

CLINICAL GUIDE TO

Gastroenterology

Edited by
Yang Chen
Maxton Pitcher

Series Editor:
Christian Fielder Camm



WILEY Blackwell

Clinical Guide to
Gastroenterology

The *Clinical Guides* series

Series Editor: Christian Fielder Camm

The *Clinical Guides* are a brand new resource for junior doctors and medical students. They provide practical and concise information on symptoms, common conditions, and day-to-day problems faced in the clinical environment. They are easy to navigate and allow swift access to information as it is needed, with step-by-step guidance on investigations, decision-making and interventions, and how to survive and thrive on clinical rotations and attachments.

Clinical Guide to Gastroenterology

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Acronyms and Abbreviations

2WW	Two-week wait	CDAD	<i>C. difficile</i> -associated diarrhoea
5-FU	5-fluorouracil	CD117	Cluster of differentiation 117
5-HIAA	5-hydroxyindoleacetic acid	CD8+	Cluster of differentiation 8
5-HT ₄	5-hydroxytryptamine-4 receptor	CDT	<i>Clostridium difficile</i> toxin
5-ASA	5-aminosalicylic acid	CEA	Carcinoembryonic antigen
A&E	Accident and emergency	CFTR	Cystic fibrosis transmembrane conductance regulator
AAA	Abdominal aortic aneurysm	CgA	Chromogranin A
Ab	Antibody	CIWA	Clinical Institute Withdrawal Assessment for Alcohol
ABG	Arterial blood gas	CKD	Chronic kidney disease
ACE	Angiotensin converting enzyme	CLD	Chronic liver disease
ACS	Acute coronary syndrome	CLO	Campylobacter-like organism
ACTH	Adrenocorticotrophic hormone	CMV	Cytomegalovirus
ADP	Adenosine diphosphate	CNS	Central nervous system
ADPKD	Autosomal dominant polycystic kidney disease	COPD	Chronic obstructive pulmonary disease
AF	Atrial fibrillation	Cr	Creatine
AFB	Acid-fast bacillus	CRP	C-reactive protein
AFP	α -fetoprotein	CRT	Capillary refill time
AIDS	Acquired immune deficiency syndrome	CSF	Cerebrospinal fluid
AIH	Autoimmune hepatitis	CT	Computed tomography
AKI	Acute kidney injury	CTAP	CT abdomen/pelvis
ALD	Alcoholic liver disease	CT CAP	CT chest/abdomen/pelvis
ALF	Acute liver failure	CTPA	Computed tomography pulmonary angiography
ALP	Alkaline phosphatase	CT PET	Computed tomography positron emission tomography
ALT	Alanine transaminase	CTVC	Computed tomography virtual colonoscopy
AMA	Antimitochondrial antibody	CXR	Chest X-ray
AMTS	Abbreviated Mental Test Score	CYP2E1	Cytochrome P450 2E1
ANA	Antinuclear antibody	D&V	Diarrhoea and vomiting
Anti-AChR	Antibodies to acetylcholine receptors	DAAAs	Directly acting antivirals
APTT	Activated partial thromboplastin time	DEXA	Dual-energy x-ray absorptiometry
ARB	Angiotensin receptor blocker	DIC	Disseminated intravascular coagulation
ARDS	Acute respiratory distress syndrome	DKA	Diabetic ketoacidosis
ASMA	Anti-smooth muscle antibody	DNA	Deoxyribonucleic acid
AST	Aspartate transaminase	DRE	Digital rectal examination
AVM	Arterio-venous malformation	dsDNA	Double-stranded deoxyribonucleic acid
AXR	Abdominal X-ray	DVT	Deep vein thrombosis
β -HCG	β -human chorionic gonadotrophin	DWI	Diffusion weighted imaging
BM	Capillary blood glucose	<i>E. coli</i>	<i>Escherichia coli</i>
BMI	Body mass index	EATL	Enteropathy-associated T-cell lymphoma
BP	Blood pressure	EBV	Epstein-Barr virus
bpm	Beats per minute	ECG	Electrocardiogram
BPPV	Benign paroxysmal positional vertigo	ECHO	Echocardiogram
<i>C. difficile</i>	<i>Clostridium difficile</i>	EEG	Electroencephalograph
Ca125	Cancer antigen 125	eGFR	Estimated glomerular filtration rate
CA-19-9	Cancer antigen 19-9	EGFR	Epidermal growth factor receptor
CBD	Common bile duct	EMA	Anti-endomyosial antibody
CBT	Cognitive behavioural therapy	EMR	Endoscopic mucosal resection
cCa	Corrected calcium level		
CCF	Congestive cardiac failure		
CCK	Cholecystokinin		

