

# Acute and Chronic pancreatitis, Splenectomy

MO teaching slides  
2<sup>nd</sup> August 2016

# Acute Pancreatitis

- Diagnosis
- At least 2 out of 3 criteria
  - *Abdominal pain ( cardinal symptom)– constant, epigastric, radiating to the back*
  - *Raised serum amylase/lipase levels 3x ULN*
  - *Radiological evidence of pancreatitis on CT scan with contrast / MRI pancreas*
- Etiology
  - *“ I GET SMASHED” , Idiopathic – most common*
  - *Hx of biliary colic, alcohol intake/dependence, trauma, drugs- steroids, ERCP, high TG, Ca, PTH*

## To note:

- *If gallstone pancreatitis, pain can be in RUQ. Others- poorly localised*
- *Painless pancreatitis presenting with hypotension e.g critically ill, post-operatively*
- *90% have nausea and vomiting that can last for several hours*
- *Imaging is discouraged esp within initial 48hrs, unless there is clinical doubt of the diagnosis.*
- *US abdomen –often done TRO gallstones*
- *Serum amylase not as sensitive and specific as lipase. Rises early and normalises by 3-5days*
- *MRI pancreas more sensitive than CT scan to characterise solid vs fluid*

# Examination

- Vitals: Fever / Tachycardia/ Hypotension
- Dyspnoeic, respiratory distress, jaundice
- Abdominal pain, guarding
- Abdominal distension, diminished bowel sounds from ileus
- Cullen and Grey Turner sign - severe necrotizing pancreatitis

# Investigations

- FBC, LFTs, RP, Calcium and Albumin, LDH, Fasting lipids, ABG
- ANA, IgG4 if suspect autoimmune pancreatitis
- Imaging- US abdo, EUS (if cause is unclear- pancreatic ductal abn, small tumors at/near ampulla, microlithiasis in GB/bile duct, early chronic pancreatitis)
- CT/MRCP – if suspect malignancy from hx ( >40y, LOW, new onset DM)
- CXR

# Predictors of severity

**Scoring systems**- none proven to be accurate in predicting severity. However, they are superior judgement for triaging pts to ICU and aggressive therapy

## ■Ranson's criteria – done at 0h and 48h

- Score 3- mortality 0-3%, score >3 11-15% mortality, score > 6 – 40% mortality

## ■APACHE II score – 12 measures. APACHE O ( inc BMI improves predictability of severe AP)

- Score < 8 – mortality < 4%, Score > 8 mortality 11-18%

## ■Glasgow score – less sensitive than Ranson's (73% vs 91%), similar specificity (71% vs 74%)

## ■CT severity Index ( Balthazar)

- Based upon the degree of necrosis, inflammation, and the presence of fluid collections
- Detection of pancreatic necrosis does not necessarily predict organ failure but it does alter treatment approach
- Internal validation study showed any degree of pancreatic necrosis was a/w mortality of 23%. Strong association with morbidity/mortality if > 30% necrosis.
- Score > 6 = severe disease

