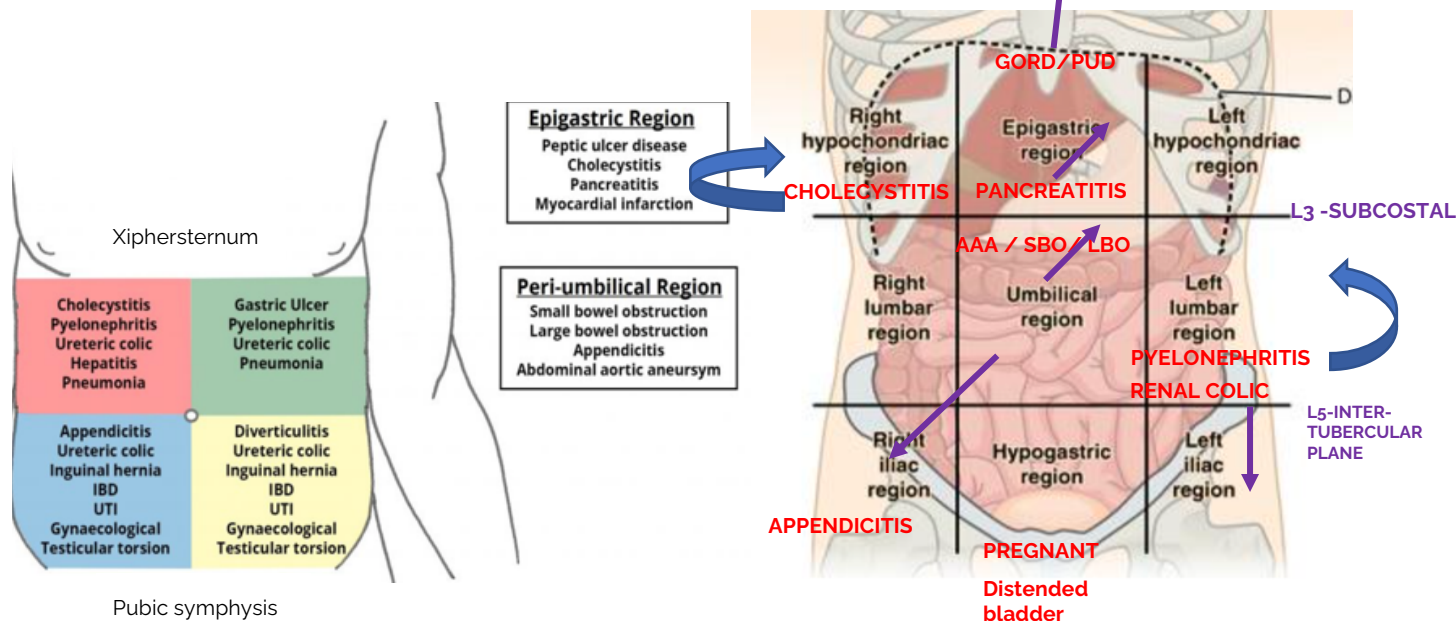


GASTROENTEROLOGY H+E:

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



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:

VOMITING <i>"contents frequency amount"</i>	<ul style="list-style-type: none"> Involuntary explosive ejection of stomach contents through mouth <table border="1"> <thead> <tr> <th>Colour</th><th>Location</th></tr> </thead> <tbody> <tr> <td>Bile (green-yellow)</td><td>SBO, duodenal atresia,</td></tr> <tr> <td>Faeculent (brown)</td><td>Small or large bowel obstruction</td></tr> <tr> <td>Coffee ground</td><td>Fe tablet, red wine & coffee ingestion</td></tr> <tr> <td>Undigested Food</td><td>Gastric outlet obstruction (level of pylorus)</td></tr> <tr> <td>Red blood/ Haematemesis</td><td> <ul style="list-style-type: none"> Upper GI Bleeding or malignancy (any bleeding proximal to Duodenal-jejunal flexure) Multiple episode of vomiting? = Mallory-Weiss Tear </td></tr> <tr> <td>Chronic unexplained</td><td>Pregnancy (AM sickness), Bullemia, Alcoholism or drugs (e.g. digoxin, opiates, chemo)</td></tr> </tbody> </table>	Colour	Location	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)	Red blood/ Haematemesis	<ul style="list-style-type: none"> Upper GI Bleeding or malignancy (any bleeding proximal to Duodenal-jejunal flexure) Multiple episode of vomiting? = Mallory-Weiss Tear 	Chronic unexplained	Pregnancy (AM sickness), Bullemia, Alcoholism or drugs (e.g. digoxin, opiates, chemo)				
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Lethargy	<ul style="list-style-type: none"> Tiredness easy fatigability = acute or chronic liver disease OR anaemia due to GI or chronic IBS 								



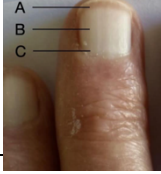



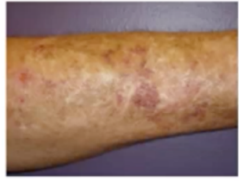



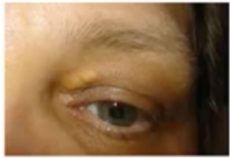




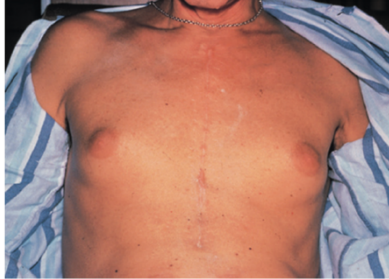
GIT examination

RED FLAG (ACUTE ABDOMEN)




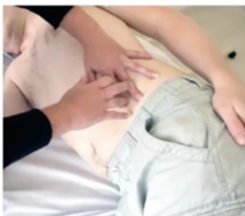
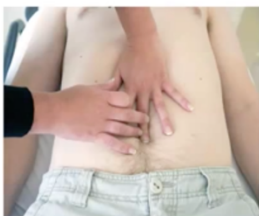
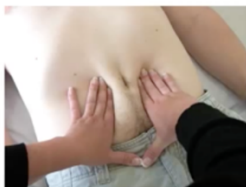


1. REBOUND TENDERNESS
2. GUARDING/RIGIDITY
3. SEPTIC SIGNS
4. ABSENT BOWEL SOUNDS

"bleed/perforation, obstruction, infarction"



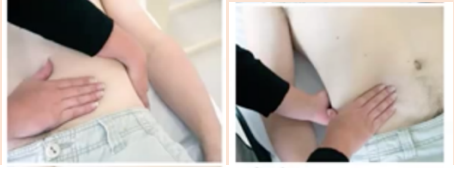
Ask patient to lie 45° (both hands on side)

General inspection	<ol style="list-style-type: none"> 1. Equipment (e.g. O₂, nasogastric feeds, medications?), Vomit bowls, IV infusions, catheters 2. Jaundice (liver disease) 3. Body habitus (obese, cachexia, ascites) 4. Well/unwell (responsiveness) → Mental state (encephalopathy) / Pain / Agitation (thyrotoxicosis) 5. Pigmentation (haemochromatosis, Whipple's disease) & Xanthomata (chronic cholestasis) 	
Nails	<ul style="list-style-type: none"> • Clubbing (cirrhosis, IBD, coeliac) • Leuconychia (transverse white lines → hypoalbuminaemia from chronic liver disease) • Koilonychia (spoon-shaped nails from severe Fe or B12/Folate deficiency) • Terry's nail (red line on tip of nail) = Chronic liver disease 	   <p>Leuconychia Clubbing</p>
Palms	<ul style="list-style-type: none"> • Pale palmar creases (anaemia) • Palmar erythema (chronic liver disease / pregnancy) • Dupuytren's contractures (alcohol) = thickened palmar fascia → fixed finger flexion usually ring or 5th finger <ul style="list-style-type: none"> ◦ EtOH, CLD, anti-epileptic • Asterixis/hepatic flap (hepatic encephalopathy / uraemia) → jerky movements of wrist due to toxic ammonia retention • Fingertip capillary glucose monitoring marks (diabetes) 	   <p>Palmar erythema Dupuytren's contracture</p>
Arms	<ul style="list-style-type: none"> • Measure BP (ask for BP in exam, don't do it) • IV Track marks (→ hepatitis C infection?) or tattoos (unregulated pallor, prison) • Bruising (clotting factor deficiency - Vit K deficiency) • Spider naevi (chronic liver disease) (> 2 is abnormal) = Blanches <ul style="list-style-type: none"> ◦ Petechiae does NOT blanch • Muscle Atrophy • Scratch marks (severe itch (pruritus) = obstructive or cholestatic jaundice = due to primary biliary cirrhosis and progressive bile duct destruction) 	  <p>Bruising</p>
Axilla	<ul style="list-style-type: none"> • lymphadenopathy • Acanthosis nigricans = darkening and thickening of skin (rarely assoc. with GIT carcinoma) <ul style="list-style-type: none"> ◦ Hyperinsulinemia → Acromegaly, PCOS, T2DM 	 <p>Acanthosis nigricans</p>
Face /eyes	<ul style="list-style-type: none"> • Uveitis = IBD, autoimmune • Sclerae: jaundice, anaemia, iritis • Xanthelasma (biliary cholangitis/cholestasis → yellowish lipid plaque in periorbital region) • Conjunctival pallor (anaemia) • SLIT-LAMP = Cornea: brown-greenish Kayser-Fleischer rings (Wilson's disease) → (abnormal Cu deposition) → low serum ceruloplasmin & Cu • Parotidomegaly = XS EtOH, MUMPS, Sjogrens, IgG4 disease <p>STIGMA OF CHRONIC LIVER DISEASE:</p> <ul style="list-style-type: none"> ➢ Jaundice → hepatic encephalopathy (asterixis) ➢ Hair loss ➢ XS E2 = palmar erythema, spider naevi ➢ Ascites → SBP ➢ Esophageal varices, caput medusae, haemorrhoids (variceal bleed) 	    <p>Kayser-Fleischer rings in Wilson's disease</p>
Mouth	<ul style="list-style-type: none"> ◦ Angular stomatitis (cracks at corner of mouth) → Fe, folate or B12 deficiency and water soluble Vit deficiencies (e.g. B2, B6, B12, B1/thiamine) ◦ Aphthous ulcers (IBD or coeliac disease) = tiny ulcers in lips and mouths ◦ Sweet Breath Smell: fetor hepaticus (end-stage chronic liver disease) • Teeth: Poor dentition (chronic liver disease, alcoholic) • Gums: gingivitis (smoking), hypertrophy (pregnancy, scurvy), • Tongue: atrophic glossitis = (Fe, folate or B12 deficiency) 	  <p>Angular stomatitis Aphthous ulceration</p>
Neck/ Chest	<ul style="list-style-type: none"> • Cervical lymphadenopathy <ul style="list-style-type: none"> • esp. Virchow's node left supraclavicular LN which is linked to GI malignancy • Troisier's signs = presence of Virchow's node + gastric cancer • Hyperpigmentation <ul style="list-style-type: none"> • heavy metals • haemochromatosis • Addison's <p>Inspect front/back</p> <ul style="list-style-type: none"> • Spider naevi Blanching test = sign of cirrhosis • Gynaecomastia (male breast enlargement and body hair loss) <ul style="list-style-type: none"> • Chronic liver disease and cirrhosis • Alcohol = ↓Leydig cell = testis atrophy + gynaecomastia • Drugs (DISCO MTV - digoxin, isoniazid, spironolactone, cimetidine, E2, methyl dopa, TCA, verapamil) • Endocrine (Hypogonadism, Hyperthyroidism) • Seminoma (testicular cancer) = ↑B-HCG, ↑E2 • Cullen's (central umbilical) and grey-turner's (flank - retroperitoneal haemorrhage) sign (pancreatitis) 	 <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

