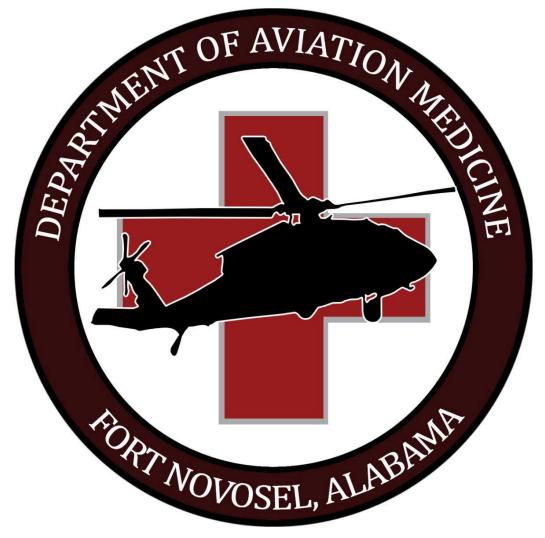
U.S. ARMY AEROMEDICAL EVACUATION STANDARD MEDICAL OPERATING GUIDELINES (SMOG)



CY24 Version

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INTRODUCTION

The CY 2024 SMOG release marks the beginning of current format (STANDARD MEDICAL OPERATING GUIDELINES & SUPPLEMENTAL HANDBOOK). The Aeromedical Evacuation community provided developmental feedback leading to a redesign of current written medical guidance and/or policy. All changes are a result of collaboration between Emergency Medicine professionals, experienced Flight Paramedics, Aeromedical Physician Assistants, Critical Care Nurses, and Flight Surgeons across the Department of Defense (DoD). There is close coordination in the development of these guidelines with the Joint Trauma System, and the Defense Committees on Trauma. Our shared goal is to ensure the highest quality enroute care possible and to standardize care across all evacuation/emergency medical pre-hospital units. It is our vision that all these enhancements will advance enroute care across the services and the Department of Defense.

Unit Medical Trainers, Medical Standardization Instructors, Medical Flight Instructors and Medical Directors will evaluate Critical Care Flight Paramedics' (CCFP), Enroute Critical Care Nurses' (ECCN), Aeromedical Physician Assistants' (APA), and Flight Surgeons' ability to follow and execute the medical instructions herein. These medical guidelines are intended for CCFPs and prehospital professionals who manage emergencies and treat patients in both garrison, humanitarian, and combat theater environments IAW the Aircrew Training Manual Task 2120. Unit medical providers are expected to adjust these guidelines to fit their unit's mission and medical air crews' training/experience. Medical directors or designated supervising physicians will endorse these guidelines upon appropriate adjustment. They will also manage individual unit medical missions within their Critical Care Flight Paramedics, Enroute Critical Care Nurses, and advanced practice aeromedical providers' scope of practice. CCFPs should administer medications as listed in these guidelines unless their medical director and/or supervising physician orders deviation. Other medications may be added, so long as the unit supervising physician and/or medical director approves them.

This manual also serves as a reference for physicians providing medical direction and clinical oversight to medical personnel. Treatment direction, which is more appropriate to the patient's condition than the guideline, should be provided by the physician so long as the medical personnel's scope of practice is not exceeded.

Any medical guideline that is out of date or has been found to cause further harm will be updated or removed immediately. The Department of Aviation Medicine (DAM) serves as the managing editor of the SMOG and is responsible for content updates, managing the formal review process, and identifying SMOG Charter members for annual review.

The Standard Medical Operating Guidelines and Supplemental Handbook provide medical procedural guidance and is in compliment to other Department of Defense and Department of the Army policies, regulations, and doctrinal guidance. Nothing herein overrides or supersedes laws, rules, regulation, or policies of the United States, DoD, or DA.

MEDICAL DIRECTOR / UNIT COMMANDER REVIEW AND APPROVAL PAGE

The Standard Medical Operating Guideline and Supplemental Handbook specify standard medical treatment guidelines to be used by all Flight Paramedics and Medical Providers performing medical care while serving in this unit in any environment. It is a guideline and not a comprehensive patient care manual.

This SMOG, Supplemental Handbook, and any attached adjustments are hereby established as standard guidelines and protocol for the following unit:

| Date of Certification and Approval by all of the | below: |
|---|--|
| <u>Unit Trainer Review:</u> | |
| This document has been reviewed by the belo | w noted individuals for correctness and mission applicability: |
| Unit Standardization Officer/NCO Signature: _ | Date: |
| Unit Training NCO Signature: | Date: |
| Authorization: | |
| The Standard Medical Operating Guideline approved for use by the undersigned. | and Supplemental Handbook have been reviewed and |
| Medical Director/Supervising Physician* | |
| Name: | - |
| Signature of Approval: | Date: |
| Unit Commander | |
| Name: | - |
| Signature of Approval: | Date: |

*Additional Medical Director comments/addenda can be attached and should contain counter signature of Unit Commander to be valid.

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Standard Medical Operating Guidelines are found at the following website:

https://www.milsuite.mil/book/groups/department-school-of-army-aviationmedicine

> Also available, along with all fillable evacuation forms and AARs on the Joint Trauma System website: <u>https://jts.health.mil/index.cfm/PI_CPGs/cpgs</u>

https://jts.health.mil/assets/docs/forms/DA4700_OP5_JTS_TACEVACAAR&PC R.pdf

All comments and/or recommendations should be sent to: <u>medcoesaamoperations@army.mil</u> with the subject line "CCFP-SMOG"

UNIVERSAL PATIENT CARE

Patient History **Key Concepts** Age of the Patient **Use MARCHES for Trauma Patients** Chief Complaint Massive Bleeding Control 0 • Airway Timing of Event / Event Factors 0 Respiratory Other Symptoms or Complaints 0 Circulation Patient's Past Medical History 0 • Hypothermia Care 0 Other Pertinent SAMPLE, OPQRST Questions Eye Injuries 0 **Spinal Motion Restriction** 0 Focused Primary Exam for Non-Traumatic Illness/Injury **Treatment/Actions** Scene Safety . o Maintain situational awareness • Utilize appropriate PPE Initial Assessment • Treat obvious and emergent life threats • MARCHES or Focused Primary Exam Utilize BLS, ALS, and/or PALS guides as necessary **Consider Spinal Immobilization** • Dangerous MOI Low risk MOI but unable to rotate neck 45° o Does not apply to situations where imminent danger exists Record vital signs and make appropriate transport decision Initial Interventions • Supplemental O2 • IV/IO (Saline Lock) as applicable Medication/fluid administration (as indicated) Secondary Assessment • 12 Lead EKG (as applicable) • ETCO2 (as applicable) Secondary interventions • Pain management Notes, Warnings, Cautions General supportive measures include airway/respiratory support, continuous hemodynamic monitoring with SPO2 and ETCO2 as appropriate, supplemental O2 PRN, IV fluid boluses, pain control PRN All patients should have complete vital signs recorded All patient encounters should be recorded on appropriate care documentation sheets per theater • policies, unit SOPs and/or in accordance with JTS Documentation CPG at end of a patient encounter. Any mishaps/errors should be brought to attention of the medical control ASAP. Contact medical control for any necessary assistance when feasible. Consider spinal immobilization if: Fall from height (versus fall from standing) 0 • Axial load to head o High speed collision, rollover, or ejection from any motorized vehicle • Explosion or blast injury Trauma resulting in temporary amnesia/loss of consciousness 0

TACTICAL EVACUATION

| Ground "Pick-Up" PhaseGround "Pick-Up" Phase• Attempt to gain info prior to landing• Goal on ground time < 5min prior to wheels up• Ensure 360-degree scene security• If the tactical situation permits, all known preventable causes of death should be addressed prior to casualty movement |
|--|
| Triage casualties Treat all preventable causes of death IAW TCCC Package and secure patients for transport Brief and guide litter teams to aircraft Load and secure patients |
| "In-Flight" Phase |
| Triage casualties as required: reassess patients and interventions Hemorrhage Control Check/add tourniquet, pack/dress wound, pressure dressing, hemostatic dressing Initiate blood (DCR) Airway/Vent Management Reposition airway, nasopharyngeal airway, RSI (intubation/BIAD), cricothyroidotomy Target SPO2 90-96% Chest Trauma Vented occlusive dressing, needle thoracostomy, finger thoracostomy, chest tube Hypothermia Management Head Injury/Altered Mental Status Monitor and treat for signs and symptoms of ICP (elevate head, 3% hypertonic saline, target ETCO2) Pain Management Consider Antibiotic Therapy |
| Document Care |
| Notes, Warnings, Cautions Damage Control Resuscitation (DCR) Order of Precedence: |
| Control hemorrhage if able Administer blood products Consider TXA 2g < 3hrs from injury Calcium administration during or after 1st unit and after every 4th unit of blood (Calcium may be given before TXA) Consider pressors (as a last resort) Replace any limb tourniquets placed over the uniform with one applied directly to the skin, 2-3 inches above the wound Maximize blood/fluid therapy prior to considering pressor administration At any time, if patient becomes pulseless and apneic go to trauma arrest protocol |
| If tactical situation allows, load deceased patients on a separate transport Consider full PSL prior to advanced airway managements to provent aspiration |
| Consider full RSI prior to advanced airway managements to prevent aspiration |

AIRWAY (ADULT/PEDIATRIC)

Signs and Symptoms of Distress and/or Failure

- SPO2 decreasing <90%, with/without supporting signs/symptoms of:
 - Tachypnea, Tachycardia, Fever, Cough, Adventitious Breath Sounds, or Shock
- Difficulty Breathing or Excess Work of Breathing as demonstrated by:
 - Pursing of Lips, Accessory Muscle Involvement, Cyanosis, Dysphasia, Diaphoresis
- Airway Obstruction due to Trauma, Edema, Excess Secretions, Foreign Body, or Tongue
- Apnea
- Decreased LOC (GCS<8)
- Pediatric patient is defined as <12 years of age.

Procedures

- Reposition airway via Jaw Thrust or Head Tilt Chin Lift. Provide shoulder padding for PEDs if required.
 Sweep (NOT BLIND) and suction as needed
 - ABD thrust or back slaps (for infants) if indicated
- Assess the need for an advanced airway (GCS <8, suspected deterioration, SpO2 < 90%, TBSA >40%, severe head injury)
- SpO2<90%
 - o Start Supplemental O2
 - Place NPA/OPA prn if no contraindications
 - Recheck q5 minutes
 - o BVM or assist with respiration prn
- Consider direct Laryngoscopy to visualize foreign body obstruction; If present remove, suction, and/or provide abdominal compressions or back slaps for pediatric patients
- Establish an advanced airway per Procedure in the following sequence (Move to the next procedure per individual competencies, contraindications, and/or attempt failures):
 - Endotracheal Intubation
 - Blind Insertion Airway Device (BIAD)
 - Cricothyroidotomy
- After failed attempt:
 - Reassess Interventions
 - Restart Protocol
 - Consider other causes
- Continuous monitoring of ETCO2, SPO2, and ventilatory waveforms and pressures
 - Repeat sedative, analgesic, and paralytic per dose and time guidelines

• Advance to FAILED AIRWAY GUIDELINE if:

- Unable to adequately open the airway
- After two (2) failed attempts by the most proficient provider on the scene to place an ET Tube and at least one (1) failed attempt with a supraglottic airway under PAI
- o Intubation contraindicated due to anatomical abnormalities or major airway trauma
- \circ Continued inability to ventilate patient with mask ventilation using a BVM

Failed Airway

- If able to ventilate with a BVM, insert NPA or OPA dependent on contraindication and continue ventilating with BVM
- If unable to ventilate, perform an age-appropriate cricothyroidotomy (>10 years of age)
- Ventilate patient per age-appropriate respiratory rate to maintain minute ventilation

| RAPID SEQUEN | CE INTUBATION |
|--|--|
| <u>History</u> Airway Compromise or Inability to Protect Airway Respiratory Failure (Hypoxic, Hypercapnic) >40% TBSA Burns, Severe Sepsis, TBI with AMS, etc. Patient or Crew Safety Combative, prolong transfer in critically sick, etc. High likelihood of failure (Distorted Anatomy) Penetrating neck trauma <u>Procedure</u> Make a plan, prepare patient and equipment (See PRE-INTUBATION CHECLIST) Conduct seven "P" pneumonic (7Ps): | Medications Induction Agents: Ketamine 1-2 mg/kg IV Hidazolam 0.2-0.4 mg/kg IV Midazolam 0.1 mg/kg IV Propofol 1-2.5 mg/kg IV Paralytics: Rocuronium 0.6-1.2 mg/kg IV Vecuronium 0.08-0.15 mg/kg IV Succinylcholine 1-1.5 mg/kg IV Ketamine 0.5-2 mg/kg IVP or 0.5-2 mg/kg bolus then 1-3mg/kg/hr. Propofol 10-75 mcg/kg/min Midazolam 0.05 mg/kg IVP or 0.05 mg/kg bolus then 0.05-0.1mg/kg/hr. Push Dose Epi: Epinephrine 5-20mcg IV q2-5min |
| Prepare | |

- Suction: available, check for function
- Oxygen: Pre-Oxygenation + Apneic Oxygenation
- Airways: ETT, SGA (iGel, King, etc.), Cricothyrotomy
- Pharmacology: Induction, Paralysis, Post-intubation Sedation
- Monitor: BP, HR, RR, SpO2%, etCO2 capnography, 4-lead
- Equipment: Bougie, Laryngoscope, Video Laryngoscope, Cric Kit
- Evaluate cricothyrotomy landmarks and assess procedural difficulty

Pre-Oxygen

 Preoxygenate / Denitrogenate ≥ 3 minutes or 8 Vital Capacity Breaths with 15 LPM NRB or BVM + PEEP, and NC 4-6 LPM, Oxygenated ≥ 94% if able.

Positioning

• 30° Head-up for Pre-Oxygenation, Ear-to-Sternal Notch for Intubation, C-Spine Consideration.

Pretreat

- Resuscitate with IVF or Blood Products. Consider Push-Dose Pressors (Epi) to ensure SBP>100mmHg.
- 3 5 Minute prior to Sedative / Paralytic
- Consider Fentanyl 3mcg/kg slow IV push to prevent Hypertension in head injury, cardiac ischemia, or aortic dissection
- Atropine 0.02 mg/kg IV to prevent bradycardia in Peds (age <1y)

Sedate/Paralyze

- Push sedative first before paralytic push (see medications above)
- Apneic Oxygenate
- Monitor SpO2% and wait for adequate paralysis.

Pass Tube

• Visualize Cords, Pass Tube, Inflate Bulb and Begin Bagging

Post-Tube Management

- Verify Tube Place with etCO2 waveform capnography and secure tube.
- Place patient on Post-intubation Maintenance Sedation (see medications above)

VENTILATOR MANAGEMENT

Clinical Indications

- Patient received from transferring facility, intubated, and requires ventilator support.
- Patient requiring intubation in the field and subsequent respiratory support.

Contraindications

Equipment malfunction / failure

Procedure

- Turn on ventilator and ensure that machine is functional, and battery is charged.
- Attached ventilator tubing and O2 tubing to machine.
- If patient is a transfer and already on a ventilator, maintain ventilator setting from medical treatment facility.

If Patient "Newly" on the Ventilator, Initial Setting Should Be

- Mode: CMV+ or Assist Control (AC)
- Tidal Volume (Vt): 6 cc/kg IBW
 - o IBW calculation
 - MEN: [(Height in inches 60) x 2.3] + 50
 - WOMEN: [(Height in inches 60) x 2.3] + 45.5
 - Tidal Volume should not be altered to fix ventilation, adjust rate instead for increased or decreased minute volumes. Vt only gets changed for lung protection (i.e. to prevent barotrauma / volutrauma)
 - \circ Reduce Vt by 1mL/kg at intervals \leq 2 hours until Vt = 6 cc/kg IBW
- Rate (RR): Initially 14, adjust based on CO2 (if CO2 >45mmHg) and ventilatory needs (do not exceed >35 BPM)
- **I:E:** 1:2 (Patients with obstructive lung diseases should have increased I:E around 1:4 or 1:5; if rate >20 (most children) will need to titrate iTime down to achieve appropriate I:E ratio)
- FiO2: 100% (then titrate FiO2 down to achieve SPO2 90-96%, SPO2>93% head injury)
- **PEEP:** 5

*NOTE: FiO2/PEEP (Should be adjusted in concert per the chart below if patient has ARDS or if desaturation is gradual and presumed to be caused by patient pathology)

To Achieve Oxygenation Goals, set the FiO2 to 30% and start titration FiO2 and PEEP collectively based on the chart. Go up every 5-10 minutes; quicker if low SpO2 sats develop.

Lower PEEP/higher FiO2

| FiO ₂ | 0.3 | 0.4 | 0.4 | 0.5 | 0.5 | 0.6 | 0.7 | 0.7 |
|------------------|-----|-----|-----|-----|-----|-------|-----|-----|
| PEEP | 5 | 5 | 8 | 8 | 10 | 10 | 10 | 12 |
| | | | | | | | | |
| FiO ₂ | 0.7 | 0.8 | 0.9 | 0.9 | 0.9 | 1.0 | | |
| PEEP | 14 | 14 | 14 | 16 | 18 | 18-24 | | |

*NOTE: Hypotensive patients (MAP <70 or SBP <90) may respond negatively to increased PEEP due to decreased venous return. Monitor for increasing hypotension and tachycardia.

Alternate Higher PEEP settings:

Higher PEEP/lower FiO2

| FiO ₂ | 0.3 | 0.3 | 0.3 | | 0.3 | | 0.3 | (|).4 | 0.4 | 0.5 |
|------------------|-----|-------|-----|---|-----|---|-----|----|-----|-----|-----|
| PEEP | 5 | 8 | 10 | | 12 | | 14 | | 14 | 16 | 16 |
| | | | | | | | | | | | |
| FiO ₂ | 0.5 | 0.5-0 | .8 | 0 | .8 | 0 | .9 | 1. | 0 | 1.0 | |
| PEEP | 18 | 20 | | 2 | 2 | 2 | 2 | 22 | | 24 | |

Oxygenation Goal: <u>Normal</u> PaO2 80-100 mmHg or SpO2 90-96%; <u>ARDS</u> PaO2 55-80 mmHg or SpO2 88-95% Plateau Pressure Goal: ≤ 30 cm H2O

- Check Pplat (0.5 second inspiratory pause), at least q 4h and after each change in PEEP or VT.
- If Pplat > 30 cm H2O: decrease VTby 1ml/kg steps (minimum = 4 ml/kg).
 - If Pplat < 25 cm H2Oand VT< 6 ml/kg, increase VT by 1 ml/kg until Pplat > 25 cm H2O or VT = 6 ml/kg.
- If Pplat < 30 and breath stacking, or dys-synchrony occurs: may increase VT in 1ml/kg increments to 7 or 8 ml/kg if
 Pplat remains < 30 cm H2O.

Alarm Settings:

- High Pressure Alarm: 10 cmH2O above peak airway pressure.
- Low Pressure Alarm: 5 cmH20 below peak airway pressure.

OR

- High Pressure Alarm: 50% above the baseline PIP (1.5 x current PIP)
- Low Pressure Alarm: 50% below the baseline PIP (0.5 x current PIP)

Pressures will be determined by placing patient on ventilator for ~ 1-2 minutes and determining intrinsic peak inspiratory pressure. (Labeled as PEAK on 754 Ventilator (top right); Labeled as Ppeak on Hamilton T1 (top left)

Monitor waveform on machine and patient to ensure no breath stacking occurs. If this occurs, a high-pressure alarm may sound. However, if breath stacking suspected even in absence of alarm – disconnect tubing and allow exhalation. Increase I:E.

Troubleshooting: Airway Compromise or Lost Airway In-Flight

- If at any time patient begins to desaturate or develop respiratory problems, immediately disconnect ventilator, and ventilate patient with BVM (with PEEP valve if available) and 100% O2 while correcting issues utilizing the D.O.P.E. algorithm:
- Displacement: ETT in place, patient not extubated/ tube did not move during transfer. If advanced pull back to
 original length and attempt to bag; if tube has pulled farther out of trachea, DO NOT ATTEMPT TO ADVANCE IT
 without placement of bougie to verify tracheal placement. When advancing bougie, feel for tracheal rings or carina
 stop. If in doubt, pull tube and attempt BVM. If this fixes problem, continue to bag patient. Upon stabilization,
 consider alternative advanced airways (extraglotic airway or cric)

**If ETT moves freely, assess for ETT bulb rupture. **

- Obstructions: Assess for secretions in ETT. Suction if indicated
- Pressure: Ensure that a tension pneumothorax / hemothorax has not developed (if chest tube in place, ensure it is functioning/ not kinked or clamped). If tension pneumothorax / hemothorax suspected, perform immediate needle thoracostomy. Assess the need for escharotomy if circumferential burn. Consider additional paralysis and sedation if patient does not tolerate ventilation
- **Equipment:** Ensure that vent did not fail; O2 tank not empty. If ventilator is operational, trace all tubes to the patient connection (airway tube, transducer line, exhalation line) ensuring patency and connections
- High Pressure Alarms / Peak Airway Pressure Alarms (Peak pressure >35 cm H2O): Correct problems
 causing increased airway resistance and decreased lung compliance, including pneumothorax or pulmonary
 edema. Check ventilator to make sure prescribed tidal volume is being delivered. Check for linked/crushed tubing.
- Air Leaks Causing Low Pressure Alarms / Volume Loss: Assess, correct air leaks in endotracheal tube, tracheostomy cuff, ventilator system; recheck ventilator to make sure prescribed tidal volume is delivered
- Ventilator Desynchrony: Agitation and respiratory distress that develop in a patient on a mechanical ventilator who has previously appeared comfortable represents an important clinical circumstance that requires a thorough assessment and an organized approach. The patient should not always be automatically re-sedated but must instead be evaluated for several potentially life-threatening developments that can present in this fashion
- Lung Hyperinflation Air Trapping and Auto-PEEP: Dynamic hyperinflation is associated with positive endexpiratory alveolar pressure, or auto-PEEP. The physiologic effects include decreased cardiac preload because of diminished venous return into the chest. The reduced cardiac output that results from the reduction in preload can lead to hypotension and, if severe, to Pulseless Electrical Activity and cardiac arrest. Dynamic hyperinflation can also lead to local alveolar overdistention and rupture. Prevent, manage lung hyperinflation by decreasing tidal volume, changing inspiratory and expiratory phase parameters, switching to another mode, and correcting physiological abnormalities that increase airway resistance
- Document Procedure, Results, and Vital Signs

Ventilator Transfer Procedure

- 1. Ensure endotracheal tube is secure, document size and position of ETT at the teeth. Clamp tube immediately before disconnecting patient from vent to maintain PEEP if conducting recruitment maneuvers or PEEP is 10 or higher, then un-clamp only after connected to new vent circuit.
- Ventilator settings should be coordinated with the transferring physician, anesthesia provider or respiratory therapist. Verify settings, review arterial blood gas (ABG) analysis, and current SPO2 and ETCO2 readings. Place those setting on transport vent and place patient on transport vent early to verify patient tolerance and compatibility.
- 3. ABG should be done within 30 minutes of flight. If time allows, patient should be on transport ventilator for at least 15 minutes prior to transport.
- 4. Ventilator settings for en-route care team should initially be matched to those of the transferring facility. Adjust settings PRN in order to maintain appropriate clinical parameters listed on first page of ventilator management protocol or transferring physician orders.
- 5. Ensure adequate sedation and analgesia medications are on hand.

CPG ID References: 92 (Mechanical Ventilation Basics); 48 (Mechanical Ventilation During Critical Care Air Transport)

BLOOD AND COMPONENT USE

IMMEDIATE INDICATIONS in Trauma Patients with SERIOUS INJURY

- Systolic BP < 100 or absent radial pulse
- Tachycardia > 100
- Amputation

Clinical Indications

- Uncontrolled hemorrhage
- Evidence of hemorrhagic shock
- Trauma patients with amputations (complete or partial with distal circulation compromise)
- Non-compressible penetrating thoracic region
- Abdominal/transitional zone injuries (significant mechanism of injury)
- Clinical signs of coagulopathy (tachycardia, tachypnea, fever, altered mentation, hypoxemia)
- Severe hypothermia associated with blood loss

<u>Treatment</u>

 Maximize hemorrhage control, treatment of suspected tension pneumothorax, patent airway or airway control, IV/IO access, hypothermia prevented and/or treated

Blood Product Order of Precedence

- Whole Blood
- Plasma, RBCs, Platelets in a 1:1:1 Ratio (no particular order)
- Plasma and RBCs in 1:1 Ratio
- Plasma (thawed, liquid, reconstituted) alone or RBCs alone
- Document all items on the SF 518 (only authorized document for blood products aboard Army Aeromedical Evacuation platforms)
 - Two-person verification of patient and blood products given matching SF 518
- Examine units of blood (look for gas, discoloration, clots, and sediment) and verify that the Safe-T-Vue is white
- Initiate large bore IV (18G min, 14G preferred) or IO access (Lidocaine 2% (2-3mL) flush in IO site provides analgesia and increases compliance)
- Blood and blood products must be administered through Y-tubing with filter, flushed with NS prior to use
- Transfuse blood through an approved fluid warming device if available
 - Rapid transfusion can be achieved via pressure bag at least 300mmHg; a 60mL syringe or manual pressure can also be utilized in the event a pressure infuser is not available
- Consider slowing all other concurrent infusions unless they are TXA or RFVIIa
- Resuscitation Goal: until palpable radial pulse, improved mental status, or SBP > 100 (>110 w/ head injury) and MAP >60mmHg
- 30mL of 10% Calcium Gluconate or 10mL of 10% Calcium Chloride IV/IO should be given to patients in hemorrhagic shock during or immediately after transfusion of the first unit of blood product and with ongoing resuscitation after every 4 units of blood products. Ionized calcium should be monitored, and calcium should be administered for ionized calcium levels less than 1mmol/L
- Monitor patient every 5 minutes and document any patient signs and symptoms consistent with transfusion reaction (monitor core temperature)
 - o If a transfusion reaction occurs, see the Transfusion Reaction protocol.

BLOOD TRANSFUSION REACTIONS

<u>Treatment</u>

• STOP THE TRANSFUSION!

If a blood transfusion reaction is suspected

- Apply O2 (if hypoxic), IV/IO, and cardiac monitor
- Establish Advanced Airway per individual competencies, contraindications, and/or attempt failures. Maintain SPO2 >93%

• Anaphylaxis

- Epinephrine 0.3mg IM (0.3mL of 1:1000)
- Diphenhydramine 25mg IV/IO/IM
- Maintain Airway
- Administer IV fluids as needed
- Consider Methylprednisolone 125mg IV/IO
- Acute Hemolytic Reaction (AHTR)
 - Diphenhydramine 25-50mg IV/IO/IM
 - Consider osmotic diuresis:
 - 20g Mannitol 20% or 250mL 3% NaCl
 - Febrile Non-Hemolytic Transfusion Reaction (FNHTR)
 - Consider Acetaminophen 500mg PO or 1G IV
- Continue to reassess the patient and ensure to document on SF 518
- Notify blood bank of all transfusion reactions.

- Reactions are very rare (less than 0.1%)
- GENERAL RULES:
 - Stop the transfusion.
 - Keep the intravenous line open with saline.
 - Identify and treat cause of the reaction.
 - Re-institute the transfusion only if it is deemed to be clinically essential.
- Before initiating IVF bolus, ensure IV tubing is new. DO NOT USE existing Y-tubing from blood administration set.
- The most common transfusion reaction is a febrile, non-hemolytic transfusion reaction. These are mostly benign with no lasting sequelae. Treatment consists of antipyretics.
- TRALI is the leading cause of transfusion-related mortality; a concern in patients who have undergone recent surgery, massive transfusion, or have an active infection. Goal of treatment is supportive: maintain oxygenation, reduced respiratory distress.
- TACO is essentially pulmonary edema secondary to congestive heart failure occurring in elderly, small children, and those with compromised cardiac function. Large volumes of fluid given rapidly is a common precursor. Goal is diuresis and treating underlying condition. Both TACO and TRALI require immediate resuscitation.

IV/IO PROTOCOL

| Assess need for IV Emergent or potentially emergent medical or trauma condition. Peripheral IV x 2 Catheter ≥ 18ga Two failed attempts or > 90 secs proceed to IO. | Intraosseous Device for Life/limb threatening. IO should only be considered first if patient is deemed difficult to gain IV access. If IV/IO access unsuccessful attempt EJ IV Cannulation. | | | | |
|--|---|--|--|--|--|
| Sternal IO Device by precedence: Fast-1[™] EZ T.A.L.O.N[™] EZ-IO[™] Locations for EZ T.A.L.O.N.[™] and EZ-IO[™] by precedence Bilateral Proximal Humerus Bilateral Proximal Tibia Bilateral Distal Tibia | Correct needle size for EZ-IOTM Yellow - 45mm for humerus and *heavy sternal Blue - 25mm for adult *sternum/tib Pink - 15mm for children and *sternal/tib *NOTE: Use of EZ-IO in sternal is off label emergency procedure only | | | | |
| Notes, Warnings, Cautions | | | | | |

GENERAL RULES

- GAIN VASCULAR ACCESS where available based upon patient. 0
- Any pre-hospital fluids or medications approved for IV use may be given through an intraosseous line, including blood products.
- All trauma patients or potentially ill patients should have at least two functioning IV/IO lines whenever possible.
- Upper extremity IV sites are preferable to lower extremity IV sites.
- Pressure infusion bag is recommended for IO starting at 300mmHg.
- o Following IV attempt failure and IO attempt failure, external jugular lines can be attempted for lifethreatening events with no peripheral access.
- Ensure open and functioning fluid bolus per specific protocol. At a minimum, maintain a slow "to-0 keep-open" (TKO) drip.

ABDOMINAL INJURY

Signs and Symptoms

- Altered Mental Status
- Tachycardia
- Absence of palpable pulses
- Pale, moist, mottled skin
- Poor peripheral pulses
- Hypotension
- Hematuria
- Pain, tenderness, distention, dissymmetry
- Absent/diminished bowel sounds
- Grey-Turner sign
- Cullen sign
- Kehr's sign

<u>Treatment</u>

- Blunt Abdominal/Pelvic Injury
 - o Serial Physical Exams/Reassessment
 - Pelvic Binder
 - Conduct FAST exam if possible*
 - Focus on resuscitation
- Penetrating Abdominal/Pelvic Injury
 - o Hemostatic Dressing/ Pack Pelvic Cavity
 - Pressure Dressing
 - Direct and Indirect pressure
 - Abdominal Dressing
 - o Pelvic Binder
 - AAJT- uncontrolled pelvic bleed
- Damage Control Resuscitation
 - Consider implementation of DCR if indications are met (SBP<100) (HR>100) (penetrating chest/abdominal injuries, amputations, pelvic injury)

* FAST exam cannot reliably exclude significant injury but may provide indication of intra-abdominal injury.

- Pregnant patient
 - o Increased risk of Aspiration and gastric acidity
 - Patient should receive max O2 due to increased O2 consumption and depleted reserves
 - Consider warm LR before crystalloids to better restore fetal oxygenation
 - >20 weeks gestation, tilt at least 15 degrees to prevent Vena Cava Syndrome
- Lateral contusions (seatbelt sign) associated with a 20% occurrence of internal injury.
- Presence of pregnant uterus should be determined. Some changes can mimic shock (heart rate can increase by 20 BPM, blood volume increases by 50% during mid-pregnancy, and can experience relative anemia from hemodilution.) Due to the increase in blood flow to the uterus, risk of massive blood loss is greatly increased with trauma to the bony pelvis

BURNS/ELECTRICAL INJURY

| History | Call the Burn Center: DSN 312-429-2876 (429- | | | | | | |
|---|--|--|--|--|--|--|--|
| How long ago was the injury? | BURN)/Comm: 210-916-2876 or 210-222-287 | | | | | | |
| Any signs of airway involvement? | , | | | | | | |
| How big/small of a space was the patient in | STOP the burning process/remove patient from | | | | | | |
| during the incident? (inhalation/carbon | electrical source. Ensure your safety first! | | | | | | |
| monoxide) | | | | | | | |
| • Are there other traumatic injuries associated? | Differential Diagnosis: cardiac arrest, environmental | | | | | | |
| Any spinal immobilization needed? | exposure, seizure, burns (chemical, electrical, | | | | | | |
| (fall from a significant height, blast, etc.) | thermal, radiation), multiple trauma, carbon monoxide | | | | | | |
| | toxicity. | | | | | | |
| Trea | atment | | | | | | |
| | ats (DO NOT overlook TRAUMA), IV/IO, O2, and monitor | | | | | | |
| (electrical injuries must have 12 lead EKG com | | | | | | | |
| Electrical: If arrythmia is present, go to appropriate the second s | · , | | | | | | |
| Remove any constricting items i.e., rings and b | • | | | | | | |
| Assess airway, if suspected airway involvement | | | | | | | |
| | Ide comatose patient, symptomatic inhalation injury, | | | | | | |
| deep facial burns, and burns over 40% Tota | al Body Surface Area (TBSA). Requires large ETT size 8 | | | | | | |
| adult. | | | | | | | |
| • Thermal/Electric burn: If able, remove burnin | g/charred clothing and cover with dry, sterile | | | | | | |
| sheets/dressings. | | | | | | | |
| Chemical burn: Brush off dry chemicals, cut o | ff contaminated clothing, flush area with saline 10-15 | | | | | | |
| minutes. | | | | | | | |
| Eyes: flush with saline for 30 minutes. | | | | | | | |
| Hydrofluoric Acid- After thorough irrigation, | apply a CaGlu gel (75mL KY Jelly + 25mL 10% CaGlu) | | | | | | |
| for 30 minutes. | | | | | | | |
| Tear Gas- rinse skin and eyes with NS. | | | | | | | |
| Alkali Burns to eye- 1-2 L of NS each eye feedback | or 30 minutes. | | | | | | |
| Determine/start fluid replacement for burn fluid | | | | | | | |
| Manage pain and prevent hypothermia. KEEP | WARM! | | | | | | |
| Monitor urinary output, if able | | | | | | | |
| Notes, Warnings, Cautions | | | | | | | |
| Urinary output is the MOST reliable guide for a | | | | | | | |
| Adult 30-50 mL/hr (75-100ml/hr electrical burn) | | | | | | | |
| | ttic airway devices such as laryngeal mask airways | | | | | | |
| (LMAs) to be inadequate. | | | | | | | |
| | OC) are the most common dysrhythmias seen with | | | | | | |
| electrical shock. | | | | | | | |
| | triage should be performed. Those victims in full arrest | | | | | | |
| should be resuscitated first. | | | | | | | |
| | spirations after airway maneuver, but no other signs of | | | | | | |
| non-survivable injury, administer ventilatory | support aggressively as resources allow | | | | | | |

BURN FLUID RESUCITATION Rules of 9's Burn % Estimation Chart 189 ADULT CHILD INFANT ADULT >40KG Burns: 10mL/hr x %TBSA; patients weighing more than 80kg, add 100 ml/hr to IV fluid rate for each 10 kg > 80 kg. Re-evaluate every 1-2 hours. Adjust IV rate to UOP goal 30-50mL (0.5-1 mL/kg in Peds). Adjust IV rate up or down by 20%. High Voltage Injury: 10mL/hr x %TBSA (estimate to nearest 10%); patients weighing more than 80kg, add 100 ml/hr to IV fluid rate for each 10 kg > 80 kg. Re-evaluate every 1-2 hours. Adjust IV rate to UOP goal 75-100mL. Adjust IV rate up or down by 20% PEDIATRIC <40KG Burns: 3 x %TBSA x body weight (kg) gives the volume for initial 24 hrs. One half is given in first 8 hours. Monitor urine output with goal of 0.5 to 1 mL/kg/hr. High Voltage Injury: Adjust IV rate to UOP goal 1-2 mL/kg. Adjust IV rate up or down by 20% Notes, Warnings, Cautions TBSA > 20%, may require acute fluid resuscitation in prehospital: LR (best)>NS (2nd • best)>Hextend (only to 1L) Notes, Warnings, Cautions It is worth your time and effort to accurately estimate burn surface area, ideal body weight, then calculate and administer appropriate fluids while the patient is under your care. Administer a maintenance rate of D5LR to children <13 years of age. Utilize the 4-2-1 rule: 4 ml/kg for

 Administer a maintenance rate of D5LR to children <13 years of age. Utilize the 4-2-1 rule: 4 ml/kg for the first 10 kg + 2 ml/kg 2nd 10 kg + 1 ml/kg over 20 kg. This maintenance rate is in addition to the isotonic infusion calculated for burn resuscitation and is not titrated.

Reference CPG ID:12 (Burn Care)

CHEST TRAUMA

Signs and Symptoms

- Difficulty breathing
- Rapid respirations with SPO2 decreasing or <93% (In flight and on O2)
- Flail chest
- Unequal rise and fall
- Open wound/impalement over the thorax
- Penetrating abdominal wound
- Bruising across chest or base of neck
- Subcutaneous emphysema or deviated trachea

<u>Treatment</u>

- Penetrating Chest Wound
 - o Open
 - Seal open wound with occlusive chest seal (vented)
 - o Impalement
 - Stabilize
 - High index of concern for Hemo-pneumothorax
 - Signs of Hemo-pneumothorax
 - Needle Thoracostomy
 - Goal: SPO2>90%; Improved RR; Equal rise/fall
 - Blunt Chest Trauma
 - o Flail Chest
 - Administer Pain Control
 - Consider Endotracheal Intubation, Pos P ventilation
 - Signs of Hemo-pneumothorax
 - Needle Thoracostomy
 - Goal: SPO2>90%; Improved RR; Equal rise/fall

- Needle Thoracostomy may need to be repeated
- Failure to improve after Needle Thoracostomy
 - o Controlled descent as able
 - Consider Finger/ Tube Thoracostomy
- Consider Spinal Immobilization for Chest Trauma Patients
- Maintain high index of suspicion for Intra-abdominal and retro-peritoneal bleeding in all chest injuries

CRUSH SYNDROME

<u>History</u>

- Entrapped extremity (as little as 1hr)
- Erythema, ecchymosis, abrasion
- Swelling, tense muscle compartment

History (Complications)

- Hyperkalemia
- Hypocalcemia
- Compartment Syndrome
- Rhabdomyolysis
- Arrhythmia
- Hypotension

<u>Treatment</u>

- Prior to Extraction
 - Consider tourniquet placement for crush injuries if the length of entrapment exceeds 2 hours and crush injury protocol cannot be initiated immediately.
 - Apply two tourniquets side by side and proximal to the site of entrapment immediately prior to extraction.
- Initiate Crush Injury Protocol before extrication if possible and before loosening tourniquets (if tourniquet conversion indicated).
- IV / IO Guideline
 - Initiate aggressive fluid administration of IV / IO crystalloids 2L initial bolus; followed by infusion rate: 1L/hr. Adjust to urine output (UOP) goal of >100-200mL/hr. (via Foley or improvised graduated cylinder)
- Monitor for life-threatening hyperkalemia (PVC's, bradycardia, peaked T-waves, decreased peripheral pulse strength, hypotension).
 - If PVCs become more frequent, the patient develops bradycardia, peripheral pulse strength decrease, or potassium levels are >5.5 mEq/L or rising, treat urgently for hyperkalemia.
 - Calcium: Administer 10 mL (10%) calcium gluconate or calcium chloride IV over 2–3 minutes.
 - Insulin and Glucose: Give 10 units of regular insulin followed immediately by 50mL of D50.Titrate PRN
 - Albuterol: Administer 12mL of albuterol sulfate inhalation solution, 0.083%(2.5mg/3mL) in nebulizer.
- If no signs of hyperkalemia develop, continue fluid administration and continuously monitor.

- Crush syndrome can occur in as little as 1 hour of entrapment
- Tourniquets may mitigate life-threatening complications in situation where fluid resuscitation and treatment cannot be immediately initiated
- Aggressive fluid resuscitation for Crush injury in the setting of noncompressible hemorrhage may increase hemorrhage. Balance the risk of uncontrolled hemorrhage against cardiotoxic effects of hyperkalemia.

EXTREMITY TRAUMA

| Signs and Symptoms | Signs and Symptoms (cont.) | | | | | |
|--|---|--|--|--|--|--|
| Pain/Swelling | Abrasion | | | | | |
| Deformity | Contusion | | | | | |
| Altered Sensation/Function | Multi-Trauma | | | | | |
| Diminished Pulse/ Cap refill | Fracture | | | | | |
| Decreased Temperature | Dislocation | | | | | |
| Bleeding Laceration | | | | | | |
| Amputation Sprain/Strain | | | | | | |
| Treatment | | | | | | |
| Heavy Active Bleeding | | | | | | |
| Check/Add Tourniquet (TQ) | | | | | | |
| Add Deliberate TQ if Hasty is in place | | | | | | |
| Pack and Dress Wound | | | | | | |
| Pressure Dressing | | | | | | |
| Hemostatic Dressing | | | | | | |
| Amputation | | | | | | |
| Follow Heavy Active Bleed Guidelines | | | | | | |
| Wrap Amputation in Sterile Dressing with No | | | | | | |
| Place in sealed container with ice slurry if av | ailable | | | | | |
| Transport with patient | | | | | | |
| | | | | | | |
| Convert Limb/Junctional TQ as soon as possible | e if: | | | | | |
| No presence of shock | | | | | | |
| , , , | · · · · · · · · · · · · · · · · · · · | | | | | |
| Not placed to control hemorrhage on amputa | | | | | | |
| Every effort should be made to convert in les | is than 2 hours if patient is not in shock | | | | | |
| Wound Care/Protection | | | | | | |
| Bandage/cover injuries | | | | | | |
| Immobilize extremity | | | | | | |
| Ice (if available) for Edema | | | | | | |
| After Diseding Controlled | igs, Cautions | | | | | |
| After Bleeding Controlled: Treat for signs and symptoms of Hypetension/ | Shool | | | | | |
| o Treat for signs and symptoms of Hypotension/S | | | | | | |
| Follow DCR protocols regarding hierarchy of flui | | | | | | |
| Carefully evaluate and document Neurovascular | | | | | | |
| Never attempt to reduce an open fracture unless | • | | | | | |
| | wer extremity injury and may be utilized in conjunction | | | | | |
| with a traction splint. | have freetures consciently the formur | | | | | |
| Blood loss can be severe and concealed in long | | | | | | |
| TQs should be used without hesitation to control | major bleeding | | | | | |
| Use only CoTCCC approved Tourniquets | | | | | | |
| Reference: CPG ID: 62 Acute Traumatic Wound Management & TCCC Guidelines | | | | | | |

Reference: CPG ID: 62 Acute Traumatic Wound Management & TCCC Guidelines

EYE INJURY/PAIN

| <u>History</u> | (Complications) |
|----------------|-----------------|
|----------------|-----------------|

- Abrasion / Laceration
- Globe Rupture / Orbital fracture
- Retinal Detachment
- Chemical / Thermal Burn
- Infection / Iritis
 - CNS Event
 - Glaucoma
 - Retinal Vessel Occlusion

<u>Treatment</u>

•

• Without Known Injury

Pain, Swelling, Blood

Deformity / Contusion

Excessive Tearing

Foreign Body

- Evaluate Pupils
- Consider unrecognized chemical exposure.

History

Decreased Visual Acuity / Blindness

• Irrigate with minimum 2L NS for chemical exposure; 30 min minimum.

• With Known Injury

- o If not isolated, move to appropriate guideline to treat life threats.
- Assess orbital stability / pupils.

• Chemical Injury

• Irrigate with minimum 2L NS for chemical exposure; 30 min minimum.

Traumatic Injury

- Remove loose debris with NS irrigation. Do not attempt to remove impaled objects or contacts.
- Cover with rigid eye shield. DO NOT PLACE ANY DRESSING/PADDING UNDERNEATH EYE SHEILD.
- o If penetrating, give moxifloxacin 400mg PO / IV.
- Refer to Pain control Guideline.
- Treat for Nausea / Anxiety
 - Nausea: Ondansetron 4-8 mg IV / IO / IM
 - Anxiety:
 - Diazepam 2-10 mg IV / IO / IM
 - Midazolam 2.5-5 mg IV

- Antiemetics are essential to prevent increased IOP. Consider Benzo for anxiety.
- Use rigid eye shields, not pads, for traumatic injuries. Can use a soft pad on unaffected eye.
- Patching both eyes to decrease sympathetic eye movements has not been shown to improve visual outcome but may increase anxiety and will render patient unable to move independently.
- If globe is out of socket do not attempt to replace. Cover with saline soaked gauze
- Copious irrigation is the cornerstone of treatment for chemical eye injuries. Some chemical injuries can require up to 10L. 30 min is the minimum amount of time to irrigate. Utilize Morgan lens if available.
 - The use of a nasal cannula across the bridge of the nose attached to 1L of NS will also work.

HEAD INJURY/TBI

Signs and Symptoms

- Head Pain, Swelling, Bleeding
- Head Deformity, Ecchymosis
 Altered Montel Status
- Altered Mental Status
- Respiratory Distress/Failure
- Vomiting
- Spinal Injury

Signs and Symptoms (cont.)

- Skull Fracture
- Epidural/Subdural Hematoma
- Subarachnoid Hemorrhage
- Abuse
 - Definitions:
 - Mild TBI GCS 13-15
 - Moderate TBI GCS 9-12
 - Severe TBI GCS 3-8

<u>Treatment</u>

- Consider Spinal Immobilization (minimize compression of neck veins)
- Resuscitate, follow Tactical Evacuation Guidelines
- TXA 2g IV/IO in *moderate to severe TBI*
- Levetiracetam 1500mg IV/IO bolus in *severe TBI* (seizure prophylaxis for transport)
- Airway Compromise
 - o Establish airway (Airway protocol) with supplemental O2
 - Monitoring Goals:
 - SPO2 > 95%
 - ETCO2 35-45mmHg (normoventilation)
 - No Obvious Airway Compromise
 - Jaw Thrust, NPA
 - Supplemental O2 via BVM (SpO2 > 95%)
 - Low threshold to RSI if deteriorating GCS/mental status
- Evidence of Elevated ICP
 - Elevate head of bed to 30 deg
 - o 3% Hypertonic Saline 250 mL IV/IO over 10 min
 - Target Vital Signs:
 - SBP>110
 - SPO2> 95%
 - ETCO2 35-45mmHg (normoventilation, do not hyperventilate)
 - CCP >60 (CCP=MAP ICP)
- Evidence of Impending Herniation (e.g., sluggish, unilateral/bilateral dilated or fixed pupil, presence of Cushing's triad) [Cushing's triad = (relative) bradycardia, hypertension/widening pulse pressures, irregular respirations]:
 - Continue elevated ICP treatment as above
 - Maintain ETCO2 35-45mmHg (normoventilation; avoid hyperventilation)
 - Request online medical control for further guidance in prolonged flight

Notes, Warnings, Cautions

- Ensure Continuous monitoring q5-10 min
- Active seizures: Lorazepam or Midazolam, see Seizure protocol
- Mannitol given as boluses: 1g/kg bolus followed by 0.25mg/kg bolus every 4 hours
- Keep SBP>110: consider LR/NS bolus
- Avoid Hypo/Hyper-capnea through dedicated closely managed ventilation
- Sedation: Ketamine is preferred over propofol due to hemodyanamic effects of propofol. Monitor SBP.
- Paralysis: Not preferred in head injury if avoidable; vecuronium preferred; ensure pain control/sedation is adequate to avoid increased ICP

Reference CPG ID: 63 (TBI Management in Prolonged Field Care) Reference CPG ID: 30 (TBI Management and Basic Neurosurgery in the Deployed Environment)

TRAUMATIC ARREST

History

- Evidence of trauma without a pulse
- Unresponsive to external stimuli

History (Differential)

- Medical cause of arrest preceding trauma
- Tension pneumothorax
- Hypovolemia
 - Cardiac Tamponade

Treatment

- Determine if injuries are incompatible with life.
 - \circ $\,$ Do not resuscitate if injuries are incompatible with life.
- Address all known points of hemorrhage.
 - Initiate transfusion with 1 unit of blood product (avoid resuscitation with crystalloid)
 - \circ TXA 2g IV/IO within 3 hours of injury
- Begin CPR
- Place advanced airway
- Start supplemental O2
- Bilateral needle thoracostomy
- Consider Advances Procedures
 - Finger Thoracostomy
 - Tube Thoracostomy
 - \circ Pericardiocentesis
- Place monitor on patient: Prepare Defibrillator
 Determine Rhythm: Pulse return?
- ROSC not Achieved
 - Continue CPR
 - Continue Blood / IV Fluids
 - Reduce Long Bone Fractures
 - Reduce Pelvic Fracture
 - Reassess known hemorrhage points
- ROSC Achieved
 - Return to Tactical Evacuation or Previous Guideline

Notes, Warnings, Cautions

- **Injuries obviously incompatible with life** include decapitation, massively deforming head or chest injury, traumatic hemi-corpectomy or total body disruption, incineration, lividity/rigor mortis.
- Casualties with torso trauma or polytrauma who have no pulse or respirations during Tactical Field Care should have bilateral needle decompression performed to ensure they do not have a tension pneumothorax prior to discontinuation of care.
- If unsure if arrest due to trauma or medical cause, initiate ALS guideline for any arrhythmias following optimization of hemostasis (in trauma patients, volume loss must be corrected 1st, consider blood admin above all else)
- CPR without addressing massive hemorrhage, blood volume resuscitation, tension pneumothorax, and pericardial tamponade will be ineffective.
- *Consider severe hypocalcemia if blood products have recently been transfused due to calcium chelation and evidence of poor cardiac activity/contractility.

