

## Achalasia

- Idiopathic loss of the normal neural structure of the lower esophageal sphincter resulting in inability to relax

### Features

- Progressive dysphagia to both solid and fluids (more to fluids)
- Regurgitation (hours after eating)
- Weight loss
- There may be history of **recurrent URTIs** or **aspiration pneumonia** as a result from untreated achalasia that leads to **nocturnal inhalation** of material lodged in the esophagus
- NO relationship with alcohol or tobacco use

### Investigations

- Barium swallow → (Sigmoid esophagus); dilation of the esophagus, which narrows into a “parrot’s beak” at the distal end
- Manometric studies (*most accurate*) → increased tone of lower esophagus

### Management

Dilation of the lower esophageal sphincter

- HELLER'S OPERATION (*myotomy*)
- Botulinum toxin injection, elderly not able to tolerate operations

#### Esophageal cancer

- dysphagia to solid first then liquids
- common in **elderly** with long history of alcohol and tobacco use

#### Schatzki ring

- Narrowing of the lower esophagus that leads to intermittent dysphagia but not associated with pain, mainly to solids and in old patients

## Esophageal spasm (CCCC)

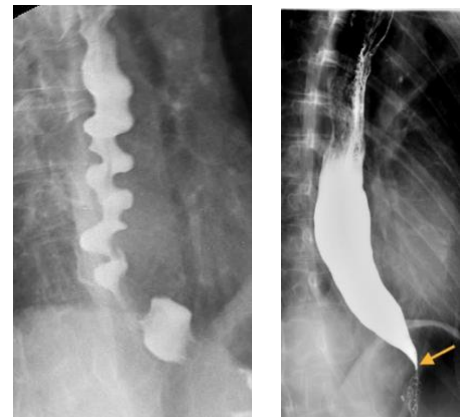
- Intermittent Chest pain and dysphagia, pain can be precipitated by Cold liquids
- Pain can simulate that of MI, but it has no relation to exertion
- Relieved after ingestion of nitrates as they are smooth muscle relaxants

### Investigations

- Manometric study → High intensity and disorganized contractions (*most accurate*)
- Barium meal → Corkscrew pattern

### Treatment

- Ca channel blockers: nifedipine



## Plummer Vinson \$

- Middle-aged woman
- Dysphagia (painless/intermittent) + IDA + post cricoid esophageal web

### Management

- Iron supplement
- Dilation of the web

## Barret's esophagus

- Occasional dysphagia
- Results from long history of **GERD**
- Replacement of Sq. epithelium to columnar epithelium
- Associated with esophageal adenocarcinoma



## Dysphagia

<b>Benign esophageal stricture (peptic stricture)</b>	<ul style="list-style-type: none"> <li>• Dysphagia to <u>both</u> solids and liquids <u>WITHOUT</u> regurgitation</li> <li>• Results from <u>scarring</u> due to: <ul style="list-style-type: none"> <li>- Acid reflux</li> <li>- Persistent <u>GERD</u> (retrosteral discomfort)</li> <li>- Ingestion of corrosives</li> <li>- Drugs: <b>Bisphosphonates</b> (alendronate) – <b>NSAIDs</b>. So patients are advised to lie down for 30min after administration</li> </ul> </li> </ul>
<b>Esophageal carcinoma</b>	<ul style="list-style-type: none"> <li>• Symptoms of cancer</li> </ul>
<b>Barrett's esophagus</b>	<ul style="list-style-type: none"> <li>• A long hx of <u>GERD</u>, <u>occasional</u> dysphagia not persistent</li> </ul>
<b>Pharyngeal pouch (Zenker's diverticulum)</b>	<ul style="list-style-type: none"> <li>• Hx of <u>halitosis</u>, regurgitation of <u>stale</u> food &amp; a throat <u>lump</u></li> <li>• <u>Barium swallow</u> may show a residual pool of contrast within the pouch</li> <li>• Endoscopy should be <u>AVOIDED</u> in fear of perforation</li> </ul>
<b>Achalasia</b>	<ul style="list-style-type: none"> <li>• Dysphagia + <u>regurgitation</u></li> </ul>
<b>Ulcers and esophageal candidiasis</b>	<ul style="list-style-type: none"> <li>• <b>Painful</b> dysphagia</li> </ul>
<b>Plummer Vinson \$</b>	<ul style="list-style-type: none"> <li>• <u>IDA</u> + <u>Esophageal web</u></li> </ul>

- *Bisphosphonates are used to treat osteoporosis but long-term use can cause esophagitis resulting in a stricture*

