Chronic Pancreatitis: Diagnosis and Treatment

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Chronic pancreatitis is an irreversible and progressive disorder of the pancreas characterized by inflammation, fibrosis, and scarring. Exocrine and endocrine functions are lost, often leading to chronic pain. The etiology is multifactorial, although alcoholism is the most significant risk factor in adults. The average age at diagnosis is 35 to 55 years. If chronic pancreatitis is suspected, contrast-enhanced computed tomography is the best imaging modality for diagnosis. Computed tomography may be inconclusive in early stages of the disease, so other modalities such as magnetic resonance imaging, magnetic resonance cholangiopancreatography, or endoscopic ultrasonography with or without biopsy may be used. Recommended lifestyle modifications include cessation of alcohol and tobacco use and eating small, frequent, low-fat meals. Although narcotics and antidepressants provide the most pain relief, one-half of patients eventually require surgery. Therapeutic endoscopy is indicated to treat symptomatic strictures, stones, and pseudocysts. Decompressive surgical procedures, such as lateral pancreaticojejunostomy, are indicated for large duct disease (pancreatic ductal dilation of 7 mm or more). Resection procedures, such as the Whipple procedure, are indicated for small duct disease or pancreatic head enlargement. The risk of pancreatic cancer is increased in patients with chronic pancreatitis, especially hereditary pancreatitis. Although it is not known if screening improves outcomes, clinicians should counsel patients on this increased risk and evaluate patients with weight loss or jaundice for neoplasm. (Am Fam Physician. 2018;97(6):385-393. Copyright © 2018 American Academy of Family Physicians.)

Chronic pancreatitis is a permanent, progressive destruction of pancreatic tissue and function. Clinical manifestations include disabling abdominal pain, steatorrhea, and diabetes mellitus. ¹⁻³ Incidence and prevalence remain low and, despite advances in medical imaging, definitive diagnosis remains challenging. ³ Primary care physicians should be familiar with presenting clinical features, as well as treatment options and long-term complications.

Etiology

Data from case series and cross-sectional studies estimate that the incidence of chronic pancreatitis is between

Additional content at http://www.aafp.org/afp/2018/0315/p385.html.

CME This clinical content conforms to AAFP criteria for continuing medical education (CME). See CME Quiz on page 370.

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Patient information: A handout on this topic is available at http://www.aafp.org/afp/2007/1201/p1693.html.

about four and 12 per 100,000 persons per year. Data on prevalence are scarce, with estimates ranging from 37 to 42 per 100,000 persons.³ Men are affected 1.5 to 3 times more than women.⁴ The average age at diagnosis is 35 to 55 years. Alcoholism is the most significant risk factor for the development of chronic pancreatitis, accounting for 70% of cases in adults. Genetic disease, especially cystic fibrosis, and anatomic abnormalities are the most common causes in children. *Table 1* includes the TIGAR-O (toxic-metabolic, idiopathic, genetic, autoimmune, recurrent and severe acute pancreatitis, obstructive) classification of common risk factors for chronic pancreatitis.^{4,5}

Pathophysiology

The pathophysiology of chronic pancreatitis involves progressive fibrotic destruction of the pancreas in response to inflammation. A simple model involving the "two-hit" theory has been proposed. The first hit is acute pancreatitis, causing injury to the pancreas. The second hit is an abnormal inflammatory response to this injury. This causes sustained activation of pancreatic profibrotic cells, including stellate cells. In at-risk individuals, this can result in

SORT: KEY RECOMMENDATIONS FOR PRACTICE

Clinical recommendation	Evidence rating	References
Contrast-enhanced computed tomography is the recommended initial imaging study in patients with suspected chronic pancreatitis.	С	3, 10, 22-24
Endoscopic ultrasonography is favored over endoscopic retrograde cholangiopancreatography for the endoscopic diagnosis of chronic pancreatitis because of its increased safety and ability to evaluate the pancreatic parenchyma and duct system.	С	3
Antioxidant therapy does not improve pain control or mortality outcomes in patients with chronic pancreatitis.	В	10, 35
Pancreatic enzyme replacement is indicated for steatorrhea and malabsorption and may help relieve pain in patients with chronic pancreatitis.	В	38-42
Endoscopic drainage of pseudocysts results in a similar rate of pain relief as surgery, with equivalent or lower mortality.	В	25, 44-47, 53, 54
Pancreatoduodenectomy (Whipple procedure, pylorus- preserving, or duodenum-preserving) is indicated in the treatment of chronic pancreatitis with pancreatic head enlargement and typically results in significant pain relief.	В	8, 49, 55
\mathbf{A} = consistent good-quality patient-oriented evidence: \mathbf{B} = incor	nsistent or lim	nited-quality nations

