This booklet aims to be guide only to the first couple of weeks in anaesthetics to give a little basis for further teaching/discussions and skill learning

- Orange call out boxes show topics that can be good to read up on in each section
- Feedback for anything you feel needs to be added/removed would be appreciated!



# Sections:

- 1. Starting your day
- 2. History taking and the anaesthetic chart
- 3. Basic Principles of anaesthesia
- 4. Recording intraoperative care
- 5. Recovery and post operative management
- 6. Anaesthetic Drugs and Reference Sheet
- 7. Machine Basics and the Circle System
- 8. Machine and Ventilation settings the basics

# Starting your day and Pre-Assessment

### Finding your list and seeing your patients

Lists for the day:

- Master copies found hanging on the window of the theatre office and you can then make copies
- Surgical Day Unit have a folder with all planned lists you cannot remove these but are a good reference
- Sometimes the theatre you are in will have an extra list you can take with you.
- CEPOD theatres have a whiteboard outside theatre 12 with the planned list for the day. Handover with night team happens there

### Seeing your patients:

- Elective patients are usually in Surgical Day Unit. This is divided into colour zones for different surgeons
- CEPOD patients will be inpatients and the wards usually written on the whiteboard. Worth double-checking on CHARTS as they sometimes get moved overnight.

# Preparing your anaesthetic room

- Introduce yourself to the ODP and let them know plans/equipment you'll need
- Machine and equipment checks
- Preparing drugs

#### **Team Brief**

WHO team brief in theatre with the whole theatre team. A chance to discuss any concerns and your plan for the anaesthetic.

#### First patient

- Safety checks
- Monitoring
- · Begin your anaesthetic!

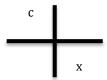
For the most part this is similar to most medical history taking you will be used to. Use a system-based approach and explore any co-morbidities in more detail. Specific anaesthetic questions to consider:

- Previous GA / any problems after
- FHx of problems with GA (Sux apneoa, Malignant hypothermia)
- GORD/aspiration risk
- Exercise Tolerance

Lots of formal ways to assess this (see callout for methods to review) Some things to include:

- Loose teeth /caps /crowns/dentures
- Neck extension
- Mouth opening

You will sometimes see documentation like this denoting the mouth and locations of crowns, broken teeth etc:



\*Tip! Use old anaesthetic charts to see intubation grade/any problems

Try and think about each stage for the patient:

- Type of anaesthetic (regional, sedation, GA?)
- Type of airway
- Any regional blocks?
- Invasive monitoring required? (Arterial line, CVC?)
- Plan for intraoperative and post op pain relief? PCA
- Antiemetic plan
- Fluid management, blood products required?

Ensure you use a chart with QR code as these allow the notes to be scanned. Some old charts on the ward do not have these

			Name Hospital Nu	mber			
Age	Hospita	al	Date of Birt	e of Birth			
Date of assessment	Time		Height	1	Weight	BMI	
Procedure planned			Asse	essing An	aesthetist		
History and Examinati	on		Pren	nedicatio	'n		
			Alle	rgies		7	
			Smo	king			
			Med	dication			
			Inve	stigation	S		
Airway		Blood Pressu	Jro.				
an way		Blood Fresso	are				
Anaesthetic plan							
1							
A							
1	ıt / consen	t					
Discussion with patien	nt / consen	t	NRI	M	LIRC	SENCY.	
1	ut / consen	t	NBI Cle- Soli	ar Fluids		GENCY eduled	

Include consent for any procedures.

Specific anaesthetic consent will vary with patients and procedures. High risk elective patients with multiple comorbidities are usually seen in clinic and this is discussed at length.

Common things to discuss with all a patients on the day are:

- Dental damage
- Sore throat
- PONV
- Anaphylaxis
- Need for blood products



### To Tube or not to tube?

Part of making plans for delivering an anaesthetic is deciding on which patients require intubation and which can be managed on an LMA. With the huge variety of surgical procedures and patient types, these are some factors that can help you decided what may be required:

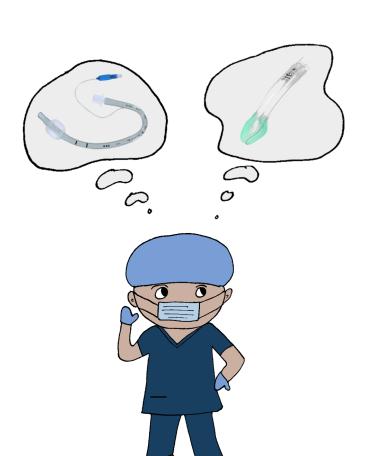
### **Patient Factors**

- History of reflux
- Pregnancy
- Recent trauma/illness (delayed gastric emptying)