## Esophageal cancer

- Adenocarcinoma is more common
- Commonly occur in the lower third, Squamous cell type is less common and it affects the upper 2/3
- More likely to develop in patients with history of GERD or Barret's
- Dysphagia to solid first then liquids + weight loss

### Risk factors

- Smoking (RF for both but mainly for SCC)
- Alcohol
- GERD
- Barret's
- Achalasia
- Plummer Vinson \$

### Diagnosis

- **Upper GI endoscope and biopsy** → *1<sup>st</sup> line*
- **Barium swallow**
  - Rat-tail appearance
  - Apple-core appearance
  - Shouldering

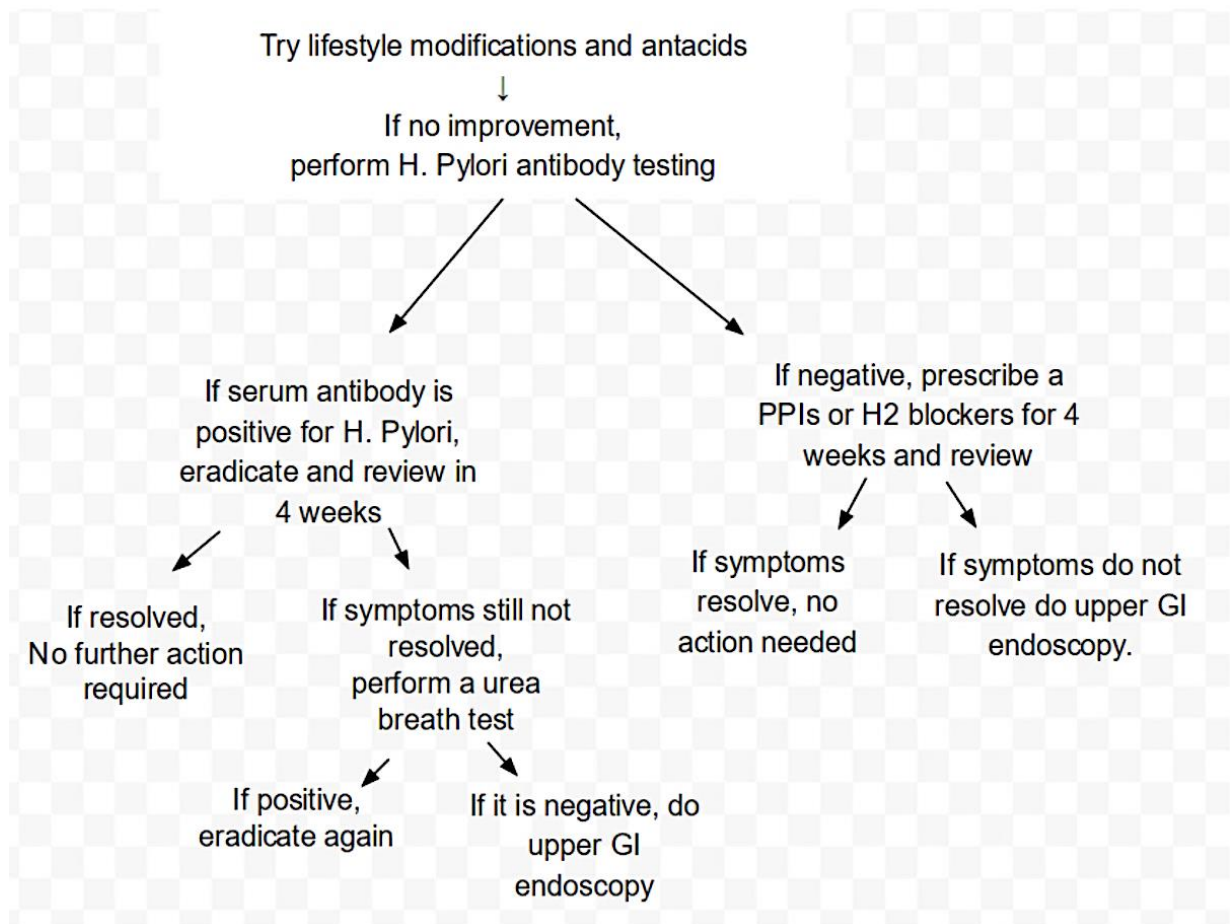


### Treatment

- Operable → surgical resection, radiotherapy
- Inoperable (e.g. metastasis) → palliative esophageal **Stent**, Percutaneous endoscopic gastrostomy (**PEG**) usually in stroke patients who are at risk of aspiration pneumonia or to decompress the stomach in cases of gastric volvulus

## Dyspepsia and H. Pylori

- < 55 with no red flags:



