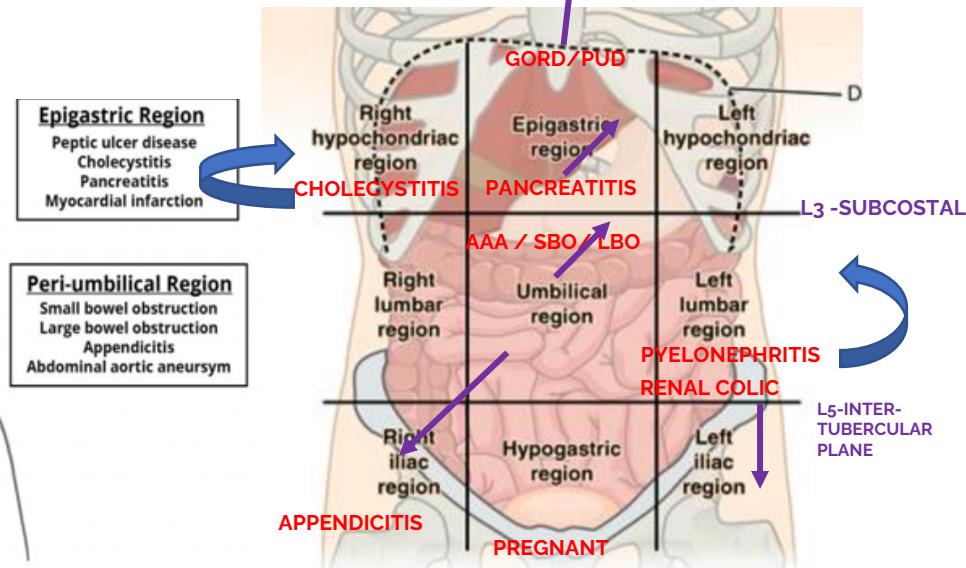
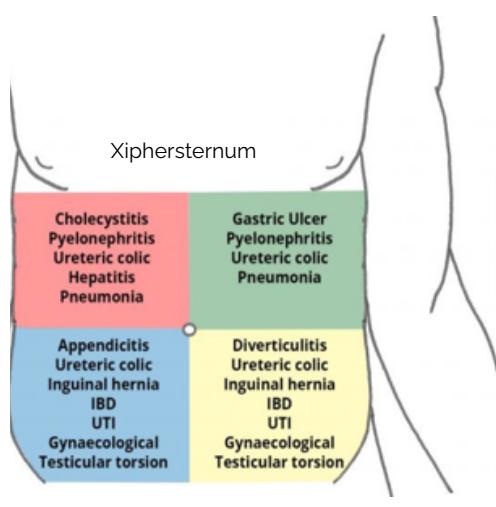


## GASTROENTEROLOGY H+E:

	Details			
History of presenting complaint [DDD's]	• Diarrhoea /contraption	• <b>Altered bowel habit:</b> Volume, smell, colour, freq, consistency. • <b>Weight gain/loss</b> • <b>Appetite/Anorexia</b>	<b>Red flags: ALARMS</b> <ol style="list-style-type: none"> <li>1. <b>A</b>naemia</li> <li>2. <b>L</b>oss of weight</li> <li>3. <b>A</b>norexia</li> <li>4. <b>R</b>ecent onset progressive symptoms</li> <li>5. <b>M</b>elaena/Haematemesis / Persistent vomiting [<b>Melaena</b> (Exc. Fe tablets, peptic ulcers, small bowel bleed)]</li> <li>6. <b>S</b>wallowing difficulty /dysphagia (esp. &gt;55 y.o., &gt;4 weeks/relapsing)</li> <li>7. <b>Painless bleed</b> = aortic enteric fistula (AAA)</li> <li>8. <b>Painful bleed</b> = fissure, IBD, abscess</li> </ol>	
	• Dysphagia	• Solids/liquids/both (which came 1 <sup>st</sup> ) • Intermittent + due to food = eosinophilic oesophagitis • Odynophagia		
	• Dyspepsia	• Acid regurgitation (GORD) • <b>Abdominal Pain (Socrates)</b> o CHRONIC + altered bowel habit = IBS		
	• General	• N +V + bloating + pruritus • Fever + NS + fatigue		
	• Urological [FUND WISE]	• <b>Storage:</b> freq, vol, urgency, nocturia, incontinence • <b>Infection:</b> dysuria, Haematuria, odour		
	• Gynaecological	• <b>PV</b> bleeding: menorrhagia, post-coital • <b>PV</b> discharge • <b>Pain:</b> pelvic, dysmenorrhoea, dyspareunia • <b>Pregnancy</b>		
Past MHx [CHOMV STAVE]	<ul style="list-style-type: none"> <li>• <b>Conditions:</b> Previous <b>H. pylori infections</b>   <b>gallstones</b>   <b>Diabetes</b>   <b>SCI/MS/Hirschsprung disease</b>   <b>Thyroid issue</b></li> <li>• <b>Medications</b> (calcium antacids, opiates, TCAs cause constipation)</li> <li>• <b>Surgeries - Abdominal</b> (appendectomy, anaesthesia)</li> <li>• <b>Tests (exploratory)</b> = (biopsies, tTg-IgA, colonoscopy, gastroscopy)</li> <li>• <b>Allergies?</b> (e.g. to aspirin, NSAIDs, intravenous contrasts)</li> </ul>			
Social Hx [WHIT SADOM]	<ul style="list-style-type: none"> <li>• Home life/accommodation + partners + single (sex – condom?)</li> <li>• <b>Occupation:</b> HCW (hepatitis exposure)</li> <li>• <b>Tattoos &amp; Illicit Drugs</b> (injections)</li> <li>• <b>Smoking</b> (pack years) + when did they quit</li> <li>• <b>Alcohol</b> (CAGE questions) = <b>ESSENTIAL</b></li> </ul>			
Family Hx	<ul style="list-style-type: none"> <li>• Family Hx of <b>colon cancer</b> (esp. family polyps)   <b>IBS</b>   <b>Coeliac disease</b>   <b>H. pylori infections</b>   <b>pancreatitis</b></li> <li>• <b>Hemolytic anemia</b> or <b>congenital hyperbilirubinemia</b> = family Hx of jaundice, anemia, splenectomy or cholecystectomy</li> </ul>			



### Acute Abdominal Pain (Visceral vs. Peritoneal (somatic) Pain)

- **Visceral** pain (non-localised | focal) starts much earlier than **peritoneal** pain (localised – somatic pain – may radiate/referred pain)
- **Visceral** pain **Creates** "MIGRATORY" pattern → DIVIDED into foregut, midgut or hindgut pain depending on blood supply

	Division	Location	Level	Somatic Pain	SNS	PSNS
Celiac trunk	Foregut	RUQ and LUQ	T12	T7-9 (epigastric)	Greater splanchnic T5-9	Vagus (CNX)
SMA	Midgut	to proximal 2/3 transverse colon (Central/paraumbilical)	L1	T10-11 (umbilical)	Lesser splanchnic T10-11	Vagus (CNX)
IMA	Hindgut	RLQ and LLQ	L3	T12-L1 (suprapubic)	Least splanchnic T12-L2	Pelvic splanchnic (S2-4)

## OTHER MAJOR SYMPTOMS IN DETAIL:

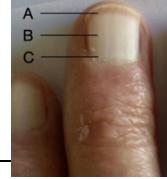
<b>VOMITING</b> "contents frequency amount"	<ul style="list-style-type: none"> <li>Involuntary explosive ejection of stomach contents through mouth</li> </ul>	
	<b>Colour</b>	<b>Location</b>
	Bile (green-yellow)	SBO, duodenal atresia,
	Faeculent (brown)	Small or large bowel obstruction
	Coffee ground	Fe tablet, red wine & coffee ingestion
	Undigested Food	Gastric outlet obstruction (level of pylorus)
	<b>Red blood/ Haematemesis</b>	<ul style="list-style-type: none"> <li><b>Upper GI Bleeding</b> or malignancy (any bleeding proximal to Duodenal-Jejunal flexure)</li> <li><b>Multiple</b> episode of vomiting? = <b>Mallory-Weiss Tear</b></li> </ul>
<b>CHANGE OF BOWEL HABIT</b>  (Wide range of normal: 3 x /day to once every 3 days)  "Have you noticed your bowel habit change? What happened?"  "Do you mean less/more frequent toilets or change in stool"	<b>DDx</b>	<b>KEY FEATURES</b>
	Colon cancer	<ul style="list-style-type: none"> <li>Elderly &gt; UWL &gt; Anaemia (SOB, fatigue)</li> <li><b>Melaena</b> (Exc. Fe tablets, peptic ulcers, small bowel bleed)</li> </ul>
	Gastroenteritis	<ul style="list-style-type: none"> <li>Acute diarrhoea</li> <li>N+V</li> </ul>
	IBS (Rome IV criteria - at least 2 of 3)	<ul style="list-style-type: none"> <li>(1) abdominal cramping pain/bloating → relieved on defecation</li> <li>(2 + 3) Altered stool frequency + appearance → Fluctuate diarrhoea/constipation "toothpaste stools"</li> <li>Anxiety/stress &gt; Jaundice &gt; Pruritus, lethargy</li> </ul>
	Coeliac disease	<ul style="list-style-type: none"> <li><b>Steatorrhoea</b> (fat malabsorption)           <ul style="list-style-type: none"> <li>excl. pancreatic disease with lipase deficiency, OR use of weight-loss drug - orlistat</li> </ul> </li> <li>Assoc with: <b>Anaemia, T1DM, Osteoporosis, autoimmune disease</b> (e.g. thyroid), <b>malignancy</b> (e.g. lymphoma, small intestinal carcinoma)</li> <li><b>Abd discomfort</b> (N+V, flatulence, cramping)</li> </ul>
	IBD (UC or crohn's)	<ul style="list-style-type: none"> <li><b>Bright red Blood (haematochezia)/mucus in stool</b> (exc. colon cancer, beetroot ingestion)</li> <li>Abd pain + weight loss           <ul style="list-style-type: none"> <li>Crohn's = affects entire GIT</li> <li>UC = affects only colon</li> <li><b>Small bowel or Right colonic bleeding</b> = Maroon-coloured blood mixed with stool</li> <li><b>Left colonic bleeding (UC)</b> = Fresh blood mixed</li> <li><b>Anal fissure, Haemorrhoids</b> = Anorectal Bleeding (NOT mixed in stool - only coated!)</li> </ul> </li> </ul>
	Anal fissure, Haemorrhoids	<ul style="list-style-type: none"> <li><b>Anorectal Bleeding</b> (COATING OF BLOOD - not mixed in stool)</li> <li>Straining/post-defaecation anal pain</li> </ul>
	Endocrine	<ul style="list-style-type: none"> <li><b>Thyrotoxicosis</b> = diarrhoea, heat intolerance, irritable, tremor, amenorrhoea</li> <li><b>Hypothyroidism</b> = constipation, cold intolerance, lethargy, menorrhagia</li> </ul>
	Other	<ul style="list-style-type: none"> <li>Drugs ( opiates, Fe, antacids, Abs), chronic infection, overflow constipation</li> <li>Lactose intolerance, dietary &amp; lifestyle changes</li> <li><b>bowel obstruction</b> (not passing flatus)</li> </ul>
	Heartburn / Regurgitation	<ul style="list-style-type: none"> <li><b>GORD</b>, but ALSO <b>oesophagitis</b> or <b>malignancy</b> Ask about <b>chronicity, frequency</b> and <b>medication</b></li> <li><b>Assoc:</b> weight loss   severity   orthopnoea   PND   Dysphagia   Malignancy   Resp. infection</li> </ul>
	Dysphagia (Difficulty swallowing)	<ul style="list-style-type: none"> <li>If just <b>solids</b> → mechanical issue /Obstruction           <ul style="list-style-type: none"> <li><b>Internal:</b> oesophageal carcinoma, eosinophilic esophagitis (intermittent), foreign body</li> <li><b>External:</b> Retrosternal goitre, mediastinal tumours, bronchial carcinoma</li> </ul> </li> <li>If <b>liquids</b> → motility issue   Or issue with <b>BOTH?</b> <ul style="list-style-type: none"> <li>MND → bulbar or pseudobulbar palsy</li> <li>Polymyositis, myasthenia gravis</li> <li><b>Progressive worsening</b> = Achalasia, scleroderma</li> </ul> </li> </ul>
	Odynophagia (Painful swallow)	<ul style="list-style-type: none"> <li><b>infectious oesophagitis</b> (e.g. candida, HSV)</li> <li><b>radiation or caustic damage to oesophagus</b></li> </ul>
<b>REFLUX / SWALLOWING DIFFICULTY</b>	<b>Personal</b>	<ul style="list-style-type: none"> <li>Intentional vs unintentional</li> <li>Collateral history from relatives about Mental stress or mood is also helpful</li> </ul>
	<b>Early Satiety</b> (post-prandial fullness)	<ul style="list-style-type: none"> <li>Do you feel hungry? Complete loss of appetite? (neurological - hypothalamus)</li> <li>indicator of <b>gastric outlet obstruction</b> or <b>impaired gastric emptying (motility related → impaired peristalsis due to diabetes)</b> causing anorexia and weight loss</li> </ul>
	<b>Rapid unexplained weight loss</b>	<ul style="list-style-type: none"> <li>What size of pants did you wear 6 months ago? How about now?</li> <li>Have you been on any diet?</li> <li>Rapid unexplained weight loss is concerning, especially if no recent dietary change</li> </ul>
	<b>Weight loss + increased appetite</b>	<ul style="list-style-type: none"> <li>Malabsorption of nutrients or hypermetabolic state (e.g. <b>thyrotoxicosis</b>)</li> <li><b>Liver disease</b> = taste disturbance (can cause smokers to give up smoking)</li> </ul>
	<b>Jaundice</b> (see under natural light)	<ul style="list-style-type: none"> <li>Excess bilirubin deposited in conjunctivae and skin → <b>LIVER FAILURE</b></li> <li>Have you noticed your skin or eye colour change?</li> <li>Have you noticed your urine or stool colour change?</li> </ul>
<b>JAUNDICE / DARK URINE / STOOL</b>	<b>Dark urine</b>	<ul style="list-style-type: none"> <li>Have you noticed your urine becoming dark or "tea" coloured?           <ul style="list-style-type: none"> <li>NOTE: concentrated urine may be called dark urine</li> </ul> </li> <li>Pale stool + dark urine = <b>obstructive or cholestatic jaundice</b> (urobilinogen cannot reach intestine)</li> </ul>
	<b>Stool</b>	<ul style="list-style-type: none"> <li>Fatty stool = steatorrhoea (nutrient malabsorption → e.g. coeliac disease, pancreatitis)</li> <li>Pale stool = <b>post-hepatic</b> jaundice</li> </ul>
	<b>Pruritis</b>	<ul style="list-style-type: none"> <li>Itching of the skin (generalised or localised) → <b>Cholestatic Liver Disease</b></li> </ul>
<b>Lethargy</b>		<ul style="list-style-type: none"> <li>Tiredness   easy fatigability = <b>acute or chronic liver disease</b> OR <b>anaemia due to GI or chronic IBS</b></li> </ul>

## GIT examination

### RED FLAG (ACUTE ABDOMEN)

1. REBOUND TENDERNESS
2. GUARDING/RIGIDITY
3. SEPTIC SIGNS
4. ABSENT BOWEL SOUNDS  
"bleed/perforation, obstruction, infarction"

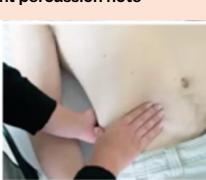
Ask patient to lie 45° (both hands on side)