Abdomen (begin here if asked to do so)	Inspect	<div>Stand back then squat down beside bed at eye level (look for distensions)</div> <ul style="list-style-type: none">Distension (Fluids, flatus, faeces, fat, foetus, "filthy" big tumours, fibroids, fake pregnancy)<ul style="list-style-type: none">liver cirrhosis → hypoalbuminemia → ascitesAcute sudden distension due to flatus: IBS (benign) or post-bowel surgeryScars/Stomas (indicates previous bowel surgery) → CHECK position of scarCaput Medusae = distended veins flowing away from umbilicus in Portal HTN (e.g. liver cirrhosis, persistent ductus venosus)IVC obstruction = distended veins flowing towards umbilicusBruising/Pigmentation/Striae (pregnancy, Cushing's)Visible peristalsis + Pulsations (AAA) <div><div> Ascites</div><div> Caput Medusae</div></div>												
	Palpate	<div><div>1. Any pain? (begin to palpate away from pain) → watch for discomfort</div><div>a. Press down with stethoscope to check for TRUE guarding</div><div>2. Palpate over 9 regions (superficial then deep) → Look for tenderness, rigidity, organomegaly</div></div> <div><table><tr><td>Painful</td><td>Inflammatory<ul style="list-style-type: none">voluntary/involuntary guarding/rigidity (peritoneal irritation – blood, pus).rebound tenderness (peritonitis)</td></tr><tr><td>Craggy</td><td>Neoplastic<ul style="list-style-type: none">Hepatomegaly = RHF, CCF, Hepatitis, PBC, PSC, Wilson's, hemochromatosis, MALFD<ul style="list-style-type: none">Big liver BUT no liver issue = GB issue, Reidel's lobe, hyperinflated lungSplenomegaly = Haematological malignancy, Portal HTN, InfiltrativeBallotting Kidney (L/R flank)</td></tr><tr><td>Tympanic</td><td>Palpable bowel loops, SBO/LBO</td></tr><tr><td>Reducible masses</td><td>Hernia – incisional, spigelian (semilunaris), direct vs indirect inguinal</td></tr><tr><td>Pulsatile</td><td>AAA</td></tr><tr><td>Special tests</td><td><div>1. Courvoisier' sign (palpable non-tender gallbladder + jaundice) → head of pancreas cancer</div><div>2. Rovsing's sign (appendicitis).</div><div>3. Murphy's sign (cholecystitis) → holds breath while palpate R subcostal area (pain = +ve test)</div><div>4. Murphy's punch (pyelonephritis) → costovertebral angle tenderness over R12 → ADPKD</div></td></tr></table><div></div></div>	Painful	Inflammatory <ul style="list-style-type: none">voluntary/involuntary guarding/rigidity (peritoneal irritation – blood, pus).rebound tenderness (peritonitis)	Craggy	Neoplastic <ul style="list-style-type: none">Hepatomegaly = RHF, CCF, Hepatitis, PBC, PSC, Wilson's, hemochromatosis, MALFD<ul style="list-style-type: none">Big liver BUT no liver issue = GB issue, Reidel's lobe, hyperinflated lungSplenomegaly = Haematological malignancy, Portal HTN, InfiltrativeBallotting Kidney (L/R flank)	Tympanic	Palpable bowel loops, SBO/LBO	Reducible masses	Hernia – incisional, spigelian (semilunaris), direct vs indirect inguinal	Pulsatile	AAA	Special tests	<div>1. Courvoisier' sign (palpable non-tender gallbladder + jaundice) → head of pancreas cancer</div> <div>2. Rovsing's sign (appendicitis).</div> <div>3. Murphy's sign (cholecystitis) → holds breath while palpate R subcostal area (pain = +ve test)</div> <div>4. Murphy's punch (pyelonephritis) → costovertebral angle tenderness over R12 → ADPKD</div>
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Percuss	<ul style="list-style-type: none">Ascites (shifting dullness) → assoc. with portal HTN, splanchnic vasodilation, cirrhosis (hypoalbuminemia), raised RAAS<ul style="list-style-type: none">Percuss for dull note in 9 regions → if dull, keep hand on position and roll patient to opposite sideKeep hand in position → wait 30s → percuss again → if resonant = ascitesPercussion tenderness <div></div>													
Auscultate (BEST HEARD IN RIF)	<table><tr><td>Normal</td><td>Borborygmia (gurgling, rumbling, or growling)</td><td></td></tr><tr><td>Absent</td><td>Functional obstruction due denervation<ul style="list-style-type: none">loss in peristalsis and GI motility</td><td><ul style="list-style-type: none">Hirschsprung diseaseparalytic ileus or peritonitis</td></tr><tr><td>High-pitched tinkling sound</td><td>Mechanical bowel obstruction</td><td><ul style="list-style-type: none">Acute SBOLeft colon carcinoma</td></tr><tr><td>Reduced</td><td>Reduction in the loudness, tone, or regularity of the sounds</td><td><ul style="list-style-type: none">Peritonitis, PancreatitisMedication Post surgery</td></tr></table>	Normal	Borborygmia (gurgling, rumbling, or growling)		Absent	Functional obstruction due denervation <ul style="list-style-type: none">loss in peristalsis and GI motility	<ul style="list-style-type: none">Hirschsprung diseaseparalytic ileus or peritonitis	High-pitched tinkling sound	Mechanical bowel obstruction	<ul style="list-style-type: none">Acute SBOLeft colon carcinoma	Reduced	Reduction in the loudness, tone, or regularity of the sounds	<ul style="list-style-type: none">Peritonitis, PancreatitisMedication Post surgery	
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OTHER	<div><div>Abdo aorta</div><div>AAW Masses</div><div>Bladder</div><div>DRE</div></div> <ul style="list-style-type: none">Between umbilicus and xiphoid process (lateral → medial)Expansile pulsation (outward movement) → AAA <p>Patient to lift their head up → intra-abdominal mass disappears or decrease in size</p> <ul style="list-style-type: none">E.g. <u>sebaceous cyst, epigastric/incisional hernia, rectus sheath haematoma or divarication</u>Smooth and firm + suprapubic dullness over pubic symphysis → bladder scan → IDC (urinary retention)Anal sphincter tone (anal wink) <div></div>													
Legs	<ul style="list-style-type: none">Skin lesions<ul style="list-style-type: none">Erythema nodosum → IBDPyoderma gangrenosum → IBDArthralgia:<ul style="list-style-type: none">Enteropathic arthritisPitting Oedema (hypoalbuminemia due to chronic liver disease)Neurological signs (alcohol) → chronic liver disease <div><div> Pitting oedema (chronic liver disease)</div><div> Erythema nodosum (inflammatory bowel disease)</div></div>													
Summary	<p>"Today I performed an abdominal exam on ____ . On inspection, any peripheral stigmata of GIT disease.</p> <p>1. On palpation (was abdomen soft/tender/masses?) → On auscultation (normal bowel sounds? Bruits?)</p> <p>• To complete the exam, check external hernial orifices, inguinal LNs, external genitalia and perform DRE</p> <ul style="list-style-type: none">DRE + PV exam → inspect (fistulae, tags, blood, mucus), palpate (masses) → look for rectal masses and haemorrhoidsSTOOL colour → pale (chronic pancreatitis), maleena (haemorrhoids, colonic pathology)Urinalysis (for bilirubin and urobilinogen, and glucose) → Temperature chart (infection)													
Further Ix	<div>Bedside:</div> <div><div>1) Urine M/C/S</div><div>2) Urine dipstick (bilirubin)</div><div>3) Urine B-HCG</div><div>4) ECG</div><div>5) Stool M/C/S</div><div>6) Bladder scan</div></div>	<div>Bloods:</div> <div><div>2. FBC</div><div>3. EUC</div><div>4. LFT</div><div>5. CRP</div><div>6. LIPASE/AMYLASE</div><div>7. COAG +/- GROUP = HOLD</div></div> <div>Imaging:</div> <div><div>➤ USS – transabdominal, transvaginal</div><div>➤ AXR</div><div>➤ ERECT CXR</div><div>➤ CT-abdo-pelvis +/- contrast</div></div>												

EXAMINATION OF THE ABDOMINAL ORGANS: (in detail)

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

DIGITAL RECTAL EXAMINATION [DRE]

