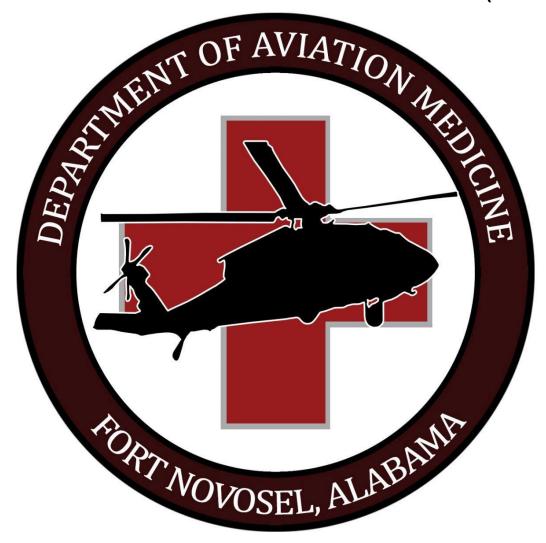
U.S. ARMY AEROMEDICAL EVACUATION STANDARD MEDICAL OPERATING GUIDELINES (SMOG)



CY24 Version

Published 01 February 2024 Revised 19 July 2024

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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 b | below: |
| <u>Unit Trainer Review:</u> | |
| This document has been reviewed by the below | v noted individuals for correctness and mission applicability: |
| Unit Standardization Officer/NCO Signature: | Date: |
| Unit Training NCO Signature: | Date: |
| Authorization: | |
| The Standard Medical Operating Guideline a 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-aviation-medicine

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

https://jts.health.mil/assets/docs/forms/DA4700 OP5 JTS TACEVACAAR&PC R.pdf

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

UNIVERSAL PATIENT CARE

Patient History

- Age of the Patient
- Chief Complaint
- Timing of Event / Event Factors
- Other Symptoms or Complaints
- Patient's Past Medical History
- Other Pertinent SAMPLE, OPQRST Questions

Key Concepts

- Use MARCHES for Trauma Patients
 - Massive Bleeding Control
 - Airway
 - Respiratory
 - Circulation
 - Hypothermia Care
 - Eye Injuries
 - Spinal Motion Restriction
- Focused Primary Exam for Non-Traumatic Illness/Injury

Treatment/Actions

- Scene Safety
 - Maintain situational awareness
 - Utilize appropriate PPE
- Initial Assessment
 - o Treat obvious and emergent life threats
 - MARCHES or Focused Primary Exam
 - Utilize BLS, ALS, and/or PALS guides as necessary
- Consider Spinal Immobilization
 - o 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
 - o 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

- 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)
 - 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

TACTICAL EVACUATION

Ground "Pick-Up" Phase

- Attempt to gain info prior to landing
- Ensure 360-degree scene security
- Collect medical information and patient documentation
- 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

Ground "Pick-Up" Phase

- Goal on ground time < 5min prior to wheels up
- If the tactical situation permits, all known preventable causes of death should be addressed prior to casualty movement
- If military working dos (MWD) are present (injured or uninjured), subdue/muzzle MWD first, then treat all human casualties before treating injured MWDs

"In-Flight" Phase

- Triage casualties as required: reassess patients and interventions
- Hemorrhage Control
 - o Check/add tourniquet, pack/dress wound, pressure dressing, hemostatic dressing
 - Initiate blood (DCR)
- Airway/Vent Management
 - o 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

- 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 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:
 - o 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%
 - Start Supplemental O2
 - Place NPA/OPA prn if no contraindications
 - Recheck q5 minutes
 - 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:
 - o Reassess Interventions
 - Restart Protocol
 - Consider other causes
- Continuous monitoring of ETCO2, SPO2, and ventilatory waveforms and pressures
 - o 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
 - Intubation contraindicated due to anatomical abnormalities or major airway trauma
 - 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 SEQUENCE INTUBATION

History

- 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.

Contraindications

- High likelihood of failure (Distorted Anatomy)
- Penetrating neck trauma

Procedure

 Make a plan, prepare patient and equipment (See PRE-INTUBATION CHECLIST) Conduct seven "P" pneumonic (7Ps):

Medications

- Induction Agents:
 - o Ketamine 1-2 mg/kg IV
 - o Etomidate 0.2-0.4 mg/kg IV
 - o 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
- Maintenance:
 - Ketamine 0.5-2 mg/kg IVP or 0.5-2 mg/kg bolus then 1-3mg/kg/hr.
 - o 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)
 - o Reduce Vt by 1mL/kg at intervals ≤ 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 | 0.4 | 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 | 22 | 22 | 22 | 24 |

Oxygenation Goal: Normal PaO2 80-100 mmHg or SpO2 90-96%; ARDS 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.
- 2. 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

Treatment

- 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
 - o Plasma, RBCs, Platelets in a 1:1:1 Ratio (no particular order)
 - Plasma and RBCs in 1:1 Ratio
 - o 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

Treatment

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

- o Epinephrine 0.3mg IM (0.3mL of 1:1000)
- o 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.
 - o Keep the intravenous line open with saline.
 - o Identify and treat cause of the reaction.
 - o 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
 - o Catheter > 18ga
 - Two failed attempts or > 90 secs proceed
- 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-1TM
 - EZ T.A.L.O.NTM
 - EZ-IOTM
- 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-IO[™]
 - Yellow 45mm for humerus and *heavy sternal
 - Blue 25mm for adult *sternum/tib
 - o 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

- o GAIN VASCULAR ACCESS where available based upon patient.
- 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.
- 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-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

Treatment

- Blunt Abdominal/Pelvic Injury
 - Serial Physical Exams/Reassessment
 - o Pelvic Binder
 - Conduct FAST exam if possible*
 - o Focus on resuscitation
- Penetrating Abdominal/Pelvic Injury
 - Hemostatic Dressing/ Pack Pelvic Cavity
 - o 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
 - o 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

- How long ago was the injury?
- Any signs of airway involvement?
- How big/small of a space was the patient in during the incident? (inhalation/carbon monoxide)
- Are there other traumatic injuries associated?
- Any spinal immobilization needed? (fall from a significant height, blast, etc.)

Call the Burn Center: DSN 312-429-2876 (429-BURN)/Comm: 210-916-2876 or 210-222-287

STOP the burning process/remove patient from electrical source. Ensure your safety first!

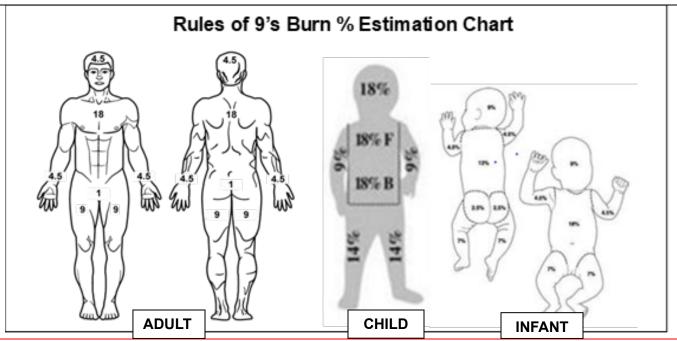
Differential Diagnosis: cardiac arrest, environmental exposure, seizure, burns (chemical, electrical, thermal, radiation), multiple trauma, carbon monoxide toxicity.

Treatment

- Access for additional injuries and treat life threats (DO NOT overlook TRAUMA), IV/IO, O2, and monitor (electrical injuries must have 12 lead EKG completed to access for Arrythmia's)
- **Electrical:** If arrythmia is present, go to appropriate Cardiac protocol
- Remove any constricting items i.e., rings and bracelets
- Assess airway, if suspected airway involvement move to Airway Guideline.
 - Indications for endotracheal intubation include comatose patient, symptomatic inhalation injury, deep facial burns, and burns over 40% Total Body Surface Area (TBSA). Requires large ETT size 8 adult.
- Thermal/Electric burn: If able, remove burning/charred clothing and cover with dry, sterile sheets/dressings.
- Chemical burn: Brush off dry chemicals, cut off 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.
 - o Alkali Burns to eye- 1-2 L of NS each eye for 30 minutes.
- Determine/start fluid replacement for burn fluid resuscitation.
- Manage pain and prevent hypothermia. KEEP WARM!
- Monitor urinary output, if able

- Urinary output is the MOST reliable guide for adequate resuscitation:
- Adult 30-50 mL/hr (75-100ml/hr electrical burn) Children: <40 kg 0.5-1ml/kg/hr.
- Edema after burn injury causes most Supraglottic airway devices such as laryngeal mask airways (LMAs) to be inadequate.
- Ventricular fibrillation (in AC) and asystole (in DC) are the most common dysrhythmias seen with electrical shock.
- MASCAL where lightning is involved reverse triage should be performed. Those victims in full arrest should be resuscitated first.
 - Specifically, if there are no spontaneous respirations after airway maneuver, but no other signs of non-survivable injury, administer ventilatory support aggressively as resources allow

BURN FLUID RESUCITATION



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 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

Treatment

- Penetrating Chest Wound
 - o Open
 - Seal open wound with occlusive chest seal (vented)
 - Impalement
 - Stabilize
 - High index of concern for Hemo-pneumothorax
 - o Signs of Hemo-pneumothorax
 - Needle Thoracostomy

Goal: SPO2>90%; Improved RR; Equal rise/fall

- Blunt Chest Trauma
 - 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
 - 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

History

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

History (Complications)

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

Treatment

- 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).
 - o 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

- Pain/Swelling
- Deformity
- Altered Sensation/Function
- Diminished Pulse/ Cap refill
- Decreased Temperature
- Bleeding
- Amputation

Signs and Symptoms (cont.)

- Abrasion
- Contusion
- Multi-Trauma
- Fracture
- Dislocation
- Laceration
- 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
 - o Follow Heavy Active Bleed Guidelines
 - o Wrap Amputation in Sterile Dressing with Normal Saline
 - o Place in sealed container with ice slurry if available
 - Transport with patient
 - o Clean amputated limb
- Convert Limb/Junctional TQ as soon as possible if:
 - No presence of shock
 - Able to monitor wound closely for bleeding
 - Not placed to control hemorrhage on amputated extremity
 - o Every effort should be made to convert in less than 2 hours if patient is not in shock
- Wound Care/Protection
 - Bandage/cover injuries
 - Immobilize extremity
 - o Ice (if available) for Edema

- After Bleeding Controlled:
 - Treat for signs and symptoms of Hypotension/Shock
- Follow DCR protocols regarding hierarchy of fluid administration
- Carefully evaluate and document Neurovascular Status in all fractures/dislocations
- Never attempt to reduce an open fracture unless you have a confirmed loss of pulse
- A pelvic binder is indicated in cases of severe lower extremity injury and may be utilized in conjunction with a traction splint.
- Blood loss can be severe and concealed in long bone fractures-especially the femur
- 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

EYE INJURY/PAIN

History

- Pain, Swelling, Blood
- Decreased Visual Acuity / Blindness
- Deformity / Contusion
- Foreign Body
- Excessive Tearing

History (Complications)

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

Treatment

• Without Known Injury

- o Evaluate Pupils
- Consider unrecognized chemical exposure.
 - 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

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

Traumatic Injury

- o 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.
- If penetrating, give moxifloxacin 400mg PO / IV.
- Refer to Pain control Guideline.
- Treat for Nausea / Anxiety
 - o 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 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

Treatment

- 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
 - 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.
 - o Do not resuscitate if injuries are incompatible with life.
- Address all known points of hemorrhage.
 - o Initiate transfusion with 1 unit of blood product (avoid resuscitation with crystalloid)
 - o 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
 - Pericardiocentesis
- Place monitor on patient: Prepare Defibrillator
 - o Determine Rhythm: Pulse return?
- ROSC not Achieved
 - Continue CPR
 - Continue Blood / IV Fluids
 - Reduce Long Bone Fractures
 - o Reduce Pelvic Fracture
 - Reassess known hemorrhage points
- ROSC Achieved
 - o 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

History

- Restlessness / Confusion
- Weakness / Dizziness
- Tachycardia
- Pale, Cool, Clammy Skin
- Delayed Cap. Refill
- Blooding
- Dehydration
- Congenital Heart Disease

History (Complications)

- Nausea / Vomiting
- Shock: Hypovolemic, Cardiogenic, Septic, Neurogenic, Anaphylactic
- Cardiac Arrhythmia
- Pulmonary Embolus
- Tension Pneumothorax
- Medication OD
- Vasovagal Episode

Treatment

- Shock Due to Hemorrhage / Trauma:
 - Control hemorrhage
 - Optimize homeostasis (See Notes, Warnings, Cautions))
 - Optimize hypothermia management
 - o 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
 - 2L or 30mL/kg IVF bolus PRN, additional crystalloid based on reassessment of clinical indication.
 - If inadequate, consider NOREPINEPHRINE 2-20 mcg/min IV/IO
 - o 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
 - o If inadequate, NOREPINEPHRINE 2-20mcg/min IV/IO

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

PEDIATRIC HYPOTENSION/SHOCK

History

- Restlessness / Confusion
- Weakness / Dizziness
- Tachycardia
- Pale, Cool, Clammy Skin
- Delayed Cap. Refill
- Blooding
- Dehydration
- Congenital Heart Disease

History (Complications)

- Nausea / Vomiting
- Shock: Hypovolemic, Cardiogenic, Septic, Neurogenic, Anaphylactic
- Cardiac Arrhythmia
- Pulmonary Embolus
- Tension Pneumothorax
- Medication OD
 - Vasovagal Episode

Treatment

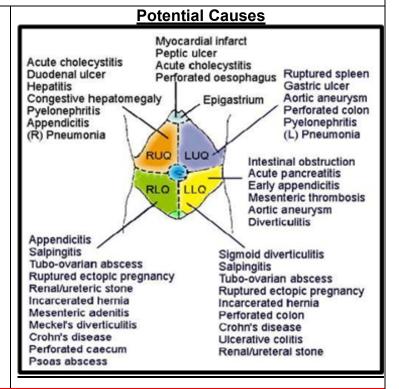
- Due to Hemorrhage / Trauma:
 - Trauma Fluid Preferences
 - Whole Blood (if available)
 - pRBC's and plasma (if available)
 - (LR/NS) 20mL/kg IVF Bolus
 - 10mL/kg Blood Product
 - If inadequate, consider (as last resort) EPINEPHRINE 1mcg/kg/min IV/IO or NOREPINEPHRINE
 .05- 0.1 mcg/kg/min slow IV push 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 Guideline:
 - Pediatric BRADYCARDIA W/ PULSE and poor perfusion
 - Pediatric TACHYCARDIA W/ PULSE and poor perfusion
 - Pediatric CARDIAC ARREST
 - Non-invasive PPV (BVM) vs. Advanced airway
 - o If rales heard on lung exam, 5 − 10mL/kg IVF over 5 − 10 min
 - If inadequate, consider EPINEPHRINE 1mcg/kg/min IV/IO

- 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.
- **Avoid Pressors** (last resort) as able, unless distributive or cardiogenic shock. Continue IVF for trauma; optimize homeostasis and correct volume.

ABDOMINAL PAIN

Signs and Symptoms

- Pain (RUQ, RLQ, LUQ, LLQ)
- Migration / Radiation)
- Tenderness
- Nausea / Vomiting
- Diarrhea (Bloody?)
- Dysuria
- Constipation
- Vaginal Bleeding / Discharge
- Distention
- Guarding / Rigidity
- Associated symptoms:
 - Fever, Headache, Weakness, Malaise / Fatigue, Myalgia, Cough, Mental Status Changes, Rash



Rule out MOI as a differential diagnosis. Universal Patient Care Guideline, O2 (if Hypoxemic), IV / IO Guideline, Cardiac Monitor12 Lead ECG (>40yo)

Trauma Considerations

- MOI- Blunt force or penetrating trauma
- Obstetric Patient trauma- follow O/B Emergency Guideline

Blood Products

- Persistent or Worsening Signs of Hypovolemic Shock (Tachycardia, Hypotension, Pulse Pressure)
- Rigid Distended Abdomen and/or Known: AAA, GI Bleed, or Ruptured Ectopic / Abruption
- Reassess unstable patients every 5 min;
 Stable patients every 15 min.

Medical Considerations

- Identify potential causes of abdominal pain (refer to chart above)
- Tachycardia / Hypotension / Orthostatic BP
 - Consider 500ml bolus per (IV/IO guideline)
- Significant Pain
 - o Pain management guideline
- Cardiac Emergencies
 - Cardiac pain guideline
- Nausea or Vomiting
 - o 500ml IVF Bolus
 - Promethazine 12.5-25mg IV
 - Ondansetron 4-8mg IV

- Maintain a high index of suspicion for ectopic pregnancy as a cause of abdominal pain in childbearing age.
- Antacids should be avoided in patients with renal disease.
- Patients older than 50 are at an increased risk for life-threatening diagnosis.
- Appendicitis presents with vague, periumbilical pain that migrates to the RLQ. This classic presentation may not be present in some patients.
- Repeat vital signs after each intervention. In non-traumatized patients, may repeat fluid bolus PRN
 depending on patient condition and vital signs. In trauma patients, fluid bolus should be used in
 accordance with hypotensive resuscitation guideline.
- Conservative approach with lower promethazine dosage for patients likely to experience sedative effects (e.g., elderly patients).

ALLERGIC REACTION

Signs and Symptoms

- Itching or Hives
- Cough / Wheeze / Resp. Distress
- Chest / Throat tightness
- Difficulty Swallowing
- Hypotension or Shock
- Edema
- Nausea / Vomiting

Differential Diagnosis

- Urticaria (rash only)
- Anaphylaxis (2 or more systems)
- Shock (other than anaphylactic)
- Angioedema
- Aspiration / Airway Obstruction
- Asthma or COPD
- Pulmonary Edema / CHF

Rule out MOI as a differential diagnosis. Universal Patient Care Guideline, O2 (if Hypoxemic), IV / IO Guideline, Cardiac Monitor (ASAP)

ADULT

Hives / Rash Only, No Respiratory Complaint

- Administer Diphenhydramine 25-50mg IV/IO/IM/PO
- Administer Methylprednisolone 125mg IV/IO

Shock / Unresponsive or Respiratory Distress / Failure

- Epinephrine-Pen or Epinephrine 1:1000;
 0.3-0.5mg IM
- o 500ml IVF if not previously started.
- Albuterol 90-180mcg 2 puffs MDI or 2.5-5mg via nebulizer
- Diphenhydramine 25-50mg IV/IO/IM/PO
- Methylprednisolone 125mg IV/IO

Worse or Unstable

 Epinephrine IV infusion at 5-15 mcg/minute (see Epinephrine infusion chart in drug card for procedure)

PEDIATRIC

Hives / Rash Only, No Respiratory Complaint

- Diphenhydramine; 1-2 mg/kg IM/IV/PO (max dose 50mg)
- Administer Methylprednisolone 2mg/kg
 IV

Shock / Unresponsive or Respiratory Distress / Failure

- Epi-Pen Jr for <30kg or Epinephrine1:1,000; 0.01mg/kg IM (max 0.3mg)
- o 20 mL/kg IVF Bolus
- Albuterol metered-dose inhaler
 - 4 8 inhalations every 20 min
- Albuterol by nebulizer
 - Children < 20 kg; 2.5 mg/dose
 - Children > 20 kg; 5 mg/dose
 - Diphenhydramine 25-50mg IV/IO/IM/PO
- Methylprednisolone 2mg/kg IV; maximum daily dose, 120 mg

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

ALTERED MENTAL STATUS

Signs and Symptoms

- Any signs of head trauma/injuries?
- Any AMS?
- Any pertinent medical conditions or medical history?
- Are there any bystanders that can provide information about the patient?
- Is this abnormal behavior?

Differential Diagnosis

- Head trauma/psychiatric disorders
- Thyroid dysfunction
- Hyper/hypoglycemia
- Diabetic ketoacidosis/toxic Ingestion
- Environment (hyper/hypothermia)
- Hypoxia

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?
 - o Refer to head injury guideline, airway guideline if applicable.
 - 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.
 - Lorazepam 2-4mg IV/IM (can use alone).
 - o 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
- o transfer (e.g., blast, motor vehicle)
- o Fall from >3 ft.
- o 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
- o Consider: Blood Product for AAA
- Suspicion of ACS/ Chest Pain
 - Chest Pain Guideline

Arrhythmia

- o Bradycardia with Pulse Guideline
- o Tachycardia with Pulse Guideline
- Cardiac Arrest Guideline (VF / Pulseless VT or Asystole / PEA)

- 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

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
 - o Ondansetron 4-8mg IV

Notes, Warnings, Cautions

- 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.

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 <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.
 - o Patients with NO AMS:
 - Administer oral glucose gel or equivalent until glucose level is >70mg/dL.

- 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

Potential Causes

- Asthma
- Anaphylaxis / Allergy
- Aspiration
- COPD
- Pleural Effusion
- Pneumonia
- Congestive Heart Failure / Cardiac
- Pulmonary Embolus
- Pneumothorax
- Pericardial Tamponade
- Hyperventilation
- Toxic Inhalation (e.g., Cyanide, CO)

<u>ADULT</u>

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 IV slow push.

Rales/CHF

- Monitor O2 and ETCO2
- Provide PPV/NIPPV (CPAP/BIPAP) with 100% oxygen support.
- Initiate IV/IO Access
- Administer nitroglycerin SL 0.4mg q5min if SBP>90.
- Failure to improve:
 - administer Furosemide 60-80mg IV slow push, place foley if possible.

PEDIATRIC

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)
 - o 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.

Rales/CHF

- Monitor O2 and ETCO2
- Provide PPV (if tolerated) with 100% oxygen support

OR

- 100% NRB if PPV is not tolerated.
- Failure to improve:
 - Administer Furosemide 1mg/kg IV slow push, place foley if possible.

- 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 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

Potential Causes

- Asthma
- Anaphylaxis / Allergy
- Aspiration
- COPD
- Pleural Effusion
- Pneumonia
- Congestive Heart Failure / Cardiac
- Pulmonary Embolus
- Pneumothorax
- Pericardial Tamponade
- Hyperventilation
- Toxic Inhalation (e.g. Cyanide, CO)

ADULT

- 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:
 - Consider Epinephrine
 - o 15- 30kg: 0.15mg IM (EPIPEN JR)
 - >30kg: 0.3mg IM (EPIPEN) OR
 - o 1:1000, 0.01mg.kg IM (max 0.3mg)
- Administer Methylprednisolone 1-2mg/kg IV.

- 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.

SEIZURE

Signs and Symptoms

- Decreased Mental Status
- Seizure Activity
- Somnolence
- Incontinence
- Evidence of Trauma
- Loss of Consciousness
- Oral Injuries (e.g., Tongue, Buccal)

Initial Treatment

- Provide oxygen support.
- Consider airway control (Definitive if necessary)
- Initiate IV/IO Access
- Place on cardiac monitor
- Monitor Blood Glucose

Adult Treatment

- Active Seizure
 - Administer Midazolam (5mg IV/IO/IN or 10mg IM)
 OR
 - Lorazepam 4mg IV/IM OR Diazepam 4mg IV/IM
 - May repeat anticonvulsants two times every 3-5 minutes if seizure has not stopped.
 - Levetiracetam (Keppra)1500mg IV over 15min; followed by maintenance dosing of 1000mg IV/IO every 12 hours.
 - Pregnancy Considerations:
 - Administer Magnesium Sulfate 4G IV over 15min. (Monitor for hypotension)
- Blood Glucose <70
 - Administer 50% Dextrose 25G IV OR
 - Glucagon 1mg IV/IM

Pediatric Treatment

- Active Seizure
 - o Administer Lorazepam 0.05-0.1 mg/kg IV (max 4mg) OR Midazolam 0.2mg/kg IM.
 - May repeat anticonvulsants two times every 3-5 minutes if seizure has not stopped. Must establish definitive airway control prior to subsequent doses.
 - o Levetiracetam (Keppra) 60mg/kg, single dose.
- Blood Glucose <40
 - Administer Thiamine 25mg IV/IM OR 25% Dextrose 2mL/kg IV OR
 - Glucagon 0.05mg/kg IM

- For seizure prophylaxis consider Levetiracetam (Keppra) 1500 mg IV over 15 min, after 12 hrs. consider maintenance dose of 1000 mg every 12 hrs.
- Status epilepticus defined as seizure >15min or two or more continuous seizures without a period of consciousness / recovery. This is a real emergency requiring rapid airway control, treatment, and transport to the nearest suitable medical treatment facility.
- Paralysis for airway control does not stop seizure activity only hides it. A seizure is a CNS electrical
 phenomenon and damage is still being done even when no muscular activity seen due to paralysis.
- In pregnant patients, Magnesium should be first line to abort non-epileptic seizures. Midazolam should only be used if this fails (pregnancy class D).
- Be prepared to assist with ventilations with the use of midazolam/lorazepam. If airway controlled and ventilating well – may give total of 4 doses of Midazolam for adults and 4 doses of Lorazepam for pediatrics.