- > 55 with unexplained/persistent symptoms (not just alarm signs) → **Endoscopy**

### ➤ Red flags (ALARMS)

- Anemia
- Loss of weight
- Anorexia
- Recent onset of progressive symptoms
- Masses, Melena or hematemesis
- Swallowing difficulty

### ➤ H. Pylori antibody testing

- Carbon-13 urea breath test
- Stool antigen test
- Serum antibody testing

### ➤ H. Pylori eradication

- PPIs
- Clarithromycin
- Amoxicillin or metronidazole

- Serological tests have no value in confirming successful eradication because antibodies persist long after successful eradication, **C-13 urea breath test** ensures successful eradication
- If the patient is taking **PPIs**, stop it 14 days prior to performing urea breath test or stool antigen test
- There must be a 28-day break after eradication with antibiotics prior to testing

## Esophageal varices

- Dilated sub-mucosal veins in the lower 1/3 of the esophagus
- Often severe and life threatening
- History of **chronic liver disease** → **portal hypertension** → **esophageal varices**

### Features

- **Hematemesis** (*most commonly*) and **melena**
- Signs of **chronic liver disease**

### Investigations

- **Endoscopy** at early stage

### Acute management of variceal bleeding

- Always with **ABC**
- Correct clotting: **FFP, vitamin K**
- **Terlipressin** (vasoactive agent) → should be offered to patients with suspected variceal bleeding at presentation
- **Antibiotic prophylaxis** → reduces mortality in patients with acute upper GI bleeding in association with chronic liver disease
- Endoscopic variceal **band ligation** → if it's not available → emergency sclerotherapy
- Sengstaken-Blakemore tube if uncontrolled hemorrhage
- Transjugular Intrahepatic Portosystemic Shunt (**TIPSS**)



### Prophylaxis of variceal hemorrhage

- **Propranolol**, given at discharge to reduce portal pressure in order to decrease the risk of repeat bleeding

## Perforated peptic ulcer

- Sever sudden localized **epigastric** pain
- May worsen with **coughing** or **moving**
- May radiate to the **shoulder tip**

### Examination

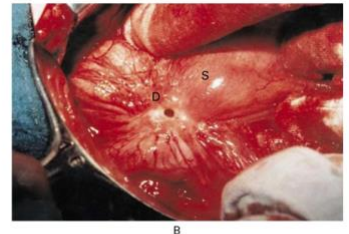
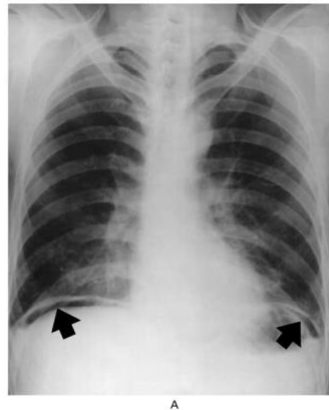
- **Absent bowel sounds**
- **Shock**
- Generalized **peritonitis** (fever, tachycardia, abdominal guarding & rigidity)

### Investigations

- **Erect X-ray** → free gas under the diaphragm
- CT scan

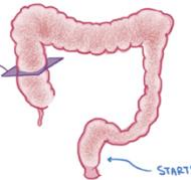
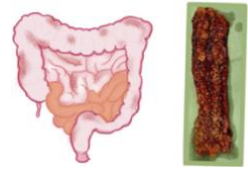
### Management

- IV analgesics
- Antiemetic (e.g. metoclopramide 10mg)
- Resuscitate with IV 0.9% saline
- IV antibiotics



For bleeding peptic ulcer without perforation → **Endoscopy**, or **IV PPIs** if endoscopy isn't available