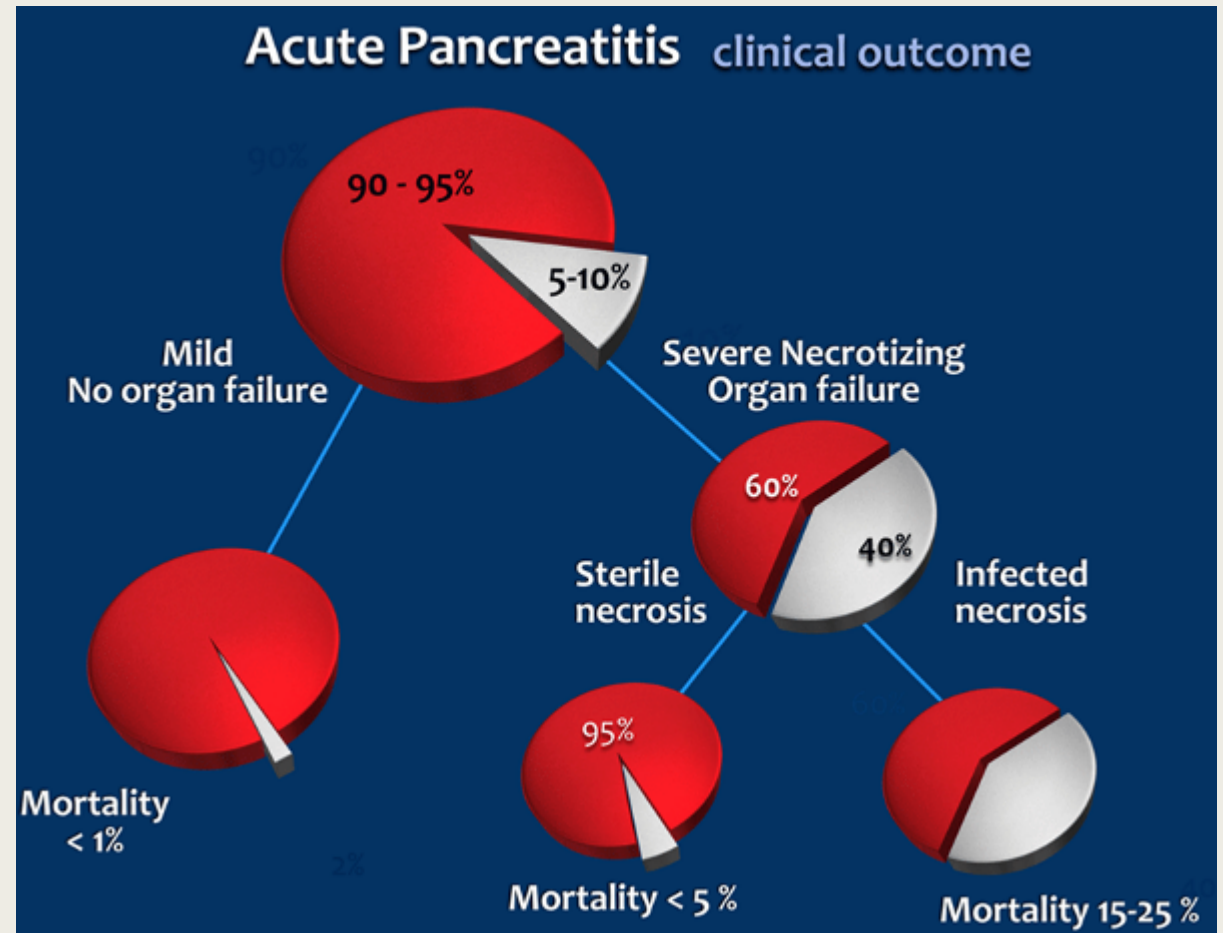
CT Severity Index		
<b>Pancreatic inflammation</b>		<b>points</b>
Normal pancreas		0
Enlargement of the pancreas		1
Peripancreatic inflammation		2
1 acute peripancreatic fluid collection		3
≥ 2 acute peripancreatic fluid collections		4
<b>Pancreatic necrosis</b>	None	0
	< 30%	2
	30% - 50%	4
	> 50%	6
		<b>Maximum 10 points</b>

# Predicting severity – American College of Gastroenterology guidelines

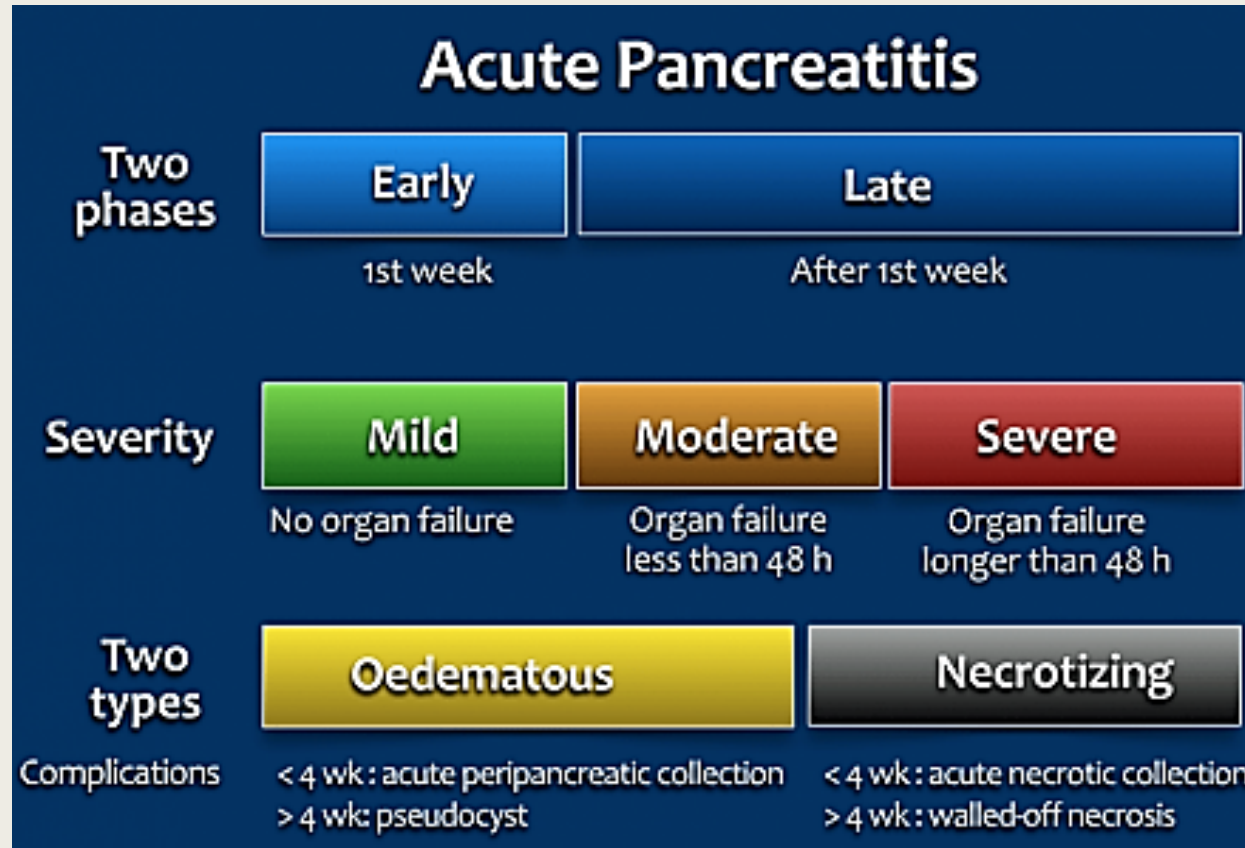
- Patient characteristics
    - Age >55 years
    - Obesity (body mass index >30 kg/m<sup>2</sup>)
    - Altered mental status
    - Comorbid disease
  - SIRS syndrome (\* early / persistent >72h)
    - Presence of >2 of the following criteria:
    - Pulse >90 beats/min
    - Respirations >20/min or PaCO<sub>2</sub> >32 mm Hg
    - Temperature >38 or <36 °C
    - White blood cell count >12,000 or <4,000 cells/mm<sup>3</sup> or >10 percent immature neutrophils (bands)
  - Laboratory findings
    - Blood urea nitrogen (BUN) >20 mg/dl
    - Rising BUN
    - Haematocrit (HCT) >44 %
    - Rising HCT
    - Elevated creatinine
  - Radiology findings
    - Pleural effusions
    - Pulmonary infiltrates
    - Multiple or extensive extrapancreatic collections
- \* CRP > 150 at 48hrs is predictive of severe pancreatitis

