

## ICU + ANAESTHETICS

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

### 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
- 2<sup>ND</sup> 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 - RNSH	
Measurements		Mortality Probability Model (24-48-72h)	
<p>Within 24 hours of admission to the ICU</p> <p>(The point score is calculated from a patient's age and these 12 routine physiological measurements)</p>	<ol style="list-style-type: none"> <li>AaDO<sub>2</sub> or PaO<sub>2</sub> (depending on FiO<sub>2</sub>)</li> <li>Temperature (rectal)</li> <li>Mean arterial pressure</li> <li>pH arterial</li> <li>Heart rate</li> <li>Respiratory rate</li> <li>Sodium (serum)</li> <li>Potassium (serum)</li> <li>Creatinine</li> <li>Hematocrit</li> <li>White blood cell count</li> <li>Glasgow Coma Scale</li> </ol>	<p>Date <input type="text" value="24h"/></p> <p>Age <input type="text" value="50"/></p> <p>Admission type <input type="text" value="Medical"/></p> <p> <input type="checkbox"/> Metastatic Cancer  <input type="checkbox"/> Cirrhosis  <input checked="" type="checkbox"/> Diuresis &lt; 150 mL/8h  <input checked="" type="checkbox"/> Creatinine &gt; 2 mg/dL  <input type="checkbox"/> Coma (GCS 3-5)  <input type="checkbox"/> Intracranial Mass Effect  <input type="checkbox"/> Vasoactive Drug &gt;= 1h  <input type="checkbox"/> Mechanical Ventilation  <input checked="" type="checkbox"/> PaO<sub>2</sub> &lt; 60 mmHg  <input type="checkbox"/> Proven Infection  <input type="checkbox"/> PT &gt; Standard +3 sec </p> <p>Calculate</p>	<p>Mortality Rate <b>23.8 %</b></p>

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: 10.1097/00003246-198510000-00009. 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:

- Hypervolaemic
- XS fluid intake
  - XS salt intake
  - IVF

Euvolaemic

Hypovolaemic  
(Dehydrated)

- Vomit / diarrhoea
- Internal bleeding
- Burns
- DI

## Examination

1. **Aids** = finished meal? Cups of water?  
IDC/drain output?

### Dehydration signs:

1. **VITALS = hypoTN, Tachycardia, CRT >2s, tachypnoea**
  2. Dry MM – stick tongue out
  3. Sunken eyeballs
  4. Cold peripheries
  5. Low urine output
  6. Low weight
- Fluid overload signs:**
1. Elevated JVP
  2. Displaced apex beat (most sensitive for HF)
  3. Pulm. Oedema (bibasal crackles, low sats)
  4. Peripheral oedema, ascites
  5. Weight gain

## Bedside Investigations:

- Wt, BMI
- **BLADDER SCAN → ?IDC**
- Urine – ↑SG, urine osmolarity
- ECG – arrhythmias?

## Blood Investigations:

- FBC (↓Hb – anaemia)
- EUC / CMP
- Serum osmolarity
- LFT – check albumin
- BNP – HF

## Imaging Investigations

- ECHO
- CXR

## Mx (conservative):

	Overload (WET)	Dry
<b>Non-pharm</b>	1L fluid restrict Na restricted diet	Hydrate
<b>Pharm</b>	20mg IV furosemide	IVF resus – crystalloid (0.9% NS or Hartman's)

## Practice note:

**What is the best fluid?** – amount/type always depends on patient's weight

**What is 3<sup>rd</sup> spacing?** Low level of fluid in intravascular space but XS fluid in interstitial space → appear hypovolemic or fluid overloaded at same time

## When to refer?

- Cannot maintain adequate fluid balance
- Cannot maintain electrolytes

Colloid			Crystalloid				
Vol. needed	Vol for vol		3x vol needed				
MoA	<ul style="list-style-type: none"><li>Molecules too large to cross capillary walls – fluid remain in intravascular space</li><li>Long half-life (hrs-days)</li></ul>		<ul style="list-style-type: none"><li>Molecules small enough to cross capillary walls → less fluid in intravascular space</li><li>Short half life (30-60 mins)</li></ul>				
<ul style="list-style-type: none"><li><b>Greater increased effect on intravascular vol. (1.5:1)</b><ul style="list-style-type: none"><li>For sepsis (NOT for TBI)</li></ul></li></ul>							
<b>Example</b>	<b>Natural</b>	<b>Synthetic</b>	<b>Example</b>	<b>Hypotonic</b>	<b>Isotonic</b>	<b>Hypertonic</b>	
	<ul style="list-style-type: none"><li>Whole blood</li><li>FFP</li><li>pRBC</li><li>Albumin</li></ul>	<ul style="list-style-type: none"><li>Geleatins</li><li>Dextrans</li><li>Hydroxyethyl starch</li></ul>	<ul style="list-style-type: none"><li>5% dextrose and 0.18% NS</li><li>Hypotonic saline</li></ul>	<ul style="list-style-type: none"><li>Normal saline (0.9%)</li><li>Lactated ringer's soln (aka Hartmann's)</li><li>plasmolyte</li></ul>	<ul style="list-style-type: none"><li>3% NaCl</li><li>5% NaCl</li></ul>		
	<b>ECF</b>	<b>Increased (esp. intravascular vol.)</b>		<b>ECF</b>	<b>Increased</b>	<b>Increased</b>	<b>Increased</b>
	<b>ICF</b>	<b>NONE</b>		<b>ICF</b>	<b>Increased</b>	<b>none</b>	<b>Decreased</b>
	<b>Use</b>	<ul style="list-style-type: none"><li>Cirrhosis</li><li>Critically ill – ARDS, burns, sepsis</li><li>Bleed → blood</li></ul>	<ul style="list-style-type: none"><li>Fluid resus</li><li>Maintenance fluid</li><li>Hypovolaemic hypoNa</li><li>IV drug solvent</li></ul>	<b>Ind. Use</b>	<ul style="list-style-type: none"><li>Correct hyperNa</li><li>Maintenance fluid</li><li>IV drug solvent</li><li>Children</li><li>Hypoglycaemia or with insulin IV</li></ul>	<ul style="list-style-type: none"><li>Fluid resus</li><li>Maintenance fluid</li><li>Hypovolaemic hypoNa</li><li>IV drug solvent</li></ul>	hypoNa (severe) cerebral oedema
<b>A/E</b>	<ul style="list-style-type: none"><li>Fluid overload → cardiac failure</li><li>Allergic</li><li>Expensive + may not be vegan</li></ul>		<b>A/E</b>	<b>FLUID OVERLOAD – cerebral, peripheral, pulmonary oedema</b> <ul style="list-style-type: none"><li>hypoNa, hypoK</li><li>hyperglycaemia</li><li>cerebral oedema</li></ul> <hr/> <div>NO dextrose in:<ul style="list-style-type: none"><li>brain haemorrhage</li><li>re-feeding syndrome</li></ul></div> <ul style="list-style-type: none"><li><b>0.9% NS</b> = HYPERchloaemia acidosis</li><li><b>Ringer's/ Hartman</b> = ++ lactate in liver failure, hyper K</li><li><b>Plasmalyte → high HCO<sub>3</sub></b></li></ul>			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, hypoTN	PERI-OP & POST-OP NBM due to bowel obstruction	POST-OP + VOMITING + DIARRHOEA	
Method	<b>Rapid fluid bolus (within 10-30mins)</b> <ul style="list-style-type: none"><li>Adults NS or LR 500-1000mL IV bolus</li><li>Children: NS or LR 10-20mL/kg IV bolus</li></ul>	NBM patients but do not have volume depletion, hypotension, or ongoing losses <ul style="list-style-type: none"><li>Adults → 1-2mL/kg/hr</li><li>Children (&gt; 28 days) → 4:2:1 rule<ul style="list-style-type: none"><li>4mL/kg/hr (1<sup>st</sup> 10 kg)</li><li>2mL/kg/hr (2<sup>nd</sup> 10 kg)</li><li>1mL/kg/hr (remaining)</li></ul></li></ul> <i>*Maintenance fluid requirement per kg of wt higher in children due to increased SA</i>	<b>Replace lost body fluids and electrolytes</b>	
			<b>Hypovol. shock</b> (N + D, burns, sepsis, fistula)	<ul style="list-style-type: none"><li>Hartmann's BEST as less Cl<sup>-</sup> to minimise risk of hyperchloremic acid (check lactate)</li></ul>
				<b>Dehydration</b> (poor intake)
Consider	<ul style="list-style-type: none"><li>vasopressors (e.g. metaraminol) → maintain BP and reduce peripheral fluid loss</li><li>Inotropes (e.g. dobutamine, levosimnadin)</li><li>Blood products (FFP, packed RBC – group + X-match)</li></ul>	<ul style="list-style-type: none"><li>Patient's weight</li><li>Check EUC before prescribing</li><li>Give oral/NG-tube fluids whenever possible → minimizes fluid overload</li></ul>	<ul style="list-style-type: none"><li>Pre-existing fluid loss (STAT bolus – 500mL 0.9% saline/Hartmann's soln)</li><li>Measure Ongoing losses (replace future losses – measure vomits, diarrhoea vs intake)</li><li>DO NOT Give K at a rate &gt; 10mM/hr or use maintenance protocols</li></ul>	