 $\mathbf{A}=$ consistent, good-quality patient-oriented evidence; $\mathbf{B}=$ inconsistent or limited-quality patient-oriented evidence; $\mathbf{C}=$ consensus, disease-oriented evidence, usual practice, expert opinion, or case series. For information about the SORT evidence rating system, go to http://www.aafp.org/afpsort.

collagen deposition and fibrosis, which can lead to chronic pancreatitis.^{2-4,6-8} However, a subset of chronic pancreatitis is caused by autoimmune and genetic factors. Chronic pancreatitis is autoimmune in 5% to 6% of cases.⁹ Autoimmune pancreatitis has a distinctive histologic appearance with lymphocytic infiltration, but also results in fibrosis and loss of pancreatic function.³

Diagnosis of Chronic Pancreatitis

CLINICAL PRESENTATION

Patients most commonly present with recurrent episodes of acute pancreatitis. This will often progress to chronic abdominal pain that is characteristically located in the epigastrium and radiates to the back. The pain is worsened by eating and relieved by leaning forward. In time, it may spontaneously resolve as the pancreas fails.^{2,10-12} Although most patients present with pain, pancreatitis is painless in roughly 10% to 20% of patients.¹ When about 90% of pancreatic function is lost, patients will have signs of exocrine dysfunction, such as steatorrhea, malabsorption, and fatsoluble vitamin deficiencies (A, D, E, K, B₁₂).^{1,10,13,14} Endocrine dysfunction, such as diabetes, will also develop.^{1,13}

DIFFERENTIAL DIAGNOSIS

Table 2 includes a comprehensive differential diagnosis. ¹⁵ Amylase and lipase levels may be elevated in patients with

chronic pancreatitis, but these elevations may occur with other conditions that also present with abdominal pain, such as mesenteric ischemia, biliary disease, renal disease, and complicated peptic ulcer disease.^{2,13,16-18}

DIAGNOSTIC TESTING

The diagnosis of chronic pancreatitis is made based on a patient's history, clinical presentation, and imaging findings. There is no single test that is diagnostic of chronic pancreatitis, especially in the early stages. If chronic pancreatitis is suspected based on history and physical examination, a diagnostic workup should be initiated. Diagnostic tests should be

chosen based on their availability after consideration of risks and benefits.^{1,2,4,9,18-20} Laboratory tests used in the evaluation of patients with suspected chronic pancreatitis are summarized in *Table 3.*¹⁵

Laboratory Testing. In acute pancreatitis, pancreatic enzymes are elevated more than three times the upper limit of normal. ^{16,18} In chronic pancreatitis, these enzymes may be only mildly elevated or normal, especially as the pancreas is replaced by increasing amounts of fibrotic tissue later in the disease process. ¹ If the biliary tract is obstructed

WHAT IS NEW ON THIS TOPIC

Chronic Pancreatitis

A meta-analysis of 43 studies that included more than 3,400 patients concluded that computed tomography, magnetic resonance imaging, endoscopic retrograde cholangiopancreatography, and endoscopic ultrasonography have comparably high diagnostic accuracy for chronic pancreatitis; therefore, a stepwise approach based on cost, invasiveness, and availability is recommended.

A Cochrane review of 10 trials involving 361 participants suggested that pancreatic enzyme replacement does not reduce pain from chronic pancreatitis.

