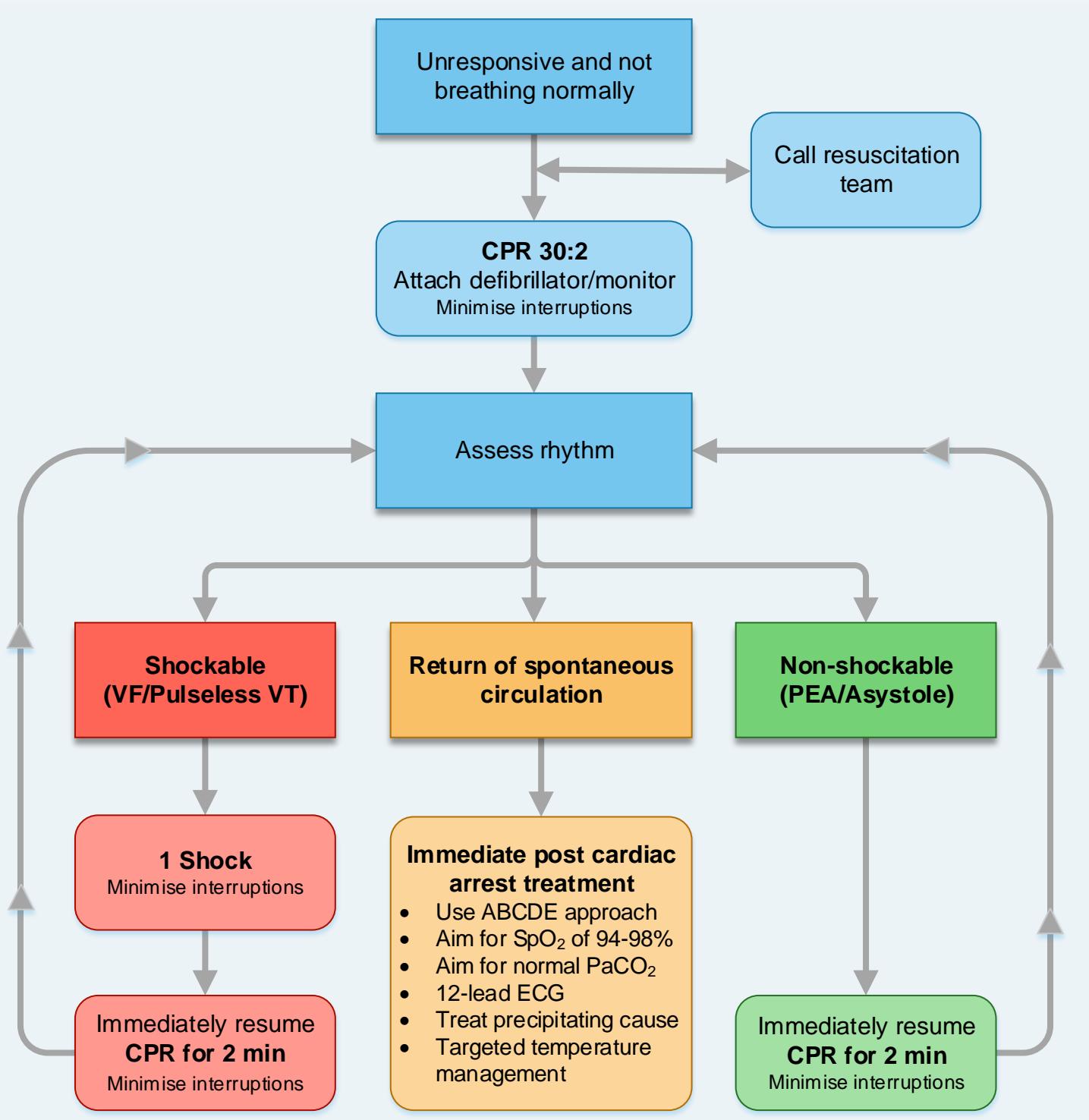
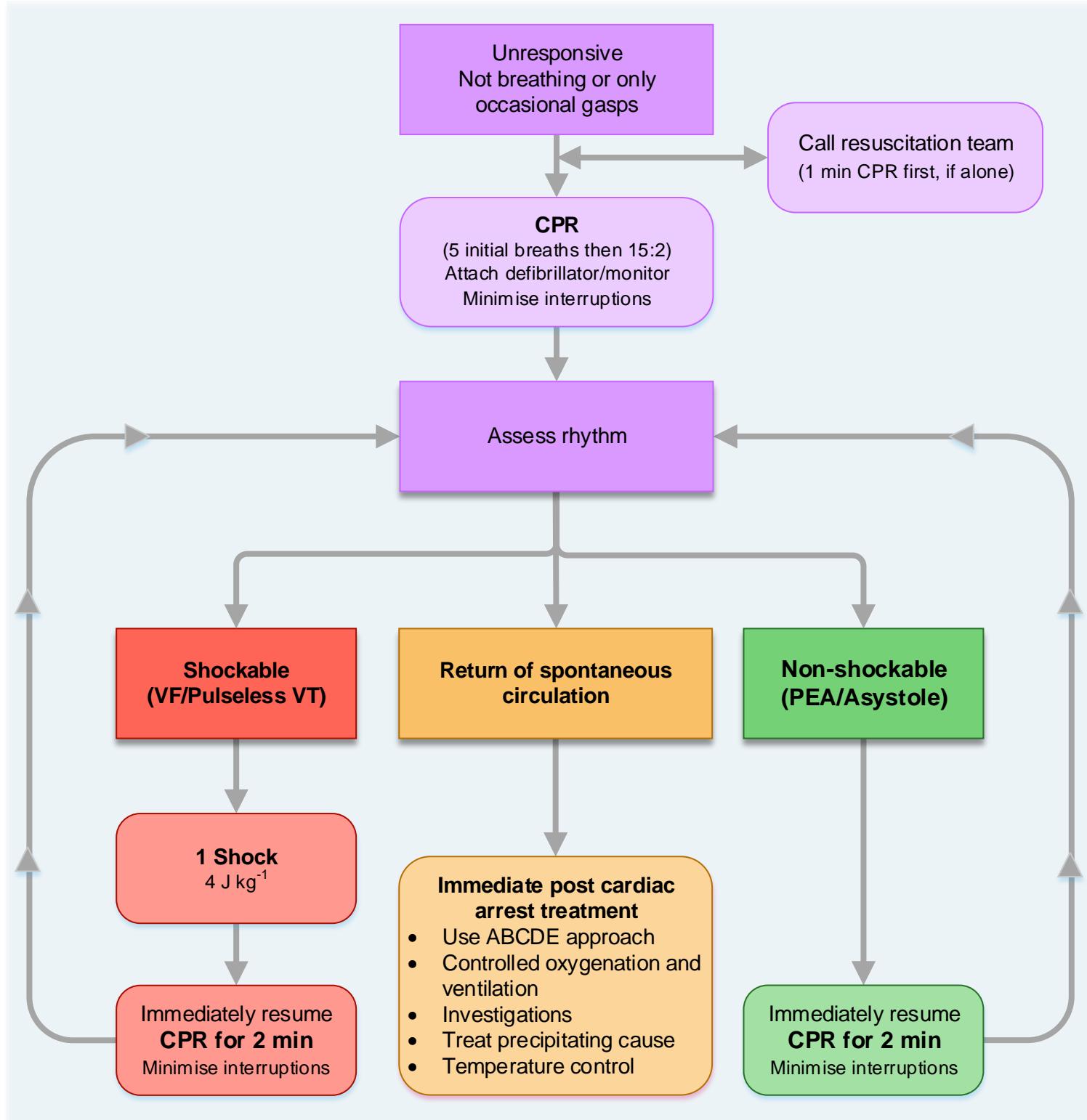




Emergency guidelines - UK based

- 2. ALS Adult**
- 3. ALS Children**
- 4. Airway management ABCD (DAS)**
- 5. Intubation checklist (NAP4)**
- 6. Unanticipated difficult intubation (DAS)**
- 7. Difficult intubation during RSI (DAS)**
- 8. Failed intubation and difficult ventilation (DAS)**
- 9. Trachy emergency – patent upper airway (NTSP)**
- 10. Trachy emergency – laryngectomy (NTSP)**
- 11. Trachy bleeding (local/ICMWK)**
- 12. Sudden hypoxia on ICU (local/ICMWK)**
- 13. Adult tachycardia (RCUK)**
- 14. Adult bradycardia (RCUK)**
- 15. Sudden hypotension (local/ICMWK)**
- 16. Post-resuscitation (RCUK)**
- 17. Anaphylaxis (RCUK)**
- 18. Extravasation (Kansas univ)**
- 19. LA toxicity (AAGBI)**
- 21. Malignant hyperthermia (AAGBI)**