SEPSIS/FEVER

History

- Wound(s) with signs of infection.
- Warm, Flush >100.4°F / 38°C
- Diaphoretic, Chills <96.8°F / 36° C
- Capillary refill time > 3 seconds
- Abnormal vital signs
 - o HR > 90
 - o BP < 90
 - o RR >20/min.
- Altered Mental Status
- Decreased Urine output
- Rash(s), Purpura
- Immunosuppressed

Definition

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

Treatment

- Initiate Monitoring: ECG, NIBP, SPO2, ETCO2, Temp
- Supplemental O2, Goal > 93% room air
- Check Glucose (<60 Hypoglycemia Protocol)
- 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?
 - o Administer Norepinephrine 2-12mcg/min IV.
 - Add Vasopressin 0.03 units/min SBP < 90 or MAP < 65.
 - o 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.

- 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.

TOXIC INGESTION

Poison Control Number (in US): 1-800-222-1222 Website: https://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 25g in 500ml NS or Glucagon 1mg/IM)
- Blood Sugar >60: Activated Charcoal 1g/kg PO (If alert and <1hr from ingestion)
- Beta Blocker Overdose: Glucagon 3-10mg IV/IM Bolus followed by 3-5mg/hr
- Opiates: Naloxone 0.4-2mg IV/IO Watch for Respiratory Depression
- TriCyclic Anti Depressant: 12 Lead
 - o QRS>100 or Hypotensive?
 - Sodium Bicarbonate 1mEg/kg
 - 100-150 mEq in 1L D5/NS @ 100-200ml/hr
- Organophosphate: Atropine 2mg IV/IO q5 + 2-PAM 600mg IV/IM
- Seizure Midazolam 2.5-5mg IV/IM

Pediatric Treatment

- Blood Sugar <65: AMS Guideline (25% Dex 2ml/kg IV or Glucagon 0.05mg/kg/IM)
- Blood Sugar >60: Activated Charcoal 1g/kg PO (If alert and <1hr from ingestion)
- Beta Blocker Overdose: Glucagon 1mg IV/IM
- Opiates: Naloxone 0.1mg/kg IV Pediatric Airway Guideline
- TriCyclic Anti Depressant: 12 Lead
 - QRS>100 or Hypotensive?
 - Sodium Bicarbonate 1mEq/kg
 - 100-150 mEq in 1L D5/NS @ 100-200ml/hr
- Organophosphate: Atropine 0.02mg/ IV/IO q5 + 2-PAM 25mg/kg IV/IM
- Seizure Lorazepam 0.1mg/kg IV

- Anticholinergic: Altered mental status, hyperthermia, mydriasis, flushing, anhidrosis, full bladder.
 - o Follow TriCylcic dosing
 - Lorazepam for agitation and seizures
- Beta Blockers watch for Hypoglycemia
- Calcium Channel Blockers watch for Hyperglycemia
- Cyclic Antidepressants signs: Hypotension, depressed mental status, respiratory depression, and cardiac arrythmias
- Opiod signs: Depressed mental status, pinpoint pupils, N/V, Respiratory depression, hypotension
- Organophosphate signs: Salivation, Lacrimation, Urination, Diarrhea, Emesis, Altered Mental Status

VOMITTING & DIARRHEA

Signs and Symptoms

- Pain
- Abdominal Distention
- Constipation
- Diarrhea
- Anorexia
- Fever
- Rash

Differential Diagnosis

- CNS Injury / Infection
- Myocardial Infection
- Drugs / Toxins
- Pregnancy
- Gastroenteritis
- Appendicitis
- Bowel Obstruction

Adult Treatment

- IV O2 Monitor
- Blood Glucose <60 w/ evidence of alcohol abuse?
 - o 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 Ondansetron 4-8mg IV

Pediatric Treatment

- IV O2 Monitor
- Blood Glucose <65 w/ evidence of malnourishment?
 - o Thiamine 100mg IV/IM
 - 25% Dextrose 2ml/kg IV or Glucagon 0.5mg IM
- Nausea / Vomiting:
 - o Promethazine (if >2yr old) 0.25mg/kg IV
 - o Ondansetron (<40kg) 0.1mg/kg IV
 - Ondansetron (>40kg) 4mg IV

- Suspicions of underlying conditions should prompt immediate referral to appropriate protocol
- In pregnant patients with N/V substitute D5 1/2NS or D5NS in place of NS
- Fluid of choice for vomiting is NS. Fluid of choice for diarrhea is LR
- Continually monitor for any decompensation

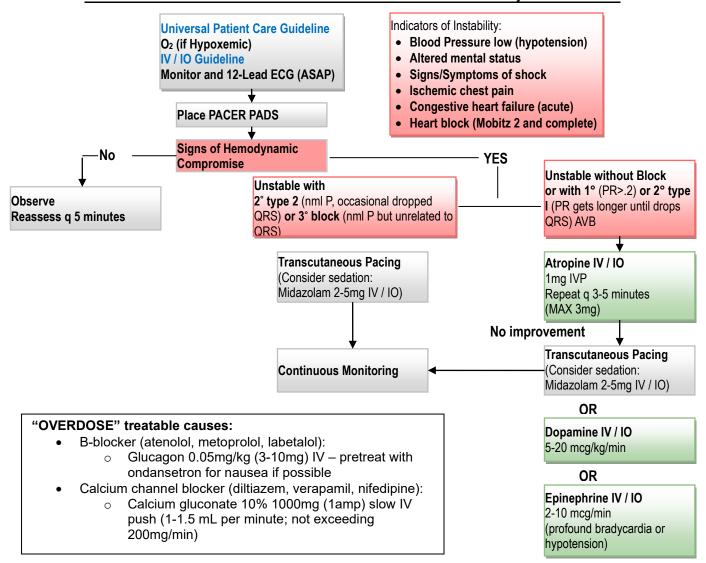
BRADYCARDIA with PULSE

Signs and Symptoms:

- HR <50bpm
- Chest Pain
- Respiratory Distress
- Hypotension / Shock
- Altered Mentation
- Syncope

Differential Diagnosis:

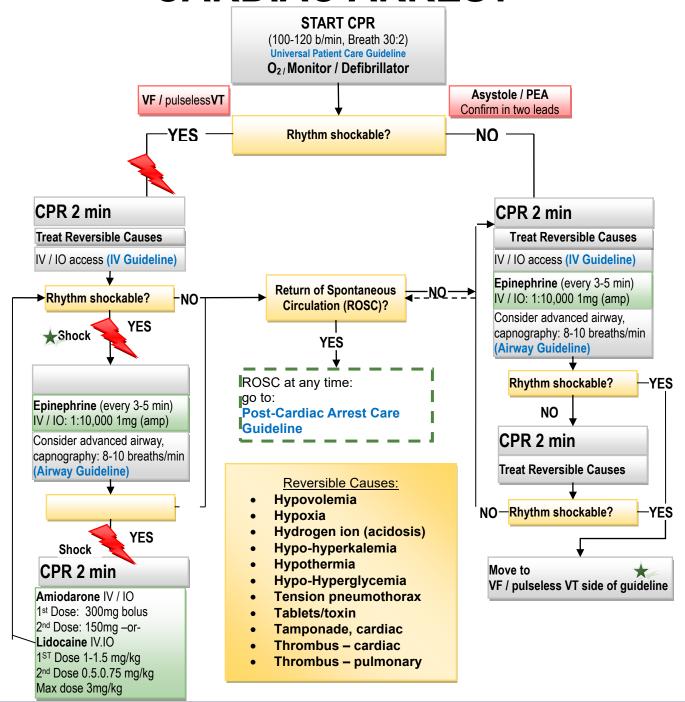
- Acute MI
- Hypoxia
- Hypothermia
- Sinus Bradycardia
- Physiologic Bradycardia (Athletes)
- Stroke
- Spinal Cord Lesion
- Toxin / Medications (B-blockers)
- AV Block / Sick Sinus Syndrome



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).

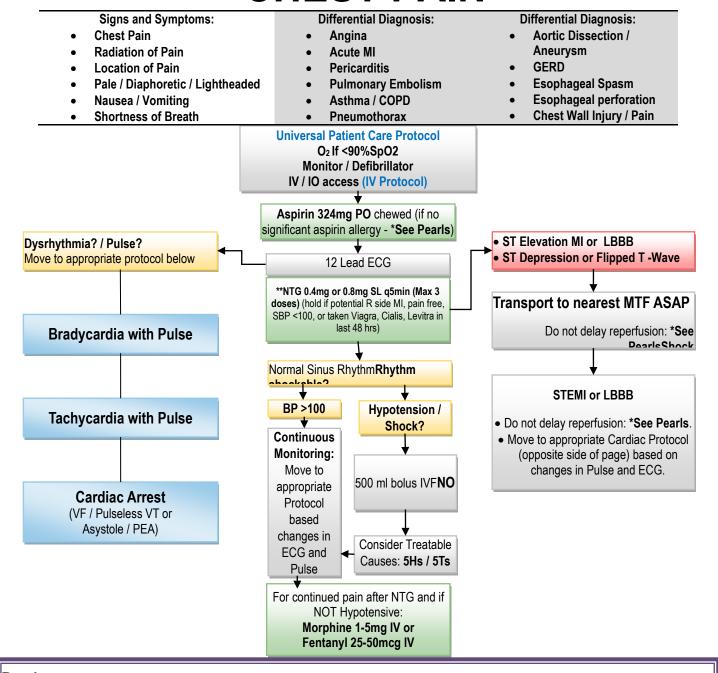
CARDIAC ARREST



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.
 - All patients should get a glucose check, at least 1L fluid bolus, and ultimately bilateral 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**

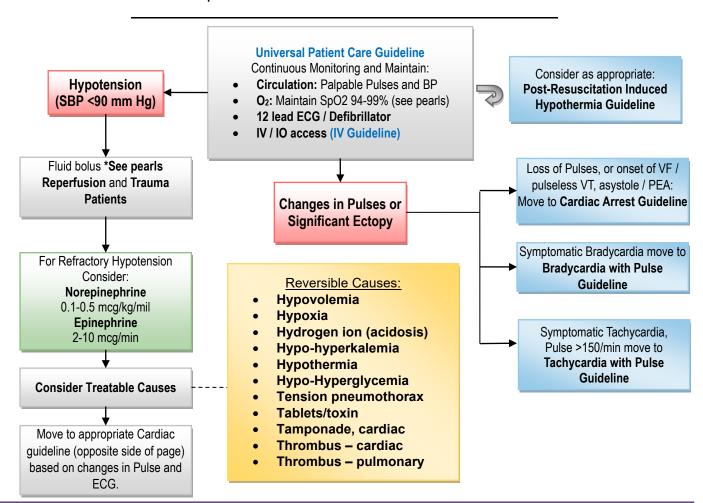
POST-CARDIAC ARREST CARE

Signs and Symptoms:

- Return of Spontaneous Circulation
- Pulse
- Respirations

Differential Diagnosis:

 Continually Address Primary Pathology Associated with Arrest



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

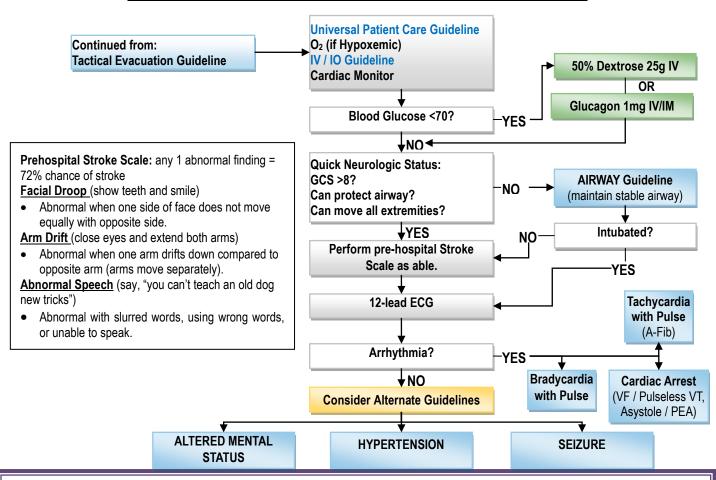
SUSPECTED STROKE / TIA

Signs and Symptoms:

- Altered Mental Status
- Weakness / Paralysis
- Blindness or Other Sensory Loss
- Aphasia / Dysarthria
- Syncope
- Vertigo / Dizziness
- Vomiting
- Headache
- Seizures

Differential Diagnosis:

- Transient Ischemic Attack
- Stroke
- Seizure
- Hypoglycemia
- CNS Infection / Mass
- Trauma
- Metabolic



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

Signs and Symptoms:

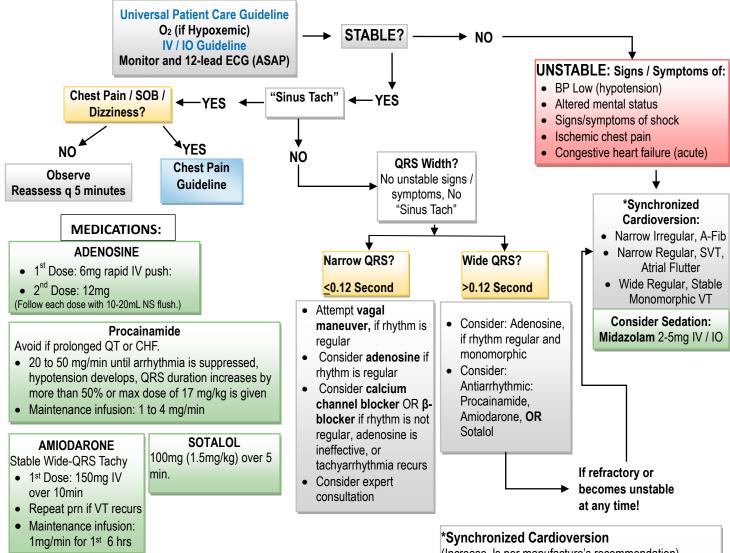
- Ventricular Tachycardia on EKG (rate typically >150/min)
- Conscious, Rapid Pulse
- Chest Pain / Shortness of Breath
- **Palpitations**
- **Dizziness**
- Anxiety

Differential Diagnosis (Wide Complex QRS >.12sec):

- Artifact / Device Failure
- Cardiac
- **Endocrine / Metabolic**
- Hyperkalemia
- Drugs
- **Pulmonary**

Differential Diagnosis (Narrow QRS):

- Wolf-Parkinson-White Syndrome
- Valvular Heart Disease
- Sick Sinus Syndrome
- **Myocardial Infarction**
- **Electrolyte Imbalance**
- Sinus Tachycardia / Atrial Flutter
- Hypoxia
- Drug Overdose / Toxin
- Hyperthyroidism



Pearls:

- **Torsades de Pointes may Benefit from early use of Magnesium: 1-2 grams IV over 60 min (Mix in 50ml D5W) Start drip of 0.5-1 gram/hr and titrate to effect.
- If hyperkalemia suspected (end-stage renal disease, dialysis) – administer Ca Chloride through central access or Ca Gluconate through peripheral IV.

(Increase Js per manufacture's recommendation)

Narrow Irregular, Atrial Fibrillation:

120-200 J biphasic

Narrow Regular, Other SVT, Atrial Flutter:

50-100 J. increase in stepwise fashion

Wide Regular, Stable Monomorphic VT:

100 J, increase in stepwise fashion

Wide Irregular:

- Defibrillate (NOT Synchronized)
- Go to: Cardiac Arrest Guideline

PEDIATRIC BRADYCARDIA with **Pulse and Poor Perfusion**

Typical HR/min

- 85 205 Newborn
- 3mth 2y/o 100 190
- 2y/o 10y/o 60 -140
- >10 y/o 60 - 100

Typical Sinus Tachycardia Rates

- Infants <220/min
- Children <180/min

Indicators of CARDIOPULMONARY COMPROMISE

- Hypotension
 - 1-10 y/o lower limit = 70+(years old x 2)mmHg
 - >10 y/o lower limit = 90mmHg
- **Acutely Altered Mental Status**
 - GCS <8, Weak Cry, Unusual Irritability, Altered Responsiveness, Lethargy, or Failure to Respond to **Painful Stimulus**
- Signs of Shock

Identify and Treat Underlying Cause! Rescue Breathing Ventilation Rate Without Advanced Airway: Continue: • **NEWBORN** = 40-60/min when performed without compressions **Universal Patient Care Guideline** • Infant / Child = 1 breath / 3 to 5 seconds Maintain Airway / Assisted Breathing Adult = 1 breath / 5 to 6 seconds • **O**₂ (Titrate to 94-99% SpO₂) CPR Rate of 100 Compressions / Min at: • IV / IO access (IV Guideline) • Monitor and 12-Lead ECG (ASAP) • One Rescuer = 30 Compressions and 2 Breaths Check Glucose Two Rescuer = 15 Compressions and 2 Breaths **CPR** Cardiopulmonary if HR <60/min Compromise YES with Poor Perfusion **Check Pulse every 2** Continues? despite O2 and Ventilation minutes during CPR **∮**NО Support ABCs Bradycardia • Continue O₂ Persists? **Continuous Monitoring** If Pulse is lost, GO TO: **PEDIATRIC Consider Consultation CARDIAC ARREST** Treatable causes: YES Check & Treat compromise in ABCs Hypoglycemia Consider: o D25 2mL/kg slow IV (max 25mL) Epinephrine 1:10,000 **Transcutaneous Pacing** o Glucagon 0.025mg/kg IM (max 1mg) 0.01mg/kg IV/IO g3-5min Tension Pneumothorax (Consider sedation: (Max single dose 1mg) "OVERDOSE (Mothers Milk)": Midazolam 0.05-0.1mg/kg IV / IO) **B-blocker** (atenolol, metoprolol, labetalol): Atropine Glucagon 0.05mg/kg (3-10mg) IV - pretreat with 0.02ma/ka IV / IO **Treat Underlying Causes** ondansetron (0.15mg/kg - max 2mg) for nausea if (Increased Vagal Tone or possible Primary AV Block) Calcium channel blocker (dilitiazem, verapamil, Support ABCs May Repeat Once after nifedipine) Continue O₂ 3-5 min Calcium chloride 10% 0.2ml/kg slow IV push **Continuous Monitoring**

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.

Naloxone 0.1mg/kg IV / IM (max 2mg) every 2-3

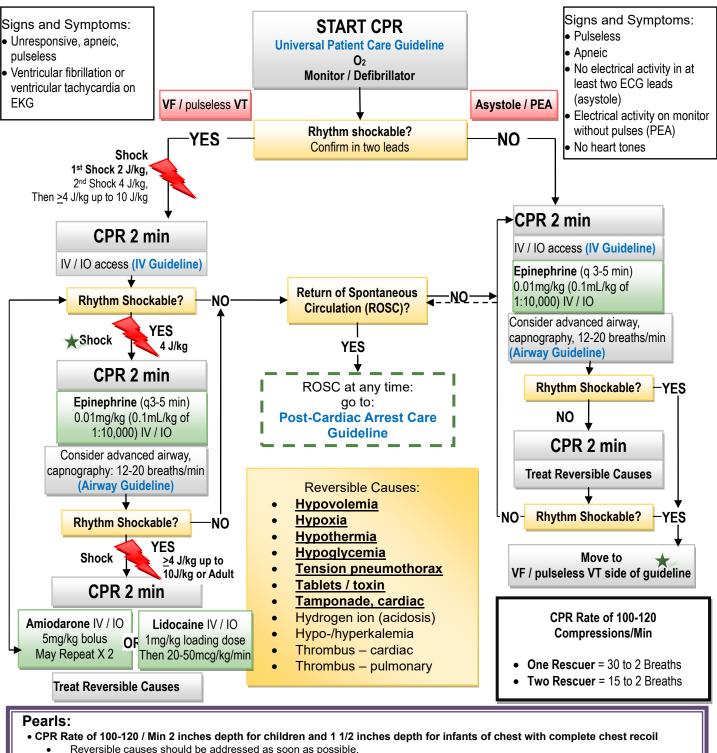
- In young, breast fed patients evaluate for mother's medications as they can cause toxicity in the infant.
- Pediatric pacer pads should be used if available. If only adult pads are obtainable they should be placed in the anterior-posterior position.

(Minimum dose 0.1mg

Max Single dose 0.5mg)

Consider Consultation

PEDIATRIC CARDIAC ARREST



- 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
 - All patients should get a glucose check, at least 20ml/kg fluid bolus of NS, and ultimately bilateral needle decompression (Trauma) before discontinuation of efforts

PEDIATRIC TACHYCARDIA with Pulse and Adequate Perfusion

Typical HR/min

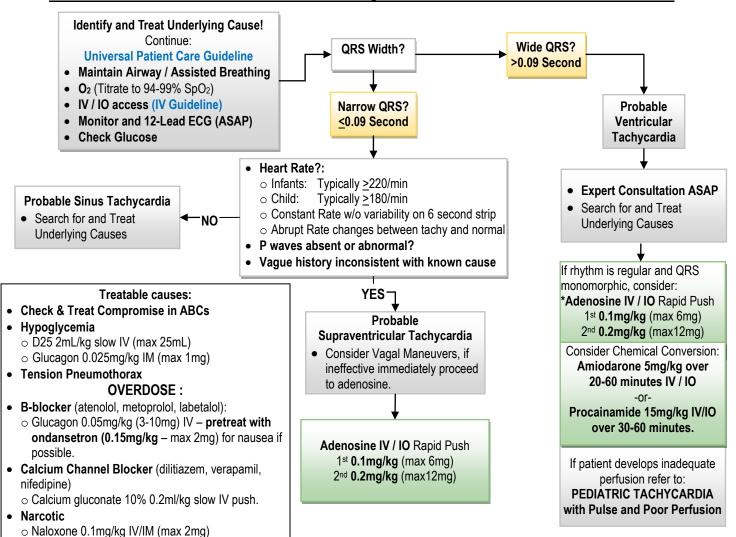
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- >10v/o 60 100

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- Hypotension
 - 1-10 y/o lower limit = 70+(years old x 2)mmHg
 - >10 y/o lower limit = 90mmHg
- Acutely Altered Mental Status
 - GCS <8, Weak Cry, Unusual Irritability, Altered Responsiveness, Lethargy, or Failure to Respond to Painful Stimulus
- Signs of Shock



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

Typical HR/min

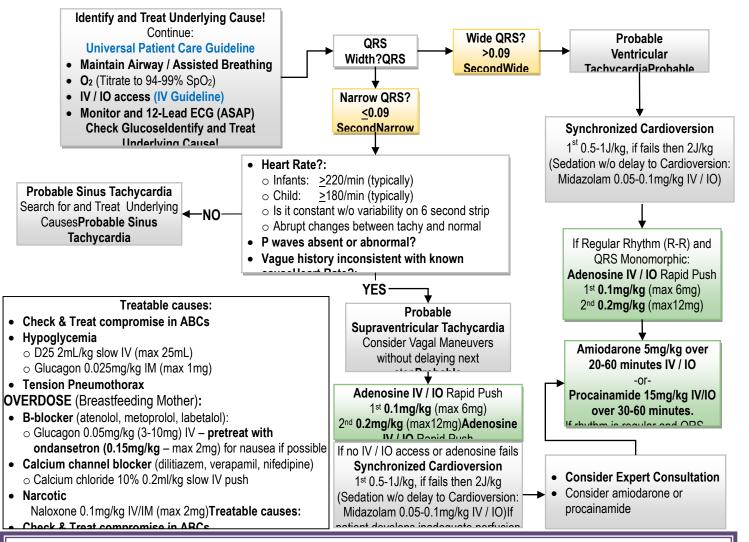
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 - > 10 y/o lower limit = 90mmHg
- Acutely Altered Mental Status
 - GCS <8, Weak Cry, Unusual Irritability, Altered Responsiveness, Lethargy, or Failure to respond to painful stimulus
- Signs of Shock



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.