TABLE 1

TIGAR-O Classification of Risk Factors Associated with Chronic Pancreatitis

Toxic-metabolic

Alcohol

Chronic renal failure

Hypercalcemia (hyperparathyroidism)

Hyperlipidemia (rare)

Medications*

Tobacco

Toxins

Idiopathic

Early and late onset

Tropical pancreatitis (tropical calcifying pancreatitis and fibrocalculous pancreatic diabetes)

Genetic

Autosomal dominant (cationic trypsinogen [codon 29 and 122 mutations])

Autosomal recessive modifier genes (*CFTR* and *SPINK1* mutations, cationic trypsinogen [codon 16, 22, and 23 mutations], alpha₁-antitrypsin deficiency)

Autoimmune

Autoimmune chronic pancreatitis associated with inflammatory bowel disease, Sjögren syndrome, primary biliary cirrhosis

Isolated autoimmune chronic pancreatitis

Recurrent and severe acute pancreatitis

Postirradiation

Postnecrotic (severe acute pancreatitis)

Recurrent acute pancreatitis

Vascular ischemia

Obstructive

Duct obstruction (pancreatic or ampullary tumors) posttraumatic pancreatic duct fibrosis)

Pancreas divisum

Sphincter of Oddi disorders

*—Drug-induced pancreatitis is mostly acute or recurrent acute pancreatitis, specifically in high-risk populations (e.g., older persons; patients with human immunodeficiency virus infection, cancer, or other immunocompromising condition). Common drugs that may induce chronic pancreatitis include angiotensin-converting enzyme inhibitors, statins, didanosine (Videx), azathioprine (Imuran), steroids, lamivudine (Epivir), hydrochlorothiazide, valproic acid (Depakene), oral contraceptives, and interferon.

Adapted with permission from Etemad B, Whitcomb DC. Chronic pancreatitis: diagnosis, classification, and new genetic developments. Gastroenterology. 2001;120(3):691, with additional information from reference 5.

(5% to 10% of cases), bilirubin, alkaline phosphatase, and hepatic transaminases may be elevated.^{1,16}

Pancreatic function tests use a catheter inserted into the pancreatic duct to directly measure pancreatic enzyme secretion. These tests are expensive and challenging to

TABLE 2

Differential Diagnosis of Chronic Pancreatitis

Pancreatitis	
More common	Less common
Acute cholecystitis	Acute appendicitis
Acute pancreatitis	Acute salpingitis
Intestinal ischemia	Crohn disease
or infarction	Ectopic pregnancy
Obstruction of common bile duct Pancreatic tumors Peptic ulcer disease Renal insufficiency	Gastroparesis
	Intestinal obstruction
	Irritable bowel syndrome
	Malabsorption
	Ovarian cyst
	Papillary cystadenocarcinoma of the ovary
	Thoracic radiculopathy

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perform. The positive predictive value of these tests is just 45%, but the negative predictive value is 97%. Therefore, although they are not recommended as part of the routine workup, they can be useful for ruling out chronic pancreatitis in patients who exhibit clinical features of pancreatitis but have normal imaging.^{1,2,4,10,19-21} Fecal elastase is an indirect measure of pancreatic exocrine function that has high sensitivity (65% to 100%) and poor specificity (55%). The high false-positive rate makes this a poor screening test, but it can be helpful in ruling out pancreatitis.¹⁹

Imaging Studies. The radiologic evaluation of a patient with suspected chronic pancreatitis should progress from least invasive to more invasive. A meta-analysis of 43 studies that included more than 3,400 patients concluded that computed tomography (CT), magnetic resonance imaging (MRI), endoscopic retrograde cholangiopancreatography, and endoscopic ultrasonography (EUS) have comparably high diagnostic accuracy; therefore, a stepwise approach based on cost, invasiveness, and availability is recommended.²² eTable A includes the sensitivity and specificity of available imaging techniques. Contrast-enhanced CT of the pancreas is the first-choice modality because it is noninvasive and readily available.3 Ductal calcifications are pathognomonic findings of chronic pancreatitis (Figure 115). CT can detect conditions that mimic chronic pancreatitis, as well as complications of chronic pancreatitis, such as pseudocysts, ductal dilation, pseudoaneurysms, necrosis, and parenchymal atrophy. 3,10,22-24

If CT findings are equivocal, patients may require referral for more focused pancreatic imaging, such as MRI or magnetic resonance cholangiopancreatography, or for endoscopic procedures, such as EUS or endoscopic retrograde cholangiopancreatography.³ Magnetic resonance

cholangiopancreatography and EUS are useful for evaluating the pancreatic parenchyma and duct system. Endoscopic retrograde cholangiopancreatography has a high risk of complications (e.g., pancreatitis, hemorrhage, infection), so it is not recommended for