ENA	Extrinsic nuclear antigen	ICD	International Classification of Disease
ENT	Ear, nose and throat	ICP	Intracranial pressure
ERCP	Endoscopic retrograde cholangiopancreatography	ICV	Ileo-caecal valve
ESR	Erythrocyte sedimentation rate	IgA	Immunoglobulin A
ESWL	Extracorporeal shock-wave lithotripsy	IgG	Immunoglobulin G
FAP	Familial adenomatous polyposis	IgG1	Immunoglobulin G1
FAST	Focused assessment with sonography for trauma	IgG4	Immunoglobulin G4
FBC	Full blood count	IgM	Immunoglobulin M
FNA	Fine needle aspirate	IHD	Ischaemic heart disease
FOB	Faecal occult blood	IL-12	Interleukin 12
FODMAP	Fermentable oligosaccharides, disaccharides, monosaccharides, and polyols	IL-23	Interleukin 23
G6PDH	Glucose-6-phosphate dehydrogenase	IM	Intramuscular
GA	General anaesthetic	IMA	Inferior mesenteric artery
GABA	γ -amino butyric acid	INR	Internationalised normalised ratio
GAVE	Gastric antral vascular ectasia	ITU	Intensive care unit
GBS	Guillain-Barré syndrome	IV	Intravenous
GCS	Glasgow coma score	IVDU	Intravenous drug user
GDH	Glutamate dehydrogenase	JVP	Jugular venouse pressure
GGT	γ -glutamyl transferase	LBO	Large bowel obstruction
GFD	Gluten-free diet	LDH	Lactate dehydrogenase
GFR	Glomerular filtration rate	LFT	Liver function test
GI	Gastrointestinal	LIF	Left iliac fossa
GIST	Gastrointestinal stromal tumour	LKM	Liver, kidney, muscle
GOJ	Gastro-oesophageal junction	LMWN	Low molecular weight heparin
GORD	Gastro-oesophageal reflux disease	LOS	Lower oesophageal sphincter
GMC	General Medical Council	LP	Lumbar puncture
GP	General practitioner	LUQ	Left upper quadrant
<i>H. pylori</i>	<i>Helicobacter pylori</i>	MALT	Mucosa-associated lymphoid tumour
HAS	Human albumin solution	MC&S	Microscopy, culture, and sensitivity
HAV	Hepatitis A virus	MCH	Mean corpuscular haemoglobin
Hb	Haemoglobin	MCV	Mean corpuscular volume
HbA1c	Glycated haemoglobin	MDCT	Multi-detector CT
HBc	Hepatitis B core	MDT	Multi-disciplinary team
HBcAg	Hepatitis B core antigen	MELD	Model for End-Stage Liver Disease
HBeAg	Hepatitis B e antigen	MEN1	Multiple endocrine neoplasia type 1
HBs	Hepatitis surface	MEN2	Multiple endocrine neoplasia type 2
HBsAg	Hepatitis surface antigen	MI	Myocardial infarction
HBV	Hepatitis B virus	MMSE	Mini mental status examination
HCC	Hepatocellular carcinoma	MR	Mitral regurgitation
HCV	Hepatitis C virus	MRCP	Magnetic resonance cholangiopancreatography
HDU	High-dependency unit	MRI	Magnetic resonance imaging
HDV	Hepatitis C virus	MSK	Musculoskeletal
HEV	Hepatitis E virus	MUSK	Muscle specific kinase
HFE	Haemochromatosis gene	M-W	Mallory-Weiss
HHS	Hyperglycaemic hyperosmolar state	MYH	MYH gene associated with polyposis
HIV	Human immunodeficiency virus	NAC	N-acetylcysteine
HLA-DQ2	Human leucocyte antigen-DQ2	NAFLD	Non-alcoholic fatty liver disease
HNPPC	Hereditary non-polyposis colorectal cancer	NAPQI	N-acetyl-p-benzoquinone imine
HR	Heart rate	NASH	Non-alcoholic steatohepatitis
HRS	Hepatorenal syndrome	NBM	Nil by mouth
HSV	Herpes simplex virus	NET	Neuroendocrine tumour
HTN	Hypertension	NEWS	National early warning score
HUS	Haemolytic uremic syndrome	NG	Nasogastric
IBD	Inflammatory bowel disease	NHS	National Health Service
IBS	Irritable bowel syndrome	NICE	National Institute for Health and Care Excellence
		NKA	Neurokinin A

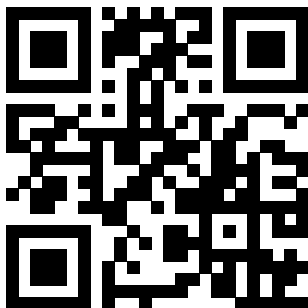
NNT	Number needed to treat	SIRS	Systemic inflammatory response syndrome
NSAIDs	Non-steroidal anti-inflammatory drugs	SLE	Systemic lupus erythematosus
NSTEMI	Non-ST elevation myocardial infarction	SLT	Speech and language therapy
OCP	Oral contraceptive pill	SMA	Superior mesenteric artery
ODP	Operating department practitioner	SPECT	Single-photon emission computed tomography
OGD	Oesophagoduodenoscopy	SPINK1	Serine protease inhibitor kazal type 1
OGJ	Oesophagogastric junction	SpO ₂	Peripheral capillary oxygen saturation
OPG	Orthopantomogram	SpR	Specialty registrar
PaO ₂	Partial pressure of oxygen	sO ₂	Saturation of oxygen level
PBC	Primary biliary cholangitis	SOB	Shortness of breath
pCO ₂	Partial pressure of carbon dioxide	SSRIs	Selective serotonin-reuptake inhibitors
PCI	Percutaneous coronary intervention	STEMI	ST elevation myocardial infarction
PCR	Polymerase chain reaction	STI	Sexually transmitted infection
PE	Pulmonary embolus	SVC	Superior vena cava
PEG	Percutaneous endoscopic gastrostomy	SVT	Supraventricular tachycardia
PET	Positron emission tomography	TNM staging	Tumour-size, lymph node disease, metastases (convention of cancer staging)
PHE	Public Health England	T2DM	Type 2 diabetes mellitus
PICC	Peripherally inserted central catheter	TB	Tuberculosis
Plts	Platelets	TDS	Three times a day
PMHx	Past medical history	TFT	Thyroid function test
PMN	Polymorphonuclear leukocyte	TIPS	Trans-jugular intrahepatic portosystemic shunting
PP	Pancreatic polypeptide	TNF- α	Tumour necrosis factor alpha
PPI	Protein pump inhibitor	TPMT	Thiopurine methyltransferase
PPM	Permanent pacemaker	TPN	Total parenteral nutrition
PR	Per rectum	TSH	Thyroid stimulating hormone
PRN	Pro re nata (defined as 'when needed')	TTG	Tissue transglutaminase
PSA	Prostate-specific antigen	UC	Ulcerative colitis
PSC	Primary sclerosing cholangitis	UDCA	Ursodeoxycholic acid
PT	Prothrombin time	U&Es	Urea and electrolytes
PTC	Percutaneous trans-hepatic cholangiogram	UGI	Upper gastrointestinal
PUD	Peptic ulcer disease	UGIB	Upper gastrointestinal bleeding
PV	Per vaginum	UKELD	United Kingdom Model for End-Stage Liver Disease
RAST	Radioallergosorbent test	ULN	Upper limit of normal
RCT	Randomised controlled trial	UOS	Upper oesophageal sphincter
RDW	Red cell distribution width	US	Ultrasound
RF	Radiofrequency	UTI	Urinary tract infection
RhF	Rhesus factor	VBG	Venous blood gas
RIF	Right iliac fossa	VEGF	Vascular endothelial growth factor
RLQ	Right lower quadrant	VKA	Vitamin K antagonist
RNA	Ribonucleic acid	VT	Ventricular tachycardia
RR	Respiratory rate	VTE	Venous thromboprophylaxis
RUQ	Right upper quadrant	VTEC	Verotoxigenic <i>Escherichia coli</i>
SAAG	Serum-ascitic albumin gradient	VZV	Varicella zoster virus
SaO ₂	Saturation of oxygen in arterial blood	WBC	White blood cells
SBAR	Situation background assessment recommendation	WCC	White cell count
SBO	Small bowel bacterial overgrowth	WHO	World Health Organization
SBP	Spontaneous bacterial peritonitis		
SC	Subcutaneous		
SHO	Senior house officer		
SIR	Systemic inflammatory response		