Ref: CPG ID:82 & TCCC Guidelines

HYPOTENSION/SHOCK

| <u>History</u> | History (Complications) |
|--|--|
| Restlessness / Confusion | Nausea / Vomiting |
| Weakness / Dizziness | Shock: Hypovolemic, Cardiogenic, Septic, |
| Tachycardia | Neurogenic, Anaphylactic |
| Pale, Cool, Clammy Skin | Cardiac Arrhythmia |
| Delayed Cap. Refill | Pulmonary Embolus |
| Blooding | Tension Pneumothorax |
| Dehydration | Medication OD |
| Congenital Heart Disease | Vasovagal Episode |
| Treatment | |

<u>Treatment</u>

- Shock Due to Hemorrhage / Trauma:
 - Control hemorrhage
 - Optimize homeostasis (See Notes, Warnings, Cautions))
 - Optimize hypothermia management
 - TXA 2G IV/IO
 - Maintain SBP >100 (>110 TBI), move to appropriate protocol for continued treatment (i.e., Blood Administration, etc.)
- Shock Due to Non-Traumatic & Non-Cardiac
 - o 2L or 30mL/kg IVF bolus PRN, additional crystalloid based on reassessment of clinical indication.
 - o If inadequate, consider NOREPINEPHRINE 2-20 mcg/min IV/IO
 - Maintain SBP >90, MAP >65
- Shock Due to Cardiac:
 - Treat per appropriate Cardiac Guideline:
 - BRADYCARDIA W/ A PULSE
 - CARDIAC ARREST
 - TACHYCARDIA W/ A PULSE
 - Non-Invasive PPV (BVM) vs. Advanced Airway
 - o 500mL IVF Bolus
 - If inadequate, NOREPINEPHRINE 2-20mcg/min IV/IO

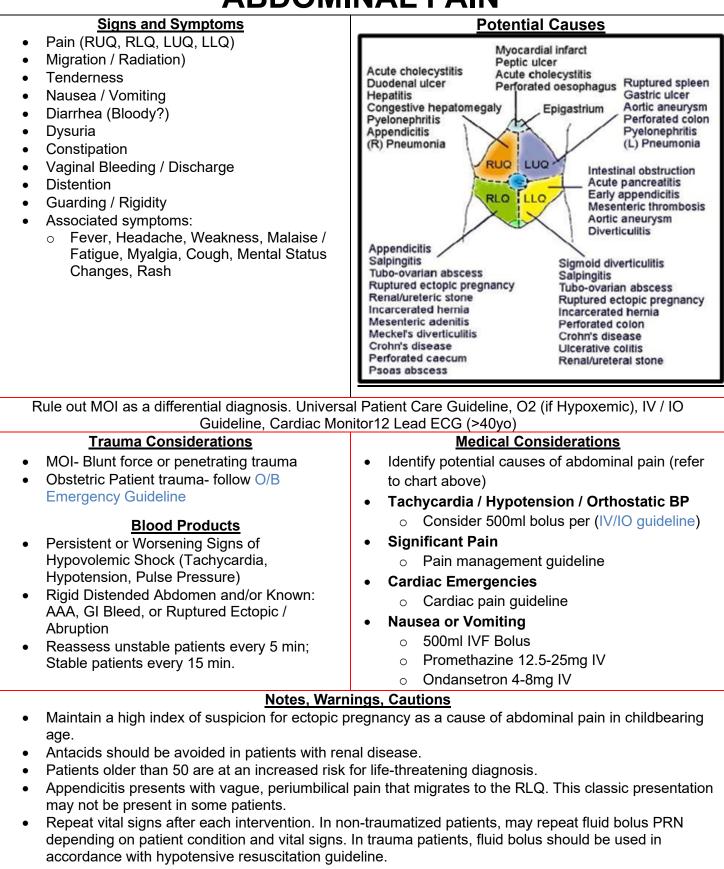
- Optimize Homeostasis:
 - <u>Hemorrhage trauma with NO significant Head Injury</u>: Should target maintaining SBP >100.
 Casualties able to maintain SBP >100 do not need immediate fluid resuscitation.
 - <u>Hemorrhage trauma WITH significant Head Injury:</u> Should target maintaining SBP >110.

PEDIATRIC HYPOTENSION/SHOCK

| : == :: | | |
|--|---|--|
| HistoryRestlessness / ConfusionWeakness / DizzinessTachycardiaPale, Cool, Clammy SkinDelayed Cap. RefillBloodingDehydrationCongenital Heart Disease | <u>History (Complications)</u> Nausea / Vomiting Shock: Hypovolemic, Cardiogenic, Septic, Neurogenic, Anaphylactic Cardiac Arrhythmia Pulmonary Embolus Tension Pneumothorax Medication OD Vasovagal Episode | |
| <u>Treat</u> | ment | |
| Due to Hemorrhage / Trauma: Trauma Fluid Preferences Whole Blood (if available) pRBC's and plasma (if available) (LR/NS) 20mL/kg IVF Bolus 10mL/kg Blood Product If inadequate, consider (as last resort) EPINEPHRINE 1mcg/kg/min IV/IO or NOREPINEPHRINE .05- 0.1 mcg/kg/min slow IV push q10-15min Due to Non-trauma & Non-cardiac: 20mL/kg (NS/LR) IVF bolus If inadequate, consider EPINEPHRINE 1mcg/kg/min IV/IO or NOREPINEPHRINE .05 – 0.1 | | |
| mcg/kg/min IV/IO (max 2mcg/kg/min) | | |
| Due to Cardiac: | | |
| Treat per appropriate Pediatric Cardiac Guid | eline: | |
| Pediatric BRADYCARDIA W/ PULSE and | | |
| Pediatric TACHYCARDIA W/ PULSE and poor perfusion | | |
| Pediatric CARDIAC ARREST | | |
| Non-invasive PPV (BVM) vs. Advanced airway | | |
| If rales heard on lung exam, 5 – 10mL/kg IVF over 5 – 10 min | | |
| If inadequate, consider EPINEPHRINE 1mcg/kg/min IV/IO | | |
| Notes, Warnings, Cautions | | |
| Hypotension in pediatric patients is defined as SBP < 70 +[2 x age(yr)] | | |
| Decreasing heart rate with worsening neuro or clinical exam may be sign of impending collapse in | | |
| pediatric patients. | | |
| Consider all causes of shock and treat per appropriate protocol. | | |
| • <u>Avoid Pressors</u> (last resort) as able, unless distributive or cardiogenic shock. Continue IVF for trauma; | | |
| optimize homeostasis and correct valume | | |

optimize homeostasis and correct volume.

ABDOMINAL PAIN



• Conservative approach with lower promethazine dosage for patients likely to experience sedative effects (e.g., elderly patients).

ALLERGIC REACTION

| | NEACTION |
|---|--|
| Signs and Symptoms | Differential Diagnosis |
| Itching or Hives | Urticaria (rash only) |
| Cough / Wheeze / Resp. Distress | Anaphylaxis (2 or more systems) |
| Chest / Throat tightness | Shock (other than anaphylactic) |
| Difficulty Swallowing | Angioedema |
| Hypotension or Shock | Aspiration / Airway Obstruction |
| • Edema | Asthma or COPD |
| Nausea / Vomiting | Pulmonary Edema / CHF |
| Rule out MOI as a differential diagnosis. Universal | Patient Care Guideline, O2 (if Hypoxemic), IV / IO |
| Guideline, Cardia | c Monitor (ASAP) |
| ADULT | PEDIATRIC |
| Hives / Rash Only, No Respiratory | Hives / Rash Only, No Respiratory |
| Complaint | Complaint |
| Administer Diphenhydramine 25-50mg IV/IO/IM/PO | Diphenhydramine; 1-2 mg/kg IM/IV/PO (max dose 50mg) |
| Administer Methylprednisolone 125mg | Administer Methylprednisolone 2mg/kg |
| IV/IO | IV |
| Shock / Unresponsive or Respiratory | |
| Distress / Failure | Shock / Unresponsive or Respiratory |
| Epinephrine-Pen or Epinephrine 1:1000; | Distress / Failure |
| 0.3-0.5mg IM | • Epi-Pen Jr for <30kg or Epinephrine |
| 500ml IVF if not previously started. Albutaral 00, 120mag 2 muffa MDL at 2.5 | 1:1,000; 0.01mg/kg IM (max 0.3mg) |
| Albuterol 90-180mcg 2 puffs MDI or 2.5- 5mg via nebulizer | 20 mL/kg IVF Bolus Albuterol metered-dose inhaler |
| | Albuterol metered-dose inhaler 4 – 8 inhalations every 20 min |
| Diphenhydramine 25-50mg IV/IO/IM/PO Methylprednisolone 125mg IV/IO | \circ Albuterol by nebulizer |
| | Children < 20 kg; 2.5 mg/dose |
| *Worse or Unstable* | Children > 20 kg; 5 mg/dose |
| • Epinephrine IV infusion at 5-15 mcg/minute | Diphenhydramine 25-50mg |
| (see Epinephrine infusion chart in drug card | IV/IO/IM/PO |
| for procedure) | Methylprednisolone 2mg/kg IV; |
| | maximum daily dose, 120 mg |
| Notes, Warnings, Cautions | |
| Use caution prior to giving epinephrine IV to patients >50yo, pregnant, have a history of cardiac | |
| disease, or have HR >150. Epinephrine can precipitate dysrhythmias / ischemia – all patients should | |
| be on monitors and have 12-lead ECG. | |
| Epinephrine can precipitate dysrhythmias / ischemia – all patients should be on monitors and have 12- lead ECG. | |
| The shorter the interval from contact to symptoms, the more severe the reaction. | |
| • The shorter the interval norm contact to symptoms, the more severe the readilon. | |

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- Arrhythmia- See appropriate pediatric cardiac guidelines. Non-arrhythmia- See hypotension guideline or respiratory distress guideline. •

ALTERED MENTAL STATUS

Signs and Symptoms **Differential Diagnosis** Any signs of head trauma/injuries? Head trauma/psychiatric disorders Any AMS? Thyroid dysfunction Any pertinent medical conditions or Hyper/hypoglycemia medical history? Diabetic ketoacidosis/toxic Ingestion Are there any bystanders that can Environment (hyper/hypothermia) provide information about the patient? Hypoxia Is this abnormal behavior? Safety of the helicopter/crew/other patients take PRIORITY! Treatment Does the patient have a head injury, unable to protect their airway (GCS<8), violent behavior, and/or AMS? Refer to head injury guideline, airway guideline if applicable. 0 Determine blood glucose, if <70 or >250 go to hypoglycemia or hyperglycemia guideline. If patient is an EPW or potential hostile, consider security escort and/or physical restraints. Attempt to calm and reassure the **combative patient** and use physical restraints if needed. Medications can be used to help calm the patient. Ensure that patient has their ETCO2 monitored after administration. Ketamine 4-5mg/kg IM/IN or 1-2 mg/kg IV/IO can repeat q 10min. 0 Lorazepam 2-4mg IV/IM (can use alone). 0 Midazolam 2.5-5mg IV/IM q15-30 min prn (larger patients may need 10mg if using IM route). If patient is still combative after the use of medications, consider RSI guideline. •

- Physical restraints such as tying down patient hands to prevent pulling lines, etc., should be limited to the least amount necessary to accomplish treatments / prevent injuries. (Kerlix gauze can be a useful restraint)
 - **Do not jeopardize the patient's airway!** Avoid hog tying, lying prone in restraints, sandwiching between spine boards, etc.
 - Check Vitals, SpO2, Pulse and Cap Refill every 5 minutes.
- Combative patients present a very real threat to the safety of themselves, the medic, and the aircrew during flight. For this reason, any patient with altered mental status and the potential for combativeness that would threaten aircrew safety or themselves should be prophylactically sedated and/or paralyzed and intubated for the flight.
- Use of sedative medications adds risk of decreasing respiratory drive and should be used with caution. However, meds should be titrated to adequate dosage to control patient. Be prepared for airway interventions/vomiting if used. Cardiac arrest in patients with excited delirium/extreme agitation following restraint is well documented. Capnography in addition to cardiac monitoring is essential.

BACK/NECK PAIN

| Signs and Symptoms • Pain • Swelling • Pain with motion • Weakness / Numbness • Bowel / Bladder Dysfunction | Differential Diagnosis• Muscle Spasm / Strain• Degenerative Disc Disease• Fracture• Kidney Stone / Infection• Abdominal Aortic Aneurysm |
|---|---|
| | Pneumonia / PE Cauda Equina Syndrome Tumor / Mass / Infection Thoracic Pain: Thoracic or abdominal aortic aneurysm |
| *Rule out MOI as a differential diagnosis. Universal Patient Care Guideline, O2 (if Hypoxemic), IV / IO Guideline (prn), Cardiac Monitor (prn) | |
| Injury Treatment • Injury/Trauma • Mechanisms that increase suspicion of possible Spinal Cord Injury: • Blunt trauma to head or neck • Injury associated with high energy • transfer (e.g., blast, motor vehicle) • Fall from >3 ft. • Fall directly onto head / neck • History of back / neck arthritis plus • Spinal Immobilization Guideline • Head Injury Guideline • Multiple Trauma Guideline | Medical Treatment • Extremity BP difference/ Suspicion of AAA • 1000mL IVF IV any trauma • Consider: Blood Product for AAA • Suspicion of ACS/ Chest Pain • Chest Pain Guideline • Arrhythmia • Bradycardia with Pulse Guideline • Cardiac Arrest Guideline (VF / Pulseless VT or Asystole / PEA) |
| Notes, Warnings, Cautions | |
| EXAMINE: Mental Status, HHENT, neck, chest, abdomen, back, extremities and neurologic. Abdominal Aortic Aneurysm is a concern in hypertensive / diabetic / >50y populations- feel for pulsatile abdominal mass. Symptoms may mimic kidney stones. | |

- Patients with trauma and midline tenderness should be immobilized.
- Any bowel/ bladder incontinence is significant and may represent a true medical emergency.

| HYPERGLYCEMIA/HYPOGLYCEMIA | | |
|--|---|--|
| Hyperglycemia S/S BLOOD GLUCOSE >250 Polyuria Polydipsia Weakness/fatigue Nausea/vomiting Change in LOC Hypotension Tachycardia Seizures/coma Fruity Breath Odor | Hypoglycemia S/S BLOOD GLUCOSE <70 (<100 in neuro injury) Diaphoresis Pallor AMS Tremor Palpitations Anxiety | |
| Treatment Place Patient on Cardiac Monitor Obtain Blood Glucose Blood Glucose >250mg/dL Initiative IV or IO Access Administer 1000ml .09%NS (10-20ml/kg) Monitor blood glucose every 30 minutes. Consider Intubation for patients with AMS. Nausea or vomiting present, administer: Promethazine 12.5-25mg IV OR Ondansetron 4-8mg IV | Treatment Place Patient on Cardiac Monitor Obtain Blood Glucose Blood Glucose <70mg/dL Initiate IV or IO Access Patients with AMS: Administer 10-25g IV Dextrose (40-100ml of 25% solution or 20-50ml of 50%) If IV access unobtainable, administer Glucagon 1mg IM. Repeat after 20 minutes as needed. Patients with NO AMS: Administer oral glucose gel or equivalent until glucose level is >70mg/dL. | |
| <u>Notes, Warnings, Cautions</u> If insulin is available, treat with low dose infusion, 0.1 units/kg/hr. Too rapid drop in blood glucose can cause hypoglycemia. Rapid drop in blood glucose levels can lead to shifts extracellular osmolality which can lead to cerebral edema. | <u>Notes, Warnings, Cautions</u> If administering Dextrose, obtain blood glucose sample from contralateral arm. Hypoglycemia may be detrimental to patients at risk for cerebral ischemia, such as victims of stroke, cardiac arrest, and head trauma. Hypoglycemic patients must be alert enough to swallow and protect airway. | |

LOWER RESPIRATORY DISTRESS

| Signs and Symptoms Shortness of Breath Pursed Lip Breathing Decreased Ability to Speak Tachypnea / Hyperpnea Wheezing / Rhonchi / Rales Use Accessory Muscles Fever / Cough Tachycardia Absent Breath Sounds Wheezes Monitor O2 and ETCO2 Place on 100% oxygen via NRB. Administer Albuterol 90-180mcg or 5mg nebulized Monitor for allergic reactions Consider Epinephrine 1:1,000 0.3-0.5mg IM (EPI PEN) Initiate IV/IO Access Administer Methylprednisolone 125mg IV Consider Magnesium Sulfate 2G IV over20 mins. Dilute in 50-100ml NS or D5W. As a last resort, administer Ketamine 1mg/kg | Potential CausesAsthmaAnaphylaxis / AllergyAspirationCOPDPleural EffusionPneumoniaCongestive Heart Failure / CardiacPulmonary EmbolusPneumothoraxPericardial TamponadeHyperventilationToxic Inhalation (e.g., Cyanide, CO) |
|---|--|
| IV slow push. | |
| Wheezes • Place on 100% oxygen via NRB. • Administer Albuterol 90mcg MDI or 5mg nebulized (Max 12 doses per 24hrs) • Monitor for allergic reactions: Consider Epinephrine (1:1000) 15-30kg give 0.15mg IM (EPIPEN JR) >30kg give 0.3mg IM (EPIPEN) OR 0.01mg/kg IM (max 0.3mg) Administer Methylprednisolone 1-2mg/kg IV. Administer Magnesium Sulfate 25-50mg/kg IV over several minutes (Max 2G) diluted in 50-100ml NS. Last resort, administer Ketamine 0.5mg/kg IV slow push. | <u>Rales/CHF</u> Monitor O2 and ETCO2 Provide PPV (if tolerated) with 100% oxygen support <i>OR</i> 100% NRB if PPV is not tolerated. Failure to improve: Administer Furosemide 1mg/kg IV slow push, place foley if possible. |
| Notes, Warnings, Cautions SPO2 <90% or respiratory status continues deteriorate, consider definitive airway control. Albuterol can be administered with spacer or short (6") section of ventilator tubing to increase delivery if patient unable to perform action appropriately. No max dose of albuterol, repeat PRN for continued wheezing. | |

• Lack of abnormal breath sounds does not always signify improvement. As respiratory status worsens, there may be inadequate air movement to produce these sounds.

UPPER RESPIRATORY DISTRESSSigns and SymptomsPotential Causes

| Signs and Symptoms | Potential Causes | | |
|---|---|--|--|
| Shortness of Breath | Asthma | | |
| Pursed Lip Breathing | Anaphylaxis / Allergy | | |
| Decreased Ability to Speak | Aspiration | | |
| Tachypnea / Hyperpnea | COPD | | |
| Wheezing / Rhonchi / Rales | Pleural Effusion | | |
| Use Accessory Muscles | Pneumonia | | |
| Fever / Cough | Congestive Heart Failure / Cardiac | | |
| 5 | Pulmonary Embolus | | |
| Tachycardia Absort Brooth Counds | Pneumothorax | | |
| Absent Breath Sounds | Pericardial Tamponade | | |
| | Hyperventilation Toxic Inhalation (e.g. Cyanide, CO) | | |
| | | | |
| View for obstructions (jaw-thrust for C-spine injury) sweep and suction prn. | | | |
| Monitor O2 and ETCO2 | | | |
| Place on 100% oxygen via NRB | | | |
| Administer Albuterol 90-180mcg MDI 2 puffs or 2.5-5mg nebulized. | | | |
| Monitor for allergic reactions: | | | |
| Consider Epinephrine 1:1,000 0.3mg IM (EPI PEN) | | | |
| Initiate IV/IO Access | | | |
| Administer Methylprednisolone 125mg IV. | | | |
| PEDIATRIC | | | |
| View for obstructions (jaw-thrust for C-spine injury) sweep and suction prn. | | | |
| Monitor O2 and ETCO2 | | | |
| Place on 100% oxygen via NRB | | | |
| Administer Nebulized Racemic Epinephrine (1:1000) 0.5mL/kg (Max dose 5mL) | | | |
| Monitor for allergic reactions: | | | |
| • <u>Consider Epinephrine</u> | | | |
| 15- 30kg: 0.15mg IM (EPIPEN JR) > 20kg: 0.3mg IM (EPIPEN) OB | | | |
| >30kg: 0.3mg IM (EPIPEN) <u>OR</u> 1:1000, 0.01mg.kg IM (max 0.3mg) | | | |
| Administer Methylprednisolone 1-2mg/kg IV. | | | |
| | ngs Cautions | | |
| | Notes, Warnings, Cautions | | |

- SPO2 <90% or respiratory status continues deteriorate, consider definitive airway control.
- Albuterol can be administered with spacer or short (6") section of ventilator tubing to increase delivery if patient unable to perform action appropriately. No max dose of albuterol, repeat as needed for continued wheezing.
- Lack of abnormal breath sounds does not always signify improvement. As respiratory status worsens, there may be inadequate air movement to produce these sounds.

CEITHDE

| SEIZURE | | | | | | | |
|--|---|--|--|--|--|--|--|
| Signs and Symptoms | Initial Treatment | | | | | | |
| Decreased Mental Status | Provide oxygen support. | | | | | | |
| Seizure Activity | Consider airway control (Definitive if necessary) | | | | | | |
| Somnolence | Initiate IV/IO Access | | | | | | |
| Incontinence | Place on cardiac monitor | | | | | | |
| Evidence of Trauma | Monitor Blood Glucose | | | | | | |
| Loss of Consciousness | | | | | | | |
| Oral Injuries (e.g., Tongue, Buccal) | | | | | | | |
| | Freatment | | | | | | |
| Active Seizure | | | | | | | |
| Administer Midazolam (5mg IV/IO/IN or 10m OR | g IM) | | | | | | |
| Lorazepam 4mg IV/IM OR Diazepam 4mg IV May report anticepy/legate two times or | | | | | | | |
| | ery 3-5 minutes if seizure has not stopped. in; followed by maintenance dosing of 1000mg IV/IO | | | | | | |
| every 12 hours. | in, followed by maintenance dealing of recently tyre | | | | | | |
| Pregnancy Considerations: | | | | | | | |
| Administer Magnesium Sulfate 4G IV over | r 15min. (Monitor for hypotension) | | | | | | |
| Blood Glucose <70 | | | | | | | |
| Administer 50% Dextrose 25G IV | | | | | | | |
| OR Clucadon 1mg IV/IM | | | | | | | |
| Glucagon 1mg IV/IM Pediatric Treatment | | | | | | | |
| Active Seizure | | | | | | | |
| Administer Lorazepam 0.05-0.1 mg/kg IV (m | | | | | | | |
| • • | ery 3-5 minutes if seizure has not stopped. Must | | | | | | |
| establish definitive airway control prior to | | | | | | | |
| Levetiracetam (Keppra) 60mg/kg, single dos Blood Glucose <40 | Э. | | | | | | |
| Administer Thiamine 25mg IV/IM OR 25% D | extrose 2ml /kg IV | | | | | | |
| OR | | | | | | | |
| Glucagon 0.05mg/kg IM | | | | | | | |
| | ings, Cautions | | | | | | |
| For seizure prophylaxis consider Levetiracetam consider maintenance dose of 1000 mg every 12 | | | | | | | |
| 3 , | two or more continuous seizures without a period of | | | | | | |
| consciousness / recovery. This is a real emerger | | | | | | | |
| transport to the nearest suitable medical treatme | | | | | | | |
| Paralysis for airway control does not stop seizure | e activity – only hides it. A seizure is a CNS electrical | | | | | | |
| | n when no muscular activity seen due to paralysis. | | | | | | |
| | line to abort non-epileptic seizures. Midazolam should | | | | | | |
| only be used if this fails (pregnancy class D). | , , , , , , , , , , , , , , , , , , , | | | | | | |
| | se of midazolam/lorazepam. If airway controlled and | | | | | | |
| pediatrics. | dazolam for adults and 4 doses of Lorazepam for | | | | | | |
| | | | | | | | |

SEPSIS/FEVER

| <u>Definition</u> Life-threatening organ dysfunction caused by a dysregulated host response to infection. Septic shock is a form of distributive shock. <u>Keys to Success</u>: Early recognition Identification of the cause of shock Early, decisive treatment of the cause and initiation of cause-specific resuscitation. |
|---|
| <u>atment</u> Temp |
| |

- Attain a minimum of 2 IV/IO sites.
- Initiate IV/IO crystalloid therapy 1000ml Bolus NS/LR achieve goal of SBP > 90 or MAP >65 or 30ml/kg.
- Initiate/Monitor Foley. UOP 0.3-0.5ml/kg/hr.
- Temperature >100.4°F / 38°C consider Acetaminophen 1gram PO/IV (if not provided in the last 6 hours)
- Persistent or Refractory Hypotension after 2L NS/LR, unable to maintain SBP > 90 or MAP > 65?
 Administer Norepinephrine 2-12mcg/min IV.
 - Add Vasopressin 0.03 units/min SBP < 90 or MAP < 65.
 - Add Epinephrine 2-20 mcg/min SBP < 90 or MAP < 65.
- Consider Antibiotic Therapy Ceftriaxone 2 g slow IV push or in 100 cc NS flow to gravity (immunocompetent) or Cefepime 2 g IV in 100 cc NS flow to gravity (immunocompromised)
- Contact Medical Control for further if able.

Notes, Warnings, Cautions

- Use vasopressin despite less than maximal norepinephrine. Consider adding it when titrating above 8-10 mcg/min IV norepinephrine. Continue it once started and decrease norepinephrine to MAP goal > 65.
- Monitor overall respiratory status. Many patients who are critically ill with sepsis will need ventilatory support at some point in their management.
- Record urine output if foley in place. Decreased urine output is an indicator of patient deterioration.
- Fever may not be present in immunocompromised, elderly, or those on immunosuppressive drugs.
- All fever is not due to infection evaluate for environmental / thyroid / toxic etiology.
- In Trauma Sepsis, Blood is Preferred
- Caution in over-resuscitation >.5ml/kg/hr UOP, wet lungs, increased work of breathing.

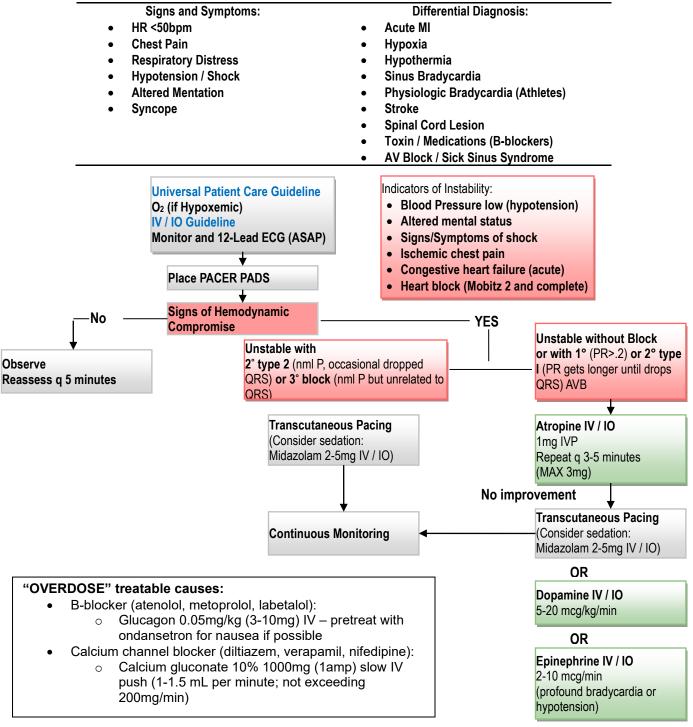
| | CESTION | | | | | | |
|--|--|--|--|--|--|--|--|
| | GESTION | | | | | | |
| | (in US): 1-800-222-1222 | | | | | | |
| | /www.poison.org | | | | | | |
| Signs and Symptoms Mental Status Changes Hypo/Hypertension Respiratory Depression Tachycardia/Arrythmias Seizure | Differential Diagnosis Cyclic Antidepressants Acetaminophen Depressants Stimulants Anticholinergic Cardiac Medications Solvents/Cleaners | | | | | | |
| Adult | Treatment | | | | | | |
| Blood Sugar <60: AMS Guideline (50%D50 25 Blood Sugar >60: Activated Charcoal 1g/kg PC Beta Blocker Overdose: Glucagon 3-10mg IV/ Opiates: Naloxone 0.4-2mg IV/IO - Watch for R TriCyclic Anti Depressant: 12 Lead QRS>100 or Hypotensive? Sodium Bicarbonate 1mEq/kg 100-150 mEq in 1L D5/NS @ 100-200m | g in 500ml NS or Glucagon 1mg/IM)) (If alert and <1hr from ingestion) IM Bolus followed by 3-5mg/hr espiratory Depression | | | | | | |
| Seizure - Midazolam 2.5-5mg IV/IM Dedictric Treatment | | | | | | | |
| Blood Sugar <65: AMS Guideline (25% Dex 2ml Blood Sugar >60: Activated Charcoal 1g/kg PO Beta Blocker Overdose: Glucagon 1mg IV/IM Opiates: Naloxone 0.1mg/kg IV - Pediatric Airwa TriCyclic Anti Depressant: 12 Lead QRS>100 or Hypotensive? Sodium Bicarbonate 1mEq/kg 100-150 mEq in 1L D5/NS @ 100-200m Organophosphate: Atropine 0.02mg/ IV/IO q5 + Seizure - Lorazepam 0.1mg/kg IV | (If alert and <1hr from ingestion) ay Guideline //hr 2-PAM 25mg/kg IV/IM | | | | | | |
| <u>Notes, War</u> | nings, Cautions | | | | | | |
| Follow TriCylcic dosing Lorazepam for agitation and seizures Beta Blockers watch for Hypoglycemia Calcium Channel Blockers watch for Hyperglyce Cyclic Antidepressants signs: Hypotension, depressants arrythmias | ressed mental status, respiratory depression, and | | | | | | |
| | pupils, N/V, Respiratory depression, hypotension , Urination, Diarrhea, Emesis, Altered Mental Status | | | | | | |

VOMITTING & DIARRHEA

| Signs and Symptoms | Differential Diagnosis | | | | | |
|---|------------------------|--|--|--|--|--|
| Pain | CNS Injury / Infection | | | | | |
| Abdominal Distention | Myocardial Infection | | | | | |
| Constipation | Drugs / Toxins | | | | | |
| Diarrhea | Pregnancy | | | | | |
| Anorexia | Gastroenteritis | | | | | |
| Fever | Appendicitis | | | | | |
| Rash | Bowel Obstruction | | | | | |
| <u>Adult Tr</u> | eatment | | | | | |
| IV - O2 - Monitor | | | | | | |
| Blood Glucose <60 w/ evidence of alcohol abuse | e? | | | | | |
| Thiamine 100mg IV/IM | | | | | | |
| 50% Dextrose 25g IV or Glucagon 1mg IM | | | | | | |
| If Glucose is outside <60 start over | | | | | | |
| N/V - Promethazine (if >2yr old) 12.5mg IV or O | ndansetron 4-8mg IV | | | | | |
| | Treatment | | | | | |
| IV - O2 - Monitor | | | | | | |
| Blood Glucose <65 w/ evidence of malnourishm | ent? | | | | | |
| Thiamine 100mg IV/IM | | | | | | |
| 25% Dextrose 2ml/kg IV or Glucagon 0.5mg | IM | | | | | |
| Nausea / Vomiting: | | | | | | |
| Promethazine (if >2yr old) 0.25mg/kg IV | | | | | | |
| Ondansetron (<40kg) 0.1mg/kg IV | | | | | | |
| Ondansetron (>40kg) 4mg IV | | | | | | |
| | nings, Cautions | | | | | |
| Suspicions of underlying conditions should prom | | | | | | |
| In pregnant patients with N/V - substitute D5 1/2 | | | | | | |
| Fluid of choice for vomiting is NS. Fluid of choice | e for diarrhea is LR | | | | | |
| | | | | | | |

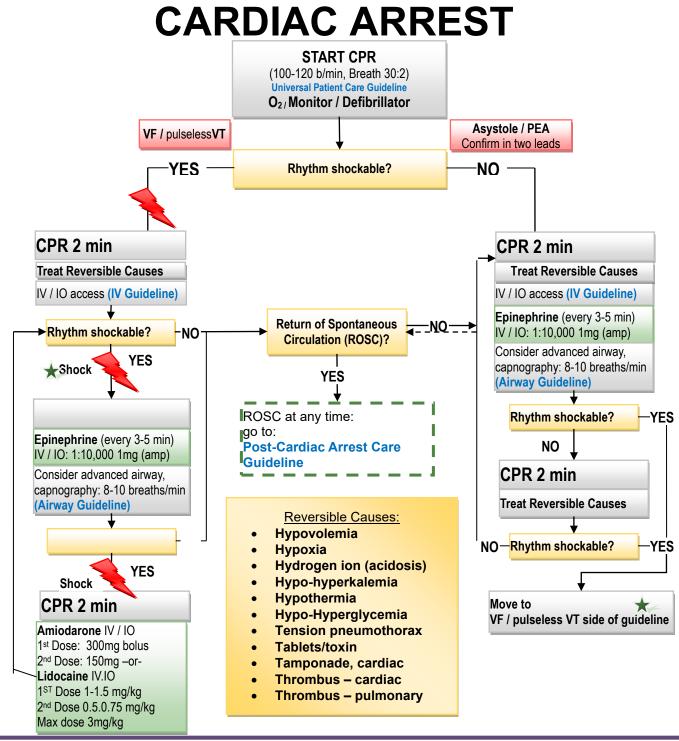
Continually monitor for any decompensation

BRADYCARDIA with PULSE



Pearls:

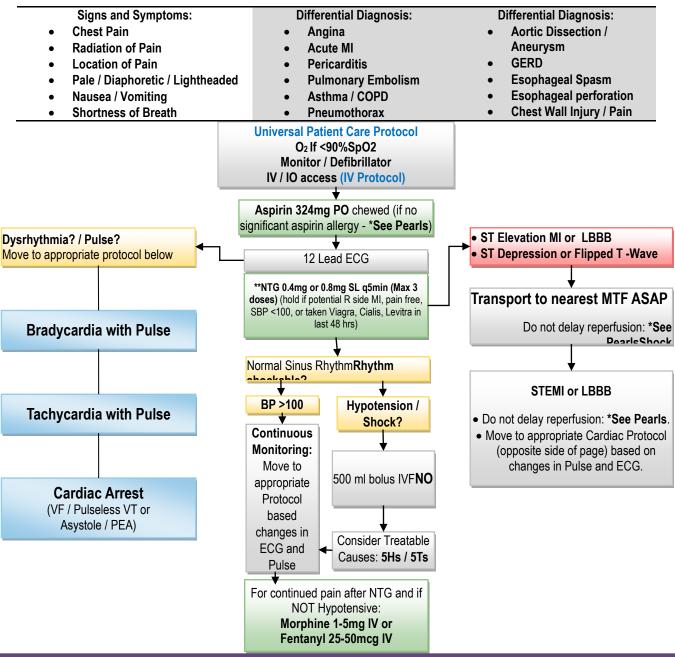
- Decompensation at any time (e.g., altered MS, hypotension) should prompt treatment as unstable patient.
- All bradycardic patients should have pacer pads in place after initial evaluation.
- Epinephrine infusion for refractory bradycardia: 2-10 mcg/min or 0.1-0.5 mcg/kg/minute (7 to 35 mcg/min in a 70 kg patient)
 - 1mg 1:10,000 in 250mL D5W / NS = 4 mcg/mL concentration
- Evaluate for treatable causes of bradycardia (B-blockade, Ca Channel blockade).



Pearls:

- Reversible causes should be addressed as soon as possible.
- Consider discontinuation of efforts if:
 - Asystole following trauma especially blunt.
 - Prolonged downtimes without resuscitative efforts > 15min.
 - Prolonged code with no response >3 rounds of medications, 30min of resuscitation.
 - <u>All patients should get a glucose check, at least 1L fluid bolus, and ultimately bilateral</u> needle decompression (especially in Trauma) before discontinuation of efforts.
 - Should take at least 1min to check for pulse in hypothermic patients.

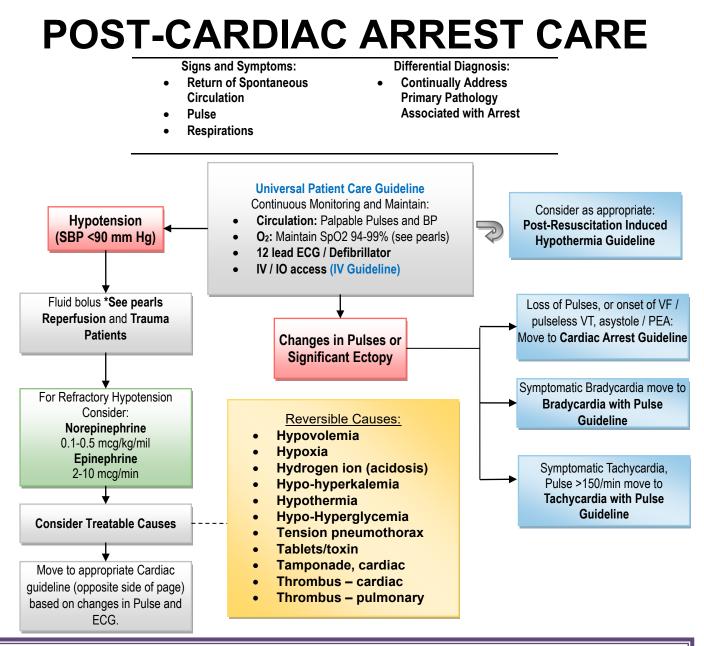
CHEST PAIN



Pearls:

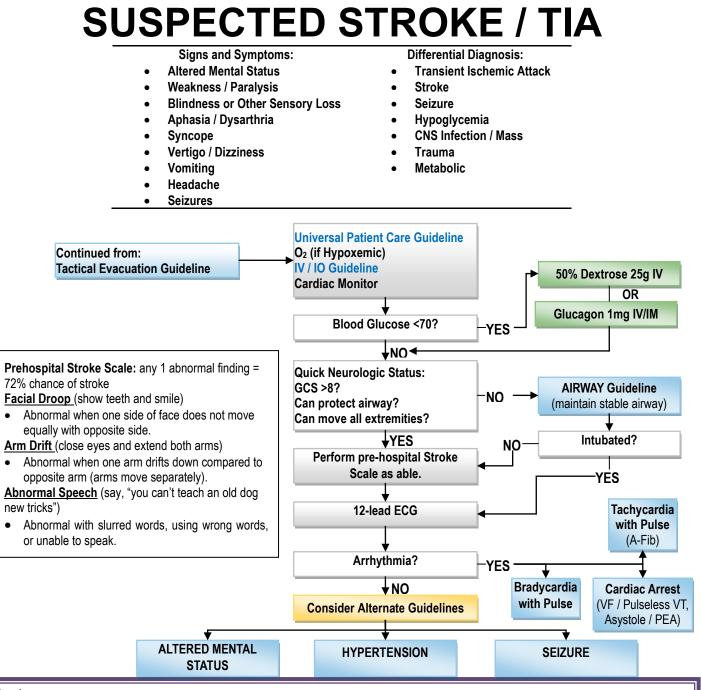
- Aspirin (4 x 81mg chewable) should be held only for patients with known significant allergy.
- Patients with suspected AMI should be transferred to the nearest MTF for further treatment / thrombolytics.
- **With right sided MI (ST Elevations in leads II, III, AvF), NTG may cause hypotension, use with caution. Add small fluid boluses for low BP.
- Ensure that you have IV access before giving SL NTG.
- Hold Morphine or Fentanyl for SBP <90.

Max dose Morphine 20mg, Fentanyl 200mcg for non-traumatic chest pain (higher doses may be required for trauma, see Pain Control algorithm).**CPR 2 min**



Pearls:

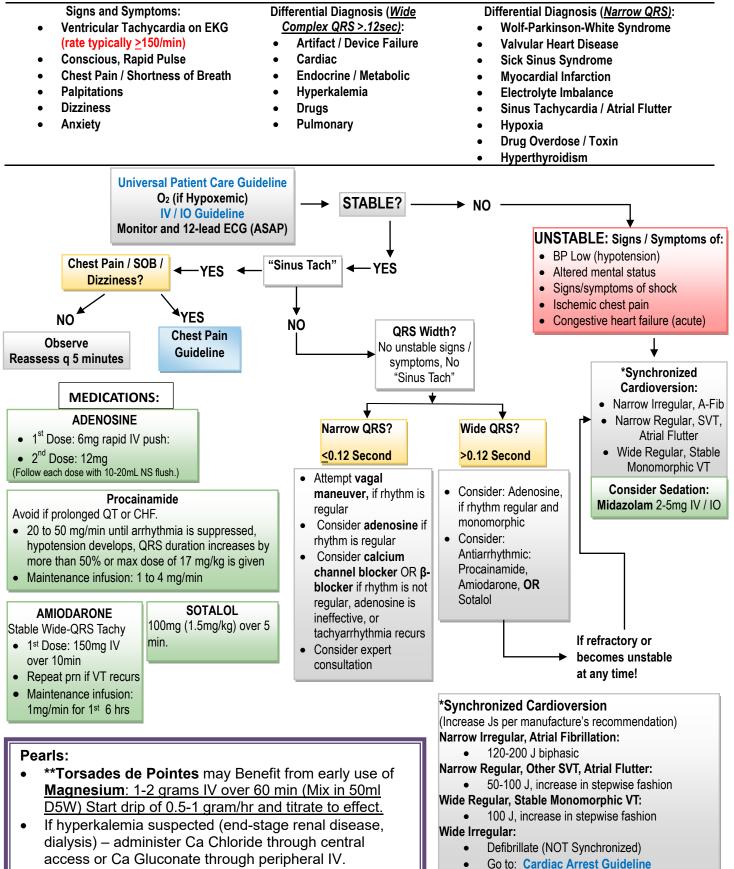
- Optimize ventilation and oxygenation. Place ETT, if not already done. Maintain SpO2 94-99% and ETCO2 35-45mmHg. Most patients will require ventilator assistance in the post-resuscitative phase.
- Hyperventilation may cause hypotension and/or recurrence of cardiac arrest in the postresuscitation phase and must be avoided.
- In non-airway controlled patients, it is important to prevent aspiration following resuscitation. For this reason, patients should be rotated onto their side (non-spinal immobilization) or be closely monitored in case vomiting occurs.
- *Reperfusion: 1-2 L IVF and consider use of a pressor IV / IO Drip EPINEPHRINE 2-10mcg/min or NOREPINEPHRINE 0.1-0.5 mcg/kg/min: 70kg adult: 7-35mcg/min.
 - **Dopamine should be started at a low dose (5mcg/kg/min)** and titrated up to maintain a SBP >90. The same applies to norepinephrine.
- ***Trauma patients** post-resuscitation should have fluid resuscitation consistent with hypotensive resuscitation guidelines. Maintain core temperature 32-36 degrees Celsius for at least 24 hours



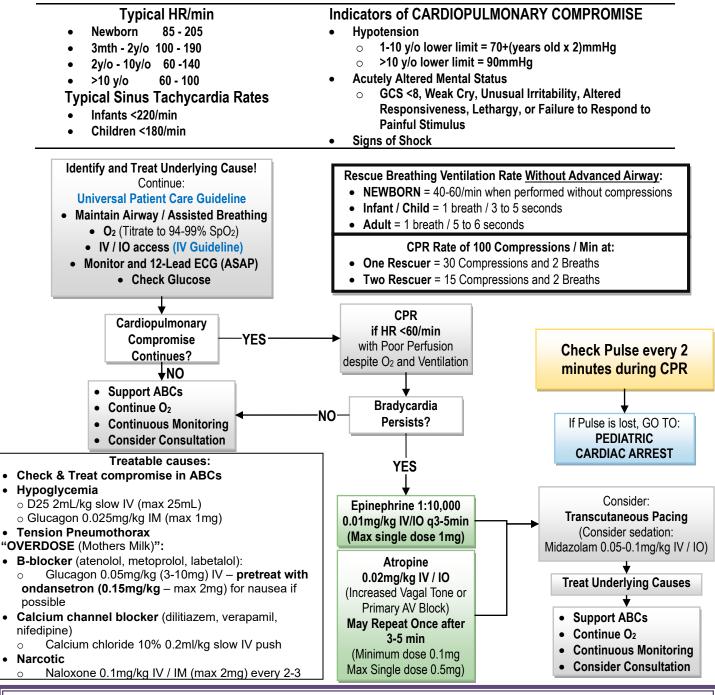
Pearls:

- Duration of symptoms should be determined as accurately as possible. Family members / colleagues can be helpful. If pt awaken with symptoms onset time est. from last time patient was seen "normal."
- Be alert for airway problem / risk of aspiration. If concerned, request intubation before departure.
- Hypoglycemia can mimic stroke / TIA. May present with focal neurologic deficit, especially in the elderly.
- EKG should be obtained in all patients to evaluate for arrhythmia especially atrial fibrillation.
- All TIAs should be transferred for evaluation, even if symptoms abated these patients have 10% risk of stroke within 30 days.
- Aspirin should not be given to patients for suspected stroke. Aspirin use is a contraindication to the use of thrombolytics for stroke.
- All strokes/TIAs are not associated with motor findings. Although uncommon, pure sensory strokes can occur. More frequently, very subtle motor abnormalities are present that the patient may not note.
- Systolic greater than 185 or Diastolic greater than 110: give Labetalol 10-20 mg IV for 1-2 minutes. May repeat 1 time.
- Aim for no more than a 20% reduction in MAP. MAP = [(2 x Diastolic) + Systolic] / 3 For additional info see: ALS Acute Coronary Syndromes and Stroke.