APPROACH TO A PATIENT WITH SUSPECTED CHRONIC PANCREATITIS

diagnosis unless all other tests are inconclusive.^{2,25}

Figure 2 shows a five-step diagnostic approach to patients with signs and symptoms of chronic pancreatitis. CT will detect involvement of large duct disease (pancreatic ductal dilation of 7 mm or more).2,4,10 CT may be inconclusive in patients with early disease or small duct disease (pancreatic ductal dilation less than 7 mm). These patients should be referred to a center with expertise in pancreatic care for an MRI or magnetic resonance cholangiopancreatography.3 These imaging modalities, with EUS, are also useful for preoperative planning in patients with stones, strictures, or pseudocysts.2-4,23,25-27 If a cystic lesion or mass is identified on imaging, malignancy must be excluded through EUS with biopsy and tumor marker analysis, specifically cancer antigen 19-9.2,3,28,29 If chronic pancreatitis is still suspected despite normal imaging findings, pancreatic function tests can be

TABLE 3

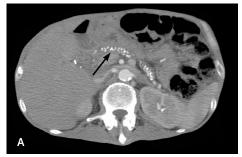
Laboratory Tests Used in the Evaluation of Patients with Suspected Chronic Pancreatitis

Tests	Comments
Complete blood count	Elevated with infection, abscess
Serum amylase and lipase	Nonspecific for chronic pancreatitis
Total bilirubin, alkaline phosphatase, and hepatic transaminase	Elevated in biliary pancreatitis and ductal obstruction by strictures or mass
Fasting serum glucose	Elevation suggests pancreatic diabetes mellitus
Pancreatic function tests	Sometimes useful in early chronic pancreatitis with normal computed tomography or magnetic resonance imaging findings
Fecal elastase	< 200 mcg per g of stool is abnormal; noninvasive, exogenous pancreatic supplementation will not alter results, requires only 20 g of stool
Fecal fat estimation	> 7 g of fat per day is abnormal; quantitative, requires 72 hours, should be on a diet of 100 g of fat per day
Secretin stimulation	Peak bicarbonate concentration < 80 mEq per L (80 mmol per L) in duodenal secretion, best test for diagnosing pancreatic exocrine insufficiency
Serum trypsinogen	< 20 ng per mL is abnormal
Lipid panel	Significantly elevated triglycerides are a rare cause of chronic pancreatitis
Calcium	Hyperparathyroidism is a rare cause of chronic pancreatitis
Immunoglobulin G4 serum antibody, antinuclear antibody, rheumatoid factor, erythrocyte sedimentation rate	Abnormality may indicate autoimmune pancreatitis

Note: Tests are listed in order of most to least commonly performed.

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FIGURE 1





Contrast-enhanced computed tomography of the upper abdomen showing (A) pancreatic calcifications (arrow) with fluid and edema around the pancreas; and (B) pancreatic calcifications (arrow) with fluid and edema around the head of the pancreas.

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FIGURE 2 Clinical signs and symptoms of chronic pancreatitis Abdominal pain Weight loss Steatorrhea Malabsorption History of alcohol abuse History of recurrent pancreatitis Perform history and physical examination Obtain laboratory testing Step 1 Computed tomography Diagnostic criteria met: no further imaging Inconclusive or nondiagnostic: proceed to step 2 Step 2 Magnetic resonance imaging/magnetic resonance cholangiopancreatography Diagnostic criteria met: no further imaging Inconclusive or nondiagnostic: proceed to step 3 Step 3 Endoscopic ultrasonography Diagnostic criteria met: no further imaging Inconclusive or nondiagnostic: proceed to step 4 Step 4 Pancreatic function tests Diagnostic criteria met: no further imaging Inconclusive or nondiagnostic: proceed to step 5 Step 5 Endoscopic retrograde cholangiopancreatography Diagnostic criteria met: chronic pancreatitis Inconclusive or nondiagnostic: monitor symptoms and repeat imaging in six months to one year A stepwise approach to imaging for chronic pancreatitis.

performed.^{2-4,27-29} A small study of 25 patients suggests that combining EUS and pancreatic function tests can improve the sensitivity of diagnosis.³⁰