Indication	<ul style="list-style-type: none">• Suspected haemorrhoids (i.e. engorged external or internal BV) → Rx: rubber bands, proctosedyl topical• Unexplained bleeding on history → bright red (distal), mixed in (left colic – UC), Maroon colour (upper GI)• Prostate screening → oliguria• Altered bowel habits (esp. >40 y.o.)	CAJ COLD for general inspect: <ul style="list-style-type: none">• Cyanosis• Anaemia• Jaundice• Clubbing• Oedema• Lymph nodes• Dehydration										
Equipment	<ul style="list-style-type: none">• Gloves → bluey → gel											
Consent & rationale	Explain examination and offer chaperone [clinically trained!!] <ul style="list-style-type: none">• "Have you had one done before?" → <u>always address patient anxiety/privacy/past experiences</u>• "Undress from waist down, then lie on your left side while bringing your knees up to your chest" NB: for prostate exam → patient standing and in the bent-over position											
Inspect	<u>Spread buttocks</u> → look for: <ul style="list-style-type: none">• Blood• Excoriations - Rash/eczema/pruritus ani (due to faecal soiling)• Fistula → may occur with Crohn's disease or perianal abscess• Fissures → IBD, malignancy or STDs• Anogenital Warts (a.k.a. condylomata acuminata) → HPV infected (needs removal)• Hemorrhoids (small <1cm tense bluish swellings)• Skin tags (collagen proliferation) → due to haemorrhoids or Crohn's	<ul style="list-style-type: none">• <u>Patient Bends down</u> → look for pelvic organ prolapse<ul style="list-style-type: none">○ Anterior (cystocele)○ Middle/Apical (uterine prolapse/ enterocoele)○ Posterior (rectocele)										
Palpate	<ol style="list-style-type: none">1. Test anal wink → stroke 4 quadrants of anus with cotton pad → observe brisk anal contraction (if none = spinal cord issue)2. Excruciating pain → ANAL FISSURE (due to primary or secondary constipation) → abandon exam											
Exam	<ol style="list-style-type: none">3. Lubricate gloved index finger → approach anus posteriorly → pause when over anus → wait until sphincter relaxes4. Warn patient → advance finger into anus<ol style="list-style-type: none">a. Faeces consistencyb. Ask patient to squeeze finger → Anal tone? [high tone = constipation]c. Ask patient to bend down → brings high Rectal lesions down5. 360° sweep for mass/wall thickenings<ol style="list-style-type: none">a. For men: feel 2 lobes of <u>anterior wall of prostate</u> and comment mass, size, symmetry, texture	<ol style="list-style-type: none">6. Remove finger → wipe on gauze → inspect mucus, blood and melaena<ol style="list-style-type: none">a. Careful of false +ve → red meet, aspirin, oral Fe, peroxidase, anticoagulants7. Clean anus	<table><tr><th>Feeling of prostate</th><th>Disease</th></tr><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></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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Very hard nodule	Carcinoma of the prostate											
Boggy and tender	Prostatitis											
Mass above the prostate or cervix	Metastatic deposit on blumer's shelf											
To complete	<ol style="list-style-type: none">1. Thank + redress + dispose waste + hand hygiene2. Document findings and name chaperone3. <u>To complete the exam I would perform:</u><ol style="list-style-type: none">a. PSA serology, Rectal US, FBC (anaemia), sigmoidoscopy and colonoscopy											



External haemorrhoids
(venous plexus engorgement)



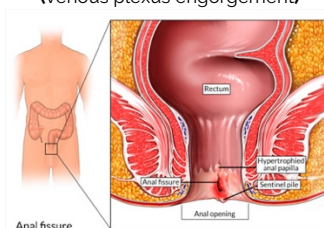
Skin tags



Anogenital warts (HPV)

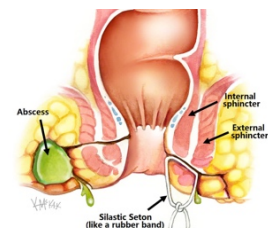


Rectal Prolapse



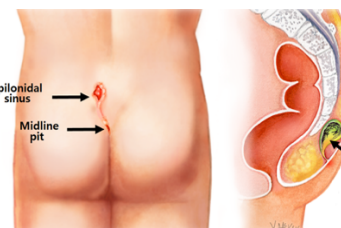
Anal fissure

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

CAJ COLD for general inspect:

- Cyanosis
- Anaemia
- Jaundice
- Clubbing
- Oedema
- Lymph nodes
- dehydration

Indication	<ul style="list-style-type: none"> • Obvious mass
Consent & rationale	<p>Explain examination and offer chaperone [clinically trained!]</p> <ul style="list-style-type: none"> • Put on gloves
Inspect	<ul style="list-style-type: none"> • General = responsive, pallor. Cachexia, Abd distension (obstruction), vomit bowls • Lumps = Size, shape, position, scrotal extension • Observe cough impulse
Palpate	<ul style="list-style-type: none"> • Scrotal contents → palpate anteriorly. If lump present, you should be able to get above it <ul style="list-style-type: none"> a. Palpable upper border = hydrocele b. Impalpable = inguinoscrotal/indirect inguinal hernia • Lump/inguinal area (bilateral) → palpate lateral to medial along inguinal ligament with one hand on patient's back <ul style="list-style-type: none"> • SITE • SIZE • SHAPE • CONSISTENCY • CONTOURS • COLOUR • TENDER • TEMP • TRANSILLUMINATION • Feel with cough impulse → compress lump/inguinal area firmly (ask patient to turn away and cough) <ul style="list-style-type: none"> a. If mass tenses + expands = positive cough impulse • Reducibility (indirect vs direct) <ul style="list-style-type: none"> a. Locate deep inguinal ring (between ASIS + pubic tubercle) b. Press on lump beginning inferiorly → compress deep inguinal ring c. Ask patient to cough: <ul style="list-style-type: none"> i. Hernia is still absent = indirect hernia ii. Hernia reappears = direct hernia d. Release and watch hernia reappear <ul style="list-style-type: none"> i. Hernia slides down obliquely = indirect ii. Hernia projects forward = direct <p><i>*If cannot reduce hernia → repeat with patient supine</i></p>
Percuss & Auscultate	Determine if bowel is present in hernia
To complete	<p>To complete the exam I would perform:</p> <ul style="list-style-type: none"> • Full abdominal exam

WHAT ARE HERNIAS?

- Abnormal protrusion of any organ through any fascia or muscle
- Abdominal, cerebellar, hiatal, lumbar (QL)

	Indirect inguinal	Direct inguinal
Location	Between inguinal ring + scrotum	Superior to pubic tubercle
Herniated Abd contents	run within inguinal canal	Come out of abd in straight line
Exits	Deep inguinal ring	Superficial ring
Reducibility	Can be contained	Cannot contain (i.e. incarcerated)

Other hernia types:

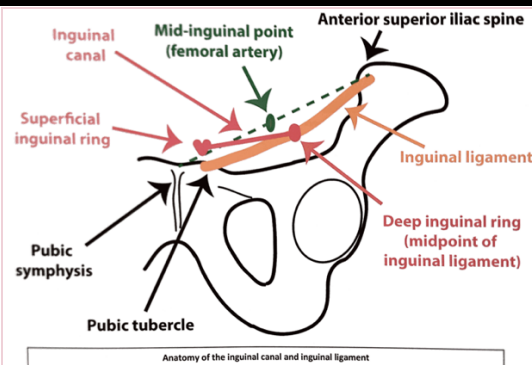
1. **Inguinal hernia** = medial and superior to pubic tubercle
2. **Femoral hernia** = **ALWAYS** lateral and inferior to pubic tubercle
commonly mistaken for enlarged inguinal LN
3. **Epigastric hernia** = hernia in epigastric region
4. **Incisional hernia** = abd scar causes abdominal weakness
5. **Umbilical hernia** = after pregnancy
6. **Hiatal hernia** = stomach herniates diaphragm

RISK FACTORS FOR HERNIA:

- 1) ++ intra-abdo pressure (obese, pregnant, abscess)
- 2) weak fascia or muscle (XS manual labour, previous abdo surgery)

RED FLAGS:

1. Inflammation
2. Irreducible
3. Obstructed bowel lumen within hernia
4. Strangulated hernia (vascular compromise of bowel) – esp. femoral hernias



Inguinal hernia



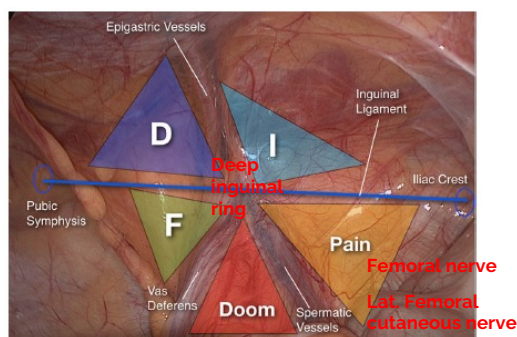
Incisional hernia

DDx: divarication of recti

RF: Obese, male

Ix: see lump when doing crunches

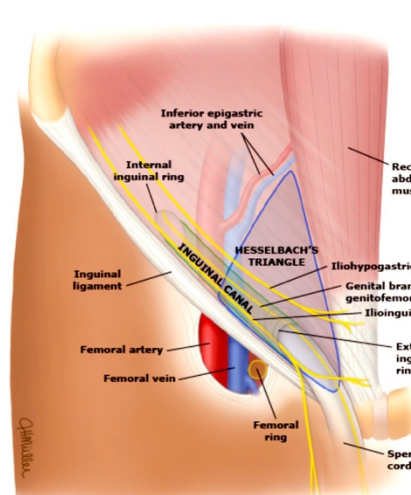
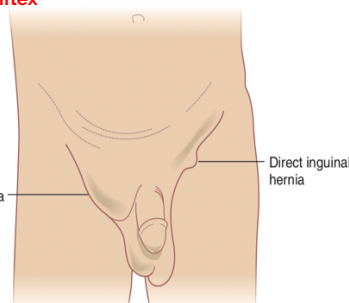
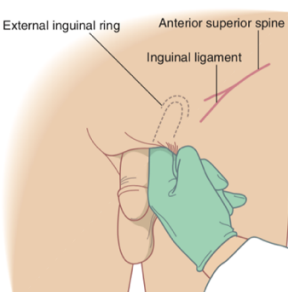
Rx: lose weight (not surgery)