**During CPR**

- Ensure high-quality CPR: rate, depth, recoil
- Plan actions before interrupting CPR
- Give oxygen
- Vascular access (intravenous, intraosseous)
- Give adrenaline every 3-5 min
- Consider advanced airway and capnography
- Continuous chest compressions when advanced airway in place
- Correct reversible causes
- Consider amiodarone after 3 and 5 shocks

Reversible Causes

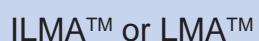
- Hypoxia
- Hypovolaemia
- Hyper/hypokalaemia, metabolic
- Hypothermia
- Thrombosis (coronary or pulmonary)
- Tension pneumothorax
- Tamponade (cardiac)
- Toxic/therapeutic disturbances

Plan A:
Initial tracheal intubation plan



failed intubation

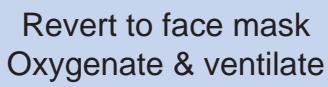
Plan B:
Secondary tracheal intubation plan



failed oxygenation

failed intubation

Plan C:
Maintenance of oxygenation, ventilation, postponement of surgery and awakening



failed oxygenation

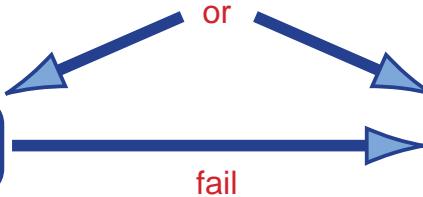
Plan D:
Rescue techniques for "can't intubate, can't ventilate" situation



increasing hypoxaemia

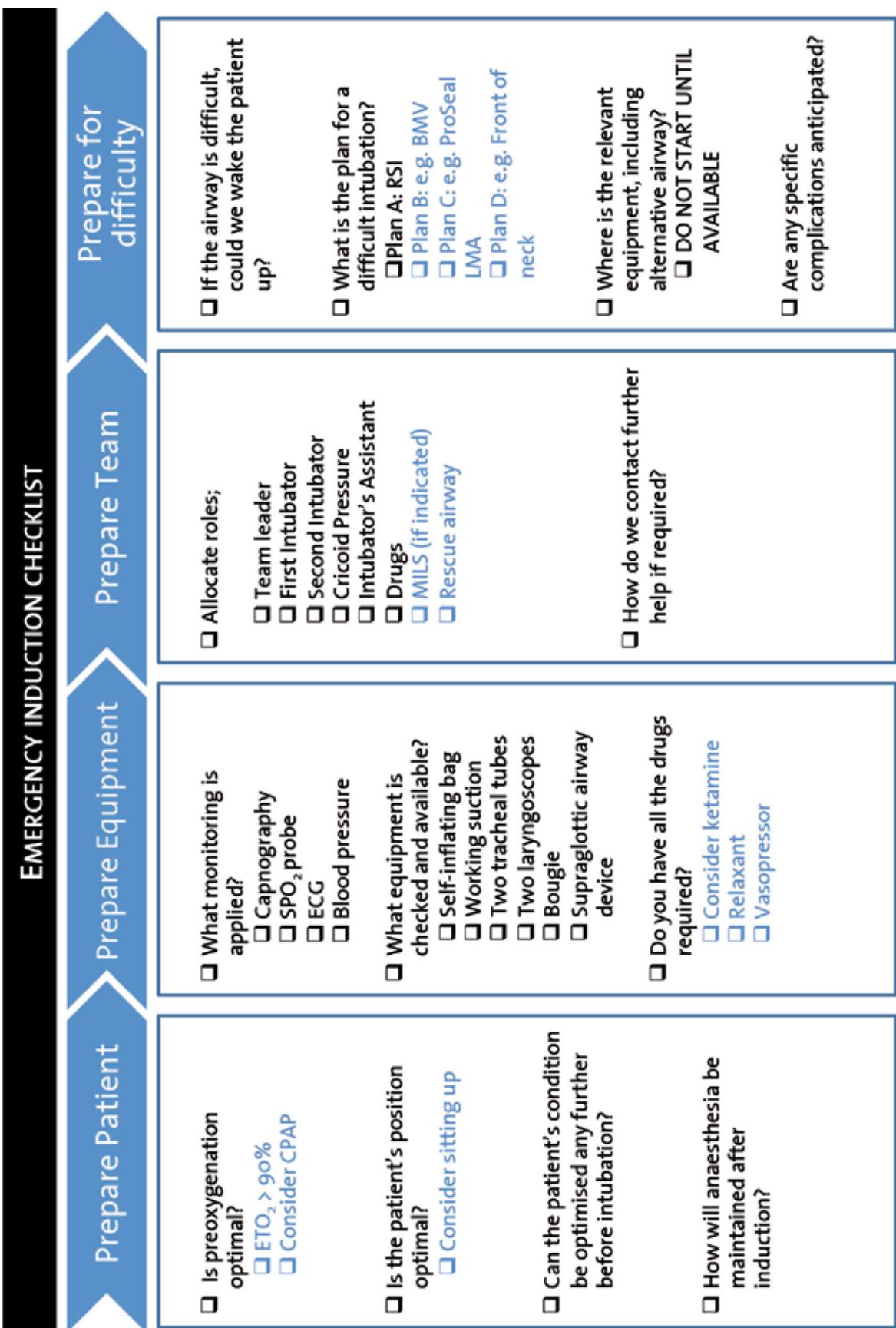
Cannula
cricothyroidotomy

Surgical
cricothyroidotomy



APPENDIX 1

Example intubation checklist for ICU and emergency department



This Checklist is not intended to be a comprehensive guide to preparation for induction



Unanticipated difficult tracheal intubation-
during routine induction of anaesthesia in an adult patient

Direct laryngoscopy → Any problems → Call for help

Plan A: Initial tracheal intubation plan

Direct laryngoscopy - check:
 Neck flexion and head extension
 Laryngoscope technique and vector
 External laryngeal manipulation - by laryngoscopist
 Vocal cords open and immobile
 If poor view: Introducer (bougie) - seek clicks or hold-up and/or Alternative laryngoscope

Not more than 4 attempts, maintaining:
 (1) oxygenation with face mask and
 (2) anaesthesia

succeed

Tracheal intubation

Verify tracheal intubation
 (1) Visual, if possible
 (2) Capnograph
 (3) Oesophageal detector
 "If in doubt, take it out"

failed intubation

Plan B: Secondary tracheal intubation plan

ILMA™ or LMA™
 Not more than 2 insertions
 Oxygenate and ventilate

succeed

Confirm: ventilation, oxygenation, anaesthesia, CVS stability and muscle relaxation - then fibreoptic tracheal intubation through ILMA™ or LMA™ - 1 attempt
 If LMA™, consider long flexometallic,nasal RAE or microlaryngeal tube
 Verify intubation and proceed with surgery

failed oxygenation
 (e.g. $\text{SpO}_2 < 90\%$ with $\text{FiO}_2 1.0$)
 via ILMA™ or LMA™

failed intubation via ILMA™ or LMA™

Plan C: Maintenance of oxygenation, ventilation, postponement of surgery and awakening

Revert to face mask
 Oxygenate and ventilate
 Reverse non-depolarising relaxant
 1 or 2 person mask technique (with oral ± nasal airway)

succeed

Postpone surgery
 Awaken patient

failed ventilation and oxygenation

Plan D: Rescue techniques for "can't intubate, can't ventilate" situation



Unanticipated difficult tracheal intubation - during rapid sequence induction of anaesthesia in non-obstetric adult patient



Plan A: Initial tracheal intubation plan

Pre-oxygenate

Cricoid force: 10N awake → 30N anaesthetised

Direct laryngoscopy - check:

Neck flexion and head extension

Laryngoscopy technique and vector

External laryngeal manipulation - by laryngoscopist

Vocal cords open and immobile

If poor view:

Reduce cricoid force

Introducer (bougie) - seek clicks or hold-up and/or Alternative laryngoscope

succeed

Tracheal intubation

Not more than 3 attempts, maintaining:

- (1) oxygenation with face mask
- (2) cricoid pressure and
- (3) anaesthesia

Verify tracheal intubation
 (1) Visual, if possible
 (2) Capnograph
 (3) Oesophageal detector
 "If in doubt, take it out"

failed intubation

Plan C: Maintenance of oxygenation, ventilation, postponement of surgery and awakening

Maintain 30N cricoid force

Plan B not appropriate for this scenario

Use face mask, oxygenate and ventilate
 1 or 2 person mask technique
 (with oral ± nasal airway)
 Consider reducing cricoid force if ventilation difficult

succeed

failed oxygenation

(e.g. $\text{SpO}_2 < 90\%$ with $\text{FiO}_2 1.0$) via face mask

LMA™

Reduce cricoid force during insertion
 Oxygenate and ventilate

failed ventilation and oxygenation

succeed

Postpone surgery and awaken patient if possible or continue anaesthesia with LMA™ or ProSeal LMA™ - if condition immediately life-threatening

Plan D: Rescue techniques for "can't intubate, can't ventilate" situation



Failed intubation, increasing hypoxaemia and difficult ventilation in the paralysed anaesthetised patient: Rescue techniques for the "can't intubate, can't ventilate" situation

failed intubation and difficult ventilation (other than laryngospasm)

Face mask
Oxygenate and Ventilate patient
Maximum head extension
Maximum jaw thrust
Assistance with mask seal
Oral ± 6mm nasal airway
Reduce cricoid force - if necessary

failed oxygenation with face mask (e.g. SpO₂ < 90% with FiO₂ 1.0)

call for help

LMA™ Oxygenate and ventilate patient
Maximum 2 attempts at insertion
Reduce any cricoid force during insertion

succeed

Oxygenation satisfactory and stable: Maintain oxygenation and awaken patient

"can't intubate, can't ventilate" situation with increasing hypoxaemia

Plan D: Rescue techniques for "can't intubate, can't ventilate" situation

or

Cannula cricothyroidotomy

Equipment: Kink-resistant cannula, e.g. Patil (Cook) or Ravussin (VBM)
High-pressure ventilation system, e.g. Manujet III (VBM)

Technique:

1. Insert cannula through cricothyroid membrane
2. Maintain position of cannula - assistant's hand
3. Confirm tracheal position by air aspiration - 20ml syringe
4. Attach ventilation system to cannula
5. Commence cautious ventilation
6. Confirm ventilation of lungs, and exhalation through upper airway
7. If ventilation fails, or surgical emphysema or any other complication develops - convert immediately to surgical cricothyroidotomy

fail

Surgical cricothyroidotomy

Equipment: Scalpel - short and rounded (no. 20 or Minitrach scalpel)
Small (e.g. 6 or 7 mm) cuffed tracheal or tracheostomy tube

4-step Technique:

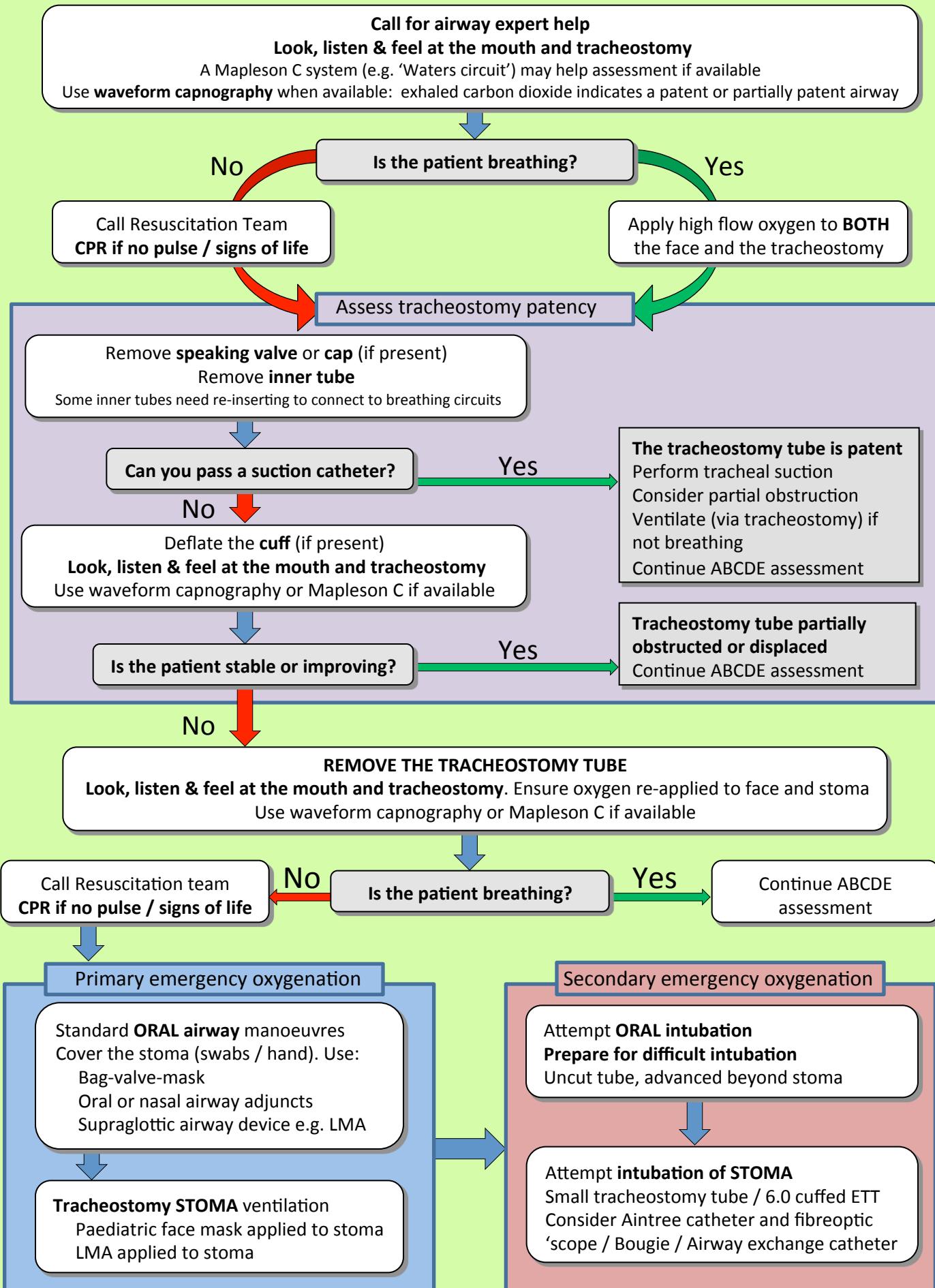
1. Identify cricothyroid membrane
2. Stab incision through skin and membrane
Enlarge incision with blunt dissection (e.g. scalpel handle, forceps or dilator)
3. Caudal traction on cricoid cartilage with tracheal hook
4. Insert tube and inflate cuff
Ventilate with low-pressure source
Verify tube position and pulmonary ventilation

Notes:

1. These techniques can have serious complications - use only in life-threatening situations
2. Convert to definitive airway as soon as possible
3. Postoperative management - see other difficult airway guidelines and flow-charts
4. 4mm cannula with low-pressure ventilation may be successful in patient breathing spontaneously



Emergency tracheostomy management - Patent upper airway



Emergency laryngectomy management

Call for airway expert help

Look, listen & feel at the mouth and laryngectomy stoma

A Mapleson C system (e.g. 'Waters circuit') may help assessment if available

Use waveform capnography whenever available: exhaled carbon dioxide indicates a patent or partially patent airway

No

Is the patient breathing?

Yes

Call Resuscitation Team
CPR if no pulse / signs of life

Apply high flow oxygen to laryngectomy stoma

If any doubt whether patient has a laryngectomy, apply oxygen to face also*

Assess laryngectomy stoma patency

Most laryngectomy stomas will NOT have a tube in situ

Remove stoma cover (if present)
Remove inner tube (if present)

Some inner tubes need re-inserting to connect to breathing circuits
Do not remove a tracheoesophageal puncture (TEP) prosthesis

The laryngectomy stoma is patent
Perform tracheal suction
Consider partial obstruction
Ventilate via stoma if not breathing
Continue ABCDE assessment

Can you pass a suction catheter?