Treatment

Treatment of chronic pancreatitis can be medical, endoscopic, or surgical (*Table 4*).¹⁵ The goals of all treatment modalities are improved pain, improved quality of life,

TABLE 4

Chronic Pancreatitis Treatment Options

Medical

Analgesics (stepwise approach)

Antidepressants (treatment of concurrent depression)

Cessation of alcohol and tobacco use

Denervation (celiac nerve blocks, transthoracic

splanchnicectomy)

Insulin (for pancreatic diabetes)

Low-fat diet and small meals

Pancreatic enzymes with proton pump inhibitors or histamine H₂ blockers

Steroid therapy (in autoimmune pancreatitis)

Vitamin supplementation (A, D, E, K, and B₁₂)

Endoscopic

Extracorporeal shock wave lithotripsy with or without endoscopy

Pancreatic sphincterotomy and stent placement for pain relief

Transampullary or transgastric drainage of pseudocyst

Surgical

Decompression

Cystenterostomy

Lateral pancreaticojejunostomy (most common)

Sphincterotomy or sphincteroplasty

Resection

Distal or total pancreatectomy

Pancreatoduodenectomy (Whipple procedure, pylorus-preserving, duodenum-preserving)

Not recommended

Allopurinol

Antioxidant therapy (vitamin C, vitamin E, selenium, methionine [no longer available in the United States])

Octreotide (Sandostatin)

Prokinetic agents (erythromycin)

Adapted with permission from Nair RJ, Lawler L, Miller MR. Chronic pancreatitis. Am Fam Physician. 2007;76(11):1684.

and decreased morbidity and mortality from endocrine and exocrine pancreatic dysfunction. Studies that compare conservative medical therapy with invasive endoscopic or surgical interventions are lacking. Therefore, treatment decisions should be based on careful consideration of the goals of treatment and the risks vs. benefits.^{31,32}

MEDICAL

Eliminating alcohol and tobacco use slows disease progression and lessens complications, such as cancer.³³ Dietary changes, particularly eating small, frequent, low-fat meals, can also help decrease pain and complications.^{2,3,10,34}

Disabling pain is the most common symptom of chronic pancreatitis. Because treatment with narcotic pain medications can lead to opioid addiction, beginning with nonsteroidal anti-inflammatory drugs or acetaminophen is

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recommended. Adding narcotic pain medications in a stepwise approach may be necessary to obtain adequate pain control. Adjunctive therapy with tricyclic antidepressants, selective serotonin reuptake inhibitors, and combined serotonin and norepinephrine reuptake inhibitors is used to act in synergy with narcotics and treat concomitant depression that is common in patients with chronic pancreatitis. 31,32

A Cochrane review of antioxidant therapy, including selenium, beta carotene, L-methionine, vitamin C, and vitamin E, did not find evidence of effectiveness.³⁵ Other studies found that octreotide (Sandostatin), allopurinol, or vitamin D provided no pain relief.¹⁰

In a three-week long study funded by the manufacturer, pregabalin (Lyrica) decreased short-term opioid use and improved short-term pain scores.^{36,37} However, studies with longer follow-up and larger sample size are needed before this becomes a standard treatment for chronic pancreatitis pain.

Pancreatic enzyme replacement is beneficial for the treatment of steatorrhea and malabsorption.³⁸⁻⁴¹ A Cochrane review of 10 trials involving 361 participants suggested that pancreatic enzyme replacement does not reduce pain from chronic pancreatitis, although results of individual studies were mixed.⁴²

ENDOSCOPY

Endoscopy can be used to treat symptomatic strictures, stones, and pseudocysts. Endoscopic drainage of pseudocysts has a similar rate of pain relief as surgery, with equivalent or lower mortality.^{25,43-48} Regression of the cyst occurs in 70% to 86% of patients after endoscopic drainage.^{25-28,43-46} Endoscopic sphincterotomy or stent placement can provide pain relief without the risks of surgery in patients with strictures.^{10,25,27} Extracorporeal shock wave lithotripsy with or without endoscopic drainage of the pancreatic duct can reduce stone burden and pain.^{10,25,49,50}

SURGERY

One-half of patients with chronic pancreatitis will eventually require surgery,¹ most commonly because of intractable, disabling pain.⁵¹ Indications for surgery are listed in *Table 5*.¹⁵ There are two types of surgeries for chronic pancreatitis. Decompression procedures are performed on patients with large duct disease, whereas resection procedures are performed on patients with small duct disease or pancreatic head enlargement.