EIA/EIV

Genital branch of genitofemoral nerve

Deep circumflex BV



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:
 - hepatic artery
 - Portal vein
 - Bile ductule
 - lymphatics
- Sinusoids** = interaction between blood and hepatocytes via Fenestrated endothelium
- Hepatocytes** = secrete 500-1200mL bile (i.e. bile salt + bilirubin + cholesterol + water etc.)
 - 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:

- Hepatic artery (25%) = oxygen rich
- Portal vein (75%) = nutrient rich
- Hepatic veins → IVC

Hepatic zones:

- Zone 1 (outer/peripheral)** = high O_2 → GNG,
- Zone 2 (middle/transitional)**
- Zone 3 (inner/pericentral)** = low O_2 → anaerobic tasks (e.g. detoxification, glycolysis, lipogenesis etc.) → prone to hypoxia

Main functions of liver:

- Bile drainage**
- BSL regulator** – produce ketones and stores glycogen
- Synthesise and store AA, vitamins, and fats
- Blood circulation & filtration**
- Immunologic** → acute phase proteins (CRP) & Detoxification
- Clotting factor production
- Albumin production

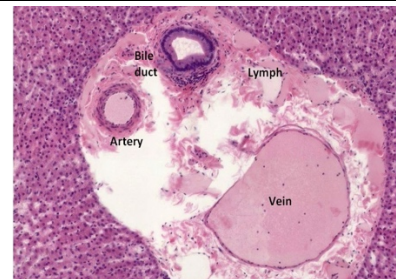
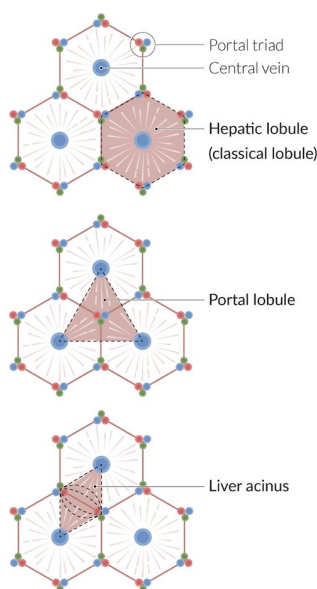
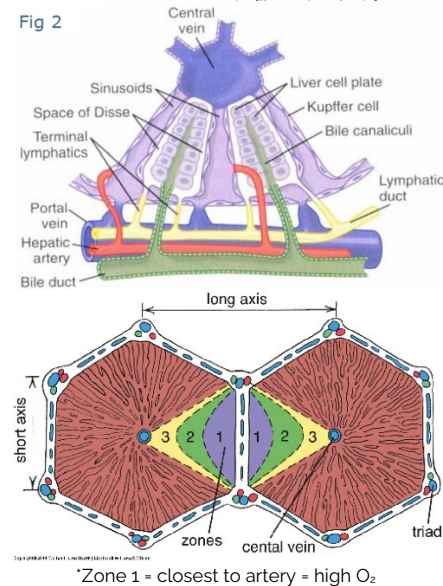


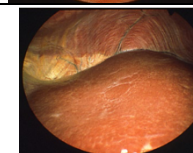
Fig 2



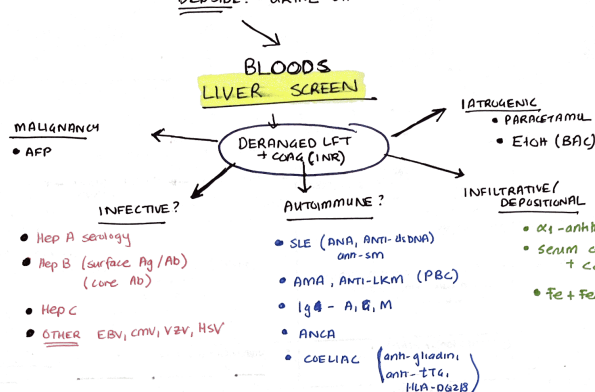
*Zone 1 = closest to artery = high O_2

INTERPRETATION OF LIVER FUNCTION TESTS

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



BEDSIDE: URINE DIPSTICK – BILIRUBIN



IMAGING: USS → GALLSTONES, CYSTS, NODULAR
ELASTOGRAPHY → MAFD
CTAP → HCC

BIOPSY INDICATION

- CONFIRM DX (WILSON, HAEMOCHROMATOSIS, HCC)
- STAGE/GRADE (HEPATITIS, HCC, POST-TRANSPLANT)
- UNKNOWN DX FOR DERANGED LFT

OR

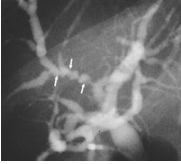

PARACENTESIS → CELL COUNT + CYTOLOGY
(ASCITIC FLUID)

- LDH
- Glucose
- Protein
- Culture
- Serum: albumin/ascite

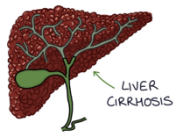
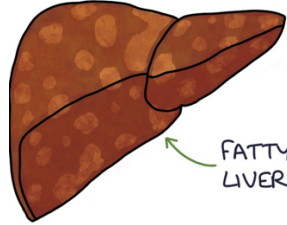
OR BIOLOGICS SCREEN

- Fe, B12, Folate, vit D (25-OH)
- MMR IgG → TPMT (myelosuppressed due to thiopurine)
- Hep B/C/HIV
- EBV, CMV
- Quantiferon Gold TB
- ASCA (cruhn)
- p-ANCA (IUC)

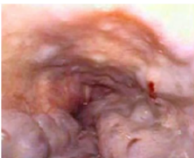
RARE HEPATIC PATHOLOGIES

	Autoimmune hepatitis	HAEMOCHROMATOSIS	WILSON'S DISEASE	ALPHA 1 ANTITRYPSIN DEF.	PRIMARY BILIARY CIRRHOSIS	PRIMARY SCLEROSING CHOLANGITIS
Def	Rare cause of chronic hepatitis	<ul style="list-style-type: none"> HFE gene mutation (Chr 6) Autosomal recessive 	<ul style="list-style-type: none"> XS copper accumulation due to mutated ATP7B copper-binding protein Autosomal recessive 	Mutated gene for protease inhibitor (a1-antitrypsin) → prevents inhibition of neutrophil elastase <ul style="list-style-type: none"> Autosomal recessive (A1AT gene Chr 14) 	Autoimmune attack on small bile ducts (1 st = intralobular ducts – canals of hering) RISK FACTORS: <ul style="list-style-type: none"> Middle aged women Autoimmune hx (e.g. thyroid, coeliac) Rheumatoid hx (e.g. SSc, Sjogren, RA) 	<ul style="list-style-type: none"> Stricture and fibrotic intrahepatic and extrahepatic ducts Sclerosis = stiffened and hardened bile ducts RISK FACTORS: <ul style="list-style-type: none"> Male aged 30-40s Ulcerative colitis (70%) FHx
Sx	<ul style="list-style-type: none"> Type 1 (adults) = women 40-50s (menopausal) Type 2 (children) – teenagers (acute hepatitis w/ high ALT/AST and jaundice) 	<ul style="list-style-type: none"> Fatigue Arthralgia Bronze diabetes Hair loss ED Amenorrhoea Cognitive (memory and mood swings) 	Neuro problem (50%) <ul style="list-style-type: none"> Dysarthria Dystonia Basal ganglia (Parkinson's) – asymmetrical motor sx Hepatic problem (40%) <ul style="list-style-type: none"> Chronic hepatitis → cirrhosis 	Lung issues: <ul style="list-style-type: none"> SOB Reduced exercise tolerance Hepatic issues: <ul style="list-style-type: none"> Fatigue jaundice 	<ul style="list-style-type: none"> Fatigue Pruritus GI disturbance + abdo pain Jaundice Pale stools Xanthoma & xanthelasma Cirrhosis signs (ascites, +SM, spider naevi) 	<ul style="list-style-type: none"> Jaudnice Chronic RUQ pain Pruritus Fatigue + HM 
Comp.	<ul style="list-style-type: none"> Cirrhosis Death <hr/> Assoc. <ul style="list-style-type: none"> Arthritis Hashimoto's 	<ul style="list-style-type: none"> T1DM (pancreatitis) Liver cirrhosis Pituitary & gonad infiltration (hypogonad, impotence, amenorrhea, infertility) Cardiomyopathy HCC Hypothyroidism Pseudogout & Chondrocalcinosis 	Psych Problem (10%) <ul style="list-style-type: none"> Depression → full psychosis Other: <ul style="list-style-type: none"> Kayser-Fleischer rings Haemolytic anaemia Renal tubular acidosis Osteopenia 	2 main complications: <ol style="list-style-type: none"> Bronchiectasis and emphysema (>30yo) Liver cirrhosis (after 50yo) leading to HCC 	Inflamed bile ducts causes obstruction of bile outflow = cholestasis <ol style="list-style-type: none"> Fibrosis Advanced liver Cirrhosis Portal HTN and liver failure <ul style="list-style-type: none"> Symptomatic pruritus Steatorrhea (greasy stools) Hypothyroidism OP HCC 	Chronic bile obstruction leads to: <ol style="list-style-type: none"> Fibrosis Advanced liver Cirrhosis Portal HTN and liver failure <hr/> SPECIFIC <ul style="list-style-type: none"> Acute bacterial cholangitis Cholangiosarcoma CRC Biliary stricture Fat soluble vit. Def (A,D,E,K)
Ix	<ul style="list-style-type: none"> Deranged LFT Elevated IgG Type 1 Autoantibodies: (ADULTS) <ul style="list-style-type: none"> ANA Anti-SM Anti-SLA/LP Type 2 Autoantibodies: (CHILDREN) <ul style="list-style-type: none"> anti-LKM1 anti-LC1 liver Biopsy to confirm Dx 	<ul style="list-style-type: none"> Fe studies (raised serum ferritin & transferrin) Liver biopsy – Perl's stain → Fe concentrated in parenchymal cells Genetic testing (H63D, c282y) CTAP MRI – Fe deposition on liver 	<ul style="list-style-type: none"> REDUCED Serum ceruloplasmin (not specific) REDUCED serum Cu RAISED 24 hr urinary Cu assay Liver biopsy (gold standard) 	<ul style="list-style-type: none"> REDUCED serum alpha-1 antitrypsin Liver biopsy (gold standard) – acid-Schiff +ve staining globules Genetic testing (A1AT gene) HRCT thorax → diagnose bronchiectasis and emphysema 	Bloods: <ul style="list-style-type: none"> Raised ESR Raised IgM Deranged LFT: <ul style="list-style-type: none"> Raised ALP (1st) ALT/AST raised later (AST:ALT >1 = CIRRHOSIS) Autoantibodies: <ul style="list-style-type: none"> AMA (most specific to PBC) ANA (present in 35% of patients) 	Deranged LFT: <ul style="list-style-type: none"> Cholestatic picture Elevated ALT/AST if progression to cirrhosis Autoantibodies (not useful) <ul style="list-style-type: none"> P-ANCA (94%) ANA (77% of patients) Anti-cardiolipin (63%) MRCP (gold standard) <ul style="list-style-type: none"> MRI of liver, bile duct and pancreas → lesions/strictures Beaded pattern stricture
AST	++++ (2x ALT)	NOT HELPFUL	++++ (2x ALT)	NOT HELPFUL	+	NOT HELPFUL
ALT	++	NOT HELPFUL	++	NOT HELPFUL	+	NOT HELPFUL
ALP	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	++++	++++
GGT	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL	++++	++++
Mx	<ol style="list-style-type: none"> High dose steroids (pred) Azathioprine (life-long) Liver transplant (may recur) 	<ol style="list-style-type: none"> Venesection (weekly) Monitor serum ferritin Avoid alcohol Genetic counsel (family) Rx complications 	Cu chelation using: <ol style="list-style-type: none"> Penicillamine Trientine 	<ul style="list-style-type: none"> Stop smoking Symptomatic Mx Lung and liver transplant Monitor for complications (e.g. HCC) 	<ol style="list-style-type: none"> Ursodeoxycholic acid (reduced intestinal absorption of cholesterol) Cholestyramine (binds to bile acids to prevent gut absorption – improve pruritus) Immunosuppression (steroids) Liver transplant (end-stage) 	<ul style="list-style-type: none"> ERCP – dilate + stent strictures (XR guided + contrasts used) Cholestyramine (binds to bile acids to prevent gut absorption – improve pruritus) Monitor comp. (cholangiosarcoma, oesophageal varices) Liver transplant (curative intent – 80% 5-year survival)