Podcast and Box Icons

Clinical	
Differential	
Scales and scores	
How to	
Podcast	

About the Companion Website

This book is accompanied by a companion website:



www.wiley.com/go/chen/gastroenterology

The website includes:

- Audio Clips
- Audio Scripts
- Self Assessment Questions
- Clinical Cases
- Online Chapters and Sections

Due to space constraints, some smaller chapters have been moved from the book to the companion website. The companion website also contains a large number of self-assessment questions in MCQ, EMQ, and SAQ format. Additionally, there are a number of clinical cases to work through.

1.1 Examination Technique

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1.1.1 CHAPTER AT A GLANCE

Box 1.1.1 Chapter at a Glance

- The abdominal examination is complex and requires practice to master
- Significant features suggestive of chronic abdominal disease are available in the peripheries
- Systematic examination of the abdomen is important to not miss findings

1.1.2 COMMON CONDITIONS

- Chronic liver disease (CLD) 
- Alcohol withdrawal 
- Viral hepatitis 
- Inflammatory bowel disease (IBD) 
- Malignancy 

1.1.3 CLINICAL EXAMINATION – PERIPHERIES

Table 1.1.1 Elements to be undertaken prior to examining the patient

Item	Detail
1. Appropriate hand hygiene	Wash hands with soap and water or alcohol hand rub
2. Introduce yourself	Full name and job title
3. Confirm patient's identity	Confirm name and date of birth, verify against wristband
4. Gain consent	Explain your role, the examination, and why you are performing it
5. Maintain dignity and comfort	Ensure that you are in a well-lit and private area where you will not be disturbed
6. Ensure a chaperone when needed	Explain reasoning and ensure the patient is happy with this
7. Be mindful of personal and cultural boundaries	Be respectful and considerate during the examination
8. Position the patient	Initially conducted with the patient at 45° On moving to the abdomen, patient should be supine
9. Expose the patient appropriately	Ideally, from the xiphoid process to the symphysis pubis. Can be done after peripheral examination

1. Chronic liver disease (CLD) ; 2. Alcohol withdrawal ; 3. Viral hepatitis ; 4. Inflammatory bowel disease (IBD) ; 5. Malignancy 

Box 1.1.2 The Use of Examination to Aid History

- Patients will often give additional history while you are examining them
- Often the silences during examination give patients time to think about things they may have forgotten to tell you previously
- Although you may be concentrating on examination, be interested in what they are saying and listen carefully

**Table 1.1.2** Examination features from the end of the bed

Item	Comments
1. Does the patient look well?	• Aggressive/agitated • Lethargic/drowsy
2. Are they alert and orientated?	• Reduced Glasgow coma score (GCS) is present in shock • Confusion (■ / ■) • Agitation (■)
3. Body mass index (BMI)	• Obesity • Underweight, e.g. malabsorption (■), cachexia (■)
4. Presentation	• Unkempt – social or mental health issues
5. Lines in and out of patient	• IV infusions • Catheters • Oxygen • Nasogastric (NG) tube
6. Patient monitoring	• Observation chart • Continuous ECG monitoring • Haemodynamic monitoring
7. Look around the patient	• Eating and drinking • Vomit bowl • Medications

Table 1.1.3 Examination findings in the hands and limbs

Item	Conditions
1. Tremor	■
2. Hepatic flap	■
3. Sweating	Sepsis / ■ / Illicit substance / ■
4. Clubbing	■ / ■ / ■
5. Palmar erythema	■
6. Dupuytren's contracture	■
7. Discoloured skin	Haemochromatosis/Addison's disease
8. Signs of intravenous drug use	■
9. Signs of arthritis	■
10. Ankle oedema	Hypoalbuminaemia / ■

1. Chronic liver disease (CLD) ■; 2. Alcohol withdrawal ■; 3. Viral hepatitis ■; 4. Inflammatory bowel disease (IBD) ■; 5. Malignancy ■

Image				
Nail Change	Koilonychia	Leukonychia	Beau's lines	Clubbing
Associated disorders	Iron deficiency anaemia	Hypoalbuminaemia	Flare of IBD GI malignancy Acute GI infection	Cirrhosis GI malignancy IBD Chronic infection

Figure 1.1.1 Nail changes in GI disease.