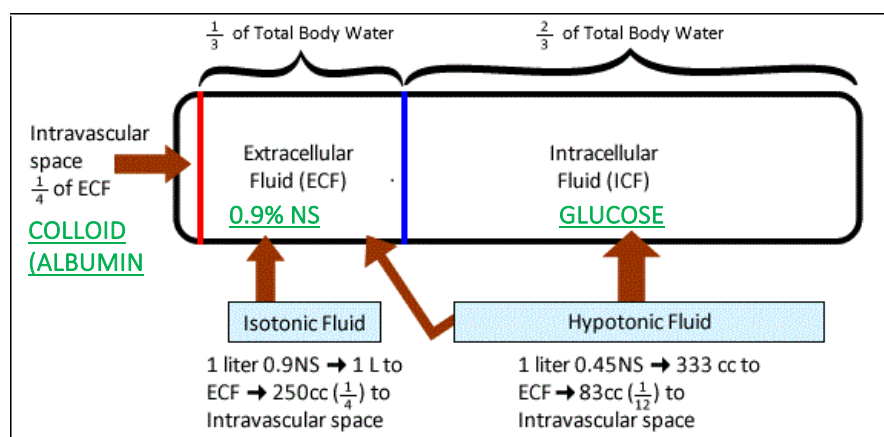
## Special cases:

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	Avoid fluid overload → monitor K → ?ECG
HF	Nb: <b>may normally be hypotensive</b> → low Na diet, furosemide, fluid restriction and record daily weights	Alcoholic	<b>Pabrinex BEFORE dextrose</b> (avoid Korsakoff) ➤ Replaced B1 (100mg thiamine)
Liver failure	<b>5% dextrose</b> – excess Na causes ascites	Brain haemorrhage	<b>Saline</b> (dextrose destroys brain)

# Crystalloid Fluids

Fluid	Na <sup>+</sup> mEq/L	Cl <sup>-</sup> mEq/L	K <sup>+</sup> mEq/L	Ca <sup>2+</sup> mEq/L	Glucose g/L	Buffer	Osmolarity mOsm/L	Tonicity	Typical Indication
Normal plasma	~ 140	~ 100	~ 4	~ 2.4	~ 0.85	HCO <sub>3</sub> <sup>-</sup> ~ 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

	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

## 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

## 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

**Anterior** (Left): 4.5%, 18%, 15%, 9%, 9%

**Posterior** (Right): 4.5%, 18%, 15%, 9%, 9%

4mL x Body Weight (kg) x TBSA (%) = Total Crystalloid Fluid in First 24 Hours

Body Part	Body Surface Area
Entire Head & Neck	9%
Entire Right Arm	9%
Entire Left Arm	9%
Entire Trunk	36%
Genit	1%
Entire Right Leg	18%
Entire Left Leg	18%

## Daily Requirements: (for 70kg man)

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

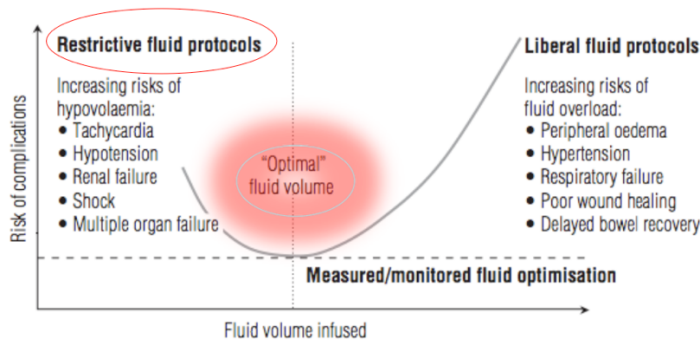
## Input vs Output:

- Daily weighing
- UO  $\approx$  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

# 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 ( $\text{Na} \times 2 + \text{urea} + \text{glucose}$ )

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

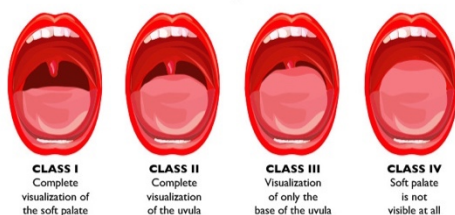


# Pre/Intra-operative Assessment

## INITIAL AIRWAY ASSESSMENT

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

The Mallampati Score



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 lx
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

and response to anaesthesia

- 1) Post-Op comp (sepsis, pain, anaesthetics)
- 2) Cardiac Hx
- 3) Resp Hx

	Concern	Pre-Op Investigations	Solution
Resp (airway + breathing)	<ul style="list-style-type: none"> <li>Beards, jaw protrusion</li> <li>Acute (E.g. PE, pneumothorax, pneumonia, infection)</li> <li>Chronic (OSA, ILD, COPD)</li> </ul>	<ul style="list-style-type: none"> <li>Auscultate + O2 sats</li> <li>CXR → check for HF, PPM</li> <li>Spirometry + ABG</li> <li>OSA checklist</li> </ul>	<ul style="list-style-type: none"> <li>Cease smoking &gt; 4 weeks</li> <li>Post-op monitor → beware breathless pt</li> <li>Immunosuppressed (steroids – COPD)</li> </ul>
CVS (IHD)	<p>AHA revised cardiac risk index</p> <ol style="list-style-type: none"> <li>4) High risk surgery</li> <li>5) Hx of <b>IHD</b> (ACS-MI, UA, decomp. HF)</li> <li>6) Hx of <b>HF + surgical</b> Hx (stents, CABG)</li> <li>7) Hx of <b>CVD</b></li> <li>8) <b>IDDM</b></li> <li>9) Renal impairment (eGFR – CKD → fluid caution)</li> </ol>	<ol style="list-style-type: none"> <li>1. <b>Bloods + products</b> → Group + save (pRBC, FFP, cryoppt, plts) <ul style="list-style-type: none"> <li>a. FBC, EUC, – <b>anaemia</b> (↑↑↑ post-op complication)</li> <li>b. COAG, B-HCG</li> </ul> </li> <li>2. <b>CXR, CTA/CTPA or V/Q scan</b></li> <li>3. <b>ECG (pre &amp; post)</b> – ? AF, ischaemic damage (inverse T waves, path Q waves, BBB) <ul style="list-style-type: none"> <li>a. <b>PERFORMED IN ALL &gt; 65 TO pre-operatively</b></li> </ul> </li> <li>4. <b>TOE/TTE, stress tests</b> → ONLY if Hx of valvular disease (MR) or heart failure (LVEF)</li> <li>5. <b>Cardiopulmonary exercise testing</b> to check for reserve – CPET, stress ECHO</li> </ol>	
CVS (HTN)	++ peri-op morbidity	24hr holter-monitor	<ul style="list-style-type: none"> <li>Target MAP &gt; 65,</li> <li>SBP &gt; 80 (higher if older w/ PVD)</li> </ul>
Diabetes	<ul style="list-style-type: none"> <li>++ infection risk</li> <li>Micro + macrovascular comp.</li> </ul>	<ul style="list-style-type: none"> <li>BSL, HbA1C,</li> <li>OGTT (new T2DM)</li> </ul>	<ul style="list-style-type: none"> <li>Optimise BSL control (target 5-10mM)</li> <li>Avoid fasting + BSL fluctuations</li> <li>Make sure 1<sup>st</sup> on list</li> </ul>
Older age + Obesity	High risk of desaturation rapidly	STOP BANG score for OSA cause – snore, tried, observed, BP, BMI, age > 50, neck cir. > 40cm, gender – male	<ul style="list-style-type: none"> <li>Minimise post-op comp</li> <li>Lose weight</li> </ul>

## Capacity and function

- 1) Previous surgeries and response to anaesthesia
- 2) Memory to take meds
- 3) SOCIAL support
- 4) Logistics (getting to hospital)

