

ICU + ANAESTHETICS

ICU principle	Indications for ICU	ICU Additions	ICU complications → solutions
<ul style="list-style-type: none"> Respiratory support Cardiovascular support Renal support Nutritional support Neurological support Dermatological support Liver support 	<p>Cannot be managed by ward !!</p> <ul style="list-style-type: none"> Threatened airway or hard to intubate Post-CPR Severe Sepsis Major trauma (closely monitor vitals + neuro obs) Post-op major surgery (e.g. AAA repair) Any Organ failure → Ventilatory/ circulatory/renal support → dialysis? Iotropes? Bleeding/massive transfusion High analgesia/sedation req. – risk of resp. depression Complex co-morbidity w/ high likelihood of early post-op complication (e.g. poorly controlled OSA, PPH patient) 	<p><u>Analgesia support:</u></p> <p><u>Nutritional support:</u></p> <ul style="list-style-type: none"> Mouth NGT (PEG) – percutaneous endoscopic gastrostomy (tube from surface of abdomen to stomach) TPN via CVC → thrombophlebitis risk so given via central line (NOT peripheral cannula) 	<p>Acute</p> <ul style="list-style-type: none"> Ventilator assoc. lung injury (barotrauma, volutrauma) → short-term APO, hypoxia → long-term: fibrosis, recurrent infection, cor-pulmonale Ventilator assoc. pneumonia (25% complication) → position semi-fowler and head elevation Catheter related blood infections (e.g. from CVC) – 25% mortality → Rx with ABx or Ag impregnated catheters Delirium → AT4 assessment (to identify delirium) → Dexmedetomidine (sedate agitated pts) + Rx cause <p>Chronic</p> <ul style="list-style-type: none"> Catheter assoc. UTI Stress related mucosal ulcers (e.g. erosion of upper GIT) → Rx: PPI and H2 antagonist VTE → calf compression +SC clexane Critical illness myopathy (due to corticosteroids or muscle relaxants) → difficult to wean off mech. Ventilation Critical illness neuropathy → optimize BSL control Transfusion related reactions (TRALI, sepsis, overload, coagulopathy)

Why is it difficult to mobilise after being ventilated?

- Patient may have decompensated from the initial event and thus, organs and muscles need to take time to adapt to the new conditions
- Critical illness will also cause both myopathy and neuropathy particularly after paralysing agents have been used
- The **long-term immobility** = muscle atrophy as deconditioned when ill → MOBILISE Early w/ PT → reactivate muscles to restore power and endurance.
- If patient has a **tracheostomy tube** still in place, the reduced air intake means there is reduced ventilation due to reduced tidal volume. This will cause a low V/Q mismatch, thus causing organ hypoperfusion.

What is the prognosis of patients who have gone through ICU?

- POOR** despite successful resuscitation and rehabilitation. Often there is some damage to organs during the entire process.
- +++ morbidity and mortality → longer rehabilitation period → lose their independence earlier relative to healthy age-matched controls.
- +++ f/u with different specialists and allied health professional to manage their condition and progress → costs, logistics, time
- +++ new meds, which can lead to further financial burden and increase the risk of drug-drug interactions, non-compliance due to polypharmacy.

Why is it important to ask about antibiotic usage during a sepsis crisis?

- Antibiotics may confound the blood culture results, particularly if there is bacterial infective source
- Antibiotics may actually be asking the severity of the condition patient is in (false sense of security when managing patient)
- Be aware of possible anaphylactic allergic reactions to certain antibiotics

What can be done when locating the source?

- Strip patient from head to toe
- 2nd survey to determine key ports of entry (e.g. IDC, wound site, central lines, cannula sites) as well as areas of excoriations, lacerations, puncture wound

ICU MORTALITY RISK CALCULATORS (APACHE-II and MPM)

What are the **2 main factors** consider during admission?

- the potential to reverse the acute condition
- the baseline physiological reserve (their baseline health).

e.g. patients with a 90% probability of dying w/ underlying terminal condition should be given palliative care approach rather than have invasive interventions

Acute Physiology And Chronic Health Evaluation II (APACHE-II)		ICU Calculators - RN SH
	Measurements	Mortality Probability Model (24-48-72 h)
Within 24 hours of admission to the ICU (The point score is calculated from a patient's age and these 12 routine physiological measurements)	1. AaDO ₂ or PaO ₂ (depending on FiO ₂) 2. Temperature (rectal) 3. Mean arterial pressure 4. pH arterial 5. Heart rate 6. Respiratory rate 7. Sodium (serum) 8. Potassium (serum) 9. Creatinine 10. Hematocrit 11. White blood cell count 12. Glasgow Coma Scale	Date 24h <input checked="" type="radio"/> 48h <input type="radio"/> 72h Age 50 Admission type <input type="checkbox"/> Metastatic Cancer <input type="checkbox"/> Cirrhosis <input checked="" type="checkbox"/> Diuresis < 150 mL/8h <input checked="" type="checkbox"/> Creatinine > 2 mg/dL <input type="checkbox"/> Coma (GCS 3-5) <input type="checkbox"/> Intracranial Mass Effect <input type="checkbox"/> Vasoactive Drug >= 1h <input type="checkbox"/> Mechanical Ventilation <input checked="" type="checkbox"/> PaO ₂ < 60 mmHg <input type="checkbox"/> Proven Infection <input type="checkbox"/> PT > Standard +3 sec Calculate Mortality Rate <input type="text" value="23.8"/> %
Interpretation:	<ul style="list-style-type: none"> Measurements are utilised in addition to information about previous health status (recent surgery, history of severe organ insufficiency, immunocompromised state). This generates a score between 0-71, higher scores correspond to higher severity of disease and higher likelihood of mortality. 	

Knaus WA, Draper EA, Wagner DP, Zimmerman JE (1985). "APACHE II: a severity of disease classification system". *Critical Care Medicine*. 13 (10): 818-29. doi:10.1097/0003246-198510000-00009. PMID 3928249. (This is the first published description of the APACHE II scoring system)

Knaus WA, Zimmerman JE, Wagner DP, Draper EA, Lawrence DE (1981). "APACHE-acute physiology and chronic health evaluation: a physiologically based classification system". *Critical Care Medicine*. 9 (8): 591-7.

FLUID Management: FLUID + ELECTROLYTES

Assessing volume levels:		Examination	Bedside Investigations:	Mx (conservative):
Hypervolaemic		1. Aids - finished meal? Cups of water? IDC/drain output?	➤ Wt, BMI ➤ BLADDER SCAN → ?IDC ➤ Urine - ↑SG, urine osmolarity ➤ ECG - arrhythmias?	Overload (WET) Non-pharm Pharm
XS fluid intake		Dehydration signs:		1L fluid restrict Na restricted diet
XS salt intake		1. VITALS - hypotension , Tachycardia , CRT >2s , tachypnoea		Hydrate
IVF		2. Dry MM - stick tongue out		
Euvolaemic		3. Sunken eyeballs		
Hypovolaemic (Dehydrated)		4. Cold peripheries		
Vomit / diarrhoea		5. Low urine output		
Internal bleeding		6. Low weight		
Burns		Fluid overload signs:		
DI		1. Elevated JVP		
		2. Displaced apex beat (most sensitive for HF)		
		3. Pulm. Oedema (bribasal crackles, low sats)		
		4. Peripheral oedema, ascites		
		5. Weight gain		

Colloid			Crystalloid			
Vol. needed	Vol for vol		3x vol needed			
MoA	<ul style="list-style-type: none"> Molecules too large to cross capillary walls - fluid remain in intravascular space Long half-life (hrs-days) 			<ul style="list-style-type: none"> Molecules small enough to cross capillary walls → less fluid in intravascular space Short half life (30-60 mins) 		
• Greater increased effect on intravascular vol. (1.5:1) <ul style="list-style-type: none"> For sepsis (NOT for TBI) 						
Example	Natural <ul style="list-style-type: none"> Whole blood FFP pRBC Albumin 		Hypotonic <ul style="list-style-type: none"> 5% dextrose and 0.18% NS Hypotonic saline Isotonic <ul style="list-style-type: none"> Normal saline (0.9%) Lactated ringer's soln (aka Hartmann's) plasmolyte Hypertonic <ul style="list-style-type: none"> 3% NaCl 5% NaCl 			
ECF	Increased (esp. intravascular vol)			ECF	Increased	
ICF	NONE			ICF	Increased	
Use	<ul style="list-style-type: none"> Cirrhosis Critically ill - ARDS, burns, sepsis Bleed → blood 			Ind. Use	none	
A/E	<ul style="list-style-type: none"> Fluid overload → cardiac failure Allergic Expensive + may not be vegan 			Ind. Use	Decreased	
FLUID OVERLOAD - cerebral, peripheral, pulmonary oedema <ul style="list-style-type: none"> hypoNa, hypoK hyperglycaemia cerebral oedema 						
A/E	<ul style="list-style-type: none"> NO dextrose in: brain haemorrhage re-feeding syndrome 			A/E	<ul style="list-style-type: none"> 0.9% NS = HYPERchloraemia acidosis Ringer's/ Hartman = ++ lactate in liver failure, hyper K Plasmalyte → high HCO₃ 	
Osmotic demyelination syndrome → cerebral oedema						

	Resus	Maintenance	Replacement fluids	
Fluid	0.9% NS	Crystalloids (NS, dextrose) or colloids	Crystalloids → selected with similar electrolyte content to fluids that are lost	
Scenario	Sepsis, hypotension	PERI-OP & POST-OP NBM due to bowel obstruction	POST-OP + VOMITING + DIARRHOEA	
Method	Rapid fluid bolus (within 10-30mins) <ul style="list-style-type: none"> Adults NS or LR 500-1000mL IV bolus Children: NS or LR 10-20mL/kg IV bolus 	NBM patients but do not have volume depletion, hypotension, or ongoing losses <ul style="list-style-type: none"> Adults → 1-2mL/kg/hr Children (> 28 days) → 4:2:1 rule <ul style="list-style-type: none"> 4mL/kg/hr (1st 10 kg) 2mL/kg/hr (2nd 10 kg) 1mL/kg/hr (remaining) *Maintenance fluid requirement per kg of wt higher in children due to increased SA	Replace lost body fluids and electrolytes	
Consider	<ul style="list-style-type: none"> vasopressors (e.g. metaraminol) → maintain BP and reduce peripheral fluid loss Ionotropes (e.g. dobutamine, levosimendan) Blood products (FFP, packed RBC - group + X-match) 	<ul style="list-style-type: none"> Patient's weight Check EUC before prescribing Give oral/NG-tube fluids whenever possible → minimizes fluid overload 	<ul style="list-style-type: none"> Pre-existing fluid loss (STAT bolus - 500mL 0.9% saline/Hartmann's soln) Measure Ongoing losses (replace future losses - measure vomits, diarrhoea vs intake) DO NOT Give K at a rate > 10mM/hr or use maintenance protocols 	<ul style="list-style-type: none"> Hartmann's BEST as less Cl- to minimise risk of hyperchloraemic acid (check lactate) Dextrose-saline (normal maintenance fluid) If hypotension → bolus Hartman/saline Colloid Transfusion (4x pRBC + 2x FFP) FFP/ platelets (stop bleed) BUT need to find source

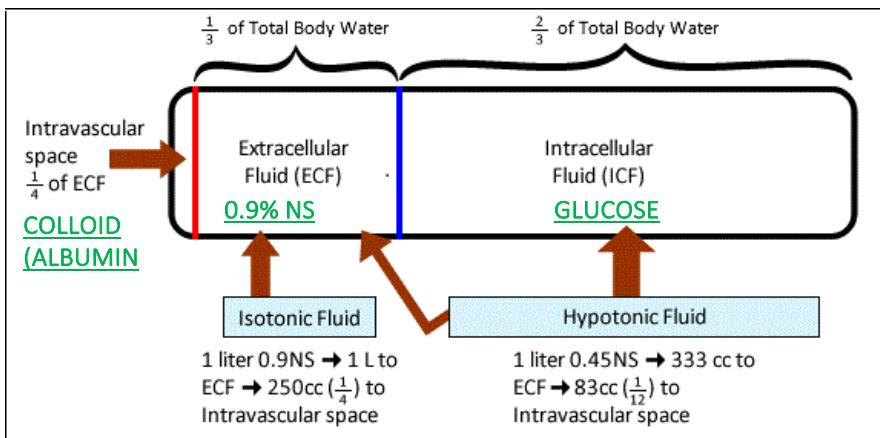
Special cases:

Post-op	Cell lysis during surgery → elevated K, AKI	No K Low protein diet (0.6g/kg/day) = ↓ hyperfiltration
Sepsis	Hartman or 0.9% saline → vasopressors + monitor?	CKD
HF	Nb: may normally be hypotensive → low Na diet, furosemide, fluid restriction and record daily weights	Alcoholic
Liver failure	5% dextrose - excess Na causes ascites	Brain haemorrhage
		Saline (dextrose destroys brain)