## IBD

	Ulcerative Colitis → Lt	Crohn's disease → Rt
<b>Def.</b>	<ul style="list-style-type: none"> <li>- BD that forms ulcers in the <u>colon and rectum only</u></li> <li>- Only affects mucosa and submucosa</li> <li>- Autoimmune</li> <li>- <u>Bloody diarrhea</u> is more prominent</li> </ul>	<ul style="list-style-type: none"> <li>- Anywhere in the GIT mostly <u>ileum &amp; colon</u></li> <li>- Extending to serosa</li> <li>- Immune-related caused by pathogens</li> <li>- <u>Weight loss</u> more prominent</li> <li>- <u>Steatorrhea</u></li> </ul>
<b>Diff.</b>	<ul style="list-style-type: none"> <li>- <u>Circumferential</u></li> <li>- <u>Continuous</u></li> <li>- <u>Crypt abscesses</u></li> <li>- 1ry sclerosing <u>Cholangitis</u></li> <li>➤ Aphthous oral ulcers</li> </ul> 	<ul style="list-style-type: none"> <li>- <u>Transmural/deep</u> ulcers</li> <li>- <u>Skip lesions</u> (cobblestone appearance) on endoscopy</li> <li>➤ Peri-anal <u>fistulas</u></li> <li>➤ Kantor's string sign</li> <li>➤ Rose thorn ulcers</li> </ul> 
<b>Symptoms</b>	<ul style="list-style-type: none"> <li>- Pain in <b>LLQ</b> (rectum)</li> <li>- LI: <b>Bloody diarrhea</b> more common</li> </ul>	<ul style="list-style-type: none"> <li>- Pain in <b>RLQ</b> (ileum)</li> <li>- LI: Diarrhea usually not bloody</li> <li>- SI: Malabsorption</li> </ul>
<b>Diagnosis</b>	<ul style="list-style-type: none"> <li>- Colonoscopy</li> <li>- Barium enema (<u>loss of haustration, drain pipe colon</u>)</li> <li>- CT \ MRI</li> <li>- Decreased goblet cells on histology</li> <li>- In children → <i>P-ANCA positive</i></li> </ul>	<ul style="list-style-type: none"> <li>- Barium swallow</li> <li>- CT</li> <li>- <b>Increased</b> goblet cells + <b>Granuloma</b> on histology</li> </ul>
<b>Treatment</b>	<ul style="list-style-type: none"> <li>➤ <b>Inducing remission</b> <ul style="list-style-type: none"> <li>- <b>1<sup>st</sup> line</b> (motions &lt;4 times) → topical aminosalicylates (e.g. <u>rectal mesalazine</u>)</li> <li>- If not responding → <u>oral mesalazine</u> (5-ASA)</li> <li>- Still not responding or motions 5/day → <u>oral prednisolone</u></li> </ul> </li> <li>If severe colitis → <b>IV steroids, Infliximab in children:</b> <ol style="list-style-type: none"> <li>1. &gt; 6 bowl movements</li> <li>2. Visible blood in large amount</li> <li>3. Pyrexia &gt; 37.8°C</li> <li>4. Tachycardia</li> <li>5. Anemic</li> <li>6. ESR &gt; 30</li> </ol> </li> <li><i>In severe cases, <u>an abdominal X-ray</u> would be very appropriate in the setting to look for features suggestive of <b>toxic megacolon</b></i></li> <li>➤ <b>Maintaining remission</b> <ul style="list-style-type: none"> <li>- <u>Mesalazine</u></li> <li>- If not well maintained → <u>oral azathioprine</u> or <u>mercaptopurine</u></li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li>➤ <b>Inducing remission</b> <ul style="list-style-type: none"> <li>- <b>1<sup>st</sup> line</b> → <u>prednisolone</u></li> <li>- <b>2<sup>nd</sup> line</b> → budesonide</li> <li>- <b>3<sup>rd</sup> line</b> → mesalazine (5-ASA)</li> </ul> </li> <li>• Add on treatment (not used as monotherapy): <b>azathioprine, mercaptopurine</b> or <b>methotrexate</b></li> <li>➤ <b>Maintaining remission after surgery</b> <ul style="list-style-type: none"> <li>- <b>1<sup>st</sup> line</b> → azathioprine, mercaptopurine or 5-ASA</li> </ul> </li> </ul>

## Toxic megacolon

- IBD or **infective colitis** characterized by total or segmental non-obstructive colonic dilatation + systemic toxicity

### Presentation

- Severe abdominal pain
- Marked toxicity (weakness, lethargy, confusion)

### Investigation

- Abdominal **X-ray**

### Treatment

- Admission to ITU, **IV fluids**
- **IV steroid** in case of IBD
- **IV antibiotics** in case of infectious cases
- Possible surgical resection (high risk of perforation and death)
- If rupture colon is suspected → **Urgent laparotomy**

## Zollinger-Ellison \$

- **Gastrinoma** (tumors found in pancreas or duodenum) → secretes **gastrin** → ↑ **gastric acid** → **peptic ulcers** at usual sites, such as 2<sup>nd</sup> part of duodenum or jejunum
- Ulcers may occur after adequate surgery

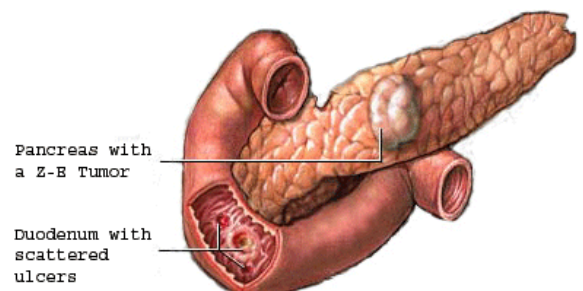
### Investigations

- **Fasting gastrin levels**
- **Secretin stimulation test** (gastrin goes up after secretin in case of Gastrinoma)