PMHx	<p>Response to anaesthetics</p> <ul style="list-style-type: none"> <li>Previous response</li> <li>Allergies to → propofol (soy, egg), Opioids,</li> <li>Check FHx – malignant hyperthermia</li> </ul> <p>Medication allergies</p> <ul style="list-style-type: none"> <li>Antibiotics: penicillin, vancomycin,</li> <li>Heparin/Protamine: HIT</li> <li>Iodine + chlorhexidine</li> </ul>	<p>Previous surgeries</p> <ul style="list-style-type: none"> <li>Difficult airways</li> <li>Stents,</li> <li>CABG</li> <li>Jt replacements</li> <li>Abdo surgery</li> </ul>
Exercise tolerance	<p>MET – Metabolic equivalent – what can they do?</p> <ul style="list-style-type: none"> <li>1 MET – eating, sitting, bath, shower and dress</li> <li>4 METs – walk up flight of stairs</li> <li>10 METs – strenuous exercise</li> </ul>	<p>Improve METs – condition/train</p> <ul style="list-style-type: none"> <li>Walk unaided?</li> <li>6-metre walk</li> </ul>
Function / mood	<ul style="list-style-type: none"> <li>Frailty score: Rockwood or Fried scale</li> <li>NSQIP risk index (surgical risk calc → age, organ fn, chronic illness, poor nutrition, cognitive dyfn, male)</li> <li>Nottingham hip # score → quicker to theatre = ↓ mortality</li> <li>Depression = longer hospital stay</li> </ul>	<p>Functional aids</p> <ul style="list-style-type: none"> <li>Sensory/hearing aids (glasses, hearing aid)</li> <li>Dentures</li> <li>Walking stick, wheelchair</li> </ul>
Alcohol	Organ damage → hepatic encephalopathy, pancreatitis	Avoid withdrawal + delirium
Smoking	<p>+++ risk of IHD, PVD, HTN →</p> <ul style="list-style-type: none"> <li>Atherosclerosis, ↓ tissue perfusion</li> <li>↓ wound healing</li> <li>Carcinogenic</li> <li>Lung disease (Bronchiectasis, COPD) → ↓ ciliary motion + sputum clearance</li> </ul>	<p>Cease smoking &gt; 4 weeks</p> <ol style="list-style-type: none"> <li>1) Improve tissue perfusion (days)</li> <li>2) ↑ wound healing</li> <li>3) ↑ pulm. Function</li> <li>4) ↑ immune system</li> </ol>

## Meds

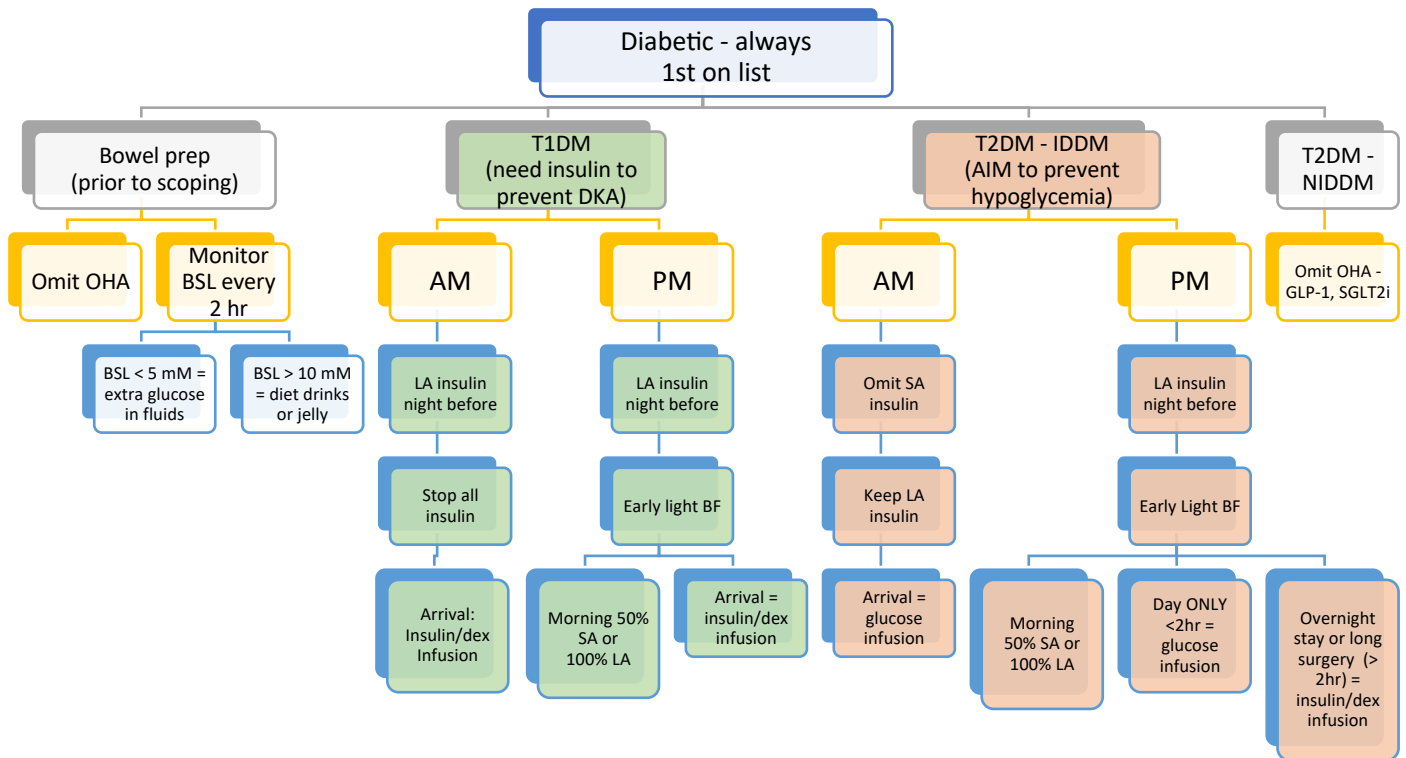
### Allowable meds:

- Anti-depress (watch for uncontrol HTN)
- Analgesics
- PD drugs (levodopa)
- Anti-anxiety (watch for ↑ SNS activity or O2 demand)
- BB, CaB
- Insulin

	Drug	Stopped?	Reason?
Antiplatelets (Stents)	Aspirin (COX 1/2 inhibitor)	5-7 days (only if high risk)	
	Clopidogrel -Plavix (P2Y12 inhibitor)	5-7 days	
	Ticagrelor (P2Y12 inhibitor)	5-7 days	
Anti-coagulants (AF, tissue valve, DVT)	Warfarin → Beware INR > 3.5 + warfarin = spinal haematoma	≈ 5 days [Vit K, prothrombinex] → restart 12-24 hours after surgery	
	Dabigatran (anti-thrombin - IIa)	1 day [Idarucizumab]	
	Clexane (LMWH)	3 days before + post-op LMWH for 3 days	
	DOACS = -xabans (Xa inhibitor)	≥ 3 days [Praxibind] + post-op LMWH for 3 days -- may not need if low risk surgery (e.g. cataracts)	
Anti-hypertensives	ACEi/ARB	1 day	HypoTN
	Diuretics	1 day	HypoK + ↓ BP
Oral Hypoglycaemics	Metformin	1 day	Lactic acidosis + AKI
	GLP-1	1 day (only if bowel prep)	GI disturbance
	SGLT2i	3-4 days (1 day if day-only procedure with no bowel prep)	Euglycaemic Ketoacidosis (monitor ketones pre and post-op)
Natural therapies	Ginger, ginkgo, garlic, ginseng + fish oil	Few days	++ bleeding risk
NSAID	Ibuprofen	Days before	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

# 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:

Skin	Ulcer	Sinus	Fistula	Abscess / cyst	Cyst and abscess drainage Hypergranulation (proud flesh)	Keloid scar
Incision	Join / fistula	Overhealing stenosis	False aneurysm Perforation / leak Haemorrhage		Types of healing: 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"> <li>Prevent / stop bleed → plt + cytokines → vasoconstrict</li> <li>1) Primary haem = during surgery</li> <li>2) Secondary haem = post-surgery</li> </ul>	1-3 hours/day	<ul style="list-style-type: none"> <li>Within "2 weeks" of surgery, anything can be repaired</li> </ul>
Inflammation	<ul style="list-style-type: none"> <li>Activated immune system to remove pathogen and necrotic tissue</li> </ul>	3-20 days	<ul style="list-style-type: none"> <li>Once &gt; 2 weeks – proliferation phase begins (highly fragile state) and anatomy changes</li> </ul>
Proliferation	<ul style="list-style-type: none"> <li>Healthy tissue regrows (ECM + collagen = granulation)</li> <li>Angiogenesis</li> </ul>	1-6 weeks	
Remodelling	<ul style="list-style-type: none"> <li>Granulation tissue matures into scar tissue (improved tensile strength)</li> </ul>	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
  - "discovered when Langer punctured numerous circular holes and noticed there were actually ellipsoid in shape"
- Enhanced recovery after surgery (ERAS) Program
  - Early mobilisation to reduce muscle breakdown, and improve muscle function
  - Early oral feeding POST-OP – avoid long periods of fasting
  - Minimise time on paralytic or ventilation

