Endocrine and Exocrine Functions of the Pancreas

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The pancreas is a 2-in-1 organ: an exocrine and endocrine gland. It is essential for digestion and the carbohydrate metabolism. Thus, a loss in pancreatic function leads to severe clinical symptoms. In this article, you will get a compact overview of the structure, functions, and diseases of the pancreas.

Macroscopic Anatomy of the Pancreas
Location of the pancreas

The pancreas is an elongated, tapered organ that is located across the back of the abdomen, right behind the stomach. It has a sigmoid shape and is organized in lobules, which are separated from each other by projections of the thin fibrous capsule that surrounds the organ, and by fine connective tissue septa.

Apart from the tail, the pancreas is a retroperitoneal organ that lies behind the peritoneal cavity, and is located deeply within the upper abdomen in the epigastrium and left hypochondrium regions. Some of its relations include the stomach superiorly and anteriorly, the colon and spleen laterally on the right and left respectively. The head of the pancreas is located on the right side of the abdomen and is connected to the duodenum through a small tube called the pancreatic duct.

The different parts can be distinguished as:

- **Caput pancreatis** (head of the pancreas): Lies in the concave aspect of the duodenum (duodenal C) and has a hook-shaped process (uncinate process), which surrounds the superior mesenteric artery and vein.
- **Corpus pancreatis** (body of the pancreas): Represents the main part of the pancreas and is contained within the posterior, retroperitoneal abdominal wall. The anterior part is covered with peritoneum and forms the posterior wall of the omental bursa.
- **Cauda pancreatis** (tail of the pancreas): Stretches to the spleen and cannot be clearly differentiated from the body.
The microscopic anatomy involves 99% exocrine and 1% endocrine tissue by weight that is organized into acini that surround multiple ducts. The ducts eventually drain into the excretory duct which is the main pancreatic duct (or Wirsung’s duct). It runs across the whole gland and leads to the major duodenal papilla along with the common bile duct.

Due to the development of the pancreas out of 2 separate anlages (ventral and dorsal anlage), another accessory pancreatic duct (or Santorini’s duct) commonly exists, which leads to the minor duodenal papilla. The pancreatic duct runs along the length of the pancreas uniting with the common bile duct, forming what is called the hepatopancreatic ampulla of Vater. This structure opens into the duodenum.

**Vascular supply of the pancreas**

The celiac trunk and the superior mesenteric artery ensure the arterial blood supply of the pancreas. The arteries reach the organ from both the cranial and the caudal side of the pancreas and form anastomoses in the area of the pancreatic head. This construction is referred to as the pancreatic arcade. It includes:

- Posterior and anterior superior pancreaticoduodenal artery (from the gastroduodenal artery, which originates from the celiac trunk)
- Inferior pancreaticoduodenal artery (out of the superior mesenteric artery)

Additionally, the splenic artery in the area of the body and the tail of the pancreas gives rise to pancreatic rami, the dorsal pancreatic artery, and the greater pancreatic artery, which forms an anastomosis with the inferior pancreatic artery at the lower margin of the pancreas.

The pancreas drains into the splenic vein and the superior mesenteric vein, which eventually form the portal vein. Due to the close relationship between the pancreatic head and these vessels, cancer of the pancreatic head may cause constriction of the portal vein resulting in portal hypertension and stasis with ascites.
The pancreas is located in the posterior abdominal wall in a retroperitoneal manner. Thus, it has close topographic relations with a lot of structures.

- **Stomach**: The omental bursa separates the 2 organs. It lies anteriorly and superiorly.

- **Small intestine and duodenum**: While the small intestine can reach the lower margin of the pancreas anteriorly and medially, the pancreatic head clings closely to the duodenum.

- **Left kidney**: The pancreatic tail runs over the kidney.

- **Spleen**: The tail of the pancreas ends at the hilum of the spleen. The spleen is located posteriorly and laterally. It is connected by ligaments to the tail of the pancreas.

- **Common bile duct**: It can be embedded in the pancreas. Cancer of the pancreatic head can block the common bile duct, which results in obstructive jaundice.

- **Superior mesenteric artery and vein**: The vessels run dorsally to the head of the pancreas and emerge on the front side at the pancreatic notch. This is where they are entwined by the uncinate process. Cancer of the pancreatic head can spread to these 2 vessels, which can lead to circulatory disorders of several organs.