Crystalloid Fluids

Fluid	Na ⁺ mEq/L	Cl ⁻ mEq/L	K ⁺ mEq/L	Ca ²⁺ mEq/L	Glucose g/L	Buffer	Osmolarity mOsm/L	Tonicity	Typical Indication
Normal plasma	~ 140	~ 100	~ 4	~ 2.4	~ 0.85	HCO ₃ ⁻ ~ 24 mEq/L	~ 290	N/A	N/A
0.9% saline (a.k.a. "normal saline" or NS)	154	154	0	0	0	0	308	"Isotonic"	Resuscitation
0.45% saline (a.k.a. ½ NS)	77	77	0	0	0	0	154	Hypotonic	Maintenance
3% saline	513	513	0	0	0	0	1026	Hypertonic	Severe Hyponatremia
D5 ½NS + 20 meq KCL	77	97	20	0	50	0	446	Hypertonic → Hypotonic	Maintenance
D5W	0	0	0	0	50	0	252	Hypotonic	Hypernatremia Hypoglycemia
Lactated Ringer's (LR) / Hartmann's solution*	130	109	4	3	0	Lactate 28 mEq/L	273	Isotonic	Resuscitation

60% TOTAL BODY WT = TOTAL WATER



DOES NOT ENTER ICF

Table 2 Daily Electrolyte Requirements

	DAILY REQUIREMENT	FOR 70-KG ADULT	FOR 10-KG CHILD
Sodium	1-2 meq/kg	70-140 meq/day	10-20 meq/day
Potassium	0.5-1.0 meq/kg	35-70 meq/day	5-10 meq/day
Calcium	0.2-0.3 meq/kg	1.4-2.1 meq/day	2.0-3.0 meq/day
Magnesium	0.35-0.45 meq/kg	24.5-31.5 meq/day	3.5-4.5 meq/day
Chloride	equal to sodium	equal to sodium	equal to sodium
Bicarbonate/Acetate	use with chloride to balance cations and help pH	use with chloride to balance cations and help pH	use with chloride to balance cations and help pH

Calculating fluids:

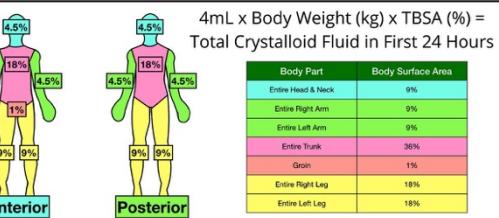
- 60% of total body weight is water
- 4:2:1 rule in children
- CHECK Hct (for haemodilution) due to XS fluid resuscitation**

PARKLAND FORMULA for burns

The volume required over 24 hours will be:

- 4 mL x % burn SA x body weight in kg
- half this volume is given in the first 8 hours
- Resus w/ NS or Hartmann's but consider colloids (albumin) if deep burns

Parkland Formula for Burns



Daily Requirements: (for 70kg man)

- Water = 25-30 mL/kg/day (2L)
- Glucose ≈ 50/100g.day (50-100g)
- Na, K and Cl = 1mM/kg/day (70mM)

Input vs Output:

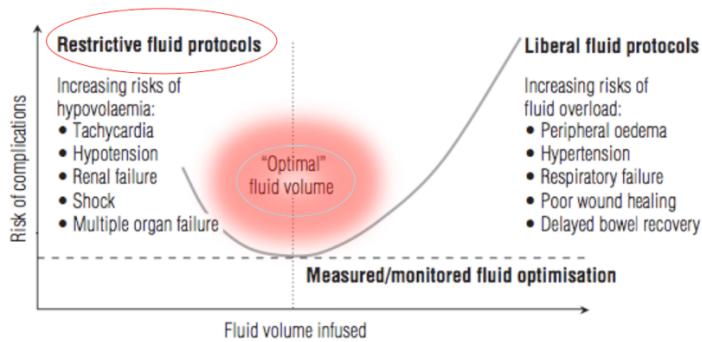
- Daily weighing
- UO ≈ 0.5mL/kg/hr
- Oral intake / IVF / NGT
- Drains/ stoma / IDC

High risk patients (need senior input)

- Elderly or frail patients
- Significant oedema
- Sodium imbalance (low or high Na)
- Heart failure
- Renal failure
- Liver failure

Classification of Shock				
	Class I	Class II	Class III	Class IV
Blood loss (mL)	< 750	750-1500	1500-2000	> 2000
Blood loss (%body vol)	< 15%	15-30%	30-40%	> 40%
HR	< 100	> 100	> 120	> 140
BP	Normal	Normal	Decreased	Decreased
Pulse pressure	Normal/ Increased	Decreased	Decreased	Decreased
RR	14-20	20-30	30-40	> 35
Urine output (mL/hr)	> 30	20-30	5-15	Negligible
CNS	Slightly anxious	Mildly anxious	Anxious and confused	Confused and lethargic
Fluid replacement	Crystalloid	Crystalloid	Crystalloid and Blood	Crystalloid and Blood
Base Excess	0 to -2	-2 to -6	-6 to -10	< -10

FLUID Management (5 R's)



Osmolarity

- mOsm/L of a solution
- measures ALL of the particles per volume in the solution
- Normal serum = **280 – 300 mOsm/L**
- **Sodium** is the biggest contributor ($Na \times 2 + \text{urea} + \text{glucose}$)

Tonicity

- The particles in a solution that CANNOT cross a plasma membrane
- Depends on membrane properties
- **Effective osmotic pressure**

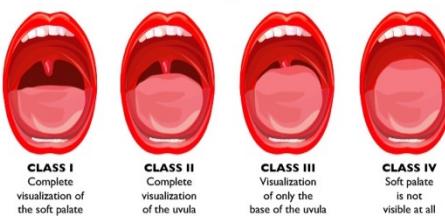
Oncotic pressure

- Osmotic pressure exerted by **proteins/colloids**

Resuscitation	<u>What do I give?</u>	Crystalloids Saline vs Hartmann's (balanced) Colloids Starches → AKI, coagulopathy, CHEST Albumin → SAFE except for TBI					
	<u>How do I give?</u>	<ol style="list-style-type: none"> 1. Have a repeatable outcome measure <ol style="list-style-type: none"> E.g. vitals Check → Warm, Wet, With it or Wactate 2. Decide on safety stopping point <ol style="list-style-type: none"> Failure to response Insipient cough OR increasing O₂ requirements PiCCO values 3. Give 500mL over < 15mins 4. Evaluate response and repeat 					
	<u>What could go wrong?</u>	<ul style="list-style-type: none"> ➤ All sick patients have risk of APO ➤ Rapid infusion of crystalloids may damage glycocalyx 					
	<u>Considerations</u>	<p>Increased fluids bolus</p> <ul style="list-style-type: none"> ➤ Increased atrial stretch → cause reduce HR and CO ➤ Increased CVP → counterintuitively reduces perfusion (despite increased preload) ➤ Increased UO → may be due to ↑ANP and NOT eGFR 					
	Routine maintenance	<ul style="list-style-type: none"> • 1L of blood = 3L of non- blood fluids • 25 mls/kg/day ≈ 1 mL/kg/hr <ul style="list-style-type: none"> ○ Na/Cl/K ≈ 1mM/kg/day ○ Glucose 50-100g/day • Less is more elderly/CCF/CKD/malnourished • Take into account OTHER sources of fluid and electrolytes <ul style="list-style-type: none"> ○ Oral intake ○ ABx ○ Sedation / vasopressors ○ Nutrition and TPN ○ Check Cl daily (if using NS) and watch for acidemia 					
Replacement	<u>What has been lost?</u>	<p>Replace Like with like</p> <ul style="list-style-type: none"> • Insensible loss = water + small amounts of NaCl • Gastric = low Na + Chloride rich (acid) • Pancreatic = Na + HCO₃ rich • Upper GI = similar to serum (Na, Cl, HCO₃ rich) • Lower GI = K rich and Cl free 					
	<u>How much is lost?</u>	<ul style="list-style-type: none"> • Easy = NGT, IDC, Ileostomy • Hard = 3rd spacing (burns, ileus, peritonitis) <table border="1"> <tr> <td>Mild (<5%)</td> <td>C/O thirst</td> </tr> <tr> <td>Mod (5-15%)</td> <td>Dry MM, skin turgor, mild tachycardia, reduced UO</td> </tr> <tr> <td>Severe (>15%)</td> <td>Altered mental state, hypoTN,</td> </tr> </table>	Mild (<5%)	C/O thirst	Mod (5-15%)	Dry MM, skin turgor, mild tachycardia, reduced UO	Severe (>15%)
Mild (<5%)	C/O thirst						
Mod (5-15%)	Dry MM, skin turgor, mild tachycardia, reduced UO						
Severe (>15%)	Altered mental state, hypoTN,						
<u>How much will continue to be lost?</u>	<p>Clinical parameters</p> <ul style="list-style-type: none"> ➤ 5, 10 and 15% assessments ➤ Daily weight ➤ Fluid balance charts 						
Redistribution	1st spacing	<ul style="list-style-type: none"> • Normal distribution within ECF and ICF 					
	2nd spacing (fluid we see)	<ul style="list-style-type: none"> • Accumulation in interstitial compartment • Available for exchange between compartments 					
	3rd spacing	<ul style="list-style-type: none"> • NOT available for exchange between compartments • e.g. ascites, oedema from burns/surgery 					
Reassessment	<ul style="list-style-type: none"> • Most important thing 						

Pre/Intra-operative Assessment

INITIAL AIRWAY ASSESSMENT			
1) Ability open mouth			
2) Subluxation of jaw			
3) Thyro-mental distance			
4) Cervical motion and stability			



ASA class	Definition	Pooled mortality (%)
I	Healthy	0-0.3
II	Mild systemic disease with no functional limitation	0.3-1.4
III	Severe systemic disease with functional limitation	1.8-5.4
IV	Severe systemic disease – constant threat to life	7.8-25.0
V	Moribund patient – unlikely to survive 24 h with or without operation	9.4-57.8
E	Suffix added to denote emergency operation	

Key tips to consider before Hx and exam?	Do YOU HAVE THE CORRECT PATIENT & Hx? 1. Who is the pt? → age, wt 2. What surgery? Why? Done by reg or consultant? 3. When is surgery? 4. Where is surgery? (radiology, OR etc.) 5. How to reduce cost? 6. Plan beyond skin incision (are they going home/wards/ICU?) → need ionotropes	Risk		Surgery	Action
		Low (<1%)	Breast + dental	No Ix	
		High (1-5%)	Vascular + thoracic (lung)		Ix to lower risk (reduce long-term health complications)
		ED (>5%)	AAA, recent MI, decomp. HF, High grade arrhythmia, valvular issue (AS, peritonitis, appendicitis)		ED surgery regardless of CVS risk

PMHx + PSHx + meds and response to anaesthesia	Concern		Pre-Op Investigations	Solution
	Resp (airway + breathing)	<ul style="list-style-type: none"> Beards, jaw protrusion Acute (E.g. PE, pneumothorax, pneumonia, infection) Chronic (OSA, ILD, COPD) 	<ul style="list-style-type: none"> Auscultate + O₂ sats CXR → check for HF, PPM Spirometry + ABG OSA checklist 	<ul style="list-style-type: none"> Cease smoking > 4 weeks Post-op monitor → beware breathless pt Immunosuppressed (steroids – COPD)
	CVS (IHD)	<ul style="list-style-type: none"> AHA revised cardiac risk index High risk surgery Hx of IHD (ACS-MI, UA, decomp. HF) Hx of HF + surgical Hx (stents, CABG) Hx of CVD IDDM Renal impairment (eGFR – CKD → fluid caution) 	<ol style="list-style-type: none"> Bloods + products → Group + save (pRBC, FFP, cryoppt. plts) <ol style="list-style-type: none"> FBC, EUC, - anaemia (↑↑ post-op complication) COAG, B-HCG CXR, CTA/CTPA or V/Q scan ECG (pre & post) – ? AF, ischaemic damage (inverse T waves, path Q waves, BBB) <ol style="list-style-type: none"> PERFORMED IN ALL > 65 TO pre-operatively TOE/TTE stress tests → ONLY if Hx of valvular disease (MR) or heart failure (LVEF) Cardiopulmonary exercise testing to check for reserve – CPET, stress ECHO 	
	CVS (HTN)	++ peri-op morbidity	24hr holter-monitor	<ul style="list-style-type: none"> Target MAP > 65 SBP > 80 (higher if older w/ PVD)
	Diabetes	<ul style="list-style-type: none"> ++ infection risk Micro + macrovascular comp. 	BSL, HbA1C, OGTT (new T2DM)	<ul style="list-style-type: none"> Optimise BSL control (target 5-10mM) Avoid fasting + BSL fluctuations Make sure 1st on list
	Older age + Obesity	High risk of desaturation rapidly	STOP BANG score for OSA cause-snore, tired, observed, BP, BMI, age > 50, neck cir. > 40cm, gender - male	<ul style="list-style-type: none"> Minimise post-op comp Lose weight