Image	Condition	Related disorder
	Palmar erythema	Liver disease
	Psoriatic patches – note typical silver scaling	Extra-intestinal IBD association
	Spider naevus	Chronic liver disease, especially alcohol (>5 spider naevi is pathological)
	Erythema nodosum (multiple lesions on legs)	IBD
	Bruising/thin skin	Steroid treatment

Figure 1.1.2 Skin changes in GI disease.

Table 1.1.4 General inspection – face and neck

Item	Conditions
1. Scleral icterus	■ / ■
2. Xanthelasma	Associated with primary biliary cholangitis
3. Kayser–Fleisher rings	Wilson’s disease
4. Anterior uveitis/scleritis	■
5. Mouth ulcers	■
6. Raw, erythematous tongue	Vitamin B12 deficiency
7. Atrophic glossitis/angular stomatitis	Iron deficiency
8. Lymphadenopathy (supraclavicular/axillary/Virchow’s)	■

Table 1.1.5 General inspection – chest

Item	Conditions
1. Spider naevi	■
2. Muscle wasting of thorax	Type 1 diabetes / ■ / ■
3. Gynaecomastia	■
4. Loss of body hair	■ / Hypogonadism or haemochromatosis

1.1.4 CLINICAL EXAMINATION – ABDOMEN

- Anatomically the abdomen can be divided into nine regions (Figure 1.1.3)

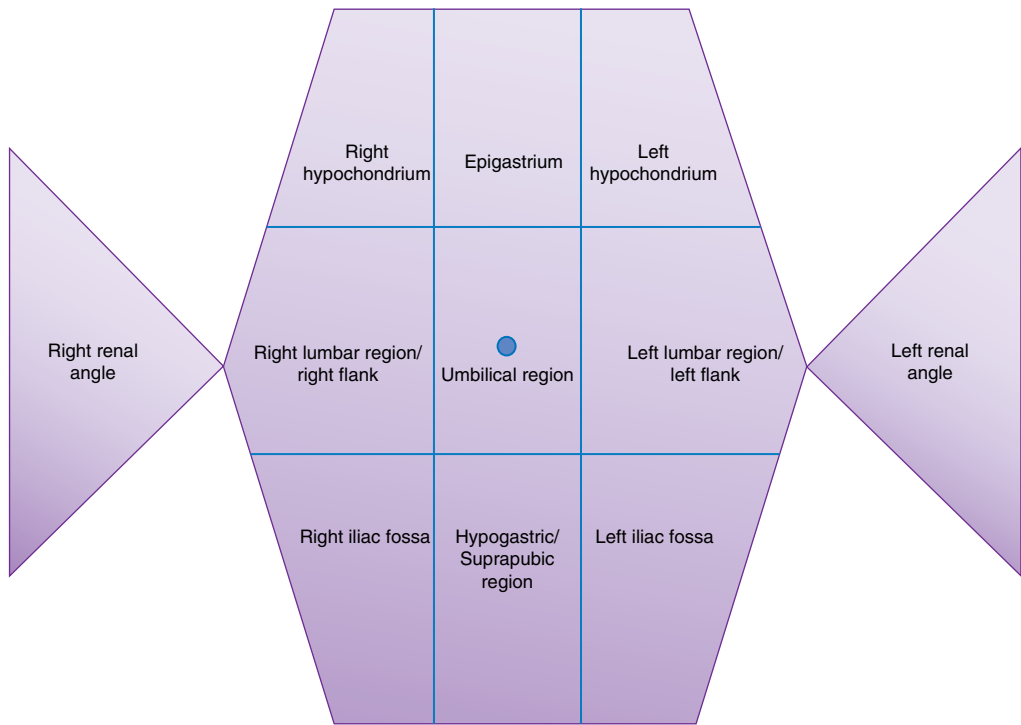


Figure 1.1.3 Division of the abdomen.

1. Chronic liver disease (CLD) ■; 2. Alcohol withdrawal ■; 3. Viral hepatitis ■; 4. Inflammatory bowel disease (IBD) ■; 5. Malignancy ■

Box 1.1.3 The 6 Fs of Distension



- Fat (obesity)
 - Faeces (constipation, obstruction)
 - Fluid (tumours, ascites, palpable bladder)
- Foetus (pregnancy)
 - Flatus (obstruction or pseudo-obstruction)
 - Function (irritable bowel syndrome, IBS)

Table 1.1.6 Inspection features on the abdomen

Item	Conditions
1. Scars	See Figure 1.1.4
2. Visible masses	Hernia / ■ / ■
3. Stoma sites	Ileostomy/colostomy (see Figure 1.1.5)
4. Hernias	Inguinal/femoral/umbilical/para-umbilical/epigastric (see Boxes 1.1.10 and 1.1.11)
5. Striae/bruising	■
6. Scratch marks	■
7. Caput medusa	■
8. Ask patient to lift head from the pillow	Sensitive sign for peritonism