General inspection	<ul style="list-style-type: none"> <li>1. Equipment (e.g. O<sub>2</sub>, nasogastric feeds, medications?), <b>Vomit bowls, IV infusions, catheters</b></li> <li>2. Jaundice (liver disease)</li> <li>3. Body habitus (obese, cachexia, ascites)</li> <li>4. Well/unwell (responsiveness) → <b>Mental state (encephalopathy) / Pain / Agitation (thyrotoxicosis)</b></li> <li>5. Pigmentation (haemochromatosis, Whipple's disease) &amp; <b>Xanthomata</b> (chronic cholestasis)</li> </ul>	
Nails	<ul style="list-style-type: none"> <li>• <b>Clubbing</b> (cirrhosis, IBD, coeliac)</li> <li>• <b>Leuconychia</b> (transverse white lines → <b>hypoalbuminaemia</b> from chronic liver disease)</li> <li>• <b>Koilonychia</b> (spoon-shaped nails from <b>severe Fe or B12/Folate deficiency</b>)</li> <li>• <b>Terry's nail</b> (red line on tip of nail) = <b>Chronic liver disease</b></li> </ul>	 
Palms	<ul style="list-style-type: none"> <li>• <b>Pale palmar creases</b> (anaemia)</li> <li>• <b>Palmar erythema</b> (chronic liver disease / pregnancy)</li> <li>• <b>Dupuytren's contractures</b> (alcohol) = thickened palmar fascia → fixed finger flexion usually ring or 5<sup>th</sup> finger <ul style="list-style-type: none"> <li>◦ EtOH, CLD, anti-epileptic</li> </ul> </li> <li>• <b>Asterixis/hepatic flap (hepatic encephalopathy) / ureaemia</b> → jerky movements of wrist due to toxic ammonia retention)</li> <li>• <b>Fingertip capillary glucose monitoring marks (diabetes)</b></li> </ul>	 
Arms	<ul style="list-style-type: none"> <li>• <b>Measure BP</b> (ask for BP in exam, don't to do it)</li> <li>• <b>IV Track marks</b> (→ hepatitis C infection?) or <b>tattoos</b> (unregulated pallor, prison)</li> <li>• <b>Bruising</b> (clotting factor deficiency – Vit K deficiency)</li> <li>• <b>Spider naevi</b> (chronic liver disease) (&gt; 2 is abnormal) = Blanches <ul style="list-style-type: none"> <li>◦ Petechiae does NOT blanch</li> </ul> </li> <li>• <b>Muscle Atrophy</b></li> <li>• <b>Scratch marks</b> (severe itch (pruritus) = obstructive or cholestatic jaundice = due to primary biliary cirrhosis and progressive bile duct destruction)</li> </ul>	 
Axilla	<ul style="list-style-type: none"> <li>• <b>lymphadenopathy</b></li> <li>• <b>Acanthosis nigricans</b> = darkening and thickening of skin (rarely assoc. with GIT carcinoma) <ul style="list-style-type: none"> <li>◦ Hyperinsulinemia → Acromegaly, PCOS, T2DM</li> </ul> </li> </ul>	
Face / eyes	<ul style="list-style-type: none"> <li>• <b>Uveitis</b> = IBD, autoimmune</li> <li>• <b>Sclerae</b>: jaundice, anaemia, iritis</li> <li>• <b>Xanthelasma (biliary cholangitis/cholestasis)</b> → yellowish lipid plaque in periorbital region</li> <li>• <b>Conjunctival pallor (anaemia)</b></li> <li>• <b>SLIT-LAMP = Cornea</b>: brown-greenish <b>Kayser–Fleischer rings (Wilson's disease)</b> → (abnormal Cu deposition) → low serum ceruloplasmin &amp; Cu</li> <li>• <b>Parotidomegaly</b> = XS EtOH, MUMPS, Sjogrens, IgG4 disease</li> </ul> <p><b>STIGMA OF CHRONIC LIVER DISEASE:</b></p> <ul style="list-style-type: none"> <li>➢ Jaundice → hepatic encephalopathy (asterixis)</li> <li>➢ Hair loss</li> <li>➢ XS E2 = palmar erythema, spider naevi</li> <li>➢ Ascites → SBP</li> <li>➢ Esophageal varices, caput medusae, haemorrhoids (variceal bleed)</li> </ul>	 
Mouth	<ul style="list-style-type: none"> <li>◦ <b>Angular stomatitis</b> (cracks at corner of mouth) → Fe, folate or B12 deficiency and water soluble Vit deficiencies (e.g. B2, B6, B12, B1/thiamine)</li> <li>◦ <b>Aphous ulcers</b> (IBD or coeliac disease) = tiny ulcers in lips and mouths</li> <li>◦ <b>Sweet Breath Smell</b>: fetor hepaticus (end-stage chronic liver disease)</li> <li>• <b>Teeth</b>: Poor dentition (chronic liver disease, alcoholic)</li> <li>• <b>Gums: gingivitis</b> (smoking), <b>hypertrophy</b> (pregnancy, scurvy),</li> <li>• <b>Tongue: atrophic glossitis</b> = (Fe, folate or B12 deficiency)</li> </ul>	 
Neck/ Chest	<ul style="list-style-type: none"> <li>• <b>Cervical lymphadenopathy</b> <ul style="list-style-type: none"> <li>◦ esp. <b>Virchow's node</b> left supraclavicular LN which is linked to GI malignancy</li> <li>◦ Troisier's signs = presence of Virchow's node + gastric cancer</li> </ul> </li> <li>• <b>Hyperpigmentation</b> <ul style="list-style-type: none"> <li>◦ heavy metals</li> <li>◦ haemochromatosis</li> <li>◦ Addison's</li> </ul> </li> </ul> <p><b>Inspect front/back</b></p> <ul style="list-style-type: none"> <li>• <b>Spider naevi</b> [Blanching test = sign of cirrhosis]</li> <li>• <b>Gynaecomastia</b> (male breast enlargement and body hair loss) <ul style="list-style-type: none"> <li>◦ <b>Chronic liver disease and cirrhosis</b></li> <li>◦ <b>Alcohol</b> = ↓Leydig cell = teste atrophy + gynecomastia</li> <li>◦ <b>Drugs</b> (DISCO MTV – digoxin, isoniazid, spironolactone, cimetidine, E2, methyldopa, TCA, verapamil)</li> <li>◦ <b>Endocrine</b> (Hypogonadism, Hyperthyroidism)</li> <li>◦ <b>Seminoma</b> (testicular cancer) = ↑B-HCG, ↑E2</li> </ul> </li> <li>• <b>Cullen's</b> (central umbilical) and <b>grey-turner's</b> (flank – retroperitoneal haemorrhage) sign (pancreatitis)</li> </ul>	 <p>Gynaecomastia with prominent breasts and unassociated with confounding obesity</p>

Ask patient to lie flat (both hands on side) + pillow under head FOR ABDO EXAM

<b>Abdomen (begin here if asked to do so)</b>	<b>Inspect</b>	<p>Stand back then squat down beside bed at eye level (look for distensions)</p> <ul style="list-style-type: none"> <li>• <b>Distension (Fluids, flatus, faeces, fat, foetus, "filthy" big tumours, fibroids, fake pregnancy,</b> <ul style="list-style-type: none"> <li>◦ liver cirrhosis → hypoalbuminemia → ascites</li> <li>◦ Acute sudden distension due to flatus: IBS (benign) or post-bowel surgery</li> </ul> </li> <li>• <b>Scars/Stomas</b> (indicates previous bowel surgery) → CHECK position of scar</li> <li>◦ <b>Caput Medusae</b> = distended veins flowing away from umbilicus in Portal HT (e.g. liver cirrhosis, persistent ductus venosus)</li> <li>◦ <b>IVC obstruction</b> = distended veins flowing towards umbilicus</li> <li>• <b>Bruising/Pigmentation/Striae</b> (pregnancy, Cushing's)</li> <li>• <b>Visible peristalsis + Pulsations (AAA)</b></li> </ul>																								
	<b>Palpate</b>	<p>1. Any pain? (begin to palpate away from pain) → watch for discomfort</p> <p>a. Press down with stethoscope to check for TRUE guarding</p> <p>2. <b>Palpate over 9 regions</b> (superficial then deep) → Look for tenderness, rigidity, organomegaly</p> <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <td style="width: 25%; text-align: center; padding: 5px;"> <b>Painful</b> </td><td style="width: 25%; text-align: center; padding: 5px;"> <b>Causes Of Abdominal Masses</b> </td><td style="width: 25%; text-align: center; padding: 5px;"> <b>Causes Of Abdominal Masses</b> </td><td style="width: 25%; text-align: center; padding: 5px;"> <b>Causes Of Abdominal Masses</b> </td></tr> <tr> <td style="text-align: center; padding: 5px;"> <b>Inflammatory</b> </td><td style="text-align: center; padding: 5px;"> <b>Right Iliac Fossa</b> </td><td style="text-align: center; padding: 5px;"> <b>Left Iliac Fossa</b> </td><td style="text-align: center; padding: 5px;"> <b>Upper Abdomen</b> </td></tr> <tr> <td style="text-align: center; padding: 5px;"> <b>Neoplastic</b> </td><td style="text-align: center; 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padding: 5px;"> <ul style="list-style-type: none"> <li>• <b>Bladder</b></li> <li>• <b>Ovarian tumour or cyst</b></li> <li>• <b>Uterus</b> (e.g. pregnancy, tumour, fibroids)</li> <li>• <b>Small-bowel obstruction</b></li> </ul> </td></tr> <tr> <td style="text-align: center; padding: 5px;"> <b>Craggy</b> </td><td style="text-align: center; padding: 5px;"> <b>Tympanic</b> </td><td style="text-align: center; padding: 5px;"> <b>Reducible masses</b> </td><td style="text-align: center; padding: 5px;"> <b>Pelvis</b> </td></tr> <tr> <td style="text-align: center; padding: 5px;"> <b>Rebound tenderness</b> (peritonitis)         </td><td style="text-align: center; padding: 5px;"> <b>Palpable bowel loops, SBO/LBO</b> </td><td style="text-align: center; padding: 5px;"> <b>Hernia</b> – incisional, spigelian (semilunaris), direct vs indirect inguinal         </td><td style="text-align: center; padding: 5px;"> <b>AAA</b> </td></tr> <tr> <td style="text-align: center; padding: 5px;"> <b>Special tests</b> </td><td style="text-align: center; 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<b>Percuss</b>	<ul style="list-style-type: none"> <li>• <b>Ascites (shifting dullness)</b> → assoc. with portal HTN, splanchnic vasodilation, cirrhosis (hypoalbuminemia), raised RAAS           <ul style="list-style-type: none"> <li>◦ Percuss for dull note in 9 regions → if dull, keep hand in position and roll patient to opposite side</li> <li>◦ Keep hand in position → wait 30s → percuss again → if resonant = ascites</li> </ul> </li> <li>• <b>Percussion tenderness</b></li> </ul>																									
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<b>Abdo aorta</b>	<ul style="list-style-type: none"> <li>• Between umbilicus and xiphoid process (lateral → medial)</li> <li>• Expansile pulsation (<b>outward movement</b>) → AAA</li> </ul>																									
<b>AAW Masses</b>	<p>Patient to lift their head up → <b>intra-abdominal mass disappears or decrease in size</b></p> <ul style="list-style-type: none"> <li>• E.g. <b>sebaceous cyst, epigastric/incisional hernia, rectus sheath haematoma or diverticulum</b></li> </ul>																									
<b>Bladder</b>	<ul style="list-style-type: none"> <li>• Smooth and firm + <b>suprapubic dullness</b> over pubic symphysis → <b>bladder scan</b> → IDC (urinary retention)</li> </ul>																									
<b>DRE</b>	<ul style="list-style-type: none"> <li>• Anal sphincter tone (anal wink)</li> </ul>																									
<b>Legs</b>	<ul style="list-style-type: none"> <li>• Skin lesions           <ul style="list-style-type: none"> <li>◦ <b>Erythema nodosum</b> → IBD</li> <li>◦ <b>Pyoderma gangrenosum</b> → IBD</li> </ul> </li> <li>• <b>Arthralgia:</b> <ul style="list-style-type: none"> <li>◦ Enteropathic arthritis</li> </ul> </li> <li>• <b>Pitting Oedema</b> (hypoalbuminemia due to chronic liver disease)</li> <li>• <b>Neurological signs</b> (alcohol) → chronic liver disease</li> </ul>	 																								
<b>Summary</b>	<p>"Today I performed an abdominal exam on _____. On inspection, any peripheral stigmata of GIT disease.</p> <ol style="list-style-type: none"> <li>1. On palpation (was abdomen soft/tender/masses?) → On auscultation (normal bowel sounds? Bruits?)</li> <li>• To complete the exam, check external hernial orifices, inguinal LNs, external genitalia and perform DRE           <ul style="list-style-type: none"> <li>• <b>DRE + PV exam</b> → inspect (fistulae, tags, blood, mucus), <b>palpate</b> (masses) → look for rectal masses and haemorrhoids</li> <li>• <b>STOOL colour</b> → <b>pale</b> (chronic pancreatitis), <b>maleena</b> (haemorrhoids, colonic pathology)</li> <li>• <b>Urinalysis</b> (for bilirubin and urobilinogen, and glucose) → <b>Temperature chart</b> (infection)</li> </ul> </li> </ol>																									
<b>Further Ix</b>	<p><b>Bedside:</b></p> <ol style="list-style-type: none"> <li>1) Urine M/C/S</li> <li>2) Urine dipstick (bilirubin)</li> <li>3) Urine B-HCG</li> <li>4) ECG</li> <li>5) Stool M/C/S</li> <li>6) Bladder scan</li> </ol>	<p><b>Bloods:</b></p> <ol style="list-style-type: none"> <li>2. FBC</li> <li>3. EUC</li> <li>4. LFT</li> <li>5. CRP</li> <li>6. LIPASE/AMYLASE</li> <li>7. COAG +/- GROUP = HOLD</li> </ol>	<p><b>Imaging:</b></p> <ul style="list-style-type: none"> <li>➤ <b>USS</b> – transabdominal, transvaginal</li> <li>➤ <b>AXR</b></li> <li>➤ <b>ERECT CXR</b></li> <li>➤ <b>CT-abdo-pelvis +/- contrast</b></li> </ul>																							

## EXAMINATION OF THE ABDOMINAL ORGANS: (in detail)

	Hepatomegaly	Splenomegaly	Kidney (Bimanual method)
Cause	<ul style="list-style-type: none"> <li>Metastasis, hepatitis, RVF, leukaemia/lymphoma,</li> <li>fatty liver, alcoholic liver disease</li> <li>portal HTN</li> </ul>	<ul style="list-style-type: none"> <li>myelofibrosis</li> <li>leukaemia/lymphoma or myeloproliferative disorder</li> <li>malaria/EBV</li> </ul>	<ul style="list-style-type: none"> <li>Hydronephrosis</li> <li>carcinoma</li> </ul>
Location	<b>RIF → RUQ (Right costal margin)</b>	<b>RIF → LUQ (left costal margin)</b>	<ul style="list-style-type: none"> <li>○ <b>Left and right flanks</b></li> </ul>
Palpate	<ul style="list-style-type: none"> <li>Push hand <b>IN</b> towards right costal margin <b>during each Inspiration</b></li> <li><b>Liver edge</b> may be palpable on lateral margin with <b>deep palpation of index finger</b> *Small liver = cirrhosis</li> </ul> 	<ul style="list-style-type: none"> <li>Patient rolls to right side with tucked legs</li> <li><b>Spleen vs kidney</b></li> <li><b>Cannot get above spleen</b></li> <li><b>Splenic notch</b></li> <li><b>Spleen not ballotable</b></li> <li><b>Spleen moves inferior-medially on inspiration</b></li> <li><b>Dull percussion note</b> (unlike kidney which is resonant)</li> </ul>	<ul style="list-style-type: none"> <li>One hand anterior, one hand posterior → push up into renal angle with posterior hand and push down with anterior hand</li> <li><b>Flex MCP joints in posterior hand to "Ballot" kidney</b> [flick → flick → stop]</li> <li>Swap hands for other kidney</li> <li><b>*Normal size kidney = NOT palpable</b></li> </ul>
Percuss	<ul style="list-style-type: none"> <li>Repeat like palpation to find <b>lower border of liver (esp. on expiration)</b></li> <li>Percuss down from chest along <b>right mid-clavicular line</b> until you hear dullness (i.e. <b>top of liver</b>)</li> <li>Measure to the palpable liver edge (with ruler or width of hand (9-10cm wide))</li> </ul>	<b>Splenomegaly</b> = DULL percussion note <ul style="list-style-type: none"> <li>Percuss over left costal margin in the left anterior axillary line during complete inspiration</li> <li>Percuss <b>Traube's space</b> (above left costal margin in mid-clavicular line → dull note = splenomegaly)</li> </ul> 	 
Auscultate	Hepatic <b>friction rub</b> (liver cancers) → rough grating sound during breathing	For <b>Splenic rub/bruit</b> (splenic infarct)	For <b>renal bruits</b> (renal artery stenosis) → 5cm superior + lateral to umbilicus

## DIGITAL RECTAL EXAMINATION [DRE]

Indication	<ul style="list-style-type: none"> <li><b>Suspected haemorrhoids</b> (i.e. engorged external or internal BV) → Rx: rubber bands, proctosedyl topical</li> <li><b>Unexplained bleeding on history</b> → bright red (distal), mixed in (left colic - UC), Maroon colour (upper GI)</li> <li><b>Prostate screening</b> → oliguria</li> <li><b>Altered bowel habits</b> (esp. &gt;40 y.o.)</li> </ul>	CAJ COLD for general inspect:								
Equipment	<ul style="list-style-type: none"> <li>Gloves → bluey → gel</li> </ul>	<ul style="list-style-type: none"> <li>Cyanosis</li> <li>Anaemia</li> <li>Jaundice</li> <li>Clubbing</li> <li>Oedema</li> <li>Lymph nodes</li> <li>Dehydration</li> </ul>								
Consent & rationale	Explain examination and offer <b>chaperone</b> [clinically trained!] <ul style="list-style-type: none"> <li>"Have you had one done before?" → always address patient anxiety/privacy/past experiences</li> <li>"Undress from waist down, then lie on your left side while bringing your knees up to your chest"</li> </ul> <b>NB:</b> for prostate exam → patient standing and in the bent-over position									
Inspect	Spread buttocks → look for: <ul style="list-style-type: none"> <li><b>Blood</b></li> <li><b>Excoriations - Rash/eczema/pruritus ani</b> (due to faecal soiling)</li> <li><b>Fistula</b> → may occur with Crohn's disease or perianal abscess</li> <li><b>Fissures</b> → IBD, malignancy or STDs</li> <li><b>Anogenital Warts</b> (a.k.a. condylomata acuminata) → HPV infected (needs removal)</li> <li><b>Hemorrhoids</b> (small &lt;1cm tense bluish swellings)</li> <li><b>Skin tags (collagen proliferation)</b> → due to haemorrhoids or Crohn's</li> </ul>	<ul style="list-style-type: none"> <li>Patient Bends down → look for <b>pelvic organ prolapse</b> <ul style="list-style-type: none"> <li>Anterior (cystocele)</li> <li>Middle/Apical (uterine prolapse/ enterocoele)</li> <li>Posterior (rectocele)</li> </ul> </li> </ul>								
Palpate	<ol style="list-style-type: none"> <li>Test anal wink + stroke 4 quadrants of anus with cotton pad → observe brisk anal contraction (if none = spinal cord issue)</li> <li>Excruciating pain → <b>ANAL FISSURE</b> (due to primary or secondary constipation) → abandon exam</li> </ol>									
Exam	<ol style="list-style-type: none"> <li>Lubricate <b>gloved index finger</b> → approach anus posteriorly → pause when over anus → wait until sphincter relaxes</li> <li>Warn patient → advance finger into anus               <ol style="list-style-type: none"> <li>Faeces consistency</li> <li>Ask patient to squeeze finger → Anal tone? [high tone = constipation]</li> <li>Ask patient to bend down → brings high Rectal lesions down</li> </ol> </li> <li>360° sweep for mass/wall thickenings               <ol style="list-style-type: none"> <li>For men: feel 2 lobes of <b>anterior wall of prostate</b> and comment mass, size, symmetry, texture</li> </ol> </li> </ol>	<ol style="list-style-type: none"> <li>Remove finger → wipe on gauze → inspect mucus, blood and melaena               <ol style="list-style-type: none"> <li><b>Careful of false +ve</b> → red meat, aspirin, oral Fe, peroxidase, anticoagulants</li> </ol> </li> <li>Clean anus</li> </ol> <table border="1"> <thead> <tr> <th>Feeling of prostate</th> <th>Disease</th> </tr> </thead> <tbody> <tr> <td>Very hard nodule</td> <td>Carcinoma of the prostate</td> </tr> <tr> <td>Boggy and tender</td> <td>Prostatitis</td> </tr> <tr> <td>Mass above the prostate or cervix</td> <td>Metastatic deposit on blumer's shelf</td> </tr> </tbody> </table>	Feeling of prostate	Disease	Very hard nodule	Carcinoma of the prostate	Boggy and tender	Prostatitis	Mass above the prostate or cervix	Metastatic deposit on blumer's shelf
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To complete	<ol style="list-style-type: none"> <li>Thank + redress + dispose waste + hand hygiene</li> <li>Document findings and name chaperone</li> <li><b>To complete the exam I would perform:</b> <ol style="list-style-type: none"> <li>PSA serology, Rectal US, FBC (anaemia), sigmoidoscopy and colonoscopy</li> </ol> </li> </ol>									