Capacity and function	PMHx	Response to anaesthetics	Previous response	Previous surgeries
		<ul style="list-style-type: none"> Allergies to → propofol (soy, egg). Opioids, Check FHx → malignant hyperthermia 		<ul style="list-style-type: none"> Difficult airways Stents , CABG Jt replacements Abdo surgery
	Exercise tolerance	MET – Metabolic equivalent – what can they do?	<ul style="list-style-type: none"> 1 MET – eating, sitting, bath, shower and dress 4 METs – walk up flight of stairs 10 METs – strenuous exercise 	Improve METS – condition/train <ul style="list-style-type: none"> Walk unaided? 6-metre walk
	Function / mood	<ul style="list-style-type: none"> Frailty score: Rockwood or Fried scale NSQIP risk index (surgical risk calc → age, organ fn, chronic illness, poor nutrition, cognitive dyfn, male) Nottingham hip # score → quicker to theatre = ↓mortality Depression = longer hospital stay 		Functional aids <ul style="list-style-type: none"> Sensory/hearing aids (glasses , hearing aid) Dentures Walking stick, wheelchair
	Alcohol	Organ damage → hepatic encephalopathy, pancreatitis		Avoid withdrawal + delirium
	Smoking	<ul style="list-style-type: none"> +++ risk of IHD, PVD, HTN → Atherosclerosis, ↓ tissue perfusion ↓wound healing Carcinogenic Lung disease (Bronchiectasis, COPD) → ↓ciliary motion & sputum clearance 		Cease smoking > 4 weeks <ol style="list-style-type: none"> Improve tissue perfusion (days) ↑ wound healing ↑ pulm. Function ↑ immune system

Meds	Drug	Stopped?	Reason?	
	Antiplatelets (Stents)	Aspirin (COX 1/2 inhibitor)	5-7 days (only if high risk)	
		Clopidogrel -Plavix (P2Y ₁₂ inhibitor)	5-7 days	
		Ticagrelor (P2Y ₁₂ inhibitor)	5-7 days	
	Anti-coagulants (AF, tissue valve, DVT)	Warfarin → Beware INR > 3.5 + warfarin + spinal haematoma	≈ 5 days IV Vit K, prothrombinex → restart 12-24 hours after surgery	
		Dabigatran (anti-thrombin - IIa)	1 day idarucuzimab	
		Clexane (LMWH)	3 days before + post-op LMWH for 3 days	
		DOACS - -xabans (Xa inhibitor)	≥3 days Praxbindi + post-op LMWH for 3 days -- may not need if low risk surgery (e.g. cataracts)	
	Anti-hypertensives	ACEi/ARB	1 day	
		Diuretics	1 day	HypoTN
				HypoK + IBP
	Oral Hypoglycaemics	Metformin	1 day	
		GLP-1	1 day (only if bowel prep)	Lactic acidosis + AKI
		SGLT2i	3-4 days (1 day if day-only procedure with no bowel prep)	GI disturbance
	Natural therapies	Ginger, gingko, garlic, ginseng + fish oil	↑ bleeding risk	
	NSAID	Ibuprofen	PUD	
	Other	OCP	4 weeks before major surgery	

Stratify stroke risk (CHADS-VASC)

- Low risk (<5%)** = no bridging needed
- Mod risk (5-10%)** = stop warfarin 5 days before + start clexane (Stop day before surgery)
- DOAC** – no bridging needed

HypoTN

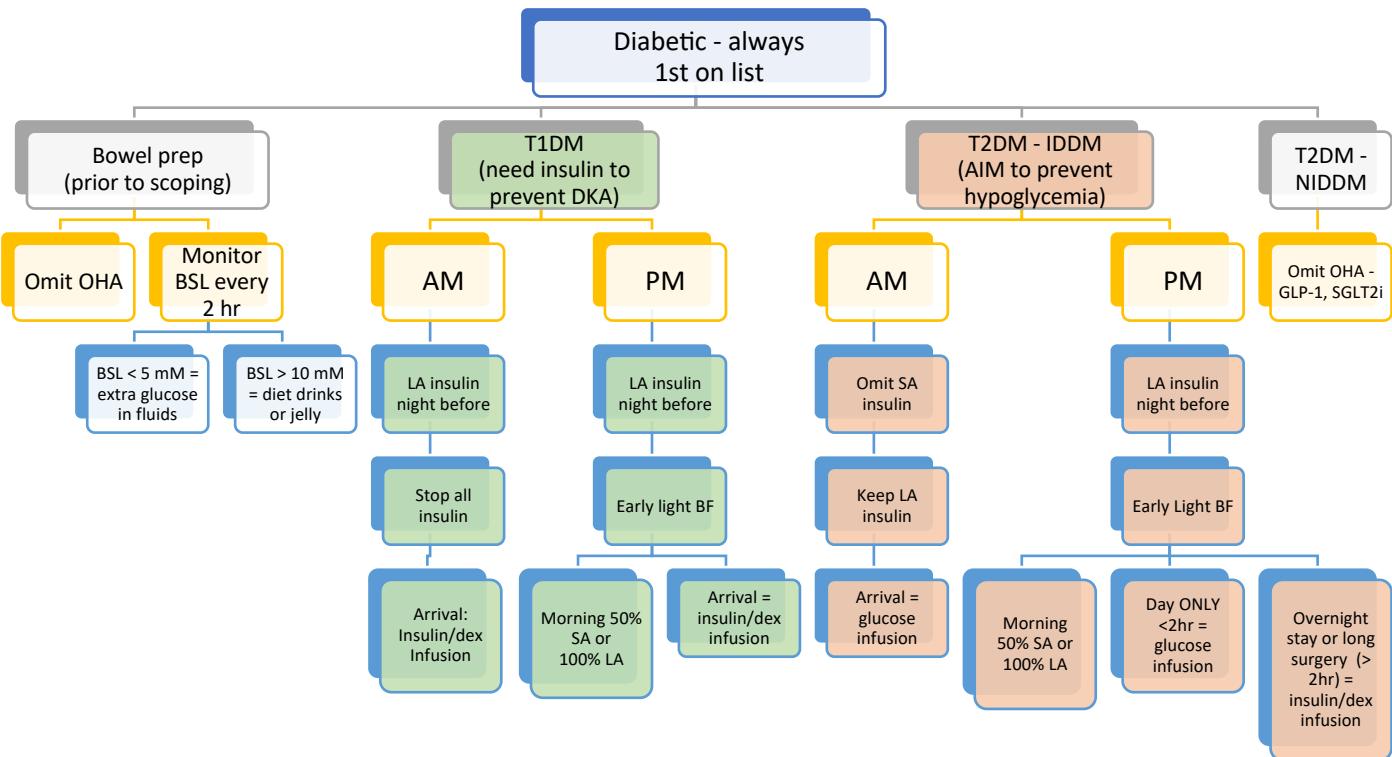
- Lactic acidosis + AKI

HypoK + IBP

- GI disturbance

Euglycaemic Ketoacidosis (monitor ketones pre and post-op)

Peri-operative Mx of the Diabetic Patient



*Make plan with endocrinology and anaesthetics

*Check ketones before and after (is it starvation ketosis, DKA or euglycemic ketoacidosis)

Key learning points

- BSL < 3mM → IV 50% dex bolus
- BSL < 6mM → IV insulin + 5% dex
- BSL 18-20mM (after intermediate insulin) → WAIT before giving actrapid to avoid hypoglycaemia
- Check ABG + ketones >1.5mM for SGLT2i (pre and post-op)
 - Can omit SGLT2i for one day (if day ONLY procedure + no bowel prep)
- Sliding scale insulin
 - BSL check qid
 - ½ U per BSL → titrate accordingly
- Steroid usage used may need to give insulin after 4-5 hrs later
- Pre-operatively
 - CHO loading and increase cardiorespiratory fitness

Intern should Plan for ERAS

- Shortened fast (2hr prior to surgery OK)
- EARLY mobility (PT)
- Fluid Mx
- ABx follow-up
- Early removal of lines
- Wean analgesia
- Nutrition step-down (**NBM → TPN → clear fluid → low residue → light diet → full**)
 - TPN (via CVC) -> given if high catabolic state (e.g. sepsis) with no gut function
- Early Transfer home, aged care, rehab?

Diabetic patients in ICU issues:

- Poor glycaemic control (hyper/hypoglycaemia)
- Use IV insulin-dextrose keep patients 6-10mM:
 - minimise tight BSL control
 - Avoid intermittent SC insulin bolus or oral hypoglycaemics
- Increased sepsis risk
- Poor wound healing
- High risk of AKI (esp. if existing renal impairment)
- Lactic acidosis
- DKA (esp. T1DM)
- Hyperosmotic -non-ketotic state (T2DM)

Principles of Surgery

Ambroise Pare's principle of surgery

"eliminate the superfluous, restore the dislocated, separate things that have united and join what has been divided and repair defects in nature"

Natural process of healing and inflammation:

<u>Skin</u>	<u>Ulcer</u>	<u>Sinus</u>	<u>Fistula</u>	<u>Abscess / cyst</u>	<u>Cyst and abscess drainage</u> <u>Hypergranulation (proud flesh)</u>	<u>Keloid scar</u>
						Along langer's lines (XS healing) Repairs in 2 years
<u>Incision</u>	<u>Join / fistula</u>	<u>Overhealing stenosis</u>	<ul style="list-style-type: none"> ➤ False aneurysm ➤ Perforation/ leak ➤ Hemorrhage 		<u>Types of healing:</u> <ol style="list-style-type: none"> 1. Healing by primary intention = primitive intension closed by sutures 2. Healing by secondary intention = wound left open to heal by granulation contraction 	

Stage	Process	Duration	Can it be repaired?
Haemostasis	<ul style="list-style-type: none"> ➤ Prevent / stop bleed → plt + cytokines → vasoconstrict 1) Primary haem = during surgery 2) Secondary haem = post-surgery 	1-3 hours/day	<ul style="list-style-type: none"> ➤ Within "2 weeks" of surgery, anything can be repaired
Inflammation	<ul style="list-style-type: none"> ➤ Activated immune system to remove pathogen and necrotic tissue 	3-20 days	<ul style="list-style-type: none"> ➤ Once > 2 weeks – proliferation phase begins (highly fragile state) and anatomy changes
Proliferation	<ul style="list-style-type: none"> ➤ Healthy tissue regrows (ECM + collagen = granulation) ➤ Angiogenesis 	1-6 weeks	
Remodelling	<ul style="list-style-type: none"> ➤ Granulation tissue matures into scar tissue (improved tensile strength) 	6 weeks to 2 years	

What factors affect surgery success – Dimension of surgery?

- **Time** → HIPR
- **Age** → older age = more fragile, longer wound healing and cut easily
- **Nutrition** → when was last proper meal, concern for surgery if BMI < 18 (ANOREXIA)
- **Chemo**
- **Redo** → within 2 weeks
- **RT** → esp. pelvic surgery → cervicitis, proctitis
- **Tissue handling** → gentle vs rough

What are the surgical "rules of thumb"

- Mishap
- Attend
- Causation
- Next operation

What are some important surgical considerations?

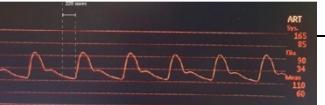
- **Langer's lines = lines of skin tension** (topological lines) to minimise skin tension across wounds + more aesthetically pleasing
 - 1) "discovered when Langer punctured numerous circular holes and noticed there were actually ellipsoid in shape"
- **Enhanced recovery after surgery (ERAS) Program**
 - 1) Early mobilisation to reduce muscle breakdown, and improve muscle function
 - 2) Early oral feeding POST-OP – avoid long periods of fasting
 - 3) Minimise time on paralytic or ventilation

How can we categorise the types of surgeries

Emergency	Elective	Semi-elective
<ul style="list-style-type: none"> ➤ E1 – (1 hr) – LSCS ED – cord prolapse, severe burn, ACS, shock 	<ul style="list-style-type: none"> ➤ CAT 1 (urgent – within 30 days) – heart valve replacement, limb amputation 	<ul style="list-style-type: none"> ➤ Breast augmentation
<ul style="list-style-type: none"> ➤ E4 – (4 hr) – testicular torsion, threatened limb 	<ul style="list-style-type: none"> ➤ CAT 2 (semi-urgent – within 3 months) – hip replacement, prostate removal, nerve decompression 	<ul style="list-style-type: none"> ➤ Removal of benign tumour
<ul style="list-style-type: none"> ➤ E8 – (8 hrs) – appendicitis 	<ul style="list-style-type: none"> ➤ CAT 3 (Non-urgent – within 1 year) – carpal tunnel release, grommets, varicose vein removal, tonsillectomy, endometriosis 	
<ul style="list-style-type: none"> ➤ E24 – (24 hrs) – abscess 		



COMMON ANAESTHESIA PROCEDURES

ECG - Check for arrhythmias	
➤ Rhythm - AF (most common), SVT	
➤ Tachycardia – pain, drugs, inadequate anaesthesia	
Pulse oximeter = O ₂ SATS AND pulse detected!!	
➤ Early warning of atelectasis, low perfusion	
Capnography = FROM GAS SAMPLER LINE	
➤ gold standard to determine patent airway	
➤ Check for bronchospasm	
NIBP ➔ Check for hypotension causes	
➤ Hypovol?, - fasting, blood loss	
➤ XS anaesthesia,	
➤ Rx: fluid bolus, vasoconstrictors	