### ZES is suspected when

1. **Multiple ulcers** that are **resistant to drugs**
2. Associated with **diarrhea + steatorrhea**
3. **Family history of peptic ulcers**

Zollinger Ellison Syndrome



## Constipation management

- **Hard stool** → Stool softeners + high fiber (residue) diet
- **Soft stool** → Senna then lactulose
- **Impacted stool** → phosphate enema
- **Constipation in pregnancy** → Lactulose then Senna

## Colorectal carcinoma

- Presents with a change in bowel habits, abdominal pain, anemia and weight loss
- Best diagnostic investigation → *Colonoscopy*
- Gold standard → *Biopsy*
- Alternatives → *Barium enema* and *CT angiography*
- CEA antigen is NOT used for diagnosis or staging but rather for monitoring relapses



## Diarrhea

- 3 loose or watery stool /day
- Acute < 14 days → **Microscopy, culture and sensitivity**
- Chronic > 14 days → **Colonoscopy**
- Most common adenoma causing electrolytes disturbances → **Villous adenoma**
- Most common electrolyte imbalance in diarrhea → **Hypokalemia**
- Acid-base imbalance in diarrhea → **Non-anion gap metabolic acidosis** (due to loss of HCO<sub>3</sub>)
- Most common cause of bloody diarrhea → **Campylobacter** (a prodrome of *headache, myalgia & fever*)
- Second most common cause of bloody diarrhea → **Shigella** → **Salmonella**
- Diarrhea after camping → **Giardia**
- Most common cause of traveler diarrhea (in less than 72h) → **E-coli**
- Traveler diarrhea lasting ≥1 week and associated e' **steatorrhea** and **weight loss** → **Giardia**
- Most common cause of diarrhea in pediatrics → **Viral (Rotavirus)**
- Diarrhea followed by weakness and areflexia (Ascending paralysis) → **GBS**
- Diarrhea followed by renal impairment → **HUS (hemolytic uremic syndrome)**
- Bloody diarrhea followed by RUQ pain → **Ameba > amoebic liver abscess**
- Chronic bloody diarrhea in young male → **IBD**
- Diarrhea after long term antibiotics → **Clostridium difficile**
- MC Antibiotic causing clostridium difficile → **Clindamycin, Cephalosporin, Co-amoxiclav**
- TTT of clostridium difficile → **Metronidazole** (**Vancomycin** for severe cases/ failure to respond to metronidazole)
- Diarrhea after eggs or chicken → **Salmonella** → **Ciprofloxacin**
- Diarrhea/vomiting just hours after meal → **Staph toxin**
- Diarrhea in bed ridden with constipation → **Fecal impaction**
- Main TTT of diarrhea → **Fluid**
- TTT of traveler diarrhea → **Fluid only**
- TTT of staph toxin → **Fluid only**
- TTT of shigella or campylobacter → **Antibiotics**
- TTT of ameba or giardia → **Metronidazole**
- 1<sup>st</sup> Inv of choice for acute diarrhea (<14 days) → **Stool C&M**
- HIV + watery diarrhea → **Cryptosporidium parvum**
- HIV + bloody diarrhea → **CMV**
- **Long standing** diarrhea after recent travel:
  1. Non-bloody / watery / steatorrhea → **Giardiasis** (1<sup>st</sup> → *Stool microscopy, then stool PCR and ELISA*)
  2. Bloody → **Campylobacter jejuni** → **Gram-negative** curved bacilli → **Erythromycin**, alternatively → **Clari/Azi**, if Macrolides are not tolerated → **Ciprofloxacin**

### Overflow diarrhea

- Severe constipation can cause fecal impaction and fecal fluid will flow around the blockage
- Common in **bedridden** elderly
- Can be due to **opioid** pain relievers → reduces GI motility

### Fecal impaction

- **Symptoms**
  - Dehydration, agitation, tachypnea
- **Management**
  - Bisacodyl suppository
  - Arachis oil retention enema
  - **Phosphate enema**

## Celiac disease

- Caused by sensitivity to the protein **gluten** (exacerbated by consumption of **wheat**)
- Repeated exposure leads to **villous atrophy** which in turn causes **malabsorption** → *buttock atrophy in children*