**External haemorrhoids**  
(venous plexus engorgement)



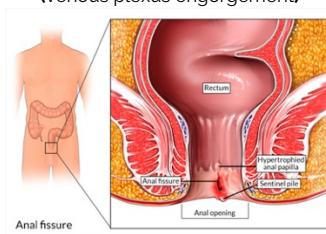
**Skin tags**



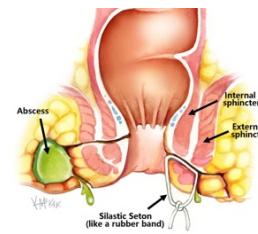
**Anogenital warts (HPV)**



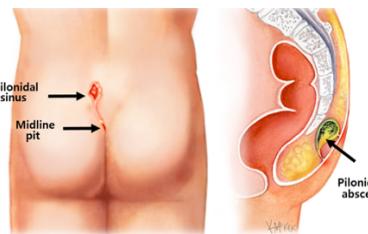
**Rectal Prolapse**



**Anal fissure**  
Rx: fibre, fluid, sitz bath and GTN ointment



**Anal fistula**  
examined under GA,  
MRI- check for fistula and abscess  
Rx: incision and drain + post-op ABx



**Pilonidal sinus**  
(hair from scalp)

### OTHER:

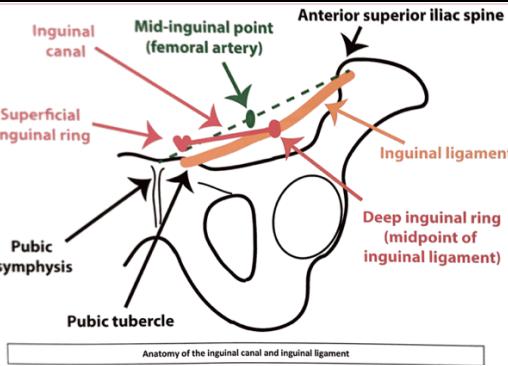
**Fournier's gangrene "wet necrotising fasciitis"**

→ severe pain disproportionate to clinical signs

**Anal Fistula:**  
Goodsall's law  
Park's law

# INGUINAL HERNIA EXAMINATION

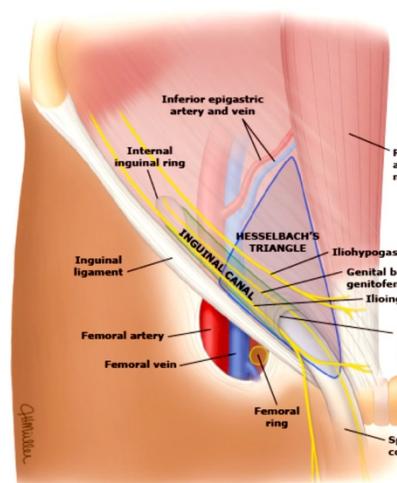
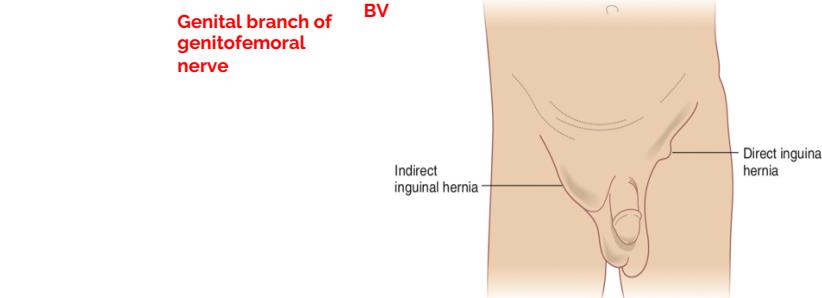
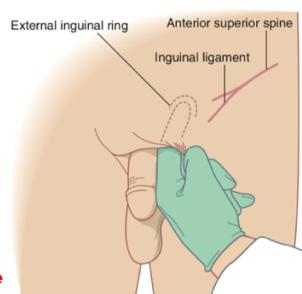
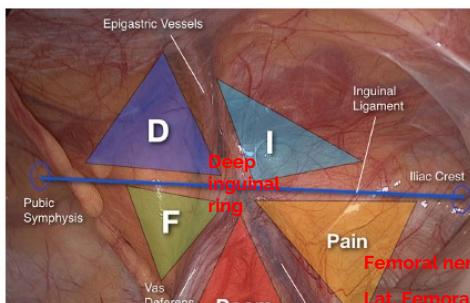
Indication	<ul style="list-style-type: none"> <li>Obvious mass</li> </ul>	CAJ COLD for general inspect: <ul style="list-style-type: none"> <li>Cyanosis</li> <li>Anaemia</li> <li>Jaundice</li> <li>Clubbing</li> <li>Oedema</li> <li>Lymph nodes</li> <li>dehydration</li> </ul>															
Consent & rationale	Explain examination and offer <b>chaperone</b> [clinically trained!]																
Inspect	<ul style="list-style-type: none"> <li><b>General</b> = responsive, pallor. Cachexia, Abd distension (obstruction), vomit bowls</li> <li><b>Lumps</b> = Size, shape, position, scrotal extension</li> <li>Observe cough impulse</li> </ul>																
Palpate	<ul style="list-style-type: none"> <li><b>Scrotal contents</b> → palpate anteriorly. If lump present, you should be able to get above it           <ul style="list-style-type: none"> <li><i>Palpable upper border</i> = <b>hydrocele</b></li> <li><i>Impalpable</i> = <b>inguinoscrotal/indirect inguinal hernia</b></li> </ul> </li> <li><b>Lump/inguinal area (bilateral)</b> → palpate lateral to medial along inguinal ligament with one hand on patient's back           <ul style="list-style-type: none"> <li><b>SITE</b></li> <li><b>SIZE</b></li> <li><b>SHAPE</b></li> <li><b>CONSISTENCY</b></li> <li><b>CONTOURS</b></li> <li><b>COLOUR</b></li> <li><b>TENDER</b></li> <li><b>TEMP</b></li> <li><b>TRANSILLUMINATION</b></li> </ul> </li> <li><b>Feel with cough impulse</b> → compress lump/inguinal area firmly (ask patient to turn away and cough)           <ul style="list-style-type: none"> <li><b>If mass tenses + expands</b> = <b>positive cough impulse</b></li> </ul> </li> <li><b>Reducibility</b> (indirect vs direct)           <ul style="list-style-type: none"> <li>Locate deep inguinal ring (between ASIS + pubic tubercle)</li> <li>Press on lump beginning inferiorly → compress deep inguinal ring</li> <li>Ask patient to cough:               <ul style="list-style-type: none"> <li>Hernia is still absent = <b>indirect hernia</b></li> <li>Hernia reappears = <b>direct hernia</b></li> </ul> </li> <li>Release and watch hernia reappear               <ul style="list-style-type: none"> <li>Hernia slides down <b>obliquely</b> = <b>indirect</b></li> <li>Hernia projects <b>forward</b> = <b>direct</b></li> </ul> </li> </ul> </li> </ul> <p>*If cannot reduce hernia → repeat with patient supine</p>	<b>WHAT ARE HERNIAS?</b> <ul style="list-style-type: none"> <li>Abnormal protrusion of any organ through any fascia or muscle</li> <li>Abdominal, cerebellar, hiatric, lumbar (QL)</li> </ul> <table border="1"> <thead> <tr> <th></th> <th>Indirect inguinal</th> <th>Direct Inguinal</th> </tr> </thead> <tbody> <tr> <td><b>Location</b></td> <td>Between inguinal ring + scrotum</td> <td>Superior to pubic tubercle</td> </tr> <tr> <td><b>Herniated Abd contents</b></td> <td>run within inguinal canal</td> <td>Come out of abd in straight line</td> </tr> <tr> <td><b>Exits</b></td> <td>Deep inguinal ring</td> <td>Superficial ring</td> </tr> <tr> <td><b>Reducibility</b></td> <td>Can be contained</td> <td>Cannot contain (i.e. incarcerated)</td> </tr> </tbody> </table> <p><b>Other hernia types:</b></p> <ol style="list-style-type: none"> <li><b>Inguinal hernia</b> = medial and superior to pubic tubercle</li> <li><b>Femoral hernia</b> = <b>ALWAYS</b> lateral and inferior to pubic tubercle (<b>commonly mistaken for enlarged inguinal LN</b>)</li> <li><b>Epigastric hernia</b> = <i>hernia in epigastric region</i></li> <li><b>Incisional hernia</b> = abd scar causes abdominal weakness</li> <li><b>Umbilical hernia</b> = after pregnancy</li> <li><b>Hiatal hernia</b> = stomach herniates diaphragm</li> </ol> <p><b>RISK FACTORS FOR HERNIA:</b></p> <ol style="list-style-type: none"> <li>++ intra-abdo pressure (obese, pregnant, abscess)</li> <li>weak fascia or muscle (XS manual labour, previous abdo surgery)</li> </ol> <p><b>RED FLAGS:</b></p> <ol style="list-style-type: none"> <li>Inflammation</li> <li>Irreducible</li> <li>Obstructed bowel lumen within hernia</li> <li>Strangulated hernia (vascular compromise of bowel) – esp. femoral hernias</li> </ol>		Indirect inguinal	Direct Inguinal	<b>Location</b>	Between inguinal ring + scrotum	Superior to pubic tubercle	<b>Herniated Abd contents</b>	run within inguinal canal	Come out of abd in straight line	<b>Exits</b>	Deep inguinal ring	Superficial ring	<b>Reducibility</b>	Can be contained	Cannot contain (i.e. incarcerated)
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Percuss & Auscultate	Determine if bowel is present in hernia																
To complete	<b>To complete the exam I would perform:</b>																
	<ul style="list-style-type: none"> <li>Full abdominal exam</li> </ul>																



**DDx: divarication of recti**  
**RF:** Obese, male

**Ix:** see lump when doing crunches

**Rx:** lose weight (not surgery)



## CONTENTS OF INGUINAL CANAL

**MALE:** spermatic cord,

**FEMALE:** round ligament

**BOTH:** ilioinguinal nerve and genital branch of genitofemoral nerve (from deep to superficial)

# Liver Failure & Hepatobiliary Injury

## Anatomy

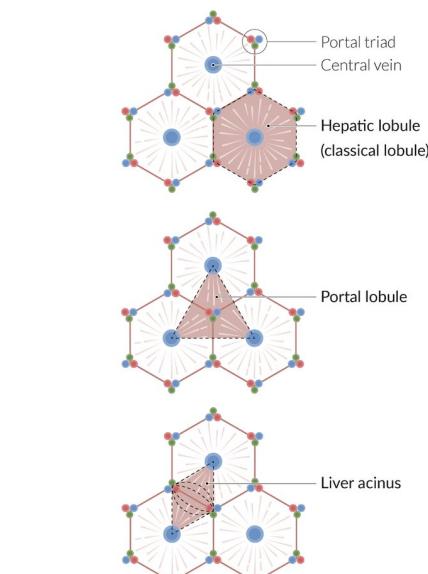
- Portal area:
  1. hepatic artery
  2. Portal vein
  3. Bile ductule
  4. lymphatics
- **Sinusoids** = interaction between blood and hepatocytes via Fenestrated endothelium
- **Hepatocytes** = secrete 500-1200ML bile (i.e. bile salt + bilirubin + cholesterol + water etc.)
  - o Bile → canaliculi → Hepatic ducts → duodenum → **fat digestion**
- **Kupffer cells (macrophages)** = remove cell debris and bacteria from GIT
- **Stellate (Ito cells)** = store Vit A within space of Disse (between sinusoid and hepatocyte) → become myofibroblasts (can develop into cirrhosis)

## Blood supply:

1. Hepatic artery (25%) = oxygen rich
2. Portal vein (75%) = nutrient rich
3. Hepatic veins → IVC

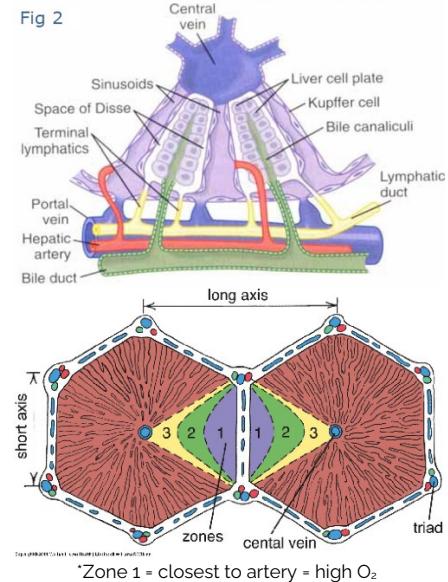
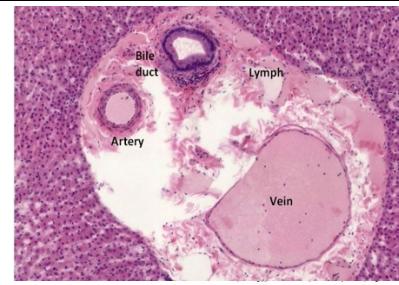
## Hepatic zones:

- **Zone 1 (outer/peripheral)** = high O<sub>2</sub> → GNG,
- **Zone 2 (middle/transitional)**
- **Zone 3 (inner/pericentral)** = low O<sub>2</sub> → anaerobic tasks (e.g. detoxification, glycolysis, lipogenesis etc.) → prone to hypoxia



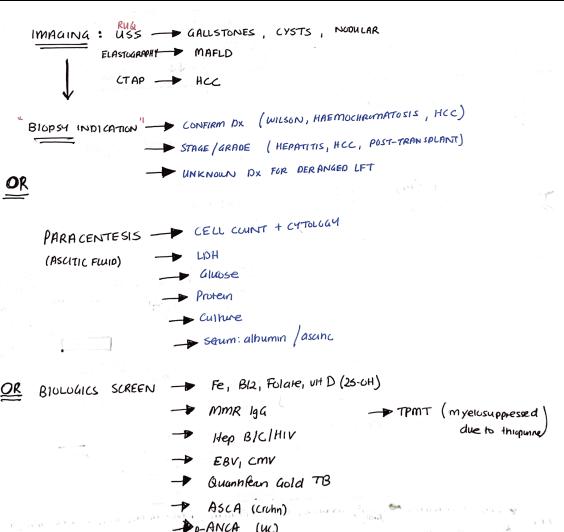
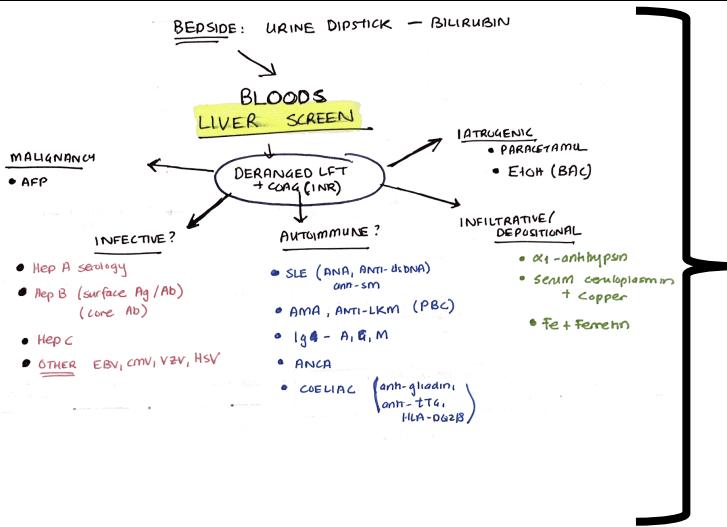
## Main functions of liver:

1. **Bile drainage**
2. **BSL regulator** – produce ketones and stores glycogen
3. Synthesise and store AA, vitamins, and fats
4. **Blood circulation & filtration**
5. **Immunologic** → acute phase proteins (CRP) & **Detoxification**
6. Clotting factor production
7. Albumin production

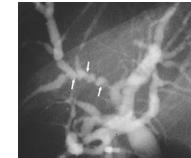


## INTERPRETATION OF LIVER FUNCTION TESTS

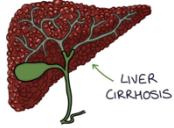
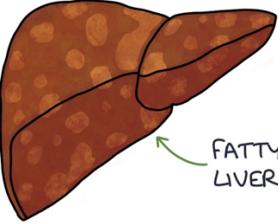
	AST	ALT	ALP	GGT	Cause	Rx
Paracetamol Toxicity	++++	++++	+	+	<ul style="list-style-type: none"> <li>XS acetaminophen saturates conjugation pathway by glutathione → ↑ reactive NAPQI</li> <li>NAPQI = <u>Irreversibly</u> binds to cysteine groups on hepatic macromolecules → oxidative hepatocyte injury</li> <li>Affect <b>Centrilobular region (Zone III)</b> → near central vein</li> <li>DDX: statin, glue sniffing, phenytoin, cocaine, carbamazepine</li> </ul>	<ul style="list-style-type: none"> <li>IV N-Ac within 4hrs (will also Rx baby if pregnant)</li> <li><b>Check for reaction (flush + wheeze)</b></li> <li><b>Cyclizine for nausea (no harm to baby)</b></li> </ul>
Liver shock or ischaemic hepatitis	+++++	+++++	+	+	<ul style="list-style-type: none"> <li>Check Coags</li> <li>Rx cause (Portal vein or hepatic artery thrombosis)</li> </ul>	<ul style="list-style-type: none"> <li><b>HIGHLY</b> vascular liver vulnerable to shock</li> </ul>
Acute hepatitis	+++++	+++++	+	+	<ul style="list-style-type: none"> <li><b>Viral</b> (Hep A/B/E) also CMV, EBV, HSV</li> <li><b>Drugs</b> (paracetamol)</li> <li><b>Fulminant Wilson's</b> → children, teens</li> <li>AIH → females</li> <li>DDX: biliary obstruction? Cholangitis?</li> </ul>	<ul style="list-style-type: none"> <li>ICU admission</li> <li>Invasive vital monitoring</li> <li>Organ replacement therapy</li> <li>Liver transplants</li> </ul>
Chronic hepatitis	+++++	+++++	+	+	<ul style="list-style-type: none"> <li><b>HBV, HCV, HDV</b> → 96% hepatitis death due to sequelae</li> <li><b>Alcoholism</b> → exacerbated by concurrent HBV, HCV infection</li> <li><b>Liver fibrosis</b></li> </ul>	<ul style="list-style-type: none"> <li>"Check <b>Factor V/VII</b> (which affects PT/INR) if fulminant hepatitis → prognostic indicator (determine suitability for liver transplant)</li> </ul>