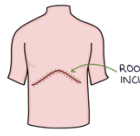
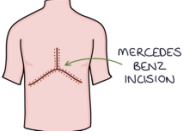
COMMON LIVER DISEASE – NAFLD, ALCOHOLIC, CIRRHOSIS

	MAFLD / NAFLD	ALCOHOLIC LIVER DISEASE	LIVER CIRRHOSIS																																				
Risk factors	<ul style="list-style-type: none"> 30% of adults Part of metabolic syndrome Risk factors <ul style="list-style-type: none"> Obese – poor diet low PA MetSyn (HTN, HC, T2DM) Smoker Middle-aged 	<ul style="list-style-type: none"> Long-term EtOH consumption Genetic predisposition 80% alcohol → metabolised by Alcohol dehydrogenase into Acetaldehyde → forms adducts with proteins in hepatocytes → activate immune response - cell injury 20% alcohol → metabolised by Cytochrome CYP2E1 into acetate releases ROS → lipid peroxidation and mitochondrial damage increased gut permeability → Endotoxin released → inflammatory cytokines → accelerated damage 	<ul style="list-style-type: none"> Chronic inflammation and damage to liver cells Damaged cells replaced by scar tissue (fibrosis) and nodules Common causes: <ul style="list-style-type: none"> Alcoholic liver disease NAFLD Hep B/C 																																				
Comp.	Step wise progression: <ul style="list-style-type: none"> Non-alcoholic fatty liver disease Non-alcoholic steatohepatitis (NASH) Fibrosis Cirrhosis Increased risk of: <ol style="list-style-type: none"> heart disease stroke diabetes mellitus 	Step wise progression: <ol style="list-style-type: none"> EtOH -related fatty liver (reversible within 2 weeks of stopping alcohol) Alcoholic hepatitis (reversible with permanent abstinence) Cirrhosis (shrunken liver w/ scar tissue – irreversible) SPECIFIC COMPLICATIONS <ul style="list-style-type: none"> B1 deficiency = Wernicke-Korsakoff syndrome <ol style="list-style-type: none"> EtOH prevents absorption of B1 Wernicke enceph (high mortality rate) = confusion, oculomotor disturbance, ataxia Korsakoff syndrome (irreversible) = retrograde and anterograde amnesia, confabulation, behaviour change Alcohol withdrawal <ol style="list-style-type: none"> 6-12 hrs = tremor, sweat, craving 12-24 hrs = hallucinations 24-48 hrs = seizures 24-72 hrs = delirium tremens (Emergency!!) → downregulation of GABA system by chronic EtOH consumption causes upregulated glutamate → acute confusion, agitation, delusions, tremor, tachycardia, HTN, hyperthermia, ataxia, arrhythmias Other complications: <ol style="list-style-type: none"> Pancreatitis Alcoholic cardiomyopathy Alcohol dependence 	Fibrotic liver increase resistance in hepatic vessels causes: <ul style="list-style-type: none"> Portal HTN → variceal bleeding, ascites → SBP Liver dysfn (malnutrition + encephalopathy) Hepato-renal syndrome HCC Child-Pugh score [AA-BEC] – severity of cirrhosis Factors: Ascites, encephalopathy, bilirubin, albumin, coagulopathy <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><34 (<2)</td><td>34-50 (2-3)</td><td>>50 (>3)</td></tr> <tr> <td>Albumin (g/dL)</td><td>>3.5</td><td>2.8-3.5</td><td><2.8</td></tr> <tr> <td>PT time (sec)</td><td><4 or <1.7</td><td>4-6 or 1.7-2.3</td><td>>6 or >2.3</td></tr> <tr> <td></td><td>A</td><td>B</td><td>C</td></tr> <tr> <td>Score</td><td>5-6</td><td>7-9</td><td>10-15</td></tr> <tr> <td>(1-5yr survival)</td><td>95%</td><td>75%</td><td>50%</td></tr> </tbody> </table> MELD score [BIC-Na] – use very 6/12 for comp. cirrhosis Factors: Bilirubin, INR, creatinine, Na + need for dialysis MELD Score = $10 \times ((0.957 \times \ln(\text{Creatinine})) + (0.378 \times \ln(\text{Bilirubin})) + (1.12 \times \ln(\text{INR}))) + 6.43$ <ul style="list-style-type: none"> Assess severity of CLD (3/12 mortality %) to determine need for liver transplant <ul style="list-style-type: none"> >25 (gravely ill) 24 to 19 = difficult transplant 18 to 11 = candidate < 10 (less ill) 	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		A	B	C	Score	5-6	7-9	10-15	(1-5yr survival)	95%	75%	50%
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(1-5yr survival)	95%	75%	50%																																				
Sx	Asymptomatic	<ul style="list-style-type: none"> Jaundice = raised bilirubin +HM = (but may shrink if more cirrhotic) +SM = due to portal HTN → hypersplenism (↓plt) Spider naevi = telangiectasia w/ central arteriole and small vessels radiating away Palmar erythema = hyperdynamic circulation (↑E2) Bruising = abnormal clotting 	<ul style="list-style-type: none"> Gynecomastia and testicular atrophy in males = endocrine dysfunction Ascites = portal HTN Caput medusae = distended periumbilical veins due to portal HTN Asterixis "flapping tremor" – decomp. Liver disease 																																				
Ix	<ul style="list-style-type: none"> LFT = +ALT/AST COAGS = raised PT/INR time Hep B/C serology Autoantibodies – ANA, SMA, AMA, LKM-1 (autoimmune hepatitis, PBC, PSC) Immunoglobulins (autoimmune hepatitis, PBC) Caeruloplasmin (Wilson's) A1-antitrypsin levels Fe studies (hereditary haem) Liver USS / = increased echogenicity to confirm fatty liver NAFLD fibrosis score (algorithm that uses age, BMI, LFTs to rule out fibrosis) Fibroscan (indication of fibrosis) 	<ul style="list-style-type: none"> FBC = raised MCV EUC = deranged (hepatorenal syndrome) LFT = +ALT/AST, +GGT <ul style="list-style-type: none"> AST:ALT ratio > 2 COAGS = raised PT/INR time Liver USS / fibroscan = increased echogenicity Endoscopy = assess and treat oesophageal varices CT/MRI = fatty infiltration of liver, HCC, +HSM, ascites, abnormal BV changes Liver biopsy - is it alcoholic hepatitis or cirrhosis 	<ul style="list-style-type: none"> EUC = hypoNa (fluid retention due to severe liver disease) LFT = often normal (until decompensated cirrhosis) AFP = MARKER FOR HCC Liver USS - nodular liver surface + <ul style="list-style-type: none"> "corkscrew" arteries w/ increased blood flow to compensate for ↓flow enlarged portal vein ascites splenomegaly Liver fibroscan (elasticity of liver) Endoscopy = assess and treat oesophageal varices CT/MRI = fatty infiltration of liver, HCC, +HSM, ascites, Liver biopsy - is it alcoholic hepatitis or cirrhosis 																																				
ALT/AST	Mildly elevated	AST:ALT ratio > 2																																					
ALP/GGT	NOT HELPFUL	NOT HELPFUL	NOT HELPFUL																																				
Mx	<ul style="list-style-type: none"> Weight loss Exercise Stop smoking Stop alcohol consumption T2DM, HC, HTN control 	<ul style="list-style-type: none"> Stop EtOH immediately and permanently Detoxication regime Nutritional support (B1 supp) + high protein diet Manage withdrawal Sx (delirium tremens) <ul style="list-style-type: none"> Benzo = Chlordiazepoxide "librium" PO every 1-4 hrs for 5-7 days IV high-dose B vitamins (pabrinex) Oral thiamine 	General Management <ul style="list-style-type: none"> High protein, low sodium diet MELD score every 6 months Liver USS and AFP levels every 6 months for hepatocellular carcinoma Endoscopy every 3 years in patients <u>without</u> known varices Consideration of a liver transplant Managing complications as below 																																				