Yes

Deflate the cuff (if present)
Look, listen & feel at the laryngectomy stoma or tube
Use waveform capnography or Mapleson C if available

Yes

Continue ABCDE assessment

No

REMOVE THE TUBE FROM THE LARYNGECTOMY STOMA if present

Look, listen & feel at the laryngectomy stoma. Ensure oxygen is re-applied to stoma
Use waveform capnography or Mapleson C if available

No

Is the patient breathing?

Yes

Continue ABCDE assessment

Call Resuscitation Team
CPR if no pulse / signs of life

Primary emergency oxygenation

Laryngectomy stoma ventilation via either
Paediatric face mask applied to stoma
LMA applied to stoma

Secondary emergency oxygenation

Attempt intubation of laryngectomy stoma
Small tracheostomy tube / 6.0 cuffed ETT
Consider Aintree catheter and fibroscopic scope / Bougie / Airway exchange catheter

* Laryngectomy patients have an end stoma and **cannot be oxygenated via the mouth or nose**
Applying oxygen to the face and stoma is the default emergency action for all patients with a tracheostomy

Tracheostomy bleeding

Lightly blood-stained secretions only?

Early (days 1-3) – likely benign - observe

Late (after day 3) – senior review

Significant frank blood?

Call for help	<input type="checkbox"/>	1. ICU consultant and senior nursing staff 2. ENT surgeon – on call consultant at John Radcliffe will coordinate. May also be one in clinic or theatre on site (put days here) 3. General surgeon 4. Senior anaesthetist
100% oxygen	<input type="checkbox"/>	Or high flow via trachy mask
Inflate cuff	<input type="checkbox"/>	Temporary over-inflation can be considered (usually at least 20 ml (for Portex tubes)
Sit nearly fully upright	<input type="checkbox"/>	45 degrees if hypotensive
Suction via tracheostomy	<input type="checkbox"/>	Large gauge catheter to retrieve clots
Attach water's circuit or self-inflating bag, and end-tidal monitoring	<input type="checkbox"/>	Consider assisting ventilation or take over
Finger pressure in sternal notch or at bleeding point if obvious	<input type="checkbox"/>	If heavily sedated, consider packing pharynx
Suction pharynx	<input type="checkbox"/>	Remove clots
Next measures	<input type="checkbox"/>	1. Kaltostat or adrenaline-soaked ribbon gauze - tuck into trachy wound. 2. Inject lidocaine + adrenaline 1:200,000 around trachy site. 3. Sedation if coughing is excessive or struggling with ventilation (+/- muscle relaxant) 4. Correct known coagulopathy or reverse anticoagulants where possible

Bleeding still through tracheostomy despite cuff inflation?

Profuse?	<input type="checkbox"/>	Consider intubation via mouth and: <ul style="list-style-type: none"> • distal balloon placement • Intentional endobronchial intubation • double-lumen tube • bronchial blocker. Or consider palliation.
Not profuse but not settling?	<input type="checkbox"/>	Adrenaline nebs IV +/- nebulized tranexamic acid. Consider plans as for profuse bleeding.
Haemostasis achieved?	<input type="checkbox"/>	Bronchoscopy and upper airway scope CXR Physiotherapy Consider possibility that it was a 'herald bleed' (re TIF below)

Causes

Early

- Suction or movement of tracheostomy

Late

- Granulation tissue
- Stoma site infection
- Tracheo-innominate fistula

Also consider:

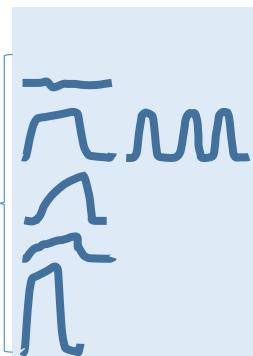
- Bleeding from non-trachy surgical site if present and nearby.
- Upper respiratory tract bleeding (eg epistaxis)
- Haemoptysis unrelated to tracheostomy

Sudden hypoxia on the ICU – cause not clear

1. Call for help

2. 100 % inspired oxygen

3. Check end tidal CO₂ trace



Flat or n/a

Normal MV or SV trace

Bronchospasm or obstructed airway

Falling CO or poor compliance

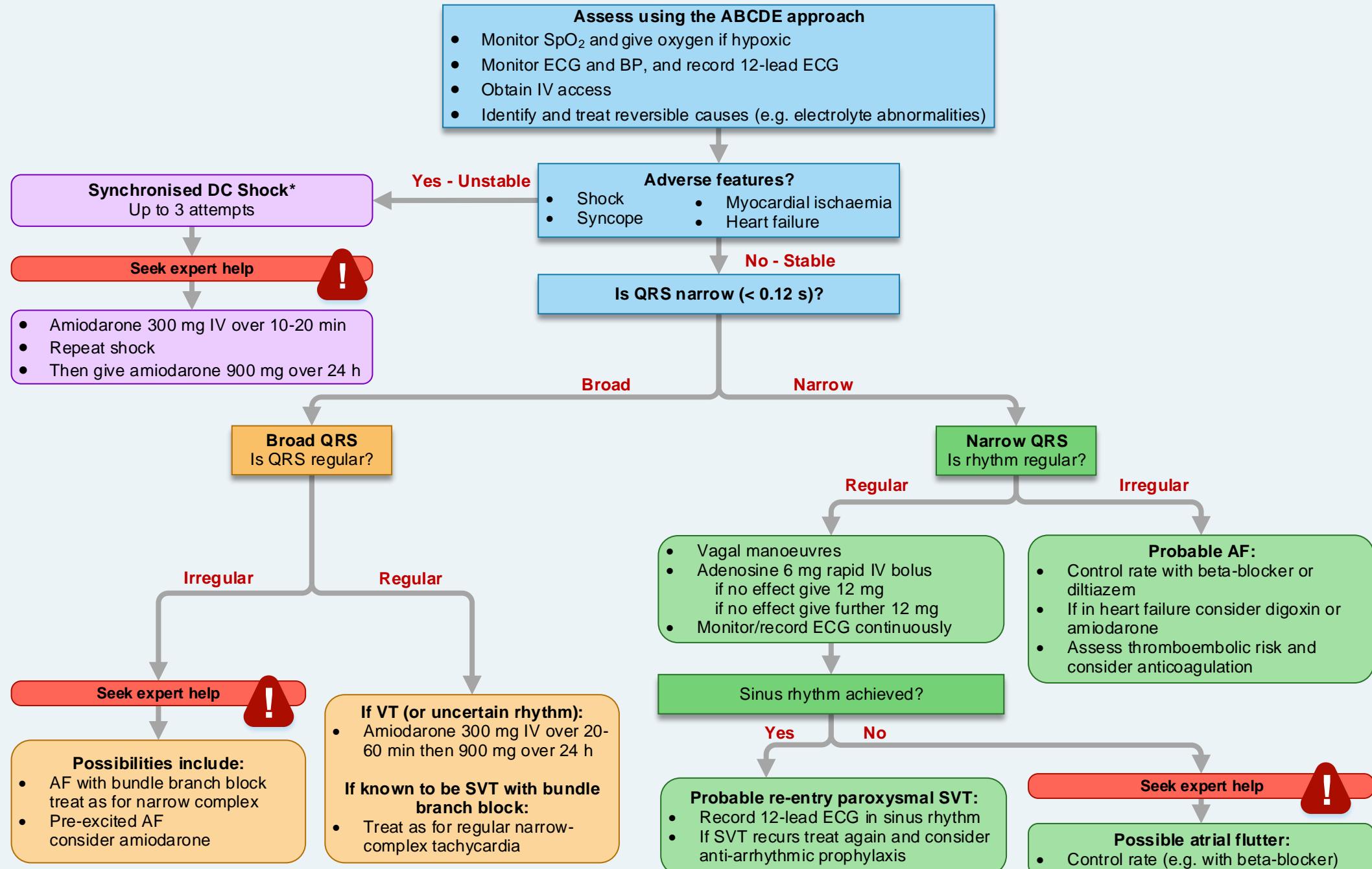
Inadequate MV (or hypermetabolic state)