## RARE HEPATIC PATHOLOGIES

Autoimmune hepatitis		HAEMOCHROMATOSIS	WILSON'S DISEASE	ALPHA 1 ANTITRYPSIN DEF.	PRIMARY BILIARY CIRRHOSIS	PRIMARY SCLEROSING CHOLANGITIS
<b>Def</b>	Rare cause of chronic hepatitis	<ul style="list-style-type: none"> <li>HFE gene mutation (Chr 6)</li> <li>Autosomal recessive</li> </ul>	<ul style="list-style-type: none"> <li>XS copper accumulation due to mutated ATP7B copper-binding protein</li> <li>Autosomal recessive</li> </ul>	<ul style="list-style-type: none"> <li>Mutated gene for protease inhibitor (a1-antitrypsin) → prevents inhibition of neutrophil elastase</li> <li>Autosomal recessive (A1AT gene Chr 14)</li> </ul>	Autoimmune attack on small bile ducts (1 <sup>st</sup> = intralobular ducts – canals of hering)	<ul style="list-style-type: none"> <li><b>Stricture and fibrotic</b> intrahepatic and extrahepatic ducts</li> <li><b>Sclerosis</b> = stiffened and hardened bile ducts</li> </ul>
<b>Sx</b>	<ul style="list-style-type: none"> <li><b>Type 1 (adults)</b> = women 40-50s (menopausal)</li> <li><b>Type 2 (children)</b> – teenagers (acute hepatitis w/ high ALT/AST and jaundice)</li> </ul>	<ul style="list-style-type: none"> <li>Fatigue</li> <li>Arthralgia</li> <li>Bronze diabetes</li> <li>Hair loss</li> <li>ED</li> <li>Amenorrhoea</li> <li>Cognitive (memory and mood swings)</li> </ul>	<p><b>Neuro problem (50%)</b></p> <ul style="list-style-type: none"> <li>Dysarthria</li> <li>Dystonia</li> <li>Basal ganglia (Parkinson's) – asymmetrical motor sx</li> </ul> <p><b>Hepatic problem (40%)</b></p> <ul style="list-style-type: none"> <li>Chronic hepatitis → cirrhosis</li> </ul>	<p><b>Lung issues:</b></p> <ul style="list-style-type: none"> <li>SOB</li> <li>Reduced exercise tolerance</li> </ul> <p><b>Hepatic issues:</b></p> <ul style="list-style-type: none"> <li>Fatigue</li> <li>jaundice</li> </ul>	<ul style="list-style-type: none"> <li>Fatigue</li> <li>Pruritus</li> <li>GI disturbance + abdo pain</li> <li>Jaundice</li> <li>Pale stools</li> <li>Xanthoma &amp; xanthelasma</li> <li>Cirrhosis signs (ascites, +SM, spider naevi)</li> </ul>	<ul style="list-style-type: none"> <li>Jaudnice</li> <li>Chronic RUQ pain</li> <li>Pruritus</li> <li>Fatigue</li> <li>+ HM</li> </ul> 
<b>Comp.</b>	<ul style="list-style-type: none"> <li>Cirrhosis</li> <li>Death</li> </ul> <p><b>Assoc.</b></p> <ul style="list-style-type: none"> <li>Arthritis</li> <li>Hashimoto's</li> </ul>	<ul style="list-style-type: none"> <li><b>T1DM</b> (pancreatitis)</li> <li><b>Liver cirrhosis</b></li> <li><b>Pituitary &amp; gonad infiltration</b> (hypogonad, impotence, amenorrhea, infertility)</li> <li><b>Cardiomyopathy</b></li> <li><b>HCC</b></li> <li><b>Hypothyroidism</b></li> <li><b>Pseudogout</b> &amp; Chondrocalcinosis</li> </ul>	<p><b>Psych Problem (10%)</b></p> <ul style="list-style-type: none"> <li>Depression → full psychosis</li> </ul> <p><b>Other:</b></p> <ul style="list-style-type: none"> <li>Kayser-Fleischer rings</li> <li>Haemolytic anaemia</li> <li>Renal tubular acidosis</li> <li>Osteopenia</li> </ul>	<p><b>2 main complications:</b></p> <ol style="list-style-type: none"> <li>Bronchiectasis and emphysema (&gt;50yo)</li> <li>Liver cirrhosis (after 50yo) leading to HCC</li> </ol>	<p>Inflamed bile ducts causes obstruction of bile outflow = cholestasis</p> <ol style="list-style-type: none"> <li>Fibrosis</li> <li>Advanced liver Cirrhosis</li> <li>Portal HTN and liver failure <ul style="list-style-type: none"> <li>Symptomatic pruritus</li> <li>Steatorrhoea (greasy stools)</li> <li>Hypothyroidism</li> <li>OP</li> <li>HCC</li> </ul> </li> </ol>	<p>Chronic bile obstruction leads to:</p> <ol style="list-style-type: none"> <li>Fibrosis</li> <li>Advanced liver Cirrhosis</li> <li>Portal HTN and liver failure</li> </ol> <p><b>SPECIFIC</b></p> <ul style="list-style-type: none"> <li>Acute bacterial cholangitis</li> <li>Cholangiosarcoma</li> <li>CRC</li> <li>Biliary stricture</li> <li>Fat soluble vit. Def (A,D,E,K)</li> </ul>
<b>Ix</b>	<ul style="list-style-type: none"> <li>Deranged LFT</li> <li>Elevated IgG</li> </ul> <p><b>Type 1 Autoantibodies: (ADULTS)</b></p> <ul style="list-style-type: none"> <li>ANA</li> <li>Anti-SM</li> <li>Anti-SLA/LP</li> </ul> <p><b>Type 2 Autoantibodies: (CHILDREN)</b></p> <ul style="list-style-type: none"> <li>anti-LKM1</li> <li>anti-LC1</li> </ul> <p><b>liver Biopsy</b> to confirm Dx</p>	<ul style="list-style-type: none"> <li><b>Fe studies</b> (raised serum ferritin &amp; transferrin)</li> <li><b>Liver biopsy</b> – Perl's stain → Fe concentrated in parenchymal cells</li> <li><b>Genetic testing (H63D, c282y)</b></li> <li><b>CTAP</b></li> <li><b>MRI</b> – Fe deposition on liver</li> </ul>	<ul style="list-style-type: none"> <li><b>REDUCED</b> Serum ceruloplasmin (not specific)</li> <li><b>REDUCED serum Cu</b></li> <li><b>RAISED</b> 24 hr urinary Cu assay</li> <li><b>Liver biopsy (gold standard)</b></li> </ul> 	<ul style="list-style-type: none"> <li><b>REDUCED</b> serum alpha-1 antitrypsin</li> <li><b>Liver biopsy (gold standard)</b> – acid-Schiff +ve staining globules</li> <li>Genetic testing (A1AT gene)</li> <li><b>HRCT thorax</b> → diagnose bronchiectasis and emphysema</li> </ul>	<p><b>Bloods:</b></p> <ul style="list-style-type: none"> <li>Raised ESR</li> <li>Raised IgM</li> </ul> <p><b>Deranged LFT:</b></p> <ul style="list-style-type: none"> <li>Raised ALP (1<sup>st</sup>)</li> <li>ALT/AST raised later</li> <li>(AST:ALT &gt;1 = CIRRHOSIS)</li> </ul> <p><b>Autoantibodies:</b></p> <ul style="list-style-type: none"> <li>AMA (most specific to PBC)</li> <li>ANA (present in 35% of patients)</li> </ul>	<p><b>Deranged LFT:</b></p> <ul style="list-style-type: none"> <li>Cholestatic picture</li> <li>Elevated ALT/AST if progression to cirrhosis</li> </ul> <p><b>Autoantibodies (not useful)</b></p> <ul style="list-style-type: none"> <li>P-ANCA (94%)</li> <li>ANA (77% of patients)</li> <li>Anti-cardiolipin (63%)</li> </ul> <p><b>MRCP (gold standard)</b></p> <ul style="list-style-type: none"> <li>MRI of liver, bile duct and pancreas → lesions/strictures</li> <li><b>Beaded pattern stricture</b></li> </ul>
<b>AST</b>	++++ (2x ALT)	NOT HELPFUL	++++ (2x ALT)	NOT HELPFUL	+	NOT HELPFUL
<b>ALT</b>	++	NOT HELPFUL	++	NOT HELPFUL	+	NOT HELPFUL
<b>ALP</b>	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	++++	++++
<b>GGT</b>	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	++++	++++
<b>Mx</b>	<ol style="list-style-type: none"> <li>High dose steroids (pred)</li> <li>Azathioprine (life-long)</li> <li>Liver transplant (may recur)</li> </ol>	<ol style="list-style-type: none"> <li>Venesection (weekly)</li> <li>Monitor serum ferritin</li> <li>Avoid alcohol</li> <li>Genetic counsel (family)</li> <li>Rx complications</li> </ol>	<p>Cu chelation using:</p> <ol style="list-style-type: none"> <li>Penicillamine</li> <li>Trientine</li> </ol>	<ul style="list-style-type: none"> <li>Stop smoking</li> <li>Symptomatic Mx</li> <li>Lung and liver transplant</li> <li>Monitor for complications (e.g. HCC)</li> </ul>	<ol style="list-style-type: none"> <li><b>Ursodeoxycholic acid</b> (reduced intestinal absorption of cholesterol)</li> <li><b>Cholestyramine</b> (binds to bile acids to prevent gut absorption – improve pruritus)</li> <li><b>Immunosuppression</b> (steroids)</li> <li><b>Liver transplant</b> (end-stage)</li> </ol>	<ul style="list-style-type: none"> <li>ERCP – dilate + stent strictures (XR guided + contrasts used)</li> <li><b>Cholestyramine</b> (binds to bile acids to prevent gut absorption – improve pruritus)</li> <li><b>Monitor comp.</b> (cholangiosarcoma, oesophageal varices)</li> <li><b>Liver transplant</b> (curative intent – 80% 5-year survival)</li> </ul>

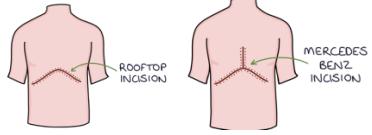
# COMMON LIVER DISEASE – NAFLD, ALCOHOLIC, CIRRHOsis

	MAFLD / NAFLD	ALCOHOLIC LIVER DISEASE	LIVER CIRRHOsis																																
Risk factors	<ul style="list-style-type: none"> <li>30% of adults</li> <li>Part of metabolic syndrome</li> </ul> <p><b>Risk factors</b></p> <ul style="list-style-type: none"> <li>Obese – poor diet low PA</li> <li>MetSyn (HTN, HC, T2DM)</li> <li>Smoker</li> <li>Middle-aged</li> </ul>	<ul style="list-style-type: none"> <li>Long-term EtOH consumption</li> <li>Genetic predisposition</li> <li><b>80% alcohol</b> → metabolised by Alcohol dehydrogenase into <b>Acetaldehyde</b> → forms <b>adducts with proteins in hepatocytes</b> → <b>activate immune response = cell injury</b></li> <li><b>20% alcohol</b> → metabolised by Cytochrome CYP2E1 into acetate releases <b>ROS</b> → <b>lipid peroxidation and mitochondrial damage</b></li> <li><b>increased gut permeability</b> → <b>Endotoxin release</b> → <b>inflammatory cytokines</b> → <b>accelerated damage</b></li> </ul>	<ul style="list-style-type: none"> <li>Chronic inflammation and damage to liver cells</li> <li>Damaged cells replaced by scar tissue (fibrosis) and nodules</li> </ul> <p><b>Common causes:</b></p> <ul style="list-style-type: none"> <li>Alcoholic liver disease</li> <li>NAFLD</li> <li>Hep B/C</li> </ul> 																																
Comp.	<p><b>Step wise progression:</b></p> <ul style="list-style-type: none"> <li><b>Non-alcoholic fatty liver disease</b></li> <li><b>Non-alcoholic steatohepatitis (NASH)</b></li> <li>Fibrosis</li> <li>Cirrhosis</li> </ul> <p><b>Increased risk of:</b></p> <ol style="list-style-type: none"> <li>1) heart disease</li> <li>2) stroke</li> <li>3) diabetes mellitus</li> </ol> 	<p><b>Step wise progression:</b></p> <ol style="list-style-type: none"> <li><b>EtOH-related fatty liver</b> (reversible within 2 weeks of stopping alcohol)</li> <li><b>Alcoholic hepatitis</b> (reversible with permanent abstinence)</li> <li><b>Cirrhosis</b> (shrunken liver w/ scar tissue – irreversible)</li> </ol> <p><b>SPECIFIC COMPLICATIONS</b></p> <ul style="list-style-type: none"> <li><b>B1 deficiency = Wernicke-Korsakoff syndrome</b> <ol style="list-style-type: none"> <li>EtOH prevents absorption of B1</li> <li><b>Wernicke enceph (high mortality rate)</b> = confusion, oculomotor disturbance, ataxia</li> <li><b>Korsakoff syndrome (irreversible)</b> = retrograde and anterograde amnesia, confabulation, behaviour change</li> </ol> </li> <li><b>Alcohol withdrawal</b> <ol style="list-style-type: none"> <li>6-12 hrs = tremor, sweat, craving</li> <li>12-24 hrs = hallucinations</li> <li>24-48 hrs = seizures</li> <li>48-72 hrs = <b>delirium tremens (Emergency!!)</b> → downregulation of GABA system by chronic EtOH consumption causes upregulated glutamate → acute confusion, agitation, delusions, tremor, tachycardia, HTN, hyperthermia, ataxia, arrhythmias</li> </ol> </li> <li><b>Other complications:</b> <ol style="list-style-type: none"> <li>Pancreatitis</li> <li>Alcoholic cardiomyopathy</li> <li>Alcohol dependence</li> </ol> </li> </ul>	<p><b>Fibrotic liver increase resistance in hepatic vessels causes:</b></p> <ul style="list-style-type: none"> <li>Portal HTN → variceal bleeding, ascites → SBP</li> <li>Liver dysfn (malnutrition + encephalopathy)</li> <li>Hepato-renal syndrome</li> <li>HCC</li> </ul> <p><b>Child-Pugh score [AA-BEC] – severity of cirrhosis</b></p> <p><b>Factors:</b> Ascites, encephalopathy, bilirubin, albumin, coagulopathy</p> <table border="1"> <thead> <tr> <th>Points</th> <th>1</th> <th>2</th> <th>3</th> </tr> </thead> <tbody> <tr> <td>Encephalopathy</td> <td>None</td> <td>Mild</td> <td>Mod-severe</td> </tr> <tr> <td>Ascites</td> <td>Absent</td> <td>Controlled</td> <td>Refractory</td> </tr> <tr> <td>Total bilirubin</td> <td>&lt;34 (&lt;2)</td> <td>34-50 (2-3)</td> <td>&gt;50 (&gt;3)</td> </tr> <tr> <td>Albumin (g/dL)</td> <td>&gt;3.5</td> <td>2.8-3.5</td> <td>&lt;2.8</td> </tr> <tr> <td>PT time (sec)</td> <td>&lt;4 or &lt;1.7</td> <td>4-6 or 1.7-2.3</td> <td>&gt;6 or &gt;2.3</td> </tr> <tr> <td>Score</td> <td>A</td> <td>B</td> <td>C</td> </tr> <tr> <td>5-yr survival</td> <td>95%</td> <td>75%</td> <td>50%</td> </tr> </tbody> </table> <p><b>MELD score [BIC-Na] – use very 6/12 for comp. cirrhosis</b></p> <p><b>Factors:</b> Bilirubin, INR, creatinine, Na + need for dialysis</p> <p><b>MELD Score</b> = <math>10 \times (0.957 \times \ln(\text{Creatinine})) + (0.378 \times \ln(\text{Bilirubin})) + (1.12 \times \ln(\text{INR})) + 6.43</math></p> <ul style="list-style-type: none"> <li><b>Assess severity of CLD (3/12 mortality %) to determine need for liver transplant</b> <ul style="list-style-type: none"> <li>&gt; 25 (gravely ill)</li> <li>24 to 19 = difficult transplant</li> <li>18 to 11 = candidate</li> <li>&lt; 10 (less ill)</li> </ul> </li> </ul>	Points	1	2	3	Encephalopathy	None	Mild	Mod-severe	Ascites	Absent	Controlled	Refractory	Total bilirubin	<34 (<2)	34-50 (2-3)	>50 (>3)	Albumin (g/dL)	>3.5	2.8-3.5	<2.8	PT time (sec)	<4 or <1.7	4-6 or 1.7-2.3	>6 or >2.3	Score	A	B	C	5-yr survival	95%	75%	50%
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Sx	Asymptomatic	<ul style="list-style-type: none"> <li><b>Jaundice</b> = raised bilirubin</li> <li>+HM = (but may shrink if more cirrhotic)</li> <li>+SM = due to portal HTN → hypersplenism (↓plt)</li> <li><b>Spider naevi</b> – telangiectasia w/ central arteriole and small vessels radiating away</li> <li><b>Palmar erythema</b> = hyperdynamic circulation (↑E2)</li> <li><b>Bruising</b> = abnormal clotting</li> </ul>	<ul style="list-style-type: none"> <li><b>Gynecomastia and testicular atrophy in males</b> = endocrine dysfunction</li> <li><b>Ascites</b> = portal HTN</li> <li><b>Caput medusae</b> = distended periumbilical veins due to portal HTN</li> <li><b>Asterixis "flapping tremor"</b> – decomp. Liver disease</li> </ul>																																
ix	<ul style="list-style-type: none"> <li>&gt; <b>LFT = + ALT/AST</b></li> <li>&gt; <b>COAGS = raised PT/INR time</b></li> <li>&gt; <b>Hep B/C serology</b></li> <li>&gt; <b>Autoantibodies</b> – ANA, SMA, AMA, LKM-1 (autoimmune hepatitis, PBC, PSC)</li> <li>&gt; <b>Immunoglobulins</b> (autoimmune hepatitis, PBC)</li> <li>&gt; <b>Caeruloplasmin</b> (Wilson's)</li> <li>&gt; <b>A1-antitrypsin levels</b></li> <li>&gt; <b>Fe studies</b> (hereditary haem)</li> <li>&gt; <b>Liver USS /</b> = increased echogenicity to confirm fatty liver</li> <li>&gt; <b>NAFLD fibrosis score</b> (algorithm that uses age, BMI, LFTs to rule out fibrosis)</li> <li>&gt; <b>Fibroscan</b> (indication of fibrosis)</li> </ul>	<ul style="list-style-type: none"> <li><b>FBC</b> = raised MCV</li> <li><b>EUC</b> = deranged (hepatorenal syndrome)</li> <li>&gt; <b>LFT = + ALT/AST, +++GGT</b> <ul style="list-style-type: none"> <li>AST:ALT ratio &gt; 2</li> </ul> </li> <li><b>COAGS</b> = raised PT/INR time</li> <li><b>Liver USS / fibroscan</b> = increased echogenicity</li> <li><b>Endoscopy</b> = assess and treat oesophageal varices</li> <li><b>CT/MRI</b> = fatty infiltration of liver, HCC, +HSM, ascites, abnormal BV changes</li> <li><b>Liver biopsy</b> – is it alcoholic hepatitis or cirrhosis</li> </ul>	<ul style="list-style-type: none"> <li><b>EUC</b> = hypoNa (fluid retention due to severe liver disease)</li> <li>&gt; <b>LFT</b> = often normal (until decompensated cirrhosis)</li> <li>&gt; <b>AFP</b> = MARKER FOR HCC</li> <li><b>Liver USS - nodular liver surface</b> + <ul style="list-style-type: none"> <li>"corkscrew" arteries w/ increased blood flow to compensate for ↓flow</li> <li>enlarged portal vein</li> <li>ascites</li> <li>splenomegaly</li> </ul> </li> <li><b>Liver fibroscan (elasticity of liver)</b></li> <li><b>Endoscopy</b> = assess and treat oesophageal varices</li> <li><b>CT/MRI</b> = fatty infiltration of liver, HCC, +HSM, ascites,</li> <li><b>Liver biopsy</b> – is it alcoholic hepatitis or cirrhosis</li> </ul>																																
ALT/AST	Mildly elevated	AST:ALT ratio > 2																																	
ALP/GGT	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL																																
Mx	<ul style="list-style-type: none"> <li>Weight loss</li> <li>Exercise</li> <li>Stop smoking</li> <li>Stop alcohol consumption</li> <li>T2DM, HC, HTN control</li> </ul>	<ul style="list-style-type: none"> <li>Stop EtOH immediately and permanently</li> <li>Detoxification regime</li> <li><b>Nutritional support</b> (B1 supp) + high protein diet</li> <li><b>Manage withdrawal Sx (delirium tremens)</b> <ul style="list-style-type: none"> <li>Benzos = Clordiazepoxide "librium" PO every 1-4 hrs for 5-7 days</li> <li>IV high-dose B vitamins (pabrinex)</li> <li>Oral thiamine</li> </ul> </li> </ul>	<p><b>General Management</b></p> <ul style="list-style-type: none"> <li>High protein, low sodium diet</li> <li>MELD score every 6 months</li> <li>Liver USS and AFP levels every 6 months for hepatocellular carcinoma</li> <li>Endoscopy every 3 years in patients <u>without</u> known varices</li> <li><b>Consideration of a liver transplant</b></li> <li>Managing complications as below</li> </ul>																																