# Clinical outcomes

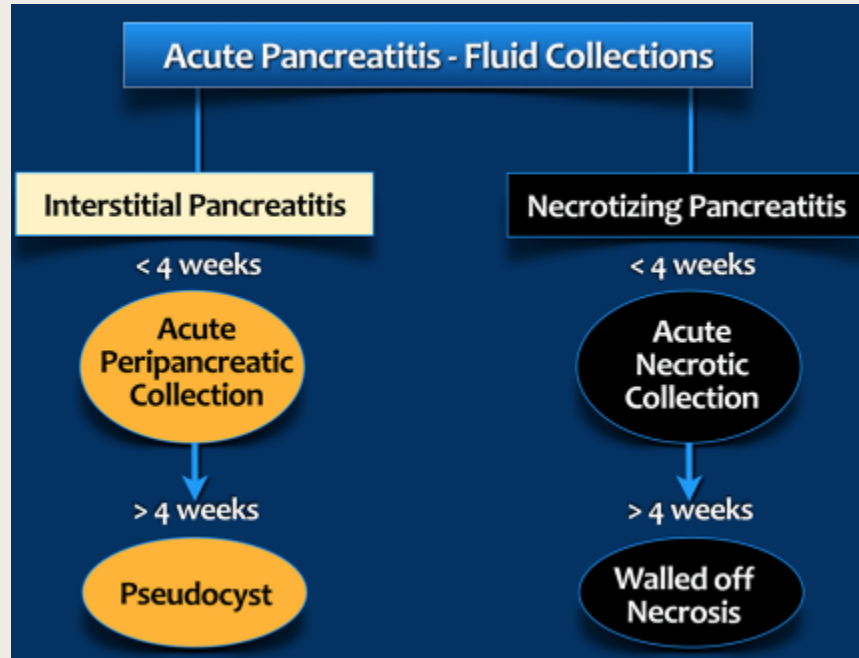
- 85% of patients with acute pancreatitis have mild disease
- 15% have necrotising pancreatitis with necrosis of pancreatic parenchyma or peripancreatic tissue
- Most patients have mild disease and recover over 3-5 days
- 20% will develop moderately severe to severe pancreatitis with either local or systemic complications or organ failure
- Overall mortality is 5%. With interstitial pancreatitis (3%) having a lower mortality than necrotising pancreatitis (17%)