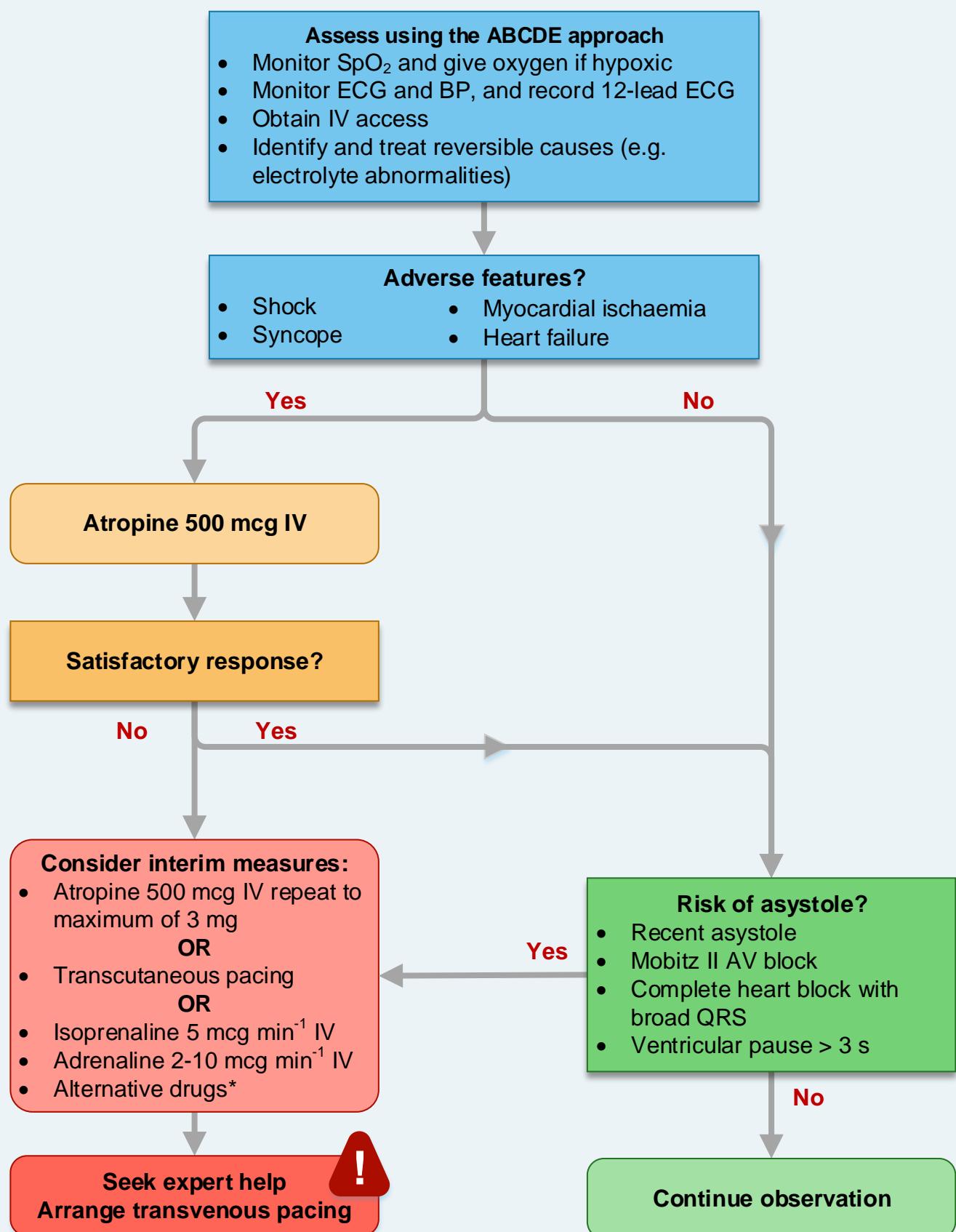
Consider cardiac arrest, ventilator failure, then tracheal or tube obstruction - follow tracheal tube algorithm

4. Tracheal suction (in-line if present, or use suction catheters)

5. Assess chest movement and listen to chest

Findings	Diagnoses to consider	Management
Obstructed appearance (see-sawing), bilaterally quiet, ↓Expansion & compliance	Consider tracheal or tube obstruction	Follow tracheal tube algorithm. Then consider these:
Quiet & reduced expansion on one side , + reduced compliance	Endobronchial intubation Has ETT moved in further (usually not more than 24cm at teeth)?	<ul style="list-style-type: none"> Consider withdrawing 1-2 cm CXR +/- bronchoscopy
	Pneumothorax or haemothorax Trauma, new line or drain, high ventilation pressure, in-situ chest drain fault.	<ul style="list-style-type: none"> Needle decompression or drain (or resolve drain issue). CXR/CT if stabilises
	Pulmonary haemorrhage Bronchiectasis, abscess, TB, or tumour?	<ul style="list-style-type: none"> Airway specialist and thoracic surgeon involvement. CXR/CT
	Collapse/atelectasis due to sputum, or aspiration	<ul style="list-style-type: none"> Suction (+/- saline) Recruitment breaths with ventilator or bag Consider bronchoscopy
New widespread wheeze +/- reduced compliance	Bronchospasm COPD/asthma Anaphylaxis Aspiration Feeding tube malposition Vomiting Pulmonary oedema New ischaemia Recent fluid bolus	<ul style="list-style-type: none"> Suction ECG/CXR +/- Echo/troponin Consider <ul style="list-style-type: none"> Bronchodilators/steroid Furosemide/nitrate Stop feeding, empty stomach Bronchoscopy
New widespread crackles +/- reduced compliance	Pulmonary oedema New ischaemia Recent fluid bolus Aspiration Feeding tube malposition Vomiting	<ul style="list-style-type: none"> Suction ECG/CXR +/- Echo/troponin Consider <ul style="list-style-type: none"> Furosemide/nitrate Stop feeding, empty stomach Bronchoscopy
Normal breath sounds and compliance	Low cardiac output state MI PE Haemorrhage Tamponade	<ul style="list-style-type: none"> ECG/Echo Assess volume status (?CO monitoring, PPV) FAST/CT





* Alternatives include:

- Aminophylline
- Dopamine
- Glucagon (if bradycardia is caused by beta-blocker or calcium channel blocker)
- Glycopyrrolate (may be used instead of atropine)

Sudden hypotension on ICU – cause not clear

Call for help!

Ensure hypotension is not due to respiratory compromise	<input type="checkbox"/> Obvious pneumothorax? <input type="checkbox"/> Ventilation difficulty - disconnection, bronchospasm, sudden drop in Vt?
--	---

SBP <50 mmHg or possible loss of cardiac output? start CPR.