TCCC MARCH

MASSIVE HEMORHORAGE

- HASTY tourniquets in the HOT ZONE
- Transition to DELIBERATE tourniquets during DECON in WARM ZONE

AIRWAY

- Assess excessive secretions indicate NERVE AGENT
- Defer most interventions consider risks in active HOT ZONE of remove mask to access airway

RESPIRATIONS

- Increased respirations consider ATNAA/CANA NERVE AGENT GUIDELINE
- Depressed respirations consider NAXOLONE INCAPACITATING AGENT GUIDELINE
- Other than antidotes respiratory interventions is best deferred to WARM ZONE

CIRCULATION

- Circulation intervention should be deferred to WARM ZONE
- Assess for shock
- IV/IO GUIDELINE
- HYPOTENSION/SHOCK GUIDELINE

MARCHE² Algorithm

CRESS Assessment

Consciousness:

Unconscious, Depressed Consciousness, Agitation

Respirations:

Present or Absent, Labored, Increased or Decreased

Eyes:

Constricted, Dilated, Normal

Secretions:

Dry, Normal, Increased

Skin:

Diaphoretic, Dry, Hot, Cyanosis

CRESS must be reassessed regularly, during zone transitions and at each transfer, to monitor for delayed onset of life threatening symptoms, and analyze antidote or countermeasure effectiveness

CBRN MARCHE²

Mask

 MASK or CHECK MASK SEAL as immediate HOT ZONE treatment

Antidote

- Utilize CRESS to differentiate chemical agent exposure RAPID IDENTIFICATION OF CHEMICAL WARFARE AGENT
- Antidotes are given in the Hot Zone if the casualty has symptoms of poisoning.
- These agents are rapid killers
 - Nerve agent (give ATNAA, CANA).
 - Cyanide (give hydroxocobalamin).
 - Pharmaceutical based sedating agent (naloxone)

Rapid Spot Decontamination

- Indicated for gross contamination on skin and/or wounds or if protective gear is breached
- Rapid exposure and decontamination of contaminated wounds is necessary lifesaving procedure in the HOT ZONE
- Apply RSDL, M100, M295, Sorbent, tech wipe, etc.

Countermeasures

- Appropriate therapy based on type of agent exposure, post initial antidote administration
- Deferred to WARM ZONE

Hypothermia Prevention & Head Injury

- Protect from lethal triad: HYPOTHERMIA, acidosis and coagulopathy through HOT/WARM/COLD ZONES
- Active warming or HPMK post decontamination and packaging for further evacuation
- Determine if altered mental status is due to chemical agent or trauma, if trauma HEAD INJURY/TBI GUIDLINE

Extricate and Evacuate

- EXTRICATE: egress patient from threat, agent contact, HOT ZONE
- Evacuate: to WARM ZONE Dirty CCP for decontamination
- COUNTERMEASURES and appropriate supportive care starts in WARM ZONE and continues during Evacuation/COLD ZONE

Notes, Warnings, Cautions

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

CHILDBIRTH

Signs and Symptoms

- Primi/Gravida/Para?
- Any pregnancy complications?
- Vaginal fluid/bleeding?
- Uterine contractions/back pain/stomach pain?
- Duration of contractions and time between
- Crowning/urge to push?
- Any complications expected with the newborn?

Possible Complications

- Preterm labor
- Placenta previa
- Prolapsed cord
- Abnormal presentations (i.e., breech)
- Spontaneous vaginal delivery (i.e. natural outcome)

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.

- 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) |
| Appearance (Skin color) | Blue, Pale | Body Pink, Extremities blue | Completely Pink |
| Respirations | Absent | Slow, irregular | Vigorous cry |

Treatment

- 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?
 - o 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?
 - 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).
 - o Consider hypoglycemia (D12.5 1/0ml/kg IV; dilute D50 to ½ 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

- Primi/Gravida/Para?
- Any pregnancy complications? (i.e. preeclampsia, gestational diabetes, etc.)
- Any abnormal bleeding?
- Uterine contractions/back pain/stomach pain?
- Blurry vision/dizziness?
- Changes in babies activity?

Differential Diagnosis

- Pre-eclampsia/eclampsia
- Placenta previa
- Abruptio placentae
- Spontaneous abortion
- Uterine rupture
- Ectopic pregnancy

Treatment

- 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
 - o 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.
 - o 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
 - o 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.

- 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:

Treatment

- 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

History

- 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

Treatment

- Inspect, wipe and/or suction mouth and pharynx.
- Ventilate with 100% oxygen.
 - o If unable, move to next step.
- Endotracheal Intubation (Size 9-11)
 - o If unable, move to next step.
- Suction Airway
 - o If airway still not clear, move to next step.
- Perform Tracheostomy
- Evaluate for pleural space and parenchymal problems.

- 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.



MWD ANALGESIA & SEDATION

Indications

- Trauma or pain
- · Need for chemical restraint
- 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

Treatment

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

- 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

- Cardiopulmonary Arrest Confirmed
- Causes:
 - Traumatic
 - Blast
 - Blunt force
 - Penetrating
 - Non-traumatic
 - Anesthesia
 - Near-drowning
 - Electrocution

Treatment

- Chest compressions (100 per min), over widest part of the chest
- · Clear airway and intubate ASAP.
 - o Perform tracheostomy if airway is obstructed.
- Manually ventilate (8-10 breaths per min) 100% O2
- Attain ECG
- Attain IV/IO access critical.
- VF or VT
 - o Defibrillate- 2-5 J/kg biphasic.
 - Immediate chest compressions x 1 cycle (2 min)
 - Defibrillate twice more, with 1 compression cycle between each defibrillation.
 - Drug therapy if defibrillation not successful
 - Epinephrine 0.01 mg/kg IV/IO or
 - Vasopressin 0.8 U/kg IV/IO once and Lidocaine 2 mg/kg IV/IO or
 - Amiodarone 5-10 mg/kg IV/IO
- Asystole/Bradycardia/PEA
 - Drug therapy
 - Atropine 0.04 mg/kg IV/IO (only if bradycardia preceded arrest)
 - Epinephrine 0.01 mg/kg IV/IO and Vasopressin 0.8 U/kg IV/IO once

- 70% of MWDs that arrest will have PEA, asystole, or sinus bradycardia as the initial arrest rhythm.
 Epinephrine or vasopressin are best choices for these rhythms or for empiric use if ECG capability is not available
- 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

History

- 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.

Treatment

- Initiate Monitoring: ECG, NIBP, SPO2, ETCO2
- Supplemental O2
- Attain (2) IV/IO sites FORELIMBS.
 - o **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
 - o Position self on left side, or lay dog on left side
 - o 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.
 - (DO NOT ATTEMPT THIRD if unsuccessful)
 - Apply gentle external pressure to abdominal wall to assist exiting air.
 - o Remove catheter once air is evacuated.
- Provide analgesia.

- 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 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 uncontrolled panting (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

Treatment

- 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
 - o 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.
 - O Give IV fluids 3-5 mL/kg/hr if not in shock
- Severe: Heat Stroke
 - 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

- 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.
- **DO NOT** 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).
- IV catheterization 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)
- IO catheterization access points are:
 - 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 4- medial tibia IO catheter location just distal to tuberosity



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
 - o 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

Clinical Signs

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

- 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.

| INJURY | PREFERRED AGENT | FREQUENCY | DURATION | |
|------------------------------|----------------------------|------------------------|------------------------------|--|
| | EXTREMET | | | |
| Skin, soft tissue, without | | | 041 | |
| open fractures | Cefazolin 2g | q 6-8hrs | 24 hours | |
| Skin, soft tissue, with open | | | 24 hours, then with each | |
| fractures, exposed bone, or | Cefazolin 2g | q 6-8hrs | subsequent I&D until soft | |
| open joints | | | tissue coverage | |
| | | WOUNDS | | |
| Penetrating chest injury | Cefazolin 2g | q 6-8hrs | 24 hours | |
| | ABDOMINA | L WOUNDS | | |
| Penetrating abdominal injury | | | | |
| with suspected/known | Cefazolin, 2g IV | | | |
| hollow viscus injury and | PLUS | q 6-8hrs | Stop 24 hours after control | |
| soilage; may apply to | metronidazole 500mg IV | q 8-12hrs | of contamination | |
| rectal/perineal injuries as | metrermudzete ecomig iv | | | |
| well | | | | |
| | MAXILLOFACIAL AN | ND NECK WOUNDS | | |
| Open maxillofacial | | | | |
| fractures, maxillofacial | Cefazolin 2g g | q 6-8hrs | 24 hours | |
| fractures with foreign body | 3 3 | • | | |
| or fixation device | CENTRAL NERVOUS | S SYSTEM WOLINDS | | |
| | Cefazolin, 2g IV | q 6-8hrs | | |
| Penetrating brain injury | PLUS (consider) | q 0-01113 | 5 days or until CSF leak is | |
| T chetrating brain injury | metronidazole 500 mg | q 8-12 hrs | closed, whichever is longer | |
| | Cefazolin, 2g IV | q 6-8hrs | | |
| Penetrating spinal cord | PLUS (consider) | 9 0 01113 | | |
| injury | metronidazole 500 mg | q 8-12 hrs | | |
| | EYE W | | | |
| | Erythromycin ophthalmic | | | |
| | ointment | q 6hrs | Until epithelium healed. | |
| Eye injury, burn or abrasion | Or | | No systemic treatment | |
| | Bacitracin ophthalmic | or PRN for symptomatic | required | |
| | ointment | relief | · | |
| | Levofloxacin 750 mg IV/PO | q 24 hrs | 7 days or until evaluated by | |
| Eye injury, penetrating | PLUS | • | 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 | X 1 dose | | |
| 1 O tolerable | dose. | 7(1 doso | Single dose therapy | |
| Not PO tolerable | Or | X 1 dose | Onlyie dose therapy | |
| 11011 0 101010010 | Ertapenem 1 g IV/ IM | 7(1 0000 | | |

COMMON LABORATORY VALUES

| LABORATORY | CHEM | CHEMISTRY CONVENTIONAL SI UNITS | | | | |
|-----------------|------|---------------------------------|----------------|------------------------|--|----------|
| Anion Gap | | 8-16 mEq/L | | 8-16 mEq/L 8-16 mmol/L | | 6 mmol/L |
| BUN | | 8-25 mg/100mL | 2.9- | 8/9 mmol/L | | |
| Calcium | 3 | 3.5-10.5 mg/100mL | 2.1- | 2.6 mmol/L | | |
| Carbon Dioxide | | 24-30 mEq/L | | 30 mmol/L | | |
| Creatine | Male | 0.2-0.5 mg/dL | Female | 0.3-0.9mg/dL | | |
| Creatine Kinase | Male | Male 17-40 U/L | | 10-79 U/L | | |
| Creatinine | | 0.6-1.5 mg/100L | 53-133 | | | |
| Glucose | | 70-110 mg/100mL | 3.9-5.6 mmol/L | | | |
| Sodium | | 135-145 mEq/L | | | | |
| Potassium | | 3.5-5.0 mEg/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

| CARDIAC MARKERS | | | | | | |
|------------------------|-------|-------------|--------|-----------------|--|--|
| Troponin I* | Onset | 4-6 hrs. | Peak | 12-24 hrs. | | |
| Troponin T* | Onset | 3-4 hrs. | Peak | 10-24 hrs. | | |
| Myoglobin | Male | 10-95 ng/ml | Female | 10-65 ng/ml | | |
| Myoglobin | Onset | 1-3 hrs. | Peak | Peak: 6-10 hrs. | | |
| INR only if Tx for DVT | | 0.8-1.2 | | 2.0-3.0 | | |

^{*}Troponin assays are becoming more analytically sensitive. Each device has different reference ranges associated. Correlate cTn with reference lab. Point of care readers are less sensitive.

Hemoglobin

Hematocrit

Platelets

| | NORMAL BLOOD GASSES | |
|-------------|---------------------|--|
| рН | 7.35-7.45 | |
| Pco2 | 35-45 mm Hg | |
| HCO3 | 22-26 mmol/L | |
| Base excess | (-2)-(+2) mEq/L | |
| CO2 | 19-24 mEq/L | |
| SaO2 | 96-100% | |

MEDICAL RECORD-SUPPLEMENTAL MEDICAL DATA For use of this form, see AR 40-66; the proponent agency is the Office of the Surgeon General REPORT TITLE JTS APPROVED (Date) Tactical Evacuation After Action Report & Patient Care Record, Page 1 (12 Jul 2018) -V4.1 Time Zone OL OZ Tail to Tail OY ON Leg# **9-Line**: Time Platform Dispatch Cat Assessed Cat Trauma MIST Report: M=Mechanism of Injury, I=Injury, S=Signs & Symptoms, T=Treatments / Disease Diagnosis: Comments Pickup: Time **Dropoff:** Time Location Capability EMT-B EMT-I EMT-P EMT-FPC RN CRNA PA MD/DO Other Circulation-Hemorrhage Control Time On ☐ CAT ☐ SOFTT ☐ Other RUE LUE RLE LLE ☐ Direct Pressure Tourniquet RUE LUE RLE LLE ☐ CAT ☐ SOFTT ☐ Other Prior TQ: Time On Hemostatic Dressing Reassess/tighten Time On ☐ CAT ☐ SOFTT ☐ Other RUE LUE RLE LLE Kerlix Dressing \bigcirc Y Time On ☐ CAT ☐ SOFTT Other ☐ RUE ☐ LUE ☐ RLE ☐ LLE \bigcirc N Pressure Dressing ☐ AAJT ☐ CRoC ☐ JETT ☐ SAM ☐ Other Junctional Time On \bigcirc N/A Other TQ Comments **Annotate Injuries** Airway Self NPA OPA Cric Trach ETT SGA Type (AMP)utation (BL)eeding Confirmed BS Vis ETCO2 (B)urn % TBSA O2 Source NC NRB BVM Vent LPM (C)repitus Intubated ☐ Prior to transport ☐ By transport crew Suction ☐ ETT ☐ Yaunker (D)eformity **Breathing** (DG)Degloving Needle Decompression Chest Equal Rise and Fall (E)cchymosis R L Mid-ax Mid-clav \bigcirc Y \bigcirc N \bigcirc N/A (FX)Fracture Time R L Mid-ax Mid-clav Respiratory Effort (GSW)Gunshot Wound Time ☐ L ☐ Mid-ax ☐ Mid-clav (H)ematoma ☐ Unlabored ☐ Labored (IMP)Impaled Object R L Mid-ax Mid-clav Time Assisted (LAC)eration \square R \square L Chest Tube (P)ain ETCO2 Vent Settings Time Mode Rate (PP)Peppering (PW)Puncture Wound Change (SQA)Subcutaneous Air Change (TBI)Suspect Other Change Circulation - Assessment **Circulation - Resuscitation** Rhythm / Ectopy Component ABO/RH Exp. Date Blood Age **Pulses Blood Infusion** Time Transfusion Indication A, D, +1, +2, +3☐ NSR ☐ SVT Amputation ☐ HR > 120 RAD ☐ ST ☐ VT ☐ SBP < 90 ☐ SB **BRAC** ☐ VF □ PEA CAR Peripheral IO Type / Site Central Line Location **Arterial Line** Paced Hand R L ga Fast-1 EZIO Other ☐ Triple lumen Wrist R L PED Asystole Humerus R L Arm 🔲 R 🔲 L ga ☐ Cordis Groin R L A-FIB Tibia R L A-FLUT □ Sternum PREPARED BY DEPARTMENT/SERVICE/CLINIC (Treating Unit) (Name, Rank & Title) PATIENT'S IDENTIFICATION (Name: last, first, middle; grade; date; hospital or medical facility) HISTORY/PHYSICAL TREATMENT Last Name DIAGNOSTIC STUDIES FLOW CHART Pt Cat OTHER EXAMINATION OR EVALUATION Sex ○ M ○ F Allergy Other OTHER, Specify

MEDICAL RECORD-SUPPLEMENTAL MEDICAL DATA For use of this form, see AR 40-66; the proponent agency is the Office of the Surgeon General REPORT TITLE JTS APPROVED (Date) Tactical Evacuation After Action Report & Patient Care Record, Page 2 (12 Jul 2018) -V4.1 RR SpO₂ ETCO₂ Temp AVPU GCS: Eyes 1-4 Verbal 1-5 Motor 1-6 Total First \bigcirc 0 \bigcirc \bigcirc Last PERRLA R Size (mm) ☐ L Size (mm) Field Ultrasound Results Other Diagnostics Additional Interventions Foley Comment Gastric Tube ☐ Eye Shield ☐ Protective Eyewear ☐ Right ☐ Left Comment **Protection** Immobilization ☐ C-Collar ☐ C-Spine ☐ Spine Board ☐ Pelvic Splint ☐ Pelvic Binder, Type Splint, Type/Location Warming ☐ Hypothermia Prevention, Product Hypothermia Prevention, Product Other Interventions Medications and Fluids Route = IM, IN, IO, IV, PO, PR, SL, SQ | Medications and Fluids Route = IM, IN, IO, IV, PO, PR, SL, SQ Time Drug / Fluid Drug / Fluid Route Route **Documents Received** ☐ TCCC Card ☐ Patient Chart ☐ None Other **Narrative Summary of Care Enroute Care Provider** Last Name First Name Rank Capability Signature Email PCR to: dha.jbsa.healthcare-ops.list.jts-prehospital@health.mil MM (PREPARED BY DEPARTMENT/SERVICE/CLINIC (Treating Unit) (Signature & Title) PATIENT'S IDENTIFICATION (Name: last, first, middle; grade; date; hospital or medical facility) **TREATMENT** HISTORY/PHYSICAL Last Name First Name ☐ DIAGNOSTIC STUDIES FLOW CHART Rank Unit Pt Cat OTHER EXAMINATION OR EVALUATION Sex () M () F Allergy Other OTHER, Specify

TACTICAL EVACUATION-AFTER ACTION REPORT & PATIENT CARE RECORD Page 3 IAW AR 40-68 (RAR) 22 May 2009 Paragraph 3-7. This page is a quality assurance document. Do not file in medical records. Casualty's Protective Equipment (Check all worn) Helmet, Ballistic ☐ Plate Front Neck Protector (Back) Groin Shield ☐ Blast Gauge Throat Protector (Front) Tactical Vest (IOTV) ☐ Plate Back Pelvic Undergarment Tier 1 ☐ Blast Sensor Helmet ☐ Plate Right Side Deltoid Right Eye Protection Pelvic Undergarment Tier 2 Blast Sensor Other ☐ Deltoid Left Plate Left Side Ear Protection **AAR Discussion** Event Date Tactical situation complicated care (Explain in discussion) **Sustains Improves** PATIENT'S IDENTIFICATION (Name: last, first, middle; grade; date; hospital or medical facility) The National Defense Authorization Act for fiscal year 1987 (Public Law (PL) No. 99-661), section 1102, Title 10, (10USC 1102) this document was created by or for the DOD in a medical QA program and is confidential and privileged. PL 99-661 and subsequent guidance predicated on this law (10 USC 1102) preclude MI _ BR# _____ Rank ____ Unit ____ disclosure of, or testimony about, any records or findings, recommendations, evaluations, opinions, or actions taken as part of a QA program except in limited situations. Under the provisions Pt Cat of 10 USC 1102, this information is exempt from release in accordance with Exemption 3 of the FOIA. Additional detailed information regarding the confidentiality of QA documents and records is contained in appendix B. Allergy ___ Other

K9 TACTICAL COMBAT CASUALTY CARE (K9TCCC) CARD

| EVAC TYPE: Fixed | Priority | SEVAC | |
|-----------------------------------|---|--|---|
| UNIT: | K9 NAME: | TATTOO: | |
| DATE: (DD-MM-YY) | TIME: SEX: M F | | - |
| Mechanism of Injury: (Mark) | (all that apply) | | |
| ☐IED ☐GSW ☐MINE | BURN GRENADE ARTILLERY | FALL MVC OTHER: | |
| Injury: (Mark all injuries that a | only with an XI | 2 | |
| | 9 9 Head Neck Thorax Abdomen Thoracic Lin Pelvic Limb Tail & Pelvic | 11% | |
| Vital Signs: (fill in the blank) | | | |
| | Time | | |
| Pulse Rate/Locatio | | | |
| Respiratory Ra | CALL MAN AND AND AND AND AND AND AND AND AND A | - - - - - - - - - - | |
| Temperature (99 | 125000000 | | |
| Capillary Refill | | + | |
| Blood Pressure | | | |
| Pulse Ox9 | ore (0-10) | | |
| . all so | | | |
| NOTES: | | | |
| FIRST RESPONDER NAME (| Last, First): | AOC/MOS: | |

K9 TACTICAL COMBAT CASUALTY CARE (K9TCCC) CARD

| luzzle - Handler prov | | | | |
|--|--|-------------------------|----------|------|
| Pressing - Hemostatic | Pressure Other: | | | |
| TQ - Wide Elastic | Extremity | Time | Other: | |
| Intact ETI/OTI | TRACH CRIC C | Other: | 540 | |
| O ² Chest Seal | NDC Chest Tube Other | | | |
| Catheter - IV IO | Location: | (बर्ग | | |
| RESUSCITATION | Name | Volume/Dose | Route | Time |
| K9 Blood Product | | | | |
| Crystalloid Fluid 500ml IV bolus, repeat only | once | 4 | | |
| | tion Hyperthermia – External Coo rate Head/Neck/Torso Other: | ling | | |
| | lark X if given and write route and time) | | 98791107 | |
| DRUG OPTIONS | DRUG NAMES | DOSE (30kg) | ROUTE | TIME |
| ANALGESIA Mild Pain: | Ketamine (analgesia) IV/IO/IM | 50mg | 11.00 | |
| ketamine + benzo OR opioid alone; Mod/Severe Pain: | Ketamine (sedation) IV/IO/IM | 100mg | | |
| ketamine + opioid OR ketamine + | Midazolam IV/IO/IM | 10mg | | |
| benzo + opioid | Hydromorphone IV/IO/IM | 3mg | | |
| SEDATION ketamine + benzo OR | Fentanyi IV/IO | 150mcg | | |
| ketamine + opioid | Morphine IM | 300mcg 10mg | | |
| | Other: | | | |
| | I Other. | | | |
| ANTIBIOTIC | Cefazolin/Ceftriaxone | 750mg | | |
| ANTIBIOTIC | Cefazolin/Ceftriaxone | 750mg 750mg | | |
| ANTIBIOTIC | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Frtapenem IV/IM | - markito. | | |
| | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Ertapenem IV/IM Other: | 750mg | | |
| ANTIBIOTIC | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Ertapenem IV/IM Other: TXA IV/IO | 750mg | | |
| | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Ertapenem IV/IM Other: | 750mg 750mg 0.5gm | | |
| | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Ertapenem IV/IM Other: TXA IV/IO Naloxone IV/IO | 750mg 750mg 0.5gm | | |
| | Cefazolin/Ceftriaxone IV/IM Cefotaxime IV/IM Ertapenem IV/IM Other: TXA IV/IO Naloxone IV/IO IM/IN Calcium | 750mg 750mg 0.5gm | | |

| | 5cc | 10cc | 20cc | 50cc | 100cc | 250cc | 500cc | 1000cc |
|----------------|----------------------|--------------------------------|--|---------------------------------------|---------------------------------------|-------------|----------------|----------------|
| 1mcg | 0.20mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.02mcg/ml | 0.01mcg/ml | 0.004mcg/ml | 0.002mcg/ml | 0.001m |
| 5mcg | 1mcg/ml | 0.5mcg/ml | 0.25mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.02mcg/ml | 0.01mcg/ml | 0.005m |
| 10mcg | 2mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.2mcg/ml | 0.1mcg/ml | 0.04mcg/ml | 0.02mcg/ml | 0.01mc |
| 25mcg | 5mcg/ml | 2.5mcg/ml | 1.25mcg/ml | 0.5mcg/ml | 0.25mcg/ml | 0.1mcg/ml | 0.05mcg/ml | 0.025m |
| 50mcg | 10mcg/ml | 5mcg/ml | 2.5mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.2mcg/ml | 0.1mcg/ml | 0.05mc |
| 100mcg | 20mcg/ml | 10mcg/ml | 5mcg/ml | 2mcg/ml | 1mcg/ml | 0.4mcg/ml | 0.2mcg/ml | 0.1mcg |
| 250mcg | 50mcg/ml | 25mcg/ml | 12.5mcg/ml | 5mcg/ml | 2.5mcg/ml | 1mcg/ml | 0.5mcg/ml | 0.25mc |
| 500mcg | 0.1mg/ml | 50mcg/ml | 25mcg/ml | 10mcg/ml | 5mcg/ml | 2mcg/min | 1mcg/ml | 0.5mcg |
| 1mg | 0.2mg/ml | 0.1mg/ml | 50mcg/ml | 20mcg/ml | 10mcg/ml | 4mcg/ml | 2mcg/ml | 1mcg/m |
| 2mg | 0.4mg/ml | 0.2mg/ml | 0.1mg/ml | 40mcg/ml | 20mcg/ml | 8mcg/ml | 4mcg/ml | 2mcg/m |
| 3mg | 0.6mg/ml | 0.3mg/ml | 0.15mg/ml | 60mcg/ml | 30mcg/ml | 12mcg/ml | 6mcg/ml | 3mcg/m |
| 4mg | 0.8mg/ml | 0.4mg/ml | 0.2mg/ml | 80mcg/ml | 40mcg/ml | 16mcg/ml | 8mcg/ml | 4mcg/n |
| 5mg | 1mg/ml | 0.5mg/ml | 0.25mg/ml | 0.1mg/ml | 50mcg/ml | 20mcg/ml | 10mcg/ml | 5mcg/n |
| 6mg | 1.2mg/ml | 0.6mg/ml | 0.3mg/ml | 0.12mg/ml | 60mcg/ml | 24mcg/ml | 12mcg/ml | 6mcg/n |
| 7mg | 1.4mg/ml | 0.7mg/ml | 0.35mg/ml | 0.14mg/ml | 70mcg/ml | 28mcg/ml | 14mcg/ml | 7mcg/m |
| 8mg | 1.6mg/ml | 0.8mg/ml | 0.4mg/ml | 0.16mg/ml | 80mcg/ml | 32mcg/ml | 16mcg/ml | 8mcg/n |
| 9mg | 1.8mg/ml | 0.9mg/ml | 0.45mg/ml | 0.18mg/ml | 90mcg/ml | 36mcg/ml | 18mcg/ml | 9mcg/m |
| 10mg | 2mg/ml | 1mg/ml | 0.5mg/ml | 0.2mg/ml | 0.1mg/ml | 40mcg/ml | 20mcg/ml | 10mcg/ |
| 15mg | 3mg/ml | 1.5mg/ml | 0.75mg/ml | 0.3mg/ml | 0.15mg/ml | 60mcg/ml | 30mcg/ml | 15mcg/ |
| 25mg | 5mg/ml | 2.5mg/ml | 1.25mg/ml | 0.5mg/ml | 0.25mg/ml | 0.1mg/ml | 50mcg/ml | 25mcg/ |
| 50mg | 10mg/ml | 5mg/ml | 2.5mg/ml | 1mg/ml | 0.5mg/ml | 0.2mg/ml | 0.1mg/ml | 50mcg/ |
| 75mg | 15mg/ml | 7.5mg/ml | 3.75mg/ml | 1.5mg/ml | 0.75mg/ml | 0.3mg/ml | 0.15mg/ml | 75mcg/ |
| 100mg | 20mg/ml | 10mg/ml | 5mg/ml | 2mg/ml | 1mg/ml | 0.4mg/ml | 0.2mg/ml | 0.1mg/r |
| 250mg | 50mg/ml | 25mg/ml | 12.5mg/ml | 5mg/ml | 2.5mg/ml | 1mg/ml | 0.5mg/ml | 0.25mg |
| 500mg | 100mg/ml | 50mg/ml | 25mg/ml | 10mg/ml | 5mg/ml | 2mg/ml | 1mg/ml | 0.5mg/r |
| 750mg | 150mg/ml | 75mg/ml | 37.5mg/ml | 15mg/ml | 7.5mg/ml | 3mg/ml | 1.5mg/ml | 0.75mg |
| 1Gram | 200mg/ml | <u> </u> | 50mg/ml | 20mg/ml | 10mg/ml | 4mg/ml | 2mg/ml | 1mg/ml |
| | <u> </u> | | - | | | | <u> </u> | |
| 750mg 1Gram | 150mg/ml 200mg/ml | 75mg/ml 100mg/ml Value e | 37.5mg/ml 50mg/ml quals amount o | 15mg/ml 20mg/ml f fluid in each | 7.5mg/ml 10mg/ml ml of dilution | 3n 4n | mg/ml mg/ml | mg/ml 1.5mg/ml |