# Revised Atlanta Classification (2012)



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Acute Peripancreatic Collection	Acute Necrotic Collection
<ul style="list-style-type: none"> <li>• &lt; 4 weeks</li> <li>• In interstitial pancreatitis</li> <li>• Homogeneous - fluid density</li> <li>• No <i>fully definable wall</i></li> <li>• Adjacent to pancreas</li> <li>• Confined by normal fascial planes</li> </ul>	<ul style="list-style-type: none"> <li>• &lt; 4 weeks</li> <li>• In necrotizing pancreatitis</li> <li>• Heterogeneous collection</li> <li>• No <i>fully definable wall</i></li> <li>• Intra- or extrapancreatic</li> </ul>
Pseudocyst	Walled-off Necrosis
<ul style="list-style-type: none"> <li>• &gt; 4 weeks</li> <li>• In interstitial pancreatitis</li> <li>• Homogeneous - fluid density</li> <li>• <i>Well defined wall</i></li> <li>• Adjacent to pancreas</li> <li>• No non-liquid component</li> </ul>	<ul style="list-style-type: none"> <li>• &gt; 4 weeks</li> <li>• In necrotizing pancreatitis</li> <li>• Heterogeneous collection</li> <li>• <i>Well-defined wall</i></li> <li>• Intra- or extrapancreatic</li> </ul>



# Management

- Supportive treatment
- Fluid resuscitation
  - *Critical for aggressive fluid replacement in initial 48hrs*
  - *Adjust according to Clinical assessment, BUN, Creatinine*
  - *Lactated Ringer's solution reduces SIRS / CRP cpd to Normal saline*
- Pain control
  - *Opioids*
  - *Fluid resuscitation ( ischaemia)*
- Active monitoring – vitals, urine output, electrolytes, hypocount – if raised, increases risk of infection
- Nutrition
  - *Mild pancreatitis – IV hydration, resume oral diet ( low fat) within 1 week if pain improving, no ileus, N/V*
  - *Moderate-severe – initiate within 48hrs*
    - *Enteral feeding preferred to parenteral :maintain the intestinal barrier and prevents bacterial translocation from the gut*
    - *High protein, low fat, semi-elemental feeding formulas (eg, Peptamen AF) because of a reduction in pancreatic digestive enzymes*
- Prophylactic antibiotics are not recommended regardless of type or disease severity
- If clinical deterioration noted after 72hrs – CT with IV contrast indicated to assess for local complications and pancreatic necrosis