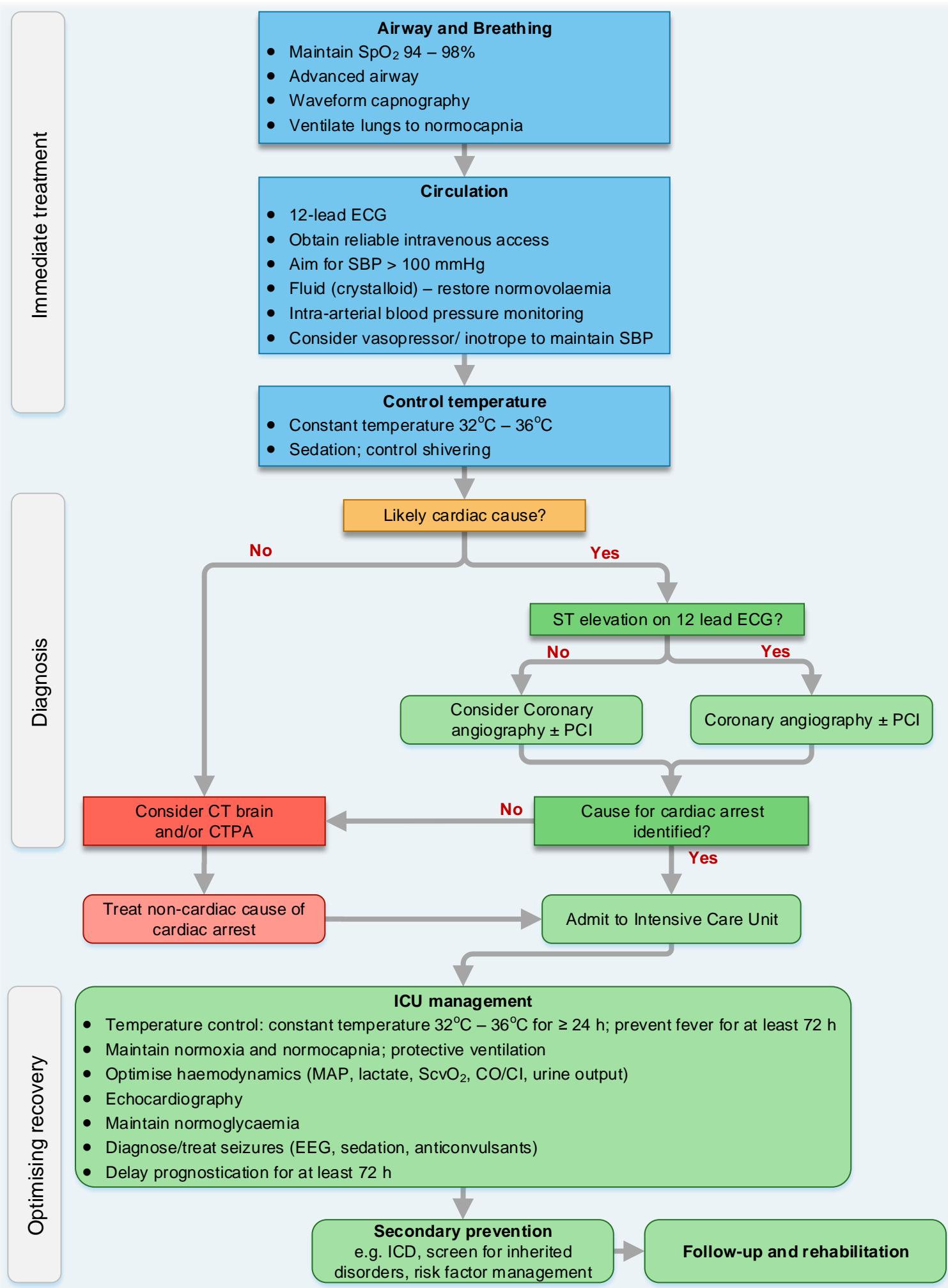
Tilt bed or raise legs	<input type="checkbox"/>
Request fluid bolus	<input type="checkbox"/> Hartmann's and plan to run 500ml stat
Request bolus dose vasopressor	<input type="checkbox"/> Metaraminol 0.5mg/ml 0.5mg (1ml) at a time Adrenaline 10 mcg/ml (1ml from a minijet into 10ml saline) 10 mcg (1ml) at a time
Is this the true BP and what is the rhythm?	<input type="checkbox"/> Feel for pulse. Look at the end-tidal CO ₂ trace. Quickly check arterial line waveform and transducer position. If there is a waveform, assume it's correct. Re-zero if necessary.
Look for blood	<input type="checkbox"/> Check for blood in the bed, bowel, stomach, new tense abdomen, or leak from haemofilter circuit
Check infusions	<input type="checkbox"/> Check for kinked line, displaced CVC, syringe driver failure?
Check skin	<input type="checkbox"/> Look for new rash and swelling
ECG	<input type="checkbox"/>
Request or do ultrasound FICE +/- FAST	<input type="checkbox"/> Decide whether this is hypovolaemic, distributive, cardiogenic (including arrhythmia) or obstructive shock.
Request CXR	<input type="checkbox"/>
Cardiac output monitoring	<input type="checkbox"/> ODM if asleep and no central access, otherwise LIDCO, or echo.

Consider :

- Overt or concealed haemorrhage
- MI
- Arrhythmia – rule out VT, if new AF consider cardioversion.
- PE
- Concealed haemorrhage – retroperitoneal, small bowel, thorax
- Tamponade (recent central venous access, pacing wire etc.)
- Drug reaction – anaphylaxis or inadvertent bolus (including epidural)
- Intracranial haemorrhage
- New sepsis – take cultures and get antibiotic advice

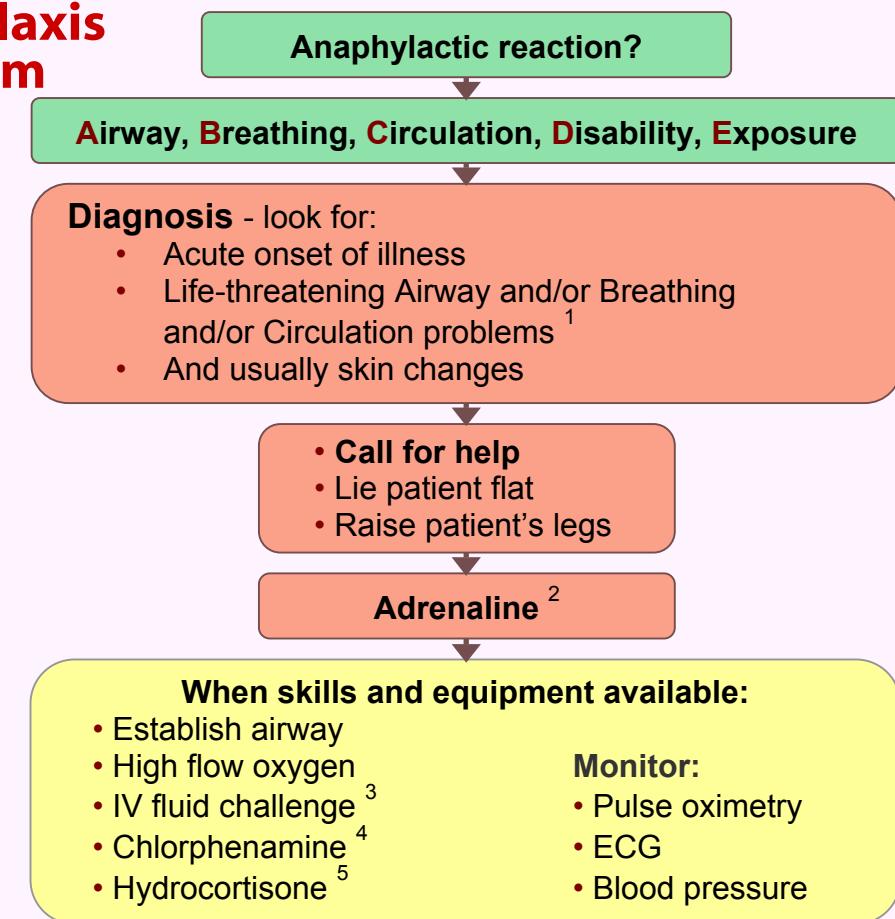
Further treatment:

- Consider increasing dose of adrenaline or noradrenaline
NOT dobutamine/dopamine/milrinone or vasopression
- Treat specific problem
order blood, plan for DCCV, CT; **call** surgeon, endoscopist, cardiologist





Anaphylaxis algorithm



1 Life-threatening problems:

Airway: swelling, hoarseness, stridor

Breathing: rapid breathing, wheeze, fatigue, cyanosis, $\text{SpO}_2 < 92\%$, confusion

Circulation: pale, clammy, low blood pressure, faintness, drowsy/coma

2 Adrenaline (give IM unless experienced with IV adrenaline)

IM doses of 1:1000 adrenaline (repeat after 5 min if no better)

- Adult 500 micrograms IM (0.5 mL)
- Child more than 12 years: 500 micrograms IM (0.5 mL)
- Child 6 -12 years: 300 micrograms IM (0.3 mL)
- Child less than 6 years: 150 micrograms IM (0.15 mL)

Adrenaline IV to be given **only by experienced specialists**

Titrate: Adults 50 micrograms; Children 1 microgram/kg

3 IV fluid challenge:

Adult - 500 – 1000 mL
Child - crystalloid 20 mL/kg

Stop IV colloid
if this might be the cause
of anaphylaxis

4 Chlorphenamine (IM or slow IV)

Adult or child more than 12 years	10 mg
Child 6 - 12 years	5 mg
Child 6 months to 6 years	2.5 mg
Child less than 6 months	250 micrograms/kg

5 Hydrocortisone (IM or slow IV)

200 mg
100 mg
50 mg
25 mg

March
2008

GUIDE TO EXTRAVASATION MANAGEMENT IN ADULT & PEDIATRIC PATIENTS

Large, well-designed, controlled clinical trials in humans are not available to support the development of extravasation management guidelines. Available data generally consists of case reports, trials utilizing animal models, and small studies with evidence of poor or inconsistent quality. This lack of evidence creates challenges in validating specific interventions and presents barriers to guideline development.

Interventions listed within this guide were derived from a consensus of the cited tertiary references. Greater consideration was given to more detailed, substance-specific references when a consensus was not apparent.