TACHYCARDIA w/ PULSE



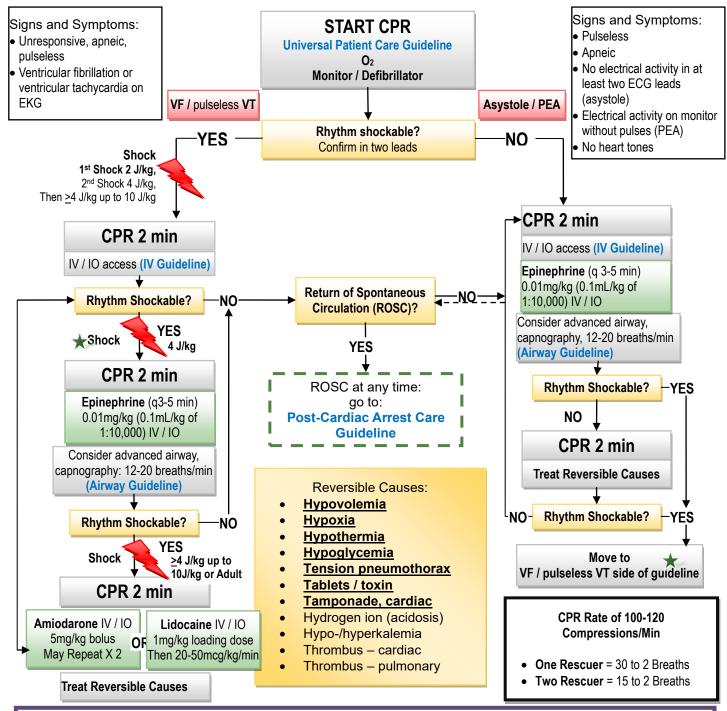
PEDIATRIC BRADYCARDIA with Pulse and Poor Perfusion



Pearls:

- Decompensation at any time (e.g., AMS, hypotension) should prompt treatment as unstable patient.
- All bradycardic patients should have pacer pads in place after initial evaluation.
- Evaluate for treatable causes of bradycardia (B-blockade, Ca channel blockade).
- The majority of pediatric cardiac problems are actually airway problems.
- In young, breast fed patients evaluate for mother's medications as they can cause toxicity in the infant.
- Pediatric pacer pads should be used if available. If only adult pads are obtainable they should be placed in the anterior-posterior position.

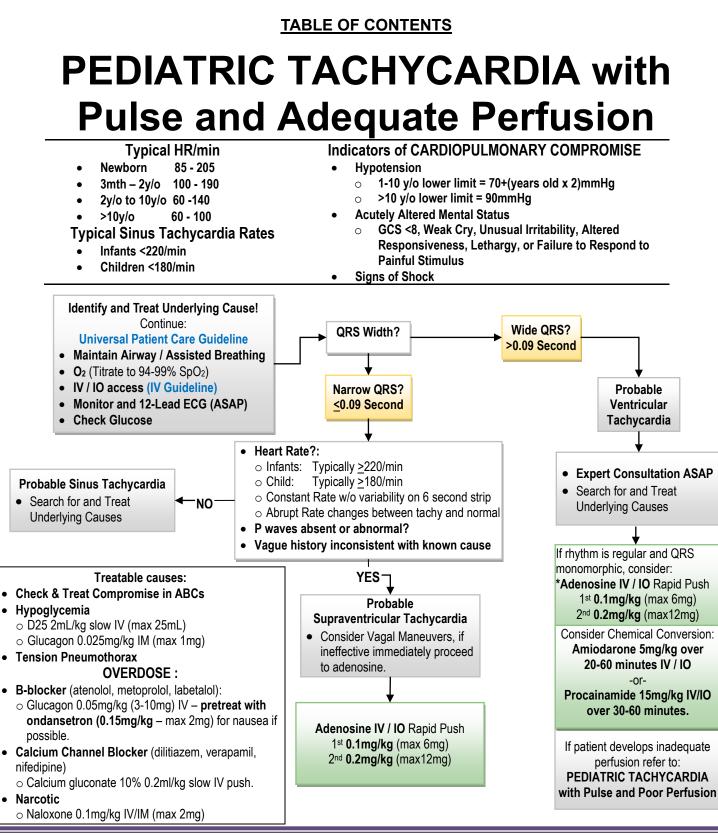
PEDIATRIC CARDIAC ARREST



Pearls:

CPR Rate of 100-120 / Min 2 inches depth for children and 1 1/2 inches depth for infants of chest with complete chest recoil
 Reversible causes should be addressed as soon as possible.

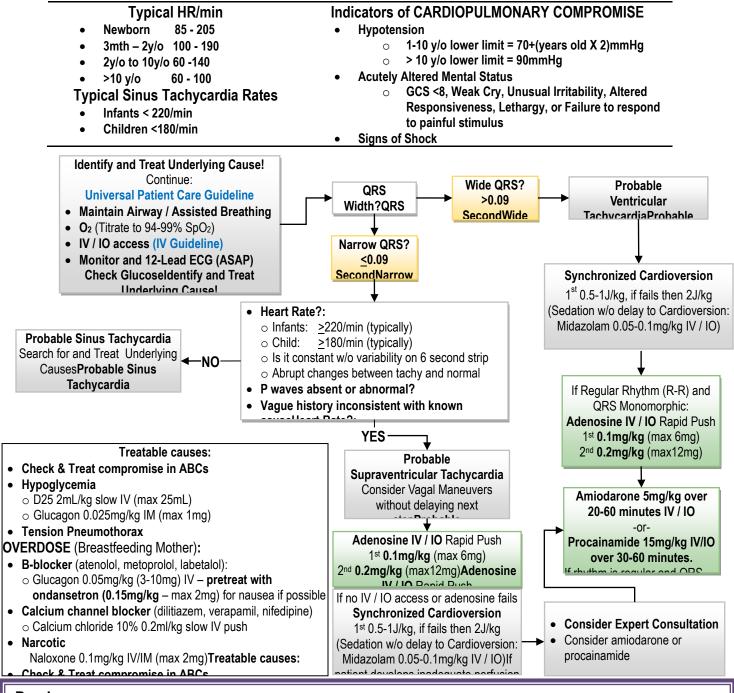
- Epinephrine Endotracheal Dose: 0.1 mg/kg (0.1mL/kg of 1:1,000 vial)
- Consider discontinuation of efforts if:
 - Asystole following trauma especially blunt
 - Prolonged downtimes > 15min
 - Prolonged code with no response >3 rounds of medications, 30min of resuscitation
 - <u>All patients should get a glucose check, at least 20ml/kg fluid bolus of NS, and ultimately</u> bilateral needle decompression (Trauma) before discontinuation of efforts



Pearls:

- **Vagal maneuvers:** blow through 18ga IV catheter, ice pack on forehead, carotid massage (unilateral only listen for bruits prior to performing), or having patient blow against closed glottis ("bear down").
- *Adenosine should be as central as possible with the "2 syringe technique" one with adenosine and the other with the saline flush. These should be attached to a 2 port IV adapter and flush should immediately follow drug.
- *Adenosine should be utilized in monomorphic and regular R-R interval type presentation.
- All patients should be warned of discomfort / feeling of heart stopping before adenosine administration.

PEDIATRIC TACHYCARDIA with Pulse and Poor Perfusion



Pearls:

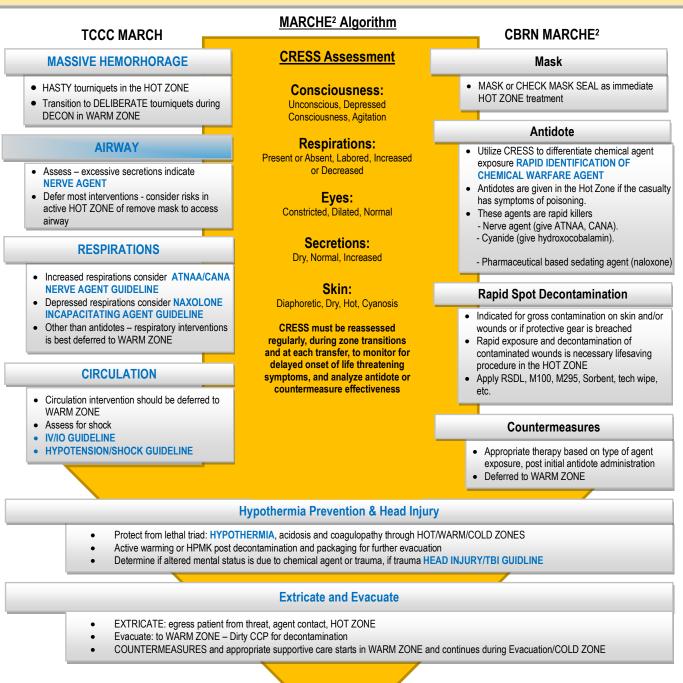
• Vagal maneuvers: blow through 18ga IV catheter, ice pack on forehead, carotid massage (unilateral only – listen for bruits prior to performing), or having patient blow against closed glottis ("bear down").

Adenosine should be given with the "2 syringe technique" – one with adenosine and the other with the saline flush. These should be attached to a 2 port IV adapter and flush should immediately follow drug.

All patients should be warned of discomfort / feeling of heart stopping before adenosine administration.

MARCHE²

After initial assessment of casualty in CBRN-threat environment for the presence or absence of CBRN symptoms using the CRESS algorithm, the integrated assessment and management of TCCC and CBRN injuries can proceed. MARCHE² integrates the TCCC MARCH algorithm with the priorities of CBRN treatment. MARCHE² is further broken down into phases similar to TCCC. The "Hot Zone" should be considered as care under fire, addressing only immediate life threats, "Warm Zone" is tactical field care and "Cold Zone" as tactical evacuation care.



Notes, Warnings, Cautions

• Treatment goals of CBRN is give antidote, extricate from exposure area, conduct spot decontamination, provide airway support.

CHILDBIRTH

Possible Complications Signs and Symptoms Preterm labor • Primi/Gravida/Para? . Placenta previa • Any pregnancy complications? • Prolapsed cord Vaginal fluid/bleeding? • Abnormal presentations (i.e., breech) Uterine contractions/back pain/stomach pain? • Spontaneous vaginal delivery (i.e. natural Duration of contractions and time between • outcome) Crowning/urge to push? Any complications expected with the newborn? Treatment Childbirth is a normal process and treatment is only supportive unless there are complications. • O2 if hypoxemic, IV/IO, cardiac monitor, and blood glucose check. Place in left lateral decubitus or pad under right hip.

- Hyper or hypotensive? Any abnormal bleeding? Refer to obstetric emergency.
- Visually inspect to see if patient is crowning, if crowning is present, assist with the birth of the child. If no, continue to monitor and transport patient to nearest MTF.
- Prep for birth: Position mother in supine supported position, prepare 2 sets of hemostats and scissors / scalpel, umbilical cord clamp if available, bulb suction.
- Suctioning of newborn nose and mouth with bulb aspirate recommended if obvious obstruction from secretions.
- Delivery: Use slight downward pressure to deliver superior shoulder, then slight upward pressure to deliver lower shoulder.
- Clamp cord after 1-3 minutes with 2 hemostats and cut between clamps. Wrap infant to prevent hypothermia and give to mother.
- Deliver placenta, do not pull. Keep placenta for evaluation by MTF.
- Externally massage uterus (through abdomen) to encourage contraction/limit bleeding.
- Continue to monitor and re-assess mother and neonate enroute to nearest MTF and refer to newborn guideline.

Notes, Warnings, Cautions

- If umbilical cord around neck, attempt to reduce manually prior to delivery of head (should feel rope-like structure around neck). As last resort, and if unable to keep pressure off of the cord, clamp and cut cord when unable to manually reduce.
- If umbilical cord seen, elevation of presenting part with vaginal hand and maintain elevation until delivery via C-section. *Do not place pressure on the cord or monitor pulse via the cord.
- If neonate appears to be stuck in the birth canal (i.e., turtling of the head), flex the mother's hips (both knees to chest).

NEWBORN CARE AND DISTRESS

Signs and Symptoms

- Full term delivery?
- Meconium staining of amniotic fluid?
- Any signs of Dehydration? (sunken fontanelles, tearless, decreased UOP, and dry mouth, skin and tongue)
- Fluid Overload? (SOB, ankle/sacral edema, increased JVP, and crackles in lungs

Determine after 1st 60 seconds of care and repeat q5 min.

Score of 6 or less=IMMEDIATE RESUSCITATION!

Severely depressed: 0-3 Moderately depressed: 4-6 Excellent condition: 7-10

| APGAR SCORING | 0 POINTS | 1 POINT | 2POINTS | | | | |
|----------------------------------|------------|--------------------------------|--|--|--|--|--|
| Activity (Muscle Tone) | Absent | Arms and Legs Flex | Active Movement | | | | |
| Pulse | Absent | Below 100 bpm | Over 100 bpm | | | | |
| Grimace (Reflex irritability) | Flaccid | Some Extremity Flexion | Active motion (<u>pull</u> away, cough) Completely Pink | | | | |
| Appearance (Skin color) | Blue, Pale | Body Pink, Extremities blue | | | | | |
| Respirations | Absent | Slow, irregular | Vigorous cry | | | | |

<u>Treatment</u>

- Does the patient have good tone? Is the airway open? (Breathing/Crying)
 - Use bulb syringe to clear mouth/nose, dry and stimulate (foot tap/back rub), keep warm, find APGAR score, and monitor SpO2 and treat hypoglycemia (glucose<40)
- Does the patient have a **HR <100**, apnea or gasping, labored breathing or persistent cyanosis?
 - Attempt to clear airway if needed, provide PPV via BVM: 30-60 breaths/min for a SpO2 of 94-99%
 - Intubate if NO chest rise
- Does the patient have a HR<60?</p>
 - Provide chest compressions and PPV (120 event/min: 90 compressions interspersed with 30 ventilations)
 - Consider intubation
 - Epinephrine 1:10,000 0.01-0.03mg/kg push q3-5min (0.1-0.3ml of 1:10,0000 10cc Cardiac Epi vial).
 - Consider hypoglycemia (D12.5 1/0ml/kg IV; dilute D50 to ¹/₄ strength or 1ml D50 in 3ml NS).
 - Consider Shock (IVF or Blood 10ml/kg IO).
 - Consider Pneumothorax (Intubation).

Notes, Warnings, Cautions

• If patient has meconium staining of amniotic fluid: suction mouth, then nose until clear with bulb syringe. Deep suction is no longer advised.

OBSTETRIC EMERGENCY

| Signs and Symptoms | Differential Diagnosis | | | | | | |
|--|--|--|--|--|--|--|--|
| Primi/Gravida/Para? | Pre-eclampsia/eclampsia | | | | | | |
| Any pregnancy complications? (i.e. pre- eclampsia, gestational diabetes, etc.) | Placenta previaAbruptio placentae | | | | | | |
| Any abnormal bleeding?Uterine contractions/back pain/stomach pain? | Spontaneous abortionUterine rupture | | | | | | |
| Blurry vision/dizziness?Changes in babies activity? | Ectopic pregnancy | | | | | | |
| Treatment O2 if hypoxemic, IV/IO, cardiac monitor, blood glucose check and place in left lateral decubitus or pad | | | | | | | |

- O2 if hypoxemic, IV/IO, cardiac monitor, blood glucose check and place in left lateral decubitus or pad under right hip
- Is the patient seizing?
 - Magnesium Sulfate 4g IV over 15 min or 5mg IM each buttocks
 - If blood glucose is WNL and patient is in Status Epilepticus move to (Seizure Guideline)
 - Midazolam 5mg IV/IO or 10mg IM or Lorazepam 2-4mg IV/IM
 - Wait 60 seconds and if not resolved, give an additional dose, if not resolved after second dose move to Altered Mental Status Guideline.
 - If blood glucose is <70 or >250 move to Altered Mental Status Guideline.
- Are they hypertensive? (>/=160/110 or 140-159/90-109) with severe headache, blurred vision, photophobia, hyperreflexia, epigastric pain?
 - Magnesium Sulfate 4g IV over 15 min
- Is the patient experiencing abdominal pain alone refer to Abdominal Pain Guideline.
- Is there vaginal bleeding present? If tachycardic/orthostatic administer blood if available or 2G TXA IV/IO and then 1000ml IVF IV/IO bolus
 - If vitals WNL, is there s/sx of labor? Refer to Childbirth Guideline.
- Continue to monitor, re access and address for changes in BP, seizures, glucose, vision changes and headaches.

Notes, Warnings, Cautions

- Use caution when using magnesium it can lead to cardiorespiratory collapse with hypotension and decreased respiratory drive.
- Treat all hypertensive patients as if they are pre-eclamptic despite any prior history of hypertension.
- The leading cause of Postpartum Hemorrhage is Uterine Atony (lack of contracting), which can be treated with uterine massage
- Seizure / headache / vision complaints: can give Midazolam 0.1mg/kg IV every 15-30 or 1mg IV every2-3min up to 5mg while waiting for magnesium to take effect.
- Seizure activity in an OB patient signifies eclampsia.
- The best life support for the fetus is to resuscitate the mother.

PATIENT REFUSAL

Indication: If a patient (or person[s] responsible for a minor) refuses treatment or transport, after pre-hospital providers have arrived on the scene, the following procedures should be carried out:

<u>Treatment</u>

- Complete primary assessment, obtain set of vitals and determine mental status.
- Any injuries or illnesses found to immediately threaten life, limb, or eyesight (or can be assumed will deteriorate enroute) should be addressed and treated immediately.
- Patients that prevent treatment of these injuries, should be encouraged. Any doubt of capacity should prompt treatment/transport under implied consent. Patient with decision- making capacity refusing treatment of life-threatening injury or illness require further clinical judgement and consultation with medical director prior to informed refusal.
- Injuries or illnesses that do not represent imminent threats to life, limb, or eyesight (or considered unlikely to deteriorate enroute) may be addressed in accordance with the following:
- Determine decision making capacity (patient/parent).
- Is the patient AOx4 and understands issues with their medical condition/status? If no, treat IAW Altered Mental Status Guideline.
- Clearly and repeatedly explain to the patient/parent of concerns and risks of refusal
- Document all findings and discussions with the patient/parent regarding refusing treatment and/or transportation.
- Obtain a signature from a witness (crewmember) and the patient/parent/parties responsible for the patient as to refusal of care.
- See Sample Patient Refusal Form in handbook.

Notes, Warnings, Cautions

• Clearly explain to **Military Personnel** why the treatment is needed. Notify them that refusal of treatment may bring judicial or administrative adverse action upon them under UCMJ.

MWD AIRWAY MANAGEMENT

| <u>History</u> Dyspnea Labored breathing Stridor Stertor Altered level of consciousness Trauma to the airway Disruption of mouth, pharynx, larynx, or trachea | Differential Diagnoses• Upper airway obstruction (foreign body)• Laryngeal paralysis• Pneumo/hemo/pyothorax• Diaphragmatic hernia• Pleural effusion• Pulmonary contusions• Pulmonary edema• Pneumonia | | | | | |
|---|---|--|--|--|--|--|
| | tment | | | | | |
| Inspect, wipe and/or suction mouth and pharynx. Ventilate with 100% oxygen. If unable, move to next step. Endotracheal Intubation (Size 9-11) If unable, move to next step. Suction Airway If airway still not clear, move to next step. Perform Tracheostomy Evaluate for pleural space and parenchymal problems. Notes, Warnings, Cautions Unconscious MWDs: Use tracheal insufflation, orotracheal intubation, or tracheostomy. If there is an obstruction, then bypass the obstruction until the patient is more stable. NOTE: intubation of the MWD is most easily performed with the dog in sternal or prone position, head and neck extended, and tongue pulled forward. Verify placement by palpating neck for 1 tube. If 2 tubes are felt, the tube is in the esophagus. Capnometer reading > 10mmHg also ensures correct placement. | | | | | | |
| 100% Oxygen Supp | lementation Examples | | | | | |
| | muzzled dogs (10-15 L/min) or tracheostomy (2 L/min) | | | | | |

MWD ANALGESIA & SEDATION

Indications

- Trauma or painNeed for chemical restraint
- Reed for chemical resulta
 Continued sedation
- Anxiety
- Fractious

Indications

- Mild Sedation
 - Relax MWD for examination, handling, reducing anxiety
 - MWD will be calm but still reactive to noise and stimulation
- Deep Sedation
 - First line protocol for fractious MWD
 - MWD will not be able to walk, may be aroused with stimulation and may maintain laryngeal and palpebral reflexes

<u>Treatment</u>

Mild Sedation

- Midazolam 0.3 mg/kg IM/IN and Hydromorphone 0.2 mg/kg or Morphine 0.2mg/kg IM/IN
- Deep Sedation
 - o Midazolam 0.3 mg/kg AND Ketamine 5 mg/kg
 - Hydromorphone 0.1 mg/kg IM or Morphine 0.1 mg/kg IM IN

Analgesia

- o Intermittent IV or IM supplementation
 - Hydromorphone 0.1-0.2 mg/kg (q2-4 hrs)
 - Morphine Sulfate 0.2-0.5 mg/kg (q4-6 hrs)
 - Fentanyl 2-10 mcg/kg (once as loading dose then must consider CRI duration ~ 30 minutes).
- Continuous Rate Infusion (CRI)
 - Fentanyl 2-10 mcg/kg/hour
 - Morphine 0.1-0.25 mg/kg/hour
 - Hydromorphone 0.02-0.05 mg/kg/hour
- Opioid Reversal
 - NALOXONE 0.01-0.02 mg/kg slow IV to effect if needed will reverse analgesia AND sedation

Notes, Warnings, Cautions

- Dosages for analgesics in dogs are significantly higher than for people.
- Opioids cause emesis, usually within 5 minutes of administration. Be prepared to remove the muzzle to minimize aspiration risk. Hydromorphone causes excessive panting; use caution with head injuries, gastric dilatation and volvulus (GDV) and respiratory disease.
- **CAUTION**: Do NOT use acetaminophen or ibuprofen in MWDs, as these drugs can cause liver toxicity. AVOID use of NSAIDs such as naproxen and aspirin in emergently ill or injured MWDs.
- Note that all protocols have analgesia incorporated into them. Additional analgesia can be provided by the IV/IM or PO route, as necessary.

MWD CPR

| Indications to Initiate CPR Pulselessness Apneic | History <u>Cardiopulmonary Arrest Confirmed</u> Causes: Traumatic Blast Blunt force Penetrating Non-traumatic Anesthesia Near-drowning Electrocution | | | |
|---|---|--|--|--|
| | Treatment | | | |
| Drug therapy if defibrillation not success Epinephrine 0.01 mg/kg IV/IO or Vasopressin 0.8 U/kg IV/IO once an Amiodarone 5-10 mg/kg IV/IO Asystole/Bradycardia/PEA Drug therapy Atropine 0.04 mg/kg IV/IO (only if br Epinephrine 0.01 mg/kg IV/IO and V | ucted. 00% O2 cycle (2 min) ression cycle between each defibrillation. sful d Lidocaine 2 mg/kg IV/IO or radycardia preceded arrest) /asopressin 0.8 U/kg IV/IO once | | | |
| 70% of MWDs that arrest will have PEA, as | <u>arnings, Cautions</u> systole, or sinus bradycardia as the initial arrest rhythm. is for these rhythms or for empiric use if ECG capability is | | | |
| | key to successful resuscitation is to SUSTAIN chest | | | |

- Avoid interrupting chest compressions! The key to successful resuscitation is to SUSTAIN chest compressions aggressively for 2-3 minutes before stopping to check status.
- Most people apply too little force when performing chest compressions! Do not be concerned with breaking ribs or injuring the heart or chest with BLS. In contrast to CPR in people, the thorax of MWDs is more compliant and fractures are rare

MWD GASTRIC DILATION-VOLVULUS

<u>History</u>

- Abdominal distention/tympany
- Non-productive retching
- Attempted vomiting without result
- Pain when palpating stomach/abdomen.
- Inability/reluctance to lay comfortably.
- Anxiety
- Signs of compensatory shock

Definition

- GDV is a rapidly life-threatening condition common in MWDs. In GDV, the stomach rapidly dilates (gastric dilation) with fluid, food, and air, and then rotates along the long axis (volvulus) and causes shock by interfering with venous return from the abdomen and pelvic limbs.
- GDV is a surgical emergency.

<u>Treatment</u>

- Initiate Monitoring: ECG, NIBP, SPO2, ETCO2
- Supplemental O2
- Attain (2) IV/IO sites FORELIMBS.
 - **Remember:** venous return is impeded from pressure in the abdomen. Hindlimb IVs will not be effective.
- Initiate IV/IO crystalloid therapy FIRST, repeat bolus every 10 20 minutes up to 4 times over the course of an hour.
 - For quick reference, ADD a ZERO to the dog's body weight (in pounds) to approximate a safe but effective bolus volume. For example, a 45# dog would need about a 450 mL bolus, and a 75# dog would need about 750 mL as a bolus.
- If refractory to crystalloids: Give Hydroxyethyl starch (HES, "Hetastarch", "VetStarch") bolus (10-20 mL/kg) to maintain BP. **Repeat** this bolus once if no response to therapy.
- If refractory to HES: Give hypertonic saline (HTS) IV bolus of 4 mL/kg over 5 minutes (if 7-7.5% HTS is available) for MWDs that fail to respond to two or three quarter-shock boluses of crystalloids and/or one or two boluses of HES.
- Decompress the Tympanic Stomach
 - Position self on left side, or lay dog on left side
 - Palpate last rib, move hand two inches caudal to the last rib, midway between the spine and the ventral border of the abdomen on the right side.
 - Forcefully insert 14-18 gauge IV over-the-needle catheter through the skin, abdominal wall, and stomach wall
 - Note gas or air escaping through the needle from the stomach to signify a successful attempt. If no gas or air, attempt once more.
 - o (DO NOT ATTEMPT THIRD if unsuccessful)
 - Apply gentle external pressure to abdominal wall to assist exiting air.
 - \circ $\;$ Remove catheter once air is evacuated.
- Provide analgesia.

Notes, Warnings, Cautions

- Goal is to treat for shock, decompress stomach, and transport for surgical intervention.
- Monitor for ventricular arrhythmias, persistent shock and recurrent dilation.
- Surgery is REQUIRED for definitive treatment to de-rotate the stomach.

MWD HEAT INJURY

MILD Heat Injury

(heat stress) - excessive thirst, discomfort associated with physical activity, mild dehydration, <u>but with</u> <u>controlled panting</u> (i.e., the patient can control or reduce panting when exposed to a noxious inhalant such as alcohol).

MODERATE Heat Injury

(heat exhaustion) - heat stress present, as well as weakness, anxiety, and <u>uncontrolled panting</u> (i.e., the patient cannot reduce panting when exposed to a noxious inhalant), but central nervous system (CNS) abnormalities are not present.

SEVERE Heat Injury

(heat stroke) – heat exhaustion are present, coupled with varying degrees of CNS abnormalities (changes in mentation and level of consciousness, seizures, abnormal pupil size, blindness, head tremors, and ataxia

<u>Treatment</u>

- Mild: Heat Stress
 - Remove patient from source of heat, discontinue exercise, cool by fans or air condition, give cold water to drink.
- Moderate: Heat exhaustion
 - Remove patient from all heat and stop all activity
 - Cool by fans or air condition. Thoroughly soak the hair coat to the skin (room-temp) in order to reduce core body temperature.
 - Give IV fluids 3-5 mL/kg/hr if not in shock
- Severe: Heat Stroke
 - o Remove patient from all heat and stop all activity
 - Establish airway, provide oxygen, establish IV for shock treatment.
 - Aggressively cool patient until rectal temp is less than 105°F. Use only room temperature fluids. Give IV fluids (shock protocol)
 - Monitor patient for vitals, blood glucose, ECG arrhythmias, Mentation / LOC, gait abnormalities, vision changes, seizures, rebound hypothermia

Notes, Warnings, Cautions

- **PANTING** is the only significant cooling mechanism for dogs.
- **NO** specific body temperature defines heat stroke in MWD's. Normal rectal temperature is 99.5° to 103° F in the MWD. Temperatures as high as 105.8°F have been associated with pathology. Most commonly, heat stroke is seen in MWDs with rectal temperatures greater than 107°F.
- **<u>DO NOT</u>** use of cold intravenous fluids, ice packs, or ice-water baths for cooling.
- Once the MWD's body temperature is = 103°F <u>CEASE</u> all cooling efforts and monitor for rebound hypothermia and prepare for rewarming measures. Actively warm the dog if the temperature <100°F
- Treat seizures with midazolam or diazepam 0.3 mg/kg IV, IO or intranasal prn
- MWDs are commonly have prolonged clotting times, and platelet abnormalities following heatstroke. Monitor for bleeding and disseminated intravascular coagulation. Given lack of canine blood products, any MWD with evidence of bleeding should be evacuated URGENTLY to a veterinary facility.

MWD NORMAL PARAMETERS

- Temperature (rectal)- 99.5° to 103° F (working temperatures up to 107° F)
- Heart Rate/ Pulse- 60 to 120 bpm
- Respiratory Rate- 16 to 32 bpm (Controlled panting is normal)
- Blood Pressure- Systolic 120 mmHg, Diastolic 80 mmHg, Mean 90 to 100 mmHg
- Average MWD weighs 30-35 kg (German shepherd dogs, Belgian Malinois, Labrador retrievers).
- <u>IV catheterization</u> access points are:
 - Cephalic vein on the cranial (superior) aspect of the forearm (figures 1 & 2)
 - Lateral saphenous vein on the lateral aspect of the hind limb at the distal tibial area (figure 3)
- <u>IO catheterization</u> access points are:
 - o Greater trochanter of the humerus
 - Medial tibia just distal to tuberosity (figure 4)
- <u>Arterial Pulse</u> is palpated at the femoral artery on the medial aspect of the proximal thigh in the inguinal area (figure 5) or at the dorsal metatarsal artery on the dorsal aspect of the proximal hind paw.
- <u>Heart sounds</u> are best auscultated over the lower left lateral thoracic wall between the 4th and 5th intercostal space.
- 3-lead <u>electrocardiograms</u> are sufficient for MWDs. Adhesive electrodes should be taped to the pads of the paws of the left forelimb (<u>black</u> lead), right forelimb (<u>white</u> lead), and left hind limb (<u>red</u> lead). (figure 6)
- <u>Pulse oximetry</u> probes can be utilized on conscious dogs using the ear pinna, lip fold, or flank skin; while not optimal for oximetry, these alternative sites are generally acceptable. For optimal reliability place probe on tongue (only in unconscious dogs)

All drug dosages should be calculated based on measured or estimated body weight. DOG HANDLER CARRIES DRUG CARD FOR THE DOG

For further reference see Clinical Practice Guidelines for Military Working Dogs, 12 Dec 2018

MWD NORMAL PARAMETERS



Figure 1- Vein best punctured on superior surface of limb toward the elbow



Figure 2- Vein occlusion proximal to elbow joint while elbow is in extension



Figure 3- lateral saphenous vein on the hind leg



Figure 5- location for palpation of the femoral arterial pulse

Figure 6- placement of adhesive ECG electrode pads on the footpads

MWD SHOCK FLUID THERAPY

| Indications ● Hypotension ○ Systolic < 90 mmHg ○ MAP < 65 mmHg ● Hypovolemia ○ Massive hemorrhage (external, cavitary) ○ Severe dehydration (heat injury, GI loss) ○ Cavitary pressure impeding arterial perfusion or venous return | <u>Clinical Signs</u> Early, compensatory shock Tachycardia, tachypnea, alert mentation, rapid pulse, normal to decreased pule quality, decreased CRT (<2 sec), normal to bright red MM. Late, decompensatory shock Bradycardia, prolonged or poor CRT (>2 seconds), pulses poor or absent, hypothermia, stupor | | | | | | |
|--|---|--|--|--|--|--|--|
| Treatment | | | | | | | |
| Attain multiple IV/IO sites. Calculate total fluid volume required (90 mL/kg) in one hour. Or see Notes below Give a Hydroxyethyl starch (HES) IV/IO bolus 10-20 mL/kg over 5-10 minutes if clinical signs of shock do not abate after the first 30 minutes (first 2 quarter-shock IV challenges) of crystalloid fluids, or response to crystalloid challenges is not sustained. Repeat this bolus if no response to therapy. Give a Hypertonic saline (HTS) IV/IO bolus 4 mL/kg over 5 minutes (if 7-7.5% HTS is available) for MWDs that fail to respond to two or three quarter-shock boluses of crystalloids and/or one or two | | | | | | | |
| boluses of HES. | V over 15 min but NOT LATED THAN 3 HOUDS need | | | | | | |

 Consider TXA 10 mg/kg in 100 mL NS or LRS, IV over 15 min but NOT LATER THAN 3 HOURS post injury.

Notes, Warnings, Cautions

- Quick calculation for a bolus shock dose: ADD a ZERO to the dog's body weight (in pounds) to approximate a safe but effective bolus volume. For example, a 45# dog would need about a 450 mL bolus, and a 75# dog would need about 750 mL as a bolus.
- **CAUTION**: Human blood products and albumin, or other animal blood products, must never be given to dogs, given the high risk of anaphylactic reactions
- Blood product transfusions for MWDs are ONLY available from Veterinary Service Support units and their administration is only authorized under the direct supervision of a veterinarian
- Clinical target for resuscitation end point is a mean arterial pressure (MAP) of > 65 mmHg or a systolic of > 90 mmHg. Neonatal or pediatric blood pressure cuffs must be used.

ANTIBIOTIC THERAPY CHART

*Post-injury antimicrobial agents are recommended to prevent early post-traumatic infectious complications, including sepsis, secondary to common bacterial flora. Selection is based on narrowest spectrum and duration required to prevent early infections prior to adequate surgical wound management. This narrow spectrum is selected to avoid selection of resistant bacteria. The antimicrobials listed are not intended for use in established infections, where multidrug-resistant (MDR) or other nosocomial pathogens may be causing infection.

| | -resistant (MDR) or other nos | | | | |
|--|---|-------------------------------|--|--|--|
| INJURY | PREFERRED AGENT | FREQUENCY | DURATION | | |
| | EXTREMET | YWOUNDS | | | |
| Skin, soft tissue, without open fractures | Cefazolin 2g | q 6-8hrs | 24 hours | | |
| Skin, soft tissue, with open fractures, exposed bone, or open joints | Cefazolin 2g | q 6-8hrs | 24 hours, then with each subsequent I&D until soft tissue coverage | | |
| | THORACIC | WOUNDS | | | |
| Penetrating chest injury | Cefazolin 2g | q 6-8hrs | 24 hours | | |
| | ABDOMINA | LWOUNDS | _ | | |
| Penetrating abdominal injury with suspected/known hollow viscus injury and soilage; may apply to rectal/perineal injuries as well | Cefazolin, 2g IV PLUS metronidazole 500mg IV | q 6-8hrs q 8-12hrs | Stop 24 hours after control of contamination | | |
| | MAXILLOFACIAL AN | ID NECK WOUNDS | | | |
| Open maxillofacial fractures, maxillofacial fractures with foreign body or fixation device | Cefazolin 2g g | q 6-8hrs | 24 hours | | |
| | CENTRAL NERVOUS | SYSTEM WOUNDS | | | |
| Penetrating brain injury | Cefazolin, 2g IV PLUS (consider) | q 6-8hrs | 5 days or until CSF leak is closed, whichever is longer | | |
| Penetrating spinal cord injury | metronidazole 500 mg Cefazolin, 2g IV PLUS (consider) | q 8-12 hrs q 6-8hrs | | | |
| injury | metronidazole 500 mg | q 8-12 hrs | | | |
| | EYE WO | DUNDS | | | |
| | Erythromycin ophthalmic ointment | q 6hrs | Until epithelium healed. | | |
| Eye injury, burn or abrasion | Or Bacitracin ophthalmic ointment | or PRN for symptomatic relief | No systemic treatment required | | |
| Eye injury, penetrating | Levofloxacin 750 mg IV/PO PLUS | q 24 hrs | 7 days or until evaluated by an ophthalmologist. | | |
| | vancomycin 15-20 mg/kg IV | q 8-12hrs | No topical agents. | | |
| | BUF | RNS | | | |
| Pre hospital | Not indicated | | | | |
| | DELAYED EVACUATION | N TO SURGICAL CARE | | | |
| PO tolerable | Moxifloxacin 400 mg PO x 1 dose. | X 1 dose | Single dose therapy | | |
| Not PO tolerable | Or Ertapenem 1 g IV/ IM | X 1 dose | | | |

| COMMON LABORATORY VALUES | | | | | | | |
|--|---|--|---|--|--|--|--|
| LABORATORY CHEMISTRY CONVENTIONAL SI UNITS | | | | | | | |
| | 8-16 mEq/L | 8- | 16 mmol/L | | | | |
| | 8-25 mg/100mL | 2.9-8/9 mmol/L | | | | | |
| | 8.5-10.5 mg/100mL | 2.1-2.6 mmol/L | | | | | |
| | 24-30 mEq/L | 24-30 mmol/L | | | | | |
| Male | 0.2-0.5 mg/dL | Female 0.3-0.9mg/dL | | | | | |
| Male | 17-40 U/L | Female | 10-79 U/L | | | | |
| | 0.6-1.5 mg/100L | | 53-133 | | | | |
| | 70-110 mg/100mL | | -5.6 mmol/L | | | | |
| | 135-145 | 5 mEq/L | | | | | |
| | 3.5-5.0 mEq/L | 3.5- | -5.0 mmol/L | | | | |
| | | | | | | | |
| | HEMATOLOGY | | | | | | |
| Male | 13-18 g/100 mL | Female | 12-16 g/100mL | | | | |
| Male | 41-50% | Female | 36-44% | | | | |
| 140,000-450,000/ml | | | | | | | |
| | | | | | | | |
| C | ARDIAC MARKERS | | | | | | |
| Onset | 4-6 hrs. | Peak | 12-24 hrs. | | | | |
| Onset | 3-4 hrs. | Peak | 10-24 hrs. | | | | |
| Male | 10-95 ng/ml | Female | 10-65 ng/ml | | | | |
| Onset | 1-3 hrs. | Peak | Peak: 6-10 hrs. | | | | |
| | 0.8-1.2 | | 2.0-3.0 | | | | |
| Ily sensitiv | e. Each device has different reference | ranges associated | . Correlate cTn with | | | | |
| | MAL BLOOD GASSES | | | | | | |
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| | CHEN Male Male Male Male Onset Onset Male Onset Male | CHEMISTRY CONVENTIONAL 8-16 mEq/L 8-25 mg/100mL 8.5-10.5 mg/100mL 24-30 mEq/L Male 0.2-0.5 mg/dL Male 17-40 U/L 0.6-1.5 mg/100L 70-110 mg/100mL 135-145 3.5-5.0 mEq/L HEMATOLOGY Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 13-18 g/100 mL Male 10-95 mg/ml Onset 4-6 hrs. Onset 1-3 hrs. Onset 1-3 hrs. O.8-1.2 Male sensitive. Each device has different reference | CHEMISTRY CONVENTIONAL S 8-16 mEq/L 8- 8-25 mg/100mL 2.9- 8.5-10.5 mg/100mL 2.1- 24-30 mEq/L 24- Male 0.2-0.5 mg/dL Female Male 17-40 U/L Female 0.6-1.5 mg/100L 70-110 mg/100mL 3.9- 135-145 mEq/L 3.5- Male 13-18 g/100 mL Female Male 13-9 Female Male 13-9 Female 0.8-1.2 0.00/ml Female Onset 3-4 hrs. Peak 0.8-1.2 0.8-1.2 0.8-1.2 Illy sensitive. Each device has different reference ranges associated nsitive. 7.35-7.45 35-45 mm Hg | | | | |

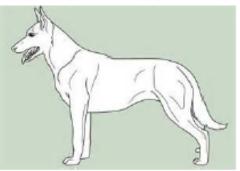
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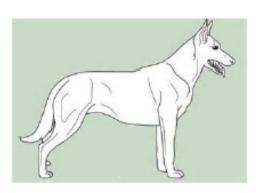
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| Last | / | | 0 | D | | | |
| PERRLA 🕅 R Siz | ze (mm) | L Size (mm) | | | | | |
| Field Ultrasound Re | sults | | | Other Diagnostics | | | |
| Additional Interven | - | | | | | | |
| | Time | | | Time | | | |
| Foley | Comment | | Gastric | | Dral 🗌 Nasal Com | ment | |
| | | | | | | | |
| Protection | | Protective Eyewear | | | | | |
| Immobilization | C-Collar | C-Spine 🗌 Spine Bo | oard 📃 Pelvic Spli | nt 🔄 Pelvic Binder | , Туре | | |
| _ | 🔄 Splint, Type | /Location | | | | | |
| Warming | Hypothermi | a Prevention, Product | | | | | |
| | | a Prevention, Product | | | | | |
| Other | | | | | | | |
| Interventions — | | | | | | | |
| _ | | | | | | | |
| Medications and Fl | uids | Route = IM, II | N, IO, IV, PO, PR, SL, SQ | Medications and F | luids | Route = IM, IN | , 10, IV, PO, PR, SL, SQ |
| Time Drug / Flui | d | Dose | Route | Time Drug / Flu | id | Dose | Route |
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| Documents Receive | ed 🗌 TCCC Card [| Patient Chart 🗌 N | lone Other | | | | |
| Narrative Summary | of Care | | | | | | |
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| Enroute Care Provid Last Name | | Name | Rank Capa | ability Signature | 2 | | |
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| | a.jbsa.healthcare-ops. | list.jts-prehospital@hea | ith.mil | T | | MM (|) |
| PREPARED BY | | | | DEPARTMENT/SERV | ICE/CLINIC (Treating U | Init) DATE | |
| (Signature & Title) | | | | | | | |
| PATIENT'S IDENTIFI | CATION (Name: last fin | st, middle; grade; date; hos | pital or medical facil | itv) | I | | |
| | • | · · · · | | () | N/I | | |
| | | First Name | | | ···· — | DIAGNOSTIC STUDIES | FLOW CHART |
| BR# Ra | ank Unit | | | Pt Cat | | OTHER EXAMINATION OR E | EVALUATION |
| SSN | DOB | Gender () M (|) F Allergy | Other | | OTHER, Specify | |

| TACTICAL EVACUATION-AFTER ACTION REPORT & PATIENT CARE RECORD Page 3 | | | | | | | |
|---|-------------|---------------------|----------------------|--------------------------|------------------------|---|---|
| IAW AR 4 | 40-68 (RAR) | 22 May 2009 Paragra | - | ge is a quality a | | | edical records. |
| | | | - | | nt (Check all wor | | |
| Helmet, Ballistic | | Plate Front | Neck Protect | | Groin Shi | | Blast Gauge |
| Tactical Vest (107 | | Plate Back | Throat Prote | , , | | dergarment Tier 1 | Blast Sensor Helmet |
| Eye Protection | | Plate Right Side | Deltoid Righ | it | Pelvic Un | dergarment Tier 2 | Blast Sensor Other |
| Ear Protection | | Plate Left Side | Deltoid Left | | | | |
| AR Discussion | Event Date | [] T | actical situation co | mplicated care <i>(E</i> | Explain in discussion) | | |
| | Sı | ustains | | | | Improves | |
| ast Name R# Rank _ | χ , | | e | facility) Pt Cat | MI | Law (PL) No. 99-661), se document was created by and is confidential and guidance predicated on disclosure of, or testim recommendations, evalua of a QA program except i | horization Act for fiscal year 1987 (P ction 1102, Title 10, (10USC 1102) or for the DOD in a medical QA prog privileged. PL 99-661 and subseq this law (10 USC 1102) prec ony about, any records or find tions, opinions, or actions taken as n limited situations. Under the provis formation is exempt from relace |
| SN | DOB | Gender 🔘 | M () F Allergy | Pt Cat Other | | of 10 USC 1102, this accordance with Exempt | nformation is exempt from release on 3 of the FOIA. Additional deta confidentiality of QA documents a |

| CANINE-TA | CTICAL COMBAT | CASUALTY | CARE CARD | <u>(cTCCC)</u> |
|--------------|-----------------------------|-------------|-------------------|----------------|
| EVAC CAT: | Urgent Priorit | y 🗌 Routine | ; | |
| EVAC TYPE: [| Fixed Rotary | Ground M | EDEVAC C | ASEVAC |
| UNIT: | NAME: | | TATTOO: | |
| DATE: (DD-MM | -YY) | TIME: | GENDER | : 🗆 M 🗆 F |
| Mechanism of | Injury: (Mark X all that ap | ply | | |
| | | | ARTILI FRY | |
| | | | | |

Injury: (Mark all injuries that apply with an X)





Signs and Symptoms: (fill in the blank)

| Time | | | |
|-----------------------------|--|--|--|
| Pain Score (0-10) | | | |
| Temperature (99-102.5) | | | |
| Pulse Rate/Location (60-80) | | | |
| Respirations (16-30) | | | |
| Blood Pressure (120/80) | | | |
| Pulse Ox% (> 95%) | | | |
| Capillary Refill (< 2 sec) | | | |

NOTES:

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| Treatments: | (Mark X all | that apply) and | fill in the blank) |
|-------------|-------------|-----------------|--------------------|
|-------------|-------------|-----------------|--------------------|

Location:

| M: Dressing - Hemostatic Pressure TQ | Other: |
|--------------------------------------|--------|
|--------------------------------------|--------|

A: Intact ET-Tube Tracheostomy

| R: | $\Box O^2$ | Needle-D | Chest-Tube | Chest-Seal |
|----|------------|----------|------------|------------|
|----|------------|----------|------------|------------|

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Total Crystalloid Shock Volume of fluids is 90 mls/kg: Administer 20ml/kg over 10-20 min. Reassess (as with human casualty): If lack of response after 2-3 boluses consider adjunct therapy (HES/HTS.)

| CRYSTALLOID | Volume | Route | Time |
|--|--------|-------|------|
| | | | |
| | | | |
| HYDROXYETHYL STARCH (HES): 5mls/kg over 5 - 10 min. After ½ shock crystalloid not effective. | | | |
| HYPERTONIC SALINE (HTS): 4mls/kg (If two or three 1/4 shock boluses and 1-2 boluses of HES not effective) | | | |
| TXA: 10 mg/kg IV in 100ml NaCl or LRS given in first 3hrs. Followed by a 10-15 mg/kg CRI over 8 hours. | | | |

- C: Splint Other Bandage

H: Hypothermia-Prevention Hypethermia-External Cooling

H: Head Injury

Pain Meds and Antibiotics (Circle if given and write the time in the notes.)

| DRUG (conc) | DOSE | RTE | 60lb/ 27.3kg | 70lb/32kg | 80lb/36.4kg |
|---------------------------|---------------|----------|---------------|---------------|---------------|
| Ketamine (100mg/ml) | 2-5mg/kg | IV/IM | 1 ml | 1.5 mls | 2 mls |
| Midazolam (5mg/ml) | 0.1-0.3mg/kg | IV/IM | 3 mls | 4 mls | 5 mls |
| Morphine (10mg auto inj.) | 0.2-0.5 mg/kg | IM | 1 <u>auto</u> | 1 <u>auto</u> | 2 <u>auto</u> |
| Meloxicam | 0.1-0.2mg/kg | IV/SQ/PO | 5 <u>mq</u> | 6 <u>mq</u> | 7 <u>mg</u> |
| Cefazolin/Ceftriaxone | 25 mg/kg | IV/IM | 600 <u>mg</u> | 800 <u>mg</u> | 900 <u>mq</u> |
| Cefotaxime | 25 mg/kg | IV/IM/SQ | 600 <u>mq</u> | 800 <u>mq</u> | 900 <u>mg</u> |
| Ertapenem (100mg/ml) | 15mg/kg | IV/SQ | 4 mls | 5 mls | 6 mls |

NOTES:

FIRST RESPONDER:

Name (Last, First):

AOC/MOS:

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| | | 5cc | 10cc | 20cc | 50cc | 100cc | 250cc | 500cc | 1000cc |
|------------|-----------|---------------------|----------------|--------------------|-----------------|------------------|-------------------|------------------|----------|
| 1 | mcg | 0.20mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.02mcg/ml | 0.01mcg/ml | 0.004mcg/ml | 0.002mcg/ml | 0.001mcg |
| 5 | mcg | 1mcg/ml | 0.5mcg/ml | 0.25mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.02mcg/ml | 0.01mcg/ml | 0.005mcg |
| 1 | 0mcg | 2mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.2mcg/ml | 0.1mcg/ml | 0.04mcg/ml | 0.02mcg/ml | 0.01mcg/ |
| | 5mcg | 5mcg/ml | 2.5mcg/ml | 1.25mcg/ml | 0.5mcg/ml | 0.25mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.025mcg |
| 5 | 0mcg | 10mcg/ml | 5mcg/ml | 2.5mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.2mcg/ml | 0.1mcg/ml | 0.05mcg/ |
| २ 1 | 00mcg | 20mcg/ml | 10mcg/ml | 5mcg/ml | 2mcg/ml | 1mcg/ml | 0.4mcg/ml | 0.2mcg/ml | 0.1mcg/m |
| 2 | 50mcg | 50mcg/ml | 25mcg/ml | 12.5mcg/ml | 5mcg/ml | 2.5mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.25mcg/ |
| 5 | 00mcg | 0.1mg/ml | 50mcg/ml | 25mcg/ml | 10mcg/ml | 5mcg/ml | 2mcg/min | 1mcg/ml | 0.5mcg/m |
| J 1 | mg | 0.2mg/ml | 0.1mg/ml | 50mcg/ml | 20mcg/ml | 10mcg/ml | 4mcg/ml | 2mcg/ml | 1mcg/ml |
| 2 | mg | 0.4mg/ml | 0.2mg/ml | 0.1mg/ml | 40mcg/ml | 20mcg/ml | 8mcg/ml | 4mcg/ml | 2mcg/ml |
| j 3 | mg | 0.6mg/ml | 0.3mg/ml | 0.15mg/ml | 60mcg/ml | 30mcg/ml | 12mcg/ml | 6mcg/ml | 3mcg/ml |
| 4 | mg | 0.8mg/ml | 0.4mg/ml | 0.2mg/ml | 80mcg/ml | 40mcg/ml | 16mcg/ml | 8mcg/ml | 4mcg/ml |
| 5 | mg | 1mg/ml | 0.5mg/ml | 0.25mg/ml | 0.1mg/ml | 50mcg/ml | 20mcg/ml | 10mcg/ml | 5mcg/ml |
| 6 | mg | 1.2mg/ml | 0.6mg/ml | 0.3mg/ml | 0.12mg/ml | 60mcg/ml | 24mcg/ml | 12mcg/ml | 6mcg/ml |
|) 7 | mg | 1.4mg/ml | 0.7mg/ml | 0.35mg/ml | 0.14mg/ml | 70mcg/ml | 28mcg/ml | 14mcg/ml | 7mcg/ml |
| 8 | mg | 1.6mg/ml | 0.8mg/ml | 0.4mg/ml | 0.16mg/ml | 80mcg/ml | 32mcg/ml | 16mcg/ml | 8mcg/ml |
|) 9 | mg | 1.8mg/ml | 0.9mg/ml | 0.45mg/ml | 0.18mg/ml | 90mcg/ml | 36mcg/ml | 18mcg/ml | 9mcg/ml |
| 1 | 0mg | 2mg/ml | 1mg/ml | 0.5mg/ml | 0.2mg/ml | 0.1mg/ml | 40mcg/ml | 20mcg/ml | 10mcg/m |
| 1 | 5mg | 3mg/ml | 1.5mg/ml | 0.75mg/ml | 0.3mg/ml | 0.15mg/ml | 60mcg/ml | 30mcg/ml | 15mcg/m |
| 2 | 5mg | 5mg/ml | 2.5mg/ml | 1.25mg/ml | 0.5mg/ml | 0.25mg/ml | 0.1mg/ml | 50mcg/ml | 25mcg/m |
| 5 | 0mg | 10mg/ml | 5mg/ml | 2.5mg/ml | 1mg/ml | 0.5mg/ml | 0.2mg/ml | 0.1mg/ml | 50mcg/m |
| 7 | 5mg | 15mg/ml | 7.5mg/ml | 3.75mg/ml | 1.5mg/ml | 0.75mg/ml | 0.3mg/ml | 0.15mg/ml | 75mcg/m |
| 1 | 00mg | 20mg/ml | 10mg/ml | 5mg/ml | 2mg/ml | 1mg/ml | 0.4mg/ml | 0.2mg/ml | 0.1mg/ml |
| 2 | 50mg | 50mg/ml | 25mg/ml | 12.5mg/ml | 5mg/ml | 2.5mg/ml | 1mg/ml | 0.5mg/ml | 0.25mg/n |
| 5 | 00mg | 100mg/ml | 50mg/ml | 25mg/ml | 10mg/ml | 5mg/ml | 2mg/ml | 1mg/ml | 0.5mg/ml |
| 7 | 50mg | 150mg/ml | 75mg/ml | 37.5mg/ml | 15mg/ml | 7.5mg/ml | 3mg/ml | 1.5mg/ml | 0.75mg/n |
| 1 | Gram | 200mg/ml | 100mg/ml | 50mg/ml | 20mg/ml | 10mg/ml | 4mg/ml | 2mg/ml | 1mg/ml |
| | | - | Value e | quals amount o | f fluid in each | ml of dilution | | | |
| Each | ml of med | lication diluted in | nto your chose | n fluid still coun | ts towards tot | al solution volu | me (i.e. 1ml of o | drug + 4ml fluid | = 5ml |

OXYGEN CYLINDER LIFE:

| Cylinder | D | E | G | Н |
|------------|------------------------|------------------------|------------------------|------------------------|
| Liters | 356 | 622 | 5260 | 6900 |
| Flow (LPM) | Length of use (min) | Length of use (min) | Length of use (min) | Length of use (min) |
| 2 | 178 | 311 | 2630 | 3450 |
| 4 | 89 | 155 | 1315 | 1725 |
| 6 | 59 | 104 | 876 | 1150 |
| 8 | 44 | 78 | 658 | 862 |
| 10 | 35 | 62 | 526 | 690 |
| 12 | 30 | 52 | 438 | 575 |
| 15 | 23 | 41 | 350 | 460 |

NOTE: Current MEDEVAC Oxygen Cylinder is "D" type.