## How can we categorise the types of surgeries

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

Wound Classification	Definition	Examples	Surgical Site Infection Risk
Clean	Sterile procedure with no entrance into the GI tract, GU tract, or respiratory tract	Hernia repair, mastectomy, thyroidectomy, AAA repair	1%-3%
Clean-contaminated	Procedure with only minor breaks in sterility with controlled entry into the GI tract, GU tract, or respiratory tract with no significant contamination	Cholecystectomy, appendectomy, small bowel resection, colon resection	5%-8%
Contaminated	Procedure with poor sterility secondary to gross spillage from GI tract, GU tract, or respiratory tract or presence of foreign debris	Cholecystectomy with bile spillage, appendectomy for perforated appendicitis, small bowel or colon resection in setting of perforation	20%-25%
Dirty/Infected	Procedure involving contamination by established infectious processes	Abscess drainage, debridement of necrotizing soft tissue infections	30%-40%



# COMMON ANAESTHESIA PROCEDURES

## ECG - Check for arrhythmias

- **Rhythm** – AF (most common), SVT
- **Tachycardia** – pain, drugs, inadequate anaesthesia

## Pulse oximeter - O2 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 hypoTN causes

- Hypovol?, - fasting, blood loss
- XS anaesthesia,
- Rx: fluid bolus, vasoconstrictors



## Oxygen analyser - FROM GAS SAMPLER LINE

- Detects failure to supply O2



## 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**



## Cerebral Oximetry

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

## AIRWAY MANAGEMENT:

Procedure	Indications
<b>Endotracheal tube (ETT)</b> <ul style="list-style-type: none"> <li>flexible plastic tube with an inflatable cuff (balloon) at one end and a connector at the other</li> <li>different sizes (diameter in mm)</li> <li>syringe inflate the cuff via the <b>pilot line</b></li> <li><b>Murphy's eye</b> 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)</li> </ul>	Assistive devices: <ul style="list-style-type: none"> <li><b>McGrath laryngoscope</b> (camera +screen attached – live feed to visualise vocal cords)</li> <li><b>bougie</b> 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</li> <li><b>stylet</b> - stiff metal wire (with a plastic coating) to bend tip of ETT <b>anteriorly</b> towards the <b>trachea</b> (to avoid going posteriorly into oesophagus)</li> </ul>
<b>Awake fibre-optic intubation</b> <ul style="list-style-type: none"> <li>ETT inserted while patient conscious under guidance of endoscope</li> </ul>	<ul style="list-style-type: none"> <li><b>restricted mouth opening or difficult anatomy</b> (e.g., after RT to the neck (since Putting patient to sleep prior to inserting ETT has +++ risk of hypoxia)</li> <li><b>Trismus</b> – makes intubation more difficult</li> </ul>
<b>Supraglottic Airway Devices</b> <ul style="list-style-type: none"> <li>tip of the SAD located at the top of oesophagus</li> <li>SADs with inflatable cuffs are called <b>laryngeal mask airways (LMA)</b>.</li> <li><b>I-gel</b> is a type of non-inflatable SAD that uses a gel-like cuff that moulds to the larynx</li> </ul>	<ul style="list-style-type: none"> <li>alternative to endotracheal intubation for ventilation</li> <li>first option if intubation fails in a difficult airway scenario</li> </ul>
<b>Nasopharyngeal airways</b> <ul style="list-style-type: none"> <li>measured from the edge of the <b>nostril</b> to the <b>tragus</b> of the ear.</li> </ul>	emergency scenarios, for example, in A&E or at cardiac arrests Cl: <b>base of skull fracture</b>
<b>Oropharyngeal (Guedel)</b> <ul style="list-style-type: none"> <li>centre of the mouth to the angle of the jaw</li> </ul>	ventilating the patient via a face mask and bag prior to inserting an SAD or ETT
<b>Tracheostomy</b> <ul style="list-style-type: none"> <li>Performed under GA or in an emergency</li> <li><b>hole is made in the front of the neck with direct access to the trachea.</b></li> <li>Tube attached via <b>"trach-tie"</b></li> <li>Tube may be temporary or permanent                             <ul style="list-style-type: none"> <li>often inserted at the end of head and neck operations, for example, after a <b>laryngectomy</b> procedure (where a permanent tracheostomy will be required)</li> </ul> </li> </ul>	<ul style="list-style-type: none"> <li><b>Respiratory failure where long-term ventilation may be required</b> (e.g., after an acquired brain injury)</li> <li><b>Prolonged weaning from mechanical ventilation</b> (e.g., ICU patients that are weak after critical illness)</li> <li><b>Upper airway obstruction</b> (e.g., by a tumour or head and neck surgery)</li> <li><b>Management of respiratory secretions</b> (e.g., in patients with paralysis)</li> <li><b>Reducing the risk of aspiration</b> (e.g., in patients with an unsafe swallow or absent cough reflex)</li> </ul>

## 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**

## ANZCA PS18 – Equipment MUSTS

1. O2 analyser
2. Pulse oximeter
3. Ventilator alarm
4. Co2 monitor
5. Inhalational anaesthetic agent monitor

## CIRCULATORY MANAGEMENT:

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

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

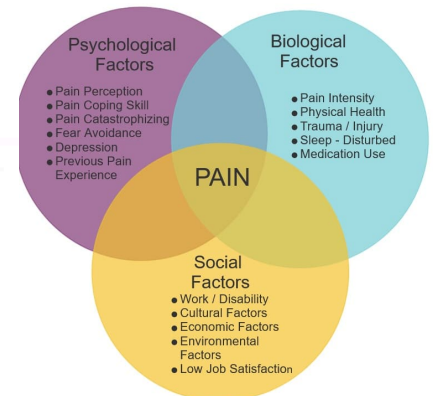
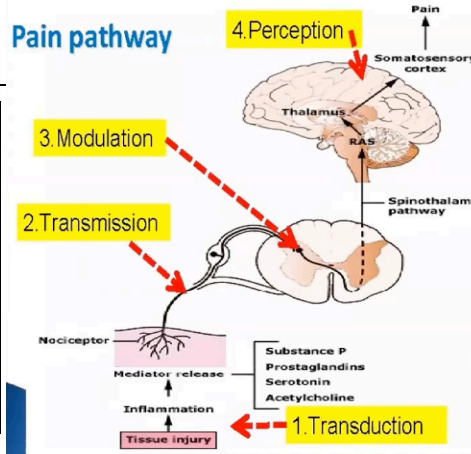


# POST-OP PAIN MANAGEMENT

## 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

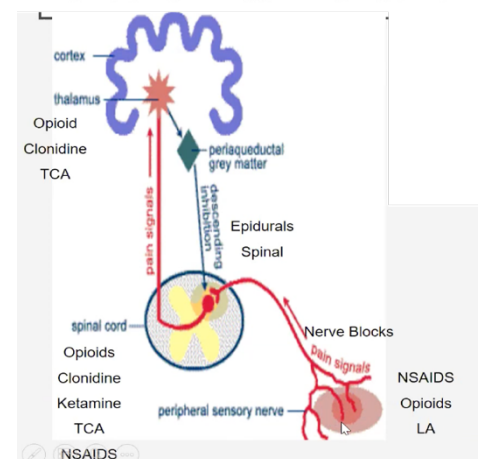
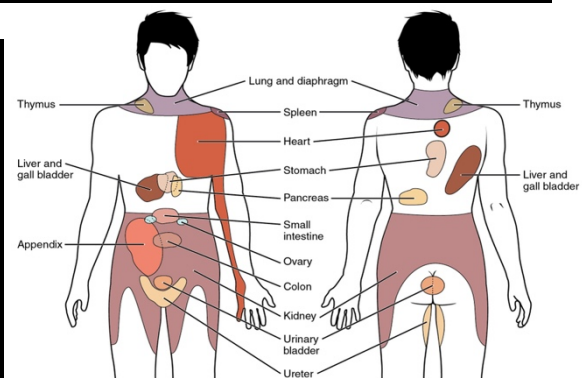
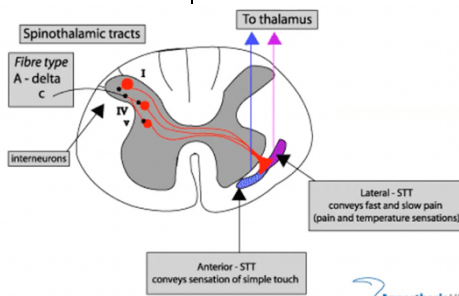
## Pain pathway