71

OXYGEN CYLINDER LIFE:

| Cylinder | D | E | G | Н |
|------------|---------------------|---------------------|---------------------|---------------------|
| Liters | 356 | 622 | 5260 | 6900 |
| Flow (LPM) | 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 | E | 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:

| 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, JTTS burn sheet initiated.) | | | | | | | | |
| | 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) | | | | | | | | |
| - | | | | | | | | | |

Facility Transfer Checklist

| | DO NOW | | TAKE WITH | | INFLIGHT |
|------------|--|------------|--|------------|---|
| M: | 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. |
| A : | 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 | ë | 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 | ö 🗆 🗆 🗆 | Vasoactive medications (dopamine, neo, norepinephrine) Pressure bags IV fluids and tubing | Ö 🗆 🗆 🗆 | Check temp, pulse BP, and cardiac rhythm Assess distal pulses and neurovascular status during transport. Assess IV access LZC Pressure Lines |
| H/l: | 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 | H/l: | 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, | H/i: | 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

| | | | T | | |
|----|--|----------------|----------------|----------------------|--------------------|
| | HYPOVOLEMIC | SEPTIC | CARDIOGENIC | NEUROGENIC | BURN |
| | SHOCK | SHOCK | SHOCK₁ | SHOCK _{2,3} | SHOCK ₄ |
| | | | | | |
| 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 [] UOmL hourly |
| Activity: [] Bed rest |
| [] Spine precautions: C-Collar/C-Spine TLS Spine |
| Nursing: [] Wound VAC dressing tomm 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) ORcm H2O Suction (circle: R L Both) [] Chest tube 2 to: water seal (circle: R L Both) ORcm H2O Suction (circle: R L Both) [] Chest tube 3 to: water seal (circle: R L Both) ORcm H2O Suction (circle: R L Both) [] Chest tube 4 to: water seal (circle: R L Both) ORcm H2O Suction (circle: R L Both) [] Keep HOB elevateddegrees [] Keep HOB flat |
| Respiratory: [] Keep O2Sat > % |
| Oxygen: [] Nasal Cannula atLPM [] 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

| | Sol | ution | ıs | s Medications | | | | | | | | | | | | | | | | | | | | | | | | | | | | | | |
|-------------------|-----|-------|----|---------------|----------|---------|------------|------------------|-----------------|-----------|------------|----------|-------------|---------|----------|------------|---------|-------------------|----------|-----------|-----------|------------|-------------|------------|----------------------------|---------------|----------------|--------------|-----------------|-----------|----------------|----------|-------------|------------|
| | 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 | റ Midazolam | റ Morphine | ೧ <mark>Nicardipine</mark> | Nitroglycerin | Norepinephrine | Pantoprazole | O Phenylephrine | Phenytoin | റ Potassium Cl | Propofol | Vasopressin | Vecuronium |
| Amiodarone | ٧ | C* | Ν | Ν | С | | | С | С | С | С | С | С | С | С | Ν | Ν | ပ | C | С | Ν | С | | | | С | С | N | | N | | | С | С |
| Calcium Chloride | С | N | ٧ | Ν | С | | С | | С | С | С | С | С | С | С | С | С | O | | С | Ν | С | С | С | С | С | С | Ν | С | Ν | U | 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 | ٧ | С | С | С | С | С | | С | С | С | С | С | Ν | ٧ | Ν | | С | С | С | С | С | С | С | С | Ν | С | Ν | С | | С | С |
| Dobutamine | С | С | O | Ν | С | | С | O | С | С | | С | С | С | С | Ν | Ν | > | C | С | С | С | Ν | С | С | С | С | Z | O | Z | O | > | С | С |
| Dopamine | С | O | C | Ν | С | | С | C | С | С | С | | С | С | С | Ν | С | Ν | O | O | O | О | О | O | O | O | О | Ν | C | Ζ | C | ٧ | O | С |
| Epinephrine | ٧ | C* | С | Ν | С | | С | С | С | С | С | С | | С | С | С | С | ٧ | С | С | С | С | С | С | С | С | С | Ν | С | Ν | С | ٧ | С | С |
| Esmolol | С | С | С | С | С | С | С | С | С | С | С | С | С | | С | N | С | С | | С | С | С | С | С | С | С | С | Ν | С | ٧ | 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 | Ν | Ν | Ν | С | Ν | С | | С | ٧ | ٧ | С | Ν | С | Ν | Ν | Ν | Ν | Ν | ٧ | Z | Ν | С | С | Ν | Ν |
| Heparin | ٧ | C* | ٧ | 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) | ٧ | N | Ν | Ν | С | | С | С | С | Ν | ٧ | Ν | ٧ | С | С | ٧ | С | | Ν | ٧ | С | С | ٧ | С | С | С | Ν | ٧ | Z | Ν | С | С | ٧ | С |
| Ketamine | С | С | Ν | Ν | С | С | С | С | | | С | С | С | | С | ٧ | Ν | Ν | | ٧ | С | | С | С | | ٧ | | | | Ν | С | С | | |
| Lorazepam | С | ٧ | Ν | С | С | С | С | С | С | С | С | С | С | С | С | С | С | ٧ | ٧ | | С | С | С | С | ٧ | С | С | Ν | С | Ν | С | С | С | С |
| Magnesium | С | С | С | Ν | С | | Ν | Z | ٧ | С | С | С | С | С | С | Ν | С | С | С | С | | С | С | С | C | С | С | Ν | С | Ν | С | ٧ | 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 | ٧ | С | С | С | С | | С | С | С | С | Ν | С | Ν | С | ٧ | С | С |
| Morphine | С | С | С | С | С | | С | С | С | С | С | С | С | С | С | N | С | С | С | С | С | С | С | | С | С | С | ٧ | С | Ν | С | ٧ | С | С |
| Nicardipine | С | С | ٧ | Ν | С | | С | С | С | С | С | С | С | С | С | Ν | Ν | С | | ٧ | 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 | ٧ | С | С | Ν | С | С | С | | С | V | С | Ν | С | ٧ | 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 | Ν | N | ٧ | Ν | ٧ | | Ν | Ν | N | Ν | ٧ | Ν | ٧ | 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 | ٧ | Ν | Ν | Ν | N | | Ν | Ν | С | N | Ν | N | Ν | V | Ν | N | Ν | N | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | Ν | | Ν | N | N | N |
| 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 | | ٧ | ٧ | ٧ | С | С | С | С | С | С | С | ٧ | | ٧ | ٧ | | V | С | Ν | Ν | Ν | С | | | ٧ |
| 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 | N | С | С | | С | С | С | С | С | С | С | С | N | С | N | С | ٧ | | |

| Ateplase (tPA) | C* | N | N | |
|---------------------|----|---|---|---|
| Tenecteplace (TNK) | C* | N | N | |
| Transexamic Acid | C* | С | С | |
| Cefazolin | O | C | C | C |
| Ceftrixazone | С | С | N | С |
| Pipercillian / Tazo | С | С | V | С |
| Meripenem | ٧ | N | N | N |
| Vancomycin | O | C | C | V |
| Levaquin (Pre-mix) | С | С | С | V |

Y- Site Compatibility

C = Compatible

C* = Compatible and perferred for reconstitution V = Variable (not preferred)

N = Non - Compatable

Vasopressors should preferably run by themselves.

Not commonly carried in MEDEVAC.

IVP through open port or with mIVF (NS/LR) not with medication line or 3% line as you will bolus medication already in the line.

VITAL FUNCTIONS ASSESSMENT REFERENCE CHART

| GLASGOW COMA SCALE | | | | | | | | | | | | | |
|--------------------|--|--|---------------------------------|--|--|--|--|--|--|--|--|--|--|
| SCORE | ADULT | CHILD | INFANT | | | | | | | | | | |
| | | Eye Opening | | | | | | | | | | | |
| 4 | Spontaneous | | | | | | | | | | | | |
| 3 | To Speech | Evo Oponing Posponso | Samo as 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 Pain | Flexion Withdrawal to Pain | Withdraws from Pain Stimulus | | | | | | | | | | |
| 3 | Abnormal Flexion (Decorticate) | Abnormal Flexion (Decorticate) | Abnormal Flexion (Decorticate) | | | | | | | | | | |
| 2 | Extension (Decerebrate) | Extension (Decerebrate) | Extension (Decerebrate) | | | | | | | | | | |
| 1 | None (Flaccid) | None (Flaccid) | None (Flaccid) | | | | | | | | | | |
| | For I | ntubated Patient use Verbal "T" | | | | | | | | | | | |
| (Ex | ample: 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 EX | TREMITIES | | | | | | | | | | | |
| INJURY to Consider | MOTOR Testing | SENSATION Testing | NERVE | | | | | | | | | | |
| Elbow Injury | Index and Little Finger Abduction | Little Finger | Ulnar | | | | | | | | | | |
| Wrist Fracture or Dislocation | Thenar Contraction with Opposition | Index Finger | Median Distal | | | | | | | | | | |
| Supracondylar Fracture of Humerus | Index Tip Extension | None | Median, Anterior Interoseous | | | | | | | | | | |
| Anterior Shoulder Dislocation | Elbow Flexion | Radial Forearm | Musculocutaneous | | | | | | | | | | |
| Distal Humeral Shaft, Anterior Shoulder Dislocation | Thumb, Finger group Extension | First Dorsal Web Space | Radial | | | | | | | | | | |
| Anterior Shoulder Dislocation, Proximal Humerus | Deltoid | Lateral Shoulder | Axillary | | | | | | | | | | |
| Fracture | | | | | | | | | | | | | |
| | LOWER EX | TREMITIES | | | | | | | | | | | |
| Pubic Rami Fractures | Knee Extension | Anterior Knee | Femoral | | | | | | | | | | |
| Obturator Ring Fractures | Hip Adduction | 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, Compartment Syndrome | Ankle / Toe Dorsiflexion | Dorsal 1st-2nd Web Space | Deep Peroneal | | | | | | | | | | |
| 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 | MUSCULAR 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 | | | | | | | | | | | | | | | | | | |
|-------------------------|-----------------|--------|--------------------|----------------------|------------------|-------------|--------|-----------|------|----------|---------------|------|---------------|-------|----------|--|--|--|
| | | (Al | ways use a Broseld | ow Pediatr | ic Emergn | есу Та | ape if | availab | ole) | | | | | | | | | |
| BROSELOW cm | <61cm | 61cm | 67cm | 75cm | 87cm | 96 | cm | 109 | cm | 12 | 122cm | | 138cm | | 149+cm | | | |
| (approx) weight | 3-5kg | 6-7kg | 8-9kg | 10-11kg | 12-14kg | 15-1 | 8kg | 19-23 | 3kg | 24- | 29kg | ; | 30-36 | Skg | 37>kg | | | |
| AGE | | МО | NTHS | | | | | YI | EARS | 3 | | | | | | | | |
| | 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/ | 'Adul | t | | | Adult | | | |
| O2 MASK | | New | born | | Pe | diatric | | | | | | | Adı | ılt | | | | |
| ORAL AIRWAY | Infa | ant/Sm | nall Child | S Child | | Child | t | | С | hild/S | S Adı | ult | | Me | ed Adult | | | |
| BAG MASK | Infant | | | Pediatric | | | | | | | | Pe | Adult | | | | | |
| LARYNGOSCOPE | 0-1 | | 1 Straight | 2 Strait 2Straight/C | | | | | b | 2-3 | 3 St/0 | Curv | v 3 St/Curved | | | | | |
| ET TUBE | 2.5-3 Uncufd | 3 | .5 Uncuffed | 4 Un cuffed | 4.5 Un cuffed | 5 l cuff | | 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 | | | | nild/ dult | | | Adult | | | | |
| IV CATHETER | | 2 | 2-24 | 20-24 | 18 | 3-22 | | | 18 | 8-20 | | | | 16 | 5-20 | | | |
| OG/NG TUBE | | ; | 5-8 | 8-10 | 10 | 10- | 12 | 12- | 14 | 14 | 4-18 | | | 18 | | | | |
| CHEST TUBE | | 10 | 0-12 | 16-20 | 20 20-24 24-32 | | | | | 28 | 28-32 32-40 | | | | 32-40 | | | |
| URINARY CATHETER | | | 5-8 | 8-10 | 10 10-12 | | | | | | | - | 12 | 2 | | | | |
| CERVICAL COLLAR | N/A | Α | | Sma | I | | | S/N | M | N | Леdiu | um | | | M/L | | | |

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

| ZO | LL DEF | IBRIL | LATI | ON ENERG | GY SE | TTING | SFOR | PEDIAT | RIC PA | TIENTS | <u> </u> |
|---------------------|--------|-------|-------|---------------|---------|---------|---------|---------|---------|---------|----------|
| BROSELOW | cm | <61cm | 61cm | 67cm | 75cm | 87cm | 96cm | 109cm | 122cm | 138cm | 149+cm |
| (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 10 |
| AGE | | | MON | THS | | | | YEARS | 3 | | |
| | | 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 |
| FLUID BOLUS | | 80ml | 130 | 170ml | 210ml | 260ml | 340ml | 420ml | | 500ml | |
| ZOLL DEFIB E 1st | NERGY | 8J | 10J | 15J | 2 | 0J | 3 | OJ | 50 | J | 75J |
| 2nd | | 15J | 20J | 30J | | 5 | 0J | 75J | 100J | 120J | 150J |
| MAXIMUM | | 30J | 50J | 75J | 100J | 120J | 15 | 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 I | FUNCT | IONS | 3 B' | Y AG | <u>E</u> | | | | | | |
|------------------|---------|-------|---------------|-------------|---------|-------|------|------|----------|-----|------|----|------|-----|--------|
| BROSELOW cm | <61cm | 61cm | 67cm | 75cm | 87cm | 96cı | m | 1090 | cm | 12 | 22cr | n | 138 | cm | 149+cm |
| (approx) weight | 3-5kg | 6-7kg | 8-9kg | 10-11kg | 12-14kg | 15-18 | 8kg | 19-2 | 3kg | 24 | -291 | kg | 30-3 | 6kg | 37>kg |
| AGE | | MON | ITHS | | | | | YE | ARS | | | | | | |
| | 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 | L | 93-161 | 88-156 | 70-14 | 2 | | 59-1 | 31 | | | 52 | -115 | | 43-108 |
| RESP RATE | 25-66 | | 22-64 | 19-53 | 17-38 | 8 | | 16-2 | 29 | | | 14 | -25 | | 12-23 |
| SYSTOLIC BP | 60 | | 79-105 | 85- | 108 | 88-1 | 10 | | 91- | 119 | | | | 97 | -137 |
| DIASTOLIC BP | | | 34-81 | 40-69 45-68 | | | | | 51-89 | | | | | 59 | -86 |
| URINE (mL/kg/hr) | | 2 | 2 | 1 | .5 | 1 | | | | | | | | | 0.5 |

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

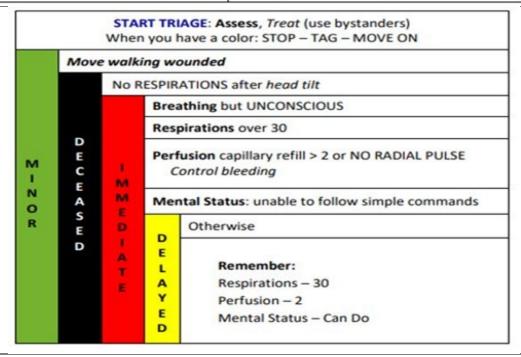
| OXYGEN SATURATION | | | | | | | | | | | |
|--------------------------|--|------------------------------|--|--|--|--|--|--|--|--|--|
| | Sea Level | 5,000 Feet MLS | | | | | | | | | |
| SpO2 (Peripheral O2 Sat) | >94% for patient with Normal Hemoglobin level | >92% | | | | | | | | | |
| StO2 (Tissue 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.

| Each Patient Triage Assessment Should Be Complete in Less Than 60 Seconds | | | | | | | | | |
|---|--|--|--|--|--|--|--|--|--|
| 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) remain armed continue to engage | Minor abrasions, burns, sprains lacerations Moderate/Mild anxiety Fractures/dislocations w/PMS Mild TBI | | | | | | | | |
| **Category IV: Expectant or Hero (blue chemlite) | Massive head or spinal injury Third degree burns > 70% BSA Injuries incompatible with life | | | | | | | | |



Notes, Warnings, Cautions

• CPG ID: 91 (Prolonged Casualty Care Guidelines)

| | | | | | | | | | | | | | | | | | <u>T</u> . | ΑE | <u>3L</u> | <u>.E</u> | C | F | C | 10 | NTE | NT | <u>S</u> | | | | | | | | | | | | | | | | |
|--|--|----------------------------|---|---|---|--|---------------------------------|---------------------------|---|---------------------------|--------------------------------|------------------------------------|--------------------|---------------------------|-----------------------------------|--------------------------------|------------------------------------|-------------|--------------------------|----------------------------------|---|-----------------------------------|---|--------------------------------------|--|---|----------|--|------------------------|----------------------------------|----------------------------------|--|--|---|---|---------------------------------------|------------------------------------|---------------------------|----------------------------|-----------|--|------------------------------|--|
| Magnesium Sulfate | Epinephrine 1mg/10mL | Atropine | Adenosine | Amiodarone (Infusion) | Amiodarone (Cardiac Arrest) | Epmephrine Img/10mL | Nitroglycerin Tablet/Spray | Aspirin, Chewable, 81mg | CARDIAC (See also Morphine or Fentanyl for AMI pain | Chrogon Kit | HYPOGLYCEMIA D50 | Epinephrine lmg/lmL | Albuterol(MDI) | Albuterol(Nebulizer) | RESPIRATORY (See also: Methylpred | Methylaredwicolome/Solm-Medral | Epinephrine lmg/lmL | ANAPHYLAXIS | Promethazine (Phenergan) | ANTI-EMETICS Zefren(Ondensetren) | Naloxone (Narcan) | Glucago | OD/Tox Ingestions (See also Sodium Bicarbonate, | Phenylephrine(NEO) | Norepinephrine (Levophed) | Epinephrine lmg/l0mL | PRESSERS | Vecuronium | Succinylcholine | Rocuronium (Zemuron) | Midazolam (Versed)** | Etomidate | Propofol (Constant Infusion) | Propofol (Bolus) | Ketamine ** HIGH DOSE | SEDATION ** = Controlled Substance | Merchine ** | Votorolos (Torodol) | Ketamine ** LOW-DOSE | ANALGESIA | Tranexamic Acid (TXA) | DRUG 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 ours | 300 mg 1" Dose 150 mg 2" Dose | Img | 0.4 mg SL | ll | nyl for AMI pain) | Date CT-01 | 10-25 Grams IV/IO | 0.3-0.5 mg SQ/IM or 0.5 ml Neb | 4-8 Puffs | 2.5-5 mg | brednisolone) | OLVI pur SCI | 0.3-0.5 mg IM | | 12.5-25 mg IV/IO/IM | 4.8 mg 11/70/11/200 | 0.4-2 mg IV/IO/IM | | <u>Ω</u> | 10 mg/10 0ml NS= 100mcg/ml or | 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 bolus IV/IO | 1-2 mg/kg IV/IO | AT BY SHIP OF AT SHIP C.T. | 2-5 mg [Wild or 0] mg/kg [W | 15 mar IV or 15 30mar IV | 0.1-0.3 mg/kg IV/IO >1 min | | 2000 mg IV/IO | STANDARD DOSING | |
| 1-2 Gram | 2-10mcg/ | | 6mg 1st d | 150mg in | 300mg IV | lmg (1 am | 11 | | | | | 0.3-0.5 | | | | | | | | | 0.4-21 | | | 50-2 | Start 2- BP) | 2-20mc | | ómg | 60-90mg | 36-72mg | 3-6mg | 12-24mg | 600-4500 mcg/min | 60-150mg | 60-120mg | Smo | ón:e | Shmoo.oo | 6-18 mg | | 2 Gran | SMALL ADULT (60KG)132LBS | |
| 1-2 Gram diluted in 50ml D5W over 15 min(Torsades w/ pulseWF/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 3-8min give 180 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) | TAXA (Smr) 101 | 1 Kit (Img) IV/IV | 20-50ml | 0.3-0.5mg SQ/IM or 0.5ml w/ 3ml NS | | 2.5-5mg (mixed in 3ml NS) | Smozz | 125mg | l Auto-Injector or 0.3mg IM | c | 12.5-25mg | d.Smig | 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) Ouce BP is appropriate, 2-4mcg/min | 2-20mcg IVP q 2-5min or 2-20mcg/min IV/IO infusion | | Smg | 80-120mg | 48-96mg | 4-8mg | 16-32mg | 800-6000 mcg/min | 80-200mg | 80-160mg | 9mc | T Sunc-et to AT Sunct | 16mg IV or 15 20mg IV | 8-24 mg | | 2 Grams in 100cc NS over 10min or slow IVP | ADULT (80KG)176LBS | |
| 15 min(Torsades | desired effect | | push w/ rapid, | wed by 360mg hrs | -5min give 150 | rest | X3 | J) | | | | ml NS | | | | | Ţ | | | | rentilation . | | | IVP/IO | y. (adjust for 4mcg/min | g/min IV/IO | | 10mg | 100-150mg | 60-120mg | 5-10mg | 20-40mg | 1000-7500 mcg/min | 100-250mg | 100-200mg | Smor | | | 10-30 mg | | or slow IVP | LARGE ADULT (100KG)220LBS | |
| Torsades De Pointes (with or without pulse) | Symptomatic Bradycardia | Symptomatic Bradycardia | Stable, narrow complex tach/PSVT | Hemodynamically unstable V-Tach/SVT | Refractory Pulseless V- Fib/V-Tach | Pulseless Arrest | Angina/ AMI | Angina/ AMI | 11) pogri cemm | Hypoglycemia | Hynoslycamia | Bronchodilator | Bronchodilator | Bronchodilator | Amiphy again Assume | Anaphriavis/Actions | Anaphylaxis | | Antiemetic/Sedation | Antiomatic | Opioid 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 Anesthesia (Non-analgesic) | Dissociative Sedation/ RSI | Fam, Augusty u., Arst. | Dain Angioktic AMI | Muscaladalahi Dia | Analgesia Analgesia AMI | | Int/Ext Hemorrhage | INDICATIONS | |
| AV Blocks | Use if refractive to Attopine Pacing | Glaucoma | May cause transient Asystole following push | Simus Bradycardia, 2nd 3rd Deg Block | Sinus Bradycardia, 2nd 3rd Deg Block | Profusing Tachycardia's | Maintain SYS BP>90 | Must be chewed | Am) is the area areas | Only if Dig not available | Intracranial bemombage | No IV use | Cardiac arrhythmia | Cardiac arrhythmia | Vir man for sen for | Do not true for head Tw | Hold for 10 sec | | Altered LOC/Vesicant | Can cause O'C and loweston | Use minimum needed | Dosage is higher than kits | AV proof 181 | Must be diluted. Max resus | Max resus w/ blood lst | Must be diluted. Max 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 | | Resn/RD draw: Head Tv | Not for Datiofald Transco | HTN, Emergence | | Give <3 hrs from injury/surgery | RESTRICTIONS/WARNINGS | |
| 30 min | Maintain drip | 5-15min | 1-2 min | Maintain drip | 3-5 min | 3-5 mm | 20-30 min | 4-6hrs | cists | TIME COME | JAK. | 5-15m | 1-4hrs | 1-4lus | | 1-0115 | 5-10min | | +6hrs | 4.600 | 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 | Valida | Variable | 4.60- | 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 | Max 2 doses | Repeat Q 3-5mm w/ CPR-No Max | May repeat up to may of 3 Doses | NO REPEAT | PARTOTA EVEL) ANIMETERA | REDEAT anary 20min DRN | Titrate: avoid over-correction | Neb over 15 min | | | TO DEFECT | NO REDEAT | Add doses Q 5-10 min until improve | | Max 25mg q 4 hrs | Max Source 6 low | Q 2-3 min PRN (Max 10mg) | Titrate infusion for hemodynamics | doses and thrate PKN | PRN to maintain SYS BP; start w/low | 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 | EVEN TOT DEMOTRAL IT THE VEST PROVIDE | DRN for sedation if RD/Ream stable | Co-co min facts | Q 10-30 min PRN for pain | | NO REPEAT | REPEATABILITY/ MAX DOSE | |