To estimate duration of use for Oxygen Cylinders:

• Duration of Flow = Contents of cylinder / Flow rate.

Cylinder Factors for Calculation of Duration of Oxygen Flow:

| Cylinder Size | D | Е | G | H and K | | |
|---------------|------|------|------|---------|--|--|
| Factor | 0.16 | 0.28 | 2.41 | 3.14 | | |

Once you have the cylinder factor and the amount of pressure remaining in the cylinder, the duration of flow can be calculated with the following equation.

Duration of flow (min) = Pressure (psig) x Cylinder Factor/Flow (L/min)

PRE-FLIGHT CHECKLIST

(for Critical Care and Post-Surgical Transfers) Once the decision is made to transfer a patient and an accepting physician has been obtained, the following steps will be taken to prepare the patient for transport:

| | prepare the patient for transport: | | | | | | | | |
|----------|---|--|--|--|--|--|--|--|--|
| Initials | Evaluation Steps | | | | | | | | |
| | 1. Sending location/physician:Accepting location/physician: Flight nurse called: name / time: | | | | | | | | |
| | 2. Anesthesia called: intubation if indicated. ETT secured/marked | | | | | | | | |
| | 3. Patient meets criteria for en route critical care transport: risk documented by sending physician | | | | | | | | |
| | (POST-OPERATIVE and CC INTRAFACILITY TRANSFER, Pre-Transfer Patient Status Requirements) | | | | | | | | |
| | Preparation Steps | | | | | | | | |
| | Positioning and Proper Monitoring: | | | | | | | | |
| | 1. Patient moved to litter (collapsible handles), positioned, padded, strapped, equipment (with necessary attachments) added and secured. | | | | | | | | |
| | 2. For head-injured patients, a pre-sedation neurologic examination will be performed. GCS and neurological exam documented on the en route care form, suggest placing patient sitting at 30°-45°. (For eye injured patients, fox shield in place. For burn patients, <u>JTTS burn sheet initiated</u> .) | | | | | | | | |
| | 3. Ventilator switched to PMI vent at least 20-30 min prior to flight and set with transfer settings ordered by physician. | | | | | | | | |
| | 4. IV / IO access verified, patent, and secured. | | | | | | | | |
| | 5. Arterial line inserted and secured, if indicated. Transducer accessible. | | | | | | | | |
| | 6. Ventilator tubing checked to be free from obstruction, with ETCO ₂ and secondary lines attached. | | | | | | | | |
| | 7. Orogastric or nasogastric tube is inserted (unless contraindicated), placement verified with chest x-ray, and attached to low-intermittent suction. | | | | | | | | |
| | 8. Chest tubes to water seal/suction (place Heimlich valve for non-atrium chest drainage systems). | | | | | | | | |
| | 9. Wound vacuum disconnected and stowed. | | | | | | | | |
| | 10. Foley catheter secured, urine output measured and documented. | | | | | | | | |
| | Equipment, Medication, Chart, and Personnel Preparation: | | | | | | | | |
| | 11. Medications needed for flight prepared and organized. | | | | | | | | |
| | 12. Flight equipment bag obtained and checked. Backup pulse oximeter readily available. | | | | | | | | |
| | 13. Complete chart photocopied (including x-ray cd), patient belongings bagged and tagged. | | | | | | | | |
| | Transfer Document, or other theater / unit approved transfer document, has been initiated. 14. Earplugs and eye protection for patient and flight nurse. | | | | | | | | |
| | 15. If facility sends medical attendant, attendant must have relevant personal protective equipment. In a combat | | | | | | | | |
| | environment this includes: Uniform, Kevlar, IBA, Weapon, ID Card, and equipment for transport. | | | | | | | | |
| | Ventilator Management: | | | | | | | | |
| | 16. Blood gas (preferably ABG) obtained, 15 min after initial settings and ventilator changes. All efforts will be made to have a documented blood gas within 30 minutes prior to flight time. | | | | | | | | |
| | 17. Adjust ventilator settings and check O ₂ tank for length of flight. Resuscitator bag under patient's head with | | | | | | | | |
| | tubing connected to O ₂ source, vent tubing free from obstruction. | | | | | | | | |
| | Final Verification: | | | | | | | | |
| | 18. Transferring Physician, Flight Paramedic, ECCN (or Flight Provider) verbally agrees to flight care plan. | | | | | | | | |
| | 19. Critical Care Transfer Orders reviewed and signed by transferring physician. (STANDARD ORDER SET for CRITICAL CARE TRANSFERS) | | | | | | | | |
| | 20. Enroute CC Transfer Document with completed preflight and enroute care data handed over to and confirmed by receiving provider / facility. (CENTCOM Transfer Document) | | | | | | | | |
| J | | | | | | | | | |

Facility Transfer Checklist

| | DO NOW | | TAKE WITH | | INFLIGHT |
|------------|---|------------|--|-----------|--|
| | Check all bandages, splints, dressings and tourniquets for placement / evidence of ongoing hemorrhage. Mark bleeding strikethrough Measurement of abdomen Request orders for type and cross-matched blood or O-negative blood from the transferring physician. | M : | Additional Blood products (1:1:1) Tubing Warmer Golden Hour Container | M: | Blood products administration Check all bandages, splints, dressings and tourniquets for placement / evidence of ongoing hemorrhage Measurement of abdomen. |
| | Assess and document ET tube size, depth, security, cuff pressure, bite block Attach ETCO2 monitor Insert/Assess and document NG/OG tube placement, size, depth, security Review chest radiograph for ET confirmation and NG/OG placement Apply C-collar for airway stability | A: | Extra ETT / King LT / IGEL Suction soft-tip 10ml syringe Bite block Tape BVM | A: | Confirm ETT is in appropriate position Look/feel for symmetric chest wall rise Verify tube position at teeth Check ETCO2 DOPE |
| R : | Setup Ventilator and confirm ventilator settings Check baseline lung compliance/resistance with BVM Auscultate heart/lung sounds Check placement and function of chest tube/drainage system/ Heimlich valve. Request arterial blood gas after transport ventilator is attached to patient | R: | O2 for transport Backup ventilator Suction Needle for decompression | R: | Look and feel for chest excursion Check Pulse Ox Check patient's color |
| C : | Setup Monitor and zero all Pressure Lines Assess distal pulses and neurovascular status Ensure IV access x 2 (Minimum). Remove air from IV bags and pressurize IV medications arranged for easy access (20ml, 10ml, 5ml, 3ml) Review CBC/Chemistry results Check Foley catheter placement, measure output amount, empty bag | | Vasoactive medications (dopamine, neo, norepinephrine) Pressure bags IV fluids and tubing | ünn nn | Check temp, pulse BP, and cardiac rhythm Assess distal pulses and neurovascular status during transport. Assess IV access LZC Pressure Lines |
| | Conduct baseline neurologic exam Provide eye and ear protection to patient; HPMK, warmed IV fluids, blankets and/or chemical heat packs. Review orders and discuss potential en-route problems with the transferring physician. Ensure patient and all equipment secured Place transport ventilator on transport O2 Assess pain control, sedation and need for paralysis. Re-dose medications if needed before flight Patient loading considerations | | Collect all labs, x-rays, pre aid- station/hospital documentation for transport Reconcile medications; verify allergies and patient's weight Secure personal effects. Sedation meds (propofol, versed, ketamine) Pain meds (fentanyl, morphine, ketamine) Paralytic meds (vecuronium, rocuronium) 3% NaCl, | | Assess neurologic and sedation Check placement of all tubes, lines and drains & ensure proper functioning Ensure all wires and tubing are accessible and have adequate slack to allow monitors and IVs to be properly positioned and secured Serial assessments Prepare patient and give report to receiving facility |

| Altitude Considerations | Medications | Patient Packaging |
|--|--|--|
| Required waiting time before transport | Type and number of patients | Additional Medical support/non-medical attendant |
| Respiratory Support | Monitoring (body systems, medical interventions, etc.) | Telephonic consultation |
| Equipment | Thermal considerations | Transport time and Route of transfer |

VASOPRESSOR PRIORITY CHART

| | HYPOVOLEMIC SHOCK | SEPTIC SHOCK | CARDIOGENIC SHOCK1 | NEUROGENIC SHOCK _{2,3} | BURN SHOCK₄ |
|----|---|-----------------|-----------------------|------------------------------------|----------------|
| 1° | Vasopressors are not recommended in the initial stabilization of hypovolemic shock | Norepinephrine | Norepinephrine | Norepinephrine | Vasopressin |
| 2° | | Vasopressin | Dobutamine | Epinephrine | Norepinephrine |
| 3° | Norepinephrine | Epinephrine | Epinephrine | Vasopressin | Epinephrine |

• Vasopressors should only be initiated with/after adequate resuscitation is provided with crystalloids, colloids, and/or blood products.

• Maintain mean arterial pressure (MAP) 65 mmHg or as needed to achieve adequate end-organ perfusion (e.g. cerebral perfusion pressure, abdominal perfusion pressure, urinary output).

1. In low output Cardiogenic Shock, dobutamine may be initiated in combination with norepinephrine.

2. Due to the physiologic nature of Neurogenic Shock, vasopressors may be initiated earlier to avoid volume overload.

3. Phenylephrine should be avoided in most Neurogenic Shock patients due to unopposed alpha activity that can result in reflex bradycardia; further worsening spinal cord injury (SCI) associated bradycardia.

4. In Burn Shock casualties at risk of burn fluid over-resuscitation (e.g. 250mL/Kg in the 1st 24 hours), a continuous, non-titratable infusion of Vasopressin at 0.04 Units/minute (2.4 Units/hour) may be initiated to avoid volume overload

EXAMPLE Standing Order Sheet for Critical Care Patient Transfers

PATIENT IDENTIFICATION

(Last, First, Middle Initial; SSN/Identification Number; grade; DOB; treatment facility)

Date:

Sending Facility:

Sending Physician:

Receiving Facility:

Diagnosis:

Condition:

Patient Category:

Allergies:

Height:

Weight (kg):

Fluids: []LR mL/hr []NS mL/hr []3% Saline mL/hr []D5W []Other_____[] PRBC []FWB []Plasma []LTOWB

Monitoring: [] Vital Signs [] Every 5 min Vital Signs [] Every 15 min Vital Signs [] Every 30 min [] Continuous cardiac monitoring, document rhythm strips pre-flight and with any rhythm changes [] ICP/CPP [] CVP [] GCS [] ETCO2 [] UO_____mL hourly

Activity: [] Bed rest

[] Spine precautions: C-Collar/C-Spine TLS Spine

Nursing: [] Wound VAC dressing to _____mm Hg suction

[] NGT to low continuous suction OR [] Clamp NGT

[] OGT to low continuous suction OR [] Clamp OGT

[] Chest tube 1 to: water seal (circle: R L Both) OR _____cm H2O Suction (circle: R L Both) [] Chest tube 2 to: water seal (circle: R L Both) OR _____cm H2O Suction (circle: R L Both) [] Chest tube 3 to: water seal (circle: R L Both) OR _____cm H2O Suction (circle: R L Both) [] Chest tube 4 to: water seal (circle: R L Both) OR _____cm H2O Suction (circle: R L Both) [] Chest tube elevated ______ degrees [] Keep HOB flat

Respiratory: [] Keep O2Sat >_____%

Oxygen: [] Nasal Cannula at _____LPM [] Non-rebreather at _____ LPM

Ventilator Settings: Mode: [] SIMV [] AC [] CPAP [] BiPAP

Rate: _____breaths per minute I:E ratio:_____

Tidal Volume: _____ mL FiO2:_____ % PEEP: _____cm H2O PIP: _____

| PATIENT IDENTIFICATION |
|--|
| (Last, First, Middle Initial; SSN/Identification Number; grade; DOB; |
| treatment facility) |
| Vasoactive Medications: |
| [] Dopaminemg/mL atmcg/kg/min IV; titrate to MAP > |
| mm Hg |
| [] Norepinephrine 4mg/mL atmcg/min IV; titrate to MAP > |
| mm Hg |
| [] Phenylephrine 10mg/mL atmcg/min IV; titrate to MAP > |
| mm Hg |
| [] Epinephrinemg (1:10,000)/mL atmcg/min IV; titrate to MAP |
| > mm Hg [] Other |
| Sedation and Analgesics: |
| [] Ketaminemg/kg Qminutes IVP PRN sedation to Riker Sedation- |
| Agitation Scale of 1-2 [] Midazolammg Qminutes IVP PRN |
| sedation to Riker Sedation-Agitation Scale of 1-2 [] Haloperidolmg |
| Qminutes IVP PRN sedation to Riker Sedation-Agitation Scale of 1-2 |
| [] Lorazepammg Qminutes IVP PRN sedation to Riker Sedation- |
| Agitation Scale of 1-2 [] Fentanylmcg Qminutes IVP PRN pain |
| [] Morphinemg Qminutes IVP PRN pain |
| [] Other |
| Paralytics: |
| [] Rocuroniummg IVP |
| [] Vecuroniummg IVP |
| Intracranial Hypertension: |
| [] 3% Hypertonic Saline 250 cc bolus for any signs of herniation |
| [] Mannitol Infusion Rate: |
| Labs: |
| [] ABG 15 minutes prior to departing sending facility |
| [] Other: |
| Additional critical information: |
| Physician Signature: |

USEFUL CALCULATIONS

PEDIATRIC FORMULAS:

- ETT Size = (Age/4)+4 (Age divided by 4 plus 4)
- **ETT Depth =** 3 x ETT Size (Endotracheal)
- Weight in kg (>1 year) = (Age (years) x 2) + 8
- Systolic Blood Pressure minimum = 70 + [2 x Age (years)]

MEDICATION FORMULAS:

- **Mcg/kg/min (micrograms/kilogram/minute)** = [16.7 X Drug Concentration (mg/ml) x infusion rate (ml/h)] Weight (kg).
- INFUSION RATE (ml/h) = [Desired mcg/kg/min x Weight (kg) x 60]/Drug concentration (mcg/mL)

HEMODYNAMIC FORMULAS:

- **MAP**: Mean Arterial Pressure = [(2 x DBP) + SBP]/3.
- **SBP** = (Systolic Blood Pressure)
- **DBP** = (Diastolic Blood pressure)
- / = (Divided by)
- PULSE PRESSURE: SBP DBP or (Systolic Blood Pressure minus Diastolic Blood pressure).
- Cerebral Profusion Pressure (CPP): MAP-ICP=CPP
- **ICP**= (Intracranial Pressure)
- Ideal CPP=>60 While ICP cannot often be measured during flights; an assumption that patients with TBI have an ICP of 15-20 will allow hemodynamic optimization in these patients to ensure adequate CPP.

COMMON CONVERSIONS:

- lbs. = kg x 2.2 or kg = lbs. x 0.45
- Fahrenheit = (Celsius x 1.8) + 32 or Celsius = (Fahrenheit -32) x 5/9
- 1 tsp. = 5 ml
- 1 tbsp. = 15 ml
- 1 oz. = 30 ml
- 1g = 1,000 mg
- 1mg = 1,000 mcg
- 1 g = 10,000 mcg

COMMON VENTILATOR FORMULAS

- Calculation to target ETCO2: (Current Rate x Current ETCO2) / Desired ETCO2 = New Rate. NOTE: You may incur a pressure limitation alarm after adjusting to New Rate for ETCO2 targeting on the Hamilton T1 vent.
- Calculation for vent adjustments due to abdominal/thoracic pressure: Current Minute Ventilation / New Rate (Current Rate + 1 or 2) = New Tidal Volume. Now set New Rate and New Tidal Volume. NOTE: Pressure alarm may not go away and may need to recalculate after several minutes if SpO2 has not improved.

Y-SITE COMPATABILITY CHART

| | Solı | utior | าร | | | | | | | | | | _ | _ | | | | | Μ | ledic | atio | ns | | | | | | _ | | | | | | |
|----------------------------|------|-------|----|---------------|----------|---------|------------|------------------|-----------------|-----------|------------|----------|-------------|---------|----------|---------------|---------|-------------------|----------|--------------|-----------|------------|-------------|------------|-------------|---------------|----------------|--------------|---------------|-----------|--------------|----------|--------------|------------|
| | | 1 | | [| [| | | | | | | | 1 | | | | | | [| [| | | | | | | | | | | | | | |
| | NS | D5W | LR | Sodium Bicarb | Mannitol | Albumin | Amiodarone | Calcium Chloride | Dexmedetomidine | Diltiazem | Dobutamine | Dopamine | Epinephrine | Esmolol | Fentanyl | Furosemide | Heparin | Insulin (regular) | Ketamine | Lorazepam | Magnesium | Metoprolol | O Midazolam | O Morphine | Nicardipine | Nitroglycerin | Norepinephrine | Pantoprazole | Phenylephrine | Phenytoin | Potassium Cl | Propofol | Vasopressin | Vecuronium |
| Amiodarone | V | C* | Ν | N | С | | | С | С | С | С | С | С | С | С | Ν | Ν | С | С | С | Ν | С | | С | С | С | С | Ν | С | Ν | С | | Ċ | C |
| Calcium Chloride | С | Ν | V | Ν | С | | С | | С | С | С | С | С | С | С | С | С | С | | С | Ν | С | С | С | С | С | С | Ν | С | Ν | С | Ν | С | С |
| Dexmedetomidine | С | С | С | С | С | | С | С | | С | С | С | С | С | С | С | С | С | | С | V | С | С | С | С | С | С | Ν | С | С | С | С | С | С |
| Diltiazem | С | С | Ν | V | С | С | С | С | С | | С | С | С | С | С | Ν | V | Ν | | С | С | С | С | С | С | С | С | Ν | С | Ν | С | | С | С |
| Dobutamine | С | С | С | Ν | С | | С | С | С | С | | С | С | С | С | Ν | Ν | V | С | С | С | С | Ν | С | С | С | С | Ν | С | Ν | С | V | С | С |
| Dopamine | С | С | С | Ν | С | | С | С | С | С | С | | С | С | С | Ν | С | Ν | С | С | С | С | С | С | С | С | С | Ν | С | Ν | С | V | С | С |
| Epinephrine | V | C* | С | Ν | С | | С | С | С | С | С | С | | С | С | С | С | V | С | С | С | С | С | С | С | С | С | Ν | С | Ν | С | V | С | С |
| Esmolol | С | С | С | С | С | С | С | С | С | С | С | С | С | | С | Ν | С | С | | С | С | С | С | С | С | С | С | Ν | С | V | С | С | С | С |
| Fetanyl | С | С | Ν | С | С | | С | С | С | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | С | С | С | Ν | С | Ν | С | С | С | С |
| Furosemide | С | С | С | С | С | С | Ν | С | С | N | Ν | Ν | С | Ν | С | | С | V | V | С | Ν | С | Ν | Ν | Ν | Ν | Ν | V | Ν | Ν | С | С | Ν | Ν |
| Heparin | V | C* | V | Ν | С | | Ν | С | | V | Ν | С | С | С | С | С | | С | Ν | С | С | С | С | С | Ν | С | С | Ν | С | Ν | С | С | С | С |
| Insulin (Regular) | V | Ν | Ν | Ν | С | | С | С | С | N | V | Ν | V | С | С | V | С | | Ν | V | С | С | V | С | С | С | Ν | V | Ν | Ν | С | С | V | С |
| Ketamine | С | С | Ν | Ν | С | С | С | С | | | С | С | С | | С | V | Ν | Ν | | V | С | | С | С | | V | | | | Ν | С | С | | |
| Lorazepam | С | V | Ν | С | С | С | С | С | С | С | С | С | С | С | С | С | С | V | V | | С | С | С | С | V | С | С | Ν | С | Ν | С | С | С | С |
| Magnesium | С | С | С | Ν | С | | Ν | Ν | V | С | С | С | С | С | С | Ν | С | С | С | С | | С | С | С | С | С | С | Ν | С | Ν | С | V | С | С |
| Metoprolol | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | | С | С | | С | С | С | Ν | | Ν | С | Ν | С | | С | С |
| Midazolam | С | С | Ν | Ν | С | Ν | С | С | С | С | Ν | С | С | С | С | Ν | С | V | С | С | С | С | | С | С | С | С | Ν | С | Ν | С | V | С | С |
| Morphine | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | Ν | С | С | С | С | С | С | С | | С | С | С | V | С | Ν | С | V | С | С |
| Nicardipine | С | С | V | Ν | С | | С | С | С | С | С | С | С | С | С | Ν | Ν | С | | V | С | С | С | С | | С | С | Ν | С | Ν | С | | С | С |
| Nitroglycerin | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | Ν | С | С | V | С | С | Ν | С | С | С | | С | V | С | Ν | С | V | С | С |
| Norepinephrine | С | C* | С | Ν | С | | С | С | С | С | С | С | С | С | С | Ν | С | Ν | | С | С | С | С | С | С | С | | Ν | С | Ν | С | С | С | С |
| Pantoprazole | С | С | С | Ν | Ν | | Ν | Ν | Ν | N | Ν | Ν | Ν | N | Ν | V | Ν | V | | Ν | Ν | Ν | Ν | V | Ν | V | Ν | | С | Ν | С | Ν | С | Ν |
| Phenylephrine | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | Ν | С | Ν | | С | С | С | С | С | С | С | С | С | | Ν | С | Ν | С | С |
| Phenytoin | V | Ν | Ν | Ν | Ν | | Ν | Ν | С | N | Ν | Ν | Ν | V | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | | Ν | Ν | Ν | Ν |
| Potassium Cl | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | С | Ν | | С | С | С |
| Propofol | С | С | С | С | С | | | Ν | С | | V | V | V | С | С | С | С | С | С | С | V | | V | V | | V | С | Ν | Ν | Ν | С | | | V |
| Vasopressin | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | Ν | С | V | | С | С | С | С | С | С | С | С | С | С | Ν | С | | | |
| Vecuronium | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | Ν | С | С | | С | С | С | С | С | С | С | С | Ν | С | Ν | С | V | | |
| Ateplase (tPA) | | | | C* | | N | N | | | 1 | · | | | | | | | | | | | | - | | | | | | | | | | | _ |
| Tenecteplace (TNK) | | | | C* | | N | N | | | | | | |) | | ite C = Co | | | | | | | | V | aso | pres | | s sh emse | | • | fera | ıbly ı | run b | ,y |
| Transexamic Acid | | | | C* | | С | С | | | | | | C* = | = Co | mpa | atible | e an | d pe | erfer | red | for | | | | Not | com | mo | nlv c | arri | ed i | n M | EDE | VAC | |
| Cefazolin | | | | С | | С | С | | С | | | | ` | v = ' | | econ able | | | | ed) | | | - | | | | | iny c | Juin | | | | <u>v/ (C</u> | · |
| Ceftrixazone | | | | С | | С | N | | С | | | | | Ν | = N | on - | Cor | mpa | table | Э | | | | | | | | | | | | | | |
| Pipercillian / Tazo | | | | С | | С | V | | С | | | | | 1) / - |) + - | | ا م | | | | - | | | | 1) // | - /• | | | <u>)</u> | | | <u></u> | | |
| Meripenem | | | | V | | N | N | | Ν | 1 | | me | | | | | | | | i po line | | | | | | | | | | | | | ady | , |
| Vancomycin | | | | С | | С | С | | V | | | | | | | | | | | | | | e lir | | | | | | | | | | | |
| Levaquin (Pre-mix) C C C V | | | | | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | _ | | | | | | | | |

VITAL FUNCTIONS ASSESSMENT REFERENCE CHART

| GLASGOW COMA SCALE | | | | | | | | | | | | |
|--|--|------------------------------------|---------------------------|--|--|--|--|--|--|--|--|--|
| SCORE | ADULT | CHILD | INFANT | | | | | | | | | |
| | | Eye Opening | | | | | | | | | | |
| 4 | Spontaneous | | | | | | | | | | | |
| 3 | To Speech | Evo Oponing Booponoo Somo oo Adult | | | | | | | | | | |
| 2 | To pain | Eye Opening Response Same as Adult | | | | | | | | | | |
| 1 None | | | | | | | | | | | | |
| | | Verbal Response | | | | | | | | | | |
| | 5 Oriented Oriented Coos and babbles | | | | | | | | | | | |
| 4 Confused Conversation Confused Conversation Irritable, Cries | | | | | | | | | | | | |
| 3 | Inappropriate Words | Inappropriate Words | Cries in Response to pain | | | | | | | | | |
| 2 | Incomprehensible | Incomprehensible Words/Sounds | Moans in Response to | | | | | | | | | |
| | Sounds | • | Pain | | | | | | | | | |
| 1 | None | None | None | | | | | | | | | |
| | | Best Motor Response | | | | | | | | | | |
| 6 | Obeys Commands | Obeys Commands | Moves Spontaneously | | | | | | | | | |
| 5 | Localizes Pain | Localizes Pain | Withdraws to Touch | | | | | | | | | |
| 4 | Flexion Withdrawal to | Flexion Withdrawal to Pain | Withdraws from Pain | | | | | | | | | |
| | Pain | | Stimulus | | | | | | | | | |
| 3 | 3 Abnormal Flexion (Decorticate) Abnormal Flexion (Decorticate) Abnormal Flexion (Decorticate) | | | | | | | | | | | |
| 2 | Extension (Decerebrate) | Extension (Decerebrate) | Extension (Decerebrate) | | | | | | | | | |
| 1 | 1 None (Flaccid) None (Flaccid) None (Flaccid) | | | | | | | | | | | |
| For Intubated Patient use Verbal "T" | | | | | | | | | | | | |
| (Ex | (Example: Eyes open to pain, Intubated, and Localizes would be E2,V1,M5, or GCS 8T) | | | | | | | | | | | |

VITAL FUNCTIONS ASSESSMENT REFERENCE CHARTS

MUSCULOSKELETAL INJURY and PERIPHERAL NERVE ASSESSMENT UPPER EXTREMITIES INJURY to Consider MOTOR Testing **SENSATION Testing** NERVE Elbow Injury Index and Little Finger Little Finger Ulnar Abduction Wrist Fracture or Thenar Contraction with Index Finger Median Distal Opposition Dislocation Index Tip Extension None Median, Anterior Supracondylar Fracture of Humerus Interoseous Anterior Shoulder Elbow Flexion Radial Forearm Musculocutaneous Dislocation Distal Humeral Shaft, Thumb, Finger group First Dorsal Web Space Radial Anterior Shoulder Extension Dislocation Anterior Shoulder Lateral Shoulder Deltoid Axillary Dislocation. **Proximal Humerus** Fracture LOWER EXTREMITIES Pubic Rami Fractures **Knee Extension** Anterior Knee Femoral Hip Adduction **Obturator Ring Fractures** Medial Thigh Obturator **Posterior Tibial** Toe Flexion Sole of Foot Knee Dislocation Fibular Neck Fracture, Knee Dislocation Ankle Eversion Lateral Dorsum of Foot **Superficial Peroneal** Fibular Neck Fracture. Ankle / Toe Dorsiflexion Dorsal 1st-2nd Web Deep Peroneal **Compartment Syndrome** Space Posterior Hip Dislocation **Plantar Flexion** Foot Sciatic Nerve Acetabular Fracture **Hip Abduction** Upper Buttocks Superior Gluteal Acetabular Fracture **Hip Extension** Lower Buttocks Inferior Gluteal

| MUS | SCULAR STRENGTH GRADING |
|-------|--|
| SCORE | EXAM RESULTS |
| 0 | Total Paralysis |
| 1 | Palpable or Visible Contraction |
| 2 | Full Range of Motion Without Gravity |
| 3 | Full Range of Motion Against Gravity |
| 4 | Full Range of Motion, but Less than Normal Strength |
| 5 | Normal Strength |
| NT | Not Testable |

VITAL FUNCTIONS ASSESSMENT REFERENCE CHART

| PEDIATRIC ALS EQUIPMENT | | | | | | | | | | | | | | | | |
|-------------------------|--|--------|--------------------|------------------|------------------|-------------|----------|--------------|------------|-------------|---------------|------|-------|-------------|----------|--|
| | | (Alv | ways use a Broselo | ow Pediatr | ic Emergne | есу Та | ape if a | availat | ole) | | | | | | | |
| BROSELOW cm | <61cm | 61cm | 67cm | 75cm | 87cm | 96cm 10 | | 109 | 109cm | | 122cm | | 138 | cm | 149+cm | |
| (approx) weight | pprox) weight 3-5kg <mark>6-7kg 8-9kg</mark> 10-11 | | | | | 15-1 | 8kg | 19-23 | 3kg | 24-29kg | | | 30-3 | 6kg | 37>kg | |
| AGE | | MO | NTHS | | | | | Y | EARS | ; | | | | | | |
| | 0 1 2 | 3 4 | 5 6 7 8 9 10 11 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12-16 | |
| RESUSCITATION BAG | Infant | | | Child | | | | | (| Child | /Adu | lt | | | Adult | |
| O2 MASK | | New | born | | Pe | diatric | : | | | | | | Ad | ult | | |
| ORAL AIRWAY | Infa | ant/Sm | all Child | S Child | | Child | ł | | С | hild/ | S Ad | lult | | Me | ed Adult | |
| BAG MASK | Infant | | | Pediatric | | | | | | Peds/A | | | | ds/Adult | | |
| LARYNGOSCOPE | 0-1 | | 1 Straight | | 2 Strait | 2Str | aight/ | Curve | b | 2-3 St/Curv | | | / | 3 St/Curved | | |
| ET TUBE | 2.5-3 Uncufd | 3 | .5 Uncuffed | 4 Un cuffed | 4.5 Un cuffed | 5 l cuff | Jn ed | 5.5 cuffe | - | 6 Cuffed | | | | 6.5 Cuffed | | |
| STYLET | | | 6 | | | | | | | | 14 | | | | | |
| SUCTION | 6-8 | | 8 | 8-10 | | | | 10 | | | | | 12 | | | |
| BP CUFF | Ne | ewborr | n/Infant | Infant/ Child | | С | hild | | | - | hild/ dult | | | A | dult | |
| IV CATHETER | 22-24 | | | 20-24 | 18 | 3-22 | | | 18 | 3-20 | | | | 16 | 6-20 | |
| OG/NG TUBE | | : | 5-8 | 8-10 | 10 | 10- | -12 | 12- | 14 | 1 | 4-18 | | | | 18 | |
| CHEST TUBE | | 10-12 | | | 20 | 20-24 2 | | | 4-32 28-32 | | | | 32-40 | | | |
| URINARY CATHETER | | 5-8 | | | 10 10-12 | | | | | | 1 | 12 | | | | |
| CERVICAL COLLAR | N/A | | | Sma | Small | | | | S/M Medium | | | um | n M/L | | | |

Weights and lengths in above chart are estimates. To achieve most accuracy, utilize Broselow tape on patient.

| ZOI | L DEF | IBRIL | LATI | | GY SE | TTING | S FOR I | PEDIAT | RIC PA | FIENTS | 5 | |
|----------------------|--------|-------|-------|---------------|---------------------------|---------|---------|---------|---------|---------------|----------|-----|
| BROSELOW | cm | <61cm | 61cm | 67cm | 75cm <mark>87cm 96</mark> | | 96cm | 109cm | 122cm | 138cm | cm 149+0 | |
| (approx) | weight | 3-5kg | 6-7kg | 8-9kg | 10-11kg | 12-14kg | 15-18kg | 19-23kg | 24-29kg | 30-36kg | 40kg | 45 |
| | pounds | 6-11 | 13-15 | 17-20lbs | 22-25 | 27-32 | 34-41 | 42-52 | 54-65 | 67-80 | 90 | 101 |
| AGE | | | MON | THS | YEARS | | | | | | | |
| | | 0 1 2 | 3 4 5 | 6 7 8 9 10 11 | 1 | 2 | 3 4 | 5 6 | 7 8 9 | 10 11 | 12-1 | 6 |
| FLUID BOLUS | | 80ml | 130 | 170ml | 210ml | 260ml | 340ml | | | | | |
| ZOLL DEFIB EN 1st | 8J | 10J | 15J | 2 | 20J 30J | | | 50 | J | 75. | J | |
| 2nd | | 15J | 20J | 30J | | 5 | i0J | 75J | 100J | 120J | 150J | |
| MAXIMUM | | 30J | 50J | 75J | 100J 120J 150 | | | юJ | | 200J | | |

Weights and lengths in above chart are estimates. To achieve most accuracy, utilize Broselow tape on patient.

VITAL FUNCTIONS ASSESSMENT REFERENCE CHARTS

| | AVERAGE VITAL FUNCTIONS BY AGE | | | | | | | | | | | | | | |
|--------------------|--------------------------------|--------------|---------------|-------------|-------------------------|------|-------|------|------------|------|-------|---------|------|------|--------|
| BROSELOW cm | <61cm | 61cm | 51cm 67cm | | 87cm | 96 | 96cm | | cm | 1 | 122cm | | 138 | cm | 149+cm |
| (approx) weight | 3-5kg | 6-7kg | 8-9kg | 10-11kg | 12-14kg | 15-1 | l8kg | 19-2 | 19-23kg 24 | | | 24-29kg | | 86kg | 37>kg |
| AGE | NTHS | YEARS | | | | | | | | | | | | | |
| | 0 1 2 3 | 3 4 5 | 6 7 8 9 10 11 | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 | 12-16 |
| HEART RATE | 107-181 | 7-181 93-161 | | | 70-14 | 2 | | 59-1 | .31 | | | 52 | -115 | | 43-108 |
| RESP RATE | 25-66 | | 22-64 | 19-53 | 19-53 17-38 16-29 14-25 | | | | | 1-25 | | 12-23 | | | |
| SYSTOLIC BP | YSTOLIC BP 60 79-105 | | | 85- | 85-108 88-110 91-1 | | | | -119 | 119 | | | | -137 | |
| DIASTOLIC BP | ASTOLIC BP 34-81 | | 40 | 40-69 45-68 | | -68 | 51-89 | | | | | | 59 | -86 | |
| JRINE (mL/kg/hr) 2 | | 1.5 | | | 1 | | | | | | | 0.5 | | | |

Weights and lengths in above chart are estimates, to achieve most accuracy utilize Broselow tape on patient

| | OXYGEN SATURATION | | | | | | | | | | | |
|--------------------------|---|------------------------------|--|--|--|--|--|--|--|--|--|--|
| | Sea Level | 5,000 Feet MLS | | | | | | | | | | |
| SpO2 (Peripheral O2 Sat) | >94% for patient with Normal Hemoglobin level | >92% | | | | | | | | | | |
| StO2 (Tissue O2 Sat) | >75-95% | Same (<75% = Poor Perfusion) | | | | | | | | | | |
| EtCO2 | 35-45 mmHg | | | | | | | | | | | |

MASCAL TRIAGE

Triage Principles

- Priorities change based on time from injury
- Activities in first hour are CRITICAL
- Don't waste time with formal triage tools
- Extricate/stop threat, stop external bleeding, clear airway
- Need for transfusion and ventilator support within the first hour identify a resource-intensive patient
- Damage control surgery has little impact after the first hour
- (COMBAT) Assume minimals will stay armed/engaged if no mental status-altering meds are given for pain
- Expectant category is ONLY used in combat operations and/or when the requirements to adequately treat these patients exceed the available resources. In peacetime, it is generally assumed that all patients have a chance of survival.

| Category | Examples |
|---|--|
| Category I: Immediate (red chemlite) | (Any MARCH issue) Airway obstruction Flail/open chest wound Tension- Pneumothorax/hemothorax Massive hemorrhage 20-70% Burns Unstable Vital Signs Severe TBI (unconscious alive Pt) |
| Category II: Delayed (green chemlite) | Open fractures w/PMS intact Soft tissue injuries Moderate TBI (stable vital signs) Open abdominal wounds |
| Category III: Minimal (no chemlite) emain armed continue to engage | Minor abrasions, burns, sprains lacerations Moderate/Mild anxiety Fractures/dislocations w/PMS Mild TBI |
| *Category IV: Expectant or Hero (blue hemlite) | Massive head or spinal injury Third degree burns > 70% BSA Injuries incompatible with life |

START TRIAGE: Assess, Treat (use bystanders) When you have a color: STOP – TAG – MOVE ON

| | | - | ounded |
|--------|---------|--------|---|
| | No R | ESPIR | ATIONS after head tilt |
| | | Bre | athing but UNCONSCIOUS |
| • | | Res | pirations over 30 |
| DECE | I M | | fusion capillary refill > 2 or NO RADIAL PULSE Control bleeding |
| A | ME | Me | ntal Status: unable to follow simple commands |
| S E | D | D | Otherwise |
| D | I A T E | ELAYED | Remember: Respirations – 30 Perfusion – 2 Mental Status – Can Do |