- **Splenic artery and vein**: The splenic artery is located on the upper margin of the pancreas, while the splenic vein runs deeper and behind the gland.
Malformations of the pancreas

The pancreas develops as 2 outpouchings from the embryonic foregut: the dorsal and ventral pancreatic buds. Both remain attached to the gut and as the duodenum rotates during development, it takes with it the dorsal anlage leading to fusion of both the dorsal and ventral anlage to form the adult pancreas. The dorsal anlage forms the head, neck, body, and tail of the pancreas while the ventral anlage forms the uncinate process of the pancreas.

Pancreas divisum

If the ventral and dorsal pancreatic anlages do not merge during pancreatic development, a pancreas divisum occurs, which is characterized by 2 separate glands. The merging of the 2 pancreatic ducts stops here so that they can flow separately into the duodenum: the major pancreatic duct at the major duodenal papilla and the minor pancreatic duct at the more cranial lying minor duodenal papilla. This is a frequently occurring abnormality, which mostly does not cause any symptoms. In some cases, this anomaly is associated with recurrent episodes of pancreatitis. However, its role as a predisposition for pancreatitis is controversial.

Annular pancreas

During the development of the pancreas, a part of the ventral anlage (the right bud) moves around the duodenum from a dorsal location, while the other part (the left bud) regresses. If this regression does not occur, the left bud rotates around the duodenum ventrally and fuses with the right bud, which results in an annular pancreas: the descending part of the duodenum is encircled by pancreatic tissue. This rare anomaly can cause duodenal atresia.

Accessory pancreatic lobe

The accessory pancreatic lobe is a rare condition that presents with an additional pancreatic lobe arising from the main pancreatic gland with its own aberrant duct. The anomaly may present with recurrent pancreatitis.

Agenesis/hypoplasia of the pancreas

Total agenesis is incompatible with life and rarely occurs. However, hypoplasia/ lack of
development of 1 anlage may be seen. It may predispose one to diabetes mellitus especially if the dorsal anlage is involved, where islet cells are located.

Microscopic Anatomy of the Pancreas

In contrast to the macroscopic anatomy, the pancreas’ organization in the exocrine and endocrine pancreas becomes apparent in histological sections.
Exocrine part of the pancreas

Histologically, the exocrine part of the pancreas shows the typical structure of serous gland and represents roughly 98% of the organ’s tissue. The acini consist of strong basophilic cells, which contain numerous secretory granules. Acini is made of zymogenic cells that surround a central lumen and is arranged in lobules.

Each lobule contains its own ductule, and many ductules join together to make intralobular ducts, which then form interlobular ducts that drain into branches of the main pancreatic duct. They contain inactive precursors of digestive enzymes, which are activated outside of the pancreas (see below) in order to prevent the organ from digesting itself. The acini are followed by intercalated ducts, which connect the acini to the ductal system.

A special feature concerning the histology of the pancreas is centroacinar cells, which represent invaginations of the intercalated ducts into the acinar lumen. With their bright coloration, they form a contrast to the acini and are easily identified in a histological picture. The pancreas lacks striated ducts, which can e.g., be found in the parotid gland. The ductal system consists of intralobular ducts with cuboidal epithelium, which fuse to form interlobular ducts with columnar epithelium. Eventually, these ducts lead to the major or minor pancreatic duct.

Endocrine part of the pancreas

About 2% of the pancreatic tissue has an endocrine function. It is represented by accumulations of cells (islets of Langerhans) within the exocrine tissue. With their weak eosinophilic coloration, they differ from the dark acini of the exocrine part. The islets can be found mainly in the tail of the pancreas.

With special immunohistochemical stains, 4 different cell populations, each producing a different hormone, can be distinguished within the islets of Langerhans: alpha cells (20%, particularly in the periphery, glucagon production), beta cells (70%, evenly spread, insulin production), delta cells (5%, somatostatin production), and PP cells (5%, pancreatic polypeptide production).

Function of the pancreas

An essential pancreas function is the conversion of food into fuel for the body’s cells. The exocrine part mainly produces digestive enzymes and helps in digestion, while the endocrine part is responsible for the regulation of the carbohydrate metabolism (blood
sugar) with its insulin and glucagon production.

Exocrine function of the pancreas

The pancreas produces roughly 1.5–2 L of digestive secretion, which leads to the duodenum via the pancreatic duct. This secretion consists of bicarbonate, which makes the fluid alkaline (pH 8), and of several enzymes, that are essential for digestion. Trypsinogen and chymotrypsinogen are inactive protease precursors (zymogens), which are activated by enterokinase in the intestinal lumen. These enzymes are secreted by zymogenic cells. The active enzymes trypsin and chymotrypsin are endopeptidases (fission of the peptide bond within a protein). This prevents the pancreatic tissue from digesting itself. Trypsin is able to activate other inactive enzymes.