### **Surgery Factors**

- Muscle relaxation required for surgery
- Surgery with increase intra-abdominal pressure (laparoscopy)
- Bowel manipulation
- Long surgical time
- Shared airway (blood/surgical debris)
- Position of patient (e.g prone)

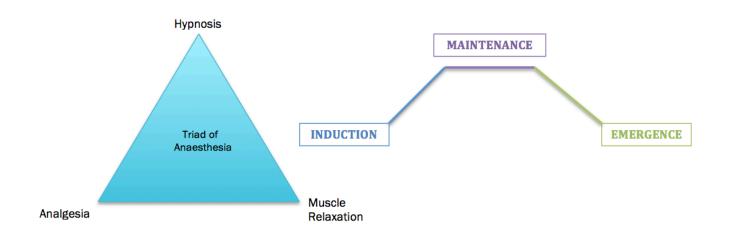




# Basic principles of Anaesthesia

There are different decisions that need to be made when conducting an anaesthetic, and as such there are many ways to think about how to divide these various components.

The classical triad of anaesthesia and the three key stages of anaesthesia are shown below:



These can be helpful in categorizing in your mind what you may need for each step and help you plan and prepare.

Type of Induction	Type of Airway	Maintenance Type	Type of Ventilation
Gas	Facemask	Gas	Spontaneous Ventilation (SV)
IV	LMA	IV	Intermittent Positive Pressure Ventilation (IPPV)
RSI	ETT		

### General sequence of events for ASA 1/2 patient

WHO safety checks AAGBI monitoring Pre-oxygenation

#### Induction:

- Small dose fentanyl, propofol and muscle relaxant
- Manual Ventilation + sevoflurane on
- Tracheal Intubation confirmed by equal chest rise, tube misting, capnography (at least 6 breaths!)

#### Maintenance:

- Oxygen/Air/Sevoflurane
- IPPV
- Analgesia

### **Emergence:**

- Pre-oxygenation
- Turn volatile off
- Reversal of neuromuscular block
- Extubation when SV and awake

- ★ Indications for RSI
- ★ Pros/Cons Gas vs IV induction and situations used
- ★ Difficult intubation and failed intubation drills
- ★ Preoxygenation
- ★ Train of Four

## **Recording Intra-operative Care**

This section documents any medications given during the anaesthetic and the times/dosages administered REMEMBER:

Antibiotics/Paracetamol should also be charted on JAC

AAGBI (Association of Anaesthetists of Great Britain and Ireland) Monitoring refers to the minimum standards of monitoring required:

- Pulse Oximetry
- NIBP
- ECG
- Insp./Exp.
   (oxygen/carbon dioxide/volatile gas)
- Peripheral nerve stimulator (if muscle relaxant use)

Additional monitoring may include:

- BIS/EEG
- Cardiac Output

Document HR and BP intra-operatively. Each line = 5 minutes

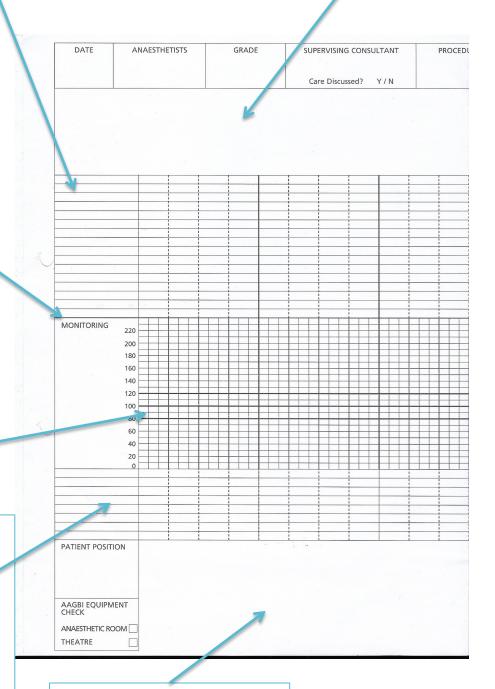
What people record in this section can vary. Parameters generally recorded in this section include:

- FiO<sub>2</sub>
- SpO<sub>2</sub>
- ETCO<sub>2</sub>
- ET (volatile agent) /MAC
- Paw (peak airway pressure)
- Temperature
- Urine output

Generally this section is used to describe your induction events. On old charts it is a useful source of information for details on airway/intubation difficulties

Good things to include

- Access/lines established
- BVM easy/difficult/adjuncts used
- Intubation grade direct/video laryngoscopy/ bougie
- Any blocks performed



Can be used to document any fluids/blood products given intra-operatively

### Recovery and Post-operative management

Following completion of surgery and emergence of anaesthesia your responsibility also lies in safe transfer, handover and review of patients in Recovery.

Before transfer to recovery ensure:

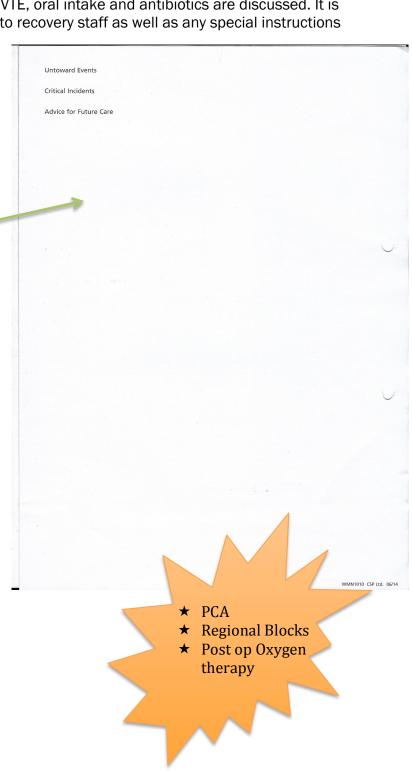
- The patient is physiologically stable and maintain breathing adequately
- Post op analgesia is prescribed (there are post-op pain protocols that can be found on JAC)
- Anti-emetics are prescribed
- All documentation on anaesthetic chart completed
- Doses of antibiotics/paracetamol given intra-operatively are charted and signed for on JAC
- During the surgical time out, plans for VTE, oral intake and antibiotics are discussed. It is important that these are handed over to recovery staff as well as any special instructions

There is space on the back of the anaesthetic chart to allow written handover to recovery staff.

Examples of things to include:

- Oxygen and monitoring requirements
- Acceptable physiological parameters
- Plan for oral intake
- Further IV fluids
- Further doses of antibiotics needed
- Can the patient E+D
- If BMs/blood gas /CXR need doing

Before Leaving Recovery – make sure the nurse is happy and knows where to find you if needed



# **Anaesthetic Drugs**

There are a plethora of drugs used in anaesthetics. As most of them are clear there is a colour coded labeling system in place



Yellow: Induction agents

Blue: Opioids

**Red**: Muscle Relaxants

Pale Orange: Antiemetics

Dark Orange: Benzodiazepines

**Grey: Local Anaesthetics** 

Green: Anticholinergic

Purple: Vasopressors

Red and White Stripes: Reversal agents

White: Miscellaneous

# **Equivalent Drug Dilutions**

Ratio	% Weight (w)/ Volume (v)	=	Grams/ 100ml	Mg/ml	Micrograms/ml
1:10	10%		10 g/ 100ml	100mg/ml	100,000mcg/ml
1:20	5%		5 g/100ml	50mg/ml	50,000 mcg/ml
1: 50	2%		2 g/ 100ml	20mg/ml	20,000mcg/ml
1:100	1%		1 g/ 100ml	10mg/ml	10,000 mcg/ml
1:200	0.5%		0.5 g/ 100ml	5mg/ml	5,000mcg/ml
1:1000	0.1%	,	0.1 g/100ml	1mg/ml	1000 mcg/ml
1: 10,000	0.01%		0.01 g/100ml	0.1mg/ml	100 mcg/ml

- ★ Pharmacology of Anaesthetic Drugs
- ★ Neuromuscular junction physiology
- ★ Detrimental effects of Suxamethonium