AIRWAY MANAGEMENT:

	Procedure	Indications
Endotracheal tube (ETT)	<ul style="list-style-type: none"> flexible plastic tube with an inflatable cuff (balloon) at one end and a connector at the other different sizes (diameter in mm) syringe inflate the cuff via the pilot line Murphy's eye provides an extra hole on the side of the tip that gas can flow through in the event that the main opening at the tip of the ETT becomes occluded (blocked) 	Assistive devices: <ul style="list-style-type: none"> McGrath laryngoscope (camera + screen attached – live feed to visualise vocal cords) bougie used (when vocal cords cannot be visualised) - ETT slides along the bougie into the correct position in the airway and remains there when bougie removed stylet - stiff metal wire (with a plastic coating) to bend tip of ETT anteriorly towards the trachea (to avoid going posteriorly into oesophagus)
Awake fibre-optic intubation	<ul style="list-style-type: none"> ETT inserted while patient conscious under guidance of endoscope 	<ul style="list-style-type: none"> restricted mouth opening or difficult anatomy (e.g., after RT to the neck (since Putting patient to sleep prior to inserting ETT has +++ risk of hypoxia) Trismus – makes intubation more difficult
Supraglottic Airway Devices	<ul style="list-style-type: none"> tip of the SAD located at the top of oesophagus SADs with inflatable cuffs are called laryngeal mask airways (LMA). I-gel is a type of non-inflatable SAD that uses a gel-like cuff that moulds to the larynx 	<ul style="list-style-type: none"> alternative to endotracheal intubation for ventilation first option if intubation fails in a difficult airway scenario
Nasopharyngeal airways	<ul style="list-style-type: none"> measured from the edge of the nostril to the tragus of the ear. 	emergency scenarios, for example, in A&E or at cardiac arrests Cl: base of skull fracture
Oropharyngeal (Guedel)	<ul style="list-style-type: none"> centre of the mouth to the angle of the jaw 	ventilating the patient via a face mask and bag prior to inserting an SAD or ETT
Tracheostomy	<ul style="list-style-type: none"> Performed under GA or in an emergency hole is made in the front of the neck with direct access to the trachea. Tube attached via "trach-tie" Tube may be temporary or permanent <ul style="list-style-type: none"> often inserted at the end of head and neck operations, for example, after a laryngectomy procedure (where a permanent tracheostomy will be required) 	<ul style="list-style-type: none"> Respiratory failure where long-term ventilation may be required (e.g., after an acquired brain injury) Prolonged weaning from mechanical ventilation (e.g., ICU patients that are weak after critical illness) Upper airway obstruction (e.g., by a tumour or head and neck surgery) Management of respiratory secretions (e.g., in patients with paralysis) Reducing the risk of aspiration (e.g., in patients with an unsafe swallow or absent cough reflex)

Difficult airway categorisation

1. Plan A – *laryngoscopy with tracheal intubation*
2. Plan B – *supraglottic airway device*
3. Plan C – *face mask ventilation* and wake the patient up
4. Plan D – *cricothyroidotomy*

CIRCULATORY MANAGEMENT:

	Procedure	Indications
Arterial Line	<ul style="list-style-type: none"> Cannula inserted into an artery (e.g., the radial artery). Cannot give medications 	<ul style="list-style-type: none"> MONITOR BP in real-time Easy ABG
Central Line (CVC)	<ul style="list-style-type: none"> long thin tube inserted into large vein (e.g. IJV, Subclavian or femoral) with tip in SVC "beware of coagulase negative staphylococci (epidermidis)" 	<ul style="list-style-type: none"> Give meds - inotropes, amiodarone or fluids with a high K⁺ concentration (all usually quite irritating if given via PIVC) Take bloods
Vas Cath (CVC)	<ul style="list-style-type: none"> Temporary long thin tube inserted into large vein (e.g. IJV or femoral) with tip in SVC 	<ul style="list-style-type: none"> short-term haemodialysis (in renal failure).
PICC Line (CVC)	<ul style="list-style-type: none"> long, thin tube is inserted into a peripheral vein up to SVC or RA 	<ul style="list-style-type: none"> useful as medium-term IV access – lower infection risk
Tunneled Central Venous Catheter (Hickman)	<ul style="list-style-type: none"> long, thin catheter that enters the skin on the chest, travels through the subcutaneous tissue ("tunneled"), then enters into the subclavian or jugular vein, with a tip that sits in the superior vena cava. 	<ul style="list-style-type: none"> cuff (sleeve) that surrounds the catheter near the skin insertion → promote adhesion to tissue = MORE PERMANENT regular IV treatment (e.g., chemotherapy or haemodialysis).
Portacath (CVC)	<ul style="list-style-type: none"> small chamber (port) under the skin at the top of the chest that is used to access the device subclavian vein, with a tip that sits in SVC or RA 	<ul style="list-style-type: none"> fully internalised under the skin, reducing the chance of infection long-term for regular IV treatment (e.g., chemotherapy).
Pulmonary Artery Catheter (Swan-Ganz catheters)	<ul style="list-style-type: none"> catheter is inserted into the internal jugular vein and ending at pulmonary artery 	<ul style="list-style-type: none"> Measure pulmonary artery wedge pressure (Indication of LA pressures)

*Nb: **coagulase negative staphylococci** (such as *Staphylococcus epidermidis*) = **most common blood isolates** that cause central line infections

Arterial line "Art-line"

- Continuous monitoring – manage unstable patient
- Easy intra-op blood sampling



Entropy (processed EEG + EMG)

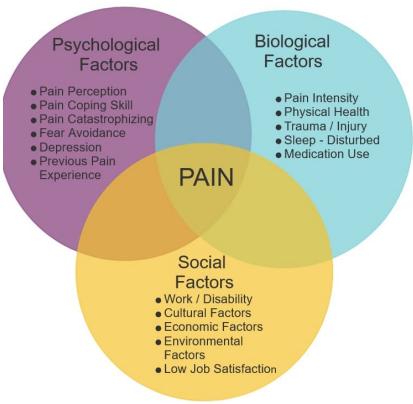
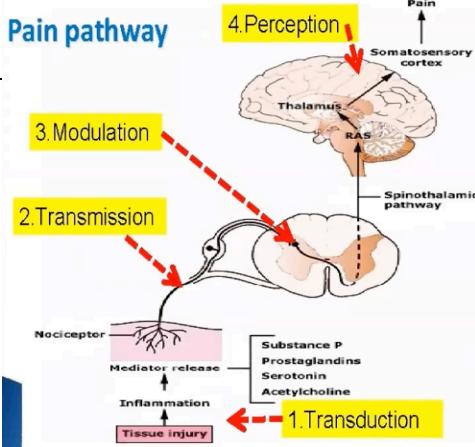
- RE- response entropy (EMG)
- SE – state entropy (EEG)
- Awareness monitoring
- **Depth of anaesthesia**

➤ Maintain pt 10-20% of baseline or flat
➤ **Improve cerebral blood flow** by → raise BP, raise CO₂, reduce head-up

POST-OP PAIN MANAGEMENT

Statements to remember:

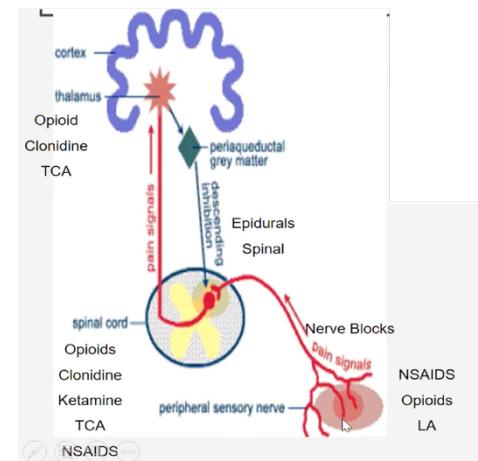
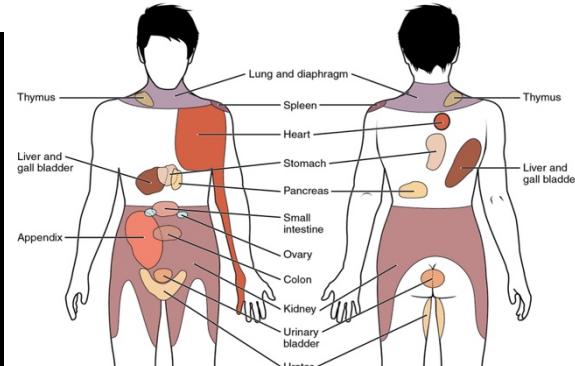
- 1) "Pain is subjective sensory and emotional experience assoc. w/ actual or potential skin damage" IASP 2020
- 2) **Poorly treated post-op pain** is assoc. w/ increased hospital stay, delayed amputation and long-term functional impairment (**Pain leads to disability**)
- 3) It is **NOT** the individual drug or therapeutic modality alone, but the combination and application that makes them effective (**NO panacea**)
- 4) Pain is **ALWAYS** an **OUTPUT** of the brain (even if nociceptors are not firing)
- 5) Rising use of opioids due to over-prescription and substance abuse



Process	Level	MoA	Pathology												
1) Transduction	Peripheral	<ul style="list-style-type: none"> Nociceptors (free nerve endings) → when damaged → release of PG, serotonin → inflammation 	<ul style="list-style-type: none"> Allodynia = pain from non-noxious stimuli 1^o Hyperalgesia = increased pain sensitivity in directly damaged tissue 												
2) Transmission	Peripheral	<ul style="list-style-type: none"> Transmitted by fast A-delta fibres (large), slow C-fibres (thin) Peripheral signal amplified 													
3) Modulation (Gate theory)	Central (spinal cord)	<ul style="list-style-type: none"> Descending pathways (impulses from brain to SC via CST) to inhibit ascending neuron <ul style="list-style-type: none"> Rostral ventral medulla → PAG PAG → release encephalins Nucleus raphe Magnus → release serotonin 	<ul style="list-style-type: none"> 2^o Hyperalgesia = increased pain sensitivity in surrounding undamaged tissues <table border="1"> <tr> <td>Physical</td> <td>Open</td> <td>Close</td> </tr> <tr> <td>Injury extent</td> <td>Medication, counter-stimulation</td> <td></td> </tr> <tr> <td>Emotional</td> <td>Anxiety, worry, tension</td> <td>Relax, + rest</td> </tr> <tr> <td>Mental</td> <td>Pain focus, boredom</td> <td>Working, distracted</td> </tr> </table>	Physical	Open	Close	Injury extent	Medication, counter-stimulation		Emotional	Anxiety, worry, tension	Relax, + rest	Mental	Pain focus, boredom	Working, distracted
Physical	Open	Close													
Injury extent	Medication, counter-stimulation														
Emotional	Anxiety, worry, tension	Relax, + rest													
Mental	Pain focus, boredom	Working, distracted													
4) Perception	Central (brain)	<p><i>Regulated by 2 main pathways + 1 interplay site:</i></p> <ul style="list-style-type: none"> Discriminatory pathway (thalamus + parietal lobe) Mesolimbic (Affective pathway) = AMYGDALA Sensory integration – prefrontal lobe (motivates behavioural change) 	<ul style="list-style-type: none"> Neural plasticity + rewiring → changes perception of pain Damaged inhibitory pathways → new neurons become excitatory (amplify pain) → allodynia, hyperalgesia 												

	Acute pain	Chronic pain = complex regional pain syndrome (CRPS)
Cause	New onset pain (within 3 weeks) <ul style="list-style-type: none"> Injury Infection surgery/trauma 	<ul style="list-style-type: none"> Pain persisting beyond expected time of healing (usu > 3 mths or 12 weeks) <ul style="list-style-type: none"> Chronic primary pain = no underlying condition can explain pain <ul style="list-style-type: none"> IBS, fibromyalgia, Chronic secondary pain = underlying condition to explain pain <ul style="list-style-type: none"> Cancer, neuropathy, migraine, endometriosis, chronic pancreatitis, post-surgical, RA
Assoc.	<ul style="list-style-type: none"> Withdrawal reflex (mainly nociceptive) NOT pathophysiological 	<ul style="list-style-type: none"> Neuropathic + Pathophysiological Assoc. w/: <ul style="list-style-type: none"> Limbic issues (depression, anxiety, PTSD, borderline personality, developmental issues) Advanced age, females, low SES, poor lifestyle Post-op
H+E	<ul style="list-style-type: none"> SOCRATES = Duration + time of day (?↑ICP) + location <ul style="list-style-type: none"> Allodynia = pain from sensory inputs not typically causing pain Pain scale <ul style="list-style-type: none"> (visual analogue 0-10 or faces pain scale Numerical rating scale (0-10) Abbey Pain scale = non-verbal/dementia pts McGill Pain Questionnaire = Pain clinic Mental health screen - ?suicidal Red flags <ul style="list-style-type: none"> Recent traumatic injury (crush fracture, cancer (active vs Rx) - UWL, NS, fever, chills immunocompromised (steroid, transplant IVDU) nocturnal pain/sweats urinary/faecal incontinence (?cauda equina) new onset weakness 	<p>Check current meds (what's worked, what hasn't)</p> <ul style="list-style-type: none"> Any tolerance + A/E Prescription habits - non-compliance, multiple Docs, private doctor scripts Shx: work stress, home environ, financial situation <p>Exams (look, feel, move) - any inconsistencies between hx and clinical features?</p> <ol style="list-style-type: none"> Well-dressed? Eccentric? Sad? Happy? Antalgic gait → Muscle wasting (disuse) Rash / Bruises / inflammation Pain on movement → ROM - reductions Sensation = brush or deep allodynia, numbness, altered cold/heat Psychosocial (beliefs, coping strategies) → DASS 21