Carboxypeptidases are also zymogens which are exopeptidases (fission of the peptide bond at the end of a protein) in their active form. Active enzymes in the pancreatic digestive secretion are pancreas lipase and phospholipase A2 which hydrolyze triglycerides and phospholipids, pancreas amylase for breaking bonds in polysaccharides, and deoxyribonucleases for the degradation of nucleic acids.

The release of these enzymes are regulated by hormones of the intestinal wall: secretin, which encourages the release of the alkaline secretion for neutralization of the acidic stomach content, and cholecystokinin, which stimulates the production and secretion of the enzymes. Gallstones in the common bile duct can cause secretion backflow. This can lead to premature activation of the enzymes with pancreatic self-digestion, which is referred to as acute pancreatitis (see below).

Endocrine function of the pancreas

The pancreas is not only digestive but also an endocrine gland. As such, it produces 2 important hormones of the carbohydrate metabolism: insulin and glucagon, amongst others. The anabolic hormone insulin is produced in the beta cells and is basally released in an oscillating manner. The half-life period in the blood is only 6–8 minutes. A high blood glucose value acts as the main stimulus for induced release. Insulin then promotes the translocation of the insulin-dependent glucose transporter GLUT 4 to the plasma membrane so that the target cells can now quickly ingest glucose which decreases the blood glucose value.

Insulin also promotes glycolysis and glycogen synthesis. Thus, the absorbed glucose is either transformed into energy or it is stored. The direct antagonist of insulin is glucagon, which is produced in the alpha cells of the pancreas. At low blood glucose levels, glucagon promotes glycoegenolysis and gluconeogenesis. It inhibits glycolysis and glycogen synthesis so that the blood glucose level rises.

Therefore, it is a catabolic hormone that has an antagonist-effect concerning insulin. Somatostatin, which is produced by the delta cells, inhibits the secretion of insulin, glucagon, pancreatic enzymes, and gastric acid. Pancreatic polypeptide is produced by PP cells and also has an inhibitory effect. It inhibits the release of pancreatic enzymes and the gallbladder release of bile.


Diseases of the Pancreas

Acute pancreatitis

Acute pancreatitis is characterized by autodigestion of pancreatic tissue due to premature activation of zymogens. The most frequent causes for this clinical picture are choledocholithiasis and alcohol abuse. The patients suffer from severe pain in the upper abdomen, which can spread to the back in a belt-like manner. Nausea and vomiting can occur, sometimes accompanied by fever. Ileus and jaundice are also possible symptoms. A circulatory shock, acute kidney failure, sepsis, respiratory arrest, heart failure, bleedings in the gastrointestinal tract and coma are feared complications.

![Image: “Acute pancreatitis” by Hellerhoff, License: CC BY-SA 3.0](image)

Up to 30% of these patients develop a pseudocyst that can regress spontaneously (see below). In the physical examination of a patient with acute pancreatitis, abdominal tenderness (elastic abdominal wall tension), reddening of the face, and sometimes bruising around the umbilicus (Cullen’s sign) and in the flank area (Grey Turner’s sign) stand out. An increase in amylase and lipase (> 3 times the normal value) in the serum can be found in the laboratory parameters, whereas lipase is specific to the pancreas.
An increase in gamma-glutamyl transferase (GGT), alkaline phosphatase (AP), serum glutamic-oxaloacetic transminase (SGOT), and bilirubin points to an obstruction of the common bile duct as a cause for pancreatitis.

**Treatment**

- Keep the patient *nil per oral* to rest the gland and avoid secretion and reactivation of the digestive function.
- Supportive therapy with fluids and electrolytes avoids dehydration and electrolyte imbalances in a patient who is *nil per oral* with diarrhea and vomiting.
- Gallstones have to be removed. If conservative therapy fails, surgery should be the next step.
Chronic pancreatitis

The most common cause (80%) for chronic pancreatitis is alcohol abuse. In 10% of the cases, no cause for the disease can be identified (idiopathic). In contrast to acute pancreatitis, gallstones are not considered to be a cause for chronic pancreatitis. The main symptom is recurrent pain in the upper abdomen, which often lasts hours to days and is present in 90% of the patients. In many cases, this leads to further abuse of alcohol and analgesics.