# MANAGEMENT OF CIRRHOSIS / CHRONIC LIVER DISEASE

Complication	Description	Management									
<b>Malnutrition + muscle wasting</b>	Liver dysfn = affects protein metabolism and stop glucose storage	<ul style="list-style-type: none"> <li>Low Salt diet (&lt;2 g)</li> <li>High protein diet → regular meals</li> <li>Avoid alcohol</li> </ul>									
<b>Ascites</b>	<ul style="list-style-type: none"> <li>Fluid accumulation in peritoneal cavity ← hypoalbuminemia + portal HTN</li> <li>Fluid wave sound</li> </ul> <table border="1"> <thead> <tr> <th></th> <th>Type</th> <th>Causes</th> </tr> </thead> <tbody> <tr> <td><b>SAAG &gt; 1.1</b></td> <td>Portal HTN (transudate)</td> <td>CCF, Cirrhosis, hepatitis budd-chiari, HCC, liver mets,</td> </tr> <tr> <td><b>SAAG &lt; 1.1</b></td> <td>Exudate</td> <td>SBP, TB, peritonitis, SBO, nephrotic syndrome, pancreas- biliary issue, ovarian cancer</td> </tr> </tbody> </table>		Type	Causes	<b>SAAG &gt; 1.1</b>	Portal HTN (transudate)	CCF, Cirrhosis, hepatitis budd-chiari, HCC, liver mets,	<b>SAAG &lt; 1.1</b>	Exudate	SBP, TB, peritonitis, SBO, nephrotic syndrome, pancreas- biliary issue, ovarian cancer	<ul style="list-style-type: none"> <li>Low Salt diet (&lt;2 g)</li> <li>Diuretics (spironolactone - K+ sparing → block RAAS)</li> <li>Paracentesis (ascitic tap)</li> <li>Prophylactic Abx (quinolones) protect against SBP if &lt; 15g/L of protein in ascitic fluid</li> <li>TIPS or transplant in <u>refractory ascites</u></li> </ul>
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<b>SAAG &gt; 1.1</b>	Portal HTN (transudate)	CCF, Cirrhosis, hepatitis budd-chiari, HCC, liver mets,									
<b>SAAG &lt; 1.1</b>	Exudate	SBP, TB, peritonitis, SBO, nephrotic syndrome, pancreas- biliary issue, ovarian cancer									
<b>Spontaneous Bacterial Peritonitis (SBP) = infection</b>	<ul style="list-style-type: none"> <li><b>Infected ascitic fluid</b> (only in cirrhosis) usu. <b>E.coli</b> cause (also Klebsiella, staph, enterococcus)</li> <li>Abd pain &amp; distension</li> <li><b>Fever (specific sx)</b></li> <li>Confusion</li> <li>Ileus</li> <li>HypoTN</li> <li>Deranged bloods (raised WBC, CRP, creatinine, met. Acidosis)</li> </ul>	<ol style="list-style-type: none"> <li>Paracentesis → Ascitic tap:             <ol style="list-style-type: none"> <li>elevated PMN &gt; 250</li> <li>SAAG &lt; 1.1 (exudate) = NOT portal HTN</li> <li>+ve bacteria culture</li> </ol> </li> <li>IV empirical Abx (cefotaxime) or oral quinolones</li> </ol>									
<b>Variceal Haemorrhage</b>	<ul style="list-style-type: none"> <li><b>Rule out first</b> → check Hb (FBC) and urea             <ul style="list-style-type: none"> <li>↓ Hb and ↑ Urea</li> </ul> </li> <li><b>Portal HTN → swollen tortuous vessels (varices)</b> <ul style="list-style-type: none"> <li>Can bleed out very quickly due to high blood flow through varices</li> </ul> </li> <li><b>Occurs at:</b> <ul style="list-style-type: none"> <li>Gastro-oesophageal junction</li> <li>Ileocecal junction</li> <li>Rectum</li> <li>Anterior abdominal wall via umbilical vein (caput medusae)</li> </ul> </li> </ul> 	<p><b>Acute (bleeding varices)</b></p> <ol style="list-style-type: none"> <li>Resus</li> <li>IV prophylactic empirical Abx (cephalosporin)</li> <li>urgent upper GI endoscopy – inject sclerosant or band ligation</li> <li>Sengstaken-Blakemore tube (tamponade bleeding varices) if endoscopy fails</li> </ol> <p><b>Stable varices → Prevention:</b></p> <ul style="list-style-type: none"> <li>Surveillance (upper GI endoscopy -2 years)</li> <li><b>Non-selective BB (propranolol)</b> → reduce variceal bleeding risk</li> <li><b>Elastic band ligation of varices</b></li> <li><b>TIPS</b> – connect hepatic vein with portal vein to reduce pressure through portal system</li> </ul>									
<b>Hepatic / portosystemic Encephalopathy</b>	<p><b>Triggers:</b></p> <ul style="list-style-type: none"> <li>Constipation</li> <li>Electrolyte disturbance</li> <li>Infection (SBP)</li> <li>GIB</li> <li>High protein diet</li> <li>Medications (sedative meds)</li> </ul> <p><b>Reversible neuropsychiatric disturbances:</b></p> <ul style="list-style-type: none"> <li>Acute = NO fever, confusion</li> <li>Chronic = asterixis, personality change, Sleep wake inversion</li> </ul>	<ul style="list-style-type: none"> <li><b>High dose Laxatives (oral lactulose)</b> <ul style="list-style-type: none"> <li>promote excretion of ammonia</li> <li>2-3x soft motions a day</li> </ul> </li> <li><b>Abx (rifaximin)</b> → reduce number of intestinal bacteria producing anaemia</li> <li><b>Nutritional support</b> → NGT feeding</li> <li><b>Identify + Rx cause</b> → infection, sedative, variceal haem</li> </ul>									
<b>Hepatocellular Carcinoma (HCC)</b>	Higher risk in cirrhosis <ul style="list-style-type: none"> <li>chronic Hep B/C infection (oncogenic virus)</li> </ul>	Rx: Hep B/C to reduce risk of malignancy <ul style="list-style-type: none"> <li><b>Every 6/12: Liver USS, check AFP levels</b></li> </ul>									
<b>Hepatorenal Syndrome</b>	Liver cirrhosis → portal HTN → reduced blood vol. through kidneys Activation of RAAS → renal vasoconstriction → worsens renal perfusion	<ul style="list-style-type: none"> <li>Fatal within a week</li> <li>Liver transplant</li> </ul>									

## LIVER CANCER AND LIVER TRANSPLANTS

LIVER CANCER (2 types)		LIVER TRANSPLANT
<b>TYPES</b>	<p><b>Benign</b></p> <ul style="list-style-type: none"> <li>Haemangiomas (incidental finding – no Rx needed)</li> <li>Focal nodular hyperplasia (fibrotic tissue) – more common in women and those on OCP (estrogen related - no Rx needed)</li> </ul> <p><b>Malignant</b></p> <ul style="list-style-type: none"> <li>Primary = HCC (80%) OR cholangiocarcinoma (20%)</li> <li>Secondary (from breast, skin, bowel or unknown primary)</li> </ul>	<p><b>Indications:</b></p> <ol style="list-style-type: none"> <li><b>Acute liver failure (immediate transplant)</b> → acute viral hepatitis or paracetamol OD</li> <li><b>Chronic liver failure</b> - may need to wait for 5/12</li> </ol> <p><b>Factors affecting unsuitability</b></p> <ul style="list-style-type: none"> <li>Significant co-morbidities (CKD, HD)</li> <li>XS weight loss + malnutrition</li> <li>Acute hep B/C</li> <li>End-stage HIV</li> <li>Active alcohol use (need 6/12 abstience)</li> </ul> <p><b>Surgery:</b></p> <ul style="list-style-type: none"> <li>Living Donor Transplant – Split Donation</li> <li>"Rooftop" or "Mercedes Benz" incision</li> </ul> 
<b>Risk factors</b>	<ul style="list-style-type: none"> <li>Viral HEPATITIS (B/C)</li> <li>Alcohol</li> <li>NAFLD</li> <li>CHRONIC LIVER DISEASE</li> </ul>	
<b>Sx</b>	<p>Asymptomatic +/-</p> <ul style="list-style-type: none"> <li><b>NON-specific sx</b> (UWL, abdo pain, anorexia, N/V, &gt; Specific sx: jaundice and pruritus)</li> </ul>	
<b>Ix</b>	<ul style="list-style-type: none"> <li><b>AFP (HCC)</b></li> <li>CA 19.9 (cholangiocarcinoma)</li> <li>Liver USS (identify tumour)</li> <li>CT + MRI (stage and diagnose cancer)</li> <li>ERCP (biopsy to diagnose cholangiocarcinoma)</li> </ul>	
<b>Mx</b>	<p><b>HCC &amp; Cholangiocarcinoma</b></p> <ul style="list-style-type: none"> <li>Both have Poor prognosis (unless diagnosed early)</li> <li>Both unresponsive to chemo and RT</li> <li>Resection cw. Curative intent</li> </ul> <p><b>SPECIFIC Rx:</b></p> <ul style="list-style-type: none"> <li>HCC → TKI (e.g. sorafenib, lenvatinib) → extend life by months</li> <li>Cholangiocarcinoma → ERCP – stent insertion in bile duct to drain bile</li> </ul>	<p><b>Post-transplantation care:</b></p> <ul style="list-style-type: none"> <li><b>Life-long immunosuppression</b> (e.g. steroids, azathioprine, tacrolimus)</li> <li><b>Avoid alcohol, smoking</b></li> <li><b>Treat opportunistic infection</b></li> <li><b>Monitor disease recurrence</b> (E.g. hepatitis, PBC)</li> <li><b>Monitor cancer recurrence</b> (higher risk in immunosuppressed patients)</li> </ul>

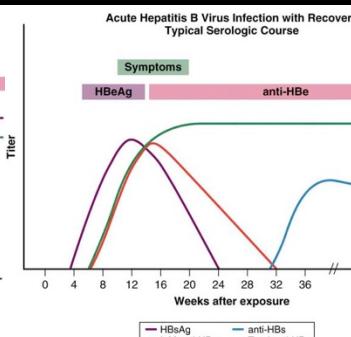
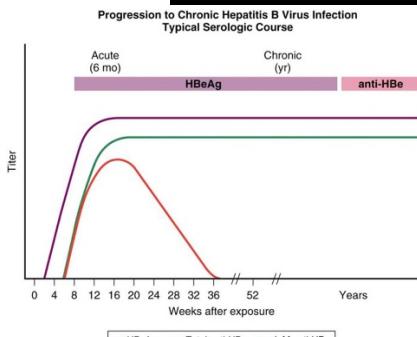
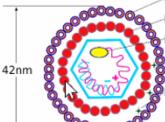
# Viral Hepatitis

► PRACTICE POINT = EBV produces same results +splenomegaly

	HAV (Picorn)	HBV (hepada-)		HCV (flavivirus)	HDV (delta)	HEV (hepe-)
Type	ssRNA	dsDNA (immune mediated killing of infected cells)		RNA (viral pol - no proof reading = higher mutations)	RNA low dose	RNA (unenveloped)
Epi	Poverty areas Most common acute viral hepatitis in children	Most common liver infection (esp. Western Pacific/Africa & ATSI)		Good prognosis = young and females Bad prognosis = male, HIV, older, EtOH usage ¾ of patients are asymptomatic	Co-infect (acute) HB/HDV = 90% recovery Superinfect Chronic hep B + acute hep D → 90% chronic	Most common acute hep in Asia, ME, North Africa Aus: from pigs (undercooked)
Tx	Faecal/oral (frozen berries, seawater) Person-to person	Vertical/Perinatal (mother-child) Horizontal (infants, contacts) Bloodborne Sex (semen)		Perinatal (mother-child) Bloodborne (IVDU, needlestick injury) Sex (semen)	Bloodborne	Oral/faecal Water/food
Acute infection	Case fatality increases with age IgM positive	Mortality increases w/ age 95% (adults) Asympt Acute viral Hep ALF	Uncommon - slowly progressive hepatitis		fulminant disease (10x more common)	High mortality = fulminant in pregnant women (esp. after recent monsoon) IgM positive
Chronic infection	No	90% (children) Chronically infected → chronic Hep B → compensated cirrhosis → decompensated cirrhosis	75% → precursor to HCC, oesophageal ca. & mixed cryoglobulinemia, porphyria cutanea tarda (rash on sun-exposed area)		Worsen hepatitis B → fulminant cirrhosis, death > mono- HBV infection	Very rare (but possible)
Comp.*	Cholestasis	HCC - Leading cause (50%) Polyarteritis Nodosa (PAN)	HCC, cirrhosis Sicca Cryoglobulinemia (palpable purpura) GN Thyroiditis			No
Serology	IgM = acute → RNA PCR IgG = chronic (protected)	PCR TESTING - VIRAL DNA HBcAB (IgM) = acute HBcAB (IgG) = previous infect Anti-HbsAg = vaccinee/protect	PCR TESTING - VIRAL RNA [seroconvert within 1 <sup>st</sup> month]		Anti-HDV IgM/IgG (past/current infection)	Elevated ALT Anti-IgM for HepE Viral Load Test
Vaccine	Yes	Yes (Essential to prevent chronic HBV infection esp. after childhood infection)	No		HBV vaccine	No
Rx for acute	None (usu. self limiting within 3/12) analgesia	(supportive care - recover within 2/12) Hep B Ig → PEP for hep B pts without 3-dose vaccination or did not properly respond to it	Available (pegIFN-alfa & ribavirin) NOT protected from re-infection		New agents?	None (lifelong immunity)
Rx for chronic		Rx • Lifetime Nucleoside/Nucleotide Analogues (e.g. Entecavir) • Goals: ↓↓ HBV DNA load + normalise ALT • Cure if HBsAg <5% of patients  If pregnant + HBV positive: ➢ Low load = vaccinate baby ➢ High load = tenofovir + hep B Ig for baby	For chronic Hepatitis C • Direct Acting Antivirals (DAs) e.g. sofosbuvir and ledipasvir (CI if cirrhotic) Elbasivir (newer more effective but more expensive) Cure = sustained virologic response • Goals: undetectable HCV load ≥ 12 weeks after completion of therapy			
Public health	Must notify	Must notify	Must notify		Must notify	Must notify

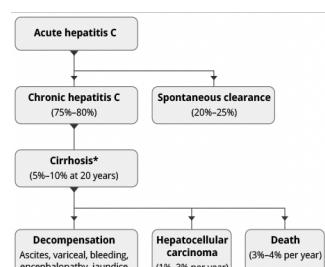
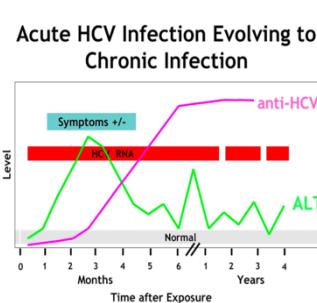
## HEP B SEROLOGY

HBsAg or HBV DNA	HBcAb (IgM)	HBcAb (IgG)	HBeAg	HbsAb	Interpretation
					Susceptible to HBV infection
+	+		+		Acute
+			+		Early acute
		+	+		Chronic
				+	Resolved acute
					Vaccinated