Notes, Warnings, Cautions

• CPG ID: 91 (Prolonged Casualty Care Guidelines)

| | | | | | | | | | | | | _ | | | | | | _ | | | _ | | | | | _ | | _ | | | | | | | | | | | | | |
|---|--|----------------------------|---|---|---|--|---------------------------------|---------------------------|--|--------------------------------|---------------------------------------|----------------------------------|---------------------------|--|--------------------------------|---|-----------------------------------|---------------------------|---------------------------|---|-----------------------------------|--|---|---|---|----------|--|------------------------|----------------------------------|----------------------------------|--|--|---|---|------------------------------------|--|----------------------------|----------------------------|-----------|--|------------------------------|
| Magnesium Sulfate | Epinephrine 1mg10mL | Atropine | Adenosine | Amiodarone (Infusion) | Amiodarone (Cardiac Arrest) | Epinephrine 1mg/10mL | Nitroglycerin Tablet/Spray | Aspirin, Che | Glucagon Kit CARDIAC (See also Morphine or Fentanyl for AMI pain) | D50 | Epineparine imgimi. | Albureroi(MDI) | Albuterol(Nebulizer) | RESPIRATORY (See also: Methylprednisolone) | Methylprednisolone/Solu-Medrol | Diphenhydramine (Benadryl) | ANAPHYLAXIS | Promethazine (Phenergan) | Zofran(Ondansetron) | Naloxone (Narcan) | Glucagon Kit | OD/Tox Ingestions (See also Sodium Bicar | Phenylephrine(NEO) | Norepinephrine (Levophed) | Epinephrine 1mg/10mL | PRESSERS | Vecuronium | Succinylcholine | Roc | Midazolam (Versed)** PARALYTICS | Etomidate | Propofol (Constant Infusion) | Propofol (Bolus) | Ketamine ** HIGH DOSE | SEDATION ** = Controlled Substance | Ketorolac (10radol) | Fentanyi (Sublimaze) | | ANALGESIA | ANTI-FIBRINOLYTIC Tranexamic Acid (TXA) | DRUG |
| 1-2 Gram IV/IO | 2-10 mcg/min | 1 mg IV/IO | 6 mg/ 12 mg IV/IO Rapid Push | 150 mg over 10-15min, followed by 1 mg/min for 6brs | 300 mg 1" Dose 150 mg 2" Dose | lmg | 0.4 mg SL | 324 mg PO | l mg | 10-25 Grams IV/IO | oan tta con na turche Statico-cro | 0.1.0.5 mm 2.0.70 ar 0.5 million | 2.5-5 mg | | 125 mg IV/IO | 25-50 mg 1M | 2 1 -11-20 CO | 12.5-25 mg IV/IO/IM | 4-8 mg IV/IO/IM/PO | 0.4-2 mg IV/IO/IM | 5 mg IV/IO/IM | See also Sodium Bicarbonate, Calcium Chloride) | 10 mg/10 0ml NS= 100mcg/ml or 10 mg/250 ml NS= 40mcg/ml | 4 mg/500 ml= 8 mcg/ml | 1 mg/500 ml NS= 2mcg/ml | | 0.1 mg/kg IV/IO push Reconstitute w/ 10 ml NS | 1-1.5 mg/kg IV/IO push | 0.6-1.2 mg/kg IV/IO push | 0.05-0.1 mg/kg IV/IO >1 min | 0.2-0.4mg/kg IV/IO push | 10-75 mcg/kg/min IV/IO | 1-2.5 mg/kg <u>bohus</u> IV/IO | 1-2 mg/kg IV/IO | AT SWEET TO DO COAT SHI C.T. | INT BUDG-CL JO AT BUD CT | 0.5-1 mcg/kg IV/IO >3-5 mm | 0.1-0.3 mg/kg IV/IO >1 min | | 2000 mg IV/IO | STANDARD DOSING |
| 1-2 Gram di | 2-10mcg/m | | 6mg 1st do | 150mg inft | 300mg IV/I | lmg (1 amp | 1 Ta | | | | men-en | 0 2 0 5 | | | | | - | | | 0.4-2m | | | 50-20 | Start 2-2 BP) O | 2-20mcg | | ómg | 60-90mg | 36-72mg | 3-6mg | 12-24mg | 600-4500 mcg/min | 60-150mg | 60-120mg | ound | Ánu: | 30-00mcg | 6-18 mg | | 2 Grams | SMALL ADULT (60KG)132LBS |
| 1-2 Gram diluted in 50ml D5W over 15 <u>mim</u> (Torsades w/ pulseVF/V-Tach) | 2-10mcg/minute infusion titrated to desired effect | 1mg q 3-5 min | 6mg 1st dose/12mg 2nd doseFast push w/ rapid large (>20cc) flush | 150mg infusion over 10-15min; followed by 360mg (1mg/min) infusion over 6hrs | 300mg IV/IO bolus. If no change in 3-5min give 150 mg IV/IO. | Img (1 amp) IV/IO Q 3-5 min for Arrest | 1 Tab/Spray SL Q 5 minMAX 3 | 4 X 81mg Tablets (Chewed) | 1 Kit (1mg) IV/IM | 20-50ml | evittes in the other stations and the | 4-8 Fulls | 2.5-5mg (mixed in 3ml NS) | , | 125mg | 25mg PO: 25-50mg IV/IO | hute Twinstor or 0 June 11 | 12.5-25mg | 4-Smg | 0.4-2mg titrated to appropriate ventilation | 5mg slow IV Push | | 50-200mcg (1-5 ml) q 5-10 min IVP/IO | Start 2-20mcg/min infusion initially. (adjust for BP) Once BP is appropriate, 2-4mcg/min | 2-20mcg IVP q 2-5min or 2-20mcg/min IV/IO infusion | | Smg | 80-120mg | 48-96mg | 4-Smg | 16-32mg | 800-6000 mcg/min | 80-200mg | 80-160mg | o III | WT BRUG-CT JO AT BRUCT | 40-80mcg | 8-24 mg | | 2 Grams in 100cc NS over 10min or slow IVP | ADULT (80KG)176LBS |
| 5 <u>min</u> {Torsades | desired effect | | push w/ rapid, | wed by 360mg hrs | -5min give 150 | est | x3 | 9 | | | III INS | m I NC | | | | | | | | entilation | | | IVP/IO | r. (adjust for hncg/min | ymin IV/IO | | 10mg | 100-150mg | 60-120mg | 5-10mg | 20-40mg | 1000-7500 mcg/min | 100-250mg | 100-200mg | Bmor | | to-100mcg | 10-30 mg | | or slow IVP | LARGE ADULT (100KG)220LBS |
| Torsades De Pointes (with or without pulse) | Symptomatic Bradycardia | Symptomatic Bradycardia | Stable, narrow compley, tach/PSVT | Hemodynamically unstable V-Tach/SVT | Refractory Pulseless V- Fib/V-Tach | Pulseless Arrest | Angina/ AMI | Angina/ AMI | Hypoglycemia | Hypoglycemia | Diopriomano | Dromchodilator | Brenchodilator | | Anaphylaxis/Asthma | Antihistamine | A secondari annia | Antiemetic/Sedation | Antiemetic | Optotid OD | Beta/Ca-ch blocker OD | | Hypotension | Hypotension | Hypotension | | Maint of paralysis | RSI | RSI/Maint of paralysis | Sedation, Seizures | RSI (Non- analgesic) | General Anesthesia Maint. (Non-analgesic) | RSI/General Amesthesia (Non-analgesic) | Dissociative Sedation/ RSI | ram, Angootyne, Ann | Dain Anniolatic AM | Analgesia, Sedation, AMI | Analgesia | | Int/Ext Hemorrhage | INDICATIONS |
| AV Elocia | Use if refractive to Attopine Pacing | Glaucoma | May cause transient Asystole following push | Sinus Bradycardia, 2nd 3rd Deg Block | Sinus Bradycardia, 2nd 3rd Deg Block | Profusing Tachycardia's | Maintain SYS BP>90 | Nfust be chewed | Only if D50 not available | Intracranial bemorrhage | NO 1 A THE | Carolac arriyunna | Cardiac arrhythmia | | Do not use for head Tx | Alinor sedation | 11-12-0 | Altered LOC/ Vesicant | Can cause OT prolongation | Use minimum needed | Dosage is higher than kits | | Must be diluted / May resus w/ blood 1st | May resus w/ blood 1st | Must be diluted/ May resuscitation w/blood 1st | | Must maintain PT airway 40-90 min | RSI Only | Must maintain PT airway 30-60min | BP Resp drop | Repeat doses can cause adrenal suppression | Hypotension (up to 30% of MAP) | Hypotension (up to 30% of MAP) | HTN, emergence, avoid sub- dissociative doses | Vr marr (dom acticate | Not for Sameriald Trauma | Kesp depression | HTN, Emergence | | Gwe<3 hrs from inturvisurgery | RESTRICTIONS/ WARNINGS |
| 30 min | Maintain drip | 5-15min | 1-2 min | Maintain drip | 3-5 min | 3-5 min | 20-30 min | 4-6lus | UNK | UNK | 0-108 | 5 15m | 1-4hrs | | | 3-10mm 4-8hrs | 5 1/hurán | 4-6hrs | 4-6lus | 20-60 min | UNK | | 5-10 min | Consistent Infusion | Infusion | | 30-60 min | 5-9 min | 25-40 min | 10-30 min | 5-10 min | Infusion | 5-10 min | 10-20 min | | Viscishin | 30-00 mm | 10-30 min | | UNK | DURATION |
| May repeat doses, MAX 4g in 1 hour | HR >60, MAP >65 | MAX 3mg (3 doses) | Give 2nd dose if no rhythm change in 1-2 min | May repeat 150mg infusion q 10 PRN. Do not exceed 15mg/min | May 2 doses | Repeat Q 3-5min w/ CPR-No Max | May repeat up to max of 3 Doses | NO REPEAT | REPEAT every 20min PRN | Titrate; avoid over-correction | Theo ones 13 mm | Male arms 15 min | | | NO REPEAT | Add doses Q 5-10 mm mm mprove Max Some | Add data 0.5 10 min trail imments | Max 25mg q 4 hrs | Max 8mg q 6 hrs | Q 2-3 mm PRN (Max 10mg) | Titrate infusion for hemodynamics | | PRN to maintain SYS BP; start w/ low doses and titrate PRN | Start at lowest dose. Titrate up by 0.5mcg/min to MAP >60 | Start low; Titrate to desired response | | PRN q 30-60 min for paralysis | NO REPEAT | PRN q 25-40 min for paralysis | PRN q 15-30min if BP/Resp stable | NO REPEAT | Titrate to effect. MAX DOSE 100 mcg/kg/min | q5-10 min PRN | 0.5-2mg/kg Q 10-20 PRN for sedation or 1-3 mg/kg/hr infusion | EVAL TO SEGMENT TO LANSA SUDDE | DDM for moduli when if DD/D one stable | Q 30-60 mm PKN | Q 10-30 min PRN for pain | | NO REPEAT | REPEATABILITY/ MAX DOSE |

| ASYSTOLE PEA BRADYCARDIA | V-FBA-TACH | Cardionalmon: | Alternate K9 S | A gitated K9 Sedation IM 1st/IV PRN | Jul - weiteres Pijjy | Analgesia-Interr Analgesia-Contin | INDICATION | 1012 TULIN | Heart Rate | Blood Pressure | Respirations | Temperature | VITALS | | | nemaworidity | | | WBC M:13-18 F:12-16 | Hemoglobin(Hgb) | / | , | HEMATOLOGY | Arefoninanken | FFP (1u=200-250ml) | PRBC (1u=250ml) | Blood Products and Management | Hypertonic Saline (3%) | Burns >20% TBSA | | Maintenance | FLUIDS Resuscitation (Crystaloid) | Mannitol (20%6) | Magnesium Sulfate | Lorazepam | Diazepam | Multi-Use/Seizures/Other | Pralidoxime Chloride (2-Pam) (DuoDote ATNAA) | Atropine | CBRNE | Calcium Chloride (100mg/ml) | Labetalol | Sodium Bicarbonate | CARDIAC-continued | DRUG |
|-------------------------------------|--|---|---------------------------------------|--|--|--|----------------|---------------------|--------------------------|--------------------------------|---|--|------------------------|---------------------------|-------------------------------|--------------|----------------------------|-----------|---------------------|-----------------|---------------------------|-----------------------|-----------------------|----------------------------------|--|--|-------------------------------|-------------------------------------|--|---|-----------------------|---|-------------------------------------|--|---|--|--------------------------|--|---|-------|--|----------------------------|---|-------------------|------------------------------|
| LADYCARDIA | ng antanana hTACH | edation 1 vito | Alternate K9 Sedation IV/IO | m IM 1st/W PRN | Mild Sadation - Df | InaigesiaIntermittent IV/IO/IM | ATION | 35-45 mmHg | 08-09 | 120/80 (avg) mm Hg; MAP 90-100 | 16-30MIN | 101-103 TF | Normal | MILITAR | / | | 450,000 | | | | | | | 500 mg DO or 1 C+ IV | 10 ml/kg | 10 mJ/kg | | 0.1-1 ml/kg/hr | LN IO III - ALDOA (Dased on 40- song auun) | TO 10 w1 # 9/ TDC A /Dared on 40, 2015 adult) | 1-2 mJ/kg | <20 ml/kz | 1 Gram/kg IV over <20 min | 1-2 Gram IV/IO | 2-4 mg IV/I0 | Anxiety: 2-10 mg IV/IM q 6trs//Seizures: 5-10mg q 5-10min (MAX 30mg)//Seizures following Nerve agent Exposure: 10-20 mg IM for seizures or if 3x Mark 1 Kits used | - | 1-3 Auto-Injectors (600 mg ea) | 1-6 mg | - | Ca Gluconate can alternatively be used @ 3x doses listed here (except for Beta Blocker OD) | 10-20 mg IV/IO over 1-2min | l mEq/kg IV/IO | | STANDARD DOSING |
| | Defib 2 | | | | | | | | | | Panting | F E01 | Excited | MILITARY WORKING DOG | .45 | | | | | 3.5 | 135 | - | 200 | 500 | 1-2 units PRN to achieve | 1-2 u | | | Hour | Inter + As DCA Dec | 75ml/hr(TKO) | 250-500 1 | 60G | Seizures = 1-2G over 30 2G over 20 min | Seizures = | es: 5-10mg q 5-10min (MA 0 mg IM for seizures or if 3 | | Inject 1-3 injectors (bas Injector contain | | | 500-1000mg over 2-5 m >5min for Ci | 1 | 60mEq | | SMALL ADULT (60KG)132LBS |
| A | یں compressions ہے -5J/kg2min CPRDefi | Commessions @ | PRO | MIDAZOLAM 0.3mg/kg AN | VUDAZO | HYDROMO | | | | | give 25% > 20 min/PR | Shock: Calculate 90ml/k | | | 35-45 / 80-100 | | , | BL | K CO' | 3.5-5.0 22-26 | 135-145 95-105 | <u>-</u> | CHI | 500ms PO or ICram IV infusion | 1-2 unit: PRN to achieve 1:1 ratio w/ PRBC's (Shelf Life(thawed)=5 days) | 1-2 units PRN to achieve Sys BP90 Life =42 days) | | 250ml bolus followed by 50-100ml/hr | Hour | 10ml + 05 DCA Day | 105ml/hr(TKO) | 250-500 ml Bolus to achieve systolic BP >90 | 80G | Seizures = 1-2G over 30 min; Wheezing/Respiratory Distress (3rd line) = 2G over 20 min; (Pre)Eclampsia = 4-6G over 15-20min | Seizures = 4mg q 3-5 prn; Agitated/Combative Patient = 2-4mg q 30-60 | X 30mg)/// <u>Seizures follow</u> x Mark 1 Kits used | | Inject 1-3 injectors (based on severity of symptoms) IM. DuoDote/ATNAA Injector contains both Atropine (2.1mg) and 2-Pam(600mg) | 1-6mg | | 500-1000mg over 2-5 min for Hyper K issues; 20mg/kg>5-10min for Beta Blocker OD; 1Gram >5min for Ca Ch Blocker OD; 1G>5min after Blood | 10-20mg IV/IO over 1-2min | S0mEq | | ADULT (80KG)176LBS |
| ATROPINE 0.04mg/kg IV/IO AND EPINEP | Defib 2-5J/kg2min CPRDefibEPI 0.01mg/kg AND Amiodarone 5-10r | 100/min: establish airway: | 9.1 | 8 | FAM ON MORE IN AND T | RPHONE 0.1-0.2mg/kg Q | K9 M | Defib 2-5 Joules/Kg | Intubate w/ 10.0 ET tube | (PRN) | N); reassess; give 25% >10 | g for total fluid to be infu: | FLUID MANAGEMENT | | 22-26/ | HCO | / | BLOOD GAS | cre | 1 | 8-25 | BLIN | CHEMISTRIES | tinm | lf Life(thawed) =5 days) | IP ~90 | | ml/hr | | (0.000) an ann an 1 | 150ml/hr(TKO) | c BP >90 | 100G | ory Distress (3rd line) = over 15-20min | Combative | ing Nerve agent | | 15) IM. DuoDote/ATNAA ad 2-Pam(600mg) | | | ug/kg>5-10min for Beta in after Blood | B | 100mEq | | LARGE ADULT (100KG)220LBS |
| 10 AND EPINEPHRINE 0.01 mg/kg | Amiodarone 5-10mg/kg2min CP | Pasnirations @ 2-10/min for 7-3 minutes | ORN to allow catheterization or intub | KETAMINE 2ms/ks AND HYDROMORPHONE 0.2ms/ks | ALLENGTH IN THE SAME THE SAME THE SAME THE SAME AND A SAME THE SAM | HYDROMORPHONE 0.1-0.2mg/kg Q2-4ms OR MORPHINE 0.2-0.5 mg/kg Q4-dms | K9 MEDICATIONS | | ube | | give 25% > 20 min(PRN); reassess; give 25% >10 min(PRN); last 25% >10 min | Shock: Calculate 90ml/kg for total fluid to be infused: Give 25% > 10 min; seases; | NT | | -100/ ±2 | O2 Sat BE | ` | | / | 011-02 | 0 ⁻ | | A COLLEGE ACCESSION | Febrile Reaction | Int/Ext Hemorrhage/ AB+ Uni Donor | Int/Ext Hemorrhage/ O- Neg Uni Donor | | ICP Reduction | full-thickness burns | SONS/ TIDE A maniful as | IV access/Homeostasis | Hypo-tension/volemia | Mod to severe head Tx | Seizures/ Wheezing in Resp Distress/ (Pre)Eclampsia | Seizures/Agitated or Combative Patient | Anxiety/ Seizures/ Nerve Agent Seizures | | Organophosphate/ Nerve Agent | Orgamophosphate/ Nerve Agent | | Hyperkalemia/ Beta&Calcium Channel Blocker OD | HTN Urgency/Emergency | TCA OD; Prolonged Cardiac Arrest | | INDICATIONS |
| 8 | ng/kg2min CPRDefib2min CPRDefib | attors hefore AT S | ation | HONE 0.2ms/kg | M OD Alexa | g/kg Q4-6lars | | | | | D.O.P.E Displacemen | | Women (hg) | Men (kg) | | PEEP | TIDAL VOLUME | IE | Fi02 | RATE | MODE | | VEN | Indus alomby | Monitor for Anaphylaxis/Hyperthemnia | Monitor for Anaphylaxis Hyperthermia HyperK | | Use only in Head Injuries | THOCK STATE THREE AND ADDREED THREE | Teach start times and amount infined | Do not over hvdrate | Blood is 1st fluid choice | Avoid in HoTN Pts | Dilute into 50-100mil NS or D5W | IM not recommended due to erratic absorption | Respiratory Depression | | Use Atropine 1st if only using single dose 2-Pam (Afark 11NAAX Kit) | Requires large amounts of Atropine (5-20 boxes) | | Not used in confine treatment of cardiac arrest | Lower MAP by <20% | Do not mix with other meds/Flush line after | | RESTRICTIONS/ WARNINGS |
| | | | | | | | | | | | it; Obstructions; Pneumo | Troubleshooting | | | Ideal Body Weight Calculation | | | | | | | Initial Vent Settings | VENTILATOR MANAGEMENT | films | PRN | PRN | | N/A | N/A | | PRN | PRN | 3-8hrs | 30 min | 30-120min | 20-30min | | 15min | 5-15min | | 30min-4 hrs | 15-60 min | 1-2hrs | | DURATION |
| | | | | | | | NOTES | | | | D.O.P.E Displacement; Obstructions; Pneumothorax; Equipment Failure | | 45.5 + 2.3 x (Ht - 60) | 50.0 + 2.3 x (Ht - 60 in) | ation | 5 cm/H2O | 6mL/kg (Ideal body weight) | 1:02 | 1.0 (100%) | 14BPM | AC/ASV (Hamilton T1 only) | | | The only for Non-Templetic reart | Ideal ratio of FFP:PRBC:Platelets is 1:1:1 | Repeat PRN to maintain SYS BP >90/ MAP >60/hemostasis | | MAX 250ml | Add 100ml/hr for each 10kg over 80kg | A DESCRIPTION OF VARIANCE | Titrate to effect | Titrate to maintain SBP >90 | Follow with 0.25 Gram/kg IVP q 4hrs | 2 Grams/hr infusion needed following loading dose for Eclampsia | Slow IV Push, max rate 2mg/min | Max dose 30mg for seizures | | If symptoms remain after 15 min, re- inject subsequent doses (Max 1800mg 2 Pam) | Double dose if previous dose does not relieve secretions(atropinization) | | 20mg/kg/hr influsion for Beta OD; 1000mg Q 10- 20 x 3 doses PRN for Ca Chan Blocker OD | Repeat one time | Maint Infusion of 100-150mEq in 1L D5W @ 100-200ml/hr for TCA OD | | MAX DOSE/ REPEATABILITY |

| Α | LTITUDE ILLNES | SS |
|--|---|--|
| | Signs and Symptoms | |
| Acute Mountain Sickness (AMS) Headache | High Altitude Cerebral Edema (HACE) | Hight Altitude Pulmonary Edema (HAPE) |
| Nausea | Worse Mountain Sickness | Cough |
| Vomiting | Ataxia | Dyspea |
| Lethargy | Altered Mental Status | Pink Frothy Sputum |
| Dizziness | | Consider DifDx |
| | Treatment | |
| • For all: | | |
| O2 (100% FiO2) / IV/IO A | ccess / Cardiac monitor | |
| | ssible (Consider Gamow bag if unabl | e) |
| Hypothermia prevention | , S | , |
| Note to aircrew – fly lowe | st allowable altitude | |
| HAPE - Pulmonary Symptom | S | |
| Nifedipine 30mg ER PO d | 12hrs (or 20mg imm. rel. PO q8hrs) | |
| Consider Assisted Ventila | | |
| HACE - Headache <u>with</u> altered | | |
| 0 | O/PO x1, then 4mg q6hrs | |
| Pediatric: 0.15 mg/kg | | |
| Acetazolamide 250mg PC | • | |
| • AMS - Headache <u>without</u> alte | | |
| | 0 q12hrs (avoid with sulfa allergy/sick | le cell) |
| Severe AMS: Add Dexam | 3 | - i |
| | e (in isolation or with AMS/HACE) con | sider |
| Acetaminophen 650-100r Ibuprofon 600 800mg PO | 0 | |
| Ibuprofen 600-800mg PO | Notes, Warnings, Cautions | |
| • The treatment of choice for a | Il altitude-related illnesses is supplem | ental O2 and descent at least 500 |
| | a hyperbaric bag (Gamow bag) can l | |
| | on as HACE or HAPE are suspected, | |
| | | 0 0 0 0 |

- actively with the PIC or other tactical commander to work the issue of descent ASAP.
- HAPE and HACE are severe and cases should be hospitalized. AMS may be managed well with outpatient treatment.
- **High-Altitude Pulmonary Edema (HAPE)** patients may have crackles / fever / hypoxia. Be prepared to consider asthma, PE, pneumonia or other diagnoses as well.
- High-Altitude Cerebral Edema (HACE) patients have AMS and may have tremors, HACE may occur along with HAPE.
 - ANY altered mental status / confusion / abnormal gait should be presumed to have cerebral edema and descent should be undertaken immediately.
 - *Descent should be done with the least amount of patient exertion possible to prevent worsening of the condition.

ANIMAL & INSECT BITES / STINGS

Signs and Symptoms

- Rash, Skin break, Wound, Retained Stinger
- Pain, swelling, erythema.
- Bleeding or Discharge
- Shortness of breath / Wheezing / throat tightness
- Hypotension or Shock

<u>Treatment</u>

- Universal patient care Guidelines
- O2 (if hypoxemic)
- IV/IO in non-effected limb
- Cardiac monitor
- Position patient supine
- Elevate bitten extremities
- Wash wound w/ soap + water
- Mark suspected bite area (Circle area affected to monitor for spreading)
- Ice/Cold packs for swelling and pain
- Follow local / surgeon policy / CPG
- Allergic reaction?
 - o If yes us allergic reaction guideline
 - If needed Pain management guideline
- Spider, Scorpion, and Snakebites
 - Confirm receiving facility has adequate supply of the appropriate regionally specific antivenoms.

Notes, Warnings, Cautions

- Never attempt to capture / transport a live animal / insect.
- Anaphylactic reactions should be treated as soon as recognized.
- Review country environmental concerns before deployment or visitation.
- All animals should be considered rabid outside the U.S. until proven otherwise. This excludes rodents, which do not carry rabies.
- Consider IV administration of Calcium Gluconate if Tetany develops.
- DO NOT apply constricting bandages or tourniquets as these may worsen local tissue injury and increase the risk of permanent disability.
- DO NOT cut, suck, electrocute, burn, or use chemicals on the envenomation site

DECOMPRESSION SICKNESS

History

- Recent history of scuba diving
- Hypobaric chamber training
- High altitude parachutist training/operational. >18,000 ft (HALO, HAHO)
- High altitude exposure

Signs and Symptoms

- The Bends (Type 1)
 - Pain in the joints, muscles, and related tissues. Initially mild and/or intermittent but can become deep, gnawing, and eventually severe.
 - Pain tends to be progressive and becomes worse during ascent.
 - Larger joints such as the knees and shoulders are most frequently affected. The hands, wrists, and ankles also are commonly involved.
 - Unusual generalized fatigue, headache, malaise (constitutional bends)
- Skin manifestations (Type 1)
 - Paresthesia tingling, itching, and cold and warm sensations.
 - A mottled red rash might appear on the skin.
 - Rarely a welt might appear and be accompanied by a burning sensation.
 - Bubbles might develop just under the skin and cause localized swelling.
 - Affected regions with excess fat beneath the skin, soreness and abnormal fluid accumulation might be present for 1 or 2 days.

Chokes (Type 2)

- Symptoms occurring in the thorax caused in part by innumerable small bubbles that block smaller pulmonary vessels.
- Burning sensation under the sternum.
- As the condition progresses, a stabbing pain is felt, chest tightness, and inhalation becomes rapid and markedly difficult.
- Uncontrollable desire to cough. Cough is ineffective and nonproductive.
- Sensation of suffocation; breathing becomes shallower.
- Cyanosis
- CNS (Type 2)
 - Brain or spinal cord is affected by nitrogen bubble formation.
 - Visual disturbance (lights are flashing/flickering when they are steady).
 - Dull to severe headache.
 - Partial paralysis / one-sided numbness and tingling
 - Loss of orientation, and Inability to hear or speak.
 - Inner ear disturbance's (vestibular DCS) vertigo, nausea, vomiting. More likely associated with diving than altitude exposure.

<u>Treatment</u>

- If DCS occurs while in flight descend to a lower altitude or to the ground level
- Place patient on 100% O2 (for denitrogenating)
- Monitor vital signs and conduct a Neuro exam. Note any changes.
- Minimize any reactor movement.
- Start IV. 250ml/hr NS or LR.
- Definitive treatment is to bring patient to a **HYPERBARIC** chamber.
 - Contact a flight surgeon for coordination.
 - Bring patient to closes Hyperbaric chamber with a low flight profile.

Notes, Warnings, Cautions

- Onset can occur as long as 48 hours after exposure to altitudes above 18,000 ft.
- De-nitrogenation (Nitrogen concentration is reduced by breathing 100% O2. This allows no new nitrogen into the body while existing nitrogen is removed from the lungs eliminating much of the nitrogen dissolved in body tissues.)
- All Types of DCS should be treated as an emergency.

HOT/COLD WEATHER INJURY

| • • • | <u>HOT Weather Signs & Symptoms</u> AMS LOC Pale or clammy skin Hypotension or shock Seizure Nausea or Vomiting | Cold, Clammy skin Cold, Clammy skin Shivering or lack of shivering Mental status change Extremity pain or numbness Bradycardia or Arrhythmia Hypotension or Shock |
|------------------|--|---|
| | HOT Weathe | |
| • • • • | Remove from heat source/ loosen or remove clo AMS - <i>AMS guidelines</i> • Glucose check Consider intubation if needed. 1L IV Bolus or PO Fluids Monitor 12 lead EKG for arrhythmias. Be prepared for Seizure – <i>Seizure guidelines</i> | |
| • | AMS & core temp > 40C /104F Start aggressive cooling (Tepid water to skin once temp <40 C /104 F) Consider benzodiazepines to block/stop shive Midazolam 0.1 mg/kg AMS & core temp < 40C/104F Tepid water or room temp water to skin | and fanning / Ice packs in groin, axilla, and neck / D/C rering & rebound hypothermia. |
| • | Continuous monitoring | |
| | <u>COLD Weath</u> | er Treatment |
| • • • | Remove wet clothing Assess: Mental status, Rectal temperature, Gluc Core Temp < 35C/95F With AMS (arrhythmia, at | cose osence of shivering) urning pt) tively shivering) |
| HOT V | Veather | |
| <u>COLD</u> | Best method to cool pt is sublimation-sprinkling Elevated risk groups: Elderly, Very young, Highly Sweating does not exclude heat stroke / heat illr <u>Weather</u> "no patient is dead until they are warm and dead Hypothermia is defined as core temp <35C/95F Pulse may be very slow in hypothermia patient - At temps < 30C/86F one defibrillation can be att 30C/86F | v active ness " |

SUBMERSION INJURY

Signs and Symptoms

- Unresponsive
- Mental Status Changes
- Hypoxia
- Cyanosis
- Hypothermia
- Vomiting
- Coughing

<u>Treatment</u>

- O2 (100% FiO2 for all injuries)
- IV/IÒ, Guideline
- Cardiac monitor / check for Arrhythmias
- Spinal immobilization protocol
- If hypothermic use Hot/Cold weather Guideline
- If patient has injuries use Multiple Trauma Guideline
- Reassess Airway

Notes, Warnings, Cautions

- If Decompression Illness or arterial gas embolism is suspected and neurological deficits (including altered mental status) are present, consider high-flow oxygen, lidocaine 1.5 mg/kg IV / IO, and aspirin 325mg. While these interventions remain unproven, the risk / benefit ratio makes them acceptable options, particularly if time to hyperbaric chamber is anticipated to be prolonged.
- Rapid hypothermia from cold water immersion in children has resulted in survival despite prolonged downtime - resuscitate per appropriate protocols and rapidly transport. This has not been seen in adults.
- All near-drowning victims should be transported for evaluation due to potential for worsening respiratory status over next several hours.
- Drowning is the leading cause of death among would-be rescuers.
- Head-first diving injuries often associated with unstable Jefferson fracture (burst fracture of C1) due to axial load. Patients found with suspicion of this type of injury should have early and careful C-spine immobilization.
- Altitude should be restricted in patients suffering from decompression illnesses to prevent worsening. Should remain <1000 ft. AGL / 10,000 ft. MSL whenever possible.
 - Aggressive pre-planning for access to hyperbaric treatment facilities is encouraged if mission requirements warrant it.

ACETAMINOPHEN

QC, Lactation Yes (Caution)

Trade Name: Tylenol

Class / Mechanism of Action

Analgesic

Blocks cyclooxygenase (COX 1 and 2) enzymes, resulting in reduced formation of prostaglandin precursors. Blocks formation of prostaglandin derivative, thromboxane A2, resulting in inhibited platelet aggregation. Has antipyretic, analgesic, and anti-inflammatory properties.

Indications

Labeled Indications: Treatment of mild to moderate pain and fever, Treatment of moderate to severe pain when provided via IV with opioid analgesia

Contraindications

- Hypersensitivity to acetaminophen or any component of the formulation
- Hepatic impairment or liver disease

Adverse Reactions / Precautions

- Use IV form cautiously in volume depleted patients
- Avoid use in patient suffering alcohol toxicity, known alcohol abuse, or renal impairment
- IV form can cause nausea and vomiting (especially in adults), headache

| Dose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|---|---------------|--|
| Pain or fever: (Limit total daily dose t PO: Regular release: 325-500 mg eve or 1000 mg 3-4 times daily (maxing) | ery 4-6 hours | Pain or fever: PO: Infants and Children <12 years: 10-15 mg/kg/dose every 4-6 hours as needed; do |
| IV: • 500-1000 mg every 6 hours | | not exceed 5 doses (4 g/day) in 24 hours IV: Infants and children <2 years 15 mg/kg every 6 hours Children 2-12 years: 15 mg/kg every 6 hours s Max single dose: 15 mg/kg/dose (≤750 mg/dose) Max daily dose: 75 mg/kg/day (≤3.75 g daily) |
| | | Note: Children ≥12 years & Adolescents: Refer to adult dosing |

| ACETAZOLAMIDE QC, Lactation | on Yes (Caution) Trade Name: Diamox |
|--|---|
| Class / Mechanism of Action | |
| Diuretic, Carbonic Anhydrase Inhibitor; Antico Inhibits carbonic anhydrase causing a decrease in | nvulsant n hydrogen ion renal secretion with increased renal water. Onset of action PO: 2 hours, IV 5-10 minutes |
| Indications | |
| Labeled Indications: Prevention or treatment of symptoms of acute Edema due to congestive heart failure | mountain sickness |
| Contraindications | |
| Hypersensitivity to acetazolamide, sulfonamid Confirmed low sodium / potassium levels other | - · · |
| Adverse Reactions / Precautions | |
| May worsen respiratory acidosis Drowsiness, deceased alertness, impairment Flushing of skin, allergic skin reaction, skin ph | otosensitivity |
| Dose and Administration: ADULT | PEDIATRIC Always Reference LB tape |
| <u>Altitude illness (Acute Mountain Sickness):</u> PO: 125-250 mg twice daily. Note: For high altitude cerebral edema (HACE), dexamethasone is the primary treatment; however acetazolamide can be used (together with dexamethasone) at the AMS dose. <u>Edema</u> (Only with referring doctor or medical director instruction): PO, IV: 250-375 mg once daily | Altitude illness (Acute Mountain Sickness): PO: (IM not recommended due to alkaline pH) • 2.5 mg/kg/dose every 8-12 hours • MAX dose 250mg/dose. T, Note: For high altitude cerebral edema (HACE), dexamethasone is the primary treatment; however, acetazolamide can be used (together with dexamethasone) at the AMS dose. |

ACETYLSALICYLIC ACID **QC, Lactation Yes** (Short Term or Trade Name: Aspirin Low Dose OK) **Class / Mechanism of Action** Nonsteroidal Anti-inflammatory Drug (NSAID) Blocks cyclooxygenase (COX 1 and 2) enzymes, resulting in reduced formation of prostaglandin precursors. Blocks formation of prostaglandin derivative, thromboxane A2, resulting in inhibited platelet aggregation. Has antipyretic, analgesic, and anti-inflammatory properties. Indications Labeled Indications: Treatment of acute coronary syndromes (ST-elevation MI, non-ST-elevation MI, unstable angina), acute ischemic stroke, and transient ischemic episodes. Contraindications Hypersensitivity to salicylates, other NSAIDs, or any component of the formulation • • Asthma, Rhinitis Inherited or acquired bleeding disorders (including factor VII and factor IX deficiency) Do not use in children less than 16 years old (Reye's syndrome) Adverse Reactions / Precautions Not for use on trauma patients in the combat environment. Risk of bleeding: Avoid use in patients with known or suspected: Bleeding disorders, GI Bleed, GI • Ulcers, patients taking Coumadin, or within 24hrs of taking Alteplase (tPA) for suspected stroke **Dose and Administration:** ADULT PEDIATRIC Always Reference BROSELOW Tape N/A: Acute coronary syndrome (ST-segment elevation myocardial infarction [STEMI], Contraindicated in children under 16 yrs. unstable angina (UA)/non-ST-segment elevation (Reye's Syndrome) myocardial infarction [NSTEMI]): (Not for use in trauma patients): PO: 162-325 mg (chew nonenteric-coated aspirin • as a single 325 mg tablet or x4 81 mg chewable tablets)

| ACTIVATED CHARCOAL QSafe, | Lactation Safe Trade Name: Actidose |
|---|--|
| Class / Mechanism of Action | |
| Antidote Non-absorbable agent that absorbs toxins within the | Cl tract inhibiting Cl character |
| Indications | Gi tract infibitung Gi absorption. |
| Labeled Indications: Management of suspected or k | nown poisonings when gastrointestinal |
| decontamination is an option. | thown poisonings when gastrointestinal |
| • Decontamination within 1 hour of ingestion of toxi | c substance |
| Contraindications | |
| • Presence of intestinal obstruction or GI tract not a | anatomically intact |
| • Patients at risk of GI hemorrhage or perforation | |
| Patients with an unprotected airway (e.g., CNS de in an and the right and accurate of a minutes) | epression without intubation) or if use would |
| increase the risk and severity of aspiration | |
| Adverse Reactions / Precautions | trol and must utilized NC/OC tube |
| If patient unconscious, must establish airway con Be prepared for possible emesis. Consider use or | |
| Avoid use in patients at risk of GI hemorrhage or | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Aguta Baiganing | |
| Acute Poisoning: PO, NG/OG: | Acute Poisoning: Children >12 years: Refer to adult dosing. |
| Single dose: 25-100 grams | PO, NG/OG: |
| | • Single dose: 1 gram/kg |
| Note: Activated Charcoal has limited efficacy if not | • Multidose: Initial dose: 1Gram/kg initially, |
| utilized within 1 hour of toxin ingestion. Risk-benefit | followed by multiple doses of 0.5 Gram/kg |
| of charcoal must seriously be considered because it does not work for all poisons, it must be given early | every 2 hours |
| when the poison is still in the stomach, it does not | Note: Some products may contain sorbitol. Co- |
| fully bind all poisons, and serious complications can | administration of a cathartic, including sorbitol, is |
| occur with aspiration. Aspiration can occur if | no longer recommended. |
| deteriorating mental status and/or vomiting. | |
| Note: Some products may contain sorbitol. Co- | Note: Activated Charcoal has limited efficacy if not utilized within 1 hour of toxin ingestion. |
| administration of a cathartic, including sorbitol, is | |
| no longer recommended. | |
| Note: Multidose charcoal is indicated if patient | |
| ingested a life-threatening amount of drug | |
| (carbamazepine, dapsone, phenobarbital, guanine, | |
| or theophylline) | |
| | |
| | |
| | |
| | |

ADENOSINE

QC. Lactation Yes (Caution)

Trade Name: Adenocard®

Class / Mechanism of Action

Antiarrhythmic Agent

Slows conduction time through the AV node, inhibits re-entry pathways through the AV node, restoring normal sinus rhythm. The half-life of under 10 seconds allows for rapid repeat dosing.

Indications

Labeled Indications: Paroxysmal supraventricular tachycardia (PSVT) when clinically advisable, vagal maneuvers should be attempted first; not effective for conversion of atrial fibrillation, atrial flutter, or ventricular tachycardia.

Unlabeled: ALS/PALS Guidelines (2020): Stable, narrow-complex regular tachycardias; unstable narrow-complex regular tachycardias while preparations are made for synchronized direct-current cardioversion; stable regular monomorphic, wide-complex tachycardia as a therapeutic (if SVT) and diagnostic maneuver.

Contraindications

- Hypersensitivity to adenosine or any component of the formulation
- Second- or third-degree AV block, sick sinus syndrome, or symptomatic bradycardia (except in patients with a functioning artificial pacemaker)
- Use in patients with atrial fibrillation/flutter with underlying Wolff-Parkinson-White (WPW) syndrome (Fuster, 2006); asthma (ARC, 2020)
- Known or suspected bronchoconstrictive (Asthma) or bronchospastic lung disease.

Adverse Reactions / Precautions

- May cause transient asystole and new arrhythmia after cardioversion (PACs, AF, PVCs) chest discomfort
- Headache, Dizziness, Flushing, Gl upset
- Dyspnea, Bronchospasm in asthmatics

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|--|---|
| Paroxysmal supraventricular tachycardia: | Paroxysmal supraventricular tachycardia: |
| I.V. (rapid push, over 1-2 seconds, via proximal peripheral line (forearm or above, large bore). | IV/IO as close to core as possible (rapid push, over 1-2 seconds, see Note): Follow each dose with 10- |
| Initial: 6 mg; if not effective within 1-2 minutes, 12 mg may be given if needed (maximum single dose: 12 mg). | 20 mL normal saline flush. Initial: 0.1 mg/kg (maximum initial dose: 6 mg); if not effective within 1-2 minutes, administer 0.2 mg/kg (maximum single dose: 12 mg). Follow |
| Notes): Follow each dose with 10-20 mL normal saline flush. | each dose with 5-10 mL normal saline flush. |
| Note: Initial dose of adenosine should be reduced to 3 mg if patient is currently receiving carbamazepine or dipyridamole, has a transplanted heart or if adenosine is administered via central line (ARC, 2020). | |
| Note: Adenosine effects are antagonized by caffeine and theophylline, and patients may require higher doses. | |

| ALBUTEROL | QC, Lactation Yes | Trade Name: Proventil / Ventolin |
|---|--------------------------|--|
| Class / Mechanism of Action | | |
| Beta ₂ Agonist (Bronchodilator) Synthetic sympathomimetic that relacing cardiac impact. Onset of action is 2 | | h muscle, causing bronchodilation, with little |
| Indications | | |
| Labeled Indications: Treatment or airway disease; prevention of exerce • Asthma • Reactive Airway / Bronchospase • COPD | ise-induced bronchos | ospasm in patients with reversible obstructive spasm |
| May also be used in Crush Syn | drome (Hvperkalemia |) |
| Contraindications | | / |
| Hypersensitivity to albuterol or a Symptomatic tachycardia | any component of the | formulation |
| Adverse Reactions / Precautions | | |
| Headache, Dizziness, Flushing, Angina, A-Fib, Arrhythmia, Che Dyspnea, Bronchospasm in ast Dose and Administration: | st Pain, Palpitations | PEDIATRIC Always Reference BROSELOW Tape |
| | | |
| Bronchospasm: | | onchospasm: |
| Metered-dose inhaler (90-180 mcg/ | - | etered-dose inhaler (90 mcg/puff): |
| • 6 puffs | • | 6 puffs every 4-6 hours as needed |
| Solution for nebulization: | Sc | olution for nebulization: |
| • 5 mg | • | 5 mg |
| Respiratory Distress (acute, seve | ere): | Children ≥12 years: Refer to adult dosing. |
| Metered-dose inhaler: | | cacerbation of asthma (acute, severe): |
| 1.0 | | |
| • 1-2 puffs | | etered-dose inhaler (90 mcg/puff): |
| 1-2 puffs Solution for nebulization: 2.5-5 mg | • | etered-dose inhaler (90 mcg/puff): Children <12 years: 4-8 puffs every 20 minutes for 3 doses, then every 1-4 hours as needed |
| Solution for nebulization: | | Children <12 years: 4-8 puffs every 20 minutes |
| Solution for nebulization: | • | Children <12 years: 4-8 puffs every 20 minutes for 3 doses, then every 1-4 hours as needed |
| Solution for nebulization: | • | Children <12 years: 4-8 puffs every 20 minutes for 3 doses, then every 1-4 hours as needed Children ≥12 years: Refer to adult dosing. |

AMIODARONE

$\ensuremath{\textcircled{}}$ **D, Lactation: Yes,** Not Recommended

| | AMIODARONE | | | | | |
|--|--------------------------------|-------------|--------------------|-----------|-----------|--|
| Initia | Initial Dose: I50mg over I0min | | | | | |
| MIX 150 mg/100 ml CONCENTRATION 1.5 mg/ml | | | | | | |
| Dose | Rate | Micro Macro | | | | |
| | | 60 | 20 gtt/m | 15 gtt/ml | 10 gtt/ml | |
| mg/min | ml/min | gtt/min | gtt/min | gtt/min | gtt/min | |
| 15 | 10 | 600 | 200 150 100 | | | |
| Macro-Drip (10gtt/ml) is set of choice for this infusion | | | | | | |
| Set rate provides complete initial infusion of 150mg over 10 minutes. May repeat Q 10 min PRN if VT recurs | | | | | | |

| Maint Dose: 1mg/min over 6 hrs (360mg over 360min) | | | | | |
|--|-----------------|-------------|-----------|-----------|-----------|
| MIX 360 mg/500 ml CONCENTRATION 0.72 mg/ml | | | | | |
| Dose | Rate | Micro Macro | | | |
| | | 60 | 20 gtt/ml | 15 gtt/ml | 10 gtt/ml |
| mg/min | ml/min | gtt/min | gtt/min | gtt/min | gtt/min |
| 1 | 1.4 84 28 21 14 | | | | |
| Macro-Drip 20gtt/ml) is set of choice for this infusion | | | | | |
| Set rate provides maintenance infusion of 360mg over 6hrs. | | | | | |

| ATROPINE Sulfate | Lactation: Yes, Use Caution | Trade Name: AtroPen |
|---|--|---|
| Class I Mechanism of Action | | |
| acetylcholine at parasympathetic site output, and dries secretions. Atropine | Antidote for Carbamate Anticholing es in smooth muscle, secretory glands, e reverses the muscarinic effects of ch on but does not affect the nicotinic rece vsis. | , and the CNS; increases cardiac nolinergic poisoning. Reverses |
| Indications | | |
| Labeled Indications: Treatment of Symptomatic Sinus Bradycardia, Antidote for anticholinesterase poinsecticides) | AV block (nodal level) oisoning (carbamate insecticides, nerve | e agents, organophosphate |
| Contraindications | | |
| Pyloric stenosis Prostatic hypertrophy Note: NO contraindications should | y component of the formulation ons between the iris and lens (ophthalm uld prevent use of atropine in setting of arbamate, or nerve agent poisoning | |
| Adverse Reactions I Precautions | | |
| Tachycardia and arrhythmia (VTa Dilated Pupils, Angle-closure glau Headache, Dry Mouth, constipatio Paradoxical Bradycardia noted | on, urinary retention, flushing. | |
| Dose and Administration: A | DULT PEDIATRIC A | Nways Reference BROSELOW |
| Symptomatic Bradycardia | Symptomatic Brady | <u>vcardia</u> |
| 1 mg every 3-5 minutes, not to ex 3 mg or 0.04 mg/kg (ARC, 2020) Organophosphate or carbamate in nerve agent poisoning: | Maximum single | nimum dose is 0.1 mg. e dose of 0.5 mg. May 8-5 minutes. Maximum total ALS, 2020) |
| IV/IO: (Nerve agent) Atropine 20mg it trate to dry respiratory secretions. IV/IM: (Used with 2-Pam Chloride aut Initial: 1-6 mg; repeat every 3-5 m needed, doubling the dose if predid not induce atropinization. Marepeat doses as needed for 2 2-based on recurrence of sympton IM (AtroPen®): Follow with 2-Pam C injector. Mild symptoms (>2 mild symptom mg once an exposure is known of strongly suspected. Severe symptoms (>1 severe symptom (>1 severe symptom severe severe severe severe severe sympton severe severe severe severe severe se | IV/IO: Initial: 0.0 minutes as need dose does not in with repeat dose based on recurred based based based on recurred based b | y 3 min. Monitor Patient for toms of atropinization, (drying s). Once clinical improvement rict to 10- 20% of original |

CALCIUM Chloride 10%

9 10% QSafe, Lactation Safe

Class I Mechanism of Action

Calcium Salt, Electrolyte Supplement

Moderates nerve and muscle contractility via action potential excitation threshold regulation

Indications

Labeled Indications: Treatment of hypocalcemia and conditions secondary to hypocalcemia (eg, tetany, seizures, arrhythmias); emergent treatment of severe hypermagnesemia; massive transfusion prophylaxis **Unlabeled:** Calcium channel blocker overdose; beta-blocker overdose (refractory to glucagon and high-dose vasopressors); severe hyperkalemia (K+ >6.5 mEq/L with toxic ECG changes) [ALS guidelines]; malignant arrhythmias (including cardiac arrest) associated with hypermagnesemia [ALS guidelines]

Contraindications

- Known or suspected digoxin toxicity
- Not recommended as routine treatment in cardiac arrest (includes asystole, ventricular fibrillation, pulseless ventricular tachycardia, or pulseless electrical activity)
- Hypercalcemia

Adverse Reactions I Precautions

- Hypokalemia: Use with caution in patients with severe hypokalemia. Acute rises in calcium can cause life-threatening arrhythmias
- Rapid push can cause: Arrhythmia, bradycardia, cardiac arrest, hypotension, syncope, vasodilation
- Use small IV I Large Vein, flush prior and after, AVOID Extravasation (will cause tissue necrosis)
 - In general, IV Calcium Gluconate is preferred over IV Calcium Chloride in nonemergency settings due to the potential for extravasation with calcium chloride
- Do not infuse calcium chloride in the same I.V. line as phosphate-containing solutions.
- Precipitates with NaHCO3 in IV Bag/Tubing

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
|---|--|--|
| Cardiac arrest or cardiotoxicity in the presence of hyperkalemia. hypocalcemia. or hypermagnesemia: IV/IO, SLOW • 500-1000 mg over 2-5 minutes Beta-blocker overdose. refractory to glucagon | <u>Cardiac arrest or cardiotoxicity in the presence</u> of hyperkalemia. hypocalcemia. or hypermagnesemia: IV/IO, SLOW 20 mglkg (maximum: 2000 mg/dose); may repeat as necessary. | |
| and high-dose vasopressors (unlabeled use): IV/IO 20 mg/kg over 5-10 minutes followed by an infusion of 20 mg/kg/hour titrated to adequate hemodynamic response. | Calcium channel blocker overdose (unlabeled use): IV/IO | |
| Calcium channel blocker overdose (unlabeled use) (CaCl preferred over Calcium Gluconate for this use): IV/IO | Initial: 20 mglkg (0.2ml/kg) (maximum: 1000 mg/dose) over 10-15 minutes; may repeat every 10-15 minutes | |
| Initial: 1000mg over 5 minutes; may repeat every 10-20 minutes with 3-4 additional doses; or a continuous infusion of 2-6 grams/hour may be initiated | Note: Adult and Pediatric dosages are expressed in terms of the <u>calcium chloride salt</u> based on a solution concentration of 100 mg/mL (10%) containing 1.4 mEq (27 mg)/mL elemental calcium. | |
| Hypocalcemia prophylaxis from massive transfusion 10ml (10cc) 10% solution over 5 minutes Damage Control Resuscitation: IV/IO, SLOW 1000 mg after 1st blood unit and after every 4th unit. | (1gram = 10cc of a 10% solution) Note: Calcium Chloride is 3X more potent than Calcium Gluconate and therefore lower doses of Calcium Chloride must be used to reach similar therapeutic doses | |
| May be given before TXA | | |