### Signs and symptoms

- **Diarrhea**, chronic or intermittent
- **Stinking stools**, difficult to flush
- **Steatorrhea**
- Persistent GI symptoms (bloating, nausea & vomiting)
- Fatigue
- **Recurrent abdominal pain**, cramping and distension
- Sudden or unexpected **weight loss**
- Unexplained iron, folate or vitamin B12 deficiency **anemia**

- One of the most common presentation of celiac disease is **Iron deficiency anemia (IDA)**.
- May be associated with **type 1 DM**
- **Folate** deficiency is more common than vitamin B12 deficiency in celiac disease

### Complications

- **Osteoporosis**
- **T-cell lymphoma of small intestine** (rare)
- **Dermatitis herpetiformis** (presented as red raised patches, often with blisters and severe itching, treated with Dapsone)

- Any patient with confirmed celiac disease who experience recurrence of the symptoms despite gluten-free diet +/- weight loss → **Intestinal lymphoma** until proven otherwise

### Investigations

#### 1. Specific auto-antibodies

- **TTG antibodies (IgA)** → *the first choice*
- Endomysial antibody (IgA)
- Alpha-gliadin antibodies

#### 2. Jejunal/duodenal biopsy (Still needed even if antibody test confirms the disease)

- Villous atrophy
- Crypt hyperplasia
- Increase in intraepithelial lymphocytes

- Patients should not start gluten free diet until diagnosis is confirmed
- If patients are already taking a gluten-free diet they should restart consuming gluten for at least 6 weeks prior to testing

**Management** → *Gluten-free diet*

## Pseudomembranous colitis

- **Clostridium difficile** is detected in stool, presented with watery diarrhea (could be bloody), abdominal pain, raised WBCs and fever

### Most common antibiotics that cause it

1. **Clindamycin**
2. Cephalosporin (2<sup>nd</sup> and 3<sup>rd</sup> gen)
3. Co-amoxiclav
4. Quinolones
5. Aminopenicillins (amoxicillin and ampicillin)

### Management (only if symptomatic)

1. Stop the causative antibiotic
2. 1<sup>st</sup> line → **Oral metronidazole**
3. If severe/not responding → Oral vancomycin (to be absorbed in the gut)

#### **Differentiating tropical sprue from celiac disease:**

1. Origin: infective in TS, autoimmune in CD
2. TS along with IDA will present with symptoms of other vitamins deficiencies (Vit D, K, A). usually begins with acute diarrhea then progresses to chronic state and malabsorption and steatorrhea
3. No dermatitis herpetiformis in tropical sprue
4. Biopsy: incomplete villous atrophy in TS while complete in CD



## Jaundice

### 1. Prehepatic

- ↑ Heme breakdown (e.g. hemolysis, G6PD deficiency, malaria) → ↑ unconjugated hyperbilirubinemia

### 2. Intrahepatic

- Inability to conjugate
  - **Gilbert's \$** → ↑ unconjugated hyperbilirubinemia
  - Crigler-Najjar \$
  - liver disease
- Inability to excrete
  - **Dubin-Johnson \$** → ↑ conjugated hyperbilirubinemia
  - PBC

### 3. Post-hepatic/obstructive (surgical)

- Gallstones
- Cholangitis

#### Gilbert's \$

- Due to ↓ UGT-1 which is the enzyme that conjugate bilirubin with glucuronic acid
- Jaundice is precipitated by: infection, illness, physical exertion, stress or fasting

#### Investigations

1. Mildly raised bilirubin (<100) with normal LFTs
2. FBC → normal reticulocyte count
3. Urine dipstick → normal

- ↑ Reticulocyte count → Hemolysis
- Total bilirubin >100 → Crigler Najjar
- Abnormal dipstick + Dark liver → Dubin-Johnson (↑ conjugated)

## Primary Biliary Cholangitis/Cirrhosis

- **Autoimmune, idiopathic**
- Associated with → Sjogren syndrome & RA
- Pruritus, ↑ALP
- **+ve AMA** (ant-mitochondrial antibodies)
- Treatment
  - Ursodeoxycholic acid
  - Cholestyramine

## Primary Sclerosing Cholangitis

- Autoimmune, idiopathic
- Fibrosis at some areas of bile ducts "beaded appearance"
- Associated with → IBD (especially UC)
- Pruritus, ↑ALP
- **ERCP** → **the most specific**
- Treatment:
  - Ursodeoxycholic acid
  - Cholestyramine

### PBC – the M rule:

- IgM
- AMA
- Middle aged female

## Ascending cholangitis

- Due to ascending bacterial infection (E. coli) as a result of choledocholithiasis
- **Charcot's triad**
  - *Fever*
  - *RUQ pain*
  - *Jaundice*
- Confirmed by US gallbladder and biliary ducts
- **Complications** → Reynold's triad = Charcot's triad + hypotension + confusion
- **Treatment** → Emergency ERCP, rehydration & antibiotics