	Nociceptive Pain		Neuropathic Pain
	Somatic	Visceral	
Location	Localised	Generalised	Radiating or specific
Character	Pinprick, stabbing, sharp	Ache, pressure, dull	Burning, prickling, electric shock like
MoA	A-delta + periphery (myelinated large diameter)	C-fibre + deep innervation (unmyelinated small diameter)	<ul style="list-style-type: none"> ► Dermatomal (periphery) ► Non-dermatomal (central)
Onset	Fast	Slow	Fast
E.g.	<ul style="list-style-type: none"> Perosteum, joints Sickle cell Superficial laceration, burns, trauma Otitis media Stomatitis 	<ul style="list-style-type: none"> Colic spasm pain (nausea + sweating) Appendicitis (anorexia) Renal colic (extreme) Chronic pancreatitis IBS Angina Periods Burning (sciatica, nerve compression) 	<ul style="list-style-type: none"> Crushing / compressing (Trigeminal, avulsion, post-traumatic neuralgia) Toxins Peripheral neuropathy (HIV, diabetes, alcohol, chemo) Limb amputation - phantom limb pain - cortical remodelling Infection Herpetic Neuralgia
Rx	<ul style="list-style-type: none"> Panadol NSAID Opioids 	<ul style="list-style-type: none"> Panadol NSAID Opioids 	<p>Try each one by itself:</p> <ul style="list-style-type: none"> Amitriptyline (TCA) Duloxetine - an SNRI Gabapentin (anticonvulsant) Pregabalin - (anticonvulsant) <p>Alternative:</p> <ul style="list-style-type: none"> Physiotherapy to maintain strength Tramadol (rescue therapy) Capsaicin - localised pain



COMMON PAIN MEDS (WHO LADDER)

	MoA	Indication	Contraindications	Adverse Effect																
Paracetamol (acetaminophen)	<ul style="list-style-type: none"> Unknown Analgesic & anti-pyretic Given PO or IV (5% glucose + 0.9% NS) 	<ul style="list-style-type: none"> Mild-mod acute pain Reduce dose in elderly or pts < 60kg 	Reduce dose in elderly, small or hepatic involvement	+++BSL (esp. if IV) OD - Hepatotoxicity (15mg/kg/dose qid)																
NSAIDs	<ul style="list-style-type: none"> COX1/2 inhibition = inhibit PG production = Analgesic & anti-pyretic & anti-inflammatory COX1/2 inhibition = Ibuprofen, Diclofenac (Voltaren) COX-2 inhibitor = Parecoxib, celecoxib (minimise GI ulcer but greater VTE risks) 	<ul style="list-style-type: none"> Mild-mod acute pain Migraine Parecoxib = renal stones/ fractures Mg as co-analgesia	<ul style="list-style-type: none"> Asthma Renal impairment (eGFR < 30) Deranged LFT Anuria Uncontrolled HTN PUD/ GIB 	<ul style="list-style-type: none"> Gastritis → PPI or w/ meals Stomach ulcers Asthma exacerbation Renal impairment HTN, CAD 																
Opiates vs Opioids	Activate μ-opioid receptor <table border="1" style="width: 100%; border-collapse: collapse;"> <tr> <th>Short-acting</th> <th>Intermediate acting</th> <th>Long- Acting (start 1st)</th> </tr> <tr> <td>Tramadol (Opioid + SNRI) PO or IV</td> <td>Oxycodone (endone)</td> <td>Targin (Opioid + naloxone)</td> </tr> <tr> <td>Hydromorphone "Jurnista" 1/6TH of total 24 hr bg dose (RESCUE DOSE)</td> <td>Morphine (ordine = soln)</td> <td>MS contin</td> </tr> <tr> <td></td> <td>Palexia (stronger than tramadol)</td> <td>Tapentadol (opioid + SSRI) (Palexia SR)</td> </tr> <tr> <td></td> <td>Buprenorphine Sl or patch (partial opioid)</td> <td></td> </tr> </table> <p>Potency levels: Fentanyl > oxycodone > morphine > codeine > tramadol NB: naloxone can cause pain exacerbation and agitation</p>	Short-acting	Intermediate acting	Long- Acting (start 1 st)	Tramadol (Opioid + SNRI) PO or IV	Oxycodone (endone)	Targin (Opioid + naloxone)	Hydromorphone "Jurnista" 1/6 TH of total 24 hr bg dose (RESCUE DOSE)	Morphine (ordine = soln)	MS contin		Palexia (stronger than tramadol)	Tapentadol (opioid + SSRI) (Palexia SR)		Buprenorphine Sl or patch (partial opioid)		Codeine PO Add prophylactic anti-emetics for nausea	Dose Panamax co Panadiene Forte 500/30 (30-60mg every 4-6 hrs)	Indications Pregnancy	CI Constipation = SBO/LBO (10% poor metabolisers) Hepatic failure (prodrug) Drowsy, confused, miosis, myoclonus, dry mouth Sedation, resp. depression
Short-acting	Intermediate acting	Long- Acting (start 1 st)																		
Tramadol (Opioid + SNRI) PO or IV	Oxycodone (endone)	Targin (Opioid + naloxone)																		
Hydromorphone "Jurnista" 1/6 TH of total 24 hr bg dose (RESCUE DOSE)	Morphine (ordine = soln)	MS contin																		
	Palexia (stronger than tramadol)	Tapentadol (opioid + SSRI) (Palexia SR)																		
	Buprenorphine Sl or patch (partial opioid)																			
Triptan	<ul style="list-style-type: none"> Serotonin (5-HT) receptor agonists Sumatriptan (oral 30%, 60% injectable) 	Migraine ONLY	MOH	MOH																
TCAs & SNRI	<ul style="list-style-type: none"> Inhibit NA and 5-HT re-uptake → inhibits nociception TCA: amitriptyline, nortriptyline SNRI: duloxetine, venlafaxine 	<ul style="list-style-type: none"> Used mainly for mood disorders Neuropathic pain Chronic primary pain 	SSRI are ineffective Serotonin syndrome	<ul style="list-style-type: none"> TCA: anti-SLUDGE SNRI: fatigue, constipation, insomnia, anorexia 																
Gabapentin & Pregabalin	<u>Neuropathic agents</u> <ul style="list-style-type: none"> Inhibit voltage gated Ca channel → reduce neurotransmitter release 	<ul style="list-style-type: none"> Post-herpetic neuralgia Diabetic neuropathy Neuropathic pain MSK fracture pain 	NOT for central neuropathic pain or HIV associated neuropathy	<ul style="list-style-type: none"> Sedation Ataxia 																
Carbamazepine	Inhibit voltage gated Na channel → reduce neuronal excitability	Mostly for trigeminal neuralgia (unproven)	pregnancy, bone marrow failure																	
Ketamine	<ul style="list-style-type: none"> NMDA receptors antagonist Anaesthesia without resp. depression 	<ul style="list-style-type: none"> ED – trauma pt and sedation Peri-op to reduce post-op pain (e.g. MSK fracture pain) 		<ul style="list-style-type: none"> Dissociation Polyuria, freq. urgency (cystitis mimic) 																
Lidocaine	<ul style="list-style-type: none"> Na channel blocker 	<ul style="list-style-type: none"> Renal colic Local, epidurals 																		

NB: good pain management involves **lifestyle changes** and **MDT** (pain specialists, physios, GPs, social workers psychologists)!! NOT just meds

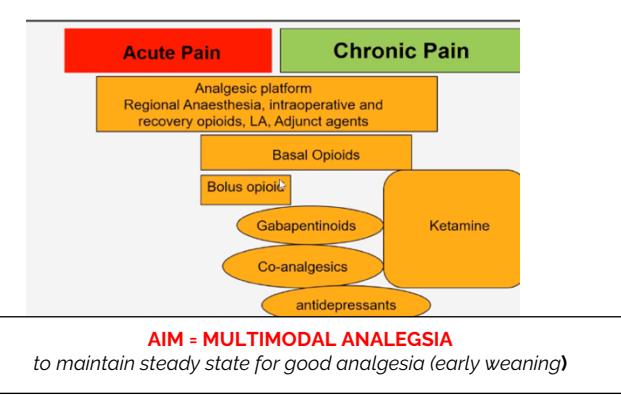
- CONTACT APS and PALL Care if unsure about baseline and breakthrough doses (ALSO time limit endone when giving script)**
- Give Mg as co-analgesia to NSAID and PCT to reduce opioid usage + Laxatives**
- Avoid NSAID, paracetamol, opiates and anti-convulsant for chronic PRIMARY pain → use TCA and SNRI**

Common exam questions:

- "This patient is on 30mg of modified-release morphine every 12 hours; what would be the correct breakthrough dose?"** In this scenario, 10mg is the correct answer, as the patient is getting 60mg background morphine every 24 hours (30mg twice a day).
- "Patient A takes the same amount of morphine as patient B but claims there is no change in their pain level? Explain?"** Patient A may be a poor acetylator, hence cannot breakdown codeine (prodrug) to get active component. Otherwise, is Patient A taking it as prescribed or is their an underlying cause

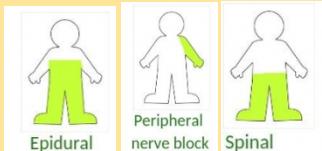
Opioid	Route	Equivalent Dose
Morphine (Mod. Release)	Oral	10mg
Morphine	IV / IM / SC	5mg (20-30mg / daily + 5mg breakthrough)
Codeine	Oral	100mg
Tramadol	Oral	100mg
Oxycodone	Oral	6.6mg
Diamorphine	IV / IM / SC	3mg
Buprenorphine	patch	5mcg/hr = 12mg / 24 hr oral morphine
Fentanyl (for CKD)	Patch Intranasal	12mcg/hr = 30mg / 24 hr oral morphine For children

ANALGESIA PROTOCOL (PERIOPERATIVE)



Types of Anaesthesia

"reversible loss of all sensation and LOC"