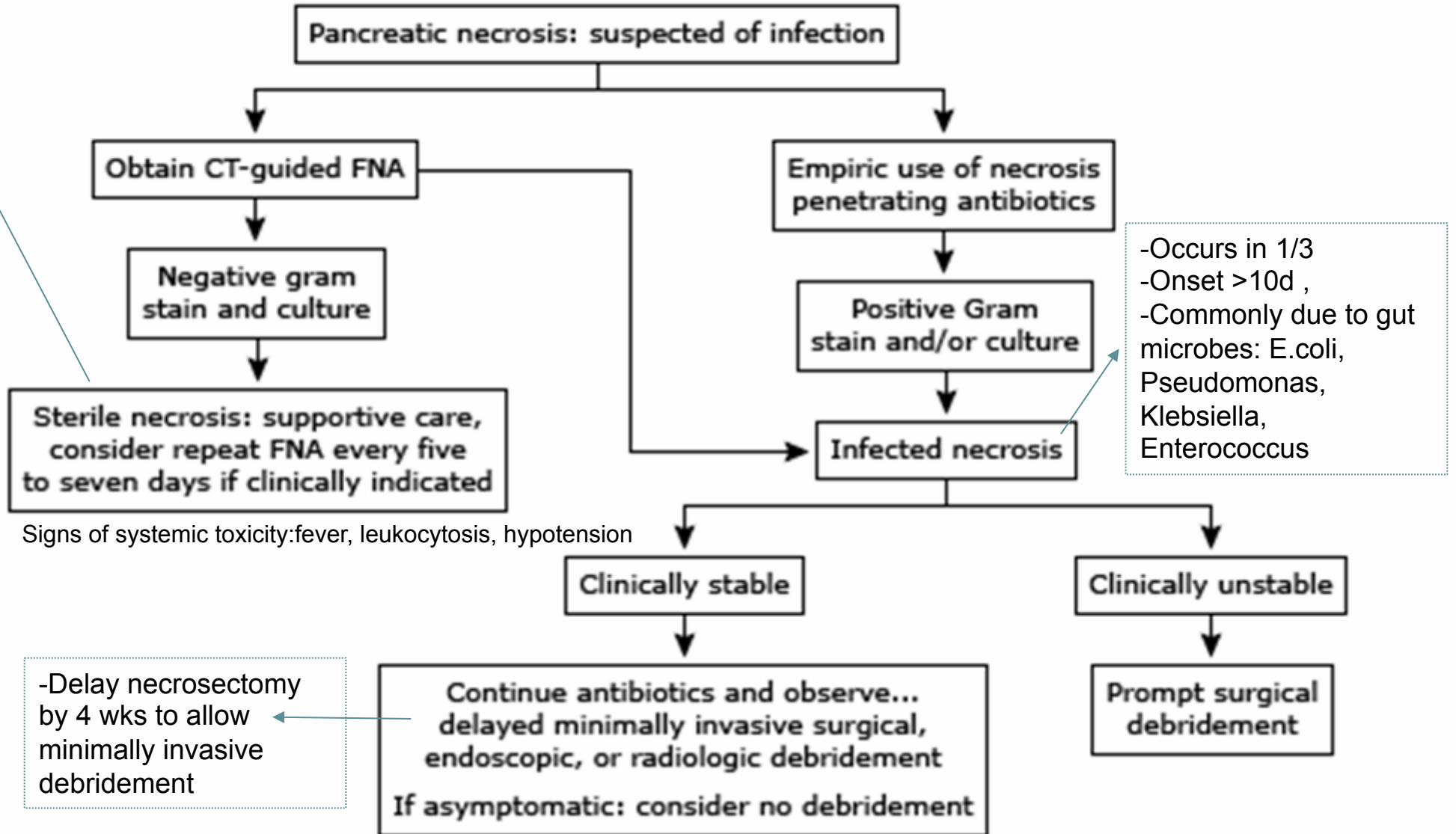
# Management of complications

- **Acute pancreatic fluid collection** – early phase. Usually asymptomatic, resolves spontaneously w/i 7-10days without need for drainage
- **Walled-off pancreatic necrosis/ Pancreatic pseudocyst** ( after 4wks)
  - Mnx depends on sx, characteristics and location of fluid collection
  - Expectant management for those w/o cystic neoplasm, pseudoaneurysm or minimal symptoms
    - Repeat CT scan every 3-6mths until cyst resolves/stabilizes at a small size
    - If pseudoaneurysm present but asymptomatic – embolization of aneurysm followed by expectant management
  - IF drainage required, 3 main options:
    - **Endoscopic drainage**
      - *Contraindicated if pseudoaneurysm present*
      - *Good for relatively small pseudocysts in communication with the main pancreatic duct for transpapillary stent placement*
      - For larger symptomatic cysts abutting the stomach, duodenum, transmural puncture to relieve symptoms possible
      - *10 to 15% morbidity, 70 to 80% fluid collection resolution, 10-15% recurrence rate.*
    - **Surgical drainage**
      - Can be open or lap. Mainly used in cases of endoscopic failures, recurrence of collection
      - Cystogastrostomy or a cystojejunostomy depending on location
      - Open approach a/w substantial morbidity and mortality ( 25%, 5% respectively)
    - **Percutaneous drainage**
      - As effective as surgery in draining and resolving sterile and infected walled-off pancreatic fluid collections
      - Stent diameter may need to be increased with time for necrotic debris to drain. Frequent irrigation required to maintain patency
      - Risk of infection, pancreaticocutaneous fistula formation

# Management of pancreatic necrosis

Indications for intervention in absence of signs of infection

- Ongoing GOO, biliary obstruction due to mass effect 4-8 wks after onset
- Persistent symptoms of abdo pain, N/V/anorexia/LOW >8 wks after onset
- Disconnected duct syndrome with persistent sx of pain and obstruction >8 wks after onset



# Management of Gallstone pancreatitis

- Gallstone pancreatitis

- *ERCP*

- ERCP shld be performed early ( w/i 24hrs) for patients with gallstone pancreatitis and cholangitis
    - ERCP also indicated if imaging shows CBD obstruction with visible stone, dilated CBD or increasing LFTs in absence of cholangitis.
    - Evidence for URGENT ERCP is controversial- *early ERCP reduced pancreatitis-related complications but not mortality in patients predicted to have severe pancreatitis*
    - ERCP NOT indicated even in severe gallstone pancreatitis WITHOUT cholangitis

- *Cholecystectomy*

- Can be done within 7 days of index hospitalization in mild pancreatitis
    - In severe necrotizing pancreatitis, cholecystectomy shld be delayed
    - Risk of recurrence acute pancreatitis \
    - Cholecystectomy shld also be offered to patients with acute pancreatitis and biliary sludge