- Investigations for **cholecystitis** → US – HIDA
- Only symptomatic Gallstones are treated with elective cholecystectomy. However, **CBD stones** may need laparoscopic cholecystectomy regardless if they're symptomatic or not
- **Choledocholithiasis** (stones in CBD) → **ERCP** (both diagnostic & therapeutic)
- One complication of ERCP → Acute pancreatitis
- **Cholangitis** → inflammation of the bile ducts

## Autoimmune hepatitis (AIH)

- Chronic disease of unknown cause characterized by continuing hepatocellular inflammation and necrosis, which tends to progress to cirrhosis. Often seen with autoimmune diseases (autoimmune thyroid disorder, Addison's or vitiligo)
- Middle-aged women

### Features

- Fever, malaise
- Rash, Polyarthritis
- *Pulmonary* infiltration, pleurisy
- *Glomerulonephritis*
- Liver enzymes are usually elevated
- Amenorrhea is common and disease tends to attenuate in pregnancy

- **Deranged LFTs + 2ry amenorrhea** → autoimmune hepatitis
  - AST & ALT are usually elevated initially
  - ALP is normal or slightly raised, > 2 folds suggests another diagnosis

### Investigations

- **ANA/SMA/LKM1** antibodies, raised **IgG** levels
- **Liver biopsy** → inflammation extending beyond limiting plate '**piecemeal necrosis**', bridging necrosis

### Management

- **Steroids**, other immunosuppressants e.g. azathioprine
- Liver transplantation

## Acute fatty liver of pregnancy

### Risk factors

- Pre-eclampsia
- First pregnancies
- Multiple pregnancies

### Features

- Begins after 30 weeks of gestation, may also appear immediately after delivery
- Presents acutely with:
  - Nausea, vomiting
  - Abdominal pain
  - Fever, headache
  - Jaundice, pruritis

### Investigations

- Elevated LFTs
- Raised bilirubin
- Hypoglycemia & ammonia
- Prolonged PT
- **Liver biopsy** → *diagnostic*

### Management

- Treat hypoglycemia
- Correct clotting disorders
- N-acetylcysteine (NAC)
- Consider early delivery

*AFL = HELLP + Hypoglycemia + Ammonia*

## Causes of elevated liver enzymes in postpartum period

### ➤ Pregnancy related liver diseases

- Obstetric cholestasis → severe pruritis due to high bile acids + x20 ALT
- Pre-eclampsia / Eclampsia
- HELLP syndrome
- Acute fatty liver of pregnancy

### ➤ Liver diseases unrelated to pregnancy

- Viral hepatitis
- Autoimmune liver disease
- Wilson's
- Budd Chiari \$
- Acute cholecystitis
- Drug-induced hepatotoxicity

## Hemochromatosis

- Autosomal recessive condition in which increased intestinal absorption of iron causes iron accumulation in tissues especially the liver which may lead to cirrhosis and HCC (hepatoma)
- Iron is accumulated mainly in peripheral hepatocytes and not in Kupffer cells, while in Hemosiderosis, iron is accumulated in Kupffer cells and more in central rather than peripheral hepatocytes

### Presentation

- Often asymptomatic until late stages
- Symptoms usually start at 40-60 years old
- Initial symptoms are vague and non-specific
- Iron overload might cause symptoms of arthropathy and gynecomastia
- Maybe diagnosed accidentally following LFTs or abnormal serum ferritin
- Symptoms of advanced disease include a triad of:
  1. **Diabetes** (bronze diabetes)
  2. **Hepatomegaly**
  3. **Bronze pigmentation**
- It may also include cardiac diseases (arrhythmias or cardiomyopathy) or neurological/psychiatric symptoms

- Hemochromatosis can lead to cardiomyopathy and it predisposes to HCC

## Acute pancreatitis

- Main causes are **gallstones** and **alcohol**

### GET SMASHED

- Gallstones
- Ethanol
- Trauma
- Steroids
- Mumps
- Alcohol, autoimmune
- Scorpion venom
- Hypertriglyceridemia, ↑Ca, hypothermia
- ERCP
- Drugs (azathioprine, mesalazine, Bendroflumethiazide, frusemide, didanosine, pentamidine, sodium valproate)

### Features

- Gradual or sudden **severe epigastric pain** or central abdominal pain (radiates to the back, relieved by sitting forward)
- **Vomiting** is prominent
- **Tachycardia**
- Fever
- Jaundice
- **Shock**
- **Rigid abdomen** with local **tenderness**
- **Periumbilical bruising** (Cullen's sign)