Local

Regional

General

Define	Numb single area	Blocking sensation to particular site	Making patient unconscious												
Pre-op	<ul style="list-style-type: none"> ➤ NO fasting needed ➤ Check allergies <table border="1" style="margin-top: 5px; border-collapse: collapse; text-align: center;"> <thead> <tr> <th></th> <th>Half-life (mins)</th> <th>Toxic Dose (mg/kg)</th> </tr> </thead> <tbody> <tr> <td>Lignocaine</td> <td>100</td> <td>2-7</td> </tr> <tr> <td>Ropivacaine</td> <td>120</td> <td>3-4</td> </tr> <tr> <td>Bupivacaine</td> <td>160</td> <td>2</td> </tr> </tbody> </table>		Half-life (mins)	Toxic Dose (mg/kg)	Lignocaine	100	2-7	Ropivacaine	120	3-4	Bupivacaine	160	2	<ul style="list-style-type: none"> ➤ NO fasting needed ➤ Check allergies 	<p>Fasting prior to operation:</p> <ul style="list-style-type: none"> ➤ 6 hours of no food or feed before operation ➤ 2 hours of no clear fluids (i.e. fully "nil by mouth") <p>Why do you need empty stomach (NBM)?</p> <ul style="list-style-type: none"> ➤ Risk of aspiration pneumonia + pneumonitis ➤ Inc. ED patients, non-fasting, full stomach
	Half-life (mins)	Toxic Dose (mg/kg)													
Lignocaine	100	2-7													
Ropivacaine	120	3-4													
Bupivacaine	160	2													
Types	<ul style="list-style-type: none"> • Topical • Infiltration <p>Indications:</p> <ul style="list-style-type: none"> ➤ Skin sutures ➤ Skin biopsies ➤ Dental procedures ➤ Minor surgery (carpal tunnel) ➤ Insertion central line ➤ Percutaneous procedures (e.g. PCI) ➤ LP 	<ol style="list-style-type: none"> 1) Peripheral nerve block (under USS guidance) <ol style="list-style-type: none"> a. Brachial plexus b. Femoral nerve (neck of femur #, TKR) 2) Neuraxial block <ol style="list-style-type: none"> a. Epidural → C/S, severe lung disease, lower limb surgery, child delivery b. Spinal – no catheter → lower limb and pelvic surgeries, LSCS c. A/E = haematoma, infection, hypoTN, local anaesthetic toxicity 3) IV regional/peripheral → Bier's block (19th century – Bohr war → IV local anaesthetic given with tourniquet to prevent systemic spread → only distal areas anaesthetized (Best for forearm injuries)) <p>Monitored anaesthetics care (MACS):</p> <ul style="list-style-type: none"> ➤ Conscious BUT calm/sedated state (e.g. benzos) 	<ol style="list-style-type: none"> 1) Pre-medication – to make intubation easier: <ol style="list-style-type: none"> a. Benzodiazepines (e.g., midazolam) = relax muscles and reduce anxiety (also causes amnesia) b. Opiates (e.g., fentanyl or alfentanil) = reduce pain and hypertensive response to the laryngoscope c. Alpha-2-adrenergic agonists (e.g., clonidine), = for sedation and pain 2) Total IV or parenteral anaesthesia (TIVA) – analgo-sedation <ol style="list-style-type: none"> a. Propofol → allosteric agonist of GABAa receptor b. Etidomide → if haem unstable (less impact on CVS) c. Barbiturates (thiopental) → brain surgery d. Benzodiazepines (midazolam) e. Opioids (fentanyl) 3) Total inhalation anaesthesia – ether analgesia <ol style="list-style-type: none"> a. Gas (NO given w/ Oxygen) - Euphoric effect, analgesia i. Indications = pregnancy, child, casualty ! b. Volatile anaesthesia (sevoflurane, isoflurane) 												
A/E	<p>A/E of Local and Regional Anaesthetics:</p> <p>Systemic toxicity (anaesthetic leak into blood NOT nerve)</p> <ul style="list-style-type: none"> ➤ Prodrome = metallic taste, facial numbness ➤ CNS (Crosses BBB) → confused, seizures, resp. depression → resp. acidosis + hypoventilation ➤ CVS toxicity → (1) initially HTN, tachycardia → (2) gradual slow rate of fast conducting pathways (purkinje) → (3) hypoTN, bradycardia, unopposed ventricular contraction → VF/VT → cardiac arrest 	<p>Commonly:</p> <ul style="list-style-type: none"> ➤ IV meds are induction agents (i.e. induce unconsciousness), ➤ inhaled medications for maintaining GA during operation (since they take time to reach effective concentration since they need to diffuse across the lung tissue and into the blood) ➤ Total intravenous anaesthesia (TIVA) uses IV meds for BOTH induction and maintenance of GA (Propofol) → nicer recovery (as they wake up) compared with inhaled options. <p>Emergence:</p> <ul style="list-style-type: none"> ➤ Wean off muscle relaxant – to avoid "awareness under anaesthesia" ➤ Nerve stimulator on either ulnar nerve (thumb twitch) or facial nerve (orbicularis oculi twitch) → if response gets weaker with more stimulation → indicates that muscle relaxant not worn off ➤ Consider reversal agent e.g. sugammadex <p>A/E of GA:</p> <ul style="list-style-type: none"> ➤ Post-op sore throat, nausea and vomiting ➤ Aspiration ➤ Dental injury ➤ Anaphylaxis ➤ Cardiovascular events (e.g. MI, stroke, arrhythmia) ➤ Death ➤ Malignant hyperthermia (rare BUT fatal) <ul style="list-style-type: none"> ○ Assoc. w/ suxamethonium and volatile anaesthetics ○ Hypermetabolic response to anaesthesia → febrile, tachycardia, acidosis, hyperK, rigidity, XS CO₂ production ○ Rx: dantrolene (interrupts muscle rigidity by disrupting Ca ion movement in SKM) 													
Rx	<p>Treatment for systemic toxicity</p> <ol style="list-style-type: none"> 1) Stop injection 2) BOLUS interlipid therapy (stop protein binding) 3) Ventilate – FiO₂ 4) Stop cerebral excitation → benzo, propofol, barbiturate 5) Correct HypoTN + arrhythmia → IVF, ionotrope, vasopressor, anti-arrhythmic 6) Monitor for hypoxia + acidosis 														

EPIDURAL VS SPINAL ANAESTHESIA

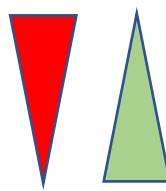
Desired dermatome level of neuraxial block

Type of surgery	Upper dermatomal block level	Anatomical landmark	Optimal insertion point
Oesophagus, lung	T1	Below clavicle	T6-7
Upper abdomen	T1	Below clavicle	T9-10
Lower abdomen	T6	Distal sternum	T9-10
Caesarean delivery	T4	Nipples	L4-5
Lower limb	L1-2	Inguinal crease	L4-5

WHAT SENSATIONS LOST FIRST? [ORDER OF BLOCKADE]

Thin nerve fibres blocked 1st before thick fibres (i.e. sensory BEFORE motor)

- 1) ANS
- 2) Pain
- 3) Temp
- 4) Touch
- 5) Deep pressure
- 6) Motor



Recovery in REVERSE ORDER

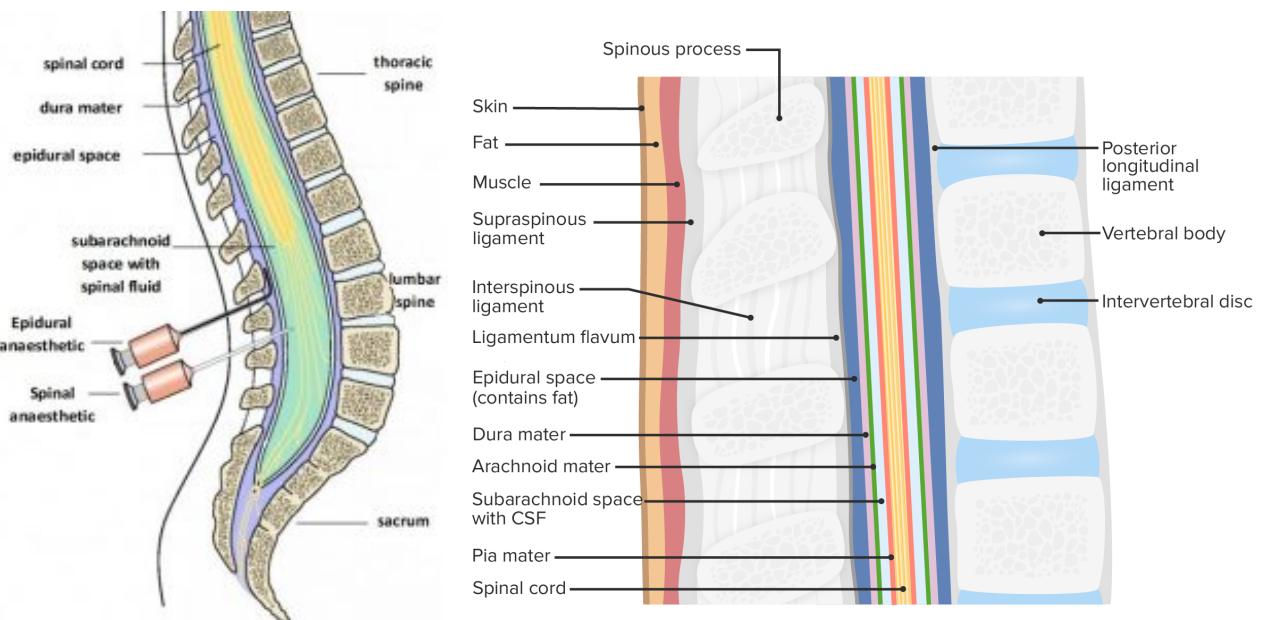
MAIN CONTRAINDICATIONS

- 5) ANTI-COAG usage (warfarin, coagulopathy)
- 6) Anaphylaxis
- 7) Thrombocytopenia
- 8) Active bleed/ infection
- 9) Raised ICP

Central Neuraxial Anaesthesia (spinal block)

Epidural Anaesthesia

Type	regional anaesthesia.	regional anaesthesia.
Indication	<ul style="list-style-type: none"> Caesarean sections – cannot feel legs Transurethral resection of the prostate (TURP) Hip fracture repairs 	<ul style="list-style-type: none"> pregnant women in labour <ul style="list-style-type: none"> can still move legs and feel contractions but pain fibres are numbed post-operatively after a laparotomy (open abdominal surgery).
Duration	<ul style="list-style-type: none"> 1-3 hours 	Continuous infusion (monitoring may be needed)
Procedure	<ul style="list-style-type: none"> Conscious patient local anaesthetic injected into CSF within the subarachnoid space (L3/4 or L4/5 spaces) Test w/ cold ice cube to determine if it is working (dermatomes) 	<ul style="list-style-type: none"> Conscious patient small tube (catheter) into the epidural space in the lower back (outside dura mater) → cervical, thoracic, lumbar or sacral Diffuse to the surrounding tissues and spinal nerve roots can be converted into spinal block if needed
Meds	<ul style="list-style-type: none"> Morphine (hydrophilic) 	Local anaesthetic meds (e.g. Levobupivacaine +/- fentanyl) <ul style="list-style-type: none"> Fentanyl - Longer acting and lipophilic → diffused into grey matter
Dosage / onset	<ul style="list-style-type: none"> Small + quick onset 	Large + slow onset
A/E	<ul style="list-style-type: none"> Numbness and paralysis of the areas innervated by the spinal nerves below the level of the injection Cardiac arrest 	<ul style="list-style-type: none"> Headache if the dura is punctured, creating a hole for CSF to leak from ("dural tap") – Rx: IVF Motor weakness in the legs (cannot straight leg raise) → incorrect site (catheter may be in SAS not epidural) → urgent anaesthetic review Infection, including meningitis Epidural Haematoma (? spinal cord compression) Prolonged 2nd stage + increased probability of instrumental delivery