# Chronic Pancreatitis

- **Definition:**
  - *Syndrome involving progressive inflammatory changes in the pancreas that result in permanent structural damage which can lead to impairment of exocrine or endocrine function*
- **Distinguishing features:**
  - *Asymptomatic over long period of time*
  - *Present with a fibrotic mass with symptoms of pancreatic insufficiency without pain.*
  - *Normal levels of amylase and lipase*
- **Morphologically:-**
  - *Patchy focal disease characterised by mononuclear infiltrate and fibrosis vs diffusely large portion of pancreas involved with a neutrophilic inflammatory response in acute pancreatitis*
- **Pancreatic insufficiency ( >90% of pancreatic function is lost)**
  - *Fat malabsorption – steatorrhoea prior to protein deficiency. Malabsorption of fat soluble vitamins*
  - *Glucose intolerance occurs in chronic pancreatitis, overt diabetes mellitus occurs late in disease*
- **Classic triad:** steatorrhoea, diabetes mellitus, pancreatic calcifications – late advance stage disease
- Can develop acute pancreatitis with sudden worsening or change in pattern of symptoms
- Watch for possibility of a pancreatic carcinoma. Increased risk in pts with chronic pancreatitis
- **Investigations**
  - *72hr faecal fat test ( >7g/day fat diagnostic of malabsorption)*
  - *Plain XR – showing calcium deposition – common in alcoholic pancreatitis*
  - *US low sensitivity (70%), specificity ( 90%) vs CT 80% spec 85%*
  - *MRCP becoming diagnostic test of choice in view of demonstrating calcifications and pancreatic duct obstruction consistent with chronic pancreatitis*
  - *ERCP –now limited for therapeutic. Characteristic beading of main pancreatic duct and ectatic side branches is diagnostic of chronic pancreatitis*

# Chronic Pancreatitis

- After establishing a secure diagnosis of pancreatitis ( i.e ruling out other causes of symptoms e.g PUD, biliary obstruction, pseudocysts, pancreatic carcinoma )
- Management
  - *Treat underlying cause e.g cessation of alcohol intake*
  - *Small portion, low fat meals ( anecdotal evidence)*
  - *Smoking cessation to reduce risk of Pancreatic CA*
  - *Pain management – some may need chronic Opioids*
  - *Pancreatic enzyme ( Creons, Vitamin ABDEK)*
- Surgical options:
  - *A meta-analysis of four trials concluded that **duodenum-preserving pancreatic head resection** was as effective as pancreatoduodenectomy for relief of pain, overall morbidity and postoperative endocrine insufficiency and was superior in some postoperative outcomes and quality of life*
  - *Total pancreatectomy is a treatment option in **carefully selected** patients with chronic pancreatitis in whom substance abuse is not a confounding factor*

# Complications of acute pancreatitis

- Splanchnic venous thrombosis
  - *Involves splenic, portal, SMV*
  - *Effective treatment of underlying pancreatitis usually leads to spontaneous resolution*
  - *Anti-coagulation shld be initiated if there is extension of clot into portal or SMV causing hepatic decompensation or reduced bowel perfusion*
- Pseudo aneurysms
  - *Rare but serious complication of acute pancreatitis*
  - *High index of suspicion for those with acute pancreatitis and unexplained GI bleeding*
- Abdominal compartment syndrome
  - *Increased risk due to aggressive fluid resuscitation, peripancreatic inflammation, ascites and ileus*

# Splenectomy – Indications

- Traumatic Splenic injury
- Non-traumatic / Medical
  - *Indications for splenectomy:*
    - ITP ( most common)
    - Hereditary spherocytosis
    - Thalassaemia
    - Sickle Cell anaemia
    - Hodgkin disease
    - Felty syndrome ( RA, splenomegaly, neutropaenia)



# Splenic injury

- Often occurs in blunt trauma due to MVA, falls
- Patient presents with trauma and is assessed with ATLS protocols
- FAST scan – hypoechoic rim of subcapsular fluid or intraperitoneal fluid around the spleen or in Morrison's pouch (hepatorenal space)
- CT scan with IV contrast – arterial phase will show a blush / extravasation, hypodensity can represent hematoma, hemoperitoneum by comparing Hounsfield units to differentiate from ascites