### Investigations

1. **Serum amylase**: >1000U/mL (or around 3-fold upper limit of normal). However, **lipase levels** are more sensitive and more specific but takes more time to rise following an attack (>24h)
2. **CT with contrast**

### Treatment

- **Fluid resuscitation, analgesia** and **nutritional** support
- **IV antibiotics** (e.g. IV imipenem) after obtaining culture
- **Laparoscopy** → only when there's infection or necrosis

## Albumin

- Made in the Liver
- Its half-life is about 20 days so it's a good parameter for **chronic liver disease**
- Its main **function** is to regulate the oncotic pressure of blood, it also binds to enzymes and hormones
- Shifts fluid into the intravascular compartment
- Produces only a **transient effect** but it's useful in cases of intravascular fluid deficit and edema
- Also, useful to obtain diuresis in hypoalbuminemic patient

## Chronic pancreatitis

**Causes** → alcohol, smoking, autoimmune

### Features

- **Epigastric pain**
  - **Episodic** with short periods of severe pain
  - **Pain free intervals** are specific to chronic pancreatitis
  - Radiates to the back
  - Relieved by sitting forward
  - Exacerbated by eating
- **Steatorrhea**
  - Due to malabsorption of fats from the lack of pancreatic lipase secretion which results on weight loss
  - Sometimes described as “loose, offensive stools which are difficult to flush”
- **Diabetes**
- **Jaundice**
  - A late presentation due to obstruction of CBD

### Investigations

1. **Serum amylase/lipase**
2. **US**
3. **CT with contrast**
  - Gold standard
  - Shows evidence of pancreatic calcification

### Management

- Pain → Analgesia
- Steatorrhea or malabsorption → Pancreatic enzymes supplements and fat-soluble vitamins
- Diabetes → Oral hypoglycemics and insulin

## Drug-induced hepatitis

- Drugs → *Co-amoxiclav, flucloxacillin, steroids, Sulphonylurea*
- Lab → Elevated bilirubin + massive increase in ALP and AST
- History of alcohol intake

## Mediastinitis

- May occur after esophageal perforation (after endoscopy)
- Anterior mediastinitis → pain located in the substernal region
- Posterior mediastinitis → pain in the epigastric region with radiation to the interscapular region
- X-ray may show **widened mediastinum** or **air in the mediastinum**

## Notes

- **Alcoholic hepatitis** → look for **GGT** in acute cases, **AST** is higher than ALT in chronic cases
- **Hematogenous** spread to the **liver** from the lungs is the most common route of metastasis
- **TTF-1** is a protein seen by immunostaining which is used as a clinical marker of **lung adenocarcinoma**
- **Gastroenteritis**
  - Very common in hospitals, especially with the spread of **norovirus**
  - They present with acute onset of diarrhea (sometimes with vomiting) and abdominal pain
  - Pain is usually central, could be epigastric
  - Patients with gastroenteritis should be isolated and given a single side room until diarrhea resolves
- **Dysentery** → infection of the intestine that leads to severe diarrhea (**blood + mucous**) and abdominal pain
- **Cancer head of pancreas** → Courvoisier sign (painless obstructive jaundice with a palpable mass)
- **Celiac disease** is associated with **lymphoma**
- **Ulcerative colitis** and **Crohn's disease** are associated with **colon cancer**
- In **IBS** → bloating, constipation alternating with diarrhea + NO blood in stool
- **Fecal calprotectin** → biochemical measurement of the protein calprotectin in the stool, if elevated → **IBD**, if normal, suspect **IBS**
- **Proctalgia fugax** → Severe recurrent rectal pain in the absence of any organic disease, may occur at night, after bowel actions or after ejaculation. Anxiety could be an associated feature
- If the patient has Ascites + bleeding → **Terlipressin**
- If the patient has Ascites without bleeding → **Perform ascitic fluid aspiration**, to detect *Neutrophil count, gram stain, culture and obtain protein level*
- Raised neutrophil count **>250** → *Start antibiotic treatment*
- Most accurate test would be a **culture** but it takes days
- **Spontaneous bacterial peritonitis**
  - One of the complications of Ascites
  - Includes abdominal pain, rebound tenderness, absent bowel sounds and fever
  - Organism usually reaches the peritoneum via hematogenous spread
- Recurrent abdominal pain (often peri-umbilical) with episodic headache in a child with no abnormal finding on examination and investigation, pain interferes with normal activities and associated with anorexia, nausea, vomiting → **Abdominal migraine** → **Reassure**