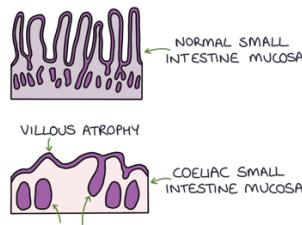


## HEP C SEROLOGY

	HCV RNA	IgM anti-HCV	IgG Anti-HCV
Acute	+	+	+
Chronic	+	-	+
Cleared	-	-	+



# COMMON GI ISSUES – GORD, PUD, UGIB and coeliac disease

	GORD	PEPTIC/DUODENAL ULCERS	COELIAC DISEASE	IRRITABLE BOWEL SYNDROME
Def	<ul style="list-style-type: none"> <li>Stomach acid refluxes through LES and irritates oesophageal lining</li> </ul>	<ul style="list-style-type: none"> <li>Ulcerated mucosa of stomach or duodenum</li> <li>Duodenal ulcers more common</li> </ul> <p>(1) <b>Breakdown protective mucosa:</b></p> <ul style="list-style-type: none"> <li>Chronic NSAID/steroids</li> <li>H. pylori</li> </ul> <p>(2) <b>Increase Acid secretion:</b></p> <ul style="list-style-type: none"> <li>Stress</li> <li>Spicy food</li> <li>Smoking + EtOH</li> <li>Caffeine</li> </ul>	<ul style="list-style-type: none"> <li>Autoimmune reaction to ingestion of gluten</li> <li>Inflamed small bowel</li> <li>Begins at any age</li> </ul> <p>Assoc. autoimmune conditions:</p> <ul style="list-style-type: none"> <li>T1DM</li> <li>Pernicious anemia</li> <li>Thyroid disease</li> <li>PBC, PSC</li> <li>Autoimmune hepatitis</li> </ul>	<ul style="list-style-type: none"> <li>Functional bowel disorder</li> <li>Diagnosis of exclusion</li> <li>20% of population</li> <li>Women &gt; men</li> <li>Young adults</li> </ul>
Comp.	<p><b>Barrett's oesophagus</b> (metaplasia from squamous to columnar) → premalignant to develop AC of oesophagus</p> <ol style="list-style-type: none"> <li>Low grade dysplasia</li> <li>High grade dysplasia</li> <li>Adenocarcinoma</li> </ol>	<ul style="list-style-type: none"> <li><b>Bleeding</b></li> <li><b>Perforation</b> → acute abdomen and peritonitis</li> <li><b>Scarring and strictures of muscle and mucosa</b> → pyloric stenosis</li> </ul>	<ul style="list-style-type: none"> <li>Vitamin def</li> <li>Anaemia</li> <li>OP</li> <li>Ulcerative jejunitis</li> <li>Enteropathy-associated T cell lymphoma</li> <li>NHL</li> </ul>	
Sx	<p><b>Dyspepsia</b> = indigestion seen as:</p> <ul style="list-style-type: none"> <li>Heartburn, acid brash</li> <li>Epigastric pain</li> <li>Bloating</li> <li>Nocturnal cough</li> <li>Hoarse voice</li> </ul>	<p><b>Eating:</b></p> <ul style="list-style-type: none"> <li><b>Worsens</b> gastric ulcer pain</li> <li><b>Improves</b> duodenal ulcer pain</li> </ul> <p><b>Common Sx</b></p> <ul style="list-style-type: none"> <li>Epigastric discomfort</li> <li>n/v</li> <li>dyspepsia</li> <li>Haematemesis / coffee brown vomitus + melaena</li> <li>Fe def anaemia</li> </ul>	<ul style="list-style-type: none"> <li><b>FTT</b></li> <li><b>Diarrhoea</b></li> <li><b>Abdo bloating</b></li> <li><b>UWL</b></li> <li><b>Mouth ulcers</b></li> <li><b>Anaemia</b> (secondary to Fe, B12, folate)</li> <li><b>Dermatitis herpetiformis</b> (itchy blistering skin rash on abdomen)</li> </ul>	<p><b>Rome IV criteria:</b></p> <p><b>Abdominal pain / discomfort:</b></p> <ul style="list-style-type: none"> <li>Relieved on opening bowels, <b>or</b></li> <li>change in bowel habit</li> </ul> <p><b>AND 2 of:</b></p> <ul style="list-style-type: none"> <li>Abnormal stool passage</li> <li>Bloating</li> <li>Worse symptoms after eating</li> <li>PR mucus</li> </ul>
IX	<p><b>H. Pylori</b> - Gram -ve aerobic</p> <ul style="list-style-type: none"> <li>Urease enzyme converts urea into ammonia damaging epithelium</li> <li><b>Urease breath test</b></li> <li><b>Stool antigen test</b></li> <li><b>Endoscopy + Rapid urease test + biopsy</b></li> </ul> <p><b>Endoscopy indications</b></p> <ul style="list-style-type: none"> <li>Peptic ulcers</li> <li>Malignancy (gastric, oesophageal)</li> <li>UGIB (Melaena or coffee brown vomitus)</li> </ul>	<p><b>Endoscopy (gold standard)</b></p> <ul style="list-style-type: none"> <li><b>Endoscopy + Rapid urease test + biopsy</b></li> </ul>	<p><b>Genetic testing</b></p> <ul style="list-style-type: none"> <li>HLA-DQ2 gene (90%)</li> <li>HLA-DQ8</li> </ul> <p><b>Autoantibodies</b></p> <ul style="list-style-type: none"> <li><b>Raised</b> anti-TTG - IgA antibodies</li> <li><b>Raised</b> Endomysial antibodies (EMAs) - IgA antibodies</li> <li><b>Raised</b> Deaminated gliadin peptides antibodies (anti-DGPs)</li> </ul> <p>Nb: patients with IgA deficiency → need to check total IgA levels to confirm this then check IgG version of anti-TTG or endoscopy</p> <p><b>Duodenal biopsy:</b></p> <ul style="list-style-type: none"> <li>Crypt hypertrophy</li> <li>Villous atrophy</li> </ul>	<ul style="list-style-type: none"> <li>Normal FBC, ESR and CRP blood tests</li> <li><b>Faecal calprotectin</b> negative to exclude IBD</li> <li>Negative coeliac disease serology (<b>anti-TTG antibodies</b>)</li> <li>Cancer is not suspected or excluded if suspected</li> </ul>
Mx	<p><b>LIFESTYLE</b></p> <ul style="list-style-type: none"> <li>Weight loss</li> <li>Reduce tea, coffee and alcohol</li> <li>Stop smoking</li> <li>Stop alcohol</li> <li>Avoid heavy meals before bed time</li> <li>Stay upright after meals</li> </ul> <p><b>Medications:</b></p> <ul style="list-style-type: none"> <li>Gaviscon (acid neutraliser)</li> <li>PPI</li> <li>Ranitidine (H2 antagoist)</li> <li><b>H. Pylori</b> → Triple therapy (PPI + amoxil + clarithromycin) for 7 days</li> <li>PO od → test of cure w/ urease breath test (stop PPI 4 wks before)</li> </ul> <p><b>Surgery</b></p> <ul style="list-style-type: none"> <li>Laparoscopic fundoplication</li> <li>Laser or cryotherapy to destroy epithelium in Barrett's oesophagus</li> </ul>	<p><b>LIFESTYLE</b></p> <ul style="list-style-type: none"> <li>Weight loss</li> <li>Reduce tea, coffee and alcohol</li> <li>Stop smoking</li> <li>Stop alcohol</li> <li>Avoid heavy meals before bed time</li> <li>Stay upright after meals</li> </ul> <p><b>Medications:</b></p> <ul style="list-style-type: none"> <li>Gaviscon (acid neutraliser)</li> <li>PPI</li> <li>Ranitidine (H2 antagoist)</li> <li><b>H. Pylori</b> → Triple therapy</li> </ul> <p><b>Acute abdomen signs:</b></p> <ul style="list-style-type: none"> <li>DR ABCD</li> <li>Urgent laparoscopic repair</li> </ul>	<p><b>LIFESTYLE</b></p> <ul style="list-style-type: none"> <li><b>Lifelong gluten free diet</b></li> <li>Monitor disease by checking coeliac antibodies</li> </ul> <p><b>Rare neurological symptoms</b></p> <ul style="list-style-type: none"> <li>Peripheral neuropathy</li> <li>Cerebellar ataxia</li> <li>Epilepsy</li> </ul> 	<p><b>LIFESTYLE</b></p> <ul style="list-style-type: none"> <li>Adequate fluid intake</li> <li>Regular small meals</li> <li>Reduced processed foods</li> <li>Limit caffeine and alcohol</li> <li>Low "FODMAP" diet (ideally with dietician guidance)</li> <li><b>Trial of probiotic supplements</b> for 4 weeks</li> </ul> <p><b>1<sup>ST</sup> LINE MEDS:</b></p> <ul style="list-style-type: none"> <li>Loperamide for diarrhoea</li> <li>Laxatives for constipation. Avoid lactulose as it can cause bloating.</li> <li>Linaclotide is a specialist laxative for patients with IBS not responding to first-line laxatives</li> <li>Antispasmodics for cramps e.g. hyoscine butylbromide (Buscopan)</li> </ul> <p><b>2<sup>ND</sup> Line Medication:</b> Tricyclic antidepressants (i.e. amitriptyline 5-10mg at night)</p> <p><b>3<sup>RD</sup> Line Medication:</b> SSRIs antidepressants</p> <p><b>Cognitive Behavioural Therapy (CBT)</b> → help patients psychologically manage the condition and reduce distress associated with symptoms.</p>

## GOR: GIT MEDICATIONS:

	Antacids (e.g. <i>Mylanta</i> And <i>Gaviscon</i> )	H <sub>2</sub> receptor antagonist ( <i>Ranitidine</i> ( <i>Zantac</i> ))	Proton Pump Inhibitors (PPI) E.g.: <i>Omeprazole</i> ( <i>Losec</i> ), <i>Esomeprazole</i> ( <i>Nexium</i> ) & <i>Pantoprazole</i> ( <i>Somac</i> )
<b>Mechanism of action</b>	<ul style="list-style-type: none"> <li>Base + acid = salt + water</li> <li>Weak bases react with gastric acid to neutralise it → increase pH</li> </ul>	<b>Competitively</b> block H <sub>2</sub> receptors on parietal cells → reducing gastric acid secretion	<ul style="list-style-type: none"> <li><b>Irreversibly inactivate</b> the H<sup>+</sup>/K<sup>+</sup> ATPase enzyme system</li> <li>Suppressing both stimulated and basal acid secretion.</li> </ul>
<b>Indications For Use</b>	<ul style="list-style-type: none"> <li>Dyspepsia</li> <li>PUD</li> <li>GORD</li> </ul>	<ul style="list-style-type: none"> <li>Dyspepsia</li> <li>PUD</li> <li>GORD</li> <li>Stress-ulcer prophylaxis</li> </ul>	<ul style="list-style-type: none"> <li>Dyspepsia</li> <li>Peptic ulcer disease (PUD)</li> <li>GORD</li> </ul> <ul style="list-style-type: none"> <li>Zollinger-Ellison syndrome</li> <li>H. Pylori eradication</li> <li>Scleroderma oesophagus</li> <li>Alongside NSAIDs</li> </ul>
<b>Adverse effects</b>	<ul style="list-style-type: none"> <li>Diarrhoea (Mg)</li> <li>Constipation (Al and Ca)</li> <li>Belching + flatulence (Ca CO<sub>3</sub>)</li> <li>Alkalosis (HCO<sub>3</sub><sup>-</sup>)</li> </ul>	well tolerated. No common AE	Well tolerated BUT common AE are: <ul style="list-style-type: none"> <li>Headache, Nausea, vomiting, rashes</li> <li>Diarrhoea, constipation, flatulence</li> <li>Risk of C. difficile infection, B12 def, OP fractures and pneumonia</li> </ul>

### H. pylori-related ulcers

**CAUSE:** Drugs inhibiting/neutralising gastric acid secretion

- Triple therapy regimen (First-line): PPI + ≥2 anti-bacterials
- improves eradication rates + reduces subsequent acquired resistance associated with partial treatment



### NSAIDs related ulcers → do not use for H. pylori

**CAUSE:** Adverse effect of long-term use of NSAIDs

- In the stomach, COX-1 produce prostaglandins (PGE<sub>2</sub> and PG<sub>I<sub>2</sub></sub>) → stimulate mucus and bicarbonate secretion to cause vasodilation
- Non-selective NSAIDs **inhibit** COX-1 enzymes → frequent upper GI side-effects (e.g. bleeding and ulceration)
- SOLUTION (if possible):** stop NSAIDs.
- If not: medication can be used to mitigate adverse effects

## DIARRHOEA: GIT MEDICATIONS:

### #1: PRIORITY - MAINTAIN HYDRATION → 1<sup>ST</sup> LINE = ORAL REHYDRATION

#### #2: Treat **reversible cause**

- 1<sup>st</sup> line = atropine (pre-med esp. for pro-cholinergics such as irinotecan)
- Immune-mediated colitis → infliximab then tapering **prednisone**
- Surgical resection → cholestyramine
- Carcinoid syndrome → octreotide
- Steatorrhoea / fat malabsorption → **Creon** + PPI
- Abx usage → stool M/C/S (C. difficile) → **metronidazole** or **vancomycin**
- Faecal impaction → **laxatives**

#### For **uncontrolled (late) diarrhoea:**

1. Loperamide (2mg after each stool) → opioid agonist → slow GI motility
2. **Codeine 30mg qid**
3. ABx if neutropenic or symptoms persist
4. **Combination**
5. **Specialist advice**

### #1: PRIORITY - MAINTAIN HYDRATION → 1<sup>ST</sup> LINE = ORAL REHYDRATION

➤ Since most cases of diarrhoea are **self-limiting**

Drug class	Drugs
Oral rehydration salts	<b>Hydralyte</b>
Anti-diarrhoeals (reduce GI motility)	<ul style="list-style-type: none"> <li><b>Opioids</b> = Loperamide (<b>Imodium</b>) → poorly crosses BBB</li> <li><b>Diphenoxylate</b> (with atropine: <b>Lomotil</b>) → combined with anti-cholinergic to <b>avoid abuse</b> of opioids due to their adverse effects</li> </ul>
Bulking agents: absorb fluid to reduce diarrhoea	<ul style="list-style-type: none"> <li><b>Psyllium</b> (Metamucil)</li> <li><b>ispaghula</b> (Fibogel)</li> </ul>
Other	<ul style="list-style-type: none"> <li><b>Probiotics</b>: conflicting evidence</li> <li><b>Zinc</b>: children (malnourished)</li> </ul>

## Constipation: GIT MEDICATIONS:

#### Definition: ROME III criteria:

- 25% straining
- 25% Bristol stool chart 1 or 2
- 25% sensation of incomplete evacuation
- < 3 bowel motions/week
- Manual manoeuvres to defecate

\*Red flag = altered bowel habit, PR bleed, UWL, fatigue

#### Bristol Stool Chart

Type 1	Separate hard lumps, like nuts (hard to pass)
Type 2	Sausage-shaped but lumpy
Type 3	Like a sausage but with cracks on its surface
Type 4	Like a sausage or snake, smooth and soft
Type 5	Soft blobs with clear-cut edges (passed easily)
Type 6	Fluffy pieces with ragged edges, a mushy stool
Type 7	Watery, no solid pieces. Entirely liquid

### CAUSES OF CONSTIPATION

Motility	Mechanical	Metabolic
<ul style="list-style-type: none"> <li>Diet</li> <li>IBS-C</li> <li>Colonic inertia</li> <li>Pelvic floor disorders</li> <li>Hirschsprung's</li> </ul>	<ul style="list-style-type: none"> <li>Stricture</li> <li>Tumour</li> <li>Prolapse / rectocele</li> <li>Diverticular disease</li> <li>Extrinsic compression</li> </ul>	<ul style="list-style-type: none"> <li>Diabetes</li> <li>Ca<sup>2+</sup>, K<sup>+</sup></li> <li>Hypothyroidism</li> <li>Hyperparathyroidism</li> <li>Chronic kidney failure</li> </ul>

#### Anal atresia

Neurogenic	Drugs	Others
<ul style="list-style-type: none"> <li>Stroke</li> <li>Multiple sclerosis</li> <li>Parkinson's disease</li> <li>Spinal cord injury</li> <li>Neuropathy / myopathy</li> </ul>	<ul style="list-style-type: none"> <li>Opioids</li> <li>TCAs / mood stabilisers</li> <li>Diuretics</li> <li>Iron</li> <li>Calcium supplements, etc</li> </ul>	<ul style="list-style-type: none"> <li>Scleroderma</li> <li>Coeliac disease</li> <li>Anal pain, e.g. fissure</li> <li>Chronic laxative use</li> <li>Psychological</li> </ul>

Motility	Drugs	Others
<ul style="list-style-type: none"> <li>Spina bifida</li> <li>Delayed passage (Hirschsprung)</li> </ul>	<ul style="list-style-type: none"> <li>Anti-chol (anti-sludge drugs)</li> <li>Verapamil</li> </ul>	<ul style="list-style-type: none"> <li>Cystic fibrosis</li> <li>Sepsis</li> </ul>

#### Complications

1. Faecal impaction = Abdo pain, distension, N/V
2. Rectal prolapse
3. Anal fissure + haemorrhoids

#### Investigations:

- Wt, BMI
- Abdo exam - look, palpate, percuss and listen
- DRE - anal tone, faeces
- Lower limb neuro
- AXR - ONLY if suspected for obstruction (faecal loading)

#### When to refer to gastro?

- Obvious obstruction
- Sepsis
- Red flags for malignancy (PR bleed, UWL, NS)

#### Mx (conservative):

1. Lifestyle (↑fibre, ↑fluid)
2. ↑PA = ↑colonic motility
3. Behavioural change = scheduled toilets
4. Review meds
5. Knees to chest = reduce rectal angle

#### Mx (meds):

1. **Bulking agents** (bran & husk, PEG, methylcellulose) = good long-term use
2. **Osmotic 2-3 days** (sugars = Movicol, lactulose, Mg) - make sure to keep hydrated as these draw water from colon to soften stool
3. **Stimulants 1 days** (coloxyl/senna) = for those who are immobile (NOT permanent fix) ➔ take PM to pass AM
4. **Stool softeners** (docusate sodium) =
5. **Suppositories/Enemas (rectal stimulants e.g. glycerin)**

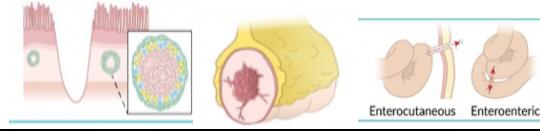
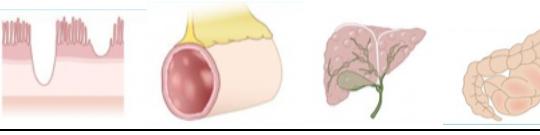
#### Practice note:

Increase in soiling/diarrhoea is common after the initiation of laxatives due to spurious overflow. If this occurs, treatment should **NOT** be reduced or stopped