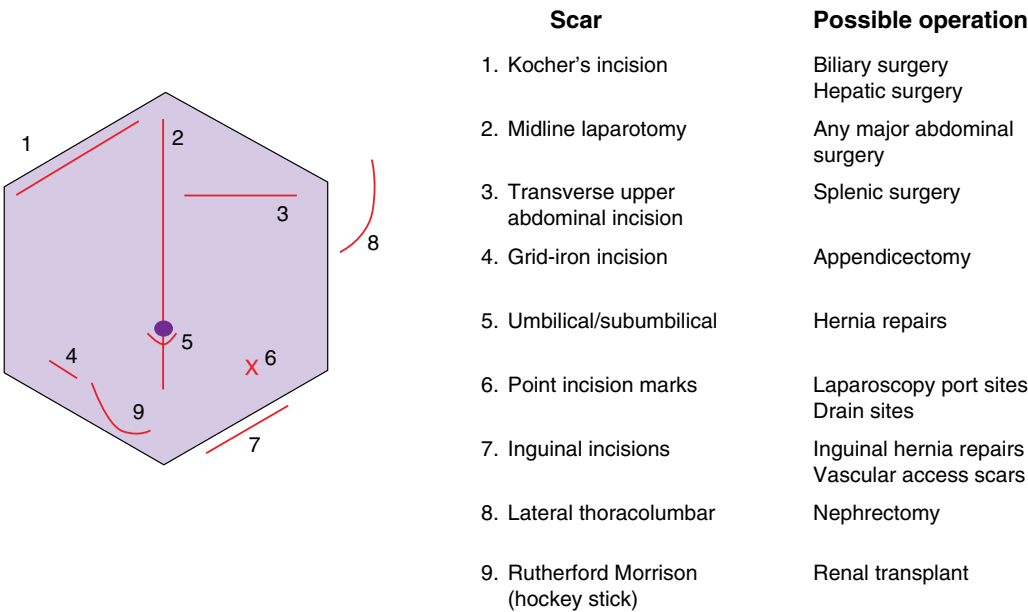


Figure 1.1.4 Common abdominal surgical scars.

Palpation and Percussion

- If there is pain, begin by palpating furthest away from the area that is tender
- Monitor the patient's facial expression at all times for signs of discomfort
- Palpate each area lightly and then repeat with deeper palpation
- Rigidity suggests peritonitis and the abdominal wall will cease to move with respiration
- Feel for masses and hernias, and characterise

1. Chronic liver disease (CLD) ■; 2. Alcohol withdrawal ■; 3. Viral hepatitis ■; 4. Inflammatory bowel disease (IBD) ■; 5. Malignancy ■

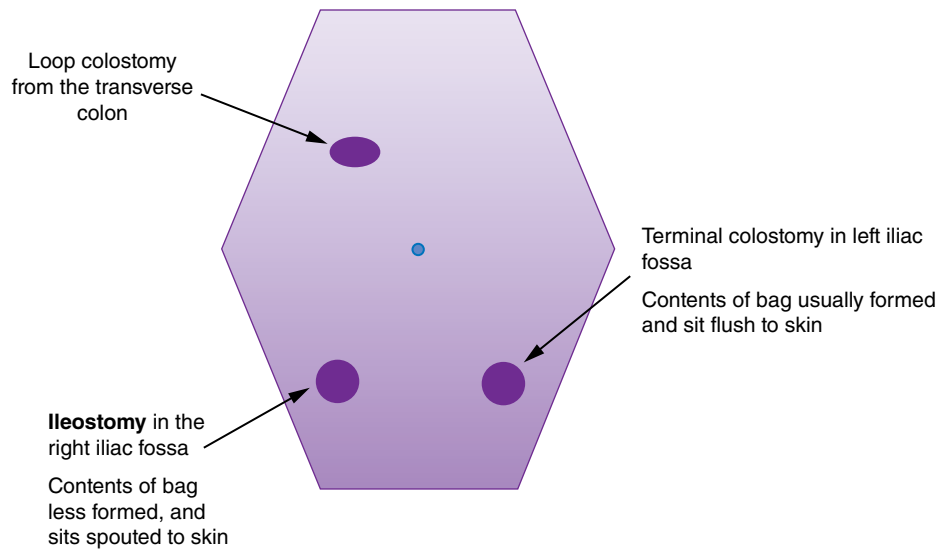


Figure 1.1.5 Common sites of stomas.