| ASYSTOLEPEABRADYCARDIA | Cardiopulmonary Arrest BLS | Alternate K.9 S | Agitated K9 Sedation – IM 1st/IV PRN | Mild Sed | Analgesia Continuous IV/IO Infusion | Analgesia-Inten | Bio. | Heart Rate | Blood Pressure | Respirations | Temperature | VITALS | | \ | Hematocrit(Hct) | 5-10 M:45-52 F:37-48 | WBC M:13-18 F:12-16 | Hemoglobin(Hgb) | / | TENTAL | Acetaminophen | FFP (lu=200-280ml) | PRBC (1u=250ml) | Blood Products and Management | Hypertonic Saline (3%) | Burns >20% TBSA | | Maintenance | FLUIDS Permettation (Creatalnis) | Mannitol (20%) | 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 |
|--|--|---|--|---|---|-----------------------------------|--|--------------------------|--------------------------------|---|--|------------------------|---------------------------|-------------------------------|-----------------------------|----------------------|---------------------|-----------------|---------------------------|-----------------------|----------------------------------|---|---|-------------------------------|-------------------------------------|---|--|-----------------------|-----------------------------------|-------------------------------------|--|--|---|----------------------------|---|---|-------|---|----------------------------|--|-------------------|------------------------------|
| RADYCARDIA | UMODRY ATTEST BLS | Alternate K.9 Sedathon IV/IO | on IM 1st/IV PRN | Mild Sedation IM | uous IV/IO Infusion | :-Intermittent IV/IO/IM | 35-45 mmHg | 60-80 | 120/80 (avg) mm Hg; MAP 90-100 | 16-30MIN | 101-103 ℉ | Normal | MILITAR | / | | 450,000 | \ | \ | \ | HEMMIOTORI | 500 mg PO or 1 G IV | 10 mJ kg | 10 mlkg | | 0.1-1 mi/kg/frr | man of man (standard but to bond menus) | IR 10 ml + %TRS4 (Based on 40- 80kg adnir) | 1-2 ml/kg | 70 ml/c | 1 Gram/kg IV over <20 min | 1-2 Gram IV/IO | 2-4 mg IV/IO | Anxien; 2-10 mg IV/IM q 6ins//Seizures; 5-10mg q 5-10min (MAXX 30mg)//Seizures following Nerve agent <u>Exposure</u> ; 10-20 mg IM for seizures or if 3x Mark 1 Kits used | | 1-3 Auto-Injectors (600 mg sa) | 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 | 1 mEq/kg IV/10 | | STANDARD DOSING |
| nem c | Dafih 2 | | | | | | | | | Panting | TO3 FF | Excited | MILITARY WORKING DOG | .45 | DH P | | | 3.5 | 135 | 2 | 5001 | 1-2 units PRN to achieve 1:1 ratio w/ PRBC's (Shelf Life(thawed)=\$ days) | 1-2 w | | 250ml | Hour | 10ml + 06 RSA Par | 75ml/hr(TKO) | 250 500 | 60G | Seizures = 1-2G over 30 : 2G over 20 min | Seizures = | <u>es</u> : 5-10mg q 5-10min (MA) 0 mg IM for seizures or if 3 ₃ | | Inject 1-3 injectors (base Injector contains | | | 500-1000mg over 2-5 mi >5min for Ca | | 60mEq | | SMALL ADULT (60KG)132LBS |
| A. A. Den | Compressions @ | PRO | MIDAZOLAM | MIDAZO: | FENTANYL 2-10 mcg/kg | HYDROMO | | | | give 25% > 20 min/PR1 | Shock: Calculate 90ml/kj | | | 8 | Paco. Pao. | BL | K co' | | 135-145 95-105 | 2 | 500mg PO or 1Gram IV infusion | 1:1 ratio w/ PRBC's (She | 1-2 units PRN to achieve Sys BP ~90 Life =42 days) | | 250ml bolus followed by 50-100ml/hr | Hour | 10ml + 06 RSA Per | KO) 105ml/hr(TKO) 150 | Dolur to achieve curtolic | 30G | Seizures = 1-2G over 30 min; Wheezing/ Respiratory Distress (3rd line) = 2G over 20 min; (Pre)Eclampsia = 4-6G over 18-20min | 4mg q 3-5 prn; Agitated/Combative Patient = 2-4mg q 30-60 | X 30mg)//Seizures followi : 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 | 80mEq | | ADULT (80KG)176LBS |
| ATROPINE 0.04mg/kg IV/IO AND EPINEP | EPI 0.01 mg/kg 4 NT | POFOL 1 mg/kg boluses i | MIDAZOLAM 0.3mg/kg AND KETAMINE 2mg/kg AND | LAM 0.3mg/kg IM AND 1 | g/hr OR MORPHINE 0.1- | RPHONE 0.1-0.2mg/kg O | Defib 2-5 Joules/Kg | Intubate w/ 10.0 ET tube | (PRN) | N); reassess; give 25% >10 | Shock: Calculate 90ml/kg for total fluid to be infused: Give 25% > 1 | FLUID MANAGEMENT | | 22-26/ | /HCO | BLOOD GAS | cre | 0.6-1.5 | \ | GILLES | infusion | If Life(thawed) =5 days) | P-90 | | ml/hr | | | 150ml/hr(TKO) | , aa - 00 | 100G | ory Distress (3rd line) = over 15-20min | Combative | ng Nerve agent | | s) IM. DuoDote/ATNAA nd 2-Pam(600mg) | | | ag/kg>5-10min for Beta in after Blood | В | 100mEq | | LARGE ADULT (100KG)220LBS |
| IO AND EPINEPHRINE 0.01 mg/kg | Compressions @ Lowmin, establish arrway, respirations @ 3-10 mm for 2-3 min (FPR _ Deft _ Deft _ Deft _ Deft _ | PROPOFOL 1 mg/kg boluses PKN to allow catheterization or intubation | NE 2mg/kg AND HYDROMORPHONE 0.2mg/kg | MIDAZOLAM 0.3mg/kg IM AND HYDROMORPHONE 0.2mg/kg IM Q2-4hrs | FENTANYL 2-10 mcg/kg/hr OR MORPHINE 0.1-0.25 mg/kg/hr OR HYDROMORPHONE 0.02-0.05 mg/kg/hr | oke 02-4hrs OR MORPHINE 0.2-0.5 m | THE PROPERTY OF THE PROPERTY O | whe | | give 25% > 20 min/PRN); reassess; give 25% >10 min(PRN); last 25% >10 min | sed: Give 25% > 10 min; reassess; | NT | | 95-100/ ±2 | | | / | OLT-02 | <u> </u> | | Febrile Reaction | Int/Ext Hemorrhage/ AB+ Uni Donor | Int/Ext Hemorrhage/ O- Neg Uni Donor | | ICP Reduction | full-thickness burns | >20% TRSA partial or | IV access/Homeostasis | Uma tancianiralamia | 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 | Organophosphate/ Nerve Agent | | Hyperkalemia/Beta&Calcium Channel Blocker OD | HTN Urgency/Emergency | TCA OD; Prolonged Cardiac Arrest | | INDICATIONS |
| S Treme The Company of the Company o | Defit 2min CPR Defit | ation | HONE 0.2mg/kg | N Q2-4hrs | HONE 0.02-0.05 mg/kg/hr | gike 04-6hrs | | | | D.O.P.E Displacemen | | Women (kg) | Men (kg) | | TIDAL VOLUME | I:E | Fi02 | RATE | MODE | 100 | Infuse slowly | Monitor for Anaphylaxis' Hyperthermia | Monitor for Anaphylaxis/ Hyperthemsia/HyperK | | Use only in Head Injuries | A DESCRIPTION OF THE PROPERTY | Track start time and amount infliced | Do not over hydrate | UTO DA ANT TO A BOOK A PROPERTY | Avoid in HoTN Pts | Dilute into 50-100ml NS or D5W | IM not recommended due to erratic dosorption | Respiratory Depression | | Use Attopine 1st if only using single dose 2-Pam (Mark 1/NAAK 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 |
| | | | | | | Ī | | | | nt; Obstructions; Pneumo | Troubleshooting | | | Ideal Body Weight Calculation | | | | | | Initial Vent Settings | Office | PRN | PRN | | N/A | NA | | PRN | TVQ0 | 3-8hrs | 30 min | 30-120min | 20-30min | | 15min | 5-15min | | 30min-4 hrs | 15-60 min | 1-2hrs | | DURATION |
| | | | | | | NOTES | 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) | | omitikg (tdeat body weight) | 1:02 | 1.0 (100%) | 14BPM | AC/ASV (Hamilton T1 only) | MELY 1 | Use only for Non-Hemolytic react | Ideal ratio of FPD-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 | | Titrate to effect | Titrate to majestic CDB CO | 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 infusion for Beta OD; 1000mg Q 10- $20\mathrm{m}^2$ 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 |

ALTITUDE ILLNESS

Signs and Symptoms

Acute Mountain Sickness (AMS)

- Headache
- Nausea
- Vomiting
- Lethargy
- Dizziness

High Altitude Cerebral Edema (HACE)

- Worse Mountain Sickness
- Ataxia
- Altered Mental Status

Hight Altitude Pulmonary Edema (HAPE)

- Cough
- Dyspea
- Pink Frothy Sputum
- Consider DifDx

Treatment

For all:

- o O2 (100% FiO2) / IV/IO Access / Cardiac monitor
- o **Descend as soon as possible** (Consider Gamow bag if unable)
- Hypothermia prevention
- Note to aircrew fly lowest allowable altitude
- HAPE Pulmonary Symptoms
 - o Nifedipine 30mg ER PO q12hrs (or 20mg imm. rel. PO q8hrs)
 - Consider Assisted Ventilation
- **HACE** Headache with altered mental status or Ataxia
 - Dexamethasone 8mg IV/IO/PO x1, then 4mg q6hrs
 - Pediatric: 0.15 mg/kg IM/IO g6hrs
 - Acetazolamide 250mg PO q12hrs
- AMS Headache without altered mental status or Ataxia
 - Acetazolamide 250mg PO q12hrs (avoid with sulfa allergy/sickle cell)
 - Severe AMS: Add Dexamethasone 4mg IV/IO/PO
- For altitude-related headache (in isolation or with AMS/HACE) consider
 - Acetaminophen 650-100mg PO
 - Ibuprofen 600-800mg PO

- The treatment of choice for all altitude-related illnesses is supplemental O2 and descent at least 500-1000m. If unable to descend, a hyperbaric bag (Gamow bag) can be utilized if available. If unable to descend immediately as soon as HACE or HAPE are suspected, the crew must begin engaging 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.
 - O 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

Treatment

- 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.

- 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.
- o 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.
- o 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.
- o Inner ear disturbance's (vestibular DCS) vertigo, nausea, vomiting. More likely associated with diving than altitude exposure.

Treatment

- 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.

- 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

HOT Weather Signs & Symptoms

- AMS
- LOC
- Pale or clammy skin
- Hypotension or shock
- Seizure
- Nausea or Vomiting

COLD Weather Signs & Symptoms

- Cold, Clammy skin
- · Shivering or lack of shivering
- Mental status change
- Extremity pain or numbness
- Bradycardia or Arrhythmia
- Hypotension or Shock

HOT Weather Treatment

- Remove from heat source/ loosen or remove clothing.
- AMS -AMS guidelines
 - Glucose check
- Consider intubation if needed.
- 1L IV Bolus or PO Fluids
- Monitor 12 lead EKG for arrhythmias.
- Be prepared for Seizure Seizure guidelines
- AMS & core temp > 40C /104F
 - Start aggressive cooling (Tepid water to skin and fanning / Ice packs in groin, axilla, and neck / D/C once temp <40 C /104 F)
 - Consider benzodiazepines to block/stop shivering & rebound hypothermia.
 - Midazolam 0.1 mg/kg
- AMS & core temp < 40C/104F
 - o Tepid water or room temp water to skin
- Continuous monitoring

COLD Weather Treatment

- Remove wet clothing
- Assess: Mental status, Rectal temperature, Glucose
- Core Temp < 35C/95F With AMS (arrhythmia, absence of shivering)
 - HPMK kit / Hypothermia blankets
 - Dry clothing
 - Hot Packs to groin, axilla, abdomen (avoid burning pt)
 - Warmed IV fluids 1L IVF Bolus
- Core Temp < 35C/95F (Alert, w/o Arrhythmia, actively shivering)
 - Drv blankets
 - Hypothermia Blanket
 - Warm PO fluids
- Monitor 12-lead EKG

Notes, Warnings, Cautions

HOT Weather

- Best method to cool pt is sublimation-sprinkling w/ water + fanning to evaporate on skin
- Elevated risk groups: Elderly, Very young, Highly active
- Sweating does not exclude heat stroke / heat illness

COLD Weather

- "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 wait at least one minute to feel pulse
- At temps < 30C/86F one defibrillation can be attempted, but withhold further attempts until temp > 30C/86F

SUBMERSION INJURY

Signs and Symptoms

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

Treatment

- O2 (100% FiO2 for all injuries)
- IV/IO, 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

- 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)

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

Trade Name: Tylenol

<u>Pain or fever:</u> (Limit total daily dose to <4 g/day) PO:

 Regular release: 325-500 mg every 4-6 hours or 1000 mg 3-4 times daily (maximum: 4 g/day)

IV:

• 500-1000 mg every 6 hours

Pain or fever:

PO:

 Infants and Children <12 years: 10-15 mg/kg/dose every 4-6 hours as needed; do 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 Yes (Caution)

Class / Mechanism of Action

Diuretic, Carbonic Anhydrase Inhibitor; Anticonvulsant

Inhibits carbonic anhydrase causing a decrease in hydrogen ion renal secretion with increased renal secretion of sodium, potassium, bicarbonate, and water. Onset of action PO: 2 hours, IV 5-10 minutes

Indications

Labeled Indications:

- Prevention or treatment of symptoms of acute mountain sickness
- Edema due to congestive heart failure

Contraindications

- · Hypersensitivity to acetazolamide, sulfonamides, or any component of the formulation
- Confirmed low sodium / potassium levels otherwise none in emergency setting

Adverse Reactions / Precautions

- May worsen respiratory acidosis
- Drowsiness, deceased alertness, impairment of coordination, nausea, headache
- Flushing of skin, allergic skin reaction, skin photosensitivity

Dose and Administration:

ADULT

PEDIATRIC Always Reference LB tape

Trade Name: Diamox

Altitude illness (Acute Mountain Sickness):

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.

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 Low Dose OK)

Trade Name: **Aspirin**

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

Acute coronary syndrome (ST-segment elevation myocardial infarction [STEMI], unstable angina (UA)/non-ST-segment elevation 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)

N/A:

Contraindicated in children under 16 yrs. (Reye's Syndrome)

ACTIVATED CHARCOAL

QSafe, Lactation Safe

Class / Mechanism of Action

Antidote

Non-absorbable agent that absorbs toxins within the GI tract inhibiting GI absorption.

Indications

Labeled Indications: Management of suspected or known poisonings when gastrointestinal decontamination is an option.

• Decontamination within 1 hour of ingestion of toxic substance

Contraindications

- · Presence of intestinal obstruction or GI tract not anatomically intact
- Patients at risk of GI hemorrhage or perforation
- Patients with an unprotected airway (e.g., CNS depression without intubation) or if use would increase the risk and severity of aspiration

Adverse Reactions / Precautions

- If patient unconscious, must establish airway control and must utilized NG/OG tube.
- Be prepared for possible emesis. Consider use of antiemetic.

ADULT

Avoid use in patients at risk of GI hemorrhage or perforation

Dose and Administration:

PEDIATRIC Always Reference BROSELOW Tape

Acute Poisoning:

PO, NG/OG:

Single dose: 25-100 grams

Note: Activated Charcoal has limited efficacy if not utilized within 1 hour of toxin ingestion. Risk-benefit of charcoal must seriously be considered because it does not work for all poisons, it must be given early when the poison is still in the stomach, it does not fully bind all poisons, and serious complications can occur with aspiration. Aspiration can occur if deteriorating mental status and/or vomiting.

Note: Some products may contain sorbitol. Coadministration 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)

<u>Acute Poisoning</u>: Children >12 years: Refer to adult dosing.

Trade Name: Actidose

PO, NG/OG:

- Single dose: 1 gram/kg
- Multidose: Initial dose: 1Gram/kg initially, followed by multiple doses of 0.5 Gram/kg every 2 hours

Note: Some products may contain sorbitol. Coadministration of a cathartic, including sorbitol, is no longer recommended.

Note: Activated Charcoal has limited efficacy if not utilized within 1 hour of toxin ingestion.

ADENOSINE

QC, Lactation Yes (Caution)

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

Trade Name: Adenocard®

Paroxysmal supraventricular tachycardia:

I.V. (rapid push, over 1-2 seconds, via proximal peripheral line (forearm or above, large bore).

Initial: 6 mg; if not effective within 1-2 minutes,
 12 mg may be given if needed (maximum single dose: 12 mg).

Notes): Follow each dose with **10-20 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.

Paroxysmal supraventricular tachycardia:

IV/IO as close to core as possible (rapid push, over 1-2 seconds, see **Note**): Follow each dose with 10-20 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 each dose with 5-10 mL normal saline flush.

ALBUTEROL QC, Lactation Yes Trade Name: Proventil / Ventolin

Class / Mechanism of Action

Beta₂ Agonist (Bronchodilator)

Synthetic sympathomimetic that relaxes bronchial smooth muscle, causing bronchodilation, with little cardiac impact. Onset of action is 2-15 minutes

Indications

Labeled Indications: Treatment or prevention of bronchospasm in patients with reversible obstructive airway disease; prevention of exercise-induced bronchospasm

- Asthma
- Reactive Airway / Bronchospasm
- COPD
- May also be used in Crush Syndrome (Hyperkalemia)

Contraindications

- Hypersensitivity to albuterol or any component of the formulation
- · Symptomatic tachycardia

Adverse Reactions / Precautions

- Risk of abortion during 1st or 2nd trimester
- Headache, Dizziness, Flushing, Diaphoresis, Tremor, Weakness
- Angina, A-Fib, Arrhythmia, Chest Pain, Palpitations
- Dyspnea, Bronchospasm in asthmatics

Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape

Bronchospasm:

Metered-dose inhaler (90-180 mcg/puff):

6 puffs

Solution for nebulization:

5 ma

Respiratory Distress (acute, severe):

Metered-dose inhaler:

1-2 puffs

Solution for nebulization:

• 2.5-5 mg

Bronchospasm:

Metered-dose inhaler (90 mcg/puff):

6 puffs every 4-6 hours as needed

Solution for nebulization:

- 5 ma
- Children ≥12 years: Refer to adult dosing.

Exacerbation of asthma (acute, severe):

Metered-dose inhaler (90 mcg/puff):

- 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.

Solution for nebulization:

- 2.5-5 mg every 20 min
- Children ≥12 years: Refer to adult dosing.

AMIODARONE

♀D. Lactation: Yes, Not Recommended

Class / Mechanism of Action

Antiarrhythmic Agent, Class III

Inhibits adrenergic stimulation (alpha and beta blocking), prolongs action potential and refractory period (prolongs PR and QT intervals); decreases AV conduction and sinus node function (decreases sinus rate)

Indications

Labeled Indications: Management of life-threatening recurrent ventricular fibrillation (VF) or hemodynamically unstable ventricular tachycardia (VT) refractory to other antiarrhythmic agents **Unlabeled:**

- Recurrent, hemodynamically unstable VT. (after other drugs have failed)
- Ventricular tachyarrhythmias (ALS/PALS): VF/VT Cardiac arrest unresponsive to CPR, Shock, and Vasopressor.

Contraindications

- Hypersensitivity to amiodarone, iodine, or any component of the formulation
- Severe sinus-node dysfunction
- 2nd and 3rd degree heart block (except in patients with a functioning artificial pacemaker)
- Bradycardia causing syncope (except in patients with a functioning artificial pacemaker)
- Cardiogenic shock

Adverse Reactions / Precautions

- Complex drug with multiple complex drug reactions! (Do not administer with procainamide)
- Hypotension
- Dizziness, fatigue, Headache, Poor coordination, Neuropathy
- Nausea, Vomiting
- Dysrhythmias, Asystole, AF, Bradycardia, AV block, Conduction abnormalities, SA node dysfunction

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Pulseless VT or VF (ALS):

IV/IO push

 300 mg rapid bolus if pulseless VT or VF continues after subsequent defibrillation attempt or recurs, administer supplemental dose of 150 mg.

Recurrent, Hemodynamically unstable VT

(ARC):2020

Initial Dose:

IV/IO slow push

 150mg IV over 1st 10 minutes (May repeat 150 mg every 10 minutes PRN if VT recurs

Maintenance Infusion following initial dosing:

• 1 mg per min over 6 hours

Pulseless VT or VF (PALS):

IV/IO push

- 5mg/kg IV bolus during cardiac arrest, May repeat twice for refractory VF/pulseless VT.
- Max single dose: 300mg

<u>Tachycardia with Pulse and poor perfusion, or symptomatic with adequate perfusion</u> (PALS):

IV/IO push

- Loading dose: 5mg/kg over 20 to 60 minutes (Fast push or bolus can precipitate cardiac failure!)
- Max single dose: 300mg

| AMIODARONE | | | | | | | | | | | | | |
|--------------------------------|---|-------|--|-------|--|--|--|--|--|--|--|--|--|
| Initial Dose: I50mg over I0min | | | | | | | | | | | | | |
| | MIX 150 mg/100 ml CONCENTRATION 1.5 mg/ml | | | | | | | | | | | | |
| Dose | Rate | Micro | | Macro | | | | | | | | | |
| | 60 20 gtt/ml 15 gtt/ml 10 gtt/ml | | | | | | | | | | | | |
| mg/min | mg/min ml/min gtt/min gtt/min gtt/min gtt/min | | | | | | | | | | | | |
| 15 | 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 | 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

Anticholinergic, Antidysrhythmic, Antidote for Carbamate Anticholinesterase poisoning Blocks acetylcholine at parasympathetic sites in smooth muscle, secretory glands, and the CNS; increases cardiac output, and dries secretions. Atropine reverses the muscarinic effects of cholinergic poisoning. Reverses bronchorrhea and bronchoconstriction but does not affect the nicotinic receptors responsible for muscle weakness, fasciculations, and paralysis.