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

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

	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"> <li>Dermatomal (periphery)</li> <li>Non-dermatomal (central)</li> </ul>
Onset	Fast	Slow	Fast
E.g.	<ul style="list-style-type: none"> <li>Periosteum, joints</li> <li>Sickle cell</li> <li>Superficial laceration, burns, trauma</li> <li>Otitis media</li> <li>Stomatitis</li> </ul>	<ul style="list-style-type: none"> <li>Colic spasm pain (nausea + sweating)</li> <li>Appendicitis (anorexia)</li> <li>Renal colic (extreme)</li> <li>Chronic pancreatitis</li> <li>IBS</li> <li>Angina</li> <li>Periods</li> <li>Burning (sciatica, nerve compression)</li> </ul>	<ul style="list-style-type: none"> <li><b>Crushing / compressing</b> (Trigeminal, avulsion, post-traumatic neuralgia)</li> <li><b>Toxins</b>: Peripheral neuropathy (HIV, diabetes, alcohol, chemo)</li> <li><b>Limb amputation</b> - phantom limb pain - cortical remodelling</li> <li><b>Infection</b>: Herpetic Neuralgia</li> </ul>
Rx	<ul style="list-style-type: none"> <li>Panadol</li> <li>NSAID</li> <li>Opioids</li> </ul>	<ul style="list-style-type: none"> <li>Panadol</li> <li>NSAID</li> <li>Opioids</li> </ul>	<p>Try each one by itself:</p> <ul style="list-style-type: none"> <li><b>Amitriptyline</b> (TCA)</li> <li><b>Duloxetine</b> - an SNRI</li> <li><b>Gabapentin</b> (anticonvulsant)</li> <li><b>Pregabalin</b> (anticonvulsant)</li> </ul> <p><b>Alternative:</b></p> <ul style="list-style-type: none"> <li>Physiotherapy to maintain strength</li> <li>Tramadol (rescue therapy)</li> <li>Capsaicin - localised pain</li> </ul>



## COMMON PAIN MEDS (WHO LADDER)

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

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

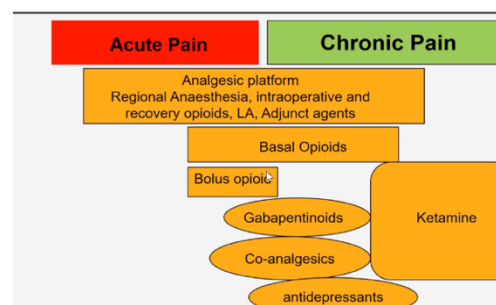
- **CONTACT APS and PALL Care if unsure about baseline and breakthrough doses (ALSO time limit endone when giving script)**
- **Give Mg as co-analgesia to NSAID and PCT to reduce opioid usage + Laxatives**
- **Avoid NSAID, paracetamol, opiates and anti-convulsant for chronic PRIMARY pain → use TCA and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

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

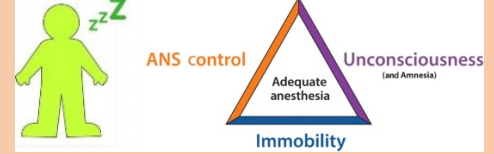
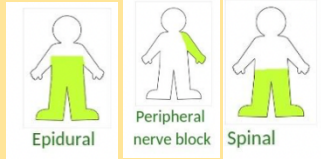
### ANALGESIA PROTOCOL (PERIOPERATIVE)



**AIM = MULTIMODAL ANALGESIA**  
to maintain steady state for good analgesia (early weaning)

# 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	➤ NO fasting needed		➤ NO fasting needed		<u><b>Fasting prior to operation:</b></u> ➤ 6 hours of no food or feed before operation ➤ 2 hours of no clear fluids (i.e. fully "nil by mouth") <u><b>Why do you need empty stomach (NBM)?</b></u> ➤ Risk of aspiration pneumonia + pneumonitis ➤ Inc. ED patients, non-fasting, full stomach	
	➤ Check allergies		➤ Check allergies			
		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"> <li>• <b>Topical</b></li> <li>• <b>Infiltration</b></li> </ul> <p><b>Indications:</b></p> <ul style="list-style-type: none"> <li>➤ Skin <b>sutures</b></li> <li>➤ Skin <b>biopsies</b></li> <li>➤ <b>Dental</b> procedures</li> <li>➤ <b>Minor surgery</b> (carpal tunnel)</li> <li>➤ Insertion <b>central line</b></li> <li>➤ <b>Percutaneous procedures</b> (e.g. PCI)</li> <li>➤ <b>LP</b></li> </ul>	<ol style="list-style-type: none"> <li><b>Peripheral nerve block</b> (under USS guidance) <ol style="list-style-type: none"> <li><b>Brachial plexus</b></li> <li><b>Femoral nerve</b> (neck of femur #, TKR)</li> </ol> </li> <li><b>Neuraxial block</b> <ol style="list-style-type: none"> <li><b>Epidural</b> → C/S, severe lung disease, lower limb surgery, child delivery</li> <li><b>Spinal</b> – no catheter → lower limb and pelvic surgeries, LSCS</li> <li><b>A/E = haematoma, infection, hypoTN, local anaesthetic toxicity</b></li> </ol> </li> <li><b>IV regional/peripheral</b> → <b>Bier's block</b> (19<sup>th</sup> century – Bohr war → IV local anaesthetic given with tourniquet to prevent systemic spread → only distal areas anaesthetized (Best for forearm injuries)</li> </ol> <p><b>Monitored anaesthetics care (MACS):</b></p> <ul style="list-style-type: none"> <li>➤ Conscious BUT calm/sedated state (e.g. benzos)</li> </ul>	<ol style="list-style-type: none"> <li><b>Pre-medication = to make intubation easier:</b> <ol style="list-style-type: none"> <li><b>Benzodiazepines</b> (e.g., midazolam) = relax muscles and reduce anxiety (also causes amnesia)</li> <li><b>Opiates</b> (e.g., fentanyl or alfentanil) = reduce pain and hypertensive response to the laryngoscope</li> <li><b>Alpha-2-adrenergic agonists</b> (e.g., clonidine), = for sedation and pain</li> </ol> </li> <li><b>Total IV or parenteral anaesthesia (TIVA) – analgo-sedation</b> <ol style="list-style-type: none"> <li><b>Propofol</b> → <b>allosteric agonist of GABA<sub>A</sub> receptor</b></li> <li><b>Etomidate</b> → if haem unstable (less impact on CVS)</li> <li><b>Barbiturates</b> (thiopental) → <b>brain surgery</b></li> <li><b>Benzodiazepines</b> (midazolam)</li> <li><b>Opioids</b> (fentanyl)</li> </ol> </li> <li><b>Total inhalation anaesthesia – ether analgesia</b> <ol style="list-style-type: none"> <li><b>Gas (NO given w/ Oxygen)</b> - Euphoric effect, analgesia <ol style="list-style-type: none"> <li><b>Indications</b> = pregnancy, child, casualty !</li> </ol> </li> <li><b>Volatile anaesthesia</b> (sevoflurane, isoflurane)</li> </ol> </li> </ol>
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A/E	<p><b>A/E of Local and Regional Anaesthetics:</b></p> <p><b>Systemic toxicity</b> (anaesthetic leak into blood NOT nerve)</p> <ul style="list-style-type: none"> <li>➤ <b>Prodrome</b> = metallic taste, facial numbness</li> <li>➤ <b>CNS (Crosses BBB)</b> → confused, seizures, resp. depression → resp. acidosis + hypoventilation</li> <li>➤ <b>CVS toxicity</b> → (1) initially HTN, tachycardia → (2) gradual slow rate of fast conducting pathways (purkinje) → (3) hypoTN, bradycardia, unopposed ventricular contraction → VF/VT → cardiac arrest</li> </ul> <table border="1"> <thead> <tr> <th>Block</th><th>Effect</th><th>Complication</th></tr> </thead> <tbody> <tr> <td rowspan="3">Spinal/epidural</td><td>Motor</td><td>Limb damage/weakness</td></tr> <tr> <td>SNS blockade</td><td>HypoTN, bradycardia, incontinence</td></tr> <tr> <td>Infection</td><td>Infection (meningitis/enceph)</td></tr> <tr> <td>High spinal/ thoacic /epidural</td><td>motor</td><td>Resp. failure Infection (meningitis/enceph)</td></tr> <tr> <td>All CNS/PNS blocks</td><td>Sensory</td><td>Thermal/pressure injury Neuropathic pain</td></tr> <tr> <td>Upper /lower limb block</td><td>Proprioception</td><td>Impaired balance</td></tr> <tr> <td>Brachial plexus</td><td>Phrenic nerve</td><td>Resp. failure</td></tr> <tr> <td>Brachial plexus</td><td>RLN</td><td>Hoarsness, aspiration</td></tr> <tr> <td>Brachial plexus</td><td>PSNS block</td><td>Horner's syndrome (enophthalmos, miosis, ptosis, anhydrosis)</td></tr> <tr> <td>Femoral nerve (within iliopsoas m.)</td><td></td><td>Inability to flex hip</td></tr> </tbody> </table>	Block	Effect	Complication	Spinal/epidural	Motor	Limb damage/weakness	SNS blockade	HypoTN, bradycardia, incontinence	Infection	Infection (meningitis/enceph)	High spinal/ thoacic /epidural	motor	Resp. failure Infection (meningitis/enceph)	All CNS/PNS blocks	Sensory	Thermal/pressure injury Neuropathic pain	Upper /lower limb block	Proprioception	Impaired balance	Brachial plexus	Phrenic nerve	Resp. failure	Brachial plexus	RLN	Hoarsness, aspiration	Brachial plexus	PSNS block	Horner's syndrome (enophthalmos, miosis, ptosis, anhydrosis)	Femoral nerve (within iliopsoas m.)		Inability to flex hip	<p><b>Commonly:</b></p> <ul style="list-style-type: none"> <li>➤ IV meds are induction agents (i.e. induce unconsciousness),</li> <li>➤ <b>inhaled</b> medications for <b>maintaining</b> GA during operation (since they take time to reach effective concentration since they need to diffuse across the lung tissue and into the blood)</li> <li>➤ <b>Total intravenous anaesthesia (TIVA)</b> uses IV meds for BOTH induction and maintenance of GA (Propofol) → nicer recovery (as they wake up) compared with inhaled options.</li> </ul> <p><b>Emergence:</b></p> <ul style="list-style-type: none"> <li>➤ Wean off muscle relaxant – to avoid "awareness under anaesthesia"</li> <li>➤ 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</li> <li>➤ Consider reversal agent e.g. <b>sugammadex</b></li> </ul> <p><b>A/E of GA:</b></p> <ul style="list-style-type: none"> <li>➤ Post-op sore throat, nausea and vomiting</li> <li>➤ Aspiration</li> <li>➤ Dental injury</li> <li>➤ Anaphylaxis</li> <li>➤ Cardiovascular events (e.g. MI, stroke, arrhythmia)</li> <li>➤ Death</li> <li>➤ Malignant hyperthermia (<b>rare BUT fatal</b>) <ul style="list-style-type: none"> <li>○ Assoc. w/ suxamethonium and volatile anaesthetics</li> <li>○ Hypermetabolic response to anaesthesia → febrile, tachycardia, acidosis, hyperK, rigidity, XS CO<sub>2</sub> production</li> <li>○ <b>Rx: dantrolene</b> (interrupts muscle rigidity by disrupting Ca ion movement in SKM)</li> </ul> </li> </ul>
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## Treatment for systemic toxicity