The information provided is intended as a general guide only. Consult additional references and product labeling for more detailed information.

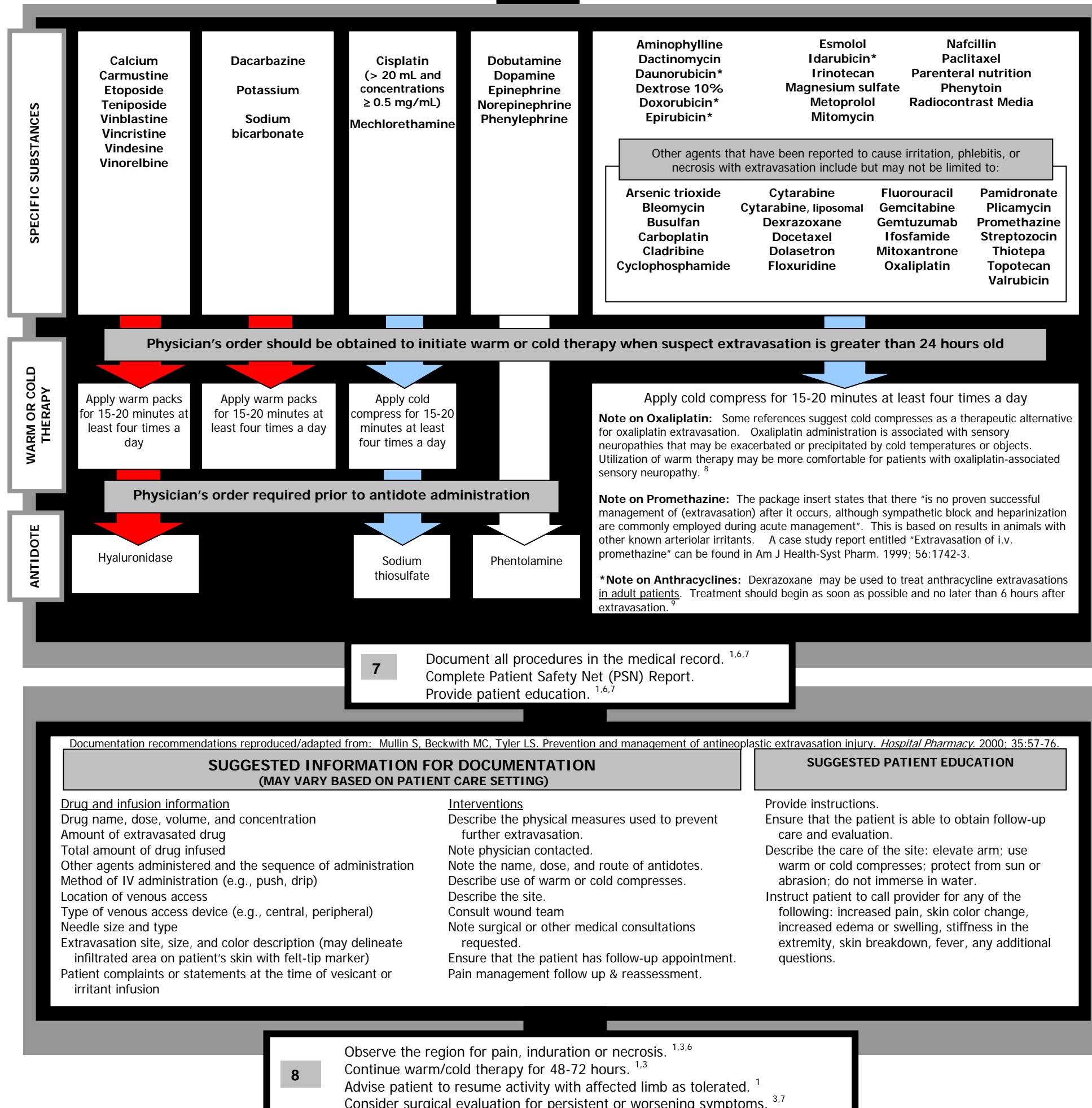
THE UNIVERSITY
OF KANSAS HOSPITAL
KUMED

RELATED POLICIES

- Nursing Standard of Practice & Procedures:
- 1) Extravasations, Patient Management of
- 2) Care of the Patient by a Non-Chemotherapy Certified RN

REFERENCES

1. Mullin S, Beckwith CM, Tyler LS. Prevention and management of antineoplastic extravasation injury. *Hospital Pharmacy*. 2000; 35:57-76.
2. eFacts [database online]. St. Louis, MO: Wolters Kluwer Health, Inc.; 2005.
3. Gahart BL, Nazareno AR. Intravenous Medications. 21st ed. St Louis: Elsevier Mosby; 2005.
4. McEvoy G, ed. American Hospital Formulary Service: Drug Information. Bethesda: American Society of Health-System Pharmacists, Inc; 2003.
5. Lexi-Comp [database online]. Hudson, OH: Lexi-Comp, Inc; 2007.
6. Polovich M, White JM, Kelleher LO, eds. *Cancer Chemotherapy and Biotherapy Guidelines and Recommendations for Practice*. 2nd ed. Pittsburgh, PA: Oncology Nursing Society; 2005.
7. Camp-Sorrell D. Developing extravasation protocols and monitoring outcomes. *J Intravenous Nursing*. 1998; 21(4):232-239.
8. Thomas, Juliana. Letter. New York, NY: Sanofi-Aventis; 2005 Sept 26.
9. Mouridsen HT, Langer SW, et al. Treatment of anthracycline extravasation with Savene (dexrazoxane): results from two prospective clinical multicentre studies. *Annals of Oncology*, 2007; 18: 546-550.



Documentation recommendations reproduced/adapted from: Mullin S, Beckwith MC, Tyler LS. Prevention and management of antineoplastic extravasation injury. *Hospital Pharmacy*. 2000; 35:57-76.

SUGGESTED INFORMATION FOR DOCUMENTATION (MAY VARY BASED ON PATIENT CARE SETTING)

SUGGESTED PATIENT EDUCATION

Drug and infusion information
Drug name, dose, volume, and concentration
Amount of extravasated drug
Total amount of drug infused
Other agents administered and the sequence of administration
Method of IV administration (e.g., push, drip)
Location of venous access
Type of venous access device (e.g., central, peripheral)
Needle size and type
Extravasation site, size, and color description (may delineate infiltrated area on patient's skin with felt-tip marker)
Patient complaints or statements at the time of vesicant or irritant infusion

Interventions
Describe the physical measures used to prevent further extravasation.
Note physician contacted.
Note the name, dose, and route of antidotes.
Describe use of warm or cold compresses.
Describe the site.
Consult wound team
Note surgical or other medical consultations requested.
Ensure that the patient has follow-up appointment.
Pain management follow up & reassessment.

Provide instructions.
Ensure that the patient is able to obtain follow-up care and evaluation.
Describe the care of the site: elevate arm; use warm or cold compresses; protect from sun or abrasion; do not immerse in water.
Instruct patient to call provider for any of the following: increased pain, skin color change, increased edema or swelling, stiffness in the extremity, skin breakdown, fever, any additional questions.

ANTIDOTE PREPARATION AND ADMINISTRATION INSTRUCTIONS

Hyaluronidase (Amphadase [bovine])²

Preparation: Use solution as provided (150 unit/1 mL vial); do not dilute further. Inject subcutaneously or intradermally into the extravasation site using a 25-gauge needle or smaller. Dosage: The dose is 150 units (1 mL) given as five 0.2 mL injections into the extravasation site at the leading edge; change the needle after each injection.

Phentolamine (Regitine)^{2,5}

Prepare by diluting 5 mg phentolamine in 10 mL of 0.9% sodium chloride. Inject subcutaneously into the extravasation area within 12 hours of extravasation. Blanching should reverse immediately; additional injections may be required if blanching returns. Do not exceed 0.1-0.2 mg/kg or 5 mg total.

Sodium Thiosulfate⁵

Mix 4 mL of sodium thiosulfate 10% with 6 mL sterile water for injection to prepare a 0.17 mol/L (4%) solution. Inject 3-10 mL subcutaneously into extravasation site; use clinical judgment and size of extravasation site to determine volume. This dosing is based on limited and varied information.