## NAUSEA: GIT MEDICATIONS:

Drug	Metoclopramide (Maxolon)	Ondansetron (Zofran)	Cyclizine	Dexamethasone
MoA	<ul style="list-style-type: none"> <li>Prokinetic</li> <li>D2 antag + 5-HT3 antag</li> </ul>	5-HT3 antag (central + peripheral) <ul style="list-style-type: none"> <li>Targets CTZ</li> </ul>	H1 antag	Steroid
Dose	10mg tds ORAL	25-50mg ORAL/SC	25-50mg	4-8 mg
Indication	<ul style="list-style-type: none"> <li>Drug Metabolic,</li> <li>Chemical</li> <li>Gastric stasis</li> </ul>	<ul style="list-style-type: none"> <li>Post RT or Post chemo</li> <li>Pregnant</li> </ul>	<ul style="list-style-type: none"> <li><b>Vertigo OR motion sickness</b></li> <li><b>raised ICP</b></li> <li>SBO/LBO</li> </ul>	<ul style="list-style-type: none"> <li>+ICP</li> <li>Early BO</li> <li>Post-chemo</li> </ul>
A/E	<ul style="list-style-type: none"> <li>Dystonia</li> <li>Avoid SBO/LBO</li> </ul>	Constipation - Avoid SBO/LBO	<ul style="list-style-type: none"> <li>SAS</li> <li>Anti-chol (anti-sludge)</li> </ul>	Steroid A/E
Other	Haloperidol (D2 antag - 0.5mg) = 2 <sup>nd</sup> line	Comes in wafer forms	2 <sup>ND</sup> LINE = <b>Stemetil</b> (D2 antag - 10mg tds) 2 <sup>ND</sup> LINE = <b>hyoscine (buscopan)</b>	

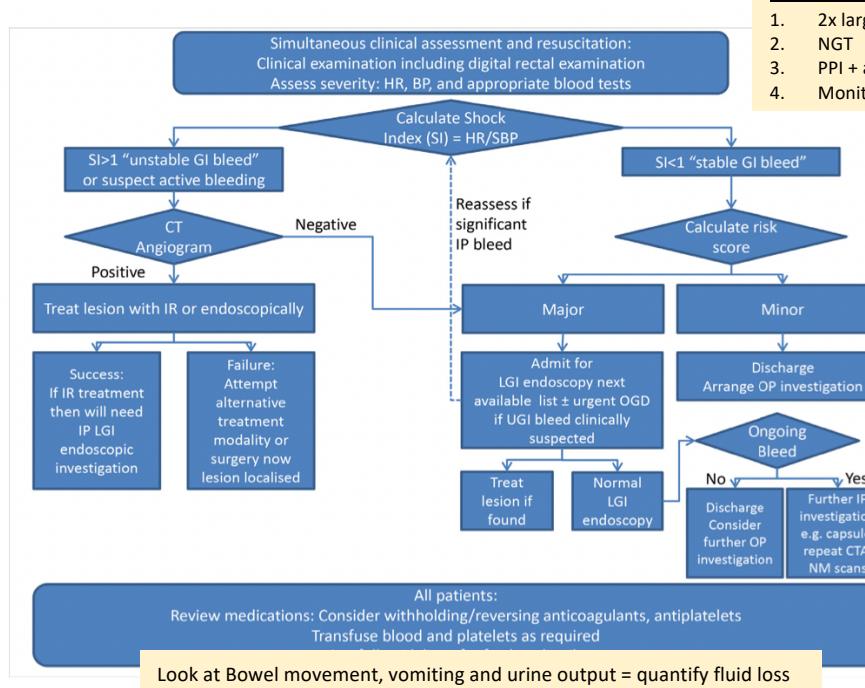
# INFLAMMATORY BOWEL DISEASE

	Crohn's	Ulcerative colitis
PP	Mediated by Th1 → IFNγ → macrophages (TNFα) and TH17 cells	Th2 mediated → IL4,13
Location	<b>Usu. terminal ileum (mouth → anus)</b> <ul style="list-style-type: none"> <li>Montreal classification → CD "Age, location, behaviour – inflammatory, structuring, perianal"</li> </ul>	<b>Sigmoid colon → rectum</b> (Montreal classification) <ul style="list-style-type: none"> <li>E1 = proctitis</li> <li>E2 = L sided UC</li> <li>E3 = Pancolitis</li> </ul>
RF	<ul style="list-style-type: none"> <li>FHx of autoimmune</li> <li>Genetics (more evident)</li> <li>Smoking (main risk factor)</li> <li>Western diet + stress + high SES</li> </ul>	<ul style="list-style-type: none"> <li>FHx of autoimmune</li> <li>Western diet + stress + high SES</li> <li>Nb: <b>smoking is a protective factor</b></li> </ul>
Clinical Sx	<ul style="list-style-type: none"> <li><b>Crampy abdominal pain</b> - NOT affected by diet or stress</li> <li><b>Malaena</b> – watery loose diarrhoea</li> <li><b>Uncontrolled bowel movements</b></li> <li><b>Fever</b></li> <li>Exhaustion</li> <li>UWL</li> </ul> 	<b>MAYO Score</b> <ul style="list-style-type: none"> <li>PR bleed + loose diarrhoea</li> <li>Tenesmus + urgency</li> <li>Chronicity of symptoms</li> <li>New constipation and diarrhoea</li> </ul> 
Extra-intestinal manifestations (EIM)	More likely to develop EIM <ul style="list-style-type: none"> <li>Arthritis (symmetrical vs asymmetrical) = 40%</li> <li>Axial arthropathy = Ank spondylitis or sacroiliitis</li> <li><b>Episcleritis</b></li> <li><b>Aphthous ulcers</b></li> <li><b>Uveitis</b></li> <li><b>Erythema nodosum (painful red papules on shin)</b></li> </ul>	<ul style="list-style-type: none"> <li>Arthritis (symmetrical vs asymmetrical) = 40%</li> <li>Axial arthropathy = Ank spondylitis or sacroiliitis</li> <li><b>PSC</b></li> <li><b>Pyoderma gangrenosum</b></li> <li><u>Less common</u> = episcleritis, aphthous ulcer, erythema nodosum (painful red papules on shin)</li> </ul>
Ix + Dx (combination)	<ul style="list-style-type: none"> <li>Clinical Hx</li> <li>Serial CRP and faecal calprotectin</li> <li>Endoscopy / colonoscopy (<b>Definitive diagnosis</b>)</li> <li>Barium X-ray = string sign (CD), lead pipe colon (UC)</li> <li>Histo findings</li> </ul> <p>Acute flare:</p> <ul style="list-style-type: none"> <li>CRP &gt; 30</li> <li>Hb &lt;100</li> <li>Raised remp</li> </ul>	<b>DDx</b> <ul style="list-style-type: none"> <li>IBD (UC, CD)</li> <li>Coeliac</li> <li>PUD, GORD</li> <li>IBS</li> <li>Colitis (ischaemic, infectious, RT, microscopic, TB-induced, NSAID)</li> </ul> <div style="display: flex; justify-content: space-between;"> <ul style="list-style-type: none"> <li>Food allergy (FPIES, cow's milk protein)</li> <li>Giardiasis</li> <li>Appendicitis</li> </ul> </div>
Histo-specific Findings	<ul style="list-style-type: none"> <li>Transmural inflammation (creeping fat)</li> <li>Skip lesions "cobblestoning"</li> <li>Granulomatous non-caseating inflammation (XS giant cells and lymphoid aggregates + ↑goblet cells)</li> <li>ASCA (yeast) antibodies</li> </ul> 	<ul style="list-style-type: none"> <li>Friable Submucosal inflammation → <b>anal region sparing</b></li> <li>Continuous lesions</li> <li>Less goblet cells</li> <li>Crypt abscess</li> <li>P-ANCA +ve</li> </ul> 
Comp.	<ul style="list-style-type: none"> <li>Fistulas (due to adhesions to viscera) <ul style="list-style-type: none"> <li>Enterourinary (UTI, air bubbles in urine)</li> <li>Enterovaginal</li> <li>Enterocutaneous</li> </ul> </li> <li>Strictures / obstructions</li> <li>Fissures</li> <li>Nephrolithiasis</li> </ul>	<ul style="list-style-type: none"> <li>Toxic megacolon → perforation → peritonitis <ul style="list-style-type: none"> <li>Subtotal colectomy + ileostomy +/- Hartmann's pouch</li> </ul> </li> <li>Haemorrhage /VTE</li> <li>CRC</li> <li>PSC → <b>cholangiosarcoma "Coursevier's signs"</b></li> </ul>
Medical Treatment	<ol style="list-style-type: none"> <li><b>Prednisolone</b> (oral pred or IV hydrocortisone)</li> <li><b>DMARDs (mod)</b> <ol style="list-style-type: none"> <li><b>Azathioprine (thiopurine)</b> - also used in RA, ALL → A/E = thrombocytopenia, liver, pancreatitis, ↑lymphoma</li> <li><b>MTX</b> (add folic acid)</li> </ol> </li> <li><b>Biologic (severe)</b> <ol style="list-style-type: none"> <li><b>Infliximab (Anti-TNF)</b> for acute flares + mod-severe disease not responding to immunomodulators</li> <li><b>Anti-IL12,23 (ustekinumab)</b> also for psoriasis</li> <li><b>Vedolizumab (anti-a4b7 integrin)</b></li> </ol> </li> </ol>	<b>Inducing remission</b> <ol style="list-style-type: none"> <li><b>DMARDs (mod)</b> <ol style="list-style-type: none"> <li>Sulfasalazine</li> <li>Azathioprine (thiopurine)</li> <li>Mercaptopurine</li> </ol> </li> <li><b>Biologic (severe)</b> <ol style="list-style-type: none"> <li><b>Infliximab (Anti-TNF)</b> for acute flares + mod-severe disease not responding to immunomodulators</li> <li><b>JAK inhibitor</b> (tofacitinib)</li> <li><b>Vedolizumab (anti-a4b7 integrin)</b></li> </ol> </li> </ol> <b>Maintaining remission:</b> <ol style="list-style-type: none"> <li>Aminosalicylate (e.g. mesalazine oral or PR)</li> <li>Azathioprine or mercaptopurine</li> <li>Clexane = ↓clot risk</li> </ol>
Surgery	<b>Only</b> for strictures and fistulas (if affecting only distal ileum)	Curative – panproctocolectomy Subtotal colectomy + permanent ileostomy or ileo-anal anastomosis (J-pouch)
Long-term	<ul style="list-style-type: none"> <li><b>Abx – metronidazole</b> (Flagyl) + cipro</li> <li><b>Biologic screen:</b> <ul style="list-style-type: none"> <li>Viral serology (Hep B/C /HIV/ CMV/EBV/VZV)</li> <li>MMR IgG +/- TB quantiferon gold</li> <li>ANCA/ASCA</li> <li>TPMT (myelosuppression due to thiopurine usage) → if low TPMT → lower dosage required</li> <li>Fe, folate, B12 and vit D</li> </ul> </li> <li><b>Endoscopy 6-9 months</b> after initial treatment</li> <li><b>MDT support to manage EIM</b> – <i>Physio, rheumatologist, gastroenterologist</i></li> <li><b>No live vaccines</b> (MMR, yellow fever) while on immunosuppressants</li> </ul>	

## APPROACH TO GI BLEEDING

	UGIB (50-150/100000)	LGIB (20-30/100000)
<b>Location</b>	Oesophagus, stomach, or proximal duodenum <ul style="list-style-type: none"> <li>Variceal?</li> <li>Non-variceal?</li> </ul>	Colon/rectum <ul style="list-style-type: none"> <li>Colonic</li> <li>PR bleeding?</li> </ul>
<b>Sx - apparent</b>	<ul style="list-style-type: none"> <li>haematemesis or coffee-brown vomitus</li> <li>bloody NGT drainage</li> </ul>	<ul style="list-style-type: none"> <li>altered blood PR</li> <li>Bright blood PR (Haematochezia)</li> </ul>
<b>Sx subtle</b>	<ul style="list-style-type: none"> <li>Epigastric pain</li> <li>anaemia</li> <li>melaena (black tarry stool suggests &gt; 150mL blood loss/day)</li> <li>faecal occult blood loss</li> </ul>	<ul style="list-style-type: none"> <li>anaemia</li> <li>melaena</li> <li>faecal occult blood loss</li> </ul>
<b>DDx</b>	<ul style="list-style-type: none"> <li><b>PUD</b> (duodenal &gt; gastric)</li> <li><b>Oesophageal variceal bleeds</b> (i.e. Portal HTN – Hx of chronic liver disease +/- encephalopathy)</li> <li><b>Mallory Weiss tear</b> (2° to vomit – often EtOH induced)</li> <li><b>Inflammation</b> (oesophagitis, gastritis)</li> <li><b>Malignancies</b> (stomach, duodenum)</li> <li><b>Dieulafoy lesion</b></li> <li><b>Vascular ectasia</b></li> <li><b>Aorto-enteric fistula</b> (post AAA repair) – painless haematochezia</li> <li><b>Boorheave syndrome</b> "oesophageal perforation" = gastrograffin oesophagography(90% dx)</li> </ul>	<ul style="list-style-type: none"> <li><b>Diverticulosis</b> (# 1 cause) – due to ↑ colonic distension <ul style="list-style-type: none"> <li>True = all bowel wall layers (Meckel's)</li> <li>False = <b>ONLY</b> mucosa and serosa (sigmoid)</li> </ul> </li> <li><b>Aorto-enteric fistula</b> (post AAA repair) – painless haematochezia</li> <li><b>Haemorrhoids</b></li> <li><b>Colitis</b> (IBD, infectious, ischaemic) → septic Sx → CT contrast → surgery</li> <li><b>Malignancy</b> (polyps, cancer)</li> <li><b>Angiodysplasia</b> – usu. R colon, painless PR bleed</li> <li><b>Dieulafoy lesion</b></li> <li><b>Post-polypectomy bleed</b></li> <li><b>Rectal ulceration</b> (irradiated proctitis)</li> </ul>
<b>Risk stratify</b>	<ul style="list-style-type: none"> <li><b>Blatchford score</b> = need for endoscopy <ul style="list-style-type: none"> <li>Raised Urea &gt; 6.5mM (blood broken down by acid in gut and reabsorbed by intestines)</li> <li>Low Hb,</li> <li>SBP &gt; 100,</li> <li>HR &lt;100</li> <li>(0-1 is low risk)</li> <li>&gt; 6 has &gt; 50% chance of requiring intervention</li> </ul> </li> <li><b>Rockall score (1/3) → risk of rebleeding and overall mortality</b> <ul style="list-style-type: none"> <li>Higher risk = shock, older age + co-morbidities</li> <li>Endoscopic stigmata of recent haem (e.g. clots, bleeding vessels)</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li><b>Oakland score</b> (≤ 8 is low risk → <b>SAFE TO DISCHARGE</b>) <ul style="list-style-type: none"> <li>Age &gt; 40</li> <li>Male</li> <li>Previous LGIB</li> <li>Blood on DRE</li> <li>HR (0-3), SBP (0-5), Hb (0-22)</li> </ul> </li> <li>Gas-colon → for Dx + Rx</li> <li>SI &gt; 1 → CT angiogram</li> <li>If CT angio -ve or SI &lt; 1 → "gas/colon"</li> </ul>
<b>Rx</b>	<b>ABATED</b> <ul style="list-style-type: none"> <li>ABCDE for immediate resus</li> <li>Bloods</li> <li>Access (2x large bore cannula)</li> <li>Transfuse if appropriate – bloods, plts, FFP (avoid XS transfusion) <ul style="list-style-type: none"> <li>Plt &lt; 10</li> <li>Prothrombinex if pt on warfarin</li> </ul> </li> <li>Endoscopy (within 24 hrs)</li> <li>Drugs (stop anti-coagulants and NSAIDs)</li> </ul>	

### Management of Acute LOWER GI bleed



#### 1<sup>st</sup> line investigations:

- Bloods** – FBC, EUC, LFT, BSL, ESR/CRP,
  - Elevated Urea!
  - Hb > 80 g/L = hold off
  - Hb < 70g/L = transfuse
- X-match (2units of blood) + save**
- Coags (INR/APTT)**
- NGT/Lavage + irrigation + aspirate contents**

#### 2<sup>nd</sup> line investigations:

- Stools in blood** – DRE / FOBT
- Endoscopy** within 24 hrs (UGIB)
- Catheter angio via brachial/femoral vein** → inject vasoconstrictive agent
- Gas-colon** (if Blatchford < 2, Oakland < 8)
- SI < 1**

#### Consider transfusion strategy:

- Patients RARELY** bleed to death → **80% deaths** – non-bleeding related e.g. **malignancy** [MAINLY], MI, pulm disease, MOF)

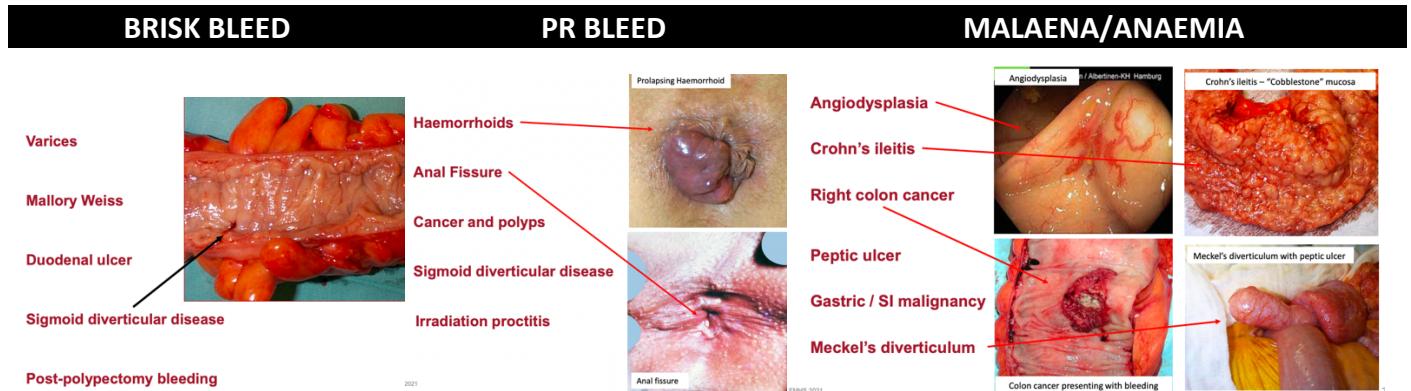
Clinical Setting	Action
INR	Sig. Bleeding
High but <4.5	No
5-9	No
>9	No
Any	Yes

IP, inpatient; IR, interventional radiology;

