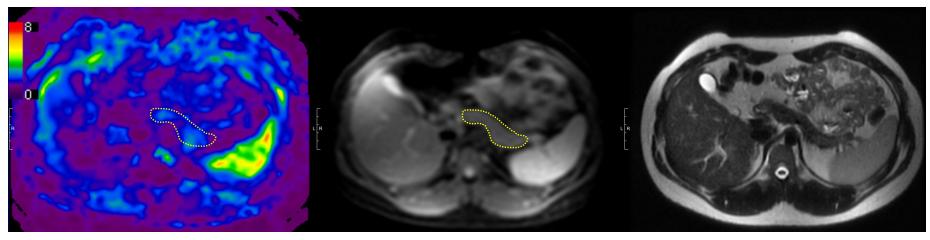
Quantitative Imaging – Panc. measurements

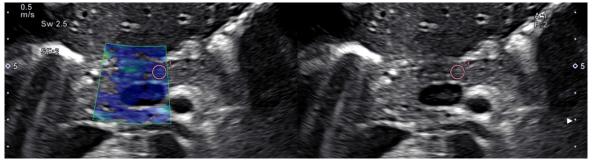






Quantitative Imaging - Elastography





Quantitative Imaging - Elastography

- MR elastography
 - Adult ↑ stiffness in tumors and CP
 - Pediatric ↓ stiffness in CP/ARP

- Ultrasound elastography
 - Adult ↑ stiffness in tumors and CP
 - Pediatric stiffness similar to healthy liver

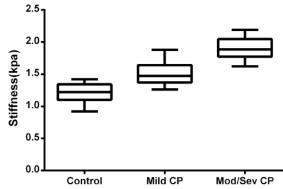
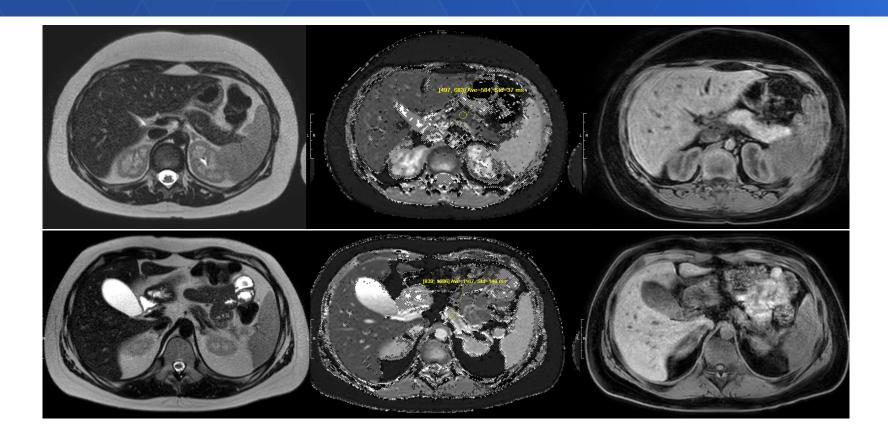


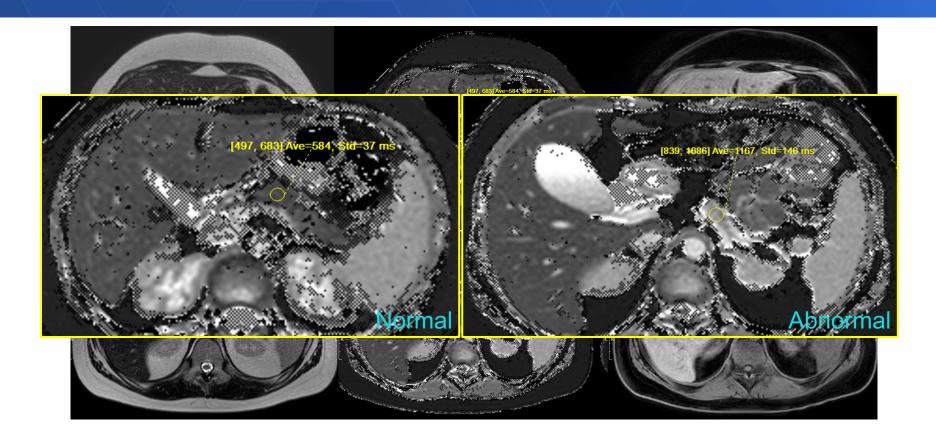
FIGURE 5: Boxplot of pancreatic stiffness in the control (mean=1.21 kPa, 95% CI: 1.17–1.26), mild CP group (mean=1.50 kPa, 95% CI: 1.44–1.56), and mod/sev CP (mean=1.90 kPa, 95% CI: 1.81–1.98). The stiffness of the pancreas in the mod/sev CP and mild CP group were significantly greater than the normal group (both P<0.001).

Wang M, et al. *J Magn Reson Imaging*. [Epub ahead of print] PubMed PMID: 29537715

Quantitative Imaging – Signal Mapping



Quantitative Imaging – Signal Mapping



Quantitative Imaging - Signal Mapping

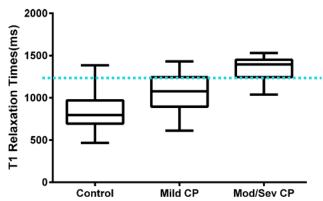
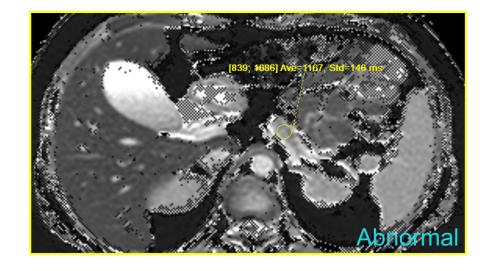


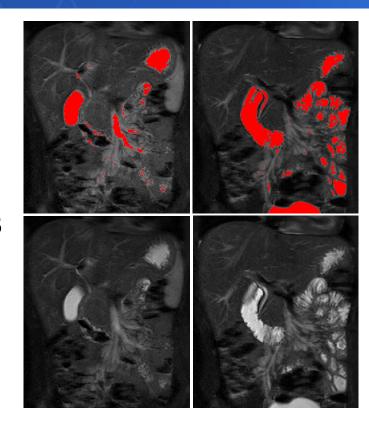
FIGURE 6: Boxplot of pancreatic T_1 relaxation times in the control (mean = 865 msec, 95% CI: 789–940) and mild CP group (mean = 1075 msec, 95% CI: 993–1158) and mod/sev CP (mean = 1350 msec, 95% CI: 1276–1423). The mean T_1 relaxation time of the pancreas in the mild CP and mod/sev CP group were significantly greater than the normal group (P=0.001 and <0.001, respectively).

Wang M, et al. *J Magn Reson Imaging*. [Epub ahead of print] PubMed PMID: 29537715



Quantitative Imaging – Exocrine fxn.

- Volumetric quantification of secreted fluid volume
- Norms exist
 - Adult: Mensel B, et al. AJR Am J
 Roentgenol. 2014 Jan;202(1):102-8
 - Peds: Trout AT, et al. Am J
 Gastroenterol. 2018
 Sep;113(9):1385



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