The patients may also have diarrhea and weight loss due to poor feeding and increased losses via diarrheal stools. Only in later stages of the disease can the pain disappear completely. Due to the constant inflammation, a decline in pancreatic tissue occurs, which results in an exocrine and endocrine insufficiency.

This leads to maldigestion and intolerance of certain kinds of food (mainly fats) with nausea, vomiting, meteorism, weight loss, and diabetes mellitus. In earlier stages, these consequences can be reversible. In later stages with a 'burnt-out' gland, however, the pancreatic insufficiency is irreversible.

The determination of elastase-1 in the stool (< 100 μg/dL at insufficiency) can be performed as an indirect measuring of exocrine pancreatic insufficiency. Calcifications in the pancreas can be seen by sonography and are highly suggestive of chronic pancreatitis.

An abdominal computed tomography (CT) scan is the gold standard method of investigation for chronic pancreatitis as calcifications are easily visualized as hyperdense lesions.

Causal therapy includes strict alcohol abstinence or the elimination of the underlying disease. In regards to symptoms, the acute episode should be treated like acute pancreatitis. Therapy of the exocrine insufficiency consists of a diet rich in carbohydrates with many small meals and substitution of pancreatic enzymes.

Due to the endocrine insufficiency, patients become insulin-dependent (CAVE: risk of hypoglycemia). Since there is a risk for addiction to analgesics, they should be enrolled in a Pain Management Center. In most cases, the pain can be reduced by eliminating obstacles to drainage. This can be accomplished by endoscopic retrograde cholangiopancreatography (ERCP) with stent implantation because this decreases the pressure in front of the stenosis. Pain that is resistant to therapy presents an indication for surgery. The prognosis of chronic pancreatitis is poor; lethality amounts to 30–40 % (within 10 years).
Pancreatic pseudocysts

Pancreatic pseudocysts can present in the course of pancreatitis and disappear spontaneously in 50% of the cases. They consist of an accumulation of fluids, necrotic tissue, pancreas enzymes, and old blood. The main symptoms are a pain in the upper abdomen, nausea, vomiting, and weight loss. Further symptoms can occur due to compression of surrounding organs (intestinal tract, spleen, and kidney).

However, pancreatic pseudocysts often are asymptomatic. Complications are a spontaneous rupture in the abdominal cavity and bleeding into the cyst. Sonography is suitable for diagnosis, in which the cysts present as anechoic structures with a hyperechoic rim. If there are cysts with a diameter of more than 5 cm, spontaneous resolution over the ensuing months is less likely and cyst infection, a local complication, may occur. In these cases, surgery with cyst-jejunostomy or partial pancreatic resection may be necessary.

Diabetes mellitus

The physician encounters diabetes very often. Since insulin is the only hormone in the body that can decrease the blood sugar level, insulin deficiency results in an increase in glucose in the blood (hyperglycemia) and increased glucose excretion in the urine (glycosuria). There may be a relative or an absolute insulin deficiency.

With type 1 diabetes, there is an absolute insulin deficiency due to the destruction of the beta cells via autoantibodies. These patients are mostly young (between ages 15–25) and have to be substituted with exogenous insulin all their lives. Typical symptoms are increased thirst (polydipsia), increased urination (polyuria), weight loss, and reduced performance.

Type 2 diabetes is by far more frequent and characterized by resistance to insulin by peripheral cells so that a relative insulin deficiency occurs. It is also referred to as adult-onset diabetes since this disease mainly affects adults. In most patients, adiposis, hypertonia, and hypertriglyceridemia can be diagnosed. Along with diabetes mellitus, this quartet is referred to as the metabolic syndrome. In regards to therapy, a balanced diet, normalization in weight, and movement are important. Oral diabetes drugs are the medication of choice.
Mucoviscidosis

Mucoviscidosis or cystic fibrosis is a frequent metabolic disease that is inherited as an **autosomal recessive** trait. It is a multi-organ disease of exocrine gland function that mainly affects the lungs and the pancreas leading to chronic respiratory tract infections and pancreatic enzyme insufficiency.

It mainly affects the **lungs** and the **pancreas**. A mutation in the cystic fibrosis transmembrane regulator (CFTR)-gene leads to defective chloride channels in the ducts of exocrine glands which makes them produce a very **viscous** secretion that can obstruct the duct system of the gland. Thus, a cystic or fibrous conversion of the affected organ takes place, which gradually leads to **loss of function**.