- 1) Stop injection
- 2) BOLUS interlepid therapy (stop protein binding)
- 3) Ventilate – FiO<sub>2</sub>
- 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

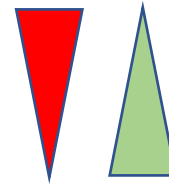
## Desired dermatome level of neuraxial block

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 1<sup>st</sup> 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

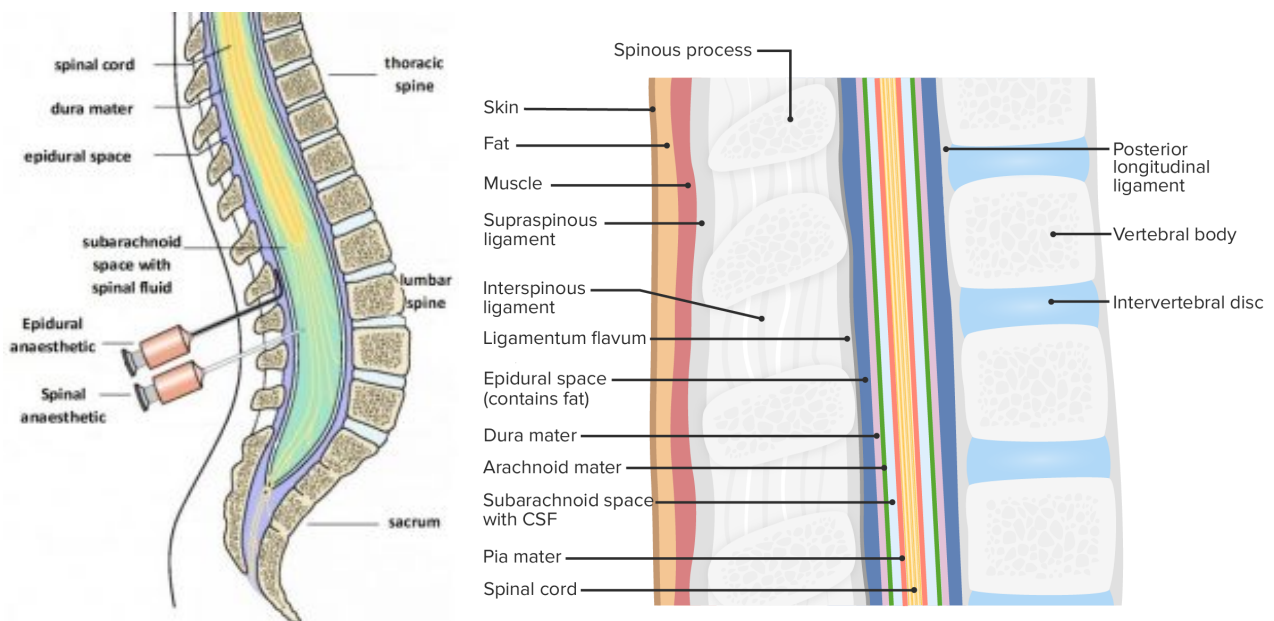
## MAIN CONTRAINDICATIONS

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





# COMMON ANAESTHESIA MEDICATIONS

	Type	MoA	Dosage	Indication	A/E
<b>Halothenes</b> 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)
<b>Nitrous Oxide</b> “beware recreational use”	Volatile sparing agent	Stops methionine synthetase to produce analgesia ➢ Minimise prolonged use	High MAC (102%)	Simple procedures	<b>Malignant Hyperthermia</b> <b>Megaloblastic anaemia (vit B12 def)</b> Agranulocytosis
<b>Oxygen</b>	Volatile agents				<b>ROP</b>
<b>Propofol</b> (white lipid soln)	IV induction or sleeping agent (emulsion / egg)	<b>Non-barbiturate</b> 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
<b>Thiopentane</b>	IV induction	<b>barbiturate</b> GABA agonist ➢ Reduces cerebral consumption	3-7mg/kg (5-10mins)	<b>Brain surgery, status epilepticus, raised ICP</b>	Tachycardia Vasodilated (↓SVR)
<b>Ketamine</b> “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
<b>Midazolam</b>	Sedative	GABA receptor agonist (benzo)	IV 0.02-0.1mg/kg	Sedation + anxiolysis	Headache, delirium, N/V, drowsy
<b>Adrenaline</b>	Vasopressor • Lower dose = vasodilate (B1, B2) • Higher dose = vasoconstrict (a1)	• <b>A1 vasoconstrictor</b> (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 110000	➢ HypoTN ➢ Anaphylaxis ➢ Severe asthma (bronchodilator)	Lactic acidosis Higher mortality in HF patients
<b>Ephedrine</b>	Vasopressor • Indirect SNS – release NORAD	• <b>Potent A1 (↑NORAD) AND B activity</b> • ↑HR, ↑CO, ↑RR, O2 cons. • bronchodilate	IV bolus <b>ONLY</b>	HypoTN	Higher mortality
<b>Phenylephrine</b>	Vasopressor • last 5 minutes	• <b>Selective A1 vasoconstrictor</b> • ↑SVR → ↑BP → ↑Coronary artery flow	IV bolus or infusion	HypoTN	reflex bradycardia (↑SVR)
<b>Metaraminol</b>	Vasopressor • Lasts 20-60mins • Longer acting	• <b>A1 vasoconstrictor (mainly)</b> • ↑SVR → ↑BP → ↑Coronary artery flow, + pulm. Vasc. pressure	IV bolus or infusion	Acute HypoTN	➢ reflex bradycardia (↑SVR) ➢ Higher mortality
<b>Vasopressin</b>	Vasopressor last 5 minutes	• <b>Selective V1 vasoconstrictor</b> • ↑SVR → ↑BP → ↑Coronary artery flow	IV via CVC		
<b>Levosimendan</b>	Positive Inotrope (increase contractility)	• PDE3 inhibitor	IV via CVC	mainly cardiac pts	
<b>Milrinone</b>	Positive Inotrope (increase contractility)	• Increases heart muscle sensitivity to calcium	IV via CVC		
<b>-curium or -curonium (e.g. rocuronium)</b>	Muscle relaxant / paralytic	Non-depolarising neuromuscular blocking agent → Competitive inhibitors against nAChR • Slower action → muscle relaxant	IV	• <b>Keep Ventilation</b> • <b>Prevent asp. pneumonia if NOT fasted</b>	Muscle pain and salivation <b>Reversal agent = sugammadex 16mg/kg</b>
<b>Suxamethonium</b>	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	<b>Hyper K, hypoTN, fever</b> <b>CI = malignant hyperthermia, glaucoma OR eye trauma</b> <b>Reversal agent = neostigmine</b>
<b>Neostigmine</b>	Reversal agent	Inhibits ACHE at NMJ → ↑ ACh	Given with anti-chol (atropine)	Reverse paralytic	<b>Cholinergic effects</b> (bronchospasm, bradycardia)
<b>Glycopyrrolate</b>	Anti-cholinergic / anti-muscarinic	↓ACh → ↑HR (mildly increase)	200mcg	Mild Bradycardia	Dry mouth, blurred vision, urinary retention, drowsy ➢ Non-opposed-tachycardia
<b>Atropine</b>	Anti-cholinergic	↓ACh (acts on SA node) → ↑HR (greater) Blocks CNX on heart	ED = 3mg	Bradycardia	
<b>Parcoxib</b>	<b>Analgesic - NSAID</b>	COX-2 selective			1. Constipation 2. Sedation/ drowsy 3. Nausea / vomit 4. Dependence
<b>Remifentanyl</b>	<b>Analgesic - opioid</b>	Ultra SA opioid • Fast onset + offset of action	IV	ICU or theatre for tube tolerance	
<b>Fentanyl</b>	<b>Analgesic – opioid</b>	SA + potent opioid • <b>Lipid soluble → diffuse</b>	5-15mcg	Analgesia → spinal blocks OR patch epidurals	
<b>Alfentanil</b>	<b>Analgesic – opioid</b>	V. short acting opioid • Less potent than fentanyl (10 min duration of action)		Analgesia for shorter cases or bridging	