Indications

Labeled Indications: Treatment of

- Symptomatic Sinus Bradycardia, AV block (nodal level)
- Antidote for anticholinesterase poisoning (carbamate insecticides, nerve agents, organophosphate insecticides)

Contraindications

- Hypersensitivity to atropine or any component of the formulation
- Narrow-angle glaucoma; adhesions between the iris and lens (ophthalmic product)
- Pyloric stenosis
- Prostatic hypertrophy
- Note: NO contraindications should prevent use of atropine in setting of lifethreatening organophosphate, carbamate, or nerve agent poisoning

Adverse Reactions I Precautions

- Tachycardia and arrhythmia (VTach, VFib), Hypotension, Palpitations
- Dilated Pupils, Angle-closure glaucoma
- Headache, Dry Mouth, constipation, urinary retention, flushing.
- Paradoxical Bradycardia noted with doses less than 0.1mg

Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape

Symptomatic Bradycardia

IV/IO

 1 mg every 3-5 minutes, not to exceed a total of 3 mg or 0.04 mg/kg (ARC, 2020)

Organophosphate or carbamate insecticide or nerve agent poisoning:

IV/IO: (Nerve agent) Atropine **20mg in 250mL**, titrate to dry respiratory secretions.

IV/IM: (Used with 2-Pam Chloride auto injector)

 Initial: 1-6 mg; repeat every 3-5 minutes as needed, doubling the dose if previous dose did not induce atropinization. Maintain with repeat doses as needed for 2 2-12 hours based on recurrence of symptoms.

IM (<u>AtroPen®</u>): Follow with 2-Pam Chloride auto injector.

- Mild symptoms (>2 mild symptoms): 2
 mg once an exposure is known or
 strongly suspected.
- Severe symptoms (>1 severe symptom): Three 2 mg doses in rapid succession.

Symptomatic Bradycardia

IV/IO

 0.02 mg/kg (Minimum dose is 0.1 mg. Maximum single dose of 0.5 mg. May repeat once in 3-5 minutes. Maximum total dose is 1 mg (PALS, 2020)

Organophosphate or carbamate insecticide:

IV/IO: Initial: 0.05-0.1 mg/kg; repeat every 5-10 minutes as needed, double dose if previous dose does not induce atropinization. Maintain with repeat doses as needed for 22-12 hours based on recurrence of symptoms.

Severe Nerve Agent Poisoning:

 IV/IO 1mg every 3 min. Monitor Patient for signs and symptoms of atropinization, (drying up of secretions). Once clinical improvement is achieved restrict to 10- 20% of original dose (approximately 2- 4mg/hr)

CALCIUM Chloride 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 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) (CaCl preferred over Calcium Gluconate for this use): IV/IO

 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

<u>Hypocalcemia prophylaxis from massive</u> <u>transfusion</u>

• 10ml (10cc) 10% solution over 5 minutes

Damage Control Resuscitation: IV/IO, SLOW

1000 mg after 1st blood unit and after every 4th unit.

May be given before TXA

Cardiac arrest or cardiotoxicity in the presence of hyperkalemia. hypocalcemia. or hypermagnesemia:

IV/IO. SLOW

• **20 mglkg** (maximum: 2000 mg/dose); may repeat as necessary.

Calcium channel blocker overdose (unlabeled use):

IV/IO

 Initial: 20 mglkg (0.2ml/kg) (maximum: 1000 mg/dose) over 10-15 minutes; may repeat every 10-15 minutes

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. (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

CALCIUM Gluconate

QSafe, Lactation Safe

Class / Mechanism of Action

Calcium Salt, Electrolyte Supplement

Moderates nerve and muscle contractility via regulation of action potential excitation threshold.

Indications

Labeled Indications: Treatment of hypocalcemia and conditions secondary to hypocalcemia (e.g, tetany, seizures, arrhythmias); cardiac disturbances secondary to hyperkalemia; magnesium sulfate overdose; massive transfusion prophylaxis

Unlabeled: Calcium channel blocker overdose; treatment of hydrofluoric acid exposure

Contraindications

- Ventricular fibrillation
- Hypercalcemia
- Concomitant use of IV calcium gluconate and ceftriaxone in neonates (risk of precipitation of calciumceftriaxone)

Adverse Reactions / 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
 - Do not exceed 200mg/min except in emergency situations
- Caution in patients receiving digoxin therapy, may cause arrhythmias
- Use small IV / Large Vein, flush prior and after, AVOID extravasation (will cause tissue necrosis)
 - In general, IV Calcium Gluconate is preferred over I.V. 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

1500-3000mg over 2-5 minutes

<u>Calcium channel blocker overdose (off-label use):</u> <u>Hypotension/conduction disturbances:</u>

IV/IO

3 Grams (3000mg) over 5 minutes; may repeat every 10-20 minutes with 3-4 additional doses.

Hypocalcemia prophylaxis from massive transfusion 30mL of 10% solution over 5 minutes

Note: Calcium Chloride is 3X more potent than Calcium Gluconate and therefore higher doses of Calcium Gluconate must be used to reach similar therapeutic doses.

<u>Hydrofluoric Acid Exposure</u> – (off-label, see Burn 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!)

Cardiac arrest or cardiotoxicity in the presence of hyperkalemia, hypocalcemia, or hypermagnesemia:

IV/IO, SLOW

60-100 mg/kg/dose (maximum: 3000 mg/dose)

<u>Calcium channel blocker overdose (unlabeled use):</u> <u>Hypotension/conduction disturbances:</u>

IV/IO

45 mg/kg (maximum 3000mg/dose) over 10-15 minutes; may repeat every 10-15 minutes

Hypocalcemia prophylaxis from massive transfusion

60mg/kg (maximum 30ml of 10% solution) over 5 minutes

Note: Calcium chloride may provide a more rapid increase of ionized calcium in critically ill children.

CEFAZOLIN QC, Lactation Yes Trade Name: Ancef

Class I Mechanism of Action

Antibiotic (Cephalosporin 1st Gen)

Bactericidal - Inhibits bacterial cell wall synthesis by binding to one or more of penicillin-binding proteins which inhibits cell wall biosynthesis, causing bacteria to eventually lyse.

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
- Some cross reactions occur in those with penicillin allergies. Use with caution.

Adverse Reactions I Precautions

- Superinfection prolonged use may result in fungal or bacterial superinfection (including C.Difficile)
- Increased INR (bleeding risk) especially in nutritionally deficient, hepatic/renal disease, prolonged treatment

Dose and Administration: ADULT PEDIATRIC

Infection Control:

Routine dosing may be based on body mass: 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

War wounds (dirty wounds), 2g in 250 mL NS IV over 5 min every 8 hours for 24 hours is adequate for most dirty wounds of the head and neck, torso, and extremities.

IV:

Adults:

- 1-2g every 6-8hrs
 - o Max daily dose: 12 g/day

Note: See antibiotic chart far dosing in accordance with injury.

Infection Control:

IV:

- 20-30 mg/kg IV q 6-8h (maximum, 100 mg/kg/day)
 - Max daily dose: 100 mg/kg/day

Class I Mechanism of Action

Systemic Corticosteroid

Anti-inflammatory, Immunosuppressant Onset of action, IV: Prompt; Duration IV: 72 hours

Indications

Labeled Indications:

- Anti-inflammatory or immunosuppressant in treatment of a variety of diseases: allergic, dermatologic, endocrine, hematologic, inflammatory, neoplastic, renal, respiratory, rheumatic, and autoimmune
- Management if cerebral edema

Unlabeled:

• Treatment of acute mountain sickness (AMS) and high-altitude cerebral edema.

Contraindications

- Hypersensitivity to dexamethasone or any component of the formulation
- Systemic fungal infection, cerebral malaria

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 | PEDIATRIC Always Reference BROSELOW |
| Tape | | |
| Acute mountain sickness (AM | S)High altitude | Acute mountain sickness (AMS)High altitude |
| cerebral edema (HACE) (unla | beled use): | cerebral edema (HACE) (unlabeled use): |
| PO, IM, IV: | | 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
- 0.15 mg/kg dose every 6 hours
 - consider use in high altitude pulmonary edema because of associated HACE with pulmonary edema

DEXTROSE 50% Lactation? Trade Name: Glutose I B-D Glucose

Class I Mechanism of Action

Antidote, Hypoglycemia

Basic source of calories (fuel) for the body and brain, regulated by insulin. Rapidly increases blood glucose, decreases protein and nitrogen loss, preventing ketosis, and promotes glycogen deposition in liver.

Onset of action: Treatment of hypoglycemia Oral dose: 10 minutes Maximum

effect: Treatment of Hyperkalemia IV: 30 minutes

Indications

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 1st.
- Tissue Necrosis if Extravasation occurs; immediately D/C and change IV site.
- Hyperglycemia
- Hypokalemia
- Hyponatremia

| Dose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW |
|--------------------------|-------|-------------------------------------|
| | | Tape |

Hvpoqlvcemia:

Oral:

 4-20 g as a single dose; may repeat if necessary.

IV:

10-25 g (40-100 mL of 25% solution or 20-50 mL of 50% solution)

Note: Society of Critical Care Medicine recommends: Treat blood glucose <70 mg/dL (<100 mg/dL in patients with neurologic injury) immediately by stopping insulin therapy (if receiving) and administering 10-20 g (20-40 mL of 50% solution) IV; repeat blood glucose measurement in 15 minutes with repeat dextrose as needed; **avoiding overcorrection**.

Hvpoglycemia:

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

Tape

Anxietv:

Oral, IV, IM: (Oral and IV doses more reliable)

• 2-10 mg 2-4 times/day if needed.

Status Epilepticus:

IV: (SLOW)

• **5-10 mg** every 5-10 minutes given over 3 minutes (maximum dose: 30 mg)

Sedation in ICU patient:

IV:

Loading dose: 5-10 mg; Maintenance dose:
 0.03-0.1 mg/kg every 30 minutes to 6 hours

Muscle Spasm:

IV:

 Initial: 5-10 mg; then 5-10 mg in 3-4 hours, if necessary. Larger doses may be required if associated with tetanus.

Nerve Agent Exposure (CBRNE)

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 casualty

Sedation | Muscle relaxation | Anxiety:

IV, IM (IV doses more reliable)

 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.

Status Epilepticus:

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.
- Children 25 years: 1 mg given slowly every 2-5 minutes (maximum total dose: 10 mg); repeat in 2-4 hours if needed.

Muscle spasm associated with tetanus:

IV, IM

- Infants >30 days and Children <5 years: 1-2 mg dose every 3-4 hours as needed.
- Children 25 years: **5-10 mg dose** every 3-4 hours as needed

DILTIAZEM

Lactation? (Not Recommended)

Class I Mechanism of Action

Calcium Channel Blocker; Antiarrhythmic Agent, Class IV

Inhibits calcium ion from entering the "slow channels" or select voltage-sensitive areas of vascular smooth muscle and myocardium during depolarization; produces relaxation of coronary vascular smooth muscle and coronary vasodilation; increases myocardial oxygen delivery in patients with vasospastic angina. Onset of action: IV: 3 minutes. Duration 1-3 hours

Indications

Labeled Indications: Atrial fibrillation or atrial flutter for acute ventricular rate control, conversion of supraventricular tachycardia, hypertension, chronic stable angina, vasospastic angina. Unlabeled: Hypertrophic cardiomyopathy; Idiopathic ventricular tachycardia; Nonsustained ventricular tachycardia or ventricular premature beats, symptomatic; Pulmonary arterial hypertension (group 1).

Contraindications

- Sick sinus syndrome (except in patients with a functioning artificial pacemaker); Second- or thirddegree AV block
- Atrial fibrillation or flutter associated with accessory bypass tract (WPW, short PR syndrome)
- Severe hypotension; Cardiogenic shock; Hypersensitivity to diltiazem or any formulation component
- Ventricular tachycardia (with wide-complex tachycardia [ORS 20.12 seconds], must determine whether origin is supraventricular or ventricular)

Adverse Reactions I Precautions

- Cardiovascular: Edema, atrioventricular block, bradycardia, hypotension, dyspnea
- Central nervous system: Headache, dizziness, pain, nervousness, vomiting, weakness, myalgia

Dose and Administration: ADULT

PEDIATRIC Always Reference BROSELOW

Trade Name: Cardizem

Tape

Atrial fibrillation or atrial flutter, rate control: Note: For rate control in hemodynamically stable patients. Do not use in patients with preexcitation associated with an accessory pathway, as this can lead to ventricular arrhythmias.

- Bolus dose: 0.25 mg/kg over 2 minutes (average dose: 20 mg); if rate control is insufficient after 15 minutes, a repeat bolus dose of 0.35 mg/kg over 2 minutes may be given (average dose: 25 mg). Patients who respond after 1 or 2 bolus doses can be started on a continuous infusion.
- Continuous infusion following bolus(es): Initial: 5 to 10 mg/hour; infusion rate may be increased in 5 mg/hour increments according to ventricular response, up to a maximum of 15 mg/hour.

Supraventricular tachvcardia (alternative agent):

Note: For hemodynamically stable patients if vagal maneuvers and/or adenosine are unsuccessful.

Bolus dose: **0.25 mg/kg** (actual body weight) over 2 minutes (average dose: 20 mg); if rate control is insufficient after 15 minutes, a repeat bolus dose of **0.35 mg/kg** over 2 minutes may be given (average dose: 25 mg). If bolus(es) do not terminate the arrhythmia, alternative therapy.

Atrial tachvarrhythmias, rate control:

Very limited data available: Infants, Children, and Adolescents

IV:

- Initial bolus: 0.25 mg/kg over 5 minutes (maximum dose: 20 mg/dose [average adult dose]) followed by a continuous IV infusion. Dose should be individualized based on patient response.
- Continuous infusion (titrated to effect): 0.05 to 0.15 mg/kg/hour (Rate control achieved =10min)

DIPHENHYDRAMINE QB, Lactation Yes (Unsafe) Trade Name: **Benadryl**

Class / Mechanism of Action

Histamine H₁ Antagonist:

Competes with histamine for H1-receptor sites within the gastrointestinal tract, blood vessels, and respiratory tract; Also produces anticholinergic and sedative effects

Indications

Labeled Indications:

- Anaphylaxis and allergy disorders
- Motion Sickness
- Antitussive

Contraindications

- Hypersensitivity to diphenhydramine or any component of the formulation
- Acute Asthma
- Use on Neonates, premature infants, Nursing mothers

Adverse Reactions / Precautions

- Normally causes sedation but may cause paradoxical excitation in children.
- May have increased sedative effects when used with other sedatives or alcohol.
- May cause hypotension (use with caution in patient with cardiovascular disease)
- Dry mouth

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:

 25-50mg once, administered after epinephrine for anaphylaxis Note: Diphenhydramine is not the 1st line medication for anaphylaxis

Acute Hemolytic reaction

(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

Motion Sickness:

Oral:

0.5-1 mg/kg every 6 hours

IV/IM:

• 1.25mg/kg every 6 hours

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 |
|--------------------------|-------|--|
| | | |

Cardiac 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/mL

Infusion Rates for Dobutamine at 1000mcg/mL

| Desired Delivery Rate | Infusion Rate |
|-----------------------|----------------------|
| (mcg/kg/min) | (mL/kg <u>hour</u>) |
| 2.5 | 0.15 |
| 5 | 0.3 |
| 7.5 | 0.45 |
| 10 | 0.6 |
| 12.5 | 0.75 |
| 15 | 0.9 |
| 20 | 1.2 |

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> after adequate fluid volume replacement

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 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):

 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.

Dopamine Dosage Efficacy:

- Renal- 1-5 mcg/kg/min= Dopaminergic effects: increased urine output, increased renal blood flow
- 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. (*phenylephrine*, *norepinephrine*, *epinephrine*.)

Hemodynamic Support:

IV:

 2-20mcg/kg/min; titrate to desired response. Infusion may be increased by 5-10mcg/kg/minute until optimal response is obtained.

Note: Dopamine is a second line medication for hemodynamic support in Pediatric patients behind Epinephrine and Norepinephrine

| Dopamine Dosing Range: 5-20mcg/kg/min (300-1200mcg/kg/hr) | | | | | Dopamine | | | | | | | | |
|---|--|------------|-----------|------------|-------------------|----------------------|--|-----------|---------------------------|---------------|---------|---------|---------|
| | Dosing Rar | nge: 5-20r | ncg/kg/mi | n (300-120 | 0mcg/kg/l | 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 | | | | | | C | CONCENT | RATION 1 | 600 mcg/n | ıL | | |
| 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 |
| 00 | 15 | 28 | 28 | 9 | 7 | 5 | | 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 |
| 33 | 15 | 31 | 31 | 10 | 8 | 5 | | 15 | 48 | 48 | 16 | 12 | 8 |
| | 20 | 41 | 41 | 14 | 10 | 7 | | 20 | 64 | 64 | 21 | 16 | 11 |
| | 5 | 11 | 11 | 4 | 3 | 2 | | 5 | 17 | 17 | 6 | 4 | 3 |
| 60 | 10 | 23 | 23 | 8 | 6 | 4 | 90 | 10 | 34 | 34 | 11 | 9 | 6 |
| 00 | 15 | 34 | 34 | 11 | 9 | 6 | 30 | 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 |
| G.F. | 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 |
| 7.5 | 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 |
| | Micro-Drip is set of choice for this infusion | | | | | T': | | | | ce for this i | | | |
| Titr | Titrate to minimum effective dose. Allow 3-5 minutes between dosing changes to assess hemodynamic effects. | | | | i itra | ate to minin chan | | dosing | Allow 3-5 m dynamic ef | | ween | | |

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.
- Hypersensitivity to sympathomimetic amines, glaucoma and non-anaphylactic shock

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.

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 | | | | | |
| Dose | Rate | Micro | | Macro | | | | |
| | | 60 gtt/mL | 20 gtt/mL | 15 gtt/mL | 10 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 | 15 450 450 150 113 75 | | | | | | | |
| Mad | cro-Drip (1 | 0gtt/ml) is s | set of choice | ce for this i | nfusion | | | |
| St | art at lowe | st dose an | d titrate to | desired eff | ect | | | |

EPINEPHRINE QC, Lactation? (Caution) 1:10,000 Trade Name: Adrenalin

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:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Asystole/pulseless arrest, pulseless VT/VF (ARC ALS 2020):

IV: 1mg/10mL (0.1mg/mL) pre-filled 10cc Syringe

 1 mg (10cc of 0.1mg/mL) every 3-5 minutes to ROSC, Follow each dose with 20mL flush

<u>Drip-Dose: Bradycardia (Symptomatic).</u> <u>Hypotension-Fluid Resistant Shock:</u>

IV Continuous Infusion:

- Bradycardia: 2-10 mcg/minute titrate to desired effect (HR >60, MAP >65)
- Hypotension/Shock: 2-10 mcg/min titrated to clinical end point (BP, end organ perfusion)

Push-Dose: Refractory Anaphylaxis:

IV/IO: Utilize the 0.1 mg/mL solution further diluted in 10mL of NS.

 0.05-0.1 mg, administered over 1-10 minutes, may repeat once after 3 minutes if patient remains unresponsive to initial dose.

Asystole, PEA, pulseless VT/VF, Unresponsive and Symptomatic Bradycardia in Infants (ARC ALS 2020):

IV:

0.01 mg/kg (0.1 mL/kg of 1mg/10mL [0.1 mg/mL]) (maximum single dose: 1 mg) every 3-5 minutes as needed or until ROSC

Severe Hypotension/shock and fluid resistant (unlabeled use):

IV: Continuous Infusion

• 0.1 - 1 mcg/kg/minute titrated to desired effect

| TABLE OF CONTENTS | | | | | | | | |
|---------------------------------|---|-------------|--------------|---------------|-----------|--|--|--|
| | Epinephrine 1mg/10ml (1:10,000) | | | | | | | |
| Pressor for Hypotension | | | | | | | | |
| Dosing Range: 2-20mcg/min (120- | | | | | | | | |
| | | | ncg/hr) | | | | | |
| | | | g/500 mL | | | | | |
| | | | TION 2 mc | | | | | |
| Dose | Rate | Micro | | Macro | | | | |
| | | (60 | 20 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 | | | |
| Ma | cro-Drip (1 | 0gtt/ml) is | set of choic | e for this ir | nfusion | | | |
| | Start at lowest dose and titrate to desired | | | | | | | |

Start at lowest dose and titrate to desired effect

ERTAPENEM QC, 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 penicillin-binding 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 hours of coverage.

IV:

Adults:

• 1 g IV every 24 hrs. for 7-14 days.

<u>Infection Control:</u> Children <12 years & ≥12 years

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 used for induction of anesthesia. Onset of action: 30-60 seconds, Duration 5-10 minutes

Indications

Labeled Indications:

• Rapid Sequence Induction

Contraindications

- Hypersensitivity to etomidate or any component of the formulation
- Labor/Delivery
- Septic Schock

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:

 0.2-0.4 mg/kg over 30-60 seconds for induction of anesthesia.

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.

RSI:

IV:

- **0.2-0.4 mg/kg** over 30-60 seconds will produce rapid sedation lasting 10-15 minutes.
 - o 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.

FENTANYL QC, Lactation Yes (not recommended) Trade Name:Sublimaze

Class / Mechanism of Action

Opioid Analgesic; General Anesthetic

Binds to opioid receptors within the CNS increasing pain threshold and altering pain reception; inhibits ascending pain pathways (blocking painful stimulus); produces CNS depression.

Onset: IV almost immediate, Duration: IV 0.5-1 hour

Indications

Labeled Indications:

- Pain relief
- Adjunct to general or regional anesthesia

Contraindications

- Hypersensitivity to fentanyl or any component of the formulation
- MAOI taken in the past 14 days.
- Hypotension
- Hypoxia
- Hypoventilation

Adverse Reactions / Precautions

- When using only as pain med and not adjunct to general anesthesia, ensure Slow IV Push (3-5 min). Rapid infusion may result in chest wall rigidity, impaired ventilation, or respiratory distress/arrest Always be prepared for use of paralytic and intubation (positive control of airway).
- 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, fatique.

Dose and Administration: ADULT PEDIATRIC

Pain Management:

IV: Slow (Unlabeled)

- 0.5-1mcg/kg PRN for breakout pain q 30-60 min IN/IM:
- 1mcg/kg mcg

Note: Patients with prior opioid exposure may have increased tolerance and require higher dosing

Sedation 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

Pretreatment for RSI:

3-5 min prior to RSI in pt's with Head injuries, Increased ICP, Cardiac Ischemia or Aortic Dissection (if situation allows):

3mcq/kq slow IV push

Non-Traumatic Chest Pain (Cardiac)

• 25-50mcg IV

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 mortality due to adrenal suppression and inability to respond to stress.

| FENTANYL (SUBLIMASE) | | | | | | | | | |
|-------------------------------|---------------------------------|----------------|--|-----------|-----------|--|--|--|--|
| Dosing Range: 0.5-1 mcg/kg/hr | | | | | | | | | |
| | MIX 1 mg/100 mL | | | | | | | | |
| | С | ONCENTRA | TION 10 m | _ | | | | | |
| 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 | 14 | 5 | 3 4 | 2 | | | | |
| 140 | | 14 | 5 | | 2 | | | | |
| 145 | 15 15 | 15 15 | 5 5 | 4 | 3 | | | | |
| 150 | | | 5 | 4 | 3 | | | | |
| 155 | 16 16 | 16 16 | 5 | 4 | | | | | |
| 160 165 | 16 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 | | | | |
| 200 | | -Drip is set o | The state of the s | | | | | | |
| | | | | | OI I | | | | |
| | Sample patient: 80kg pt at 0.5- | | | | | | | | |

Sample patient: 80kg pt at 0.5-1mcg/kg/hr = 40mcg/hr-80mcg/hr dosing range

FUROSEMIDE

C, Lactation Yes (Caution)

Class / Mechanism of Action

Antihypertensive; Loop Diuretic

Inhibits reabsorption of sodium and chloride in the kidney, causing increased loss of water, sodium, chloride, magnesium, and calcium within urine. When given IV it also causes rapid venous dilation. Symptomatic improvement of acute pulmonary edema approximately 15-20 minutes

Indications

Labeled Indications: Management of edema associated with heart failure and hepatic or renal disease.

- Management of edema associated with heart failure and hepatic or renal disease; acute pulmonary edema.
- Hypertension (alone or in combination with other antihypertensives)

Contraindications

- Hypersensitivity to furosemide or any component of the formulation
- Anuria (No pre-hospital utility in hypovolemic shock)

Adverse Reactions / Precautions

- Can cause profound diuresis with resulting shock and electrolyte depletion. Monitor closely!
 - o May cause: Hypovolemia, Hypotension, hyponatremia, hypokalemia
- May potentiate effect of additional antihypertensives

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Lasix

Acute pulmonary edema:

IV

 40 mg over 1-2 minutes. If response not adequate within 1 hour, may increase dose to 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/dose and 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 infusion doses of 10-40 mg/hour. If urine output is <1 mL/kg/hour, double as necessary to a maximum of 80-160 mg/hour.