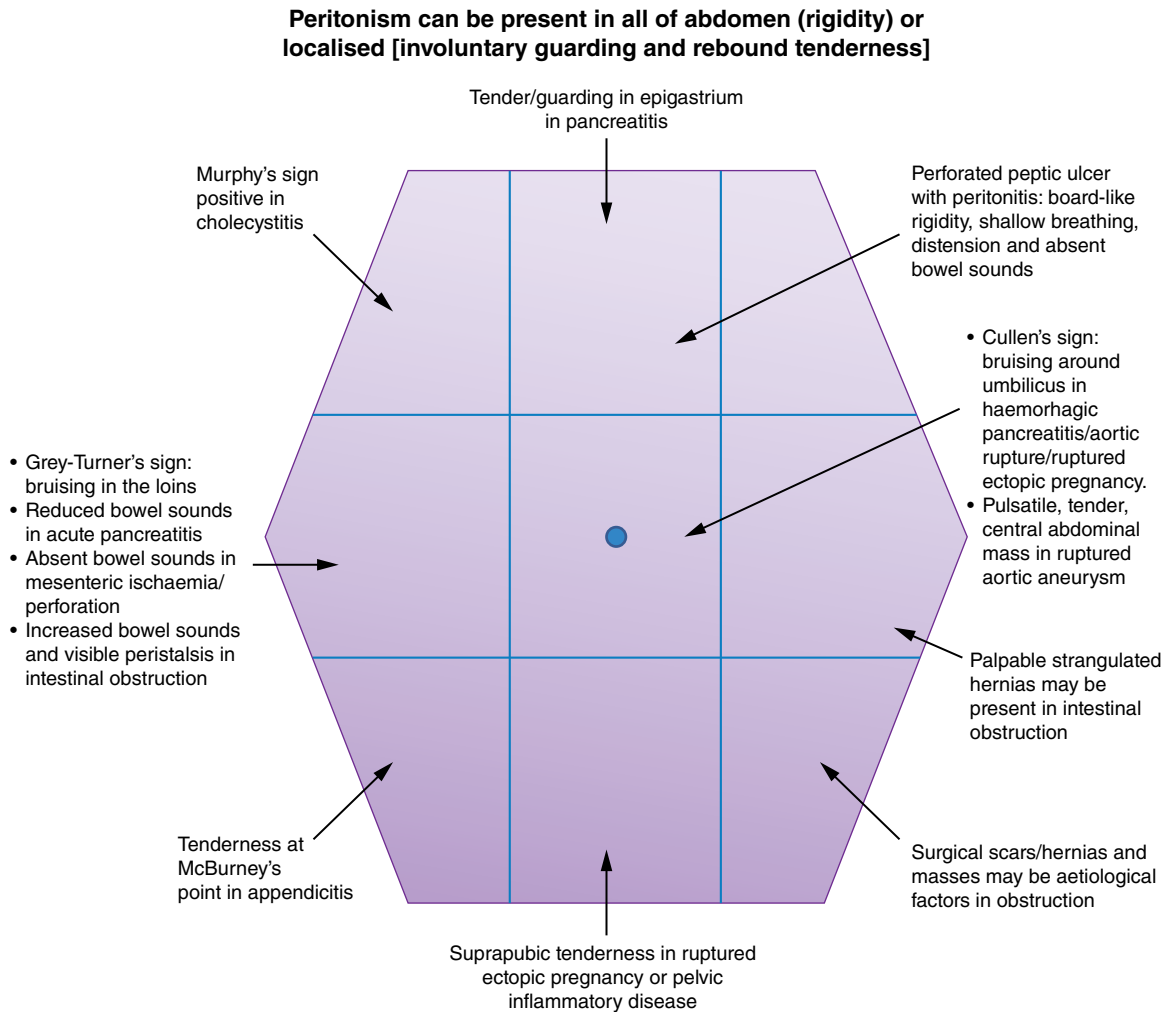


Figure 1.1.6 Possible findings in acute abdominal pain.

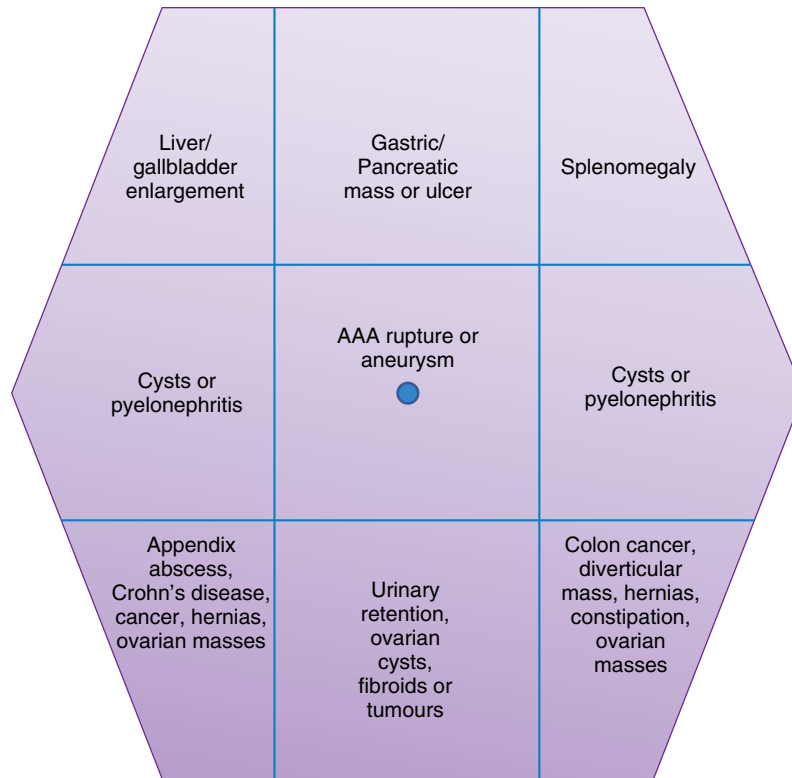


Figure 1.1.7 Location of palpable abnormalities.

Box 1.1.4 Characterising a Lump

- Site
- Size
- Shape
- Solitary
- Skin
- Tenderness
- Temperature
- Tethered/mobility
- Consistency
- Firmness
- Associated lymphadenopathy



Box 1.1.5 Guarding

- Throughout palpation, feel for voluntary or involuntary guarding (tensing) of the abdominal wall muscles
- In involuntary guarding, the abdominal muscles contract as a reflex due to inflammation of the peritoneum
- In voluntary guarding the patient flexes due to pain



Specific Organomegaly

- Palpate for enlarged organs during deep inspiration by the patient
- This can be difficult in patients with limited communication or who are in severe pain
- Start at the right iliac fossa (RIF) and move towards the right (liver) and left (spleen) hypochondria

Liver

- A palpable liver is not always a sign of liver disease
- Normal variant, especially in children and thin adults, e.g. Riedel's lobe
- A pulsatile liver may be found in tricuspid regurgitation

Box 1.1.6 Palpating the Liver

- Located in the right upper quadrant (RUQ) of the abdomen
- Normally sits between the fifth rib and costal margin
- Place your right palm firmly on the abdomen at the RIF (Figure 1.1.4)
- With each expiration, move your hand towards the RUQ; as the patient inspires, keep your hand still and feel for the liver edge against your hand
- If palpable, note the number of finger-breaths below the costal margin, whether the surface is smooth or nodular, whether it is soft or hard, and whether tender

**Box 1.1.7 Causes of Hepatomegaly**

- 1. Hepatitis:**
 - Infections (e.g. viral hepatitis A-E, EMV, CMV, HIV)
 - Autoimmune
- 2. Alcoholic liver disease (ALD)**
- 3. Non-alcoholic fatty liver disease (NAFLD)**
- 4. Neoplastic**
 - Metastases
 - Hepatocellular carcinoma (HCC)
- 5. Infiltrative**
 - Amyloidosis
 - Sarcoidosis
- 6. Metabolic**
 - Glycogen storage disorders
 - Haemochromatosis
- 7. Haematological**
 - Leukaemia
 - Lymphoma
 - Haemolytic anaemia
- 8. Other**
 - Congestive cardiac failure (CCF)
 - Cysts or abscesses
 - Primary biliary cholangitis (PBC)
 - Budd–Chiari syndrome