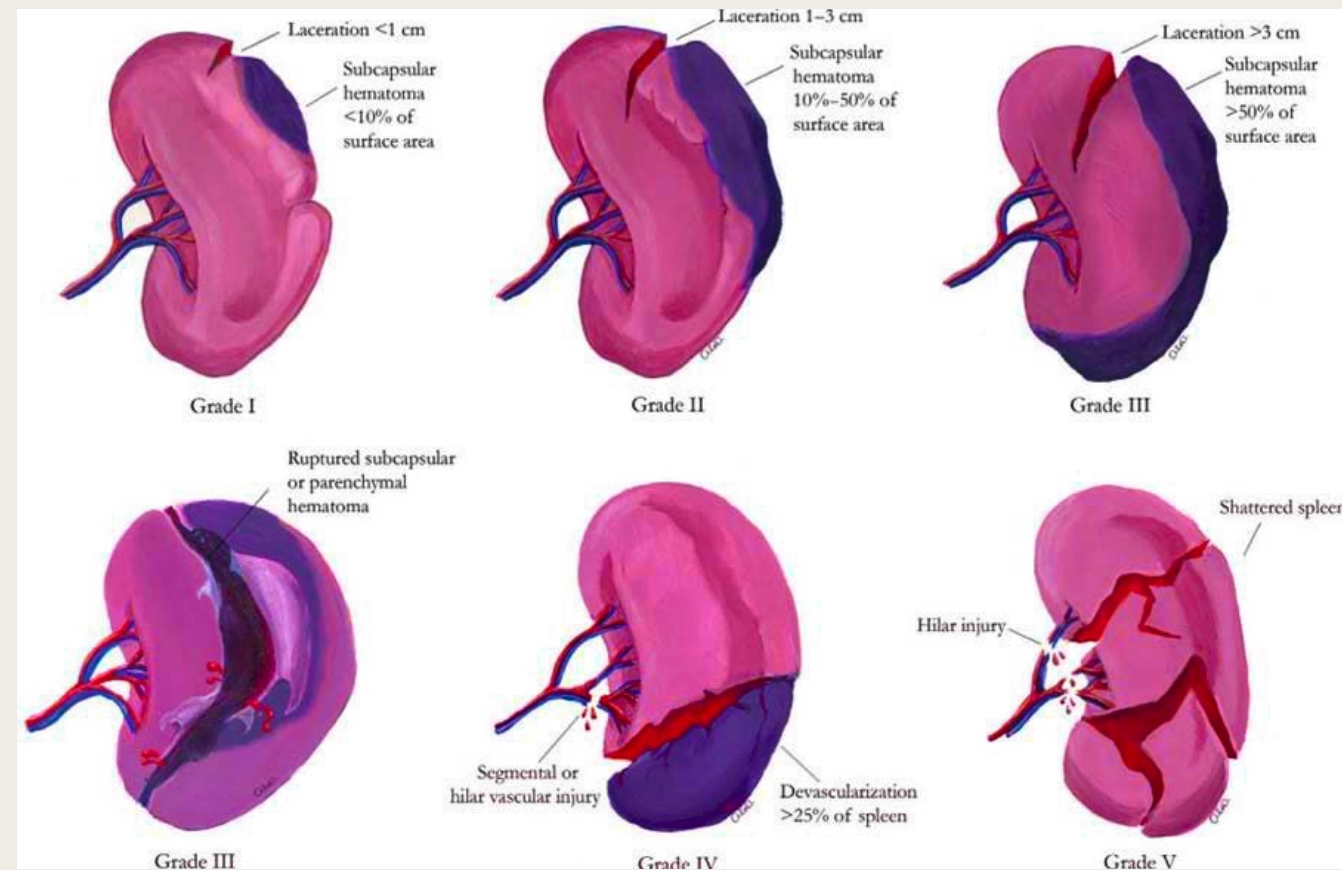
# Grading of splenic injury – AAST

## Splenic CT Injury Grading Scale

Grade I	Laceration(s) < 1 cm deep Subcapsular hematoma < 1cm diameter
Grade II	Laceration(s) 1-3 cm deep Subcapsular or central hematoma 1-3cm diam
Grade III	Laceration(s) 3-10 cm deep Subcapsular or central hematoma 3-10 cm diam
Grade IV	Laceration(s) > 10 cm deep Subcapsular or central hematoma > 10cm diam
Grade V	Splenic tissue maceration or devascularization

### The shortcomings of this grading scale are:

- Underestimation of injury extent
- Significant inter-observer variability.
- Does not include:
  - Active bleeding
  - Contusion
  - Post-traumatic infarcts
- Most importantly: **no predictive value** for non-operative management (NOM).



# Management of traumatic splenic injuries

- **Non-operative management**
  - *Hemodynamically stable*
  - *Low grade splenic injury ( I-III)*
  - *Without evidence of intra-abdominal injury*
- No evidence of active contrast extravasation/ blush on CT
  - *Even those who require extra-abdominal injuries e.g fracture stabilization*
  - *Contraindications: hemodynamic instability, generalized peritonitis, other intra-abd injuries requiring surgical exploration. Portal hypertension –relative contraindication*
- **Angio-embolisation**
- Indications: Active contrast extravasation on CT, intra-parenchymal pseudoaneurysm formation
- Variable success rates – institution dependent
- Relative contraindication: Grade IV/V injuries due to vascular disruption, age>55yrs due to splenic capsule thinning out and associated with higher failure rates
- If contrast extravasation is from splenic parenchyma supplied by short gastric vessels – then operative intervention prompted as they are less amenable to embolization
  