MANAGEMENT OF CIRRHOSIS / CHRONIC LIVER DISEASE

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

LIVER CANCER AND LIVER TRANSPLANTS

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

Viral Hepatitis

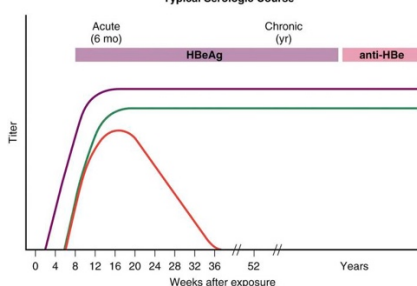
➤ PRACTICE POINT = EBV produces same results +splenomegaly

	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 = vaccinate/protect	PCR TESTING - VIRAL RNA Iseroconvert within 1 st 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 (DAAs) 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

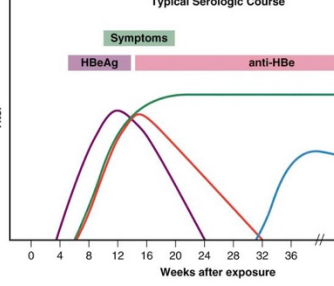
HBsAg or HBV DNA	HBcAb (IgM)	HBcAb (IgG)	HbeAg	HbsAb	Interpretation
					Susceptible to HBV infection
					Acute
					Early acute
			+/-		Chronic
					Resolved acute
					Vaccinated

Progression to Chronic Hepatitis B Virus Infection Typical Serologic Course



— HBsAg — Total anti-HBc — IgM anti-HBc — anti-HBe

Acute Hepatitis B Virus Infection with Recover Typical Serologic Course

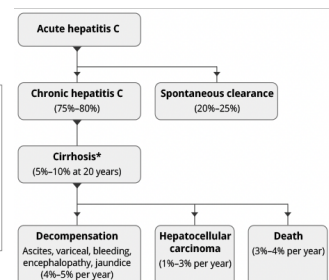
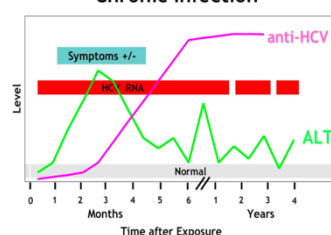


— HBsAg — anti-HBs — IgM anti-HBc — Total anti-HBc

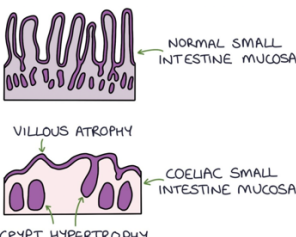
HEP C SEROLOGY

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

Acute HCV Infection Evolving to Chronic Infection



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

GOR: GIT MEDICATIONS:

	Antacids (e.g. Mylanta And Gaviscon)	H ₂ receptor antagonist (Ranitidine (Zantac))	Proton Pump Inhibitors (PPI) E.g.: Omeprazole (Losec), Esomeprazole (Nexium) & Pantoprazole (Somac)
Mechanism of action	<ul style="list-style-type: none"> Base + acid = salt + water Weak bases react with gastric acid to neutralise it → increase pH 	Competitively block H ₂ receptors on parietal cells → reducing gastric acid secretion	<ul style="list-style-type: none"> Irreversibly inactivate the H⁺/K⁺ ATPase enzyme system Suppressing both stimulated and basal acid secretion.
Indications For Use	<ul style="list-style-type: none"> Dyspepsia PUD GORD 	<ul style="list-style-type: none"> Dyspepsia PUD GORD Stress-ulcer prophylaxis 	<ul style="list-style-type: none"> Dyspepsia Peptic ulcer disease (PUD) GORD Zollinger-Ellison syndrome H. Pylori eradication Scleroderma oesophagus Alongside NSAIDs
Adverse effects	<ul style="list-style-type: none"> Diarrhoea (Mg) Constipation (Al and Ca) Belching + flatulence (Ca CO₃) Alkalosis (HCO₃⁻) 	well tolerated. No common AE	Well tolerated BUT common AE are: <ul style="list-style-type: none"> Headache, Nausea, vomiting, rashes Diarrhoea, constipation, flatulence Risk of C. difficile infection, B12 def, OP fractures and pneumonia

H. pylori-related ulcers	NSAIDs related ulcers → do not use for H. pylori
CAUSE: Drugs inhibiting/neutralising gastric acid secretion <ul style="list-style-type: none"> Triple therapy regimen (First-line): PPI + ≥2 anti-bacterials improves eradication rates + reduces subsequent acquired resistance associated with partial treatment) 	CAUSE: Adverse effect of long-term use of NSAIDs <ul style="list-style-type: none"> In the stomach, COX-1 produce prostaglandins (PGE₂ and PGI₂) → stimulate mucus and bicarbonate secretion to cause vasodilation Non-selective NSAIDs <i>inhibit</i> 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** → 1ST LINE = ORAL REHYDRATION

#2: **Treat reversible cause**

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

For uncontrolled (late) diarrhoea:

- Loperamide (2mg after each stool) → opioid agonist → slow GI motility
- Codeine 30mg qid
- ABx if neutropenic or symptoms persist
- Combination
- Specialist advice

#1: **PRIORITY - MAINTAIN HYDRATION** → 1ST LINE = ORAL REHYDRATION

➢ Since most cases of diarrhoea are self-limiting

Drug class	Drugs
Oral rehydration salts	Hydralyte
Anti-diarrhoeals (reduce GIT motility) *Do not give to children!! [CTG guidelines] -ileus!	<ul style="list-style-type: none"> Opioids = Loperamide (Imodium) → poorly crosses BBB Diphenoxylate (with atropine: Lomotil) → combined with anti-cholinergic to avoid abuse of opioids due to their adverse effects
Bulking agents: absorb fluid to reduce diarrhoea	<ul style="list-style-type: none"> Psyllium (Metamucil) ispaghula (Fibogel)
Other	<ul style="list-style-type: none"> Probiotics: conflicting evidence Zinc: children (malnourished)

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 <ul style="list-style-type: none"> Diet IBS-C Colonic inertia Pelvic floor disorders Hirschsprung's 	Mechanical <ul style="list-style-type: none"> Stricture Tumour Prolapse / rectocele Diverticular disease Extrinsic compression 	Metabolic <ul style="list-style-type: none"> Diabetes Ca²⁺, K⁺ Hypothyroidism Hyperparathyroidism Chronic kidney failure

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

Complications

- Faecal impaction = Abdo pain, distension, N/V
- Rectal prolapse
- 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):

- Lifestyle (↑fibre, ↑fluid)
- ↑PA = ↑colonic motility
- Behavioural change = scheduled toilets
- Review meds
- Knees to chest = reduce rectal angle

Mx (meds):

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




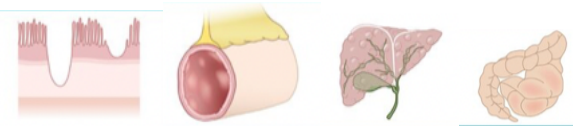
Practice note:

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

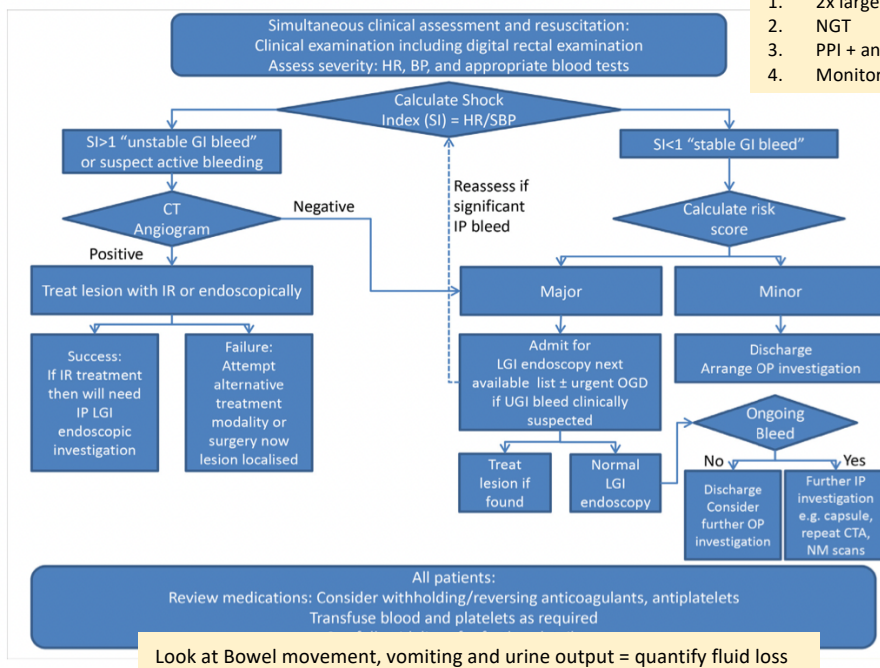
INFLAMMATORY BOWEL DISEASE

	Crohn's	Ulcerative colitis
PP	Mediated by Th1 → IFN γ → macrophages (TNFa) and TH17 cells	Th2 mediated → IL4,13
Location	Usu. terminal ileum (mouth → anus) <ul style="list-style-type: none"> Montreal classification → CD "Age, location, behaviour – inflammatory, structuring, perianal" 	Sigmoid colon → rectum (Montreal classification) <ul style="list-style-type: none"> E1 = proctitis E2 = L sided UC E3 = Pancolitis
RF	<ul style="list-style-type: none"> FHx of autoimmune Genetics (more evident) Smoking (main risk factor) Western diet + stress + high SES 	<ul style="list-style-type: none"> FHx of autoimmune Western diet + stress + high SES Nb: smoking is a protective factor
Clinical Sx	<ul style="list-style-type: none"> Crampy abdominal pain - NOT affected by diet or stress Malaena – watery loose diarrhoea Uncontrolled bowel movements Fever Exhaustion UWL 	MAYO Score <ul style="list-style-type: none"> PR bleed + loose diarrhoea Tenesmus + urgency Chronicity of symptoms New constipation and diarrhoea 
Extra-intestinal manifestations (EIM)	More likely to develop EIM <ul style="list-style-type: none"> Arthritis (symmetrical vs asymmetrical) = 40% Axial arthropathy = Ank spondylitis or sacroiliitis Episcleritis Aphthous ulcers Uveitis Erythema nodosum (painful red papules on shin) 	<ul style="list-style-type: none"> Arthritis (symmetrical vs asymmetrical) = 40% Axial arthropathy = Ank spondylitis or sacroiliitis PSC Pyoderma gangrenosum Less common = episcleritis, aphthous ulcer, erythema nodosum (painful red papules on shin)
Ix + Dx (combination)	<ul style="list-style-type: none"> Clinical Hx Serial CRP and faecal calprotectin Endoscopy / colonoscopy (Definitive diagnosis) Barium X-ray = string sign (CD), lead pipe colon (UC) Histo findings <hr/> Acute flare: <ul style="list-style-type: none"> CRP > 30 Hb <100 Raised remp 	DDx: <ul style="list-style-type: none"> IBD (UC, CD) Coeliac PUD, GORD IBS Colitis (ischaemic, infectious, RT, microscopic, TB-induced, NSAID) <div style="border-left: 1px solid black; padding-left: 10px;"> <ul style="list-style-type: none"> Food allergy (FPIES, cow's milk protein) Giardiasis Appendicitis </div>
Histo-specific Findings	<ul style="list-style-type: none"> Transmural inflammation (creeping fat) Skip lesions "cobblestoning" Granulomatous non-caseating inflammation (XS giant cells and lymphoid aggregates + ↑ goblet cells) ASCA (yeast) antibodies 	<ul style="list-style-type: none"> Friable Submucosal inflammation → anal region sparing Continuous lesions Less goblet cells Crypt abscess P-ANCA +ve 
Comp.	<ul style="list-style-type: none"> Fistulas (due to adhesions to viscera) <ul style="list-style-type: none"> Enterourinary (UTI, air bubbles in urine) Entero-vaginal Enterocutaneous Strictures / obstructions Fissures Nephrolithiasis 	<ul style="list-style-type: none"> Toxic megacolon → perforation → peritonitis <ul style="list-style-type: none"> Subtotal colectomy + ileostomy +/- Hartmann's pouch Haemorrhage /VTE CRC PSC → cholangiosarcoma "Coursevier's signs"
Medical Treatment	<ol style="list-style-type: none"> Prednisolone (oral pred or IV hydrocortisone) DMARDS (mod) <ol style="list-style-type: none"> Azathioprine (thiopurine) - also used in RA, ALL → A/E = thrombocytopenia, liver, pancreatitis, ↑ lymphoma MTX (add folic acid) Biologic (severe) <ol style="list-style-type: none"> Infliximab (Anti-TNF) for acute flares + mod-severe disease not responding to immunomodulators Anti-IL12,23 (ustekinumab) also for psoriasis Vedoluzimab (anti-a4b7 intregin) 	Inducing remission <ol style="list-style-type: none"> DMARDS (mod) <ol style="list-style-type: none"> Sulfasalazine Azathioprine (thiopurine) Mercaptopurine Biologic (severe) <ol style="list-style-type: none"> Infliximab (Anti-TNF) for acute flares + mod-severe disease not responding to immunomodulators JAK inhibitor (tofacitinib) Vedoluzimab (anti-a4b7 intregin) Maintaining remission: <ol style="list-style-type: none"> Aminosalicyclate (e.g. mesalazine oral or PR) Azathioprine or mercaptopurine Clexane = ↓ clot risk
Surgery	Only 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"> Abx – metronidazole (Flagyl) + cipro Biologic screen: <ul style="list-style-type: none"> Viral serology (Hep B/C /HIV/ CMV/EBV/VZV) MMR IgG +/- TB quantiferon gold ANCA/ASCA TPMT (myelosuppression due to thiopurine usage) → if low TPMT → lower dosage required Fe, folate, B12 and vit D Endoscopy 6-9 months after initial treatment MDT support to manage EIM – Physio, rheumatologist, gastroenterologist No live vaccines (MMR, yellow fever) while on immunosuppressants 	

APPROCH TO GI BLEEDING

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

Management of Acute LOWER GI bleed



Immediate Mx:

- 2x large bore cannula
- NGT
- PPI + anti-emetic
- Monitor UO

1st 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**

2nd 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)

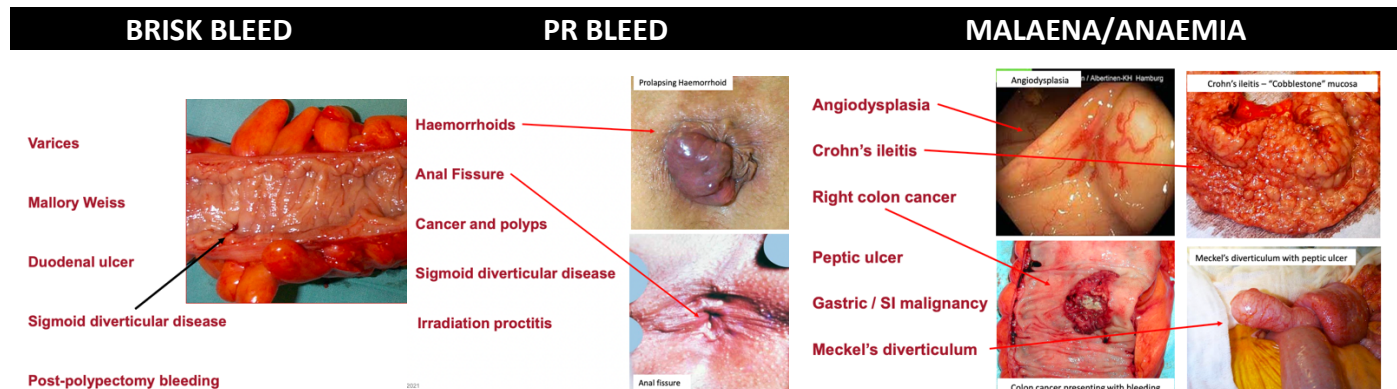
Clinical Setting		Action
INR	Sig. Bleeding	
High but <4.5	No	Lower dose/omit warfarin
5-9	No	If high bleed risk give vit. K
>9	No	Low risk – vit. K High risk – PTX
Any	Yes	Vitamin K + PTX +/- FFP

IP, inpatient; IR, interventional radiology;

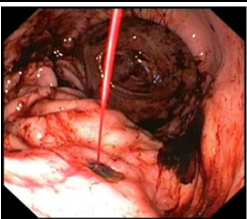
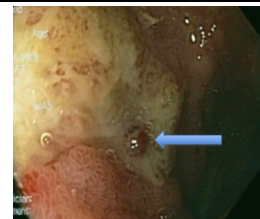

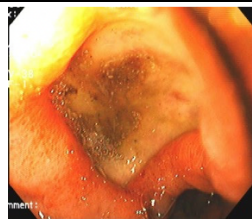

OGD, oesophagogastrroduodenoscopy; OP, outpatient

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">1/3 = cirrhosis → 1/3 fatal → 1/3 = survive (> 1 year)
Risk factors	<ul style="list-style-type: none">Use of NSAIDs and low-dose aspirin (LDA)H. pylori infection	Predictors of <u>large</u> oesophageal varices <ul style="list-style-type: none">Severity of liver disease (Child-Pugh Classification)Hypersplenism (splenomegaly) → Thrombocytopenia < 100 x10⁹ /L
Mx	Pre-endoscopic <ul style="list-style-type: none">Resus + ABCD (blood transfusion if Hb < 70 + HypoTN)<u>Risk stratification</u> Glasgow Blatchford<ul style="list-style-type: none">< 1 = outpt endoscopy> 6 = need urgent endoscopyPPI (8mg/hr) → ↑ clot formation + "downstages" lesion) <u>BUT</u> does NOT reduce bleeding or mortalityErythromycin = prokinetic agentIf cirrhosis → vasoactive drugs and ABxSTOP anti-coag or reverse (e.g. PTX for warfarin)	Vasoconstrictor therapy <ul style="list-style-type: none">➤ Octreotide (somatostatin analogue) → reduce splanchnic blood flow➤ <u>Terlipressin/somatostatin</u>➤ <u>Vit K/FFP</u>
		Antibiotics Prophylactic broad spectrum ABx→sig. reduces early rebleeding + infection/sepsis
		Resuscitation Minimise rapid expansion → can remove clot IVF bolus beware!!!
		ICU level care
		Endoscopy (OGD) Within 12 hrs + ETT required (band ligation preferred)
		Alternatives <ul style="list-style-type: none">Early placement <u>within 24-72 hrs</u> of TIPS (Transjugular Intrahepatic Portosystemic Shunt)SB Tube [Provide time for TIPS] ➔ Gastric balloon for immediate temporary control BUT high complication rate (e.g. necrosis, oesophageal perforation)
		Beta Blockade (prevention) <ul style="list-style-type: none">Non-selective BB (propranolol) → ↓↓CO, reduces splanchnic vasoconstrict (i.e. vasodilate) = ↓↓ blood flow portal veinReduce risk for recurrent variceal haemorrhage
	Early Endoscopy <ul style="list-style-type: none">Within 24 hours if <u>acute UGIB</u><u>Indications:</u> Ulcers w/ active bleeding or non-bleeding visible vesselsInjection therapy (e.g. adrenaline) – temporary measure →still need to stop rebleeding w/ endoscopy and embolisationThermal probes, and clipsOesophageal Variceal bleeds → tissues gluesRefractory bleeding → TIPS	
	Post-endoscopy [NOT Dying > NOT rebleeding] <ul style="list-style-type: none">Check <u>H. pylori status</u> in all ulcer patients<u>High ulcer risk</u> = High-dose PPI for 72 h<u>low risk ulcers</u> = fed promptly+ oral PPI<u>Cirrhosis pts</u> = ABx for 7 days (regardless of bleeding)<u>Variceal bleed</u> = vasoactive drugs for 5 daysRestart low dose aspirin in CV-patients once bleeding resolved	