<b>Morphine</b>	<b>Analgesic – opioid</b>	Gold standard analgesic • Hydrophilic	100-300mcg	analgesia	
<b>Procaines (cocaine, tetracaine, benzocaine)</b>	<b>Ester</b> -linkage LA	➢ Short-acting (broken down by plasma esterase) ➢ Vasodilatation		Short duration analgesia w/ slow onset of action	
<b>Bupivacaine</b>	<b>Amide</b> -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	
<b>Lidocaine</b>	<b>Amide</b> -linkage LA + anti-arrhythmic			RA onset of action for Short procedures: ➢ Lower pKa = more ionised = faster action	
<b>Penicillins</b>	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
<b>Cephalosporins</b>	Antibiotic	Cefuroxime = 2 <sup>nd</sup> gen Cephalosporin (only 2 <sup>nd</sup> gen that <b>crosses BBB</b> )	IV 2g	Broad spectrum (g +ve and g -ve)	ATN
<b>Linosamides</b>	Antibiotic	Clindamycin		2 <sup>nd</sup> line if penicillin or cephalosporin allergy	
<b>Aminoglycosides</b>	Antibiotic	Gentamicin		UTI? UROLOGY	Nephrotoxic, ototoxic
<b>Oxytocin</b>	Obstetric (syntocinon)	• ↑uterine contractions		induce labour or post-LCSC bleed (PPH)	
<b>Tranexamic acid</b>	Antifibrinolytic	• Stop clot breakdown		Trauma, surgery, ortho, menorrhagia	VTE
<b>Ondansetron</b>	Anti-emetic (PONV)	<b>5HT3 receptor in CTZ</b> → vomiting centre ➢ Do ECG prior (?prolonged QT)	4mg IV	N+ V	Fatigue, drowsy, constipation, <b>prolonged long QT</b>
<b>Metoclopramide</b>	Anti-emetic (PONV)	<b>D2 receptor antag</b> + prokinetic			Caution in <b>sedation</b> , constipation, <b>prolonged QT</b>
<b>Dexamethasone</b>	Anti-emetic (PONV)	40-50x more potent than hydrocortisone and longer lasting		N+ V, odynophagia, systemic illness	Steroid issues – caution in DM and immunocomp.
<b>Cyclizine</b>	Anti-emetic (PONV)	<b>H1 anatag</b>			Caution w/ HF and elderly

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

	Concern	Solutions
Pain	<b>CVS</b> – HTN, tachycardia, DVT <b>Resp</b> – atelectasis, impaired cough, pneumonia <b>GI</b> – reduced gastric motility <b>Neuroendocrine</b> – impaired wound healing, increased catabolic hormones (** BSL)	➤ <b>Choose appropriate analgesia</b> → dosage and duration
Temperature	<b>Anaesthesia causes loss of thermoregulation</b> ➤ Heat loss by conduction, radiation, convection	➤ <b>Pre-warming = best</b> (bair hugger, warm blankets) ➤ <b>Warmed fluids</b> (esp. blood products, fluids) ➤ <b>Theatre temperature control</b> (esp. in paediatrics!!)
Pressure area	<b>Long surgical time</b>	➤ <b>Patient positioning</b> – protect skin and nerves ➤ <b>Padding</b> – surgical table, heels ➤ <b>Protect vulnerable nerves</b> (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

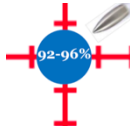

	Cause	Sx	Rx
<b>Neuroleptic malignant syndrome</b>	➤ <b>dopamine receptor-blocking drugs</b> → antipsychotic agents (e.g. chlorpromazine, haloperidol) and some antiemetics (metoclopramide). ➤ <b>OR sudden withdrawal from dopaminergic drug</b> (e.g. levodopa, bromocriptine)	➤ muscle rigidity, ➤ bradyreflexia, ➤ bradykinesia, ➤ altered mental status, ➤ EPSE ➤ hyperthermia.	<b>bromocriptine</b> (a dopamine agonist) PO or NGT ➤ Start at 2.5 mg 8-hourly, > ➤ increasing to 5 mg every 4 h (maximum 30 mg/day) in moderate to severe cases
<b>Malignant hyperthermia syndrome</b>	➤ autosomal dominant disorder after receiving an inhalational anaesthetic or suxamethonium	➤ muscular rigidity, ➤ tachypnoea, ➤ tachycardia, ➤ hypertension, ➤ mottled diaphoretic skin ➤ cardiac arrhythmias.	➤ <b>dantrolene 1 mg/kg IV for</b> severe muscle rigidity and hyperthermia. ➤ 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)

- Acute onset < 1 week
- Predisposing condition
- bilateral opacities consistent with APO must be present (detected via chest CT or CXR)
- PF ratio <300mmHg with a minimum of 5 cmH<sub>2</sub>o PEEP
- must not be fully explained by cardiac failure or fluid overload,

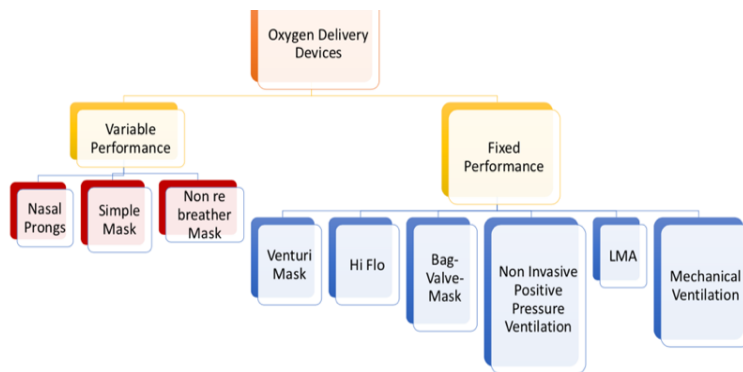
## RESPIRATORY SUPPORT

### Targeted Oxygen Therapy

In the <b>absence</b> of COPD		In the <b>presence</b> of COPD	
Where SpO <sub>2</sub> is <b>&lt;92%</b>		With a SpO <sub>2</sub> <b>&lt;88%</b>	
<ul style="list-style-type: none"> <li>titrate oxygen to maintain SpO<sub>2</sub> <b>92-96%</b></li> </ul>		<ul style="list-style-type: none"> <li>titrate oxygen to achieve a SpO<sub>2</sub> <b>88-92%</b> using a <b>Venturi mask delivering a FiO<sub>2</sub> 24%-28%</b></li> </ul>	