- **Surgical intervention**
- Indication: hemodynamically unstable, unable to tolerate significant hypotension, failure of non-surgical management
- Splenic salvage:
  - *Splenorrhaphy refers to the suture repair of the spleen with or without splenic wrapping, and is generally supplemented by electrocautery techniques for control of parenchymal haemorrhage*
  - *Partial splenectomy is a form of splenic salvage and refers to the removal of a portion of the spleen based upon its segmental blood supply.*
- Total splenectomy is the safest option and is often done in emergency setting where the risk of blood loss outweighs the risk of OPSI
- Replantation of splenic tissue – If splenectomy for injury is deemed necessary, heterotopic autotransplantation of splenic tissue into omental pockets may provide some splenic function, although this has not been proven conclusively

# Post-splenectomy related issues

## ■ Activities

- *Restricted activities and avoidance of high risk activities for up to 3 months*
- *No clinical studies to support duration*

## ■ Vaccinations

- *Shld be given 14 days before elective surgery*
- *Given on POD 14 for emergency surgeries*
- *Inactivated Influenza vaccines– NOT live attenuated vaccines*

## ■ Antibiotics

- *In Children, daily penicillin ( amoxicillin) up to 5yrs and 1 yr after splenectomy*
- *In Adults – no indication for daily antibiotics unless in hypogammaglobulinemia, HIV infection, solid organ transplant recipients, and patients with advanced liver disease*
- *However, if febrile, early administration of antibiotics reduces risk of severe infection*

**Vaccination of persons with asplenia or sickle cell disease**

Vaccine	Asplenia or sickle cell disease	
	Recommendation	Strength, evidence quality
<i>Haemophilus influenzae b conjugate</i>	U: age <5 years	Strong, moderate
	R: age ≥5 years	Weak, low
Hepatitis A	U	Strong, moderate
Hepatitis B	U	Strong, moderate
Diphtheria toxoid, tetanus toxoid, acellular pertussis; tetanus toxoid, reduced diphtheria toxoid; tetanus toxoid, reduced diphtheria toxoid, and reduced acellular pertussis	U	Strong, moderate
Human papillomavirus	U	Strong, moderate
Influenza-inactivated (inactivated influenza vaccine)	U	Strong, moderate
Influenza-live attenuated (live attenuated influenza vaccine)	X	Weak, very low
Measles, mumps, and rubella-live	U	Strong, moderate
Measles, mumps, and rubella-varicella-live	U	Strong, moderate
Meningococcal conjugate*	R <sup>¶</sup>	Strong, low
Pneumococcal conjugate (PCV13)	U: age <6 years <sup>‡</sup>	Strong, moderate
	R: age ≥6 years <sup>°</sup>	Strong, very low
Pneumococcal polysaccharide (PPSV23)	R: age ≥2 years <sup>§</sup>	Strong, low
Polio-inactivated (inactivated poliovirus vaccine)	U	Strong, moderate
Rotavirus-live	U	Strong, moderate
Varicella-live	U	Strong, moderate
Zoster-live	U	Strong, moderate

R: recommended—administer if not previously administered or not current; such patients may be at increased risk for this vaccine-preventable infection; U: usual—administer if patient not current with recommendations for dose(s) of vaccine for immunocompetent persons in risk and age categories; X: contraindicated.

Thank you!

### Ranson criteria to predict severity of acute pancreatitis

0 hours	
Age	>55
White blood cell count	>16,000/mm <sup>3</sup>
Blood glucose	>200 mg/dL (11.1 mmol/L)
Lactate dehydrogenase	>350 U/L
Aspartate aminotransferase (AST)	>250 U/L
48 hours	
Hematocrit	Fall by $\geq 10$ percent
Blood urea nitrogen	Increase by $\geq 5$ mg/dL (1.8 mmol/L) despite fluids
Serum calcium	<8 mg/dL (2 mmol/L)
pO <sub>2</sub>	<60 mmHg
Base deficit	>4 MEq/L
Fluid sequestration	>6000 mL

The presence of 1 to 3 criteria represents mild pancreatitis; the mortality rate rises significantly with four or more criteria.

**A score  $\geq 3$  indicates Acute Severe Pancreatitis**  
**A score = 2 indicates Acute Moderate Pancreatitis**  
**A score < 2 indicates Acute Mild Pancreatitis**

#### Assessing the severity of acute pancreatitis

Glasgow prognostic score: (NOTE PANCREAS ACRONYM)

- PaO<sub>2</sub> < 8kPa (60mmhg)
- Age > 55 years
- Neutrophils: (WBC >15 x10<sup>9</sup>/l)
- Calcium < 2mmol/l
- Renal function: (Urea > 16mmol/l)
- Enzymes: (AST/ALT > 200 iu/L or LDH > 600 iu/L)
- Albumin < 32g/l
- Sugar: (Glucose >10mmol/L)

**Any 3 factors means acute severe pancreatitis**