COMMON ANAESTHESIA MEDICATIONS

	Type	MoA	Dosage	Indication	A/E
Halothanes Sevoflurane (pleasant odour) Desflurane (pungent odour)	Volatile agents	Unknown ➤ Reduces minute ventilation ➤ Bronchodilator	Sevo (2% MAC) – 2% metabolised Des (6% MAC) – wake up quicker	MAC = min alveolar conc. to prevent reaction to surgical stimulus in 50% of patients at 1 atm pressure	HTN, tachycardia Malignant Hyperthermia Bronchoconstrict (if Des)
Nitrous Oxide "beware recreational use"	Volatile sparing agent	Stops methionine synthetase to produce analgesia ➤ Minimise prolonged use	High MAC (102%)	Simple procedures	Malignant Hyperthermia Megaloblastic anaemia (vit B12 def) Agranulocytosis
Oxygen	Volatile agents				ROP
Propofol (white lipid soln)	IV induction or sleeping agent (emulsion / egg)	Non-barbiturate GABA modulator ➤ ↓SVR + ↓ myocardial contractility ➤ ↓ CO (25%) ➤ ↓ Laryngeal reflex	1 - 2mg/kg (rapid onset)	➤ GA ➤ Anti-emetic (5-HT3 Receptor)	➤ Anaphylaxis to egg/soy ➤ Reduced airway reflexes ➤ HypoTN ➤ Stings when injected
Thiopentane	IV induction	barbiturate GABA agonist ➤ Reduces cerebral consumption	3-7mg/kg (5-10mins)	Brain surgery, status epilepticus, raised ICP	Tachycardia Vasodilated (↓SVR)
Ketamine "special K" or "K-hole"	IV induction dissociative /conscious anaesthesia + analgesic	• Phencyclidine derivative created from Vietnam War → (NMDA receptor antag) • ↑ SNS → ↑HR, ↑CO, ↑BP, ↑ Cardiac O2 cons.	1-2mg/kg 0.2-0.5 mg/kg analgesia	➤ Trauma pt ➤ Seizures	HypoTN → reduce dose NOW
Midazolam	Sedative	GABA receptor agonist (benzo)	IV 0.02-0.1mg/kg	Sedation + anxiolysis	Headache, delirium, N/V, drowsy
Adrenaline	Vasopressor • Lower dose = vasodilate (B1, B2) • Higher dose = vasoconstrict (α1)	• A1 vasoconstrictor (2-3 min half life) • ↑HR, ↑BP, ↑RR, O2 cons., Glycogenolysis, Lipolysis	0.01-0.1 mg/kg/min IV in ICU Or 1mg ED (ALS) Or IM 1:10000	➤ HypoTN ➤ Anaphylaxis ➤ Severe asthma (bronchodilator)	Lactic acidosis Higher mortality in HF patients
Ephedrine	Vasopressor • Indirect SNS – release NORAD	• Potent A1 (NORAD) AND B activity • ↑HR, ↑CO, ↑RR, O2 cons., bronchodilate	IV bolus ONLY	HypoTN	Higher mortality
Phenylephrine	Vasopressor • last 5 minutes	• Selective A1 vasoconstrictor • ↑SVR → ↑BP → ↑Coronary artery flow	IV bolus or infusion	HypoTN	reflex bradycardia (↑SVR)
Metaraminol	Vasopressor • Lasts 20-60mins • Longer acting	• A1 vasoconstrictor (mainly) • ↑SVR → ↑BP → ↑Coronary artery flow, + pulm. Vasc. pressure	IV bolus or infusion	Acute HypoTN	➤ reflex bradycardia (↑SVR) ➤ Higher mortality
Vasopressin	Vasopressor last 5 minutes	• Selective V1 vasoconstrictor • ↑SVR → ↑BP → ↑Coronary artery flow	IV via CVC		
Levosimendan	Positive Inotrope (increase contractility)	• PDE3 inhibitor	IV via CVC	mainly cardiac pts	
Milrinone	Positive Inotrope (increase contractility)	• Increases heart muscle sensitivity to calcium	IV via CVC		
-curium or -curonium (e.g. rocuronium)	Muscle relaxant / paralytic	Non-depolarising neuromuscular blocking agent → Competitive inhibitors against nACh _{Rs} • Slower action → muscle relaxant	IV	• Keep Ventilation • Prevent asp. pneumonia if NOT fasted	Muscle pain and salivation Reversal agent = sugammadex 16mg/kg
Suxamethonium	Muscle relaxant/ paralytic • Check K levels • Caution in burns pt	• Rapid onset • Depolarising NMJ blocking agent Metabolised by pseudocholinesterase	IV	Initially causes twitches and fasciculations before relaxation • Raises IOP	Hyper K, hypoTN, fever CI = malignant hyperthermia, glaucoma OR eye trauma Reversal agent = neostigmine
Neostigmine	Reversal agent	Inhibits AChE at NMJ → ↑ACh	Given with anti-chol (atropine)	Reverse paralytic	Cholinergic effects (bronchospasm, bradycardia)
Glycopyrrolate	Anti-cholinergic / anti-muscarinic	↓Ach → ↑HR (mildly increase)	200mcg	Mild Bradycardia	Dry mouth, blurred vision, urinary retention, drowsy ➤ Non-opposed-tachycardia
Atropine	Anti-cholinergic	↓Ach (acts on SA node) → ↑HR (greater) Blocks CNX on heart	ED = 3mg	Bradycardia	
Parcoxib	Analgesic - NSAID	COX-2 selective			
Remifentanil	Analgesic - opioid	Ultra SA opioid • Fast onset + offset of action	IV	ICU or theatre for tube tolerance	
Fentanyl	Analgesic – opioid	SA + potent opioid • Lipid soluble → diffuse	5-15mcg	Analgesia → spinal blocks OR patch epidurals	
Alfentanil	Analgesic – opioid	V. short acting opioid • Less potent than fentanyl (10 min duration of action)		Analgesia for shorter cases or bridging	
Morphine	Analgesic – opioid	Gold standard analgesic • Hydrophilic	100-300mcg	analgesia	
Procaines (cocaine, tetracaine, benzocaine)	<u>Ester</u> -linkage LA	➤ Short-acting (broken down by plasma esterase) ➤ Vasodilatation		Short duration analgesia w/ slow onset of action	
Bupivacaine	<u>Amide</u> -linkage LA	Long-acting (broken down by liver enzymes) ➤ High plasma protein binding = longer duration of action ➤ High potency (as high solubility)		Nerve block or Regional anaesthetics	
Lidocaine	<u>Amide</u> -linkage LA + anti-arrhythmic			RA onset of action for Short procedures: ➤ Lower pKa = more ionised = faster action	
Penicillins	Antibiotic	• Amoxil = B-lactam (prevent PG synthesis by cross-linking) • Co-amoxiclav = + B-lactamase inhibitor	500mg qid PO/IV 1.2 g IV	Skin- cellulitis, impetigo, diabetic foot, mastitis Otitis media, URTI, UTI	ATN
Cephalosporins	Antibiotic	Cefuroxime = 2 nd gen Cephalosporin (only 2 nd gen that crosses BBB)	IV 2g	Broad spectrum (g +ve and g -ve)	ATN
Linosamides	Antibiotic	Clindamycin		2 nd line if penicillin or cephalosporin allergy	
Aminoglycosides	Antibiotic	Gentamicin		UTI? UROLOGY	Nephrotoxic, ototoxic
Oxytocin	Obstetric (syntocinon)	• ↑uterine contractions		induce labour or post-LCSC bleed (PPH)	
Tranexamic acid	Antifibrinolytic	• Stop clot breakdown		Trauma, surgery, ortho, menorrhagia	VTE
Ondansetron	Anti-emetic (PONV)	5HT3 receptor in CTZ → vomiting centre ➤ Do ECG prior (?prolonged QT)	4mg IV	N+ V	Fatigue, drowsy, constipation, prolonged long QT
Metoclopramide	Anti-emetic (PONV)	D2 receptor antag + prokinetic			Caution in sedation, constipation, prolonged QT
Dexamethasone	Anti-emetic (PONV)	40-50x more potent than hydrocortisone and longer lasting		N+ V, odynophagia, systemic illness	Steroid issues – caution in DM and immunocomp.
Cyclizine	Anti-emetic (PONV)	H1 antag			Caution w/ HF and elderly

INTRA-OP CONSIDERATIONS: Pain, TEMP, PRESSURE SORES,

Concern		Solutions
Pain	CVS – HTN, tachycardia, DVT Resp – atelectasis, impaired cough, pneumonia GI – reduced gastric motility Neuroendocrine – impaired wound healing, increased catabolic hormones (++) BSL)	<ul style="list-style-type: none"> ➤ Choose appropriate analgesia → dosage and duration
Temperature	Anaesthesia causes loss of thermoregulation <ul style="list-style-type: none"> ➤ Heat loss by conduction, radiation, convection 	<ul style="list-style-type: none"> ➤ Pre-warming = best (bair hugger, warm blankets) ➤ Warmed fluids (esp. blood products, fluids) ➤ Theatre temperature control (esp. in paediatrics!!)
Pressure area	Long surgical time	<ul style="list-style-type: none"> ➤ Patient positioning – protect skin and nerves ➤ Padding – surgical table, heels ➤ Protect vulnerable nerves (e.g. median, ulnar – thumb's up)

POST-OP NAUSEA & VOMITING (PONV) MANAGEMENT

PONV risk factors: **APFEL criteria**

- Assume all patients have 25% have post-operative
- Each 1 will increase by 25% = (1) **Hx of PONV**, (2) **female**, (3) **non-smoker**, (4) **Patient will need opiate post-op**

Patient	Female > Male, ratio 2.5 : 1	Surgical	ENT, especially middle ear operations, adenoids and tonsillectomy
	Anxiety		Squint surgery
	Previous history of PONV		Gynaecological surgery
	History of motion sickness		Gastrointestinal surgery
	Non-smoker		Laparoscopic procedures
	Pain		Intestinal obstruction
	Presence of gastric contents		
Anaesthetic	Volatile agents	Medical	Hypoxia
	Nitrous oxide		Uraemia
	Opioids		Metabolic disorders, e.g. hypoglycaemia, hypercalcaemia
	Intravenous anaesthetics (ketamine, etomidate)		
	Neostigmine		
	Stomach insufflation		
	Spinal anaesthetic (with hypotension)		

RARE BUT IMPORTANT MALIGNANT SYNDROMES

	Cause	Sx	Rx
Neuroleptic malignant syndrome	<ul style="list-style-type: none"> ➤ dopamine receptor-blocking drugs → antipsychotic agents (e.g. chlorpromazine, haloperidol) and some antiemetics (metoclopramide). ➤ OR sudden withdrawal from dopaminergic drug (e.g. levodopa, bromocriptine) 	<ul style="list-style-type: none"> ➤ muscle rigidity, ➤ bradypreflexia, ➤ bradykinesia, ➤ altered mental status, ➤ EPSE ➤ hyperthermia. 	bromocriptine (a dopamine agonist) PO or NGT <ul style="list-style-type: none"> ➤ Start at 2.5 mg 8-hourly, > ➤ increasing to 5 mg every 4 h (maximum 30 mg/day) in moderate to severe cases
Malignant hyperthermia syndrome	➤ autosomal dominant disorder after receiving an inhalational anaesthetic or suxamethonium	<ul style="list-style-type: none"> ➤ muscular rigidity, ➤ tachypnoea, ➤ tachycardia, ➤ hypertension, ➤ mottled diaphoretic skin ➤ cardiac arrhythmias. 	dantrolene 1 mg/kg IV for severe muscle rigidity and hyperthermia. <ul style="list-style-type: none"> ➤ Further doses of 1–2.5 mg/kg (up to a maximum of 10 mg/kg/24 h) may be required

ACUTE RESPIRATORY DISTRESS SYNDROME (ARDS)

1. Acute onset < 1 week
2. Predisposing condition
3. bilateral opacities consistent with APO must be present (detected via chest CT or CXR)
4. PF ratio < 300mmHg with a minimum of 5 cmH₂O PEEP
5. must not be fully explained by cardiac failure or fluid overload.

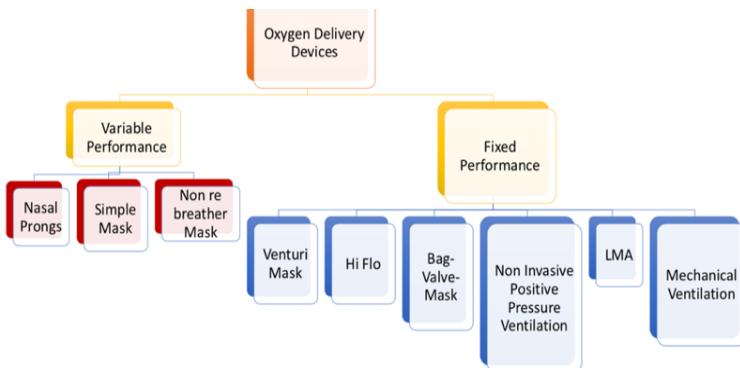
RESPIRATORY SUPPORT

Targeted Oxygen Therapy

In the absence of COPD	In the presence of COPD
Where SpO ₂ is <92% ➤ titrate oxygen to maintain SpO ₂ 92-96%	With a SpO ₂ <88% ➤ titrate oxygen to achieve a SpO ₂ 88-92% using a Venturi mask delivering a FiO ₂ 24%-28%

What/When to use?

- **Mainly for acute respiratory distress syndrome (ARDS)**
- Respiratory support ONLY buys time until underlying problems can be managed
- Chest physiotherapy and suction = helps to clear secretions and improve respiratory function
- **Remember O₂ is a drug (XS oxygen → can lead to ROP)**



Acute respiratory Distress:

- Atelectasis (alveolar collapse)
- APO (NOT due to fluid overload)
- Reduced lung compliance
- Fibrosis of lung tissue

Investigations

- ABG - Hypoxia
- CXR: bilateral infiltrates

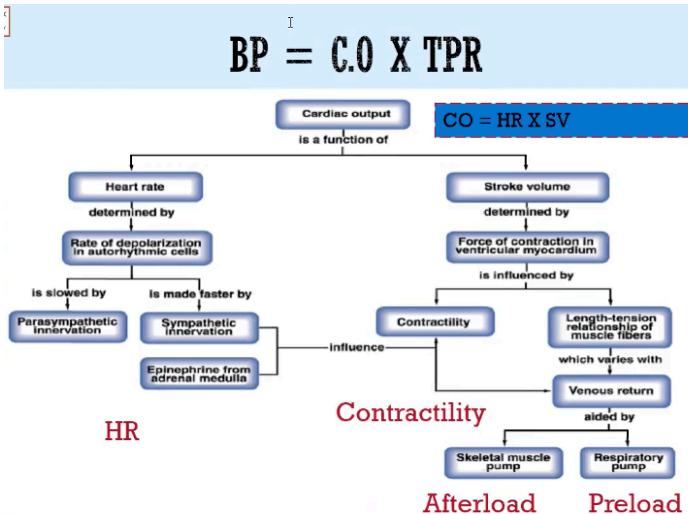
General Mx:

- 1) Respiratory support
- 2) Prone positioning (lying on front)
- 3) Fluid and electrolyte management (avoid XS fluid in lung)