**Anaesthetic Drug Reference Sheet** 

Dr	ug	Dose	Comes as/Preparation				
Induction Dr	ugs						
Propofol	<u> </u>	Induction dose 1-2mg/kg	1% 10mg/ml (20ml vial)				
·			2% 20mg/ml				
Ketamine		IV induction 1-2mg/kg	10mg/ml				
		IM induction 5-10mg/kg	50mg/ml				
			100mg/ml				
Thiopental		Induction dose 3-5mg/kg	Yellow Powder reconstituted to				
			2.5% solution				
0 - 1 - 1 - 1 -							
<u>Opioids</u>		1					
Fentanyl		1mcg/kg	50mcg/ml (small = 2 ml vial, large				
Morphine		*titrate to effect	= 10ml vial) 10mg/ml (1 ml vial) diluted to 10ml				
Morphine		"titrate to effect	saline				
Oxycodone		*titrate to effect	10mg/ml (1ml vial) diluted to 10ml				
OAYCOUOTIC			saline				
Muscle Rela	xants						
Atracurium		0.5mg/kg	10mg/ml				
Cisatracurium		0.1-0.2mg/kg	2mg/ml				
2.300.000.70111		1.2 0.2	5mg/ml				
Rocuronium		0.6mg/kg	10mg/ml (5ml vial)				
rtodaromam		RSI: 1mg/kg	Tomig ini (onii viai)				
Suxamethonium		1-1.5mg/kg	50mg/ml (2ml vial)				
Vecuronium		0.1mg/kg	10mg powder (Reconstituted to				
Vecaroniani		J. Zing Ng	1mg/ml)				
			<del></del>				
Reversal Age	ente						
		Neo 2.5mg/Glyco 500mcg (1ml)	(lei)				
Neostigmine/Glycopyrronium Suggamadex		Routine reversal: 2-4mg/kg (TOF					
		twitches dependent) 500mg vial					
		I TWITCHES MENENMENT!					
			Jooning vidi				
		Immediate post RSI: 16mg/kg	Journal Viai				
Local Appear	thotics		Journal Viai				
	thetics	Immediate post RSI: 16mg/kg					
	thetics		0.25% - 2.5mg/ml				
Bupivacaine		Immediate post RSI: 16mg/kg  Max 2mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml				
Bupivacaine		Immediate post RSI: 16mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml				
Bupivacaine		Immediate post RSI: 16mg/kg  Max 2mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml				
Bupivacaine Levobupivacaine		Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml				
Bupivacaine Levobupivacaine		Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml				
Bupivacaine Levobupivacaine		Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml				
Bupivacaine Levobupivacaine Lidocaine		Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml				
Bupivacaine Levobupivacaine Lidocaine  Emergency [	)rugs	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml				
Bupivacaine Levobupivacaine Lidocaine Emergency [ Atropine	Drugs Bradycardia	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml				
Bupivacaine  Levobupivacaine  Lidocaine  Emergency [ Atropine Glycopyrronium	Drugs Bradycardia Bradycardia	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)  20mcg/kg  20mcg/kg  200mcg bolus	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 600mcg/ml 200mcg/ml				
Bupivacaine Levobupivacaine Lidocaine Emergency [ Atropine Glycopyrronium	Drugs Bradycardia	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 600mcg/ml 200mcg/ml 30mg diluted in 10ml saline				
Local Anaes: Bupivacaine Levobupivacaine Lidocaine  Emergency [ Atropine Glycopyrronium Ephedrine	Drugs Bradycardia Bradycardia Hypotension	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)  20mcg/kg  20mcg/kg  20mcg bolus  3mg bolus	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 200mcg/ml 30mg diluted in 10ml saline (pre-formed syringe)				
Bupivacaine  Levobupivacaine  Lidocaine  Emergency [ Atropine Glycopyrronium Ephedrine	Drugs Bradycardia Bradycardia	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)  20mcg/kg  20mcg/kg  200mcg bolus	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 600mcg/ml 200mcg/ml 30mg diluted in 10ml saline (pre-formed syringe) 0.1mg/ml				
Bupivacaine Levobupivacaine Lidocaine  Emergency E Atropine Glycopyrronium Ephedrine Phenylephrine	Drugs Bradycardia Bradycardia Hypotension Hypotension	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)  20mcg/kg  200mcg bolus  3mg bolus  bolus	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 30mg/ml 30mg diluted in 10ml saline (pre-formed syringe) 0.1mg/ml 10mg/ml				
Bupivacaine Levobupivacaine Lidocaine Emergency [ Atropine Glycopyrronium	Drugs Bradycardia Bradycardia Hypotension	Immediate post RSI: 16mg/kg  Max 2mg/kg  Max 2mg/kg  Max 3mg/kg  Max 6mg/kg (with adrenaline)  20mcg/kg  20mcg/kg  20mcg bolus  3mg bolus	0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.25% - 2.5mg/ml 0.5% - 5mg/ml 0.75% - 7.5mg/ml 1% 10mg/ml 2% 20mg/ml 600mcg/ml 200mcg/ml 30mg diluted in 10ml saline (pre-formed syringe) 0.1mg/ml				