Edema. heart failure: Infants and Children

IV, IM:

 Initial: 1 mg/kg/dose; if response not adequate, may increase dose in increments of 1 mg/kg/dose and administer not sooner than 2 hours after previous dose, until a satisfactory response is achieved; may administer maintenance dose at intervals of every 6-12 hours; maximum dose: 6 mg/kg/dose

GLUCAGON

PB Lactation? (Caution)

Class / Mechanism of Action

Antidote, Hypoglycemia Antidote, Diagnostic agent

Hyperglycemic agent, pancreatic hormone, insulin antagonist; Raises blood glucose levels by stimulating increased production of cyclic AMP, promoting hepatic glycogenolysis and gluconeogenesis

Indications

Labeled Indications: Management of hypoglycemia (Glucose <70 in adults or <60 in children) **Unlabeled:**

- Beta-blocker or calcium channel blocker induced myocardial depression (with or without hypotension) unresponsive to standard measures.
- Hypoglycemia secondary to insulin or sulfonylurea overdose (as adjunct to dextrose)

Contraindications

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

Adverse Reactions / Precautions

- Should NOT be used as 1st line medication for hypoglycemia or Altered mental status
 - Hypoglycemia patients should receive dextrose. If IV access cannot be established or if dextrose is not available, glucagon may be used as alternate until dextrose can be given.
- Thiamine should precede use in patient with suspected alcoholism or malnutrition

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Hypoglycemia:

IV, IM, IN, SubQ:

1 mg; may repeat in 20 minutes as needed.

<u>Beta-blocker / Calcium channel blocker</u> <u>overdose (myocardial depression)</u> 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 QC, Lactation No Trade Name:

Class / Mechanism of Action

Anticoagulant

Inactivates thrombin and activated coagulation factors (IX, X, XI, XII, and plasmin) and prevents conversion of fibrinogen to fibrin.

Indications

Labeled Indications: Treatment of thromboembolic disorders.

Unlabeled: ST elevation MI (STEMI) as an adjunct to thrombolysis; unstable angina/non-STEMI

Contraindications

- Hypersensitivity to heparin or any component of the formulation
- Active Bleeding (Trauma Patient)

Adverse Reactions / Precautions

- Continuously monitor for bleeding: Stop immediately if any bleeding occurs.
- Urticarial reactions and anaphylaxis can occur

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

<u>Acute coronary syndromes: STEMI/Unstable</u> <u>Angina as an adjunct to fibrinolysis (full dose</u> alteplase:

IV:

- Initial bolus of 60 units/kg (MAX: 4000 units)
 - Maintenance: 12 units/kg/hour (MAX: 1000 units/hour) as continuous infusion.

Treatment of venous thromboembolism:

IV: (unlabeled dosing)

- <u>DVT/PE:</u> 80 units/kg or 1mg/kg (or alternatively 5000 units) IV push followed by continuous infusion of 18 units/kg/hour.
- COVID VTE Prophylaxis: 7,500 iu SQ every 8 hours

Note: Heparin is ONLY for use only under written direction of referring provider or direct consultation with medical director.

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

| Н | ETASTARCH | ♀C, Lactation \ | es (caution) | Trade Name: Hextend |
|-----|--|------------------------|-----------------------------|-----------------------------|
| CI | ass / Mechanism of Action | , | | |
| | asma Volume Expander, Colle | | Onest of Astions commercia | mataly 20 mainsuta a |
| | blloidal starch producing plasma | volume expansion | . Onset of Action: approxir | nately 30 minutes |
| | dications | 1 1 1 | | |
| | beled Indications: Volume exp | ander used in trea | tment of hypovolemic / her | morrhagic shock |
| | ontraindications | | | |
| • | Hypersensitivity to hydroxyeth | | | 1 |
| • | Renal failure with oliguria and Fluid overload conditions, (pul | • | , | |
| | Pre-existing bleeding or coagu | • | • | e). Hee caution in bleeding |
| | disorders; may increase risk o | | g, von villebrand 3 diseas | e). Ose caution in bleeding |
| Ac | Iverse Reactions / Precautions | | | |
| • | Anaphylactoid reactions (aller | gies to corn) | | |
| Do | ose and Administration: | ADULT | PEDIATRIC Alwa | ays Reference BROSELOW Tape |
| Pla | asma volume expansion: | | | |
| IV | . | | | |
| • | 250-500ml Bolus. May repeat | PRN (up to | | |
| | 1500 mL/day). Titrate to indivi | dual ` . | | |
| | hemodynamic needs (Sys BP | >90). | | |
| No | otes: | | | |
| • | May be administered via infus pressure infusion. | ion pump or | | |
| • | Do not administer with blood t same line / tubing | nrough the | | |
| • | Change tubing or flush extens before administering blood thr same line. | | | |
| • | Use only for burns as an adjur preferred until hour 8-12; may difficult resuscitation. | | | |

Class / Mechanism of Action

Opioid Analgesic

Binds to opioid receptors within the CNS increasing pain threshold and altering pain reception; inhibits ascending pain pathways (blocking painful stimulus); produces CNS depression Onset: IV 10-20 minutes. Duration 2-4 hours

Indications

Labeled Indications: Moderate to severe pain.

Contraindications

- Hypersensitivity to hydromorphone or any component of the formulation
- Severe respiratory depression (in absence of resuscitative equipment or ventilator support)
- · 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 paralytic and intubation (maintain positive control of airway).
- Head trauma: Use with extreme caution in head injury, or suspected increased ICP;
 exaggerated increase in ICP may occur.
- May cause Hypotension, Use with caution in hypovolemic patients.
- May cause life-threatening Reparatory depression
- CNS depression: Impairs physical and mental abilities
- Dizziness
- Headache
- Syncope

| • | Syncope | | |
|----|--|---------|--|
| Do | ose and Administration: | ADULT | PEDIATRIC Always Reference BROSELOW Tape |
| A | cute pain (moderate-to-severe): | | Acute pain (moderate-to-severe): |
| IV | : (Slow) | | IV: (Slow) |
| • | 0.5mg (range 0.25-2mg) IV/IO q 1 prn | -6hr as | Children: 0.015mg/kg IV q 4-6 PRN Adolescents >50kg: Refer to adult |
| • | Critically ill require lower dose, o tolerant may require higher dose | | , and the second |
| • | Continuous infusion: Usual dosage 0.5- 3 mg/hour (See infusion cha page) | | |

| HYDROMORPHONE (D | DILAUDID) |
|------------------|-----------|
|------------------|-----------|

Dosing Range: 0.5-3mg/hr (8.3-50mcg/min)

MIX 2 mg/100 mL CONCENTRATION 20 mcg/mL

| 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-Drip (20gtt/ml) or Micro-Drip is set of choice for this infusion

Start at lowest dose and increase rate by 0.5mg/hr PRN for appropriate pain management

HYDROXOCOBALAMIN

QC, Lactation? (Caution)

Class / Mechanism of Action

Antidote: Vitamin

Precursor to Vitamin B₁₂ (cyanocobalamin). Binds cyanide ion to form nontoxic cyanocobalamin which is excreted within urine

Indications

Labeled Indications:

- IM: Treatment of pernicious anemia and B12 deficiencies
- IV: (Cyanokit®) Treatment of known or suspected cyanide poisoning

Contraindications

• No contraindications when treating for suspected or known cyanide poisoning

Adverse Reactions / Precautions

- May cause transient hypertension (≥180mmHg systolic, ≥110mmHg diastolic)
- Will cause persistent red colored urine and skin, rendering pulse oximetry values inaccurate

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Cyanokit®

Cyanide Poisonings:

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

<u>Smoke Inhalation / Fire victims:</u> (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

Cvanide Poisonings:

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.

<u>Smoke Inhalation / Fire victims: (Closed space exposure with evidence of airway injury: soot in mouth / nose / sputum)</u>

 May present with both cyanide and carbon monoxide poisoning. Hydroxocobalamin is the agent of choice for treating cyanide toxicity in this setting.

KETAMINE

C, Lactation Yes (Risk not ruled out)

Trade Name: Ketalar

Class / Mechanism of Action

General Anesthetic

Dissociative anesthetic; produces a cataleptic-like state acting directly on the cortex and limbic system. Onset of action IV: 30-60 seconds; Duration is dose dependent averaging 10-20 minutes

Indications

Labeled Indications: Induction and maintenance of general anesthesia

Unlabeled: Analgesia and sedation

Contraindications

- Hypersensitivity to ketamine or any component of the formulation
- Conditions that cannot tolerate sustained increases in blood pressure (non-traumatic intracerebral hemorrhage, hypertension associated with acute coronary syndrome; NOT contraindicated in TBI)
- Children <3 mo. age

Adverse Reactions / Precautions

- Rapid IV administration may cause hypotension, apnea, or laryngospasm. Large doses (>1 mg/kg) may cause hypotension and respiratory depression
- Use with caution in patients with cardiovascular disease. Continuously monitor cardiac function
- Preferred general anesthetic / sedative (pre-hospital) for head injury patient (does not raise ICP)
- Dosing between 0.5-0.9 mg/kg IV (and equivalent IM dose) can give patients the feeling of unreality leading to agitation and should be avoided

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

LOW DOSE:

Analgesia:

IV/IO Push (over 1 min)

- **0.1-0.3 mg/kg,** repeat q 10-30 prn IM/IN
- 0.5 1.0 mg/kg, repeat q 10-30 prn
 HIGH DOSE:

RSI / Induction of anesthesia; Combative Patients:

IV Push

1-2 mg/kg

IM

4-5 mg/kg

Maintenance of anesthesia:

IV:

- **0.5-2 mg/kg** dose every 10-20 minutes IV Continuous Infusion
- 0.5-2mg/kg bolus then 1-3 mg/kg/hr. Titrate levels by 0.25mg/kg/hr. PRN to achieve appropriate sedation. (See infusion chart next page)

Analgesia:

IM:

0.5 mg/kg, repeat q 10-30 prn

IV:

• 0.1 - 0.2 mg/kg, repeat q 10-30 prn

Induction of anesthesia (unlabeled dosing):

IV:

1-2 mg/kg (3-5mg/kg for procedural sedation)

Maintenance of anesthesia:

IV:

- ½ to Full induction dose every 20-30 minutes IV Continuous Infusion:
- 0.5-1 mg/kg/hr. Titrate levels by 0.25mg/kg/hr PRN to achieve appropriate sedation.

NOTE Avoid sub-dissociative doses to prevent emergence phenomenon.

NOTE: If patient experiences Ketamine Induced Agitation (Emergence Phenomena) give Midazolam 2-5mg IV x1 for adults and 0.05 mg/kg for children not hypotensive or in danger of being

| | KETAMINE (KETALAR) | | | | | | | |
|------------|--------------------|------------|--------------------|------------|-------------|-----------|--|--|
| | Dosing | Range: 1- | 3mg/kg/hr (1 | 17-50mcg/l | (g/min) | | | |
| | | | 00 mg/500 n | | | | | |
| | | CONC | ENTRATION mg/mL | N 1 | | | | |
| Pt. Weight | Dose | Rate | Micro | | Macro | | | |
| | | | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL | | |
| kg | mcg/kg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | |
| | 15 20 | 45 60 | 45 60 | 15 20 | 11 15 | 8 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 | 34 | 23 | | |
| | 55 | 165 | 165 | 50 55 | 38 41 | 25 28 | | |
| | 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 35 | 99 116 | 99 116 | 33 39 | 25 29 | 17 19 | | |
| 55 | 40 | 132 | 132 | 44 | 33 | 22 | | |
| | 45 | 149 | 149 | 50 | 37 | 25 | | |
| | 50 | 165 | 165 | 55 | 41 | 28 | | |
| | 55 | 182 | 182 | 61 | 46 | 30 | | |
| | 60 15 | 198 54 | 198 54 | 66 18 | 50 14 | 33 9 | | |
| | 20 | 72 | 72 | 24 | 18 | 12 | | |
| | 25 | 90 | 90 | 30 | 23 | 15 | | |
| | 30 | 108 | 108 | 36 | 27 | 18 | | |
| 60 | 35 | 126 | 126 | 42 | 32 | 21 | | |
| 00 | 40 | 140 | 140 | 47 | 35 | 23 | | |
| | 45 50 | 162 180 | 162 180 | 54 60 | 41 45 | 27 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 30 | 98 117 | 98 117 | 33 39 | 25 29 | 16 20 | | |
| | 35 | 137 | 137 | 46 | 34 | 23 | | |
| 65 | 40 | 156 | 156 | 52 | 39 | 26 | | |
| | 45 | 176 | 176 | 59 | 44 | 29 | | |
| | 50 55 | 195 215 | 195 215 | 65 72 | 49 | 33 36 | | |
| | 60 | 234 | 234 | 78 | 54 59 | 39 | | |
| | 15 | 63 | 63 | 21 | 16 | 11 | | |
| | 20 | 84 | 84 | 28 | 21 | 14 | | |
| | 25 | 105 | 105 | 35 | 26 | 18 | | |
| | 30 | 126 | 126 | 42 | 32 | 21 | | |
| 70 | 35 40 | 147 168 | 147 168 | 49 56 | 37 42 | 25 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 | | |
| | 25 | 113 | 113 | 38 | 28 | 19 | | |
| | 30 | 135 | 135 | 45 | 34 | 23 | | |
| 75 | 35 | 158 | 158 | 53 | 40 | 26 | | |
| 75 | 40 | 180 | 180 | 60 | 45 | 30 | | |
| | 45 50 | 203 | 203 | 68 | 51 | 34 | | |
| | 50 55 | 225 248 | 225 248 | 75 83 | 56 62 | 38 41 | | |
| | 60 | 270 | 270 | 90 | 68 | 45 | | |
| | | | et of choice for | | | | | |
| Sam | nple patient: 80 | | | | | ml/hr\ | | |
| Sall | ipic patietit. 00 | ong prar I | omg/kg/III = | JU-Z4UIIIQ | iii (00-240 | 1111/111) | | |

| | KETAMINE (KETALAR) Dosing Range: 1-3mg/kg/hr (17-50mcg/kg/min) | | | | | | | | |
|-------------|---|-------------|----------------|------------|--------------------------|-----------|--|--|--|
| | MIX 500 mg/500 mL CONCENTRATION 1 | | | | | | | | |
| Pt. Weight | Dose | Rate | mg/mL Micro | | Macro | | | | |
| i t. Weight | Dose | rate | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/mL | | | |
| kg | mcg/kg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min | | | |
| | 15 20 | 72 96 | 72 96 | 24 32 | 18 24 | 12 16 | | | |
| | 25 | 120 | 120 | 40 | 30 | 20 | | | |
| | 30 | 144 | 144 | 48 | 36 | 24 | | | |
| 80 | 35 40 | 168 192 | 168 192 | 56 64 | 42 | 28 32 | | | |
| | 45 | 216 | 216 | 72 | 54 | 36 | | | |
| | 50 | 240 | 240 | 80 | 60 | 40 | | | |
| | 55 | 264 | 264 | 88 | 66 | 44 | | | |
| | 60 15 | 288 77 | 288 77 | 96 26 | 72 19 | 48 13 | | | |
| | 20 | 102 | 102 | 34 | 26 | 17 | | | |
| | 25 | 128 | 128 | 43 | 32 | 21 | | | |
| | 30 | 153 | 153 | 51 | 38 | 26 | | | |
| 85 | 35 40 | 179 204 | 179 204 | 60 68 | 45 51 | 30 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 | | | |
| | 25 | 135 | 135 | 45 | 34 | 23 | | | |
| | 30 | 162 | 162 | 54 | 41 | 27 | | | |
| 90 | 35 | 189 | 189 | 63 | 47 | 32 | | | |
| 30 | 40 45 | 216 243 | 216 243 | 72 81 | 54 61 | 36 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 25 | 114 143 | 114 143 | 38 48 | 29 36 | 19 24 | | | |
| | 30 | 171 | 171 | 57 | 43 | 29 | | | |
| 05 | 35 | 200 | 200 | 67 | 50 | 33 | | | |
| 95 | 40 | 228 | 228 | 76 | 57 | 38 | | | |
| | 45 50 | 257 285 | 257 285 | 86 95 | 64 71 | 43 48 | | | |
| | 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 30 | 150 180 | 150 180 | 50 60 | 38 45 | 25 30 | | | |
| | 35 | 210 | 210 | 70 | 53 | 35 | | | |
| 100 | 40 | 240 | 240 | 80 | 60 | 40 | | | |
| | 45 | 270 | 270 | 90 | 68 | 45 | | | |
| | 50 55 | 300 | 300 330 | 100 110 | 75 83 | 50 55 | | | |
| | 60 | 360 | 360 | 120 | 90 | 60 | | | |
| | 15 | 95 | 95 | 32 | 24 | 16 | | | |
| | 20 | 126 | 126 | 42 | 32 | 21 | | | |
| | 25 | 158 | 158 | 53 | 40 | 26 | | | |
| | 30 35 | 189 221 | 189 221 | 63 74 | 47 55 | 32 37 | | | |
| 105 | 40 | 252 | 252 | 84 | 63 | 42 | | | |
| | 45 | 284 | 284 | 95 | 71 | 47 | | | |
| | 50 | 315 | 315 | 105 | 79 | 53 | | | |
| | 55 60 | 347 | 347 | 116 | 87 | 58 | | | |
| | 60 378 378 126 95 63 Macro-Drip is set of choice for this infusion | | | | | | | | |
| | 111GOI | - 2 | | | | | | | |
| San | nple patient: 80 | Okg pt at 1 | -3mg/kg/hr = | 80-240mg | <mark>/hr (80-240</mark> | ml/hr) | | | |

KETOROLAC

QC, Lactation Yes (Caution)

Class / Mechanism of Action

Nonsteroidal Anti-inflammatory Drug (NSAID)

Inhibits cyclooxygenase (COX 1 & 2) enzymes, which decreases production of prostaglandin precursors. Provides antipyretic, analgesic, and anti-inflammatory action.

Indications

Labeled Indications: Short term management of moderate to severe acute pain as an opioid alternative.

Contraindications

- Hypersensitivity to ketorolac, aspirin, other NSAIDs, or any component of the formulation.
- High risk of bleeding, recent history of GI bleeding or perforation, known history of peptic ulcer disease.
 - Not for use as pain management for battlefield trauma patient!
- Suspected cerebrovascular bleeding
- Risk of renal failure secondary to volume depletion
- Concurrent use with other NSAIDs

Adverse Reactions / Precautions

- Inhibits platelet function
- Associated with an increased risk of adverse cardiovascular thrombotic events, including MI and stroke
- May increase risk of GI irritation, inflammation, ulceration, bleeding, and perforation.
- May cause severe bronchospasm in patients with asthma
- May cause new onset hypertension or worsening of existing hypertension.

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Toradol

Pain management (acute; moderately severe):

Patients ≥50 kg

IM:

 15-30 mg every 6 hours (maximum daily dose: 120 mg)

IV:

 15 mg every 6 hours (maximum daily dose: 120 mg)

Adults <65 years and/or adults <50 kg

IM:

• **15-30 mg** every 6 hours (maximum daily dose: 60 mg)

IV:

• **15 mg** every 6 hours (maximum daily dose: 60 mg)

Pain management (acute; moderately severe):

Children 2-16 y/o:

- Moderate discomfort: 1 mg/kg IM
- Severe discomfort: 0.5 mg/kg IV
- Febrile seizure: 1 mg/kg IV

Adolescents >17:

· Refer to adult dose

LABETALOL QC, Lactation Yes (Caution) Trade Name: Trandate

Class / Mechanism of Action

Beta Blocker with alpha blocking activity

Blocks alpha and beta1/beta2 adrenergic receptor sites. Onset IV: 2-5 minutes

Indications

Labeled Indications: Treatment of hypertension.

IV: Treatment of severe hypertension and hypertensive emergencies

Unlabeled:

 Pre-eclampsia and severe hypertension in pregnancy, hypertension during acute ischemic stroke, and Pediatric hypertension

Contraindications

- Hypersensitivity to labetalol or any component of the formulation
- Bradycardia <60bpm, heart block >1st degree
- Uncompensated heart failure, Cardiogenic shock
- Asthma

Adverse Reactions / Precautions

- Symptomatic hypotension with or without syncope, Monitor EKG closely
- Use with extreme caution in patients with compensated heart failure and Bradycardia
- Patient with bronchospastic diseases (reactive airway) should not use Beta blockers

Dose and Administration: Tape

ADULT

PEDIATRIC Always Reference BROSELOW

Acute Hypertension (hypertensive

emergency/urgency: Hypertensive Crisis (Sys: >185/Dia: >110)

Intermittent IV:

10-20 mg IV over 1-2 minutes. May repeat one time.

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

*Hypertension emergencies:

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.

*Not 1st Line medication for children

Pearls:

For inter-facility transports with confirmed Ischemic CVA, Intraparenchymal Hemorrhagic CVA, or Spontaneous Non-traumatic SAH manage Hypertension according to diagnosis or guidance from sending facility

- Ischemic CVA Lytic ineligible: SBP < 220 and DBP <120
- 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 ÇC, 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 | PEDIATRIC Always Reference BROSELOW Tape |
| Traumatic brain injury (severe a seizure prophylaxis): Loading dose: 1500 mg over 1 Maintenance dose: 1000 mg of 12 hours | 5 minutes | Status epilepticus, refractory: (Limited data available) 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)

- 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 (2nd or 3rd) 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 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 1-4mg/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
 - o 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.

Decompression Illness/ Arterial Gas Embolism:

• 1.5mg/kg IV/IO

VF/Pulseless VT, Wide Complex Tachycardia (with 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 therapy)

LORAZEPAM QD, Lactation Yes (not recommended) Trade Name: Ativan

Class / Mechanism of Action

Benzodiazepine

Acts as an Anxiolytic/Hypnotic, anticonvulsant and sedative.

Onset of action: IV Sedation 2-3 minutes; IM hypnotic, 15-30 minutes. Duration: IV, 8-12 hours.

Indications

Labeled Indications: Anesthesia premedication, Status epilepticus **Unlabeled:**

- Rapid tranquilization of the combative / agitated patient
- Alcohol withdrawal delirium / syndrome
- Seizures
- Induce Sedation and Amnesia (Midazolam is primary medication)

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 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: 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:

0.5-2 mg slow IV push

Rapid tranquilization of agitated / combative patient (Off-label use):

IV, IM:

• **2-4mg** every 30-60 minutes; may be used alone or administered with an antipsychotic (i.e., haloperidol)

<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.

Agitation:

IV/IM

• **0.02-0.1 mg/kg/dose** q 20-30 min PRN. (Maximum dose: 2mg/dose).

MAGNESIUM SULFATE

QD, 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 pregnancies with severe pre-eclampsia or eclampsia
- Torsades de Pointes: Cardiac arrhythmias (VT/VF) cause by low serum magnesium

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 depression, and hypotension in large doses
- Hypomagnesaemia is often joined by hypokalemia and requires correction in order to normalize potassium.

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

<u>Torsades de pointes or VF/pulseless VT</u> <u>associated with torsades de pointes (unlabeled</u> 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.

Wheezing in Respiratory Distress (3rd line drug):

IV:

2 Grams single dose over 20min

Seizure (Refractory to Benzodiazepines):

IV/·

• **1-2 Grams** over 30 min

Eclampsia/pre-eclampsia, severe (unlabeled):

IV:

• **4-6 g** over 15-30 minutes followed by 1-2 g/hour continuous infusion for 24 hrs

Torsades de pointes: (PALS 2020)

IV, IO:

• 25-50 mg/kg/dose over several minutes

o maximum single dose: 2000 mg

Respiratory Distress (Status Asthmaticus):

IV:

• 25-50 mg/kg over 30 min (max 2 grams)

Magnesium Sulfate should be diluted into 50-100ml NS or D5W for all Adult and Pediatric infusions

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 <u>under-resuscitated or hypotensive casualties</u>)

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.

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Moderate to severe head injury, Patient continuing to deteriorate or showing signs of herniation despite adjustment to ventilation and starting hypertonic saline.

IV

- 1 g/kg IV bolus over <20 minutes.
- 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

<u>Increased intracranial pressure</u> (unlabeled <u>dosing):</u>

IV:

- 0.25-1 g/kg/dose.
- Maintenance dose of 0.25-0.5 g/kg IV q 4-6hrs prn to maintain serum osmolality <300-320 mOsm/kg

METHYLPREDNISOLONE QC. Lactation Yes(Caution) Trade Name: SoluMedrol

Class / Mechanism of Action

Systemic Corticosteroid

Anti-inflammatory, Immunosuppressant, shock

Indications

Labeled Indications: Treatment of a variety of diseases: allergic, inflammatory, hematologic, neoplastic, and autoimmune;

Unlabeled:

None identified unless added by medical direction.

Contraindications

- Hypersensitivity to methylprednisolone or any component of the formulation
- No other in emergency setting

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

<u>Asthma exacerbations, including status</u> asthmaticus

IV:

• **125mg** x 1 dose

Allergic Reaction:

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• 125mg x 1 dose

can be used for IV doses.

Note: Only methylprednisolone sodium succinate

<u>Asthma exacerbations, including status</u> 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 x 1 dose

Note: Only methylprednisolone sodium succinate can be used for IV doses.