	Minimally invasive	Flow Rate (L/min)	FiO ₂ (%)	Indication for use
	Nasal Cannula/Prongs	1-4	25-35	<ul style="list-style-type: none"> • Minimal resp. distress (~90%) • Less claustrophobic, low cost, can eat and speak
	Simple Face Mask	5-10	40-60	<ul style="list-style-type: none"> • Moderately hypoxic • Risk of CO₂ retention
	Non-Rebreathing Mask *Reservoir bag MUST be inflated + tight fit needed	15	> 60%	<ul style="list-style-type: none"> • Post-cardiac or respiratory arrest • Severely hypoxic patients <p>*Patients have adequate ventilation but require high O₂ concentrations</p>
	Venturi Mask (blue - white- orange - yel -red-green) (2 -----4 -----6 -----8 --10 --15)	2-10 [adaptor]	24-60% (fixed)	<ul style="list-style-type: none"> • COPD patients (i.e. avoid CO₂ retention) or patients using hypoxic drive to breath • Transition to non-rebreathing mask in emergency or if patient is not well
	Hi Flow Nasal Cannula "humidified O ₂ delivery under pressure"	High flow nasal cannula: ≤60	0.24-0.8	<ul style="list-style-type: none"> • Low level prep → applied nasally or via tracheostomy • Increases FiO₂ + provides PEEP
	Bag-Valve- Mask Resuscitator	15	>0.8	<ul style="list-style-type: none"> • Need adjunct → nasopharyngeal or oropharyngeal (Guedel) airway or BOTH <ul style="list-style-type: none"> ◦ Place upright horizontally in (avoid pushing tongue back into oropharynx) → rotate when at oropharynx • Need to optimize patient position, bed height etc. <div style="text-align: right;">  <p>(oropharyngeal) (guedel)</p> </div>
	Non-invasive positive end expiratory pressure (PEEP) (CPAP, BPAP)			<p>Prevent airway collapse → reduce atelectasis and improve alveolar ventilation to recruit more alveoli for more gas exchange and decrease WoB</p> <ul style="list-style-type: none"> • CPAP = T₁RF - constant pressure to keep airways expanded) • BPAP = T₂RF = inspiratory positive airway pressure (air forced into lungs) and expiratory PEEP (prevent airways collapse)
	Mechanical ventilation			<ul style="list-style-type: none"> • Endo-tracheal tube = definitive airway to maintain airway patency • Control FiO₂, RR, TV, Peak flow rate, PEEP • Tracheostomy (emergency)
Adjuncts	LMA (laryngeal mask airway)			<ul style="list-style-type: none"> • Non-definitive airway
Dynamic manoeuvres	Prone Position			<ul style="list-style-type: none"> • Reduce compression of lungs by adjacent organs • Improve blood flow to lungs - esp. to well-ventilated areas • Improve overall oxygenation and secretion clearance • Reduce assistance needed from Mechanical ventilation

Fraction of Inspired Oxygen (FiO₂) concentration ≈ (Flow Rate x 4) + 21 (FiO₂ in room air)

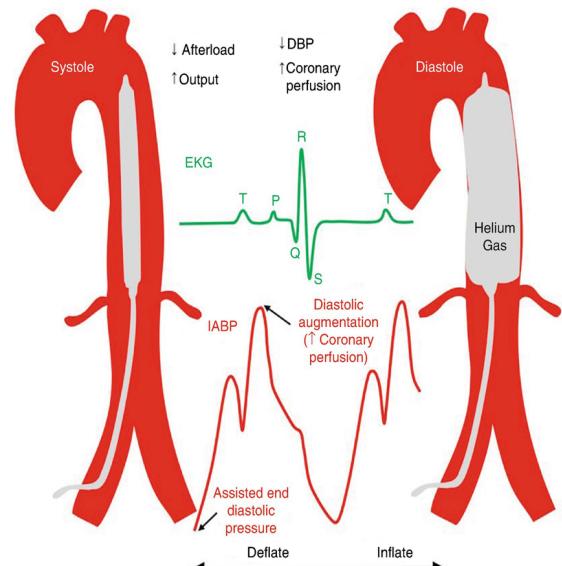
CARDIOVASCULAR SUPPORT



	Decrease	increased
HR	PSNS = opiate	SNS = Shock, stress
	Inotropes	BB
Preload (CVP = RAP) = heart muscle tension or pressure filling ventricle (central venous pressure)	Volume loss = bleeding	↑ vol to heart = ↑RA = ↑EDV (e.g. pregnancy)
	IV fluid or MTP	Diuresis
Afterload (≈TPR or intrathoracic pressure) = force needed to pump against aorta (aortic pressure)	MR Vasodilating (e.g. Anaphylaxis)	Viscous blood L-side (systemic HTN, AR, MR) R side (PHTN)
	Vasopressors	Anti-coags,
Contractility = affected by HR, preload, afterload	Ischaemia	SNS
	Inotropes	BB, CaB
Venous return	Vasopressors	Resp. pump SKM pump SNS systemic veins Abdo compression reflex

Monitoring:	
➤ Non-invasive	
1) HR	
2) Peripheral BP	
3) Pulse Oximeter	
4) Continuous ECG	
➤ Invasive	
1) ABG (arterial line)	
2) Central venous pressure (preload) and O ₂ sats (via CVC)	
3) Pulmonary artery O ₂ sats and wedge pressure (pulmonary artery catheter)	
4) ECHO (TTE, vs TOE)	
➤ Intensive:	
1) Oesophageal doppler monitor (assess blood flow through thoracic aorta to measure SV and CO)	
Other cardiac interventions:	
• Impella	
• Swan ganz catheter	
• ECMO → replace heart + lung	
• Cardio-pulm bypass circuit	

General Treatment for Hypotension:	
➤ Increase contractility → positive inotropes (e.g. milrinone, levosimendan, dobutamine, adrenaline, isoprenaline)	
➤ Increase afterload → vasopressors (e.g. adrenaline, vasopressin)	
General treatment for bradycardia:	
➤ Reduce PSNS – e.g. atropine, glycopyrrolate	
General treatment for tachycardia:	
➤ BB, CaB, Flecainide	
When do we use an intra-aortic balloon pump?	
➤ Cardiogenic shock	
➤ ACS (UA, NSTEMI, STEMI)	
➤ Immediately after heart surgery	
How does it work?	
➤ Inserted via femoral artery up to descending thoracic aorta → linked to special machine synchronized with heart contractions	
➤ Inflate balloon during diastole (relaxed heart) push blood into coronary artery to (1) improve coronary perfusion	
➤ Deflate during systole (2) reduce afterload and (3) increase cardiac output	



RENAL SUPPORT – RENAL FAILURE

INDICATIONS:

- **A** – acidosis (pH < 7.1)
- **E** – electrolyte (hypeK)
- **I** – ingestion of drugs
- **O** – overload fluids
- **U** – ureamic encephalopathy

AIM: Removing excess fluid, solutes and waste products

Types of dialysis used in ICU:

- **Continuous renal replacement therapy (RRT)** involves continuously performing dialysis 24 hours a day.
 - 1) Most patients requiring haemodialysis in ICU will be on CRRT.
- **Intermittent haemodialysis** involves running the machine and performing dialysis for set periods, for example, 3-12 hours, before taking a break from dialysis.

Considerations for dialysis:

- Need 2-lumen CVC (Vas Cath) to have abundant blood supply access
- Anti-coag with citrate of heparin
- Dialysate (the fluid where solutes filter out from blood) into

POST-OP COMPLICATIONS

Immediate	Early
<p>1. Anaesthetic</p> <ul style="list-style-type: none"> ○ <u>CVS</u>: arrhythmia, HTN, HypoTN, MI/Stroke, ○ <u>RESP</u>: irregular RR ○ <u>Hypothermia (lost due to conduction, radiation, convection)</u> ○ <u>General</u>: teeth/lip/tongue damage, hyperthermia, allergies, <p>2. Haemorrhage – check drain, obs, FBC, HCT</p>	<p>1. Dehydration</p> <p>2. Electrolyte imbalance</p> <p>3. Infection – systemic Vs. local</p> <p>4. VTE</p> <p>5. Wound or anastomotic breakdown</p> <p>6. Bed sores</p> <p>7. Delirium (4AT – identify cause)</p> <p>8. Post-op ileus</p> <p>9. Post-op falls</p>

SPECIFIC SURGICAL COMPLICATIONS

GENERAL SURGERY			VASCULAR SURGERY		ENDOCRINE	
<u>Gastrectomy</u>	<u>SBO/LBO</u>	<u>Biliary / cholecystectomy</u>	<u>CABG/stent</u>	<u>Graft/stent/bypass</u>	<u>Thyroidectomy</u>	<u>Parotidectomy</u>
<ul style="list-style-type: none"> > Dumping syndrome > Malabsorption > Anastomotic ulcer > PUD / gastric cancer > Bacterial overgrowth > Gas bloating 	<ul style="list-style-type: none"> > Ileus > Anastomotic leak (usu 5-10 days post-op) > Stoma retraction > Intra-abdo collection / abscess > Adhesions → SBO/LBO > Pre-sacral plexus damage 	<ul style="list-style-type: none"> > CBD injury / stricture or bile leak > Anastomotic leak > Bleeding into biliary tree (jaundice) > pancreatitis 	<ul style="list-style-type: none"> > reperfusion arrhythmias > post-op ACS > ionotropes usage may cause peripheral organ hypoperfusion 	<ul style="list-style-type: none"> > graft vs host disease > haemorrhage / haematoma > infection > re-thrombosis > limb/organ ischaemia > AV fistula > Cholesterol embolism (trash foot) > Contrast (anaphylaxis, AKI)) 	<ul style="list-style-type: none"> > Airway obstructed 2^o to hemorrhage → URGENT opening of wound > hypoCa (removal of PTH glands) > recurrent laryngeal palsy = hoarseness 	<ul style="list-style-type: none"> > CN7 palsy
Orthopaedics			Urology		Other	
ANY ORTHO	THR	Cystoscopy / TURP	Endovascular		LN dissection	Neck dissection
<ul style="list-style-type: none"> > Prosthetic infection > Loss of position / failure to fix > Non-union/mal-union/delayed union > Neurovascular injury > Compartment syndrome 	<ul style="list-style-type: none"> > Sciatic nerve damage > Dislocation > LLD > Loosening + wear → repeat surgery > Pain + hip # = delirium 	<ul style="list-style-type: none"> > UTI > Absorbed glycine irrigation fluid = hypoNa > Impotence / retrograde ejaculation > Sphincter dysfunction > Urethral stricture 	<ul style="list-style-type: none"> > Retroperitoneal haemorrhage (e.g. SADPUCKER) > Any vascular surgery = bleeding and dementia 		<ul style="list-style-type: none"> > Lymphodema (e.g. axillary nodes) <p>NB: any plastics surgery = high level of immobilisation</p>	<ul style="list-style-type: none"> > CN 11/12 palsy

THE UNWELL POST-OP PATIENT

		Post-op	Cause	Ix	Rx	
Post-op Fever	Wind	< 24 HRS	atelectasis, URTI/ pneumonia • Pseudomonas (ventilator) 3-7 days • Chest infections causing suboptimal ventilation inducing basal atelectasis, UTIs	CXR, sputum culture	Abx (benzyl + doxy) OR (azithro + cef)	
	Water	3-5 days	UTI, Fluid depletion → check IDC, fluid balance	UA (M/C/S)	Abx (trimethoprim) + IVF	
	Walking	4-6 days	➢ DVT/PE ➢ Pressure ulcers ➢ Ileus (post-abdo surgery)		➢ Mobilise + calf compressor ➢ SC clexane → warfarin → DOAC) – check INR 2.5-3.5	
	Wound infection	5-7 days	wound site infection → lines, chest drain, surgical site or anastomotic leak		➢ Remove lines and Clean wound +/- incision or debride ➢ IV ABx	
	Wonder drugs	>7 days	transfusion/drug reactions (e.g. serotonin syndrome, thyroid storm)		➢ Stop drug ➢ DIC – meds, ➢ CT scan → drainage/abscess	
Post-op HypoTN	Shock type	Pulm. Artery pressure	CVP	SVR	CO Cause	
	Septic	Low	Low	Low	High Infection	
	Cardiogenic (pump failure)	High	High	High	Low • Surgical stress = ↑ACS risk (. 48 hrs post-op) • Epidural analgesia or high block (T5 and above) = lose SNS outflow = vasodilatation + cardio shock • Fluid overload + HF	
	Hypovolaemic	Low	Low	High	Low ➢ Long operations, XS sweating. ➢ haemorrhage, reduced PO intake	
	Obstructive	High /normal	High	High	Low PE, Pneumothorax, tamponade	
Low UO	Physiological Cause	Cause				
	Pre-renal (most common)	Hypovolaemia – haemorrhage? Reduced CO?				
	Renal	Nephrotoxic drugs (e.g. Contrast, aminoglycosides, salicylates, metformin)				
	Post-renal	BPH or ++ intra-abdo pressure causing ureter compression				
Pain	Physiology			Clinical Signs		
	CVS	↑HR, ↑BP, ↑hypercoagulability		ACS		
	RESP	↓ Lung vol. ↓ cough (retain secretion)		VTE (DVT/PE)	➢ Atelectasis ➢ Pneumonia ➢ Hypoxemia	
	GIT	↓ bowel motility ↓ gastric emptying			➢ Constipation, anorexia ➢ Paralytic ileus (causes hypoTN)	
	General	Medication overuse headache for any drugs (regular overuse for > 3 mths)				
	Check for: 1) Poor perfusion signs a. Prolonged CRT b. Pallor c. Cold peripheries d. Tachycardia 2) Organ dysfunction a. pH< 7.1 b. lactate > 2 c. reduced UO (< 0.5mL/kg/hr) d. confusion					
	Normal > 0.5mL/kg/hr Check for: 1) Fluid status (input VS output) a. 3 rd space losses into bowel or tissues b. Med review c. Examine IDC + bladder scan					
	Consult pain specialists / anaesthetics ➢ WHO step ladder for analgesia					