### Machine Basics and the Circle System

The anaesthetic machine is a complex multicomponent system. The following section aims only to give the general principles of how the machine functions and outlines the principles of the circle system

Broadly, the function of the machine can be divided into

- 1. Supply
- 2. Processing and pressure regulation
- 3. Delivery
- 4. Disposal

### Breathing systems:

The function of any breathing system is to deliver oxygen, anaesthetic gases and to eliminate CO<sub>2</sub>.

They can be:

- Non rebreathing systems
- Rebreathing (circle) system \*most common\*

The key feature of the circle system is that it uses one-way valves to ensure unidirectional flow and a CO<sub>2</sub> absorber (soda lime) to capture CO<sub>2</sub>. This allows for low fresh gas flows and anaesthetic gas usage without toxic build up of carbon dioxide.

#### The Anaesthetic Machine

# 1. Supply

Gas supply can come from:

- Pipelines
  - Vacuum Insulated Evaporator
  - Cylinder Manifold
- Cylinders

#### 2. Processing and Pressure Regulation

• Both pipeline and cylinder gas enter the system at very high pressure so to prevent damage to the machine; pressure regulation valves reduce this to a suitable supply pressure.

Pipeline supply is used preferentially

- Within the system there is a low oxygen pressure alarm with oxygen reservoir. This kicks in if pressure <200kPa. There is a N<sub>2</sub>O shut off valve to prevent hypoxic mixture in absence of low oxygen.
- Gas Flow then enters the rotameters that further reduce the pressure to just above atmospheric (100Kpa). Rotameters also have a hypoxic guard mechanism to prevent hypoxic gas mixtures when using  $N_2O$
- Gas flow then pass through vapourisers, which can be mechanically, or electronically controlled. Each is calibrated for a particular vapour.

### 3. Delivery

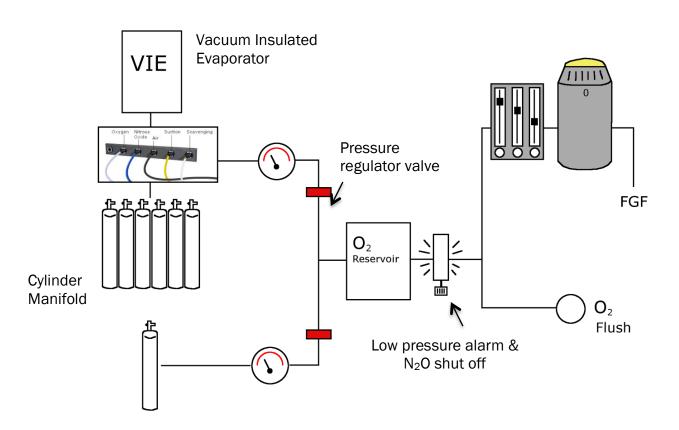
\*Note the Oxygen Flush is outside of the flow meter pressure reduction – therefore using this give high-pressure oxygen into the system – Should therefore not be used routinely.



All gas mixtures (Fresh Gas Flow FGF) then pass into the common gas outlet and enter the breathing system. Below – the circle system is discussed.

### 4. Disposal

Anaesthetic Gas Scavenging (AGSS) actively pipes waste anaesthetic gases and vents into atmosphere



### The Circle System

- FGF enters the Inspiratory limb via the one-way valve to the patient.
- Exhaled gases travel in the expiratory limb. This also has a one-way valve to prevent back flow of expired gases to the patient
- During SV both expiratory tubing and the reservoir bag contains expired gases. When the
  pressure limit is reached, these escape via the adjustable pressure limiting valve (APL) and are
  scavenged by AGSS (prevents bag explosions!)
- The APL should be open at minimal pressure during SV (otherwise imagine trying to breath out through narrow straws)