TABLE 5

Chronic Pancreatitis: Indications for Surgery

Biliary or pancreatic stricture

Duodenal stenosis Fistulas (peritoneal or pleural

effusion) Hemorrhage

Intractable chronic abdominal pain

Pseudocysts

Suspected pancreatic neoplasm Vascular complications

Adapted with permission from Nair RJ, Lawler L, Miller MR. Chronic pancreatitis. Am Fam Physician. 2007;76(11):1684. The most common decompression procedure is lateral pancreaticojejunostomy (*Figure 3A*¹⁵). This procedure is more effective at providing long-term pain relief (60% to 91% of patients) than endoscopic approaches. ^{2,52-54} Surgery may also be performed to drain a symptomatic pseudocyst; however, because endoscopic approaches are as effective at relieving pain with fewer risks, surgery is reserved for when endoscopic intervention is ineffective or for complicated pseudocysts. ^{47,48}

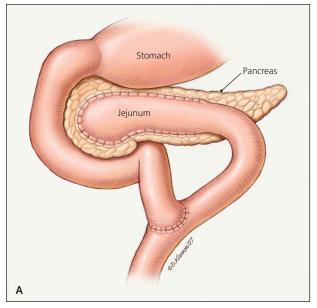
Resection procedures^{55,56} include pancreatoduodenectomy (Whipple procedure, *Figure 3B*¹⁵), pyloruspreserving pancreatoduodenectomy (*Figure 3C*¹⁵), duodenum-preserving

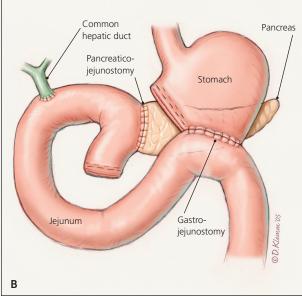
pancreatic head resection (Beger or Frey procedures), and total pancreatectomy (Figure 3D15). The Whipple procedure is the most commonly performed surgery in patients with chronic pancreatitis because of its high success rate (pain relief in 85% of patients) and low mortality (less than 3%). 8,55,56 Pancreatoduodenectomy is indicated with pancreatic head enlargement and typically results in significant pain relief.⁵¹ A Cochrane review of five trials including 292 participants compared the Whipple procedure with duodenumpreserving pancreatic head resection. Although hospital stay was shorter for duodenum-preserving pancreatic head resection, there was no statistical difference in mortality or quality of life. 57,58 Total pancreatectomy is rarely performed because of high rates of complications and low rates of pain relief. Because of the high risk of diabetes in patients with chronic pancreatitis, total pancreatectomy should always be performed with an autologous islet cell transplant.^{56,59}

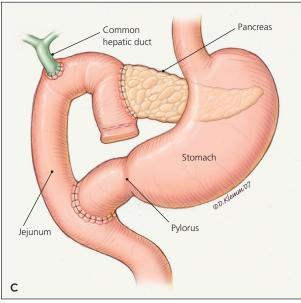
Complications

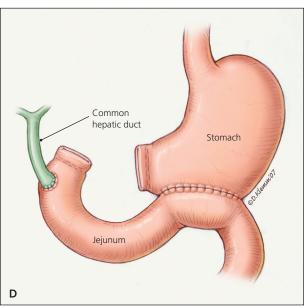
Complications of chronic pancreatitis are summarized in *Table* 6.13,15,55,60 The most common causes of morbidity and mortality are chronic debilitating pain, diabetes, pancreatic pseudocysts, and pancreatic cancer. Although pain is often present at diagnosis, it takes roughly five years for diabetes to develop. 12,13,55,61 Pancreatic pseudocysts are usually asymptomatic but may cause serious complications in 25% to 30% of patients with pancreatitis. Pseudocysts can lead to rupture, infection, bleeding, and obstruction. 10,13,54,55 Recurrent attacks of acute pancreatitis can cause pancreatic abscesses and necrosis, sepsis, and multiorgan failure. Palmost one-fourth of patients (23.4%) with chronic pancreatitis have osteoporosis and almost two-thirds (65%) have osteoporosis or osteopenia. 60 There is

FIGURE 3









Most patients with chronic pancreatitis undergo surgery when initial medical and endoscopic treatments do not relieve pain. Surgical interventions include (A) lateral pancreaticojejunostomy, (B) pancreatoduodenectomy (Whipple procedure), (C) pylorus-preserving pancreatoduodenectomy, and (D) total pancreatectomy.