**Spleen****Box 1.1.8 Palpating the Spleen**

- In contrast to a palpable liver, a palpable spleen is always pathological
- Located superior to the left upper quadrant (LUQ) of the abdomen, beneath the 9th to the 11th ribs
- Place your right hand in RIF (Figure 1.1.5)
- With each expiration, move your hand towards the LUQ; as the patient inspires, keep your hand still and feel for an edge against your hand
- You can place your other hand beneath the patient's left flank and ask the patient to roll slightly towards you to accentuate any mass
- If palpable, note the number of finger-breaths below the costal margin, whether the surface is smooth or nodular, whether it is soft or hard, and whether any tenderness
- The mass should move with the respiratory cycle
- The mass may have a notch to it

**Box 1.1.9 Causes of Splenomegaly**

- 1. Haematological**
 - Lymphoma
 - Leukaemia
 - Myeloproliferative
- 2. Infection:**
 - TB
 - Malaria
 - Leishmaniasis
 - Schistosomiasis
- 3. Portal hypertension**
- 4. Autoimmune**
 - Systemic lupus erythematosus (SLE)
 - Felty's syndrome
- 5. Infiltrative disorders**
 - Sarcoidosis
 - Gaucher's disease



Table 1.1.7 Other masses palpable during an abdominal examination

Finding	Condition
1. Ballotable kidneys	Autosomal dominant polycystic kidney disease (ADPKD)/simple cyst / ■
2. Palpable + pulsatile epigastric/umbilical mass	Abdominal aortic aneurysm (AAA)
3. Suprapubic mass	Palpable bladder (urinary retention)
4. RIF mass	TB ileitis / ■ / ■/faeces
5. Left iliac fossa (LIF) mass	■ / diverticular mass/faeces

Shifting Dullness

- Identifies the presence of ascites (fluid in the peritoneal cavity)

Box 1.1.10 Examination Technique for Shifting Dullness



- Percuss laterally from the umbilicus away from you, noting any change in resonance
- When you reach an area of dullness towards the flanks, keep your finger on the site and ask the patient to turn towards you and lie on their side
- Allow up to 30s before percussing again
- If the area of dullness has become resonant then shifting dullness is present

Auscultation

- Continue to listen for at least 2 minutes before concluding bowel sounds are absent
- Consider paralytic ileus or peritonitis in the case of absent bowel sounds

Table 1.1.8 Additional sounds heard on auscultation

Location	Extra sounds	Cause
Variable	Tinkling or high pitched bowel sounds	Intestinal obstruction
Above the umbilicus	Aortic bruits	AAA/superior mesenteric artery stenosis
2–3 cm laterally above the umbilicus	Renal artery bruits	Renal artery stenosis
Over the liver	Hepatic bruits	■

To Complete the Examination

Examination of Hernia Orifices

Box 1.1.11 Hernias



- A hernia is the protrusion of tissue through the wall of a cavity
- Common finding in abdominal examination but it is important to exclude serious complication of obstruction or strangulation
- Abdominal hernias are often caused by a weakness in the abdominal wall or area of previous surgical excision
- When a patient contracts the abdominal wall muscles, coughs, sneezes or strains, the hernias become more visible

1. Chronic liver disease (CLD) ■; 2. Alcohol withdrawal ■; 3. Viral hepatitis ■; 4. Inflammatory bowel disease (IBD) ■; 5. Malignancy ■

Box 1.1.12 Hernia Examination

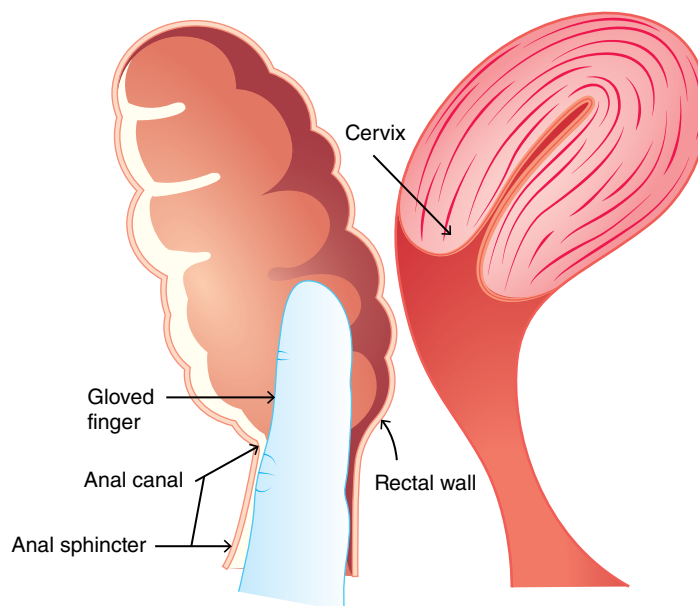
- Examine the groin while the patient stands upright
- Look for any visible masses within the scrotum, inguinal and femoral canals
- Look again while you ask the patient to cough (making the hernia more visible)
- Ask the patient to lie down and inspect for abdominal hernias
- In the groin, if a bulge is identified, palpate it and identify whether the hernia is femoral or inguinal
- Ask the patient to try and reduce the hernia themselves; if they are unable to do this, press two fingers over the hernia and gently try to reduce it
- Examine for other hernias, especially bilaterally in the groin
- Feel for a cough impulse and auscultate for bowel sounds over the hernia
- Most hernias are reducible but some are irreducible and permanently displaced from their correct anatomical location:
 - 1. Incarcerated hernia:** a hernia that is irreducible
 - 2. Obstructed hernia:** incarcerated + evidence of bowel obstruction – pain, vomiting, constipation, distension; requires immediate investigation and surgical team review
 - 3. Strangulated hernia:** incarcerated + evidence of compromise of bowel loop and impending necrosis – exquisite pain; requires immediate investigation and surgical team review