### What/When to use?

- Mainly for acute respiratory distress syndrome (ARDs)
- Respiratory support ONLY buys times until underlying problems can be managed
- Chest physiotherapy and suction = helps to clear secretions and improve respiratory function
- Remember O<sub>2</sub> 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:

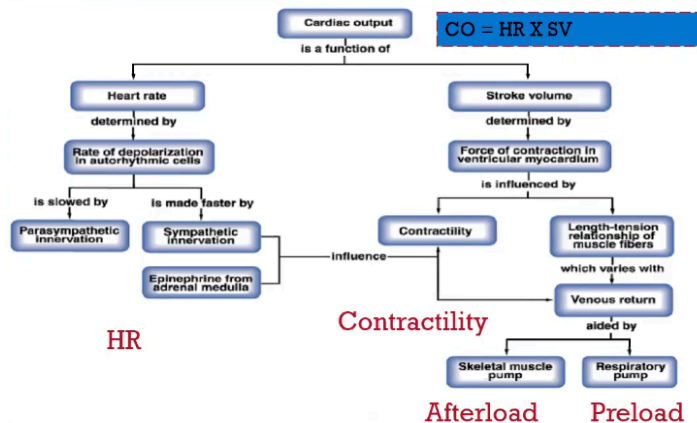
- Respiratory support
- Prone positioning (lying on front)
- Fluid and electrolyte management (avoid XS fluid in lung)

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

**Fraction of Inspired Oxygen (FiO<sub>2</sub>) concentration ≈ (Flow Rate x 4) + 21 (FiO<sub>2</sub> in room air)**

## CARDIOVASCULAR SUPPORT

$$BP = CO \times TPR$$



### DETERMINANTS OF CARDIAC OUTPUT

Low MAP = tissue **hypoperfusion**, leading to **hypoxia, anaerobic respiration, lactate** production and damage to the tissue.

	Decrease	increased
<b>HR</b>	PSNS - opiate Inotropes	SNS = Shock, stress BB
<b>Preload (CVP = RAP) = heart muscle tension or pressure filling ventricle (central venous pressure)</b>	Volume loss = bleeding IV fluid or MTP	↑ vol. to heart = ↑ RA = ↑ EDV (e.g. pregnancy) Diuresis
<b>Afterload (≈TPR or intrathoracic pressure) = force needed to pump against aorta (aortic pressure)</b>	MR Vasodilating (e.g. Anaphylaxis) Vasopressors	Viscous blood L-side (systemic HTN, AR, MR) R side (PHTN) Anti-coags.
<b>Contractility = affected by HR, preload, afterload</b>	Ischaemia Inotropes	SNS BB, CaB
<b>Venous return</b>	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<sub>2</sub> sats (via CVC)
- 3) Pulmonary artery O<sub>2</sub> 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 HypoTN:

- Increase contractility → positive inotropes (e.g. milirone, levisemindan, dobutamine, adrenaline, isoprenaline)
- Increase afterload → vasopressors (E.g. adrenaline, vasopressin)

#### General treatment for bradycardia:

- Reduce PSNS - e.g. atropine, glycopyrrolate

#### General treatment for tachycardia:

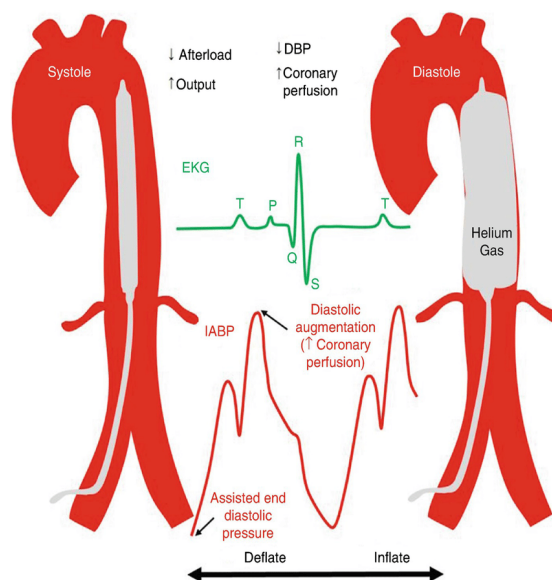
- BB, CaB, Flecainide

#### 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** – uremic encephalopathy

**AIM:** Removing excess fluid, solutes and waste products

#### Types of dialysis used in ICU:

- **Continuous renal replacement therapy (CRRT)** 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 or heparin
- Dialysate (the fluid where solutes filter out from blood) into



# POST-OP COMPLICATIONS

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

## SPECIFIC SURGICAL COMPLICATIONS

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

## THE UNWELL POST-OP PATIENT

Post-op Fever		Post-op	Cause			Ix	Rx	
	Wind	< 24 HRS	atelectasis, URTI/ pneumonia <ul style="list-style-type: none"><li>• <b>Pseudomonas (ventilator) 3-7 days</b></li><li>• Chest infections causing suboptimal ventilation inducing basal atelectasis, UTIs</li></ul>			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 ( <b>trimethorprim</b> ) + IVF	
	Walking	4-6 days	<ul style="list-style-type: none"><li>➢ DVT/PE</li><li>➢ Pressure ulcers</li><li>➢ Ileus (post-abdo surgery)</li></ul>				<ul style="list-style-type: none"><li>➢ <b>Mobilise + calf compressor</b></li><li>➢ SC clexane → warfarin → DOAC) – check INR 2.5-3.5</li></ul>	
	Wound infection	5-7 days	wound site infection → lines, chest drain, surgical site or anastomotic leak				<ul style="list-style-type: none"><li>➢ Remove lines and Clean wound +/- incision or debride</li><li>➢ IV ABx</li></ul>	
	Wonder drugs	>7 days	transfusion/drug reactions (e.g. serotonin syndrome, thyroid storm)				<ul style="list-style-type: none"><li>➢ Stop drug</li><li>➢ DIC – meds,</li><li>➢ CT scan → drainage/abscess</li></ul>	
Post-op HypoTN	Shock type		Pulm. Artery pressure	CVP	SVR	CO	Cause	<b>Check for:</b> 1) Poor perfusion signs <ul style="list-style-type: none"><li>a. Prolonged CRT</li><li>b. Pallor</li><li>c. Cold peripheries</li><li>d. Tachycardia</li></ul> 2) Organ dysfunction <ul style="list-style-type: none"><li>a. pH&lt; 7.1</li><li>b. lactate &gt; 2</li><li>c. reduced UO (&lt; 0.5mL/kg/hr)</li><li>d. confusion</li></ul>
	Septic	Low	Low	Low	High	Infection		
	Cardiogenic (pump failure))	High	High	High	Low	<ul style="list-style-type: none"><li>• <b>Surgical stress</b> = ↑ACS risk (. 48 hrs post-op)</li><li>• <b>Epidural analgesia or high block</b> (T5 and above) = lose SNS outflow = vasodilatation + cardio shock</li><li>• <b>Fluid overload + HF</b></li></ul>		
	Hypovolaemic	Low	Low	High	Low	<ul style="list-style-type: none"><li>➢ Long operations, XS sweating,</li><li>➢ haemorrhage, reduced PO intake</li></ul>		
	Obstructive	High /normal	High	High	Low	PE, Pneumothorax, tamponade		
Low UO	Physiological Cause		Cause					Normal > 0.5mL/kg/hr <b>Check for:</b> 1) Fluid status (input VS output) <ul style="list-style-type: none"><li>a. 3<sup>rd</sup> space losses into bowel or tissues</li><li>b. Med review</li><li>c. Examine IDC + bladder scan</li></ul>
	Pre-renal (most common)		Hypovolaemia – heamorrhage? Reduced CO?					
	Renal		Nephrotoxic drugs (e.g. Contrast, aminoglycosides, salicyclates, metformin)					
	Post-renal		BPH or ++ intra-abdo pressure causing ureter compression					
Pain	Physiology			Clinical Signs			Consult pain specialists / anaesthetics ➢ WHO step ladder for analgesia	
	CVS	↑HR, ↑BP, ↑hypercoagulability		ACS VTE (DVT/PE)				
	RESP	↓ Lung vol. ↓ cough (retain secretion)		<ul style="list-style-type: none"><li>➢ Atelectasis</li><li>➢ Pneumonia</li><li>➢ Hypoxemia</li></ul>				
	GIT	↓ bowel motility ↓ gastric emptying		<ul style="list-style-type: none"><li>➢ Constipation, anorexia</li><li>➢ Paralytic ileus (causes hypoTN)</li></ul>				
	General	Medication overuse headache for any drugs (regular overuse for > 3 mths)						