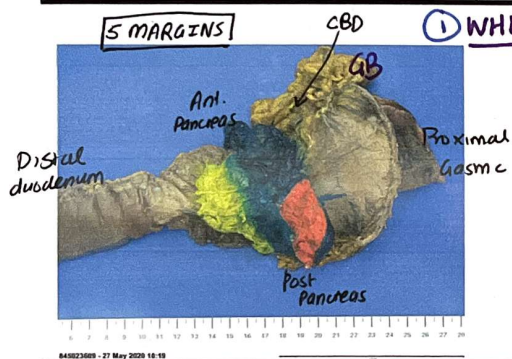


SURGICAL IMAGES OF ACTIVE VS INACTIVE BLEEDS

Major stigmata			Minor stigmata	
o High-bleeding risk endoscopic therapy			o Low rebleeding risk – NO endoscopic therapy needed	
				
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 – spurt FORREST 1B – oozing	FORREST 2A	FORREST 2B	FORREST 2C	FORREST 3

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

HPS	Examination	
<ul style="list-style-type: none">56 year old male Presents with abdominal pain<ul style="list-style-type: none">Localised → parietal peritoneumPoorly localised → visceral peritoneum/organsPain is constant, not associated with eating → exc. <u>GORD, achalasia</u>Radiates to the back → <u>pancreatitis, AAA, gastritis (PUD), costochondritis?</u>Sudden onset 3 hours agoHas been progressively becoming worseHas noticed a dull epigastric pain for many weeks prior which he put down to refluxNot associated with N&V or diarrhoea → exc. impaired gastric emptyingNo fever, night sweats → exc. <u>infection, lymphoma</u>No mucous, melaena or blood in the stool → exc. <u>cancer?, Upper /Lower GI bleed</u>Is passing wind → <u>important to exc. bowel obstruction</u>Not constipated (last past stool yesterday, appeared normal)Has lethargy8kg weight loss in last 6 months → <u>cancer? Parasitic organisms – tapeworm?</u>No rashes but has intensely itchy skin → <u>suggests cholestasis</u> → jaundice /scleral icterus	<ul style="list-style-type: none">Appears cachecticHas palmar erythemaScleral icterus → <u>yellow eyes (Hepatic dysfunction)</u><ul style="list-style-type: none">Best inspected under natural light (by window)Mouth normalUmbilical hernia noted with ascites	<ul style="list-style-type: none">No caput medusa or spider naevi → <u>No portal HTN</u>Liver enlarged and tender to palpation → <u>hepatomegaly</u>Epigastrium tenderNo guardingTestes normal → exc. <u>testicular torsion</u>DRE normal
	Investigations	
	<ul style="list-style-type: none"><u>FBC</u> – WCC raised → check <u>infection</u><u>EUC</u> – normal → check <u>renal</u> function<u>LFT</u> – GGT and LDH high → check <u>liver</u> dysfunction<u>COAGS</u> – normal → <u>liver dysfunction</u> → <u>loss of vitamin K derived clotting factors</u><u>Lipase</u> - High → check pancreas<u>Amylase</u> – High → check pancreas<u>CEA (carcinoembryonic antigen)</u> – High<ul style="list-style-type: none">sign of certain cancers (e.g. colon and rectum, prostate, ovary, lung, thyroid, or liver)may also be a sign of some noncancerous conditions, such as cirrhosis, noncancerous breast disease, and emphysema<u>CA19.9 (Carbohydrate antigen)</u>- High<ul style="list-style-type: none">differentiate cancer of the pancreas or bile ducts and other conditions OR monitor response to pancreatic cancer treatment and watch for recurrence	
	Detailed investigations	
	<ul style="list-style-type: none"><u>CT abdomen</u> – liver, kidneys and spleen<u>Endoscopic-ultrasound-guided FNA</u> (minimise excision – as pancreas is highly vascular endocrine)	



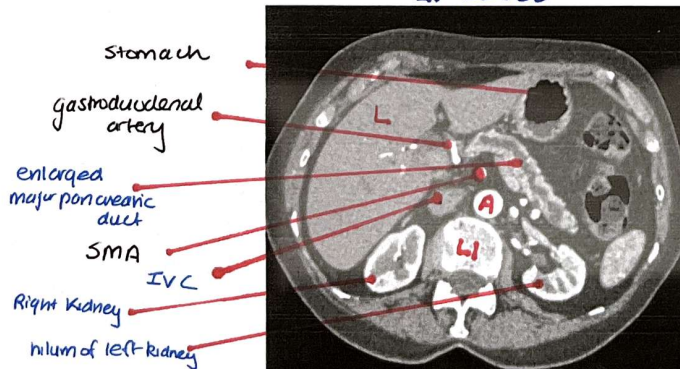
- Red** = pancreatic neck region
- Green-blue** = portal vein bed
- Yellow** = peri-uncinate soft tissue (lymph nodes)

① WHIPPLES PROCEDURE (ONLY FEW PANCREATIC Ca RESECTABLE!)

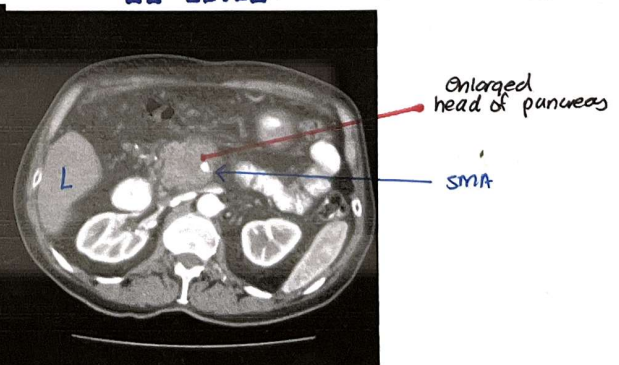


- Mucosal ulceration at head of pancreas (*most common*)
- ② ENDOSCOPIC STENT – SX MIX FOR JAUNDICE** (= AC)

L1 LEVEL



L2 LEVEL

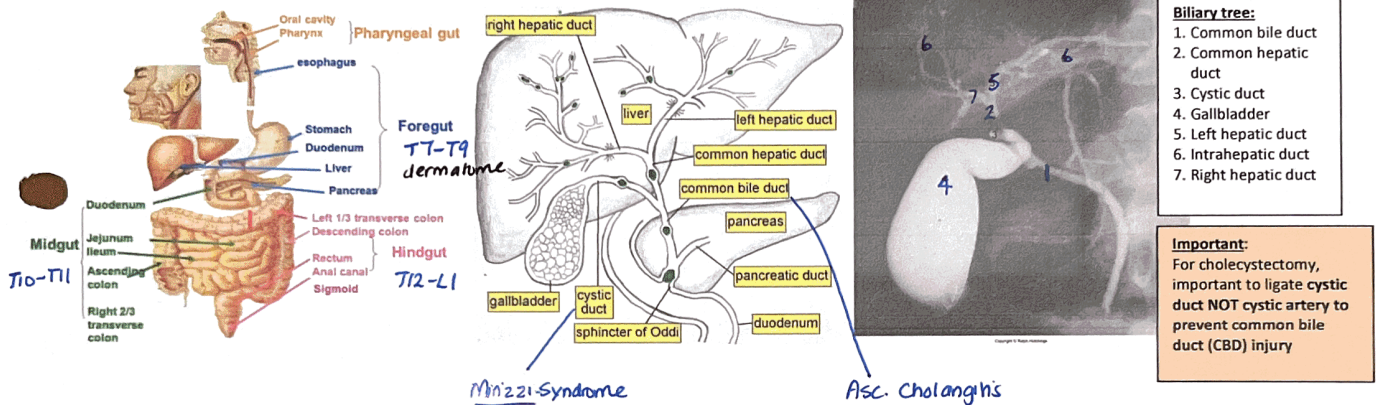


Interpretation: 40mm head of pancreas mass with **double duct sign** = simultaneous dilatation of CBD + pancreatic duct

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

DDx for pancreatic mass causes	Final Dx:	Treatment MOST UNRESECTABLE
<ul style="list-style-type: none"> 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) → ↑ amylase lipase Lymphoma Adenocarcinoma of duodenum Vascular lesion (thrombus in SMV) 	<ul style="list-style-type: none"> Pancreatic ductal adenocarcinoma Involves portal vein bed PNI and LVI (i.e. perineural and lymphovascular invasion) Metastatic carcinoma present in umbilical hernia <p>Indicates stage IV</p>	<ul style="list-style-type: none"> Completion pancreatectomy — depends if margins have been exceeded FOLFOX chemotherapy → consider severe debilitation on patient (tox'city) Radiotherapy? → relatively easier as retroperitoneal location (for pancreas exc. tail) Palliation (if all fails) Poor prognosis → 7.1% 5-year survival rate

Describe anatomy & relationships of organs derived from embryonic foregut



Retroperitoneal	Intraperitoneal	Portosystemic anastomoses
<ul style="list-style-type: none"> Supra-adrenal Aorta/IVC Duodenum (2nd and 3rd part) Pancreas (exc. tail) Ureter Ascending and descending colon Kidneys Esophagus Rectum <p>SADPUCKER</p>	<ul style="list-style-type: none"> Liver Stomach Jejunum and ileum Spleen TC SC Upper rectum 	<p>Affected during portal HTN (e.g. caused by cirrhosis (liver scar tissue) leading to increased blood flow resistance:</p> <ul style="list-style-type: none"> Backflow into left gastric vein causing dilatation of oesophageal veins resulting in oesophageal varices (fragile to rupture → haematemesis) [VARICEAL BLEED] <ul style="list-style-type: none"> Oesophageal veins → hemiazygos → azygos → SVC → RA Caput medusae (varices in umbilical vein) Haemorrhoids (varices in superior rectal vein that drain into IVC)

Vascular supply				Nerve supply		
	Foregut	Midgut	Hindgut	PSNS	SNS	
BV	Coeliac trunk	SMA	IMA	<ul style="list-style-type: none">• Vagus nerve (duo to mid-TC)<ul style="list-style-type: none">◦ Left (hepatic branch) anterior to stomach◦ Right (coeliac branch): posterior to stomach• Pelvic splanchnic – S2-S4 (mid-TC – anal canal)	Thoracolumbar SNS ganglia (T5-L2) <ul style="list-style-type: none">• Greater splanchnic: T5-T9• Lesser splanchnic = T10-11• Least splanchnic = T12-L2	
Level	T12	L1	L3			
Somatic pain region	T7-T9 (epigastric)	T10-T11 (umbilical)	T12-L1 (suprapubic)			