**Rectal Examination**

- Position the patient on their side facing away from you
- Ask them, as much as possible, to pull their knees up towards their chest, and relax
- Inspect the anus externally (skin tags/fistulae/external haemorrhoids)
- Insert index finger
- Feel for sphincter tone and ask the patient to squeeze down on your finger
- Rotate finger anteriorly and palpate prostate or cervix
- Assess for stool (soft/impacted) or masses
- On removing your finger, examine the glove for stool (consistency, colour) and evidence of blood or mucus

Box 1.1.13 Maintaining Dignity During Rectal Examination

- Ensure to consent the patient for this very personal examination and communicate exactly what the examination will involve
- A chaperone will be required who should ideally be the same sex as the patient; make sure that this is also explained and consented. Familiarise yourself with up-to-date GMC guidance
- Cultural differences should be respected – ask about this if unsure
- Ensure environment is appropriate – curtains drawn, privacy sign, locked door

**Figure 1.1.8** Palpable anatomy during a female rectal examination.

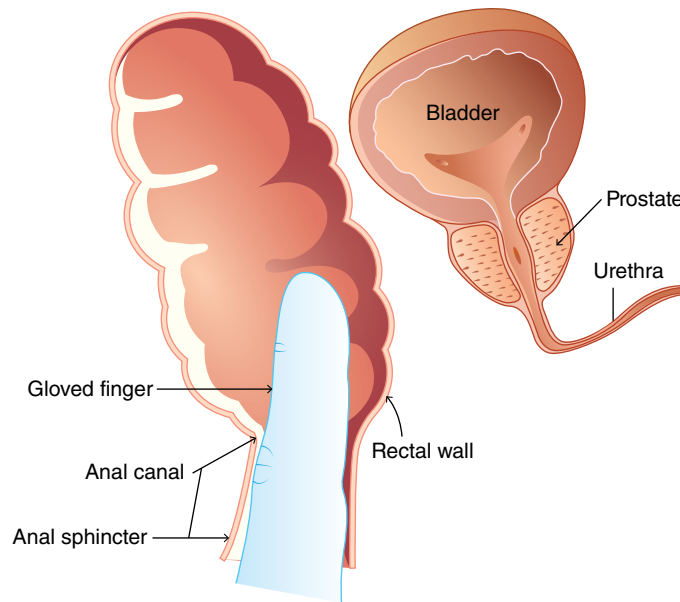


Figure 1.1.9 Palpable anatomy during a male rectal examination.

1.1.5 HOW TO PRESENT YOUR FINDINGS

Safety First Approach

Details

- Give initial overview of how patient looks, e.g. well vs unwell
- Comment on positive and relevant negative findings that can eliminate serious conditions
- Give a potential diagnosis after presenting findings and always offer a differential

Example

I examined a 52-year-old gentleman. He was alert and comfortable at rest and from the end of the bed I noted no relevant medications or therapies such as oxygen. On inspection of the hands, there was palmar erythema but no evidence of track marks or excoriations. There was no lymphadenopathy in the neck or scleral icterus.

On examination of the abdomen, it was soft and non-tender throughout. A fullness was present in his RUQ and on further characterisation there was a liver edge palpable 2 cm below the costal margin. This was smooth and non-tender. There were no other masses and no shifting dullness.

In conclusion, this patient presents with a palpable liver edge – likely hepatomegaly – and peripheral stigmata suggestive of CLD. However, my differential diagnosis would include other causes of an enlarged liver, including malignancy through spread of a gastrointestinal (GI) cancer or cardiac failure.

Ward-Based Presenting

- Often involves other elements to consider – concise and thorough history, investigations that have been sanctioned (and their results), response to therapy and discharge planning
- Present your findings using SBAR to be efficient
- Especially useful in communicating time-critical information, e.g. patients with an 'acute abdomen' or those who have deteriorated
- Trajectory (response to therapy) becomes much easier to observe

1.1.6 EPONYMOUS SIGNS AND SYMPTOMS

Table 1.1.9 Eponymous signs and symptoms

Eponym	Comments
Cullen's sign	Bruising around the umbilicus, suggest retroperitoneal haemorrhage Typically secondary to acute pancreatitis
Grey Turner's sign	Bruising around the flanks, suggests retroperitoneal haemorrhage
Murphy's sign	Ask patient to take deep breaths as you palpate the RUQ As the liver and gallbladder move down with inspiration, pain will be elicited if there is inflammation A positive sign occurs when the discomfort causes termination of the inspiration Only positive if the corresponding sequence on the LUQ is negative Suggests cholecystitis (sensitive but not specific)
McBurney's point	One third of the distance from the right anterior superior iliac spine to the umbilicus Tenderness at this point may indicate appendicitis (see Box 1.1.12)
Rovsing's sign	Palpation of the LIF causes pain in the RIF (may indicate appendicitis)
Troisier's sign	Enlarged/hardening of Virchow's node strongly associated with intra-abdominal malignancy
Virchow's node	Lymph node in the left supraclavicular fossa, common point of lymphatic drainage from the abdomen

Box 1.1.14 The Role of McBurney's Point in Appendicitis

- Tenderness in the area of **McBurney's point** is not caused by direct irritation of the appendix
- Indicates that the peritoneum, which is in contact with the appendix, has become inflamed
- This is a late stage in appendicitis and can give warning that the appendix is at risk of rupture