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a markedly increased risk of pancreatic cancer in patients with chronic pancreatitis, particularly among those with long-term alcohol use or hereditary pancreatitis. 2,3,12,14,25-28

Surveillance

Patients with hereditary pancreatitis have a 10-fold increased risk of pancreatic cancer compared with the

general population. 61,62 Therefore, some experts suggest offering counseling and screening to patients with hereditary pancreatitis beginning at 40 years of age. The most recommended screening methods are EUS, CT, and endoscopic retrograde cholangiopancreatography; there is no consensus on screening interval. Additionally, patients with chronic pancreatitis who exhibit a change

TABLE 6

Complications of Chronic Pancreatitis

Complication	Incidence (%)
Acute pancreatitis	Recurrent
Chronic pain	80 to 90
Osteoporosis or osteopenia	65
Diabetes mellitus	> 40
Weight loss	> 40
Pseudocyst	25 to 30
Pancreatic cancer	15 to 40
Malabsorption and steatorrhea	10 to 15
Bile duct, duodenal, or gastric obstruction	5 to 10
Pancreatic ascites or pleural effusion	< 10
Pseudoaneurysm, especially of splenic artery	< 1
Splenic or portal vein thrombosis	< 1
Vitamin deficiency (A, D,* E, K, and B_{12})	Rare

^{*—}Vitamin D deficiency has recently been reported more often with pancreatic exocrine dysfunction.

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in clinical symptoms (pain, weight loss, jaundice) should be evaluated for neoplasm.⁶³ All patients at increased risk of cancer should be counseled on mitigating risk factors, especially alcohol and tobacco use.^{62,63} If a malignancy is suspected, a triphasic, pancreas protocol CT is the standard for diagnosis. If this test is not available or if the patient has an allergy to contrast media, MRI with gadolinium may be used.⁶²

This article updates a previous article on this topic by Nair, et al. 15

Data Sources: A PubMed search was completed in Clinical Queries using the key terms chronic pancreatitis and pancreatitis. The search included meta-analyses, randomized controlled trials, clinical trials, and reviews. Also searched were the Agency for Healthcare Research and Quality evidence reports, Clinical Evidence, the Cochrane database, the Institute for Clinical Systems Improvement, the National Guideline Clearinghouse database, Dynamed, and Essential Evidence Plus. Search dates: March 18, 2016, and September 20, 2017.

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BONUS DIGITAL CONTENT

eTABLE A

Imaging and Endoscopic Studies for Diagnosing Chronic Pancreatitis

Test	Sensitivity (%)	Specificity (%)	Comment
Contrast-enhanced CT	75 to 90	85 to 91	Initial radiologic test of choice for evaluation of suspected chronic pancreatitis; can visualize calcifications, pseudocysts, thrombosis, pseudoaneurysms, necrosis, and atrophy
Endoscopic retrograde cholangiopancreatog-raphy	75 to 95	90 to 94	Reference standard in many studies, invasive and associated with complications, mainly used in diagnosis of early chronic pancreatitis with normal CT and pancreatic function tests
Endoscopic ultrasonography	81 to 97	60 to 90	Useful in evaluation of early chronic pancreatitis, pancreatic mass, and cystic lesions; can be combined with fine-needle aspiration biopsy
Magnetic resonance imaging or magnetic resonance cholan-giopancreatography	78	96 to 100	Noninvasive and nonionizing radiation or contrast media, less sensitive than endoscopic retrograde cholangiopancreatography for evaluation of side branches, can be combined with secretin test
Plain abdominal radiography	NA	NA	Not routinely recommended; calcifications may be visible
Ultrasonography	67	98	Rarely diagnostic; may be useful for ultrasonography-guided aspiration of cyst

CT = computed tomography; NA = not available.

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Additional information from:

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