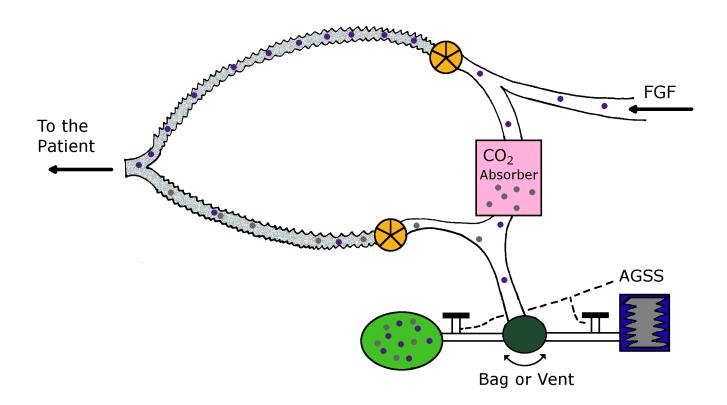
This can feature can be utilised however if required to give some PEEP

• Expired gas that does not escape via the APL travels in the expiratory tubing to the CO<sub>2</sub> absorber (soda lime) where chemical change occurs allowing return of gases (now CO<sub>2</sub> free!) to return to the inspiratory limb and into the patient

• The patient then breaths in again using gases contained in the reservoir bag (which has passed through the soda lime) in addition to any FGF from the machine

#### POSITIVE PRESSURE VENTAILATION

- When the patient no longer drives inspiration through negative pressure, ventilation can be maintained either through manual positive pressure ventilation by squeezing reservoir bag or through the ventilator driving bellows
- When squeezing the reservoir bag to deliver gases if the APL if set to minimum, it would let the
  gas mixture escape. Therefore to ventilate the patient, the APL valve needs to be set to open
  only at maximal pressure. This is problematic however as you could only manual ventilate a few
  breaths before the reservoir bag expands and pressure in the system would increase.
- You would need to continually flip between maximal pressure when delivering a breath and minimal pressure during expiration. When manually ventilating a patient you therefore compromise with "middle" APL setting
- The ventilator mechanically controls its own ventilator APL valve and so it able to manually switch these pressure more efficiently. When the ventilator is in use, the manual APL valve and reservoir bag are disconnected from the system



## Machine and Ventilation Settings - THE BASICS

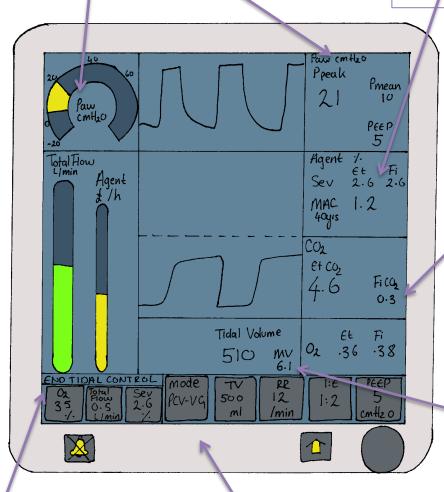
Peak airway Pressure (Paw) = Relates to the resistive airway pressure Should be kept <30cmH<sub>2</sub>O to prevent barotrauma. Causes of high peak airway pressure can be thought

 Machine causes: kinked tubing, waterlogged HME

of as:

 Patient Causes: Bronchospasm, Tube biting, Mucus plug

- Minimum alveolar concentration (MAC)
   of an inhaled anaesthetic = the alveolar
   concentration at which 50% of patients
   show no motor response to a standard
   midline surgical incision
- MAC changes with age and so is often adjusted
- Is one method of gauging depth of anaesthesia



In addition to ensuring ETCO<sub>2</sub> is maintained at an appropriate level, the capnography trace itself can be a useful determinant of ventilation Increased Inspired FiCO<sub>2</sub> can indicate problems – e.g exhausted soda lime

Tidal Volume (aim 7-9 ml/kg)

 $MV = TV \times RR$ 

End Tidal Control: set the ETO<sub>2</sub>, flow rate, ET volatile .The machine then adjusts the levels of FGF to meet this requirement

Manual gas control: Controlled and adjusted manually by the anaesthetist

Ventilator modes commonly used:

- Pressure control ventilation (PCV): Set the pressure and TV delivered varies on lung compliance
- Volume control ventilation (VCV): set the TV peak airway pressure varies depending on patient lung compliance
- Pressure controlled Volume Guaranteed (PCV-VG): Newer ventilation method where you set the TV and the ventilator adjusts inspiratory pressure breath by breath to deliver the volume set