Besides the lungs and the pancreas, the intestine, liver, bile ducts, and gonads can be affected so that the symptoms heavily depend on the affected organ. For the pancreas, the exocrine insufficiency is most important: the consequences are indigestion with diarrhea and weight loss up to cachexia. **Diabetes mellitus** can possibly result due to the destruction of the islet tissue and, hence, a deficient insulin production. In the respiratory tract, they present with a persistent cough, wheezing, and restlessness.

The **pilocarpine iontophoresis sweat test** (measures the chloride content in sweat; pathological at values > 60 mmol/L) can be performed to ensure the correct diagnosis. Since mucoviscidosis is **not curable**, only symptomatic therapy can be carried out. Life expectancy is mainly limited by respiratory insufficiency and is roughly 40 years.

**Note:** An early diagnosis (if possible in the 1st half of the 1st year of life) is important in order to increase life expectancy.

Pancreatic carcinoma

Pancreatic carcinoma is the **3rd most frequent** tumor of the digestive tract, affecting men more frequently than women. The peak in disease onset is between the ages of 60–80. The causes are unknown, but **smoking, adiposis, and heavy alcohol consumption** are suspected to be risk factors. This **adenocarcinoma** originates from the epithelium of the duct and, in roughly 70% of the cases, it is located in the head of the pancreas.
The absence of early symptoms makes pancreatic cancer hard to diagnose and this has a negative effect on the prognosis as it is recognized very late. Symptoms like upper abdominal pain, nausea, and weight loss make the differential diagnosis of chronic pancreatitis possible. Many patients report pain spreading to the back and jaundice, which points to a progressed stage of the disease.

Another symptom that is rather rare is a recurrent risk for thrombosis. If the body or the tail of the pancreas is affected, symptoms generally occur even later so that the prognosis is worse compared to cancer in the head of the pancreas.

Normally, the tumor is not palpable. However, the gall bladder is palpable if the tumor obstructs the common bile duct. Jaundice also results in this situation (Courvoisier’s sign). Tumor markers CA 19-9 and CEA should not serve for early diagnosis but are helpful in the post-surgery follow-up.

Only a few patients can be cured with the resection of the tumor since metastases are, in most cases, already there when the diagnosis is made.

The results of chemo and radiation therapy are also not satisfactory. The latter is helpful,
however, in the therapy of the pancreatic pain. In most patients, **palliative interventions** (biliodigestive anastomosis, percutaneous stenting, and biliary drainage) are indicated. They can prolong life for 6–9 months.

**Note:** Pancreatic carcinoma = difficult diagnosis, difficult therapy, and poor prognosis!

### Examination of the Pancreas

Due to its location in the retroperitoneal space, the pancreas is neither palpable nor can it be auscultated or imaged using a conventional X-ray. Pain expresses itself in the upper abdomen in a belt-like manner.

**Note:** The pancreas is not accessible for a simple physical examination! Often, a pancreatic disease is noticed when symptoms occur. With a carcinoma of the pancreatic head, jaundice can result due to an obstruction of the common bile duct (see above).

| **Sonography** | The main structure for the localization of the pancreas is the splenic vein, which lies dorsally of it. The echogenicity is similar to the liver's; echogenicity increases with adiposis and increasing age (=pancreatic lipomatosis). Pancreatic tumors are normally hypoechogenic. Calcifications are evidence of chronic pancreatitis. For better imaging of the tail of the pancreas, the ultrasound head should be rotated slightly clockwise. |
| **CT/MRT (magnetic resonance tomography)** | In the sectional views, the pancreas lies in front of the vertebras in a hoof-shape. |
| **ERCP** | Imaging of the bile ducts and pancreatic ducts with an application of contrast agents via the major duodenal papilla and screening afterward. If needed, a direct therapeutic intervention (papillotomy, stone extraction, or drainage placement) is possible. |
| **MRCP (magnetic resonance cholangiopancreatography)** | Non-invasive and does not involve radiation exposure. Imaging of fluids in the pancreatic ducts and the bile ducts as areas with high signal intensity, but no direct therapeutic intervention possible. |

### Patients frequently ask

**Can you live without a pancreas?**

Medically, it is possible to survive after the complete removal of the pancreas. However, in the absence of the pancreas, body is incapable to produce digestive enzymes, insulin, and other hormones to digest food and control blood sugar levels. Thus, the patient develops diabetes and he has to completely rely on exogenous insulin supply and other medicines to maintain sugar levels.

### References


Hoboken: John Wiley & Sons.


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