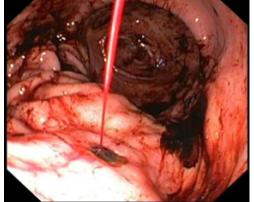
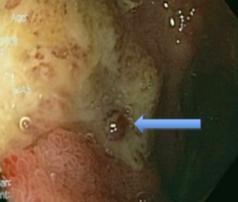
OGD, oesophagogastroduodenoscopy; OP, outpatient

# Upper Gastrointestinal Bleeding (UGIB) → Nonvariceal and Variceal [80% resolve naturally]

	Non-Variceal UGIB	Variceal [high pressure bleeds]
Def	any etiology of UGIB other than varices	Caused by esophageal or gastric varices <ul style="list-style-type: none"> <li>1/3 = cirrhosis → 1/3 fatal → 1/3 = survive (&gt; 1 year)</li> </ul>
Risk factors	<ul style="list-style-type: none"> <li>Use of NSAIDs and low-dose aspirin (LDA)</li> <li>H. pylori infection</li> </ul>	<b>Predictors of large oesophageal varices</b> <ul style="list-style-type: none"> <li>Severity of liver disease (Child-Pugh Classification)</li> <li>Hypersplenism (splenomegaly) → Thrombocytopenia &lt; 100 x10<sup>9</sup> /L</li> </ul>
Mx	<p><b>Pre-endoscopic</b></p> <ul style="list-style-type: none"> <li>Resus + ABCD (blood transfusion if Hb &lt; 70 + Hypotension)</li> <li><u>Risk stratification</u> Glasgow Blatchford <ul style="list-style-type: none"> <li>&lt; 1 = outpatient endoscopy</li> <li>&gt; 6 = need urgent endoscopy</li> </ul> </li> <li>PPI (8mg/hr) → ↑ clot formation + "downstages" lesion <u>BUT</u> does NOT reduce bleeding or mortality</li> <li><b>Erythromycin</b> = prokinetic agent</li> <li>If <b>cirrhosis</b> → vasoactive drugs and ABx</li> <li>STOP anti-coag or reverse (e.g. PTX for warfarin)</li> </ul> <p><b>Early Endoscopy</b></p> <ul style="list-style-type: none"> <li>Within <b>24 hours</b> if <u>acute UGIB</u></li> <li><u>Indications</u>: Ulcers w/ active bleeding or non-bleeding visible vessels</li> <li><b>Injection therapy (e.g. adrenaline)</b> – temporary measure → still need to stop rebleeding w/ endoscopy and embolisation</li> <li><b>Thermal probes, and clips</b></li> <li>Oesophageal Variceal bleeds → <b>tissues glues</b></li> <li>Refractory bleeding → <b>TIPS</b></li> </ul> <p><b>Post-endoscopy [NOT Dying &gt; NOT rebleeding]</b></p> <ul style="list-style-type: none"> <li>Check <u>H. pylori status</u> in all ulcer patients</li> <li><u>High ulcer risk</u> = High-dose PPI for 72 h</li> <li><u>low risk ulcers</u> = fed promptly + oral PPI</li> <li><u>Cirrhosis pts</u> = ABx for 7 days (regardless of bleeding)</li> <li><u>Variceal bleed</u> = vasoactive drugs for 5 days</li> <li>Restart low dose aspirin in CV-patients once bleeding resolved</li> </ul>	<p><b>Vasoconstrictor therapy</b></p> <ul style="list-style-type: none"> <li><b>Octreotide</b> (somatostatin analogue) → reduce splanchnic blood flow</li> <li><b>Terlipressin/somatostatin</b></li> <li><b>Vit K/FFP</b></li> </ul> <p><b>Antibiotics</b></p> <p>Prophylactic broad spectrum ABx → sig. <b>reduces</b> early rebleeding + infection/sepsis</p> <p><b>Resuscitation</b></p> <p>Minimise rapid expansion → can remove clot IVF bolus beware!!!</p> <p><b>ICU level care</b></p> <p><b>Endoscopy (OGD)</b></p> <p>Within 12 hrs + ETT required (band ligation preferred)</p> <p><b>Alternatives</b></p> <ul style="list-style-type: none"> <li><b>Early</b> placement within 24-72 hrs of <b>TIPS</b> (Transjugular Intrahepatic Portosystemic Shunt)</li> <li><b>SB Tube</b> [<b>Provide time for TIPS</b>] → <b>Gastric balloon</b> for immediate temporary control <b>BUT</b> high complication rate (e.g. necrosis, oesophageal perforation)</li> </ul> <p><b>Beta Blockade (prevention)</b></p> <ul style="list-style-type: none"> <li><b>Non-selective BB (propantheline)</b> → ↓ CO, reduces splanchnic vasoconstrict (i.e. vasodilate) = ↓ blood flow portal vein</li> <li>Reduce risk for recurrent variceal haemorrhage</li> </ul>

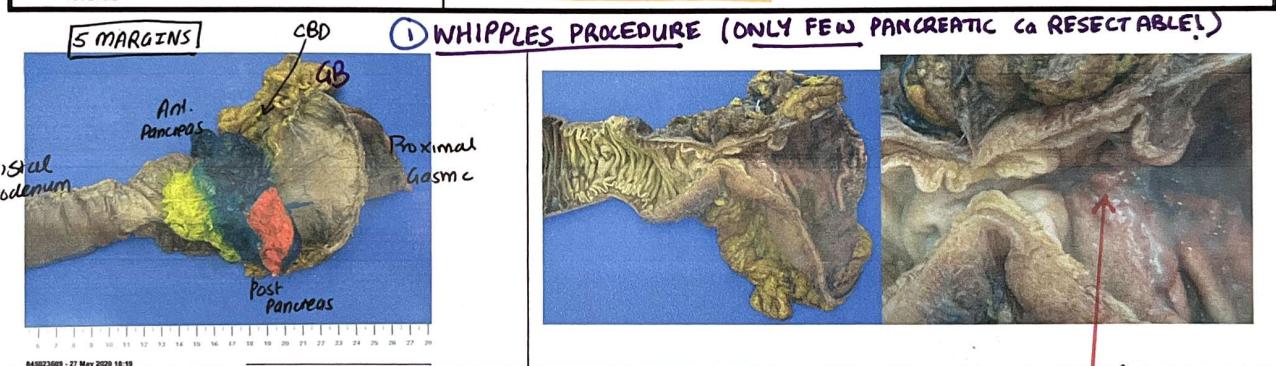


## SURGICAL IMAGES OF ACTIVE VS INACTIVE BLEEDS

Major stigmata			Minor stigmata		
<ul style="list-style-type: none"> <li>High-bleeding risk endoscopic therapy</li> </ul>			<ul style="list-style-type: none"> <li>Low rebleeding risk – NO endoscopic therapy needed</li> </ul>		
					
Active Spurting [active bleeding]	Non-Bleeding Visible Vessel	Adherent clot → usually removed with underlying lesion assessed	Flat pigmented spot	Clean base ulcer	
55-90%	40-50%	0-35%	7-10%	< 5%	
FORREST 1A – spurting	FORREST 2A	FORREST 2B	FORREST 2C	FORREST 3	
FORREST 1B – oozing					

## #A: Anatomy and Pathology of Upper GI and Pathology of the Pancreas

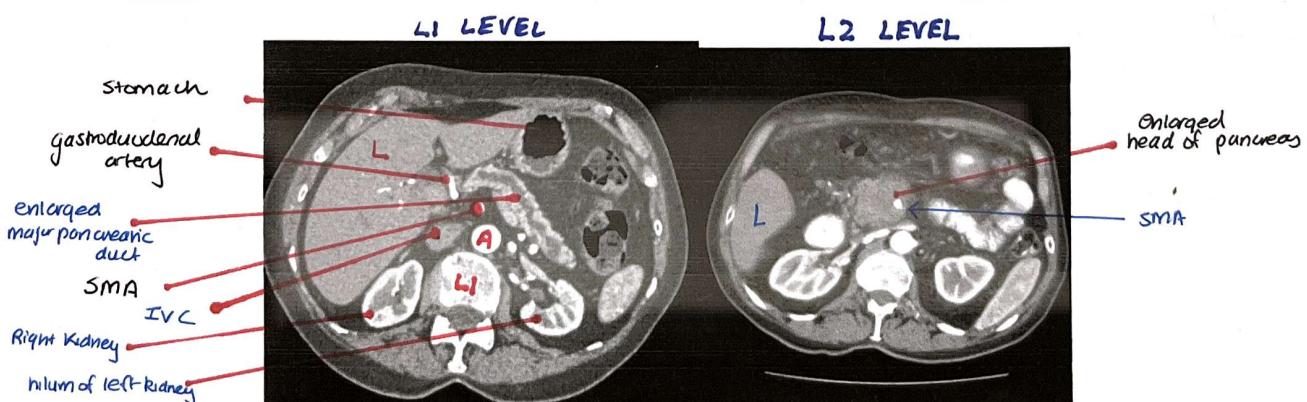
HPS	Examination	Investigations
<ul style="list-style-type: none"> <li>56 year old male Presents with abdominal pain           <ul style="list-style-type: none"> <li>Localised → parietal peritoneum</li> <li>Poorly localised → visceral peritoneum/organs</li> </ul> </li> <li>Pain is constant, not associated with eating → exc. <i>GORD, achalasia</i></li> <li>Radiates to the back → <i>pancreatitis, AAA, gastritis (PUD), costochondritis?</i></li> <li>Sudden onset 3 hours ago</li> <li>Has been progressively becoming worse</li> <li>Has noticed a dull epigastric pain for many weeks prior which he put down to reflux</li> <li>Not associated with N&amp;V or diarrhoea → exc. impaired gastric emptying</li> <li>No fever, night sweats → exc. infection, lymphoma</li> <li>No mucus, melaena or blood in the stool → exc. cancer?, Upper /Lower GI bleed</li> <li>Is passing wind → important to exc. bowel obstruction</li> <li>Not constipated (last past stool yesterday, appeared normal)</li> <li>Has lethargy</li> <li>8kg weight loss in last 6 months → cancer?</li> <li>Parasitic organisms - tapeworm?</li> <li>No rashes but has intensely itchy skin → suggests cholestasis → jaundice /scleral icterus</li> </ul>	<ul style="list-style-type: none"> <li>Appears cachectic</li> <li>Has palmar erythema</li> <li>Scleral icterus → yellow eyes (<b>Hepatic dysfunction</b>)           <ul style="list-style-type: none"> <li>Best inspected under natural light (by window)</li> </ul> </li> <li>Mouth normal</li> <li>Umbilical hernia noted with ascites</li> </ul>	<ul style="list-style-type: none"> <li>No caput medusa or spider naevi → No portal HTN</li> <li>Liver enlarged and tender to palpation → hepatomegaly</li> <li>Epigastrium tender</li> <li>No guarding</li> <li>Testes normal → exc. testicular torsion</li> <li>DRE normal</li> </ul>



- Red = pancreatic neck region
- Green-blue = portal vein bed
- Yellow = peri-uncinate soft tissue (lymph nodes)

• Mucosal ulceration at head of pancreas (most common) = AC

**ENDOSCOPIC STENT - 5x MX FOR JAUNDICE**



**Interpretation:** 40mm head of pancreas mass with **double duct sign** = simultaneous dilatation of CBD + pancreatic duct

- Nb: Arterial contrast used (left) → as seen by hyperattenuating aorta, SMA and
- Enlarged heterogeneous pancreatic duct suggestive of blockage → causing compression of gastroduodenal artery (bright line)
- Blockage cause: either gallstone OR mass

### DDx for pancreatic mass causes

- Stone with obstruction and chronic pancreatitis (check alcohol)
- Intraductal papillary mucinous neoplasm
- Mucinous cystic neoplasm
- Neuroendocrine tumour (glucagonoma, insulinoma)
- Solid pseudopapillary neoplasm
- Acinar cell carcinoma (exocrine) → *Tamylose Lipase*
- Lymphoma
- Adenocarcinoma of duodenum
- Vascular lesion (thrombus in SMV)

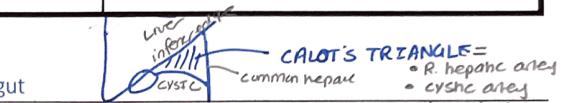
### Final Dx:

- Pancreatic ductal adenocarcinoma
- Involves portal vein bed
- PNI and LVI (i.e. perineural and lymphovascular invasion)
- Metastatic carcinoma present in umbilical hernia

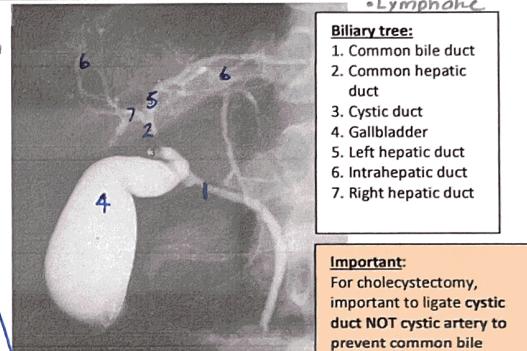
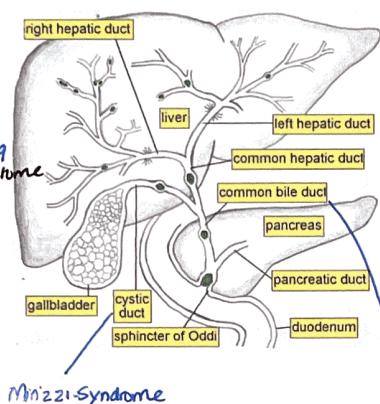
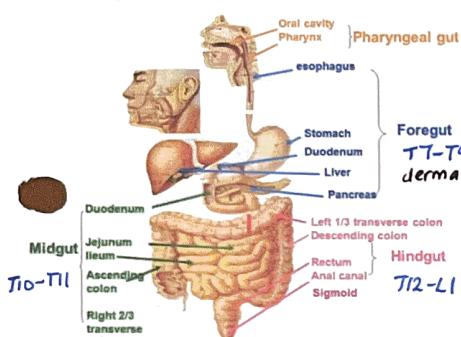
Indicates Stage IV

### Treatment MOST UNRESECTABLE

- Completion pancreatectomy — depends if margins have been exceeded
- FOLFOX chemotherapy → consider severe debilitation on patient (toxicity)
- Radiotherapy? → relatively easier as retroperitoneal location (for pancreatic tail)
- Palliation (if all fails)
- Poor prognosis → 7.1% 5-year survival rate



Describe anatomy & relationships of organs derived from embryonic foregut



**Biliary tree:**

- Common bile duct
- Common hepatic duct
- Cystic duct
- Gallbladder
- Left hepatic duct
- Intrahepatic duct
- Right hepatic duct

**Important:**  
For cholecystectomy, important to ligate cystic duct NOT cystic artery to prevent common bile duct (CBD) injury

Retroperitoneal	Intraperitoneal
<ul style="list-style-type: none"> <li>Supra-adrenal</li> <li>Aorta/IVC</li> <li>Duodenum (2<sup>nd</sup> and 3<sup>rd</sup> part)</li> <li>Pancreas (exc. tail)</li> <li>Ureter</li> <li>Ascending and descending colon</li> <li>Kidneys</li> <li>Esophagus</li> <li>Rectum</li> </ul> <p><b>SADPUCKER</b></p>	<ul style="list-style-type: none"> <li>Liver</li> <li>Stomach</li> <li>Jejunum and ileum</li> <li>Spleen</li> <li>TC</li> <li>SC</li> <li>Upper rectum</li> </ul>

### Intraperitoneal

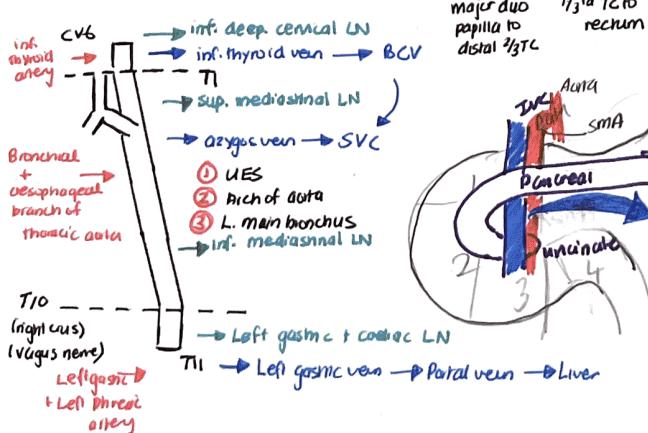
### Portosystemic anastomoses

Affected during portal HTN (e.g. caused by cirrhosis (liver scar tissue) leading to increased blood flow resistance:

- Backflow into left gastric vein causing dilatation of oesophageal veins resulting in **oesophageal varices** (fragile to rupture → haematemesis) [VARICEAL BLEED]
  - Oesophageal veins → hemiazygos → azygos → SVC → RA)
- Caput medusae** (varices in umbilical vein)
- Haemorrhoids** (varices in superior rectal vein that drain into IVC)

### Vascular supply

	Foregut	Midgut	Hindgut
BV	Coeliac trunk	SMA	IMA
Level	T12	L1	L3
Somatic pain region	T7-T9 (epigastric)	T10-T11 (umbilical)	T12-L1 (suprapubic)



### PSNS

- Vagus nerve (duo to mid-TC)**
  - Left (hepatic branch) anterior to stomach
  - Right (coeliac branch): posterior to stomach
- Pelvic splanchnic — S2-S4 (mid-TC — anal canal)**

### SNS

**Thoracolumbar SNS ganglia (T5-L2)**

- Greater splanchnic: T5-T9
- Lesser splanchnic = T10-11
- Least splanchnic = T12-L2 (sacral)

*(splenic flexure)*

\***Parietal peritoneum** = supplied by anterior rami of lower thoracic spinal nerves

R myenteric plexus = motor (SNS + PSNS) to **extensor** muscle → **motor** → **extensor**  
K Meissner plexus = PSNS secretomotoric in all **submucosa** of GIT

**SMA Syndrome** = SMA + aorta compress 3<sup>rd</sup> part of duodenum → biliary vomiting + extreme UWL  
NB **Nutcracker syndrome** = SMA + aorta compress L renal vein

**PROXIMAL**

- Sup. mes. LN
- Pancreato-duodenal LN

### Nerve supply

#### SNS

**Thoracolumbar SNS ganglia (T5-L2)**

- Greater splanchnic: T5-T9
- Lesser splanchnic = T10-11
- Least splanchnic = T12-L2 (sacral)

#### PSNS

**Pelvic splanchnic — S2-S4 (mid-TC — anal canal)**

**GASTRIC LN**

**ANTRUM** → **subpyloric LN** → **suprapyloric LN** → **distal**  
**LESSER CURVATURE** → **Ligament of Treitz** → **GASTRO-ORENTAL LN**