METOCLOPRAMIDE ₽В 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: ADULT** PEDIATRIC Always Reference BROSELOW Tape Not recommended or approved for routine 5-10 mg IV/IO slow push; repeat dose x1 every 20pediatric use 30 minutes PRN for a max of 20mg.

Ideal administration to Prevent Agitation/Adverse Effects: Dilute in a 50 or 100ml NS bag and infuse over 10-15 minutes.

METOPROLOL

QC, Lactation?(Not Recommended)

Class / Mechanism of Action

Beta-1 Selective Beta-Blocker; Antihypertensive; Antianginal Agent

Selective inhibitor of beta₁-adrenergic receptors; competitively blocks beta₁-receptors, with little or no effect on beta₂-receptors at oral doses <100 mg (in adults); does not exhibit any membrane stabilizing or intrinsic sympathomimetic activity.

Onset of action: IV: 5 minutes, Duration 3-5 hours

Indications

Labeled Indications: Angina, Hypertension, Myocardial infarction

Unlabeled: Atrial fibrillation/flutter; Hypertrophic cardiomyopathy; Marfan syndrome with aortic aneurysm; Migraine prophylaxis; Supraventricular tachycardia (AVNRT, AVRT, focal atrial tachycardia); Thyrotoxicosis; Ventricular arrhythmias

Contraindications

- Hypersensitivity to metoprolol, any component of the formulation, or other beta-blockers; second- or third-degree heart block
- Severe sinus bradycardia (heart rate <45 beats/minute); significant first-degree heart block (P-R interval ≥0.24 seconds); systolic blood pressure <100 mm Hg; moderate to severe cardiac failure

Adverse Reactions / Precautions

- Cardiovascular: Hypotension, bradycardia, first degree atrioventricular block, arterial insufficiency, cardiac failure, CVA, cold extremities, palpitations, peripheral edema, claudication
- Central nervous system: Dizziness, fatigue, depression, vertigo, confusion, disturbed sleep, hallucination, headache, insomnia, nightmares, temporary amnesia, tinnitus

Dose and Administration: ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Lopressor

Atrial fibrillation or atrial flutter (off-label use): Acute ventricular rate control:

IV:

• **2.5 to 5 mg** over 2-5 minutes; repeat dose every 5 minutes as needed; maximum total dose: 15 mg.

<u>Supraventricular tachycardia/Ventricular arrhythmias (off-label use):</u>

Note: For hemodynamically stable patients if vagal maneuvers and/or adenosine are unsuccessful.

IV:

2.5 to 5 mg over 2-5 minutes; repeat dose every 5 minutes as needed to achieve a ventricular rate of 90 – 100; maximum total dose: 15 mg.

Note: For sustained ventricular tachycardia, Betablockers are generally administered in addition to an antiarrhythmic drug (eg, Amiodarone) for these indications. A beta-blocker is also used to reduce shocks in patients who receive an implantable cardioverter defibrillator for these indications; propranolol may be the preferred beta-blocker in these situations

Note: Guidelines do not recommend betablockers as initial therapy in pediatric patients; beta-blockers should be reserved for use in patients who have contraindications to preferred agents or after ≥2 preferred agents have failed in patients with hypertension and chronic kidney disease, proteinuria, or diabetes mellitus.

MIDAZOLAM

♀D, Lactation Yes (Caution)

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:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Versed

<u>Induction for RSI; Continued sedation;</u> <u>Hyperthermia:</u>

IV:

- Induction 0.1mg/kg IV/IO
- Continued Sedation .05-.1 mg/kg IV/IO
- Infusion sedation 0.05 mg/kg bolus IV, then titrate 0.05-0.1mg/kg/hr. IV gtt

<u>Transcutaneous Pacing /</u> <u>Cardioversion, Anxiety, Agitation.</u>

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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

<u>Procedural sedation; Transcutaneous Pacing;</u> 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 nare

Induction/RSI (Not preferred drug)

IV:

0.1-0.3 mg/kg

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

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MORPHINE QC, Lactation Yes(Caution)

Class / Mechanism of Action

Opioid Analgesic

Binds to opioid receptors within the CNS increasing pain threshold and altering pain reception; inhibits ascending pain pathways (blocking painful stimulus); produces CNS depression Onset: IV variable but rapid, Duration variable, patient dependent.

Indications

Labeled Indications: Moderate to severe acute and chronic pain; pain of myocardial infarction; preanesthetic medication

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

- Always be prepared for use of paralytic and intubation (maintain positive control of airway).
- 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:

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.

IV/IO: (Slow)

 5mg (0.1 mg/kg, range 2.5 – 10mg) every 1-6 hours 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.

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

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| 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: | | |
| PO: Adults: • 400 mg once daily • Max daily dose: 400 mg/day | Pediatrics: <15yrs old: • 10 mg/kg/day PO • Max daily dose: 400 mg/day | | |

>15yrs old:

• 400 mg once daily

o Max daily dose: 400 mg/day

NALOXONE QC, Lactation (Caution) Trade Name: Narcan

Class / Mechanism of Action

Antidote, Opioid Antagonist

Competes and displaces opioids at opioid receptor sites, reversing narcotic effects.

Indications

Labeled Indications: Reversal of opioid drug effects, including respiratory depression

Contraindications

• Hypersensitivity to naloxone or any component of 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

<u>Opioid overdose</u> (with standard ACS protocols): IV, IM, SubQ:

- 0.4-2 mg; may dose every 2-3 minutes if needed.
 - If no response after 10 mg total, look for other cause of respiratory depression.
 - Following reversal, may need to readminister after 20-60 minutes.

Reversal of respiratory depression with therapeutic opioid doses:

IV, IM, SubQ:

 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.

<u>Opioid overdose</u> (with standard PALS protocols): IV, IM, SubQ:

- <5 years or ≤20 kg (unlabeled dose): 0.1
 mg/kg/dose (maximum dose: 2 mg); repeat
 every 2-3 minutes PRN
 </p>
- ≥5 years or >20 kg: Adult Dosing

Reversal of respiratory depression with therapeutic opioid doses:

IV, IM, SubQ:

0.001-0.015 mg/kg/dose; repeat as needed.

NIFEDIPINE QC, Lactation Yes(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, nauseaDose and Administration:

Bradycardia, nausea

ADULT

<u>High altitude pulmonary edema (unlabeled use):</u> PO:

• 10 mg every 4-6 hours

Pulmonary hypertension (unlabeled use) PO:

 30 mg (Extended Release) twice daily; may increase cautiously to 120-240 mg/day

Note: Do not use for acute anginal episodes; may precipitate myocardial infarction

High altitude pulmonary edema (Not FDA approved for use in children) (unlabeled use):

PEDIATRIC Always Reference BROSELOW Tape

PO:

 Immediate release: 0.5 mg/kg/dose (maximum: 20 mg/dose) every 8 hours

Note: Treatment is needed only necessary if response to oxygen and/or descent is poor.

NITROGLYCERIN

QC, Lactation (Caution)

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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

Trade Name: NitroMist/Nitrostat

Angina/coronary artery disease:

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

CHF related Respiratory Distress:

PO:

 <u>Sublingual</u>: **0.4 mg** every 5 minutes for maximum of 3 doses in 15 minutes as long as SBP>90

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
 - o 30% reduction in MAP if hypertensive.
 - Max dose: 400mcg/minute)

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

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)
- 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

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.
- 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 with infusion pump. Do not use in same line as sodium bicarbonate. It will inactivate norepinephrine.

- Initial: 2-20 mcg/minute: titrate to SBP goal.
 - o Maintenance: 2-4 mcg/minute

Post ROSC Hypotension:

Initial: 0.1-0.5 mcg/kg/minute titrate to effect.

If unable to maintain MAP >60mmHg, add **Epinephrine** infusion.

Use in Burn Patient:

For Burn patients, norepinephrine is only used when target MAP (\geq 55) and UOP (\geq 30mL/hr) fail to be reached with fluid resuscitation alone. Its sequence of use follows administration of **Vasopressin**.

(See infusion chart next page for mix and dosage information)

Hypotension/shock:

IV: Continuous infusion

- Initial: 0.05-0.1 mcg/kg/minute; titrate to effect
 - Max dose: 2 mcg/kg/minute

| NOREPINEPHRINE (LEVOPHED) | | | | | |
|--|---|-----------|---------|-----------|---------|
| Dosing Range: 2-20mcg/min (120-1200mcg/hr) | | | | | |
| MIX 4 mg/500 mL | | | | | |
| | C | CONCENTRA | • | | |
| 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 congestive 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)

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)

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

Trade Name: Zofran

Nausea and Vomiting:

IV/IO/IM/PO

4-8 mg

<u>Treatment of severe or refractory hyperemesis</u> <u>gravidum (unlabeled use):</u>

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8 mg administered over 15 minutes every 12 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? (Caution)

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 component of the formulation
- Ventricular Tachycardia and Hypertension
- Bradvcardia

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.
- 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

Trade Name: Neosynephrine

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 NS for a concentration of 100mcg/mL

IV Infusion:

- **40 200 mcg/min**; titrate to MAP > 60 mm Hg.
 - To titrate, increase rate by 10 mcg/min every 2 minutes.
 - Maximum dose is 200 mcg/min.
 - Mix 10mg phenylephrine in 250mL D5W/NS for a concentration for 40mcg/mL

If unable to maintain MAP >60mmHg, add Epinephrine infusion.

Hypotension / Shock:

IV Push:

5-20 mcg/kg/dose every 10-15 minutes as needed

IV Infusion:

0.1-0.5 mcg/kg/minute

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 able.

PROMETHAZINE

QC, Lactation?(Not Recommended)

Class / Mechanism of Action

Phenothiazine derivative Antiemetic, Histamine H₁ Antagonist, Sedative

Blocks postsynaptic dopaminergic receptors in the brain; strong alpha adrenergic blocking effect and depresses release of hypothalamic and hypophyseal hormones; reduces stimuli to the reticular system Onset of action IV: 5 minutes, Duration 4-6 hours

Indications

Labeled Indications: Symptomatic treatment for allergic conditions; antiemetic; motion sickness; sedative; adjunct to postoperative analgesia and anesthesia

Unlabeled: Treatment of nausea and vomiting of pregnancy

Contraindications

- Hypersensitivity to promethazine, phenothiazine allergy, or any component of the formulation
- Coma
- Children <2 years old
- Intra-arterial and SubQ administration

Adverse Reactions / Precautions

- May cause Bradycardia, hyper-/hypotension, nonspecific QT changes, orthostatic hypotension, tachycardia: Life threatening arrhythmias have occurred with normal dosage
- May cause extrapyramidal symptoms (pseudoparkinsonism, acute dystonic reactions, akathisia, etc.)
- Avoid use in severe respiratory disease (asthma, COPD), and in patients using other sedatives or depressants: may lead to respiratory depression
- Vesicant: can cause severe tissue injury regardless of route of delivery
 - Deep IM injection; or IV in line. Slow IVP over 1 minute
 - For IV, ensure proper needle/catheter venous placement; avoid extravasation

Dose and Administration:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

Trade Name: Phenergan

Antiemetic:

IV push over >1 minute

- 12.5 mg, not to exceed 25 mg
 - May repeat 12.5mg once after 10 minutes if first dose ineffective
 - Subsequent dose of 25mg may be given every 4 hours
 - o Can dilute with 10-20mL of NS

Sedation, analgesia/hypnotic adjunct:

IM, IV:

 25-50 mg in combination with analgesic or hypnotic (at reduced dosage)

<u>Allergic conditions</u> (including allergic reactions to blood or plasma):

IM, IV:

• 25 mg, may repeat in 2 hours when necessary

Antiemetic:

IM. IV:

 Children ≥2 years: 0.25 mg/kg 4-6 times/day as needed (maximum: 12.5 mg/dose)

Preoperative analgesia/hypnotic adjunct: IM, IV:

 Children ≥2 years: 1.1 mg/kg in combination with an analgesic or hypnotic (at reduced dosage) and with an atropine like agent (at appropriate dosage).

Note: Promethazine dosage should not exceed half of suggested adult dosage.

PROPOFOL PRO

Class / Mechanism of Action

General Anesthetic

Lipophilic intravenous general anesthetic.

Onset of action IV bolus: 9-51 seconds (average 30 seconds), Duration is dose and rate dependent: 3-10 minutes, prolonged with continued doses

Indications

Labeled Indications: Induction of anesthesia in patients ≥3 years of age; maintenance of anesthesia in patients >2 months of age; sedation in intubated, mechanically ventilated ICU patients

Contraindications

- Hypersensitivity to propofol or any component of the formulation
- Allergy to eggs, egg products, soybeans, soy products, and peanuts.

Adverse Reactions / Precautions

- May cause Hypotension especially in hypovolemic patients or if bolus dosing is used.
 - Hypotension may result in reduction of MAP exceeding 30%
 - Head Injury patients or those with suspected / known increased intracranial pressure are at increased risk of decreased cerebral perfusion pressure.
- Do not use in pre-hospital trauma environment or in burn transfer patients unless directed by medical director or provided written orders by referring provider.
- No Analgesic properties. Must supplement with analgesic agents.

Dose and Administration: ADULT PEDIATRIC Always Reference BROSELOW Tape

Sedation/ RSI:

IV Push:

• 1-2.5 mg/kg every 5-10min PRN.

Maintenance of general anesthesia:

IV Infusion:

- 10-75 mcg/kg/min via infusion pump or Dial-a-Drip. Titrate to minimum effective dose. (See infusion chart next page)
 - o 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 changes to clinically assess drug effects. Smaller doses are required when used with opioids.

Note: Not preferred in Burn patients in the first 48-72 hrs

Sedation/ RSI:

IV Push:

• 1-2.5 mg/kg every 5-10min PRN.

Maintenance of general anesthesia,

IV Infusion:

Healthy children 2 months to 16 years:

• **125-300 mcg/kg/minute** (or 7.5-18 mg/kg/hour)

| osing Rang | e: 10-75mcg/kg | | | g/hr) | | |
|------------|----------------|----------|-----------------------------|-----------|-----------|-----------|
| | | |) mg/50 mL ITRATION 10 r | ng/ml | | |
| . Weight | Dose | Rate | Micro | IS/IIIL | Macro | |
| | | | (60 gtt/mL) | 20 gtt/mL | 15 gtt/mL | 10 gtt/ml |
| kg | mcg/kg/min | mL/hr | gtt/min | gtt/min | gtt/min | gtt/min |
| | 10 | 3 | 3 | 1 | 1 | 1 |
| | 15 20 | 5 6 | 5 | 2 | 2 | 1 |
| | 25 | 8 | 6 8 | 3 | 2 | 1 |
| | 30 | 9 | 9 | 3 | 2 | 2 |
| | 35 | 11 | 11 | 4 | 3 | 2 |
| 50 | 40 | 12 | 12 | 4 | 3 | 2 |
| 50 | 45 | 14 | 14 | 5 | 3 | 2 |
| | 50 | 15 | 15 | 5 | 4 | 3 |
| | 55 60 | 17 18 | 17 18 | 6 | 4 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 |
| | 15 | 5 | 5 | 2 | 1 | 1 |
| | 20 | 7 | 7 | 2 | 2 | 1 |
| | 25 30 | 8 10 | 8 10 | 3 | 2 | 2 |
| | 30 35 | 10 | 10 | 4 | 3 | 2 |
| | 40 | 13 | 13 | 4 | 3 | 2 |
| 55 | 45 | 15 | 15 | 5 | 4 | 2 |
| | 50 | 17 | 17 | 6 | 4 | 3 |
| | 55 | 18 | 18 | 6 | 5 | 3 |
| | 60 | 20 | 20 | 7 | 5 | 3 |
| | 65 70 | 21 23 | 21 23 | 7 | 5 | 4 |
| | 70 75 | 23 | 25 | 8 | 6 | 4 |
| | 10 | 4 | 4 | 1 | 1 | 1 |
| | 15 | 5 | 5 | 2 | 1 | 1 |
| | 20 | 7 | 7 | 2 | 2 | 1 |
| | 25 | 9 | 9 | 3 | 2 | 2 |
| | 30 | 11 | 11 | 4 | 3 | 2 |
| | 35 | 13 | 13 | 4 | 3 | 2 |
| 60 | 40 45 | 14 16 | 14 16 | 5 | 4 | 2 |
| | 50 | 18 | 18 | 6 | 5 | 3 |
| | 55 | 20 | 20 | 7 | 5 | 3 |
| | 60 | 22 | 22 | 7 | 5 | 4 |
| | 65 | 23 | 23 | 8 | 6 | 4 |
| | 70 | 25 | 25 | 8 | 6 | 4 |
| | 75 | 27 | 27 | 9 | 7 | 5 |
| | 10 | 4 | 4 | 1 | 1 | 1 |
| | 15 20 | 6 8 | 6 8 | 3 | 2 | 1 |
| | 25 | 10 | 10 | 3 | 2 | 2 |
| | 30 | 12 | 12 | 4 | 3 | 2 |
| | 35 | 14 | 14 | 5 | 3 | 2 |
| 65 | 40 | 16 | 16 | 5 | 4 | 3 |
| 05 | 45 | 18 | 18 | 6 | 4 | 3 |
| | 50 | 20 | 20 | 7 | 5 | 3 |
| | 55 60 | 21 23 | 21 | 7 | 5 6 | 4 |
| | 65 | 25 | 25 | 8 | 6 | 4 |
| | 70 | 27 | 27 | 9 | 7 | 5 |
| | 75 | 29 | 29 | 10 | 7 | 5 |
| | 10 | 4 | 4 | 1 | 1 | 1 |
| | 15 | 6 | 6 | 2 | 2 | 1 |
| | 20 | 8 | 8 | 3 | 2 | 1 |
| | 25 30 | 11 13 | 11 13 | 4 | 3 | 2 |
| | 35 | 15 | 15 | 5 | 4 | 2 |
| | 40 | 17 | 17 | 6 | 4 | 3 |
| 70 | 45 | 19 | 19 | 6 | 5 | 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 75 | 29 32 | 29 32 | 10 11 | 8 | 5 5 |
| | 10 | 5 | 5 | 2 | 1 | 1 |
| 75 | 15 | 7 | 7 | 2 | 2 | 1 |
| | 20 | 9 | 9 | 3 | 2 | 2 |
| | 25 | 11 | 11 | 4 | 3 | 2 |
| | 30 | 14 | 14 | 5 | 3 | 2 |
| | 35 | 16 | 16 | 5 | 4 | 3 |
| | 40 | 18 | 18 | 6 | 5 | 3 |
| | 45 50 | 20 | 20 | 7 | 5 6 | 3 |
| | 55 | 25 | 25 | 8 | 6 | 4 |
| | 60 | 27 | 27 | 9 | 7 | 5 |
| | 65 | 29 | 29 | 10 | 7 | 5 |
| | 70 | 32 | 32 | 11 | 8 | 5 |
| | 75 | 34 | 34 | 11 | 8 | 6 |

| Macro-Drip (20gtt/ml) or Micro-Drip is set of choice for this infusion |
|--|
| Titrate to minimum effective dose. Allow 3-5 minutes between dosing changes to |
| sedative and hemodynamic effects. |

| | | | 0 mg/50 mL NTRATION 10 r | ng/m! | | |
|--------|------------|----------|-----------------------------|-----------|-----------|----------|
| Weight | Dose | Rate | Micro | ng/mL | Macro | |
| weignt | Dose | Nate | (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/mir |
| | 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 60 | 26 29 | 26 29 | 9 | 7 | 4 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 |
| or. | 40 | 20 | 20 | 7 | 5 | 3 |
| 85 | 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 20 | 8 11 | 8 11 | 3 | 3 | 2 |
| | 25 | 14 | 14 | 5 | 3 | 2 |
| | 30 | 16 | 16 | 5 | 4 | 3 |
| | 35 | 19 | 19 | 6 | 5 | 3 |
| | 40 | 22 | 22 | 7 | 5 | 4 |
| 90 | 45 | 24 | 24 | 8 | 6 | 4 |
| | 50 | 27 | 27 | 9 | 7 | 5 |
| | 55 | 30 | 30 | 10 | 7 | 5 |
| | 60 | 32 | 32 | 11 | 8 | 5 |
| | 65 | 35 | 35 | 12 | 9 | 6 |
| | 70 | 38 | 38 | 13 | 9 | 6 |
| | 75 | 41 | 41 | 14 | 10 | 7 |
| | 10 | 6 | 6 | 2 | 1 | 1 |
| | 15 | 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 |
| 95 | 40 45 | 23 | 23 26 | 8 | 6 | 4 |
| | 50 | 26 29 | 26 | 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 |
|) | 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 25 | 13 16 | 13 | 5 | 3 | 3 |
| | | + | 16 | 6 | 5 | 3 |
| | 30 35 | 19 22 | 22 | 7 | 6 | 4 |
| | 40 | 25 | 25 | 8 | 6 | 4 |
| 5 | 45 | 28 | 25 | 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 |

Macro-Drip (20gtt/ml) or Micro-Drip is set of choice for this infusion
Titrate to minimum effective dose. Allow 3-5 minutes between dosing changes to sedative and hemodynamic effects.

ROCURONIUM QC, Lactation? (Caution) Trade Name: Zemuron

Class / Mechanism of Action

Nondepolarizing Neuromuscular Blocking Agent (Paralytic)

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

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 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: |
| 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) | Maintenance bolus dosing: (unlabeled and unreferenced dose) 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 succinvlcholine 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.)

Adverse Reactions / Precautions

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 mEg/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: 1-1.5 mg/kg Note: Pretreatment with 10% dosag depolarizing agents prior to neuromublockade with Succinylcholine is NO ADVISED | uscular- | RSII / Neuromuscular blockade: IV: • <10kg: • lnitial: 1.5-2 mg/kg/dose • >10kg: • lnitial: 1-1.5 mg/kg/dose Note: Pretreatment with 10% dosage of non-depolarizing agents prior to neuromuscular-blockade with Succinylcholine is NO LONGER ADVISED |

| THIAMINE | QA, LactationYes (Caution) Trade Name: Vitamin B | | | | |
|--|--|-----------------------|------------------------------|--|--|
| Class / Mechanism of Action | | | | | |
| Vitamin, water soluble Essential coenzyme in carbohydrate metabolism. Onset of action IV/IM: Rapid | | | | | |
| Indications | 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 thiamine or any component of the formulation | | | | | |
| Adverse Reactions / Precautions | | | | | |
| Administration of dextrose may worsen acute symptoms of thiamine deficiency; use caution when low thiamine is suspect | | | | | |
| Dose and Administration: | ADULT | PEDIATRIC AI | ways Reference BROSELOW Tape | | |
| AMS; Seizure; Syncope; Ma and Diarrhea; w/ Hx of ETO IM/IV: • 100mg/day | | AMS or Seizure w/ sig | ns of Malnutrition: | | |

TRANEXAMIC ACID ♀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)

Adverse Reactions / Precautions

- 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:

ADULT

PEDIATRIC Always Reference BROSELOW Tape

<u>Trauma-associated hemorrhage (unlabeled use):</u>

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.
- If patient received 1 gram of TXA prior and <3hrs from time of injury: 1-gram TXA IV/IO push over 1 minute or mixed in 100cc NS or LR Bolus. If >3hr from time of injury: <u>DO NOT</u> administer TXA.

<u>Suspected Post-Operative Hemorrhage by dissection, enteric staples or suspected internal bleeding:</u>

 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 push) but NOT later than 3 hours after start of suspected hemorrhage.

<u>Trauma-associated hemorrhage (unlabeled use):</u>

IV:

• Initial Dose: **15mg/kg** via IV/IO Bolus (goal within 1 minute),

VASOPRESSIN

QC, Lactation?(Caution)

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

Trade Name: Vasostrict

BROSELOW Tape

Hypotension / Shock:

Vasopressors should be used if patient is hypotensive after fluid resuscitation to maintain mean arterial pressure (MAP) ≥65 mmHg.

Use in addition to norepinephrine for raising MAP to target or to decrease norepinephrine dosage.

Titrate to lowest effective dose.

IV Infusion:

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

Hypotension / Shock:

Limited data available; efficacy results have varied.

IV Infusion:

• 0.17 to 8 <u>milliu</u>nits/kg/<u>minute</u> (0.01 to 0.48 units/kg/hour)

VECURONIUM QC, Lactation? (Caution) Trade Name: Norcuron

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 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) and maintenance of paralysis:

IV Push:

- Induction: 0.1 mg/kg Dose range (0.08-0.15 mg/kg)
- Maintenance: 0.1 mg/kg Dose range (0.08-0.15 mg/kg) every 30-60 minutes PRN

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.

RSI and maintenance of paralysis:

IV Push:

- Induction: 0.1-0.15 mg/kg
- Intermittent bolus dosing: 0.1 mg/kg every 30-60 minutes PRN

IV Continuous infusion:

• 1-2.5 mcg/kg/minute

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WE WILL NEVER FORGET



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