- -

| CALCIUM GIUCONATE QSafe, Lactatio | n Safe |
|--|--|
| Class / Mechanism of Action | |
| Calcium Salt, Electrolyte Supplement Moderates nerve and muscle contractility via regulation of a | action potential excitation threshold. |
| Indications | · · · · · |
| Labeled Indications: Treatment of hypocalcemia and cond arrhythmias); cardiac disturbances secondary to hyperkale prophylaxis Unlabeled: Calcium channel blocker overdose; treatment of | mia; magnesium sulfate overdose; massive transfusion |
| Contraindications | |
| Ventricular fibrillation | |
| Hypercalcemia | |
| | triaxone in neonates (risk of precipitation of calcium- |
| Adverse Reactions / Precautions | |
| Hypokalemia: Use with caution in patients with se life-threatening arrhythmias | evere hypokalemia. Acute rises in calcium can cause |
| Rapid push can cause: Arrhythmia, bradycardia, o Do not exceed 200mg/min except in eme | cardiac arrest, hypotension, syncope, vasodilation |
| Caution in patients receiving digoxin therapy, may | 0 , |
| | , AVOID extravasation (will cause tissue necrosis) |
| • | erred over I.V. calcium chloride in nonemergency |
| • Do not infuse calcium chloride in the same I.V | . line as phosphate-containing solutions. |
| Precipitates with NaHCO₃ in IV Bag/Tubing | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Cardiac arrest or cardiotoxicity in the presence of hyperkalemia, hypocalcemia, or hypermagnesemia: IV/IO, SLOW 1500-3000mg over 2-5 minutes | Cardiac arrest or cardiotoxicity in the presence of hyperkalemia, hypocalcemia, or hypermagnesemia: IV/IO, SLOW 60-100 mg/kg/dose (maximum: 3000 mg/dose) |
| Calcium channel blocker overdose (off-label use): Hypotension/conduction disturbances: | Calcium channel blocker overdose (unlabeled use): Hypotension/conduction disturbances: |
| IV/IO | IV/IO |
| 3 Grams (3000mg) over 5 minutes; may repeat every 10-20 minutes with 3-4 additional doses. | 45 mg/kg (maximum 3000mg/dose) over 10-15 minutes; may repeat every 10-15 minutes |
| Hypocalcemia prophylaxis from massive transfusion 30mL of 10% solution over 5 minutes | Hypocalcemia prophylaxis from massive transfusion |
| Note: Calcium Chloride is 3X more potent than Calcium Gluconate and therefore higher doses of Calcium | 60mg/kg (maximum 30ml of 10% solution) over 5 |
| Gluconate must be used to reach similar therapeutic | minutes |
| doses. <u>Hydrofluoric Acid Exposure</u> – (off-label, see Burn | Note: Calcium chloride may provide a more rapid increase of ionized calcium in critically ill children. |
| SMOG) | |
| Topical therapy: After thorough irrigation , a CaGlu gel (75mL KY Jelly + 25mL 10% CaGlu) can be made and | |
| applied to the affected area, left on for 30 minutes, | |
| cleaned off, and repeated every 4 hours. Assess for pain relief and monitor EKG. (NO Calcium Chloride!) | |

| CEFAZOLIN | C , Lactation Yes | Trade Name: Ancef |
|---|--|--------------------------------|
| Class I Mechanism of Action | | |
| Antibiotic (Cephalosporin 1 st Gen) Bactericidal - Inhibits bacterial cell wall which inhibits cell wall biosynthesis, ca | | of penicillin-binding proteins |
| Indications | | |
| Labeled Indications: Used for infection prophylaxis. | n control prophylaxis for traumatic o | pen injuries and surgical |
| Contraindications | | |
| the formulation | cephalosporin antibiotics, other beta with penicillin allergies. Use with ca | |
| Adverse Reactions I Precautions | | |
| | y result in fungal or bacterial superinf ecially in nutritionally deficient, hepa | |
| Dose and Administration: ADUL | т | PEDIATRIC |
| Infection Control: Routine dosing may be based on body 1g if weight <80kg 2g if weight 81-160 kg (177-352 lbs), 3g if weight > 160 kg (>352 lbs) Max dose is 12g per day | IV: • 20-30 mg/kg IV (maximum, 100 | |
| War wounds (dirty wounds), 2g in 250 over 5 min every 8 hours for 24 hours i for most dirty wounds of the head and and extremities. | is adequate | |
| IV: | | |
| Adults: | | |
| 1-2g every 6-8hrs Max daily dose: 12 g/day Note: See antibiotic chart far dosing in a injury. | ccordance with | |

| DEXAMETHASONE QC, Lactation ? | (Not Recommended) Trade Name: Decadron | | |
|---|--|--|--|
| Class I Mechanism of Action | | | |
| Systemic Corticosteroid | | | |
| Anti-inflammatory, Immunosuppressant Onset of actio | n, IV: Prompt; Duration IV: 72 hours | | |
| Indications Labeled Indications: | | | |
| • Anti-inflammatory or immunosuppressant in treat | ment of a variety of diseases: allergic, dermatologic, c, renal, respiratory, rheumatic, and autoimmune | | |
| • Treatment of acute mountain sickness (AMS) and | high-altitude cerebral edema. | | |
| Contraindications | | | |
| Hypersensitivity to dexamethasone or any compoSystemic fungal infection, cerebral malaria | nent of the formulation | | |
| Adverse Reactions I Precautions | | | |
| Not for use in treatment of head injury, increased mortality has occurred in head injury patients treated with high dose IV methylprednisolone. Corticosteroids should not be used in head injuries. | | | |
| Dose and Administration: ADULT Tape | PEDIATRIC Always Reference BROSELOW | | |
| Acute mountain sickness (AMS)High altitude cerebral edema (HACE) (unlabeled use): PO, IM, IV: AMS: 4 mg every 6 hours HACE: 8 mg as a single dose; followed with: 4 mg every 6 hours until symptoms resolve | Acute mountain sickness (AMS)High altitude cerebral edema (HACE) (unlabeled use): PO, IM, IV: 0.15 mg/kg dose every 6 hours consider use in high altitude pulmonary edema because of associated HACE with pulmonary edema | | |

| ? Trade Name: Glutose I B-D Glucose | | | | |
|---|--|--|--|--|
| Class I Mechanism of Action | | | | |
| n, regulated by insulin. Rapidly increases blood enting ketosis, and promotes glycogen deposition in se: 10 minutes Maximum | | | | |
| | | | | |
| Labeled Indications: Treatment of: Hypoglycemia: Doses may be repeated in severe cases Hyperkalemia: (Must be used in combination WITH Insulin) Contraindications Known Hyperglycemia, otherwise None in the Pre-hospital setting Adverse Reactions I Precautions Most adverse effects associated with excessive dose or infusion rate. If evidence of malnutrition or alcohol abuse, thiamine should be given 1^{st.} Tissue Necrosis if Extravasation occurs; immediately D/C and change IV site. Hyperglycemia Hypokalemia | | | | |
| PEDIATRIC Always Reference BROSELOW | | | | |
| Таре | | | | |
| Hypoglycemia: Oral: 4-20 g as a single dose; may repeat if necessary. IV: Newborns: 5ml/kg D10 (Max 25 G/dose) Infants and Children: 2ml/kg D25 (Max 25 g/dose) Adolescents: Refer to adult dosing Note: D25= 25ml NS + 25ml D50 (12.5g in 50ml's solution) D10= 100ml NS + 25ml D50 (12.5g in 125ml's solution) or 40ml NS + 10ml D50 (5g in 50ml's solution) | | | | |
| | | | | |

DIAZEPAM

QD Lactation Yes (Unsafe)

Trade Name: Valium

Class I Mechanism of Action

Benzodiazepine:

Acts as an Anxiolytic/Hypnotic, anticonvulsant, and sedative - Long Half Life (25-100hrs) Onset of action: IV, Almost Immediate

Duration: IV, 20-30 minutes

Indications

Labeled Indications:

- Anxiety Disorders
- Convulsive Disorders and Alcohol Withdrawal Symptoms
- Skeletal Muscle Relaxant
- Induce Sedation and Amnesia (Midazolam is primary medication)

Contraindications

- Hypersensitivity to diazepam or any component of the formulation or other benzodiazepines
- Acute narrow angle glaucoma, Acute Alcohol Intoxication
- Respiratory Insufficiency/Depression (Overdose Reversal: FLUMAZENIL can be used; however, it carries elevated risk. Respiratory support until the medication is metabolized is traditionally the best care in Benzodiazepine overdose)
- Neurologic Depression (Head Trauma)

Adverse Reactions I Precautions

- No Analgesic properties (Narcotic pain control is needed for RSI'd / Intubated trauma patients)
- May Cause Respiratory depression: Do not give without stable IV line and BVM (airway control) ready.
- Hypotension, vasodilation
- Amnesia, confusion, drowsiness, slurred speech (Paradoxical Reactions possible: aggressiveness, agitation, anxiety, inappropriate behavior)

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW |
|---|---|
| Таре | |
| Anxiety: | Sedation I Muscle relaxation I Anxiety: |
| Oral, IV, IM: (Oral and IV doses more reliable) | IV, IM (IV doses more reliable) |
| • 2-10 mg 2-4 times/day if needed. <u>Status Epilepticus:</u> IV: (SLOW) | Children: 0.04-0.3 mg/kg dose every 2-4 hours to a maximum of 0.6 mg/kg within an 8-hour period if needed. |
| 5-10 mg every 5-10 minutes given over 3 minutes (maximum dose: 30 mg) Sedation in ICU patient: | <u>Status Epilepticus:</u> IV: |
| IV: Loading dose: 5-10 mg; Maintenance dose: 0.03-0.1 mg/kg every 30 minutes to 6 hours Muscle Spasm: IV: | Infants >30 days and Children <5 years: 0.2- 0.5 mg given slowly every 2-5 minutes (maximum total dose: 5 mg); repeat in 2-4 hours if needed. |
| Initial: 5-10 mg; then 5-10 mg in 3-4 hours, if necessary. Larger doses may be required if associated with tetanus. | Children 25 years: 1 mg given slowly every 2-5 minutes (maximum total dose: 10 mg); repeat in 2-4 hours if needed. |
| <u>Nerve Agent Exposure (CBRNE)</u> IM: | Muscle spasm associated with tetanus: IV, IM |
| 10-20mg for seizures associated with Nerve Agent exposure (up to 40mg may be needed); or if 3 MARK 1 kits were used on a | Infants >30 days and Children <5 years: 1-2 mg dose every 3-4 hours as needed. |
| casualty | Children 25 years: 5-10 mg dose every 3- 4 hours as needed |

| DILTIAZEM | Lactation? (Not R | ecommended) | Trade Name: Cardizem |
|---|--|--|---|
| Class I Mechanism of Action | | | |
| Calcium Channel Blocker; Antiarrhy Inhibits calcium ion from entering the ' muscle and myocardium during depol and coronary vasodilation; increases Onset of action: IV: 3 minutes, Duratio | "slow channels" or s larization; produces myocardial oxygen | select voltage-sensitive relaxation of coronary | vascular smooth muscle |
| Indications | | | |
| Labeled Indications: Atrial fibrillation supraventricular tachycardia, hyperter Unlabeled: Hypertrophic cardiomyopa tachycardia or ventricular premature b | nsion, chronic stabl athy; Idiopathic vent | e angina, vasospastic ricular tachycardia; Nor | angina. nsustained ventricular |
| Contraindications | | | |
| Sick sinus syndrome (except in padegree AV block Atrial fibrillation or flutter associate Severe hypotension; Cardiogenic s Ventricular tachycardia (with widedetermine whether origin is suprav | d with accessory by shock; Hypersensitiv -complex tachycard | pass tract (WPW, short vity to diltiazem or any f ia [ORS 20.12 seconds | PR syndrome) ormulation component |
| Adverse Reactions I Precautions | | , | |
| Cardiovascular: Edema, atrioventri Central nervous system: Headache Dose and Administration: ADUL Tape | e, dizziness, pain, n | ervousness, vomiting, v | |
| Atrial fibrillation or atrial flutter. Note: For rate control in hemodyn patients. Do not use in patients wit associated with an accessory pathw lead to ventricular arrhythmias. IV: Bolus dose: 0.25 mg/kg over 2 m (average dose: 20 mg); if rate cor insufficient after 15 minutes, a rep dose of 0.35 mg/kg over 2 minute given (average dose: 25 mg). Pat respond after 1 or 2 bolus doses of on a continuous infusion. Continuous infusion following bolu 10 mg/hour; infusion rate may be mg/hour increments according to response, up to a maximum of 15 Supraventricular tachvcardia (altern Note: For hemodynamically stable pat maneuvers and/or adenosine are uns Bolus dose: 0.25 mg/kg (actual over 2 minutes (average dose: 22 control is insufficient after 15 min bolus dose of 0.35 mg/kg over 2 be given (average dose: 25 mg). not terminate the arrhythm alternative therapy. | amically stable th preexcitation yay, as this can hinutes htrol is beat bolus es may be tients who can be started us(es): Initial: 5 to e increased in 5 ventricular o mg/hour. ative agent): tients if vagal successful. body weight) 20 mg); if rate putes, a repeat 2 minutes may If bolus(es) do | [average adult do continuous IV infuindividualized based ba | ilable: Infants, cents mg/kg over 5 im dose: 20 mg/dose ose]) followed by a usion. Dose should be |

| DIPHENHYDRAMINE QB , Lactation Y | Yes (Unsafe) Trade Name: Benadryl |
|--|--|
| Class / Mechanism of Action | |
| Histamine H ₁ Antagonist: Competes with histamine for H1-receptor sites within respiratory tract; Also produces anticholinergic and s | |
| Indications | |
| Labeled Indications: Anaphylaxis and allergy disorders Motion Sickness Antitussive | |
| Contraindications | |
| Hypersensitivity to diphenhydramine or any comp Acute Asthma Use on Neonates, premature infants, Nursing mod | |
| Adverse Reactions / Precautions | |
| Normally causes sedation but may cause parado May have increased sedative effects when used May cause hypotension (use with caution in patie Dry mouth | with other sedatives or alcohol. |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Anaphylaxis/Allergic Reactions and Motion Sickness: Oral: 25mg every 4-6 hours or 50mg every 6-8 hours- motion sickness IV Push: | Motion Sickness:Oral:0.5-1 mg/kg every 6 hoursIV/IM:• 1.25mg/kg every 6 hours |
| 25-50mg once, administered after epinephrine for anaphylaxis Note: Diphenhydramine is not the 1st line medication for anaphylaxis <u>Acute Hemolytic reaction</u> (Rapid onset of itching, chills, flushing, nausea/vomiting, coughing, wheezing, laryngeal edema, dyspnea, hypotension hemoglobinuria, rise in venous pressure, distended neck veins, crackles in lung bases): IV/IM: 25mg once | Anaphylaxis reaction: Adolescents: IV, IM, Oral: 25 - 50 mg/dose Allergic reaction: Children Ages 2 to <6 years: Oral: 6.25 mg every 4-8 hours Ages ≥6 to <12 years: Oral: 12.5 to 25 mg every 4-8 hours Adolescents: IV, IM, Oral: 25 to 50 mg/dose |

DOBUTAMINE

QB, Lactation? (Caution)

Trade Name: Dobutrex

Class / Mechanism of Action

Adrenergic Agonist

Positive Inotropic agent. Stimulates beta1 adrenergic receptors: Increases HR and contraction force while sparing beta2 and alpha receptors. Onset IV: 1-2 minutes

Indications

Labeled Indications: Short term management of cardiac decompensation.

Contraindications

- Hypersensitivity to dobutamine or sulfites (some contain sodium metabisulfite), or any component of the formulation.
- Hypertrophic cardiomyopathy with outflow tract obstruction

Adverse Reactions / Precautions

- Always attempt to correct Hypovolemia 1st when using vasopressors and/or inotropes.
 - May be combined with Dopamine or Norepinephrine for hypotension not responding to fluid administration.
 - No applicable use in hemorrhagic shock until fluid replacement therapy maximized!
- Increase in BP is common but does have a rare incidence of causing hypotension.
- Increases HR
- Hypotension and ventricular ectopy

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|---|--|
| Dose and Administration:ADUL1Cardiac Decompensation: IV: Dobutamine may be combined with dopamine or norepinephrine for hypotension not responsive to fluid therapy.•2-5 mcg/kg/min, start low and titrate to targeted MAP > 60 mmHg.•Usual dosage is 2-10mcg/kg/min- max dose 20mcg/kg/min•Preparation: Mix 250mg Dobutamine in 250mL D5W or NS for a concentration of 1000mcg/mLInfusion Rates for Dobutamine at 1000mcg/mLDesired Delivery Rate (mcg/kg/min)0.15 550.37.50.45100.612.50.75150.9201.2 | Cardiac Decompensation: Continuous IV or intraosseous infusion: • Initial: 0.5 to 1 mcg/kg/minute; titrate gradually every few minutes until desired response achieved; usual range: 2 to 20 mcg/kg/minute |

| DOPAMINE |
|----------|
|----------|

QC, Lactation? (Use Caution)

Trade Name: Intropin

Class / Mechanism of Action

Adrenergic Agonist; Vasopressor

Stimulates adrenergic and dopaminergic receptors. High doses stimulate dopaminergic and beta1 adrenergic receptors, producing cardiac stimulation and renal vasodilation. Very large doses stimulate alpha adrenergic receptors.

Indications

Labeled Indications:

Treatment of non-hemorrhagic shock (e.g., neurogenic, renal failure, cardiac decompensation) <u>persisting</u> <u>after adequate fluid volume replacement</u>

Unlabeled: Symptomatic bradycardia or heart block unresponsive to atropine

Contraindications

- Hypersensitivity to sulfites
- Ventricular Fibrillation
- Pheochromocytoma
- Uncorrected tachyarrhythmias

Adverse Reactions / Precautions

- No applicable use in hemorrhagic shock unless fluid replacement therapy maximized! Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shock.
- Tachycardia and/or Arrhythmias: May increase HR and worsen arrhythmias.
- Vesicant: Avoid extravasation, will cause tissue damage/necrosis
- Assure adequate circulatory volume to minimize need for vasoconstrictors. Monitor BP closely, <u>avoid</u> <u>hypertension</u> and adjust infusion rate as needed.

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | | | |
|---|---|--|--|--|
| Hemodynamic Support: IV(Use micro drip chamber only): | Hemodynamic Support: Ⅳ: | | | |
| • 5-20mcg/kg/min ; titrate to desired response. Infusion may be increased by 1- 4mcg/kg/minute at 10-to-30-minute intervals until optimal response is obtained. | • 2-20mcg/kg/min ; titrate to desired response. Infusion may be increased by 5- 10mcg/kg/minute until optimal response is obtained. | | | |
| Dopamine Dosage Efficacy: | | | | |
| Renal- 1-5 mcg/kg/min= Dopaminergic effects: increased urine output, increased renal blood flow | Note : Dopamine is a second line medication for hemodynamic support in Pediatric patients behind Epinephrine and Norepinephrine | | | |
| • Cardiac - 5-10 mcg/kg/min= Beta1 effects: Increased CO, HR, and contractility | | | | |
| • Vasoconstriction- >10 mcg/kg/min= Alpha1 effects: Increased BP | | | | |
| Note: Doses >20 mcg/kg/minute likely do not have a beneficial effect on blood pressure and may increase risk of tachyarrhythmias | | | | |
| Add additional vasopressor if Dopamine doses of 20 mcg/kg/min are inadequate. (<i>phenylephrine, norepinephrine, epinephrine</i> .) | | | | |

| | Dopamine | | | | | | Dopamine | | | | | | |
|--------|--|--------------|--------------|--------------|-------------|---------|--|--|------------|---------|---------------|------------|---------|
| | Dosing Range: 5-20mcg/kg/min (300-1200mcg/kg/hr) | | | | | | | Dosing Range: 5-20mcg/kg/min (300-1200mcg/kg/hr) | | | | | |
| | MIX 800 mg/500 mL | | | | | | | MIX 800 mg/500 mL | | | | | |
| | CONCENTRATION 1600 mcg/mL | | | | | | | CONCENTRATION 1600 mcg/mL | | | | | |
| Pt. | Dose | Rate | Micro | | Macro | | Pt. | Dose | Rate | Micro | | Macro | |
| Weight | | | (60 | 20 | 15 gtt/mL | 10 | Weight | | | (60 | 20 | 15 | 10 |
| | | | , | gtt/mL | | gtt/mL | | | | | gtt/mL | gtt/mL | gtt/mL |
| kg | mcg/kg/mi | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | kg | mcg/kg/mi | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min |
| | 5 | 9 | 9 | 3 | 2 | 2 | | 5 | 15 | 15 | 5 | 4 | 3 |
| 50 | 10 | 19 | 19 | 6 | 5 | 3 | 80 | 10 | 30 | 30 | 10 | 8 | 5 |
| 50 | 15 | 28 | 28 | 9 | 7 | 5 | 00 | 15 | 45 | 45 | 15 | 11 | 8 |
| | 20 | 38 | 38 | 13 | 10 | 6 | | 20 | 60 | 60 | 20 | 15 | 10 |
| | 5 | 10 | 10 | 3 | 3 | 2 | | 5 | 16 | 16 | 5 | 4 | 3 |
| 55 | 10 | 21 | 21 | 7 | 5 | 4 | 85 | 10 | 32 | 32 | 11 | 8 | 5 |
| 55 | 15 | 31 | 31 | 10 | 8 | 5 | 00 | 15 | 48 | 48 | 16 | 12 | 8 |
| | 20 | 41 | 41 | 14 | 10 | 7 | | 20 | 64 | 64 | 21 | 16 | 11 |
| | 5 | 11 | 11 | 4 | 3 | 2 | 2 4 6 8 | 5 | 17 | 17 | 6 | 4 | 3 |
| 60 | 10 | 23 | 23 | 8 | 6 | 4 | | 10 | 34 | 34 | 11 | 9 | 6 |
| 60 | 15 | 34 | 34 | 11 | 9 | 6 | | 15 | 51 | 51 | 17 | 13 | 9 |
| | 20 | 45 | 45 | 15 | 11 | 8 | | 20 | 68 | 68 | 23 | 17 | 11 |
| | 5 | 12 | 12 | 4 | 3 | 2 | | 5 | 18 | 18 | 6 | 5 | 3 |
| 05 | 10 | 24 | 24 | 8 | 6 | 4 | 95 | 10 | 36 | 36 | 12 | 9 | 6 |
| 65 | 15 | 37 | 37 | 12 | 9 | 6 | 95 | 15 | 53 | 53 | 18 | 13 | 9 |
| | 20 | 49 | 49 | 16 | 12 | 8 | | 20 | 71 | 71 | 24 | 18 | 12 |
| | 5 | 13 | 13 | 4 | 3 | 2 | | 5 | 19 | 19 | 6 | 5 | 3 |
| 70 | 10 | 26 | 26 | 9 | 7 | 4 | 100 | 10 | 38 | 38 | 13 | 10 | 6 |
| 70 | 15 | 39 | 39 | 13 | 10 | 7 | 100 | 15 | 56 | 56 | 19 | 14 | 9 |
| | 20 | 53 | 53 | 18 | 13 | 9 | | 20 | 75 | 75 | 25 | 19 | 13 |
| | 5 | 14 | 14 | 5 | 4 | 2 | | 5 | 20 | 20 | 7 | 5 | 3 |
| | 10 | 28 | 28 | 9 | 7 | 5 | 105 | 10 | 39 | 39 | 13 | 10 | 7 |
| 75 | 15 | 42 | 42 | 14 | 11 | 7 | 105 | 15 | 59 | 59 | 20 | 15 | 10 |
| | 20 | 56 | 56 | 19 | 14 | 9 | | 20 | 79 | 79 | 26 | 20 | 13 |
| | | | | | | | | | | | | | |
| | Mic | ro-Drip is s | set of choic | e for this i | nfusion | | | | | | ce for this i | | |
| Titr | rate to minin | num effect | ive dose. A | Allow 3-5 m | ninutes bet | ween | Titra | ate to minin | num effect | | Allow 3-5 m | inutes bet | ween |
| | dosing | | | | | | dosing | | | | | | |
| I | changes to assess hemodynamic effects. | | | | | | changes to assess hemodynamic effects. | | | | | | |

EPINEPHRINE

QC, Lactation? (Caution)

1:1000

Trade Name: EpiPen / EpiPen Jr

Class / Mechanism of Action

Alpha & Beta Agonist

Sympathomimetic, stimulates both alpha- and beta-adrenergic receptors, causing relaxation of the bronchial tree, induces systemic vasoconstriction and increases heart rate and contractility

Indications

- Allergic Reactions, Anaphylaxis •
- Asthma (Bronchoconstriction) •

Contraindications

- Not for IV use, must first dilute into 10mL NS syringe for Cardiac / IV use. ٠
- Hypersensitivity to sympathomimetic amines, glaucoma and non-anaphylactic shock ٠

Adverse Reactions / Precautions

- No applicable use in hemorrhagic shock unless fluid replacement therapy maximized! Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shock.
- Chest Pain, Tachycardia, Arrhythmias, Palpitations, Sudden death •
- Anxiety, Cerebral Hemorrhage, Headache
- Vesicant: Avoid extravasation, will cause tissue damage/necrosis .
- Use with caution in patients taking tricyclic antidepressants; effects of epinephrine may be increased .

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | | |
|---|---|--|--|
| Bronchodilator: SubQ/IM: • 0.3-0.5 mg every 5-10 min Nebulization: • Add 0.5 mL to nebulizer and dilute with 3 mL of NS; administer over 15 min Anaphylaxis / Hypersensitivity reaction: IM: • 0.3-0.5 mg in mid-outer thigh every 5-10 minutes until clinical improvement IV Infusion: • Initiate with an infusion at 5-15 mcg/minute (with crystalloid) (See infusion chart next page) Acute Hemolytic reaction IM: • 0.5mg IM in lateral thigh • Repeat every 5-15min for moderate bronchospasm or facial/laryngeal edema. | PEDIATRIC Always Reference BROSELOW Tape Bronchodilator: infants and Children: 0.01 mg/kg (0.01 mL/kg) (maximum single dose: 0.5 mg) every 3-5 min for 3 doses Nebulization: Children <4 years: Croup: 0.05-0.1 mL/kg (maximum dose: 0.5 mL); dilute in 2-3 mL of NS. May repeat dose every 20 min Children ≥4 years: Adult dosing Anaphylaxis / Hypersensitivity reaction: Infants and Children: SubQ/IM: 0.01 mg/kg (0.01 mL/kg of 1mg/mL solution) (maximum single dose: 0.3 mg) every 5-15 minutes Autoinjector, Children <15kg: 0.15 mg; if anaphylactic symptoms persist, dose may be repeated in 5-15 minutes using an additional EpiPen Jr Autoinjector, Children ≥15 kg: 0.3 mg; if anaphylactic symptoms persist, dose may be repeated in 5-15 minutes using an additional EpiPen | | |

| Epinephrine 1mg/1ml (1:1,000) | | | | | | | |
|-------------------------------|-------------|---------------|--------------|---------------|---------|--|--|
| Anaphylaxis | | | | | | | |
| Do | sing Ran | ge: 5-15m | cg/min (15 | 0-450mcg | /hr) | | |
| | | | g/500 mL | | | | |
| | COI | NCENTRA | TION 2 mc | :g/mL | | | |
| Dose | Rate | Micro | | Macro | | | |
| | | 60 gtt/mL | 20 | 15 gtt/mL | 10 | | |
| | | | gtt/mL | | gtt/mL | | |
| mcg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | |
| 5 | 150 | 150 | 50 | 38 | 25 | | |
| 6 | 180 | 180 | 60 | 45 | 30 | | |
| 7 | 210 | 210 | 70 | 53 | 35 | | |
| 8 | 240 | 240 | 80 | 60 | 40 | | |
| 9 | 270 | 270 | 90 | 68 | 45 | | |
| 10 | 300 | 300 | 100 | 75 | 50 | | |
| 11 | 330 | 330 | 110 | 83 | 55 | | |
| 12 | 360 | 360 | 120 | 90 | 60 | | |
| 13 | 390 | 390 | 130 | 98 | 65 | | |
| 14 | 420 | 420 | 140 | 105 | 70 | | |
| 15 | 450 | 450 | 150 | 113 | 75 | | |
| Mao | cro-Drip (1 | 0gtt/ml) is : | set of choic | ce for this i | nfusion | | |
| St | art at lowe | st dose an | d titrate to | desired eff | ect | | |

EPINEPHRINE

QC, Lactation? (Caution)

Trade Name: Adrenalin

1:10,000

Class / Mechanism of Action

Alpha & Beta Agonist

Sympathomimetic, stimulates both alpha- and beta-adrenergic receptors, causing relaxation of the bronchial tree, cardiac stimulation, and dilation of skeletal muscle blood vessels

Indications

- Cardiac Arrest (VF, pulseless VT, asystole, PEA) •
- Drip-Dose: Bradycardia (Symptomatic), Fluid Resistant Shock •
- Push-Dose: Refractory Anaphylaxis •

Contraindications

Uncontrolled hypertension is a relative contraindication, otherwise none •

Adverse Reactions / Precautions

.

- No applicable use in hemorrhagic shock unless fluid replacement therapy maximized! Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shock.
- Chest Pain, Tachycardia, Arrhythmias, Palpitations, Sudden death .
- Anxiety, Cerebral Hemorrhage, Headache
- Vesicant: Avoid extravasation, will cause tissue damage/necrosis
- Use with caution in patients taking tricyclic antidepressants; effects of epinephrine may be increased •

| Dose and Administration: ADUL | T PEDIATRIC Always Reference BROSELOW Tape |
|---|---|
| Asystole/pulseless arrest, pulseless VT/VF (ALS 2020): IV: 1mg/10mL (0.1mg/mL) pre-filled 10cc Syrin | and Symptomatic Bradycardia in Infants (ARC |
| • 1 mg (10cc of 0.1mg/mL) every 3-5 minute: ROSC, Follow each dose with 20mL flush | 0.01 mg/kg (0.1 mL/kg of 1mg/10mL [0.1 |
| Drip-Dose: Bradycardia (Symptomatic), Hypotension-Fluid Resistant Shock: IV Continuous Infusion: | mg/mL]) (maximum single dose: 1 mg) every 3- 5 minutes as needed or until ROSC |
| • Bradycardia: 2-10 mcg/minute titrate to desired effect (HR >60, MAP >65) | Severe Hypotension/shock and fluid resistant (unlabeled use): IV: Continuous Infusion |
| Hypotension/Shock: 2-10 mcg/min titrate clinical end point (BP, end organ perfusion) | |
| Push-Dose: Refractory Anaphylaxis: IV/IO: Utilize the 0.1 mg/mL solution further dilu in 10mL of NS. | ted |
| 0.05-0.1 mg, administered over 1-10 minute may repeat once after 3 minutes if patient remains unresponsive to initial dose. | es, |
| | |

| Epinephrine 1mg/10ml (1:10,000) | | | | | | | |
|---------------------------------|---|---------------|-----------------------|---------------|-----------|--|--|
| Pressor for Hypotension | | | | | | | |
| | Dosing Range: 2-20mcg/min (120- 600mcg/hr) | | | | | | |
| | | | g/500 mL TION 2 mc | a /m.l | | | |
| Deee | | | | Macro | | | |
| Dose | Rate | Micro (60 | 20 att/ml | | 10 att/ml | | |
| | | , | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL | | |
| mcg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | |
| 2 | 60 | 60 | 20 | 15 | 10 | | |
| 3 | 90 | 90 | 30 | 22.5 | 15 | | |
| 4 | 120 | 120 | 40 | 30 | 20 | | |
| 5 | 150 | 150 | 50 | 37.5 | 25 | | |
| 6 | 180 | 180 | 60 | 45 | 30 | | |
| 7 | 210 | 210 | 70 | 52.5 | 35 | | |
| 8 | 240 | 240 | 80 | 60 | 40 | | |
| 9 | 270 | 270 | 90 | 67.5 | 45 | | |
| 10 | 300 | 300 | 100 | 75 | 50 | | |
| 11 | 330 | 330 | 110 | 82.5 | 55 | | |
| 12 | 360 | 360 | 120 | 90 | 60 | | |
| 13 | 390 | 390 | 130 | 97.5 | 65 | | |
| 14 | 420 | 420 | 140 | 105 | 70 | | |
| 15 | 450 | 450 | 150 | 112.5 | 75 | | |
| 16 | 480 | 480 | 160 | 120 | 80 | | |
| 17 | 510 | 510 | 170 | 127.5 | 85 | | |
| 18 | 540 | 540 | 180 | 135 | 90 | | |
| 19 | 570 | 570 | 190 | 142.5 | 95 | | |
| 20 | 600 | 600 | 200 | 150 | 100 | | |
| Ма | cro-Drip (1 | 0gtt/ml) is : | set of choic | e for this in | nfusion | | |
| | Start at lo | | and titrate fect | to desired | | | |

ERTAPENEM

♀**C,** Lactation Yes

Trade Name: Invanz

Class / Mechanism of Action

Antibiotic (Carbapenem),

Bactericidal – broad spectrum, inhibits bacterial cell wall synthesis by binding to one or more of penicillinbinding proteins which inhibits cell wall biosynthesis, causing bacteria to eventually lyse.

Labeled Indications: Used for infection control prophylaxis for traumatic open injuries and surgical prophylaxis.

Contraindications

• Hypersensitivity to cefazolin, other cephalosporin antibiotics, other beta-lactams, or any component of the formulation

Adverse Reactions / Precautions

- Superinfection prolonged use may result in fungal or bacterial superinfection (including C.Difficile)
- Gastrointestinal: Diarrhea (Adults 9-12%).

| Dose and Administration: ADULT | PEDIATRIC |
|---|---|
| Infection Control: Give 1g in 250 mL NS IV over 5 min, provides 24 | Infection Control: Children <12 years & ≥12 years |
| hours of coverage. IV: Adults: • 1 g IV every 24 hrs. for 7-14 days. | IV: Pediatrics: • <12years old: 15 mg/kg IV every 12 hrs. ○ Max daily dose: 500mg/dose • ≥12 years old: 1000mg once daily ○ Max daily dose: 1000mg once daily. |

| ETOMIDATE QC, Lactation | ? (caution) Trade Name: Amidate |
|--|--|
| Class / Mechanism of Action | |
| General Anesthetic Ultra short acting non-barbiturate sedative/hypnotic u 60 seconds, Duration 5-10 minutes | sed for induction of anesthesia. Onset of action: 30- |
| Indications | |
| Labeled Indications:Rapid Sequence Induction | |
| Contraindications | |
| Hypersensitivity to etomidate or any component of Labor/Delivery Septic Schock | of the formulation |
| Adverse Reactions / Precautions | |
| NO Analgesic properties! Apnea/Respiratory Depression Hypo/hyperventilation Dysrhythmias Hypo/hypertension Nausea/Vomiting Transient involuntary skeletal muscle movement Pain at injection site Inhibits adrenal steroid production; may increase | mortality if repeat dosing is required |
| Dose and Administration: ADULT | PEDIATRIC |
| RSI: IV: | <u>RSI:</u> Ⅳ: |
| • 0.2-0.4 mg/kg over 30-60 seconds for induction of anesthesia. | 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. Max dose: 20 mg |
| Note: Limit to single dose for anesthesia/induction. Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortality due to adrenal suppression and inability to respond to stress. | Note: Limit to single dose for anesthesia/induction. Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortality due to adrenal suppression and inability to respond to stress. |

| FE | ENTANYL QC, Lactation recommende | | | | | | |
|---|---|--|--|--|--|--|--|
| Cla | ess / Mechanism of Action | | | | | | |
| Bin asc On | ioid Analgesic; General Anesthetic ds to opioid receptors within the CNS increasing p cending pain pathways (blocking painful stimulus); set: IV almost immediate, Duration: IV 0.5-1 hour lications | pain threshold and altering pain reception; inhibits ; produces CNS depression. | | | | | |
| | | | | | | | |
| • | b eled Indications: Pain relief | | | | | | |
| • | Adjunct to general or regional anesthesia | | | | | | |
| Со | ntraindications | | | | | | |
| • | Hypersensitivity to fentanyl or any component of | the formulation | | | | | |
| ٠ | MAOI taken in the past 14 days. | | | | | | |
| • | Hypotension | | | | | | |
| • | Нурохіа | | | | | | |
| • | Hypoventilation | | | | | | |
| Ad | verse Reactions / Precautions | general anesthesia, ensure Slow IV Push (3-5 min). | | | | | |
| • | Rapid infusion may result in chest wall rigidity, impaired ventilation, or respiratory distress/arrest <u>Always be prepared for use of paralytic and intubation (positive control of airway).</u> Head trauma: Use with extreme caution in head injury, or suspected increased ICP; exaggerated increase in ICP may occur if patient management is inadequate. CNS depression, Confusion Paradoxical excitation, delirium, drowsiness, apnea/dyspnea, bradycardia, dysrhythmias, hypotension, syncope, nausea/vomiting, abdominal pain, dehydration, | | | | | | |
| | bradycardia, dysrhythmias, hypotension, syncop | | | | | | |
| • | bradycardia, dysrhythmias, hypotension, syncop fatigue. | e, nausea/vomiting, abdominal pain, dehydration, | | | | | |
| • Do | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT | e, nausea/vomiting, abdominal pain, dehydration, PEDIATRIC | | | | | |
| • Do Pai | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: | e, nausea/vomiting, abdominal pain, dehydration, | | | | | |
| • Do Pai | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT | PEDIATRIC RSI: IV: | | | | | |
| • Do | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce | | | | | |
| • <u>Pai</u> IV: • No | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortali | | | | | |
| • Pai IV: • No incl | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction. Repeat dosing and continuous infusion | | | | | |
| • Pai IV: • No incl | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • • • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) (Combine with Midazolam for best effect) | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • • • • • • • • • • • • • • • • • • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) (Combine with Midazolam for best effect) 0.5-2mcg/kg IVP q 20-60min | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • • • • • • • • • • • • • • • • • • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) (Combine with Midazolam for best effect) 0.5-2mcg/kg IVP q 20-60min treatment for RSI: min prior to RSI in pt's with Head injuries, reased ICP, Cardiac Ischemia or Aortic | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • • • • • • • • • • • • • • • • • • • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) (Combine with Midazolam for best effect) 0.5-2mcg/kg IVP q 20-60min etreatment for RSI: min prior to RSI in pt's with Head injuries, reased ICP, Cardiac Ischemia or Aortic section (if situation allows): 3mcg/kg slow IV push | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |
| • Pai IV: • No incl • • • • • • • • • • • • • • • • • • • | bradycardia, dysrhythmias, hypotension, syncop fatigue. se and Administration: ADULT in Management: Slow (Unlabeled) 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM: 1mcg/kg mcg te: Patients with prior opioid exposure may have reased tolerance and require higher dosing dation during mechanical ventilation: IV: Initial Bolus: 1-2mcg/kg Continued Sedation: 0.5-1mcg/kg/hr. infusion (See Infusion chart next page) (Combine with Midazolam for best effect) 0.5-2mcg/kg IVP q 20-60min treatment for RSI: min prior to RSI in pt's with Head injuries, reased ICP, Cardiac Ischemia or Aortic section (if situation allows): | PEDIATRIC PEDIATRIC RSI: IV: • 0.2-0.4 mg/kg over 30-60 seconds will produce rapid sedation lasting 10-15 minutes. • Max dose: 20 mg Note: Limit to single dose for anesthesia/induction Repeat dosing and continuous infusion (maintenance dosing) may increase patient mortalid due to adrenal suppression and inability to responde | | | | | |

| FENTANYL (SUBLIMASE) | | | | | | | | |
|-------------------------|-----------------|----------------|---------------|----------------|-----------|--|--|--|
| | | Dosing Rang | _ | ncg/kg/hr | | | | |
| | MIX 1 mg/100 mL | | | | | | | |
| CONCENTRATION 10 mcg/mL | | | | | | | | |
| Dose | Rate | Micro | Macro | | | | | |
| | | (60 | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL | | | |
| | | gtt/mL) | | | | | | |
| mcg/hr | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | | |
| 25 | 3 | 3 | 1 | 1 | 0 | | | |
| 30 | 3 | 3 | 1 | 1 | 1 | | | |
| 35 | 4 | 4 | 1 | 1 | 1 | | | |
| 40 | 4 | 4 | 1 | 1 | 1 | | | |
| 45 | 5 | 5 | 2 | 1 | 1 | | | |
| 50 | 5 | 5 | 2 | 1 | 1 | | | |
| 55 | 6 | 6 | 2 | 1 | 1 | | | |
| 60 | 6 | 6 | 2 | 2 | 1 | | | |
| 65 | 7 | 7 | 2 | 2 | 1 | | | |
| 70 | 7 | 7 | 2 | 2 | 1 | | | |
| 75 | 8 | 8 | 3 | 2 | 1 | | | |
| 80 | 8 | 8 | 3 | 2 | 1 | | | |
| 85 | 9 | 9 | 3 | 2 | 1 | | | |
| 90 | 9 | 9 | 3 | 2 | 2 | | | |
| 95 | 10 | 10 | 3 | 2 | 2 | | | |
| 100 | 10 | 10 | 3 | 3 | 2 | | | |
| 105 | 11 | 11 | 4 | 3 | 2 | | | |
| 110 | 11 | 11 | 4 | 3 | 2 | | | |
| 115 | 12 | 12 | 4 | 3 | 2 | | | |
| 120 | 12 | 12 | 4 | 3 | 2 | | | |
| 125 | 13 | 13 | 4 | 3 | 2 | | | |
| 130 | 13 | 13 | 4 | 3 | 2 | | | |
| 135 | 14 | 14 | 5 | 3 | 2 | | | |
| 140 | 14 | 14 | 5 | 4 | 2 | | | |
| 145 | 15 | 15 | 5 | 4 | 2 | | | |
| 150 | 15 | 15 | 5 | 4 | 3 | | | |
| 155 | 16 | 16 | 5 | 4 | 3 | | | |
| 160 | 16 | 16 | 5 | 4 | 3 | | | |
| 165 | 17 | 17 | 6 | 4 | 3 | | | |
| 170 | 17 | 17 | 6 | 4 | 3 | | | |
| 175 | 18 | 18 | 6 | 4 | 3 | | | |
| 180 | 18 | 18 | 6 | 5 | 3 | | | |
| 185 | 19 | 19 | 6 | 5 | 3 | | | |
| 190 | 19 | 19 | 6 | 5 | 3 | | | |
| 195 | 20 | 20 | 7 | 5 | 3 | | | |
| 200 | 20 | 20 | 7 | 5 | 3 | | | |
| | Micro | -Drip is set o | f choice fo | r this infusio | on | | | |
| | S | ample patier | nt: 80kg pt a | at 0.5- | | | | |
| | | 1mcg | g/kg/hr = | | | | | |
| | 40 | mcg/hr-80mo | cg/hr dosin | g range | | | | |

| FUROSEMIDE QC, Lact | ation Yes (Caution) | Trade Name: Lasix | |
|--|--|--|--|
| Class / Mechanism of Action | | | |
| Antihypertensive; Loop Diuretic Inhibits reabsorption of sodium and chloride in chloride, magnesium, and calcium within urin Symptomatic improvement of acute pulmonar | e. When given IV it also causes ra | apid venous dilation. | |
| Indications | | | |
| Labeled Indications: Management of edema Management of edema associated with he edema. Hypertension (alone or in combination with | eart failure and hepatic or renal dis | | |
| Contraindications | | | |
| Hypersensitivity to furosemide or any com Anuria (No pre-hospital utility in hypovo | • | | |
| Adverse Reactions / Precautions | | | |
| Can cause profound diversis with resulting May cause: Hypovolemia, Hypote May potentiate effect of additional antihyperity and the second se | nsion, hyponatremia, hypokalemia | Monitor closely! | |
| Dose and Administration: ADU | LT PEDIATRIC Alwa | ys Reference BROSELOW Tape | |
| Acute pulmonary edema: IV 40 mg over 1-2 minutes. If response not adequate within 1 hour, may increase dose 80 mg. Edema. heart failure: IV, IM: Initial: 20-40 mg/dose; if response is not adequate, may repeat the same dose or increase dose in increments of 20 mg/dos administer 1-2 hours after previous dose (maximum dose: 200 mg/dose). Continuous IV Infusion: Initial: IV bolus dose 20-40 mg over 1-2 minutes, followed by continuous IV infusio doses of 10-40 mg/hour. If urine output is mL/kg/hour, double as necessary to a maximum of 80-160 mg/hour. | adequate, may increa 1 mg/kg/dose and ad hours after previous of response is achieved maintenance dose at hours; maximum dos e and | ; if response not ase dose in increments of minister not sooner than 2 dose, until a satisfactory l; may administer i intervals of every 6-12 | |

| GLUCAGON QB Lactation? | (Caution) |
|---|---|
| Class / Mechanism of Action | |
| Antidote, Hypoglycemia Antidote, Diagnostic agen Hyperglycemic agent, pancreatic hormone, insulin an increased production of cyclic AMP, promoting hepat | tagonist; Raises blood glucose levels by stimulating |
| Indications | |
| Labeled Indications: Management of hypoglycemia Unlabeled: Beta-blocker or calcium channel blocker induced unresponsive to standard measures. Hypoglycemia secondary to insulin or sulfonylureation. | myocardial depression (with or without hypotension) |
| Contraindications | |
| Hypersensitivity to glucagon or any component of Insulinoma / Pheochromocytoma Hyperglycemia Adverse Reactions / Precautions | the formulation |
| • Should NOT be used as 1 st line medication for hyp | ooglycemia or Altered mental status |
| • Hypoglycemia patients should receive | dextrose . If IV access cannot be established or if be used as alternate until dextrose can be given. |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Hypoglycemia: IV, IM, IN, SubQ: 1 mg; may repeat in 20 minutes as needed. Beta-blocker / Calcium channel blocker overdose (myocardial depression) unresponsive to standard measures (unlabeled use): IV: • 5mg bolus slow IV push | *Hypoglycemia: IV, IM, IN, SubQ: Children <20 kg: 0.5 mg repeated in 20min prn. Children ≥20 kg: Adult dosing. Note: IV dextrose should be given ASAP; if patient fails to respond to glucagon, IV dextrose must be given *Only use if hyperinsulinemia thought to be cause of hypoglycemia (rare in kids). If hypoglycemic without glycogen stores, Glucagon will be ineffective. |
| | Beta-blocker / Calcium channel blocker overdose (myocardial depression) unresponsive to standard measures (unlabeled use): IV: Children <25kg: 0.5mg slow IV push q5min prn Children >25 kg: 1mg slow IV push q5min prn |

| HEPARIN | ♀ C, Lactation No | D Trade Name: |
|---|---|--|
| Class / Mechanism of Action | , | |
| Anticoagulant | coagulation factors | (IX, X, XI, XII, and plasmin) and prevents |
| Indications | | |
| · · · · · · · · · · · · · · · · · · · | | isorders. thrombolysis; unstable angina/non-STEMI |
| Contraindications | | |
| Hypersensitivity to heparin or a Active Bleeding (Trauma Patie) | • • | ne formulation |
| Active Bleeding (Trauma Patie Adverse Reactions / Precautions | , | |
| Continuously monitor for bleed Urticarial reactions and anaphy | ing: Stop immediate | ely if any bleeding occurs. |
| Dose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Acute coronary syndromes: STE Angina as an adjunct to fibrinolysis alteplase: IV: Initial bolus of 60 units/kg (MA units) Maintenance: 12 units/kg/hd 1000 units/hour) as continue Treatment of venous thromboen IV: (unlabeled dosing) <u>DVT/PE:</u> 80 units/kg or 1mg/l alternatively 5000 units) IV pus continuous infusion of 18 units. <u>COVID VTE Prophylaxis:</u> 7,500 8 hours Note: <u>Heparin is ONLY for use onl</u> written direction of referring provide consultation with medical director. | s (full dose AX: 4000 our (MAX: ous infusion. hbolism: kg (or sh followed by /kg/hour. 0 iu SQ every y under | Treatment of venous thromboembolism: IV: (unlabeled dosing) >1 year DVT/PE: 75 units/kg IV push followed by continuous infusion of 20 units/kg/hour |