Dexrazoxane⁹

Mix each 500mg vial with 50mL of diluent (provided by manufacturer); mixed solution should be further diluted in 1000mL NS and begin administration within 4 hours. Infuse over 1 to 2 hours in a large caliber vein in an extremity/area other than the one affected by the extravasation. Cooling procedures such as ice packs should be removed from the area at least 15 minutes before administration in order to allow sufficient blood flow to the area of extravasation. ADULT Dose: 1000mg/m² (maximum 2000mg) on Days 1 and 2, 500mg/m² (maximum 1000mg) on day 3. Adjust dose for renal impairment.

AAGBI Safety Guideline

Management of Severe Local Anaesthetic Toxicity



1 Recognition	<p>Signs of severe toxicity:</p> <ul style="list-style-type: none">• Sudden alteration in mental status, severe agitation or loss of consciousness, with or without tonic-clonic convulsions• Cardiovascular collapse: sinus bradycardia, conduction blocks, asystole and ventricular tachyarrhythmias may all occur• Local anaesthetic (LA) toxicity may occur some time after an initial injection
2 Immediate management	<ul style="list-style-type: none">• Stop injecting the LA• Call for help• Maintain the airway and, if necessary, secure it with a tracheal tube• Give 100% oxygen and ensure adequate lung ventilation (hyperventilation may help by increasing plasma pH in the presence of metabolic acidosis)• Confirm or establish intravenous access• Control seizures: give a benzodiazepine, thiopental or propofol in small incremental doses• Assess cardiovascular status throughout• Consider drawing blood for analysis, but do not delay definitive treatment to do this
3 Treatment	<p>IN CIRCULATORY ARREST</p> <ul style="list-style-type: none">• Start cardiopulmonary resuscitation (CPR) using standard protocols• Manage arrhythmias using the same protocols, recognising that arrhythmias may be very refractory to treatment• Consider the use of cardiopulmonary bypass if available <p>GIVE INTRAVENOUS LIPID EMULSION (following the regimen overleaf)</p> <ul style="list-style-type: none">• Continue CPR throughout treatment with lipid emulsion• Recovery from LA-induced cardiac arrest may take >1 h• Propofol is not a suitable substitute for lipid emulsion• Lidocaine should not be used as an anti-arrhythmic therapy <p>WITHOUT CIRCULATORY ARREST</p> <p>Use conventional therapies to treat:</p> <ul style="list-style-type: none">• hypotension,• bradycardia,• tachyarrhythmia <p>CONSIDER INTRAVENOUS LIPID EMULSION (following the regimen overleaf)</p> <ul style="list-style-type: none">• Propofol is not a suitable substitute for lipid emulsion• Lidocaine should not be used as an anti-arrhythmic therapy
4 Follow-up	<ul style="list-style-type: none">• Arrange safe transfer to a clinical area with appropriate equipment and suitable staff until sustained recovery is achieved• Exclude pancreatitis by regular clinical review, including daily amylase or lipase assays for two days• Report cases as follows:<ul style="list-style-type: none">in the United Kingdom to the National Patient Safety Agency (via www.npsa.nhs.uk)in the Republic of Ireland to the Irish Medicines Board (via www.imb.ie)If Lipid has been given, please also report its use to the international registry at www.lipidregistry.org. Details may also be posted at www.lipidrescue.org

Your nearest bag of Lipid Emulsion is kept

This guideline is not a standard of medical care. The ultimate judgement with regard to a particular clinical procedure or treatment plan must be made by the clinician in the light of the clinical data presented and the diagnostic and treatment options available.

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IMMEDIATELY

Give an initial intravenous bolus injection of 20% lipid emulsion $1.5 \text{ ml}.\text{kg}^{-1}$ over 1 min

AND

Start an intravenous infusion of 20% lipid emulsion at $15 \text{ ml}.\text{kg}^{-1}.\text{h}^{-1}$

AFTER 5 MIN

Give a **maximum of two** repeat boluses (same dose) if:

- cardiovascular stability has not been restored **or**
- an adequate circulation deteriorates

Leave **5 min** between boluses

A maximum of **three** boluses can be given (including the initial bolus)

AND

Continue infusion at same rate, but:

Double the rate to $30 \text{ ml}.\text{kg}^{-1}.\text{h}^{-1}$ at any time after 5 min, if:

- cardiovascular stability has not been restored **or**
- an adequate circulation deteriorates

Continue infusion until stable and adequate circulation restored or maximum dose of lipid emulsion given

Do not exceed a maximum cumulative dose of $12 \text{ ml}.\text{kg}^{-1}$

An approximate dose regimen for a 70-kg patient would be as follows:

IMMEDIATELY

Give an initial intravenous bolus injection of 20% lipid emulsion 100 ml over 1 min

AND

Start an intravenous infusion of 20% lipid emulsion at $1000 \text{ ml}.\text{h}^{-1}$

AFTER 5 MIN

Give a **maximum of two** repeat boluses of 100 ml

AND

Continue infusion at same rate but **double rate to $2000 \text{ ml}.\text{h}^{-1}$** if indicated at any time

Do not exceed a maximum cumulative dose of 840 ml



This AAGBI Safety Guideline was produced by a Working Party that comprised:
Grant Cave, Will Harrop-Griffiths (Chair), Martyn Harvey, Tim Meek, John Picard, Tim Short and Guy Weinberg.

This Safety Guideline is endorsed by the Australian and New Zealand College of Anaesthetists (ANZCA).

Malignant Hyperthermia Crisis



AAGBI Safety Guideline

Successful management of malignant hyperthermia depends upon early diagnosis and treatment; onset can be within minutes of induction or may be insidious. The standard operating procedure below is intended to ease the burden of managing this rare but life threatening emergency.

1 Recognition	<ul style="list-style-type: none">• Unexplained increase in ETCO₂ AND• Unexplained tachycardia AND• Unexplained increase in oxygen requirement (Previous uneventful anaesthesia does not rule out MH)• Temperature changes are a late sign	
2 Immediate management	<ul style="list-style-type: none">• STOP all trigger agents• CALL FOR HELP. Allocate specific tasks (action plan in MH kit)• Install clean breathing system and HYPERVENTILATE with 100% O₂ high flow• Maintain anaesthesia with intravenous agent• ABANDON/FINISH surgery as soon as possible• Muscle relaxation with non-depolarising neuromuscular blocking drug	
3 Monitoring & treatment	<ul style="list-style-type: none">• Give dantrolene• Initiate active cooling avoiding vasoconstriction• TREAT:<ul style="list-style-type: none">• Hyperkalaemia: calcium chloride, glucose/insulin, NaHCO₃⁻• Arrhythmias: magnesium/amiodarone/metoprolol AVOID calcium channel blockers - interaction with dantrolene• Metabolic acidosis: hyperventilate, NaHCO₃⁻• Myoglobinaemia: forced alkaline diuresis (mannitol/furosemide + NaHCO₃⁻); may require renal replacement therapy later• DIC: FFP, cryoprecipitate, platelets• Check plasma CK as soon as able	<p>DANTROLENE 2.5mg/kg immediate iv bolus. Repeat 1mg/kg boluses as required to max 10mg/kg</p> <p>For a 70kg adult</p> <ul style="list-style-type: none">• Initial bolus: 9 vials dantrolene 20mg (each vial mixed with 60ml sterile water)• Further boluses of 4 vials dantrolene 20mg repeated up to 7 times. <p>Continuous monitoring Core & peripheral temperature ETCO₂ SpO₂ ECG Invasive blood pressure CVP</p> <p>Repeated bloods ABG U&Es (potassium) FBC (haematocrit/platelets) Coagulation</p>
4 Follow-up	<ul style="list-style-type: none">• Continue monitoring on ICU, repeat dantrolene as necessary• Monitor for acute kidney injury and compartment syndrome• Repeat CK• Consider alternative diagnoses (sepsis, phaeochromocytoma, thyroid storm, myopathy)• Counsel patient & family members• Refer to MH unit (see contact details below)	

The UK MH Investigation Unit, Academic Unit of Anaesthesia, Clinical Sciences Building, Leeds Teaching Hospitals NHS Trust, Leeds LS9 7TF. Direct line: 0113 206 5270. Fax: 0113 206 4140. Emergency Hotline: 07947 609601 (usually available outside office hours). Alternatively, contact Prof P Hopkins, Dr E Watkins or Dr P Gupta through hospital switchboard: 0113 243 3144.

Your nearest MH kit is stored

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