| Η | ETASTARCH | C , Lactation Y | es (caution) | Trade Name: Hextend |
|-----|---|------------------------|-------------------------------|--------------------------|
| Cla | ass / Mechanism of Action | , | | |
| | asma Volume Expander, Colloi | | | |
| Co | lloidal starch producing plasma | olume expansion | . Onset of Action: approximat | ely 30 minutes |
| | dications | | | |
| | beled Indications: Volume expa | inder used in trea | tment of hypovolemic / hemo | rrhagic shock |
| Co | ontraindications | | | |
| ٠ | Hypersensitivity to hydroxyethy | • | - | |
| ٠ | Renal failure with oliguria and a | · · | , | |
| • | Fluid overload conditions, (puln Pre-existing bleeding or coagul | | 8 | Line coution in blooding |
| • | disorders; may increase risk of | | g, von villebrand's disease). | Ose caution in pleeding |
| Ad | Iverse Reactions / Precautions | y | | |
| ٠ | Anaphylactoid reactions (allerg | es to corn) | | |
| Do | ose and Administration: | ADULT | PEDIATRIC Always F | Reference BROSELOW Tape |
| Pla | asma volume expansion: | | | |
| IV | - | | | |
| • | 250-500ml Bolus. May repeat F | PRN (up to | | |
| | 1500 mL/day). Titrate to individ | | | |
| | hemodynamic needs (Sys BP > | ·90). | | |
| No | otes: | | | |
| • | May be administered via infusion pressure infusion. | n pump or | | |
| • | Do not administer with blood th same line / tubing | rough the | | |
| • | Change tubing or flush extensive before administering blood throws ame line. | | | |
| • | Use only for burns as an adjund preferred until hour 8-12; may u difficult resuscitation. | | | |

| HYDROMORPHONE 9 | C, Lactation Yes (not recommended) | Trade Name: Dilaudio |
|--|---|---|
| Class / Mechanism of Action | , | |
| Opioid Analgesic Binds to opioid receptors within the inhibits ascending pain pathways (blo Onset: IV 10-20 minutes. Duration 2-4 | ocking painful stimulus); produc | |
| Indications | | |
| Labeled Indications: Moderate to se | evere pain. | |
| Contraindications | | |
| Hypersensitivity to hydromorphon Severe respiratory depression (in Acute or severe asthma Paralytic ileus MAOI use in past 14 days GI obstruction Hypotension Hypoxia Head injury Hypoventilation Adverse Reactions / Precautions Always be prepared for use of pa Head trauma: Use with extreme exaggerated increase in ICP ma May cause Hypotension, Use with May cause life-threatening Repar CNS depression: Impairs physica Dizziness Headache | absence of resuscitative equip ralytic and intubation (maintain caution in head injury, or su ay occur. h caution in hypovolemic patier atory depression | positive control of airway). |
| Syncope Dose and Administration: | ADULT PED | NATRIC Always Reference BROSELOW Tape |
| Acute pain (moderate-to-severe): | | noderate-to-severe): |
| IV: (Slow) | IV: (Slow) | |
| 0.5mg (range 0.25-2mg) IV/IO q prn Critically ill require lower dose, tolerant may require higher dos Continuous infusion: Usual dosag 0.5- 3 mg/hour (See infusion ch page) | 1-6hr as , opioid se ge range: | 0.015mg/kg IV q 4-6 PRN nts >50kg: Refer to adult |

| | HYDROMORPHONE (DILAUDID) | | | | | | |
|---------|---|----------------------------|--------------|-----------|--------------|--|--|
| D | Dosing Range: 0.5-3mg/hr (8.3-50mcg/min) | | | | | | |
| | MIX 2 mg/100 mL | | | | | | |
| | CON | ICENTRAT | ION 20 m | - | | | |
| Dose | Rate | Micro | | Macro | | | |
| | | 60 gtt/mL | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL | | |
| mg/hr | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | |
| 0.5 | 25 | 25 | 8 | 6 | 4 | | |
| 1 | 50 | 50 | 17 | 13 | 8 | | |
| 1.5 | 75 | 75 | 25 | 19 | 13 | | |
| 2 | 100 | 100 | 33 | 25 | 17 | | |
| 2.5 | 125 | 125 | 42 | 31 | 21 | | |
| 3 | 150 | 150 | 50 | 38 | 25 | | |
| Macro | Macro-Drip (20gtt/ml) or Micro-Drip is set of choice for this infusion | | | | | | |
| Start a | | ose and ind appropriate | | , , | /hr PRN | | |

| HYDROXOCOBALAMIN QC, Lactation | on? (Caution) Trade Name: Cyanokit® |
|--|--|
| Class / Mechanism of Action | |
| Antidote; Vitamin Precursor to Vitamin B ₁₂ (cyanocobalamin). Binds cy excreted within urine | anide ion to form nontoxic cyanocobalamin which is |
| Indications | |
| Labeled Indications: | |
| IM: Treatment of pernicious anemia and B12 defice | |
| IV: (Cyanokit®) Treatment of known or suspected Contraindications | cyanide poisoning |
| | or known ovenide neisening |
| No contraindications when treating for suspected Adverse Reactions / Precautions | |
| May cause transient hypertension (≥180mmHg s) | vetolic >110mmHa diastolic) |
| Will cause persistent red colored urine and skin, r | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| IV/IO: (Note: If cyanide poisoning is suspected, antidotal therapy must be given immediately) Initial: 5 grams as single infusion given over 15 min Repeat a second 5-gram dose based on severity and clinical response. Maximum cumulative dose: 10 grams Smoke Inhalation / Fire victims: (Closed space exposure with evidence of airway injury: soot in mouth / nose / sputum) May present with both cyanide and carbon monoxide poisoning. Hydroxocobalamin is the agent of choice for treating cyanide toxicity in this setting. | IV/IO: (Unlabeled Use) Initial: 70mg/kg (max 5 grams) as single infusion given over 15 min Repeat a second dose of 35mg/kg based on severity and clinical response. Smoke Inhalation / Fire victims: (Closed space exposure with evidence of airway injury: soot in mouth / nose / sputum) May present with both cyanide and carbon monoxide poisoning. Hydroxocobalamin is the agent of choice for treating cyanide toxicity in this setting. |
| Preparation: Cyanokit®: Reconstitute each vial with 200 mL of NS (LR and D5W also OK). Do not shake vial (gently mix) Do not use if solution is not dark red | |

| | \mathcal{C} , Lactation Yes | (Risk not ruled out) | Trade Name: Ketala |
|--|---|---|---|
| Class / Mechanism of Actio | n | | |
| General Anesthetic Dissociative anesthetic; produ Onset of action IV: 30-60 sec | | | |
| Indications | | | |
| Labeled Indications: Inducti | | general anesthesia | |
| Unlabeled: Analgesia and se Contraindications | alion | | |
| Hypersensitivity to ketam Conditions that cannot to hemorrhage, hypertensio Children <3 mo. age | lerate sustained increas | es in blood pressure (no | |
| Adverse Reactions / Precau | utions | | |
| mg/kg) may cause hypUse with caution in patierPreferred general anesth | otension and respirate nts with cardiovascular o etic / sedative (pre-hosp ng/kg IV (and equivalen | ory depression disease. Continuously mo bital) for head injury patie | |
| Dose and Administration: | ADULT | PEDIATRIC Alway | vs Reference BROSELOW Tape |
| LOW DOSE: Analgesia: IV/IO Push (over 1 min) • 0.1-0.3 mg/kg, repeat q 7 IM/IN • 0.5 - 1.0 mg/kg, repeat q HIGH DOSE: <u>RSI / Induction of anesthes</u> <u>Patients:</u> IV Push • 1-2 mg/kg IM • 4-5 mg/kg <u>Maintenance of anesthesia</u> IV: • 0.5-2 mg/kg dose every IV Continuous Infusion • 0.5-2mg/kg bolus then 7 | 10-30 prn ia; Combative 10-20 minutes | IV: 1-2 mg/kg (3-5mg/sedation) Maintenance of anestl IV: 1⁄₂ to Full induction IV Continuous Infusion: 0.5-1 mg/kg/hr. Time PRN to achieve application | peat q 10-30 prn ia (unlabeled dosing): kg for procedural hesia: n dose every 20-30 minutes trate levels by 0.25mg/kg/h propriate sedation. ssociative doses to prever |

2-5mg IV x1 for adults and 0.05 mg/kg for children not hypotensive or in danger of being

| | Dosing | | 3mg/kg/hr (1 | | (g/min) | |
|------------|-------------------|--------------|--------------------------|---------------------|---------------|--------------|
| | | | 00 mg/500 m ENTRATION | | | |
| | | | mg/mL | | | |
| Pt. Weight | Dose | Rate | Micro | 00 11/ 1 | Macro | 40 11/ |
| lea. | mcg/kg/min | mL/hr | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/ml |
| kg | nicg/kg/min 15 | 45 | gtt/min 45 | gtt/min 15 | gtt/min 11 | gtt/min 8 |
| | 20 | 60 | 60 | 20 | 15 | 10 |
| | 25 | 75 | 75 | 25 | 19 | 13 |
| | 30 | 90 | 90 | 30 | 23 | 15 |
| 50 | 35 | 105 | 105 | 35 | 26 | 18 |
| 50 | 40 | 120 | 120 | 40 | 30 | 20 |
| | 45 50 | 135 150 | 135 150 | 45 50 | 34 38 | 23 25 |
| | 55 | 165 | 165 | 55 | 41 | 23 |
| | 60 | 180 | 180 | 60 | 45 | 30 |
| | 15 | 50 | 50 | 17 | 13 | 8 |
| | 20 | 66 | 66 | 22 | 17 | 11 |
| | 25 | 83 | 83 | 28 | 21 | 14 |
| | 30 | 99 | 99 | 33 | 25 | 17 |
| 55 | 35 40 | 116 132 | 116 132 | 39 44 | 29 33 | 19 22 |
| | 40 45 | 132 | 132 149 | 44 50 | 33 | 22 |
| | 43 50 | 149 | 149 | 55 | 41 | 23 |
| | 55 | 182 | 182 | 61 | 46 | 30 |
| | 60 | 198 | 198 | 66 | 50 | 33 |
| | 15 | 54 | 54 | 18 | 14 | 9 |
| | 20 | 72 | 72 | 24 | 18 | 12 |
| | 25 | 90 | 90 | 30 | 23 | 15 |
| | 30 | 108 | 108 | 36 | 27 | 18 |
| 60 | 35 40 | 126 140 | 126 140 | 42 47 | 32 35 | 21 23 |
| | 40 45 | 140 | 140 | 47 54 | 41 | 23 |
| | 50 | 180 | 180 | 60 | 45 | 30 |
| | 55 | 198 | 198 | 66 | 50 | 33 |
| | 60 | 216 | 216 | 72 | 54 | 36 |
| | 15 | 60 | 60 | 20 | 15 | 10 |
| | 20 | 78 | 78 | 26 | 20 | 13 |
| | 25 | 98 | 98 | 33 | 25 | 16 |
| | 30 35 | 117 137 | 117 137 | 39 46 | 29 34 | 20 23 |
| 65 | 40 | 156 | 156 | 52 | 39 | 25 |
| | 45 | 176 | 176 | 59 | 44 | 29 |
| | 50 | 195 | 195 | 65 | 49 | 33 |
| | 55 | 215 | 215 | 72 | 54 | 36 |
| | 60 | 234 | 234 | 78 | 59 | 39 |
| | 15 | 63 | 63 | 21 | 16 | 11 |
| | 20 | 84 | 84 | 28 | 21 | 14 18 |
| | 25 30 | 105 126 | 105 126 | 35 42 | 26 32 | 21 |
| | 35 | 147 | 147 | 49 | 37 | 25 |
| 70 | 40 | 168 | 168 | 56 | 42 | 28 |
| | 45 | 189 | 189 | 63 | 47 | 32 |
| | 50 | 210 | 210 | 70 | 53 | 35 |
| | 55 | 231 | 231 | 77 | 58 | 39 |
| | 60 | 252 | 252 | 84 | 63 | 42 |
| | 15 20 | 68 90 | 68 90 | 23 30 | 17 23 | 11 15 |
| | 20 | 113 | 90 113 | 30 | 23 | 15 |
| | 30 | 135 | 135 | 45 | 34 | 23 |
| | 35 | 158 | 158 | 53 | 40 | 26 |
| 75 | 40 | 180 | 180 | 60 | 45 | 30 |
| | 45 | 203 | 203 | 68 | 51 | 34 |
| | 50 | 225 | 225 | 75 | 56 | 38 |
| | 55 | 248 | 248 | 83 | 62 | 41 |
| | 60 Maara | 270 | 270 | 90 or this infus | 68 | 45 |
| | wacro | is se חויט-י | et of choice fo | unis intus | | |

| | KETAMINE (KETALAR) Dosing Range: 1-3mg/kg/hr (17-50mcg/kg/min) | | | | | |
|-------------|---|--------------|--------------------|---------------|-----------|----------|
| | MIX 500 mg/500 mL | | | | | |
| | | | ENTRATION mg/mL | | | |
| Pt. Weight | Dose | Rate | Micro | | Macro | |
| i t. weight | Dose | TALE | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/m |
| kg | mcg/kg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min |
| 0 | 15 | 72 | 72 | 24 | 18 | 12 |
| | 20 | 96 | 96 | 32 | 24 | 16 |
| | 25 | 120 | 120 | 40 | 30 | 20 |
| | 30 | 144 | 144 | 48 | 36 | 24 |
| 80 | 35 | 168 | 168 | 56 | 42 | 28 |
| | 40 | 192 | 192 | 64 | 48 | 32 |
| | 45 50 | 216 240 | 216 240 | 72 80 | 54 60 | 36 40 |
| | 55 | 240 | 240 | 88 | 66 | 40 |
| | 60 | 288 | 288 | 96 | 72 | 48 |
| | 15 | 77 | 77 | 26 | 19 | 13 |
| | 20 | 102 | 102 | 34 | 26 | 17 |
| | 25 | 128 | 128 | 43 | 32 | 21 |
| | 30 | 153 | 153 | 51 | 38 | 26 |
| 05 | 35 | 179 | 179 | 60 | 45 | 30 |
| 85 | 40 | 204 | 204 | 68 | 51 | 34 |
| | 45 | 230 | 230 | 77 | 58 | 38 |
| | 50 | 255 | 255 | 85 | 64 | 43 |
| | 55 | 281 | 281 | 94 | 70 | 47 |
| | 60 | 306 | 306 | 102 | 77 | 51 |
| | 15 20 | 81 108 | 81 108 | 27 36 | 20 27 | 14 18 |
| | 20 25 | 106 | 106 | 45 | 34 | 23 |
| | 30 | 162 | 162 | 43 54 | 41 | 23 |
| | 35 | 189 | 189 | 63 | 47 | 32 |
| 90 | 40 | 216 | 216 | 72 | 54 | 36 |
| | 45 | 243 | 243 | 81 | 61 | 41 |
| | 50 | 270 | 270 | 90 | 68 | 45 |
| | 55 | 297 | 297 | 99 | 74 | 50 |
| | 60 | 324 | 324 | 108 | 81 | 54 |
| | 15 | 90 | 90 | 30 | 23 | 15 |
| | 20 | 114 | 114 | 38 | 29 | 19 |
| | 25 | 143 | 143 | 48 | 36 | 24 |
| | 30 | 171 | 171 | 57 | 43 | 29 |
| 95 | 35 | 200 | 200 | 67 | 50 | 33 |
| | 40 45 | 228 257 | 228 257 | 76 86 | 57 64 | 38 43 |
| | 43 50 | 285 | 285 | 95 | 71 | 43 |
| | 55 | 314 | 314 | 105 | 79 | 52 |
| | 60 | 342 | 342 | 114 | 86 | 57 |
| | 15 | 90 | 90 | 30 | 23 | 15 |
| | 20 | 120 | 120 | 40 | 30 | 20 |
| | 25 | 150 | 150 | 50 | 38 | 25 |
| | 30 | 180 | 180 | 60 | 45 | 30 |
| 100 | 35 | 210 | 210 | 70 | 53 | 35 |
| 100 | 40 | 240 | 240 | 80 | 60 | 40 |
| | 45 | 270 | 270 | 90 | 68 | 45 |
| | 50 55 | 300 330 | 300 | 100 | 75 | 50 55 |
| | 55 | | 330 360 | 110 120 | 83 | 55 60 |
| | 60 15 | 360 95 | 360 95 | 120 32 | 90 24 | 60 16 |
| | 20 | 95 126 | 126 | 42 | 32 | 21 |
| | 25 | 158 | 120 | 53 | 40 | 26 |
| | 30 | 189 | 189 | 63 | 47 | 32 |
| | 35 | 221 | 221 | 74 | 55 | 37 |
| 105 | 40 | 252 | 252 | 84 | 63 | 42 |
| | 45 | 284 | 284 | 95 | 71 | 47 |
| | 50 | 315 | 315 | 105 | 79 | 53 |
| | 55 | 347 | 347 | 116 | 87 | 58 |
| | 60 | 378 | 378 | 126 | 95 | 63 |
| | | o Drin io oc | et of choice fo | or this infus | ion | |

| K | ETOROLAC QC, Lactation Yes (Caution)Trade Name | | | |
|--------------------------------|---|--|--|--------------------------|
| Cla | ss / Mechanism of Actio | n | | |
| Inh | nsteroidal Anti-inflamma ibits cyclooxygenase (CO. ovides antipyretic, analges | X 1 & 2) enzymes, which | n decreases production of pros y action. | staglandin precursors. |
| Ind | lications | | | |
| La | beled Indications: Short t | erm management of mo | derate to severe acute pain as | s an opioid alternative. |
| Со | ntraindications | | | |
| • | High risk of bleeding, rec disease. | ent history of GI bleeding | Ds, or any component of the fo g or perforation, known history attlefield trauma patient! | |
| • | Suspected cerebrovascu | | | |
| • | Risk of renal failure second | - | n | |
| • | Concurrent use with othe | | | |
| Ad | verse Reactions / Precau | itions | | |
| • | stroke | ritation, inflammation, uld hospasm in patients with | | - |
| Do | se and Administration: | ADULT | PEDIATRIC Always Refe | rence BROSELOW Tape |
| <i>Pa</i> : IM: ● IV: | 15-30 mg every 6 hours 120 mg) 15 mg every 6 hours (ma 120 mg) <i>ults ≥65 years and/or ad</i> 15-30 mg every 6 hours | (maximum daily dose: aximum daily dose: ults <u><</u>50 kg | Pain management (acute;Children 2-16 y/o:• Moderate discomfort: 1 r• Severe discomfort: 0.5 r• Febrile seizure: 1 mg/kgAdolescents >17:• Refer to adult dose | mg/kg IM ng/kg IV |
| IV: | 60 mg) 15 mg every 6 hours (ma mg) | iximum daily dose: 60 | | |

| LABETALOL QC, Lactation Yes (Caution | on) Trade Name: Trandat e |
|--|---|
| Class / Mechanism of Action | |
| Beta Blocker with alpha blocking activity Blocks alpha and beta1/beta2 adrenergic receptor sites | s Onset IV: 2-5 minutes |
| Indications | |
| Labeled Indications: Treatment of hypertension. IV: Treatment of severe hypertension and hyperter Unlabeled: Pre-eclampsia and severe hypertension in pregnar Pediatric hypertension | nsive emergencies ncy, hypertension during acute ischemic stroke, and |
| Contraindications | |
| Hypersensitivity to labetalol or any component of th Bradycardia <60bpm, heart block >1st degree Uncompensated heart failure, Cardiogenic shock Asthma | |
| Adverse Reactions / Precautions Symptomatic hypotension with or without syncope, | |
| Use with extreme caution in patients with compens Patient with bronchospastic diseases (reactive airw | • |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELO |
| Таре | |
| Acute Hypertension (hypertensive emergency/urgency: | *Hypertension emergencies: |
| Hypertensive Crisis (Sys: >185/Dia: >110) Intermittent IV: | IV Continuous Infusion 0.25-3 mg/kg/hour; administration requires the use of an infusion pump. Intermittent bolus doses of 0.2-1 mg/kg/dose have been reported. Maximum dose 40mg/dose. |
| • 10-20 mg IV over 1-2 minutes. May repeat one time. | *Not 1 st Line medication for children |
| Continuous Infusion : Initial loading dose: 10-20 mg over 2 minutes, followed by 0.5-2.0 mg/min | |
| Note: Goal to lower MAP by no more than 25% within minutes to one hour | |
| Pearls: For inter-facility transports with confirmed Ischemic CVA | A, Intraparenchymal Hemorrhagic CVA, or Spontaneou |

- Ischemic CVA Lytic eligible: SBP <185 and DBP <110 Intraparenchymal Hemorrhagic CVA: SBP <180 Non-traumatic SAH: SBP <160 •
- •
- •

Ringer's Lactate (Lactated Ringers)

Class: Isotonic crystalloid solution.

Mechanism of Action: Replaces water and electrolytes.

Indications: Hypovolemic shock; keep open IV. Standard burn resuscitation

Contraindications: Should not be used in the same line with blood components. Use with caution for intravascular volume replacement for hemorrhagic shock due to hemodilution and exacerbation of coagulopathy. Use with caution in patients with known congestive heart failure and kidney disease. Can cause lactic acidosis. Should not use in head injury patients.

Adverse Reactions: Rare

Drug Interactions: Few in the pre-hospital emergency setting.

How Supplied: 250mL, 500mL, and 1,000mL bags. IV infusion.

Dosage and Administration: Hypovolemic shock; titrate according to the patient's physiologic

response. Burn resuscitation use Rule of 10's to calculate infusion rate.

(See appropriate Guidelines)

Dextrose 5% in Water (D5W)

Class: Hypotonic dextrose-containing solution.

Mechanism of Action: D5W provides nutrients in the form of dextrose as well as free water.

Indications: IV diluent for certain emergency drugs; for dilution of concentrated drugs for intravenous infusion.

Contraindications: Not for use as fluid replacement for hypovolemic states.

Adverse Reactions: Rare

Drug Interactions: Phenytoin (Dilantin)

How Supplied: Supplied in 50mL, 100mL, 150mL, 250mL, 500mL, and 1,000mL bags.

Dosage and Administration: Normally administered through a mini-drip (60 gtt/mL) set at a rate of "to keep open" (TKO).

LEVETIRACETAM

^QC, Lactation Yes (Caution)

Trade Name: KEPPRA

Class / Mechanism of Action

Anticonvulsant

Causes modulation of synaptic neurotransmitter release through binding to the synaptic vesicle protein SV2A in the brain.

Indications

Labeled Indications

• Treatment of focal (partial) onset seizures

Unlabeled:

Traumatic brain injury, severe acute (short-term seizure prophylaxis); Status epilepticus; Craniotomy, • seizure prophylaxis; Subarachnoid hemorrhage (short-term seizure prophylaxis)

Contraindications

- Hypersensitivity to any component of the formulation •
- **Adverse Reactions / Precautions**

May cause CNS depression •

- Dermatologic reactions, possibly severe (TEN, SJS, etc.) •
- Hypertension has been reported in children <4 years
- Hematologic effects: Decreases in red blood cell counts, hemoglobin, hematocrit, white blood cell counts, and neutrophils and increases in eosinophils have been observed

| Dose and Administration: | ADULT |
|--------------------------|-------|
| | |

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|---|---|
| <u>Traumatic brain injury (severe acute) (short-term</u> seizure prophylaxis): | Status epilepticus, refractory: (Limited data available) |
| Loading dose: 1500 mg over 15 minutes Maintenance dose: 1000 mg over 15 minutes every 12 hours | Infants, Children, and Adolescents: IV: 60 mg/kg over 15 minutes as a single dose; 4,500mg max dose. |
| Status epilepticus: • IV: 2000 mg over 15 minutes | Traumatic brain injury, Seizure prophylaxis: IV: 20-55 mg/kg/day in divided doses twice daily. |

| LIDOCAINE QB, Lactation Yes (Caution) Trade Name: Xylocaine (Cardiac) Class / Mechanism of Action Antiarrhythmic Suppresses automaticity of cardiac conduction tissue. Indications Labeled Indications: Acute treatment of ventricular arrhythmias from myocardial infarction (alternate to amiodarone when amiodarone not available) Unlabeled: (ACLS, 2015) Hernodynamically stable monomorphic VT and polymorphic VT Pulseless VT /VF (unresponsive to defibrillation, CPR, and vasopressor administration) Monomorphic VT secondary to drug, when amiodarone is not available Contraindications Outraindications Intracessed ventricular rate may be seen when given to a patient in A Fib. Prophylactic use in AMI Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions Continuous EKG monitoring is necessary Increased ventricular rate may be seen when given to a patient in A Fib. At high doeser, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Doca mad Administration 1. Does and Administration: ADULT PEIDATRIC Aweys Retence BROSELOW Tage Cardiac A | IABLE OI | CONTENTS |
|---|--|--|
| Antiarrhythmic Suppresses automaticity of cardiac conduction tissue. Indications Labeled Indications: Acute treatment of ventricular arrhythmias from myocardial infarction (alternate to amiodarone when amiodarone not available) Unlabeled: (ACLS, 2015) • Hemodynamically stable monomorphic VT and polymorphic VT • Pulseless VT /VF (unresponsive to defibrillation, CPR, and vasopressor administration) • Monomorphic VT secondary to drug, when amiodarone is not available Contraindications • Hypersensitivity to lidocaine or any component of the formulation • Prophylactic use in AMI Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block • Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions • Continuous EKG monitoring is necessary • Increased ventricular rate may be seen when given to a patient in A Fib. • At high doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. o D/C immediately if toxicity develops • The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT PEIATRIC Always Reference BROSELOW Tape • For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes o Maximum of 30 doses or total of 3mg/k | LIDOCAINE QB, Lactation Yes (C | Caution) Trade Name: Xylocaine (Cardiac) |
| Suppresses automaticity of cardiac conduction tissue. Indications Labeled Indications: Acute treatment of ventricular arrhythmias from myocardial infarction (alternate to amiodarone when amiodarone to available) Unlabeled: (ACLS, 2015) • Hemodynamically stable monomorphic VT and polymorphic VT • Pulseless VT / VF (unresponsive to defibrillation, CPR, and vasopressor administration) • Monomorphic VT secondary to drug, when amiodarone is not available Contraindications • Hypersensitivity to lidocaine or any component of the formulation • Prophylactic use in AMI • Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block • Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions • Continuous EKG monitoring is necessary • Increased ventricular rate may be seen when given to a patient in A Fib. • A high doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. • D/C immediately if toxicity develops • The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT • Arest from VF/VT, (if Amiodarone is not available): Stable VT, wide complex tachycardia, significant ectopy; • No: • Initial dose: 1 to 1.5mg/kg • To Isomg/kg. R | Class / Mechanism of Action | |
| Indications Labeled Indications: Acute treatment of ventricular arrhythmias from myocardial infarction (alternate to amiodarone when amiodarone of available) Unlabeled: (ACLS, 2015) Hemodynamically stable monomorphic VT and polymorphic VT Pulseless VT / VF (unresponsive to defibrillation, CPR, and vasopressor administration) Monomorphic VT secondary to drug, when amiodarone is not available Contraindications • Hypersensitivity to lidocaine or any component of the formulation • Prophylactic use in AMI Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block • Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions • Continuous EKG monitoring is necessary • Increased ventricular rate may be seen when given to a patient in A Fib. • At high doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. • D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT Perfusing Arrhythmia (if amiodarone is not available): (ARC 2020); IV, IO Initial dose: 1 to 1.5mg/kg • Forlow with continuous infusion 1-5 to 0.75mg/kg every 5 to 10 minutes • Maximum cumulative dose 3 mg/kg | Antiarrhythmic | |
| Labeled Indications: Acute treatment of ventricular arrhythmias from myocardial infarction (alternate to amiodarone when amiodarone not available) Unlabeled: (ACLS, 2015) Hemodynamically stable monomorphic VT and polymorphic VT Pulseless VT / VF (unresponsive to defibrillation, CPR, and vasopressor administration) Monomorphic VT secondary to drug, when amiodarone is not available Contraindications Hypersensitivity to lidocaine or any component of the formulation Prophylactic use in AMI Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions Continuous EKG monitoring is necessary Increased ventricular rate may be seen when given to a patient in A Fib. At high doses, monitor closely for CNS toxicity, seizure, depression, o D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT VE/Pulseless VT, Wide Complex Tachycardia (with auilable): (ARC 2020): IV, IO Initial dose: 1 to 1.5mg/kg Forterfactory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 0 minutes of Maximum of 3 doses or total of 3mg/kg o Follow with continuous infusion 1-4 mg/min (2.2020); IV, IO <td< td=""><td>Suppresses automaticity of cardiac conduction tissue</td><td></td></td<> | Suppresses automaticity of cardiac conduction tissue | |
| amiodarone when amiodarone not available) Unlabeled: (ACLS, 2015) Hemodynamically stable monomorphic VT and polymorphic VT Pulseless VT / VF (unresponsive to defibrillation, CPR, and vasopressor administration) Monomorphic VT secondary to drug, when amiodarone is not available Contraindications Hypersensitivity to lidocaine or any component of the formulation Prophylactic use in AMI Bradycardia, severe degrees (2 nd or 3 rd) of SA, AV, or intraventricular heart block Wolff-Parkinson-White syndrome, Adam-Stokes syndrome Adverse Reactions / Precautions Continuous EKG monitoring is necessary Increased ventricular rate may be seen when given to a patient in A Fib. A thigh doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. o D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape Cardiac Arrest from VF/VT, (if Amiodarone is not available): (ARC 2020): IV, IO: Initial dose: 1 to 1.5mg/kg Follow with continuous infusion 1-4 mg/min agionificant ectopy: IV, IO 1 to 1.5mg/kg, Repeat 0.5 to 0.75mg/kg every 5 to 10 minutes O Maximum or 3 doses 3 mg/kg Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg preat in 5 to 0.75mg/kg every 5 to 10 minutes O Maximum culualitive dose 3 mg/kg Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg preat 0.5 to 0.75mg/kg every 5 to 10 minutes O Maximum culualitive dose 3 mg/kg Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg preat 0.5 to 0.75mg/kg every 5 to 10 minutes O Maximum culualitive dose 3 mg/kg Disent Infusion initiatel > 15 min after initial bolus therapy) Elush after initiation of ID: May add 2-3 mLidocaine 2% (without epinephrine) to Sml NS flush Local Anesthesia during Tube/Finger Thoracostomy Draw fund 2% Lidocaine and locally anesthetize | Indications | |
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| Continuous EKG monitoring is necessary Increased ventricular rate may be seen when given to a patient in A Fib. At high doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape Cardiac Arrest from VF/VT, (if Amiodarone is not available): (ARC 2020): IV, IO: Initial dose: 1 to 1.5mg/kg For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes Maximum of 3 doses or total of 3mg/kg Follow with continuous infusion 1.4 mg/min (20-50 mcg/kg/minute) Fulsh after initiation of IO: Max add 2-3 ml Lidocaine 2% (without epinephrine) to 5ml NS flush Local Anesthesia during Tube/Finger Thoracostomy Tor a moloni 2% Lidocaine and locally anesthetize | Prophylactic use in AMI Bradycardia, severe degrees (2nd or 3rd) of SA, A^N | V, or intraventricular heart block |
| Increased ventricular rate may be seen when given to a patient in A Fib. At high doses, monitor closely for CNS toxicity, seizure, depression, and respiratory depression. D/C immediately if toxicity develops The elderly may have increased chance of CNS and cardiovascular side effects. Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape Cardiac Arrest from VF/VT, (if Amiodarone is not available): (ARC 2020): IV, IO: Initial dose: 1 to 1.5mg/kg For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes Maximum of 3 doses or total of 3mg/kg Follow with continuous infusion 1-4 mg/min (aff amiodarone is not available): Stable VT, wide complex tachycardia, significant ectopy: IV, IO I to 1.5mg/kg. Repeat 0.5 to 0.75mg/kg every 5 to 10 minutes | • | |
| Cardiac Arrest from VF/VT, (if Amiodarone is not available): (ARC 2020): VF/Pulseless VT, Wide Complex Tachycardia (with pulses): (PALS, 2020) IV, IO: Initial dose: 1 to 1.5mg/kg • For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes • Maximum of 3 doses or total of 3mg/kg • Follow with continuous infusion 1-4mg/minute after ROSC Maintenance Infusion (Peds): Perfusing Arrhythmia (if amiodarone is not available): Stable VT, wide complex tachycardia, significant ectopy: IV, IO: IV, IO • 1 to 1.5mg/kg. Repeat 0.5 to 0.75mg/kg every 5 to 10 minutes • 0 Maximum cumulative dose 3 mg/kg • Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg/minute) • Maximum cumulative dose 3 mg/kg Fulsh after initiation of IO: • May add 2-3 ml Lidocaine 2% (without epinephrine) to 5ml NS flush Local Anesthesia during Tube/Finger Thoracostomy • Draw 10ml 2% Lidocaine and locally anesthetize | At high doses, monitor closely for CNS toxicity, see D/C immediately if toxicity develops The elderly may have increased chance of CNS and the second se | eizure, depression, and respiratory depression. and cardiovascular side effects. |
| available): (ARC 2020): IV, IO: Initial dose: 1 to 1.5mg/kg For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes Maximum of 3 doses or total of 3mg/kg Follow with continuous infusion 1- 4mg/minute after ROSC Perfusing Arrhythmia (if amiodarone is not available): Stable VT, wide complex tachycardia, significant ectopy: IV, IO 1 to 1.5mg/kg. Repeat 0.5 to 0.75mg/kg every 5 to 10 minutes Maximum cumulative dose 3 mg/kg Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg/minute) Flush after initiation of IO: May add 2-3 ml Lidocaine 2% (without epinephrine) to 5ml NS flush Local Anesthesia during Tube/Finger Thoracostomy Draw 10ml 2% Lidocaine and locally anesthetize | Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| Decompression Illness/ Arterial Gas Embolism: | <u>available): (ARC 2020):</u> IV, IO: Initial dose: 1 to 1.5mg/kg For refractory VF may give additional 0.5 to 0.75mg/kg IV push, repeat in 5 to 10 minutes Maximum of 3 doses or total of 3mg/kg Follow with continuous infusion 1- 4mg/minute after ROSC Perfusing Arrhythmia (if amiodarone is not available): Stable VT, wide complex tachycardia, significant ectopy: IV, IO 1 to 1.5mg/kg. Repeat 0.5 to 0.75mg/kg every 5 to 10 minutes Maximum cumulative dose 3 mg/kg Follow with continuous infusion 1-4 mg/min (20-50 mcg/kg/minute) Flush after initiation of IO: May add 2-3 ml Lidocaine 2% (without epinephrine) to 5ml NS flush Local Anesthesia during Tube/Finger Thoracostomy Draw 10ml 2% Lidocaine and locally anesthetize incision area. | pulses): (PALS, 2020) IV, IO: Initial dose: 1mg/kg loading dose. Maintenance Infusion (Peds): IV, IO: Continuous Infusion 20 to 50 mcg/kg per min infusion (repeat bolus dose if infusion initiated > 15 min after initial bolus |
| | • 1.5mg/kg IV/IO | |

| LORAZEPAM QD, Lactation Yes (n | ot recommended) Trade Name: Ativan | |
|---|--|--|
| Class / Mechanism of Action | | |
| Benzodiazepine Acts as an Anxiolytic/Hypnotic, anticonvulsant and se Onset of action: IV Sedation 2-3 minutes; IM hypnoti | | |
| Indications | | |
| Labeled Indications: Anesthesia premedication, Statunation Rapid tranquilization of the combative / agitated p Alcohol withdrawal delirium / syndrome Seizures Induce Sedation and Amnesia (Midazolam is print) | patient | |
| Contraindications | | |
| Hypersensitivity to Lorazepam or any component of the formulation or other benzodiazepines Acute narrow angle glaucoma, Acute Alcohol Intoxication, Sleep apnea Respiratory Insufficiency/Depression (except during mechanical ventilation) Overdose Reversal: <u>FLUMAZENIL</u> can be used; however, it carries elevated risk. Respiratory support until the medication is metabolized is traditionally the best care in Benzodiazepine overdose | | |
| Neurologic Depression (Head Trauma) (unless had Adverse Reactions / Precautions | aving active seizure) | |
| <u>No Analgesic properties</u> (Narcotic pain control is needed for RSI'd / Intubated trauma patients) May Cause Respiratory depression: Do not give without stable IV line and BVM (airway control) ready Hypotension, vasodilation Amnesia, confusion, drowsiness, slurred speech (Paradoxical Reactions possible: aggressiveness, agitation, anxiety, inappropriate behavior) | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
| Acute Seizures: IV: • 4 mg at a maximum rate of 2mg/min; may repeat at 3-5 min. Note: Not recommended IM for seizure due to erratic absorption. Anxiety: IV: | <u>Acute Seizures / Status epilepticus (unlabeled use):</u> IV/IO: 0.1 mg/kg; slow push. May repeat x1 dose in 5-10 minutes. Maximum dose is 4mg/dose. <u>Agitation:</u> IV/IM 0.02-0.1 mg/kg/dose q 20-30 min PRN. (Maximum dose: 2mg/dose). | |
| 0.5-2 mg slow IV push <u>Rapid tranquilization of agitated / combative</u> <u>patient (Off-label use):</u> IV, IM: 2-4mg every 30-60 minutes; may be used alone or administered with an antipsychotic (i.e., haloperidol) | | |

| MAGNESIUM SULFATE | D, Lactation Yes(Caution) | |
|---|--|--|
| Class / Mechanism of Action | | |
| Anticonvulsant, Electrolyte Supplement IV magnesium decreases acetylcholine in motor nerve terminals and slows rate of SA node impulse formation and prolongs conduction time. Magnesium functions to facilitate the movement of calcium, sodium, and potassium in and out of cells. | | |
| Indications | | |
| Labeled Indications: Prevention and treatment of seizures in pregnance Torsades de Pointes: Cardiac arrhythmias (VT/V | | |
| Contraindications | | |
| Hypersensitivity any component of the formulation Myocardial damage and heart blocks Use for pre-eclampsia / eclampsia during 2-hour period before delivery | | |
| Adverse Reactions / Precautions | | |
| Possible cardiovascular arrest, respiratory depres Hypomagnesaemia is often joined by hypokalem potassium. | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
| Torsades de pointes or VF/pulseless VT associated with torsades de pointes (unlabeled use): IV, IO: 1-2 g over 15 minutes; if no response or torsades de pointes recurs, may repeat dose to a total of 4g in 1 hour. | Torsades de pointes: (PALS 2020) IV, IO: • • 25-50 mg/kg/dose over several minutes • maximum single dose: 2000 mg Respiratory Distress (Status Asthmaticus): IV: | |
| Wheezing in Respiratory Distress (3rd line drug): | • 25-50 mg/kg over 30 min (max 2 grams) | |
| 2 Grams single dose over 20min | | |
| Seizure (Refractory to Benzodiazepines): IV: | Magnesium Sulfate should be diluted into 50-100ml NS or D5W for all Adult and Pediatric infusions | |
| • 1-2 Grams over 30 min | | |
| Eclampsia/pre-eclampsia, severe (unlabeled): | | |
| IV: | | |

MANNITOL 20%

QC, Lactation? (Caution)

Class / Mechanism of Action

Osmotic Diuretic

Increases osmotic pressure of glomerular filtrate. This reduces kidney reabsorption of water and electrolytes and increases urinary output. Decreases cerebral blood volume and intracranial pressure (ICP) while increasing cerebral blood flow and O2 transport. Onset of action is 15-30 minutes

Indications Labeled Indications: Reduction of increased ICP secondary to cerebral edema Reduction of elevated intraocular pressure • • Urinary excretion of toxic substances **Contraindications** Hypersensitivity to mannitol or any component of the formulation Active intracranial bleeding • Pulmonary congestion and edema . Severe renal disease, or renal dysfunction after mannitol use Severe dehydration: (Do NOT use in under-resuscitated or hypotensive casualties) • **Adverse Reactions / Precautions** Chest pain, CHF, tachycardia, circulatory overload (with rapid administration), peripheral edema • Headache, seizure • Fluid and electrolyte imbalance, dehydration and hypovolemia • Keep in a temperature-controlled climate. Will crystalize at low temperatures. PEDIATRIC Always Reference BROSELOW Tape Dose and Administration: ADULT Moderate to severe head injury, Patient Increased intracranial pressure (unlabeled continuing to deteriorate or showing signs of dosing): herniation despite adjustment to ventilation and IV: starting hypertonic saline. 0.25-1 g/kg/dose. • IV Maintenance dose of 0.25-0.5 g/kg IV q 4-6hrs • prn to maintain serum osmolality <300-320 1 g/kg IV bolus over <20 minutes. • mOsm/kg • Follow with 0.25 g/kg IVP every 4 hours Note: Always have urinary catheter in place and monitor output. Note: 3% Hypertonic Saline is preferred over Mannitol

| METHYLPREDNISOLONE QC, Lactati | ion Yes(Caution) Trade Name: SoluMedrol | |
|--|--|--|
| Class / Mechanism of Action | | |
| Systemic Corticosteroid Anti-inflammatory, Immunosuppressant, shock | | |
| Indications | | |
| Labeled Indications: Treatment of a variety of disea and autoimmune; Unlabeled: None identified unless added by medical direction. | ses: allergic, inflammatory, hematologic, neoplastic, | |
| Contraindications | | |
| Hypersensitivity to methylprednisolone or any cor No other in emergency setting | nponent of the formulation | |
| Adverse Reactions / Precautions | | |
| Not for use in treatment of head injury; increased mortality has occurred in head injury patients treated with high dose IV methylprednisolone. No immediate effect will be observed while treating in the pre-hospital environment. Onset of action may take several hours | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
| Asthma exacerbations, including status asthmaticus IV: • 125mg x 1 dose Allergic Reaction: IV: • 125mg x 1 dose | Asthma exacerbations, including status asthmaticus IV: • Children <12 years: 1-2 mg/kg initial dose; followed by 0.5-1 mg/kg q 6 hrs. (maximum: 60 mg/day) Allergic Reaction IV • 2 mg/kg y 1 does | |
| Note: Only methylprednisolone sodium succinate can be used for IV doses. | 2 mg/kg x 1 dose Note: Only methylprednisolone sodium succinate can be used for IV doses. | |

METOCLOPRAMIDE **QB**

Trade Name: Reglan

Class / Mechanism of Action

Prokinetic Agent: Antiemetic, Upper GI Stimulant Potent dopamine-receptor antagonist. At higher doses blocks serotonin receptor in chemoreceptor trigger zones of CNS. Increases GI tract motility and gastric emptying. Onset of action 1-5 minutes via IV with a duration of 1-2 hours.

Indications

Labeled Indications: Prevention of postoperative nausea and vomiting; Acid Reflux/Heartburn/GERD; Migraine Headache

Contraindications

- Hypersensitivity to glucagon or any component of the formulation
- Insulinoma / Pheochromocytoma

Adverse Reactions / Precautions

• Hypersensitivity, History of tardive dyskinesia or dystonic reaction to Metoclopramide in the past, GI Obstruction or Hemorrhage, and seizure disorder (epilepsy).

| Dose and Administration: AD | PEDIATRIC Always Reference BROSELOW Tape | |
|---|--|--|
| 5-10 mg IV/IO slow push; repeat dose x1 ev 30 minutes PRN for a max of 20mg. | very 20- Not recommended or approved for routine pediatric use | |
| *Rapid IV push may cause intense feelings of anxiety and restlessness* | of | |
| Ideal administration to Prevent Agitation/Adv Effects: Dilute in a 50 or 100ml NS bag and over 10-15 minutes. | | |

| METOPROLOL | QC, Lactation? (Not Re | ecommended) | Trade Name: Lopressor |
|---|--|---|---|
| Class / Mechanism of Action | T- , -actailent (1.501 (1. | | |
| Beta-1 Selective Beta-Blocker Selective inhibitor of beta ₁ -adre effect on beta ₂ -receptors at oral | Beta-1 Selective Beta-Blocker; Antihypertensive; Antianginal Agent Selective inhibitor of beta ₁ -adrenergic receptors; competitively blocks beta ₁ -receptors, with little or no effect on beta ₂ -receptors at oral doses <100 mg (in adults); does not exhibit any membrane stabilizing or intrinsic sympathomimetic activity. | | |
| Indications | | | |
| Labeled Indications: Angina, H Unlabeled: Atrial fibrillation/flut Migraine prophylaxis; Supraver Thyrotoxicosis; Ventricular arrh | er; Hypertrophic cardiom tricular tachycardia (AVN | yopathy; Marfan syn | |
| Contraindications | | | |
| Hypersensitivity to metopro third-degree heart block Severe sinus bradycardia (l interval ≥0.24 seconds); systematical seconds | neart rate <45 beats/minu | te); significant first-c | degree heart block (P-R |
| Adverse Reactions / Precaution | ons | | |
| Cardiovascular: Hypotensic cardiac failure, CVA, cold e Central nervous system: Dia hallucination, headache, inst | xtremities, palpitations, pe zziness, fatigue, depression | eripheral edema, cla on, vertigo, confusio orary amnesia, tinni | nudication n, disturbed sleep, itus |
| | ADULT | | lways Reference BROSELOW Tape |
| Atrial fibrillation or atrial flutter Acute ventricular rate control IV: 2.5 to 5 mg over 2-5 minuter minutes as needed; maximer Supraventricular tachycardiaa arrhythmias (off-label use): Note: For hemodynamically state maneuvers and/or adenosine at IV: | : es; repeat dose every 5 um total dose: 15 mg. Ventricular ble patients if vagal re unsuccessful. | blockers as initia patients; beta-blo for use in patients contraindications after ≥2 preferred patients with hyp | do not recommend beta- I therapy in pediatric ickers should be reserved is who have is to preferred agents or agents have failed in ertension and chronic proteinuria, or diabetes |
| 2.5 to 5 mg over 2-5 minut minutes as needed to achie 90 – 100; maximum total do Note: For sustained ventricular blockers are generally administ antiarrhythmic drug (eg, Amioda indications. A beta-blocker is al shocks in patients who receive cardioverter defibrillator for thes propranolol may be the preferre situations | eve a ventricular rate of ose: 15 mg. tachycardia, Beta- ered in addition to an arone) for these so used to reduce an implantable se indications; | | |

MIDAZOLAM

QD, Lactation Yes (Caution)

Trade Name: Versed

Class / Mechanism of Action

Benzodiazepine

Acts as an Anxiolytic/Hypnotic, anticonvulsant and sedative.

Onset of action: Sedation; IV: 1-5 minutes, IM: 15 minutes, Intranasal: 4-8 minutes

Duration: IV, less than 2 hours. (20-30 Minutes per ECCN Nurse Protocols, May 2012)

Indications

Labeled Indications: Preoperative sedation, induction and maintenance of general anesthesia **Unlabeled:** Anxiety / agitation, status epilepticus, conscious sedation (intranasal)

Contraindications

- Hypersensitivity to midazolam or any component of the formulation or other benzodiazepines
- Acute narrow angle glaucoma, Acute Alcohol Intoxication
- Respiratory Insufficiency/Depression (except during mechanical ventilation)
- (Overdose Reversal: <u>FLUMAZENIL</u> can be used; however, it carries elevated risk. Respiratory support until the medication is metabolized is traditionally the best care in Benzodiazepine overdose)
- Should not be used in shock
- Neurologic Depression (Head Trauma) (unless having active seizure)

Adverse Reactions / Precautions

- No Analgesic properties (Narcotic pain control is needed for RSI'd / Intubated trauma patients)
- May Cause Respiratory depression: Do not give without stable IV line and BVM (airway control) ready
- Hypotension, vasodilation
- Amnesia, confusion, drowsiness, slurred speech (Paradoxical Reactions possible: aggressiveness, agitation, anxiety, inappropriate behavior)

| Dose and Administration: ADUL | T PEDIATRIC Always Reference BROSELOW Tape |
|---|--|
| Induction for RSI; Continued sedation; Hyperthermia: IV: Induction 0.1mg/kg IV/IO Continued Sedation .051 mg/kg IV/IO Infusion sedation 0.05 mg/kg bolus | Procedural sedation; Transcutaneous Pacing; Cardioversion: IV: 0.05-0.1mg/kg q 15-30 min PRN Intranasal (unlabeled route): 0.2-0.5 mg/kg (maximum total dose: 10 mg or 5 mg per |
| IV, then titrate 0.05-0.1mg/kg/hr. IV gtt <u>Transcutaneous Pacing /</u> <u>Cardioversion, Anxiety, Agitation.</u> IV | nare Induction/RSI (Not preferred drug) IV: • 0.1-0.3 mg/kg Seizure |
| 2.5-5mg q 15-30 min PRN Seizure Dosage: If no IV/IO access, 5mg IN (repeat in 10 minutes in opposite nostril if still seizing (preferred) or 10mg IM (alternate) 5mg IV/IO, may repeat After 3 doses should consider addition of another agent. | Seizure IV, IM: 0.2 mg/kg Q 15-30 min PRN Status epilepticus, prehospital treatment (unlabeled use): IV: Infants: 1-2 mg 13-40 kg: 4 mg once >40 kg: Refer to adult dosing |

| MORPHINE QC, Lactation Yes(Caution) | | |
|---|---|--|
| Class / Mechanism of Action | | |
| Opioid Analgesic Binds to opioid receptors within the CNS increasing p ascending pain pathways (blocking painful stimulus); Onset: IV variable but rapid, Duration variable, patient | produces CNS depression | |
| Indications | 1 | |
| Labeled Indications: Moderate to severe acute and preanesthetic medication | chronic pain; pain of myocardial infarction; | |
| Contraindications | | |
| Hypersensitivity to morphine sulphate or any component of the formulation Severe respiratory depression Acute or severe asthma (in an unmonitored setting or without resuscitative equipment) Paralytic ileus | | |
| Adverse Reactions / Precautions | | |
| Head trauma: Use with extreme caution in head injury, or suspected increased ICP; exaggerated increase in ICP may occur. Some formulations are specifically contraindicated. May cause Hypotension, Use with caution in hypovolemic patients. May worsen Bradycardia May cause life-threatening hypoventilation and Reparatory depression CNS depression: Impairs physical and mental abilities | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
| Chest Pain/AMI: IV/IO: | Acute pain (moderate-to-severe): | |
| 2-5 mg q 5-15 min PRN Acute pain (moderate-to-severe): | IM, SubQ: The use of IM/ SubQ injections is no longer recommended especially for repeated administration due to painful administration, variable absorption and lag time to peak effect. | |
| IM, SubQ: The use of IM/ SubQ injections is no longer recommended especially for repeated administration due to painful administration, variable absorption and lag time to peak effect. IV/IO: (Slow) 5mg (0.1 mg/kg, range 2.5 – 10mg) every 1-6 hours PRN | IV: (Slow) 0.1-0.2 mg/kg q 2-4 hr. PRN, not to exceed 10 mg per dose Continuous infusion: 10-30 mcg/kg/hour; titrate PRN for pain | |

MOXIFLOXACIN

QC, Lactation Yes

Trade Name Avelox

Class / Mechanism of Action

Antibiotic (Fluoroquinolone)

Bactericidal - DNA gyrase inhibitor and topoisomerase IV inhibitor – which is an essential enzyme that maintains the superhelical structure, replication, transcription, and repair of bacterial DNA.

Indications

Labeled Indications: Used for infection control prophylaxis for traumatic open injuries and surgical prophylaxis.

Contraindications

- Hypersensitivity to cefazolin, other cephalosporin antibiotics, other beta-lactams, or any component of the formulation
- May cause QT prolongation.
- Avoid use in known aortic aneurysm or dissection

Adverse Reactions / Precautions

• Superinfection – prolonged use may result in fungal or bacterial superinfection (including C.Difficile)

| Dose and Administration: ADULT | PEDIATRIC |
|---|--|
| Infection Control: For PO tolerable patients | Infection Control: |
| PO: Adults: • 400 mg once daily o Max daily dose: 400 mg/day | PO: Pediatrics: <15yrs old: • 10 mg/kg/day PO ○ Max daily dose: 400 mg/day >15yrs old: • 400 mg once daily ○ Max daily dose: 400 mg/day |

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| NALOXONE QC, Lactation (| Caution) Trade Name: Narcan | |
| Class / Mechanism of Action | | |
| Antidote, Opioid Antagonist Competes and displaces opioids at opioid receptor s | ites, reversing narcotic effects. | |
| Indications | | |
| Labeled Indications: Reversal of opioid drug effects | , including respiratory depression | |
| Contraindications | | |
| Hypersensitivity to naloxone or any component of | f the formulation | |
| Adverse Reactions / Precautions | | |
| When correcting for respiratory depression in a postoperative (intubated patient), carefully titrate the dose to reverse hypoventilation; do not fully awaken patient or reverse analgesic effect. Recurrence of respiratory depression is possible continue to watch for respiratory depression until patient hand-off. May cause narcotic withdrawal effects | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | |
| Opioid overdose (with standard ACS protocols): IV, IM, SubQ: 0.4-2 mg; may dose every 2-3 minutes if needed. | Opioid overdose (with standard PALS protocols): IV, IM, SubQ: <5 years or ≤20 kg (unlabeled dose): 0.1 mg/kg/dose (maximum dose: 2 mg); repeat | |
| If no response after 10 mg total, look for other cause of respiratory depression. | every 2-3 minutes PRN • ≥5 years or >20 kg: Adult Dosing | |
| Following reversal, may need to readminister after 20-60 minutes. | Reversal of respiratory depression with therapeutic opioid doses: IV, IM, SubQ: | |
| Reversal of respiratory depression with therapeutic opioid doses: IV, IM, SubQ: | 0.001-0.015 mg/kg/dose; repeat as needed. | |
| • 0.1-0.4 mg titrated to adequate respiratory rate. If not improved after 0.8 mg total, look for other cause of respiratory depression. | | |

| NIFEDIPINE QC. | Lactation Ves(Not | Recommended) | Trade Name: Procardia |
|---|------------------------------|--|---|
| | | | |
| Class / Mechanism of Action | | | |
| Antianginal Agent, Calcium Channel Blocker Inhibits movement of calcium ion across cell membranes of smooth muscle and myocardium resulting in relaxation of coronary vascular smooth muscle and vasodilation as well as reduced peripheral vascular resistance (reducing blood pressure). | | | |
| Indications | | | |
| Labeled Indications: Chronic stable or vasospastic angina Unlabeled: Prevention and treatment of high altitude pulmonary edema | | | |
| Contraindications | | | |
| Hypersensitivity to nifedipine or any component of the formulation Cardiogenic Shock Acute MI | | | |
| Adverse Reactions / Precautions | | | |
| Symptomatic hypotension:Bradycardia, nausea | | | |
| Dose and Administration: | ADULT | | ays Reference BROSELOW Tape |
| High altitude pulmonary edema PO: 10 mg every 4-6 hours Pulmonary hypertension (unlabe PO: 30 mg (Extended Release) two increase cautiously to 120-24 | eled use) rice daily; may | High altitude pulmonar approved for use in child PO: Immediate release: 0 (maximum: 20 mg/do Note: Treatment is need response to oxygen and/ | ren) (unlabeled use): 0.5 mg/kg/dose ose) every 8 hours ed only necessary if |
| Note: Do not use for acute angina precipitate myocardial infarction | l episodes; may | | |

NITROGLYCERIN

QC. Lactation (Caution)

Trade Name: NitroMist/Nitrostat

Class / Mechanism of Action

Antianginal agent, Vasodilator

Induces smooth muscle relaxation and vasodilation of peripheral veins and arteries and coronary arteries thus improving collateral blood flow to ischemic regions of the myocardium. Reduces cardiac oxygen demand by decreasing preload. Onset of action: Sublingual tablet and spray, 1-3 minutes. Duration: 25 minutes

Indications

Labeled Indications: Treatment or prevention of angina pectoris

Contraindications

- Hypersensitivity to nitrates or any component of the formulation
- Use with phosphodiesterase-5 inhibitors (Sildenafil, Levitra, Cialis) in previous 48hrs
- Increased intracranial pressure
- Hypotension (SBP <90mmHg or >30mmHg below baseline), Bradycardia <50bpm, Tachycardia without heart failure (>100bpm), and Right ventricular infarction.

Adverse Reactions / Precautions

• IV/IO access should be placed and SBP should be > 110.

- Use cautiously in cases of chest pain unless inferior wall / right-ventricular MI can be ruled-out by ECG prior to administration
- Can cause severe hypotension with associated paradoxical bradycardia and increased angina
- Use with caution in volume depleted patients
- Do not use for inferior wall MI and suspected right ventricular involvement

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | | |
|--|---|--|--|
| <u>Angina/coronary artery disease:</u> PO: <u>Sublingual</u>: 0.4 mg every 5 minutes for maximum of 3 doses in 15 minutes <u>Translingual</u>: 1 spray (0.4mg per spray) onto or under tongue every 3-5 minutes for maximum of 3 doses in 15 minutes <u>CHF related Respiratory Distress:</u> PO: | Not indicated in most children, even with heart failure, as their heart failure is not usually due to coronary artery disease. Could cause significant problems in those with depressed myocardial function. Consult Medical Direction (if able) before use in Pediatrics.CHF related Respiratory Distress: PO:• 0.4mg q 5min if SBP > 70 + 2 x Age | | |
| <u>Sublingual</u>: 0.4 mg every 5 minutes for maximum of 3 doses in 15 minutes as long as SBP>90 | CHF or Cardiogenic Shock: IV Drip: Children: 0.25 - 0.5 mcg/kg/min; titrate by 1 mcg/kg/min q 15-20 min as tolerated (Typical dose=1-5mcg/kg/min)(Max 10mcg/kg/min) | | |
| IV Drip: (Only used at written direction of referring provider or consultation with medical director) Start at 10 mcg/min, titrate up or down to: 10% reduction in MAP if normotensive 30% reduction in MAP if hypertensive. Max dose: 400mcg/minute) | Adolescents: 5-10 mcg/min (not per kg) (max 200 mcg/min) | | |

NOREPINEPHRINE

QC, Lactation? (Caution)

Trade Name: Levophed

Class / Mechanism of Action

Alpha and Beta Agonist

Stimulates beta₁ and alpha-adrenergic receptors increases contractility, heart rate, and vasoconstriction. Increases systemic blood pressure and coronary blood flow. Effects on vasoconstriction (alpha receptors) are greater than inotropic (beta receptors). Onset of action: IV very rapid. Duration: 1-2 minutes

Indications

Labeled Indications: Treatment of shock persisting after adequate fluid volume replacement; severe hypotension.

ALS 2020: Severe cardiogenic shock and hemodynamically significant hypotension (SBP <70mmHg) with low total peripheral resistance. Agent of last resort for management of ischemic heart disease and shock.

Contraindications

- Hypersensitivity to norepinephrine, bisulfites or any component of the formulation
- Hypotension from hypovolemia except as an emergency measure to maintain coronary and cerebral perfusion until volume can be replaced

- No applicable use in hemorrhagic shock unless fluid replacement therapy maximized! Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shock.
- Strong Vesicant: ensure proper catheter placement and avoid extravasation, use a large vein (preferably a central line) and avoid leg veins.
- Assure adequate circulatory volume to minimize need for vasoconstrictors. Monitor BP closely, <u>avoid</u> <u>hypertension</u> and adjust infusion rate as needed.

| Dose and Administration: | ADULT | PEDIATRIC |
|--|---|--|
| Hypotension/shock: IV: Administer as continuous infusion wit same line as sodium bicarbonate. It will i Initial: 2-20 mcg/minute; titrate to SE Maintenance: 2-4 mcg/minute Post ROSC Hypotension: Initial: 0.1-0.5 mcg/kg/minute titrate | nactivate norepinephrine. 3P goal. te | Hypotension/shock: IV: Continuous infusion Initial: 0.05-0.1 mcg/kg/minute; titrate to effect Max dose: 2 mcg/kg/minute |
| If unable to maintain MAP >60mmHg, add <u>Use in Burn Patient:</u> For Burn patients, norepinephrine is only and UOP (<u>></u> 30mL/hr) fail to be reached w sequence of use follows administration of | v used when target MAP (<u>></u> 55) vith fluid resuscitation alone. Its f <u>Vasopressin</u> . | |
| (See infusion chart next page for n | nix and dosage information) | |

| | NOREPINEPHRINE (LEVOPHED) | | | | | |
|--|--|---------|-----------|-----------|---------|--|
| Dos | Dosing Range: 2-20mcg/min (120-1200mcg/hr) | | | | | |
| | MIX 4 mg/500 mL | | | | | |
| | C | ONCENTR | ATION 8 m | icg/mL | | |
| Dose | Rate | Micro | | Macro | | |
| | | (60 | 20 | 15 gtt/mL | 10 | |
| | | gtt/mL) | gtt/mL | | gtt/mL | |
| mcg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | |
| 2 | 15 | 15 | 5 | 4 | 3 | |
| 3 | 23 | 23 | 8 | 6 | 4 | |
| 4 | 30 | 30 | 10 | 8 | 5 | |
| 5 | 38 | 38 | 13 | 10 | 6 | |
| 6 | 45 | 45 | 15 | 11 | 8 | |
| 7 | 53 | 53 | 18 | 13 | 9 | |
| 8 | 60 | 60 | 20 | 15 | 10 | |
| 9 | 68 | 68 | 23 | 17 | 11 | |
| 10 | 75 | 75 | 25 | 19 | 13 | |
| 11 | 83 | 83 | 28 | 21 | 14 | |
| 12 | 90 | 90 | 30 | 23 | 15 | |
| 13 | 98 | 98 | 33 | 25 | 16 | |
| 14 | 105 | 105 | 35 | 26 | 18 | |
| 15 | 113 | 113 | 38 | 28 | 19 | |
| 16 | 120 | 120 | 40 | 30 | 20 | |
| 17 | 128 | 128 | 43 | 32 | 21 | |
| 18 | 135 | 135 | 45 | 34 | 23 | |
| 19 | 143 | 143 | 48 | 36 | 24 | |
| 20 | 150 | 150 | 50 | 38 | 25 | |
| Macro | Macro-Drip (20gtt/ml) or Micro-Drip is set of choice for | | | | | |
| this | | | | | | |
| infusion | | | | | | |
| Start at lowest dose and increase rate by 0.5mcg/min | | | | | | |
| every 2 minutes PRN to target MAP >60mmHg | | | | | | |
| | | | | | | |

0.9% Sodium Chloride (Normal Saline)

Class: Isotonic crystalloid solution.

Mechanism of Action: Replaces water and electrolytes.

Indications: Hypovolemia, Shock, Heat-related injuries, diabetic ketoacidosis, TKO IV, a diluent of choice for blood product transfusion.

Contraindications: Avoid for intravascular volume replacement for hemorrhagic shock due to hemodilution and hyperchloremic metabolic acidosis. Use with caution in patients with known congestive heart failure.

Adverse Reactions: Rare

Drug Interactions: Few in the pre-hospital emergency setting.

How Supplied: 250mL, 500mL, and 1,000mL bags.

Dosage and Administration: The specific situation being treated will dictate the rate in which normal saline will be administered. Hypovolemic shock requires rapid bolus (see relevant guidelines). In other cases, it is advisable to administer the fluid at a moderate rate (for example, 100 mL/h).

Hypertonic Saline 3% Sodium Chloride

Class: Hypertonic crystalloid solution.

Mechanism of Action: Replaces water and electrolytes, reduces the amount of fluid in the cranial cavity, decreases ICP, increases intravascular sodium concentration, may induce diuresis.

Indications: Refractory elevated intracranial pressure (ICP) due to various etiologies (eg, subarachnoid hemorrhage, neoplasm); traumatic brain injury with elevated ICP (can be used in place of mannitol). **Contraindications:** Do not use in the same line as blood products – cause crenation and lysis of RBC. Caution or avoid use in patients with known concestive heart failure and kidney disease.

Adverse Reactions: Rare.

Drug Interactions: Few in the pre-hospital emergency setting.

How Supplied: 250mL, 500mL, bags.

Dosage and Administration:

- · Dosing (Adult):
 - o Bolus: 250mL IV Bolus over 15 min.
 - o Infusion: 50-100 cc/hr
- Dosing (Pediatrics):
 - o Bolus: 5 cc/kg IV Bolus over 15 min.
 - o Infusion: 0.5 cc/kg/hr
- Should be administered through a central line due to its high osmolarity and tonicity.

ONDANSETRON

QB, Lactation? (Caution)

Trade Name: Zofran

Class / Mechanism of Action

Antiemetic

Blocks serotonin, peripherally on vagus nerve terminals and centrally. Onset of action is 5-30 minutes dependent on route.

Indications

Labeled Indications: Prevention of postoperative nausea and vomiting **Unlabeled:** Hyperemesis gravidarum (severe or refractory)

Unlabeled: Hyperemesis gravidarum (severe or reir

Contraindications

• Hypersensitivity to ondansetron or any component of the formulation

Adverse Reactions / Precautions

• Dose dependent QT interval prolongation occurs and IV doses >16mg are not recommended.

 In most patients, QT changes are not clinically relevant; however, if used with other medications that prolong QT intervals (antiarrhythmics) or in those at risk for QT prolongation, arrhythmia can occur. Torsades de points has been reported.

| Dose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|---|-------|---|
| Nausea and Vomiting: IV/IO/IM/PO • 4-8 mg Treatment of severe or refractory hy gravidum (unlabeled use): IV: • 8 mg administered over 15 minute hours | | Nausea and Vomiting (Children 1 month to 12 years): IV: ≤40 kg: 0.1 mg/kg as a single dose over 2-5 minutes >40 kg: 4 mg as a single dose over 2-5 Minutes |

Oxygen

Class: Atmospheric gas.

Mechanism of Action: Reverses hypoxemia

Duration of action: Onset: immediate. Peak effect: not applicable. Duration: less than 2 minutes. **Indications:** All causes of decreased tissue oxygenation and/or decreased level of

consciousness.(Confirmed or expected hypoxemia, ischemic chest pain, respiratory insufficiency, prophylactically during air transport, as an antidote for confirmed or suspected carbon monoxide poisoning).

Contraindications: Coincidental paraquat (herbicide) inhalation rare, supplemental oxygen enhances the toxicity damaging the alveolar cells; COPD patients may become hypopneic with high O2 flow rates due to "oxygen baroreceptor respiratory drive (relative contraindication).

Adverse Reactions: Retinopathy of prematurity (prolonged use); potential oxygen toxicity in hyperbaric environments; cerebral vasoconstriction.

Drug Interactions: None

Dosage and Administration:

 Assure adequate ventilation (spontaneous or supported) supplemental oxygen therapy, ideally by end- tidal CO₂ measurement (Goal EtCO2 35-45).

- All critically ill and injured transport patients will receive supplemental oxygen to maintain SPO2 of >93%
- Administer oxygen 2-6 LPM via nasal cannula.

o If O2 Saturation remains < 95%, apply non-rebreather face mask with oxygen at 15 LPM. o If O2 Saturation remains < 90%, refer to **Airway guideline**.

- Patient on Ventilator:
 - o Adjust ventilator settings based on ventilatory goals for patient: ETCO2, peak pressures, SpO2, and patient clinical condition.
 - o Adjust FiO2 to maintain pulse oxygen saturations > 93% / tissue oxygen saturation (STO₂) > 70%, if applicable.
- When planning for available O₂ during non-pressurized, aeromedical transfer, ensure adequate resources to provide 1.5 to 2 times the ground transport volume of O₂ to compensate for increased consumption associated with altitude related physiological impact.

PlasmaLyte A

Class: Isotonic crystalloid solution.

Mechanism of Action: Replaces water and electrolytes.

Indications: Hypovolemic shock; compatible with blood or blood components. It may be administered before or following the infusion of blood through the same administration set (i.e., as a priming solution), added to or infused concurrently with blood components, or used as a diluent in the transfusion of packed erythrocytes. PLASMALYTE A and 0.9% Sodium Chloride Injection are equally compatible with blood or blood components.

Contraindications: Use with caution for intravascular volume replacement for hemorrhagic shock due to hemodilution and exacerbation of coagulopathy. Use with caution in patients with known congestive heart failure and kidney disease. Excess administration may result in metabolic alkalosis.

Adverse Reactions: Rare

Drug Interactions: Few in the pre-hospital emergency setting.

How Supplied: 500mL, and 1,000mL bags IV infusion.

Dosage and Administration: Hypovolemic shock; titrate according to the patient's physiologic response. (See appropriate Guidelines)

| PHENYLEPHRINE QC, Lactation | 1? (Caution) Trade Name: Neosynephrine | | | | |
|---|---|--|--|--|--|
| Class / Mechanism of Action | | | | | |
| Alpha Adrenergic Agonist Potent, direct acting alpha adrenergic agonist with virtually no beta-adrenergic activity; causes systemic arterial vasoconstriction. Onset of action IV: Immediate, Duration: approximately 5-10 minutes. | | | | | |
| Indications | | | | | |
| Labeled Indications: Treatment of hypotension, | , vascular failure in shock | | | | |
| Contraindications | | | | | |
| Hypersensitivity to phenylephrine or any com Ventricular Tachycardia and Hypertension Bradycardia | ponent of the formulation | | | | |
| Adverse Reactions / Precautions | | | | | |
| Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shoc Not recommended for routine use in the treatment of septic shock Reflexive Bradycardia. Assure adequate circulatory volume to minimize need for vasoconstrictors. Monitor BP closely, avoid hypertension and adjust infusion rate as needed. Vesicant: Avoid extravasation, will cause tissue damage/necrosis, ensure proper needle placement | | | | | |
| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape | | | | |
| Hypotension / Shock: IV Push: • 50-200 mcg/dose q 5-10 minutes ○ Max 1000 mcg ○ Titrate to blood pressure, use as temporary support or bridge to Vasopressor drip ○ Mix 10mg phenylephrine in 100mL N for a concentration of 100mcg/mL IV Infusion: • 40 - 200 mcg/min; titrate to MAP > 60 mm H ○ To titrate, increase rate by 10 mcg/me every 2 minutes. ○ Maximum dose is 200 mcg/min. ○ Mix 10mg phenylephrine in 250mL D5W/NS for a concentration for 40mcg/mL | Note: Almost never used in pediatric shock. Isolated increased afterload usually causes significant problems in this population. Use with caution and contact Medical Direction if | | | | |
| If unable to maintain MAP >60mmHg, add Epinephrine infusion. | | | | | |

TADIE OF CONTENTS

|] | ABLE OF C | CONTENTS | |
|--|---|---|---|
| PROPOFOL P | B, Lactatior Recommended | | Trade Name: Diprivan |
| Class / Mechanism of Action | | | |
| General Anesthetic Lipophilic intravenous general anesthet Onset of action IV bolus: 9-51 seconds minutes, prolonged with continued dose | (average 30 s | econds), Duration is | s dose and rate dependent: 3-10 |
| Indications | | | |
| Labeled Indications: Induction of anese patients >2 months of age; sedation in | | | |
| Contraindications | | | |
| Hypersensitivity to propofol or any of the second sec | • | | |
| Allergy to eggs, egg products, soyb | eans, soy proc | ducts, and peanuts. | |
| Adverse Reactions / Precautions May cause Hypotension especially | | | |
| Hypotension may result in reduction. Head Injury patients or those increased risk of decreased Do not use in pre-hospital trauma e director or provided written orders to No Analgesic properties. Must supplementation of the second second | with suspecte cerebral perfu- nvironment or by referring pro | d / known increased sion pressure. in burn transfer pati ovider. | · |
| Dose and Administration: | ADULT | PEDIATR | RIC Always Reference BROSELOW Tape |
| Sedation/ RSI: | | Sedation/ RSI: | |
| IV Push: | | IV Push: | every 5-10min PRN. |
| • 1-2.5 mg/kg every 5-10min PRN. | | | eneral anesthesia, |
| Maintenance of general anesthesia: | | IV Infusion: | |
| IV Infusion: | | | months to 16 years: / /kg/minute (or 7.5-18 |
| • 10-75 mcg/kg/min via infusion pump or Dial-a- Drip. Titrate to minimum effective dose. (See infusion chart next page) | | mg/kg/hour) | |
| MAX DOSE: 100 mcg/kg/min. | | | |
| Use of Dial-a-Drip tubing in the absence of an infusion pump will increase accuracy of infusion dosage. | | | |
| Note: Wait 3-5 minutes between dosage clinically assess drug effects. Smaller d required when used with opioids. | | | |
| Note: Not preferred in Burn patients in t 72 hrs | he first 48- | | |

| | | |) mg/50 mL ITRATION 10 n | ng/mL | | |
|----------|------------------|------------|-----------------------------|--------------|--------------|--------------|
| . Weight | Dose | Rate | Micro | Ċ, | Macro | |
| | | | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL |
| kg | mcg/kg/min 10 | mL/hr 3 | gtt/min 3 | gtt/min 1 | gtt/min 1 | gtt/min 1 |
| | 10 | 5 | 5 | 2 | 1 | 1 |
| | 20 | 6 | 6 | 2 | 2 | 1 |
| | 25 | 8 | 8 | 3 | 2 | 1 |
| | 30 | 9 | 9 | 3 | 2 | 2 |
| | 35 | 11 | 11 | 4 | 3 | 2 |
| 50 | 40 45 | 12 14 | 12 14 | 4 | 3 | 2 |
| | 45 50 | 14 | 14 | 5 | 4 | 3 |
| | 55 | 17 | 17 | 6 | 4 | 3 |
| | 60 | 18 | 18 | 6 | 5 | 3 |
| | 65 | 20 | 20 | 7 | 5 | 3 |
| | 70 | 21 | 21 | 7 | 5 | 4 |
| | 75 | 23 | 23 | 8 | 6 | 4 |
| | 10 | 3 | 3 | 1 | 1 | 1 |
| | 20 | 5 | 5 | 2 | 2 | 1 |
| | 25 | 8 | 8 | 3 | 2 | 1 |
| | 30 | 10 | 10 | 3 | 2 | 2 |
| | 35 | 12 | 12 | 4 | 3 | 2 |
| 55 | 40 | 13 | 13 | 4 | 3 | 2 |
| | 45 | 15 | 15 | 5 | 4 | 2 |
| | 50 55 | 17 18 | 17 18 | 6 | 4 | 3 |
| | 55 60 | 18 20 | 18 20 | 6 | 5 | 3 |
| | 65 | 20 | 20 | 7 | 5 | 4 |
| | 70 | 23 | 23 | 8 | 6 | 4 |
| | 75 | 25 | 25 | 8 | 6 | 4 |
| | 10 | 4 | 4 | 1 | 1 | 1 |
| | 15 | 5 | 5 | 2 | 1 | 1 |
| | 20 | 7 | 7 | 2 | 2 | 1 |
| | 25 30 | 9 11 | 9 11 | 3 | 2 | 2 |
| | 35 | 13 | 11 | 4 | 3 | 2 |
| | 40 | 14 | 14 | 5 | 4 | 2 |
| 60 | 45 | 16 | 16 | 5 | 4 | 3 |
| | 50 | 18 | 18 | 6 | 5 | 3 |
| | 55 | 20 | 20 | 7 | 5 | 3 |
| | 60 65 | 22 23 | 22 23 | 7 | 5 | 4 |
| | 70 | 25 | 25 | 8 | 6 | 4 |
| | 75 | 27 | 27 | 9 | 7 | 5 |
| | 10 | 4 | 4 | 1 | 1 | 1 |
| | 15 | 6 | 6 | 2 | 1 | 1 |
| | 20 | 8 | 8 | 3 | 2 | 1 |
| 65 | 25 | 10 | 10 | 3 | 2 | 2 |
| | 30 35 | 12 14 | 12 | 4 | 3 | 2 |
| | 40 | 14 | 14 | 5 | 4 | 3 |
| | 45 | 18 | 18 | 6 | 4 | 3 |
| | 50 | 20 | 20 | 7 | 5 | 3 |
| | 55 | 21 | 21 | 7 | 5 | 4 |
| | 60 | 23 | 23 | 8 | 6 | 4 |
| | 65 | 25 | 25 | 8 | 6 | 4 |
| | 70 75 | 27 29 | 27 29 | 9 10 | 7 | 5 |
| | 10 | 4 | 4 | 10 | 1 | 1 |
| | 15 | 6 | 6 | 2 | 2 | 1 |
| | 20 | 8 | 8 | 3 | 2 | 1 |
| | 25 | 11 | 11 | 4 | 3 | 2 |
| | 30 | 13 | 13 | 4 | 3 | 2 |
| | 35 40 | 15 17 | 15 17 | 5 | 4 | 2 |
| 70 | 40 | 1/ | 17 | 6 | 4 | 3 |
| | 50 | 21 | 21 | 7 | 5 | 4 |
| | 55 | 23 | 23 | 8 | 6 | 4 |
| | 60 | 25 | 25 | 8 | 6 | 4 |
| | 65 | 27 | 27 | 9 | 7 | 5 |
| | 70 | 29 | 29 | 10 | 7 | 5 |
| | 75 | 32 | 32 | 11 | 8 | 5 |
| | 10 15 | 5 | 5 | 2 | 2 | 1 |
| | 20 | 9 | 9 | 3 | 2 | 2 |
| | 25 | 11 | 11 | 4 | 3 | 2 |
| | 30 | 14 | 14 | 5 | 3 | 2 |
| 75 | 35 | 16 | 16 | 5 | 4 | 3 |
| | 40 | 18 | 18 | 6 | 5 | 3 |
| 15 | 45 | 20 | 20 | 7 | 5 | 3 |
| | 50 | 23 | 23 | 8 | 6 | 4 |
| | 55 60 | 25 | 25 | 8 | 6 7 | 4 |
| | 60 65 | 27 29 | 27 29 | 9 10 | 7 | 5 |
| | 70 | 32 | 32 | 10 | 8 | 5 |
| | 75 | 34 | 34 | 11 | 8 | 6 |
| | | | | | | |

| t. Weight | Dose | Rate | Micro | ng/mL | Macro | |
|-----------|------------|----------|-------------|----------------|-----------|----------|
| . Height | Dusc | . acc | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/m |
| kg | mcg/kg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min |
| | 10 | 5 | 5 | 2 | 1 | 1 |
| | 15 | 7 | 7 | 2 | 2 | 1 |
| | 20 | 10 | 10 | 3 | 2 | 2 |
| | 25 | 12 | 12 | 4 | 3 | 2 |
| | 30 | 14 | 14 | 5 | 4 | 2 |
| | 35 | 17 | 17 | 6 | 4 | 3 |
| 80 | 40 | 19 | 19 | 6 | 5 | 3 |
| 00 | 45 | 22 | 22 | 7 | 5 | 4 |
| | 50 | 24 | 24 | 8 | 6 | 4 |
| | 55 | 26 | 26 | 9 | 7 | 4 |
| | 60 | 29 | 29 | 10 | 7 | 5 |
| | 65 | 31 | 31 | 10 | 8 | 5 |
| | 70 | 34 | 34 | 11 | 8 | 6 |
| | 75 | 36 | 36 | 12 | 9 | 6 |
| | 10 | 5 | 5 | 2 | 1 | 1 |
| | 15 | 8 | 8 | 3 | 2 | 1 |
| | 20 | 10 | 10 | 3 | 3 | 2 |
| | 25 | 13 | 13 | 4 | 3 | 2 |
| | 30 | 15 | 15 | 5 | 4 | 3 |
| | 35 | 18 | 18 | 6 | 4 | 3 |
| 85 | 40 | 20 | 20 | 7 | 5 | 3 |
| | 45 | 23 | 23 | 8 | 6 | 4 |
| | 50 | 26 | 26 | 9 | 6 | 4 |
| | 55 | 28 | 28 | 9 | 7 | 5 |
| | 60 | 31 | 31 | 10 | 8 | 5 |
| | 65 | 33 | 33 | 11 | 8 | 6 |
| | 70 | 36 | 36 | 12 | 9 | 6 |
| | 75 | 38 | 38 | 13 | 10 | 6 |
| | 10 | 5 | 5 | 2 | 1 | 1 |
| | 15 | 8 | 8 | 3 | 2 | 1 |
| | 20 | 11 | 11 | 4 | 3 | 2 |
| | 25 | 14 | 14 | 5 | 3 | 2 |
| | 30 | 16 | 16 | 5 | 4 | 3 |
| | 35 | 19 | 19 | 6 | 5 | 3 |
| 90 | 40 45 | 22 24 | 22 24 | 7 | 5 | 4 |
| | 45 50 | 24 | 24 | 8 | 6 | 4 |
| | | _ | | - | | |
| | 55 60 | 30 32 | 30 32 | 10 11 | 7 | 5 |
| | 65 | 35 | 32 | 11 | 9 | 6 |
| | 70 | 38 | 35 | 12 | 9 | 6 |
| | 70 | 41 | 41 | | | 7 |
| | 10 | | 41 6 | 14 | 10 | / |
| | 10 | 6 9 | 9 | 3 | 2 | 1 |
| | 20 | 11 | 11 | 4 | 3 | 2 |
| | 25 | 14 | 14 | 5 | 4 | 2 |
| | 30 | 17 | 17 | 6 | 4 | 3 |
| | 35 | 20 | 20 | 7 | 5 | 3 |
| | 40 | 23 | 23 | 8 | 6 | 4 |
| 95 | 45 | 26 | 26 | 9 | 6 | 4 |
| | 50 | 29 | 29 | 10 | 7 | 5 |
| | 55 | 31 | 31 | 10 | 8 | 5 |
| | 60 | 34 | 34 | 11 | 9 | 6 |
| | 65 | 37 | 37 | 12 | 9 | 6 |
| | 70 | 40 | 40 | 13 | 10 | 7 |
| | 75 | 43 | 43 | 14 | 11 | 7 |
| | 10 | 6 | 6 | 2 | 2 | 1 |
| | 15 | 9 | 9 | 3 | 2 | 2 |
| | 20 | 12 | 12 | 4 | 3 | 2 |
| | 25 | 15 | 15 | 5 | 4 | 3 |
| | 30 | 18 | 18 | 6 | 5 | 3 |
| | 35 | 21 | 21 | 7 | 5 | 4 |
| | 40 | 24 | 24 | 8 | 6 | 4 |
| .00 | 45 | 27 | 27 | 9 | 7 | 5 |
| | 50 | 30 | 30 | 10 | 8 | 5 |
| | 55 | 33 | 33 | 11 | 8 | 6 |
| | 60 | 36 | 36 | 12 | 9 | 6 |
| | 65 | 39 | 39 | 13 | 10 | 7 |
| | 70 | 42 | 42 | 14 | 11 | 7 |
| | 75 | 45 | 45 | 15 | 11 | 8 |
| | 10 | 6 | 6 | 2 | 2 | 1 |
| | 15 | 9 | 9 | 3 | 2 | 2 |
| | 20 | 13 | 13 | 4 | 3 | 2 |
| | 25 | 16 | 16 | 5 | 4 | 3 |
| | 30 | 19 | 19 | 6 | 5 | 3 |
| | 35 | 22 | 22 | 7 | 6 | 4 |
| 05 | 40 | 25 | 25 | 8 | 6 | 4 |
| .05 | 45 | 28 | 28 | 9 | 7 | 5 |
| | 50 | 32 | 32 | 11 | 8 | 5 |
| | 55 | 35 | 35 | 12 | 9 | 6 |
| | 60 | 38 | 38 | 13 | 9 | 6 |
| | 65 | 41 | 41 | 14 | 10 | 7 |
| | 70 | 44 | 44 | 15 | 11 | 7 |
| | 75 | 47 | 47 | 16 | 12 | 8 |
| | | | | e for this inf | | Ŭ |

ROCURONIUM

QC. Lactation? (Caution)

Trade Name: Zemuron

Class / Mechanism of Action

Blocks acetylcholine from binding to motor neuron receptors inhibiting depolarization.

Onset of action IV: 1-2 minutes (can be fast with higher doses and/or faster pushes), Duration:

approximately 25-40 minutes (increases with higher doses)

Indications

Labeled Indications: Rapid Sequence Intubation/Paralysis and routine endotracheal intubation, facilitates mechanical ventilation in ICU patients

Contraindications

• Hypersensitivity (e.g., anaphylaxis) to rocuronium, other neuromuscular-blocking agents, or any component of the formulation

- Resistance may occur in burn patients (>30% of body) for period of 5-70 days after injury
- High potential for interactions: Numerous drugs either antagonize (e.g., acetylcholinesterase inhibitors) or potentiate (e.g., calcium channel blockers, certain antimicrobials, inhalation anesthetics, lithium, magnesium salts, procainamide, and quinidine) the effects of neuromuscular blockade; use with caution in patients receiving these agents.
- Provides NO analgesia or sedation!
 - Must provide appropriate sedation and analgesia prior to paralytic use and throughout maintenance.

| Dose and Administration: ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|---|---|
| RSI: IV Push: | RSI: IV: |
| • 1mg/kg (Dosing ranges from 0.6-1.2 mg/kg) | • 1mg/kg (Dosing ranges from 0.6 - 1.2 mg/kg .) |
| Note: In adult patients with morbid obesity (BMI >40 kg/m2), use dose of 1.2 mg/kg using ideal body weight (IBW) | <u>Maintenance bolus dosing: (unlabeled and unreferenced dose)</u> IV Push: |
| Maintenance dosing: (unlabeled and unreferenced dose) IV Push: | • 1 mg/kg every 30-45 minutes (Dosing ranges from 0.6 - 1.2 mg/kg .) |
| 1 mg/kg IV/IO q30-45min PRN or 8-12 mcg/kg/min IV/IO (Dosing ranges from 0.6-1.2 mg/kg) | |

SUCCINYLCHOLINE

QC. Lactation?(Caution)

Trade Name: Anectine

Class / Mechanism of Action

Depolarizing Neuromuscular Blocking Agent (Paralytic)

Acts like acetylcholine, produces myoneural depolarization causing sustained flaccid skeletal muscle paralysis. Onset of action IV: 30-60 seconds, Duration 5-9 minutes with single dose

Indications

Labeled Indications: Rapid Sequence Intubation and routine endotracheal intubation

Contraindications

- Hypersensitivity to succinylcholine or any component of the formulation
- Acute phase of injury following major burns, multiple trauma (greater than 5 days after injury)
- Myopathies associated with elevated serum creatine phosphokinase and myasthenia gravis
- DO NOT USE IN PATIENTS WITH BURNS, CRUSH INJURIES, OR HYPERKALEMIA
- Re-Dosing is not advised due to increased risk of Hyperkalemia
- Neuromuscular disease (Muscular dystrophy, Spinal Muscular Atrophy, etc.)

- May cause Bradycardia, Malignant hyperthermia, and increased intraocular pressure
- Severe hyperkalemia can develop in cases of chronic abdominal infection, burn injury, children with skeletal muscle myopathy, subarachnoid hemorrhage, or conditions which cause degeneration of the nervous system commonly greater than 5 days old. Potassium increase of 0.5 mEq/L is expected with use.
- Provides NO analgesia or sedation!
 - Must provide appropriate sedation and analgesia prior to paralytic use and throughout maintenance.

| Dose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW Tape |
|--|----------|---|
| RSI/ Neuromuscular blockade: IV: IV: • 1-1.5 mg/kg Note: Pretreatment with 10% dosag depolarizing agents prior to neurometed blockade with Succinylcholine is NO ADVISED | uscular- | RSII / Neuromuscular blockade: IV: IV: • <10kg: |

| THIAMINE | QA, LactationYes (Caution) | | Trade Name: Vitamin B1 | |
|--|-----------------------------------|--------------------------|--------------------------------|--|
| Class / Mechanism of Action | | | | |
| Vitamin, water soluble Essential coenzyme in carbo | hydrate metabolism. On | set of action IV/IM: Rap | vid | |
| Indications | | | | |
| Labeled Indications: Treatment of thiamine deficiency including beriberi, Wernicke's encephalopathy, Korsakoff's syndrome, neuritis associated with pregnancy, or in alcoholic patients | | | | |
| Contraindications | | | | |
| Hypersensitivity to thiam | ine or any component of | the formulation | | |
| Adverse Reactions / Preca | utions | | | |
| Administration of dextrose may worsen acute symptoms of thiamine deficiency; use caution when low thiamine is suspect | | | | |
| Dose and Administration: | ADULT | PEDIATRIC | Always Reference BROSELOW Tape | |
| AMS; Seizure; Syncope; M and Diarrhea; w/ Hx of ETC IM/IV: • 100mg/day | | | igns of Malnutrition: | |

TRANEXAMIC ACID (TXA)

♀B, **Lactation: Yes (**Caution)

TradeName: Cyklokapron/Lysteda

Class / Mechanism of Action

Antifibrinolytic Agent, Hemostatic Agent

Displaces plasminogen from fibrin resulting in inhibition of fibrinolysis and inhibits the proteolytic activity of plasmin

Indications:

- Trauma-associated hemorrhage: Casualty likely needing blood transfusion (hemorrhagic shock, elevated lactate, one or more major amputations, penetrating torso trauma, or evidence of severe bleeding)
- Post-Operative Hemorrhage by dissection, enteric staples or suspected internal bleeding
- Signs or symptoms of moderate or severe TBI or altered mental status associated with trauma
- Postpartum Hemorrhage (continued bleeding despite Oxytocin and fundal massage)

Contraindications

- TXA is contraindicated in trauma if dose is not given within first 3 hours following Traumatic event (Ideal dosing timeframe is as soon as possible)
- Hypersensitivity to tranexamic acid
- Non-traumatic subarachnoid hemorrhage
- Thromboembolic disease (Cerebral Thrombosis, DVT, PE)

- Disseminated intravascular coagulation (DIC): Use with extreme caution in patients with DIC requiring antifibrinolytic therapy; patients should be under strict supervision of a physician experienced in treating this disorder. TXA should be used in Pt.'s with trauma related DIC however.
- Thrombosis (especially when given after 3hr from injury)
- Seizure

| Dose and Administration: AI | DULT | PEDIATRIC Always Reference BROSELOW Tape | |
|---|-------------------|---|--|
| Trauma-associated hemorrhage (unlabeled use): IV: Initial Dose: 2 grams of TXA in 100 cc NS or LR via IV/IO Bolus, or 2 gram IV/IO push (1 gram over 1 minute per push) but NOT later than 3 hours after injury. | | Trauma-associated hemorrhage (unlabeled use): IV: Initial Dose: 15mg/kg via IV/IO Bolus (goal within 1 minute), | |
| If patient received 1 gram of TXA prior <3hrs from time of injury: 1-gram TXA push over 1 minute or mixed in 100cc N Bolus. If >3hr from time of injury: <u>DO N</u> administer TXA. | IV/IO NS or LR | | |
| Suspected Post-Operative Hemorrhage dissection, enteric staples or suspected internal bleeding: | | | |
| Initial Dose: 2 grams of TXA in 100 cc LR via IV/IO Bolus or 2 gram IV/IO pus gram over 1 minute push) but NOT late hours after start of suspected hemorrha | h (1 er than 3 | | |

VASOPRESSIN

QC, Lactation?(Caution)

Trade Name: Vasostrict

Class / Mechanism of Action

Antidiuretic Hormone Analog-Vasopressor

Vasopressin, at therapeutic doses used for vasodilatory shock, stimulates the AVPR1a (or V1) receptor and increases systemic vascular resistance and mean arterial blood pressure; in response to these effects, a decrease in heart rate and cardiac output may be seen. Onset of action IV: Rapid with peak effect occurring within 15 minutes of initiation of continuous IV infusion. Duration: Within 20 minutes after IV infusion terminated. Indications **Labeled Indications:** Treatment of hypotension, vascular failure in shock Contraindications Hypersensitivity to Vasopressin or any component of the formulation • Use with caution in patients with asthma, cardiovascular disease, renal disease, or a history of seizure disorder **Adverse Reactions / Precautions** No applicable use in hemorrhagic shock unless fluid replacement therapy maximized! Maximize use of Blood products / Crystalloids before considering use in hemorrhagic shock. Assure adequate circulatory volume to minimize need for vasoconstrictors. Monitor BP • closely, avoid hypertension and adjust infusion rate as needed. Vesicant: Avoid extravasation, will cause tissue damage/necrosis, ensure proper needle • placement Cardiac arrhythmias are possible, monitor with 12 lead EKG **Dose and Administration:** ADULT **PEDIATRIC** Always Reference BROSELOW Tape Hypotension / Shock: Hypotension / Shock: Limited data available; efficacy results have Vasopressors should be used if patient is varied. hypotensive after fluid resuscitation to maintain mean arterial pressure (MAP) ≥65 mmHg. IV Infusion: Use in addition to norepinephrine for raising MAP 0.17 to 8 milliunits/kg/minute (0.01 to to target or to decrease norepinephrine dosage. 0.48 units/kg/hour)

Titrate to lowest effective dose.

IV Infusion: **4 Unit bolus IV/IO** followed by **0.04 U/min** infusion to maintain MAP>65 mmHg

VECURONIUM

QC, Lactation? (Caution)

Class / Mechanism of Action Nondepolarizing Neuromuscular Blocking Agent (Paralytic) Blocks acetylcholine from binding to motor neuron receptors inhibiting depolarization. Onset of action IV: 1.5-3 minutes, Duration: approximately 30-60 minutes Indications Labeled Indications: Endotracheal intubation, facilitates mechanical ventilation in ICU patients Contraindications Hypersensitivity to vecuronium or any component of the formulation **Adverse Reactions / Precautions** Resistance may occur in burn patients (>30% of body) for period of 5-70 days after injury High potential for interactions: Numerous drugs either antagonize (e.g., acetylcholinesterase • inhibitors) or potentiate (e.g., calcium channel blockers, certain antimicrobials, inhalation anesthetics, lithium, magnesium salts, procainamide, and guinidine) the effects of neuromuscular blockade; use with caution in patients receiving these agents. **Provides NO analgesia or sedation!** • Must provide appropriate sedation and analgesia prior to paralytic use and 0 throughout maintenance. Dose and Administration: ADULT PEDIATRIC Always Reference **BROSELOW** Tape RSI) and maintenance of paralysis: RSI and maintenance of paralysis: IV Push: IV Push: Induction: 0.1 mg/kg Dose range (0.08-Induction: 0.1-0.15 mg/kg • 0.15 mg/kg) Intermittent bolus dosing: 0.1 mg/kg every • Maintenance: 0.1 mg/kg Dose range 30-60 minutes PRN (0.08-0.15 mg/kg) every 30-60 minutes IV Continuous infusion: PRN 1-2.5 mcg/kg/minute IV Continuous infusion: 1 mcg/kg/min and titrate to 2:4 train of four (TOF) if stimulation devise is available. Note: Paralytic use and management: If available, utilize the train of four stimulation device with either the temple or radial/ulnar nerve placement. Maintain paralysis at a level of 2/4 twitches with TOF stimulation.

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WE WILL NEVER FORGET



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