Sierra – Sacramento Valley EMS Agency Treatment Protocol			
Non-Traumatic Pulseless Arrest			
Approval: Troy M. Falck, MD – Medical Director	Effective: 12/01/20)24	
Approval: John Poland – Executive Director	Next Review: 10/2	2027	
MANUAL CHEST COMPRESSIONS	MECHANICAL CHEST COMPRESSION DEVICES		
 Rate: 100-120/min Depth: 2 inches – allow full chest recoil Minimize interruptions (≤10 secs) Rotate compressors every 2 mins 	IndicationsContraindication• Adult pt (≥15 yo)• Pt does not fit in th • 3 rd trimester pregn	ions ne device ancy	
 Perform CPR during AED/defibrillator charging Resume CPR immediately after shock 	 Use in accordance with manufacturer indications contraindications Apply following completion of at least one manu CPR cycle, or at the end of a subsequent cycle 		
DEFIBRILLATION & GENERAL PT MANAGEMENT	ADVANCED AIRWAY MANAGEMENT		
 Analyze rhythm/check pulse after every 2 min CPR cycle Biphasic manual defibrillation detail: Follow manufacturer recommendations If unknown, start at 200 J (subsequent doses should be equivalent or higher) Movement of pt may interrupt CPR or prevent adequate depth and rate of compressions Consider resuscitation on scene up to 20 mins Go to ROSC protocol (C-2) if ROSC is obtained 	 Consider/establish advanced airway at app time during resuscitation Do not interrupt chest compressions to est advanced airway Waveform capnography (if available) shall on all pts with an advanced airway in place An abrupt increase in PETCO₂ is indicati ROSC Persistently low PETCO₂ levels (<10 mm suggest ROSC is unlikely 	oropriate ablish an be used ve of iHG)	
TREAT REVERSIBLE CAUSES	TERMINATION OF RESUSCITATION		
 Hypovolemia Hypoxia Hydrogen Ion (acidosis) Hypo-/hyperkalemia Hypothermia Thrombosis, pulmonary Thrombosis, cardiac Toxins (1) Refer to Hypothermia & Avalanche/Snow Immersion Suffocation Resuscitation Protocol (E-2) or Traumatic Pulseless Arrest Protocol (T-6) as appropriate (2) Contact the base/modified base hospital for consultation & orders as appropriate (2) Consider early transport of pts who have reversible causes that cannot be adequately treated in the prehospital setting	 Base/Modified Base Hospital Physician (If resuscitation attempts do not obtain ROS consider termination of resuscitation efforts BLS termination of resuscitation criteria (al (1) Arrest not witnessed by EMS (2) No AED shocks delivered (3) No ROSC after 3 rounds of CPR/AED at ALS Termination of Resuscitation Criteria (ALS Termination of Resuscitation Criteria ((1) Arrest not witnessed by EMS (2) No effective bystander CPR was provide effective CPR cannot be maintained (3) No AED shocks or defibrillations delivered (4) No ROSC after full ALS care **In the event of communication failure, EMS personnel may terminate resuscitation withor base/modified base hospital physician order who meets ALS termination of resuscitation 	Drder** SC, SC, I): analysis (all): led, or red Sout a on a pt criteria.	

SEE PAGE 2 FOR TREATMENT ALGORITHM















General Medical Treatment

Nausea/Vomiting

- Nausea/vomiting can be symptoms of a multitude of different causes. If possible, the specific underlying cause should be determined and treated. The use of an antiemetic may relieve symptoms while leaving the cause untreated, and possibly, more difficult to detect. EMS personnel should weigh the benefits of antiemetic use against the possible risk of making an accurate diagnosis more difficult, and the possible side effects of the antiemetic agent.
- Treatment of nausea/vomiting is indicated for pts where it may contribute to a worsening of their medical condition, or where the pt's airway may be endangered.
- EMS personnel may consider administering Zofran (Ondansetron) prophylactically, prior to or immediately after opioid administration, for a pt with a history of nausea/vomiting secondary to opioid administration. Zofran (Ondansetron) may also be administered prior to transport to a pt with a history of motion sickness.



Zofran (Ondansetron)

- 4 8 mg oral disintegrating tablet, **OR** 4 8 mg IM, **OR** 4 8 mg slow IV/IO (over 30 seconds)
- May repeat as needed (max total dose: 16 mg)

Zofran (Ondansetron) is contraindicated during the first 8 weeks of pregnancy





N-3

Suspected Stroke Effective: 12/01/2024 Approval: Troy M. Falck, MD – Medical Director Approval: John Poland – Executive Director Next Review: 10/2027 Cincinnati Prehospital Stroke Scale (CPSS) **Normal Result** Component **Abnormal Result Facial Droop** One side of face does not move as Both sides of face move equally (Ask pt to show teeth or smile) well as the other side Arm Drift Both arms move the same, or both One arm does not move, or one arm (Ask pt to close eyes & hold both arms do not move drifts down compared with the other arms out with palms up) Speech Pt slurs words, uses the wrong (Ask pt to say "you can't teach an old Pt uses correct words with no slurring words, or is unable to speak dog new tricks") BLS • Assess V/S, including SpO₂ • O₂ at appropriate rate if hypoxemic (SpO₂ <94%) or short of breath Perform CPSS assessment Suspect stroke for any of the following: New onset symptoms with abnormal CPSS • New onset altered state (GCS <14) with unidentifiable etiology CPSS is normal, but patient/bystander report stroke symptoms within previous 24 hrs If stroke suspected: Determine time of onset of symptoms (pt last known normal) - When possible, obtain and relay to the receiving hospital the name/contact information of the individual who can verify the time of onset of symptoms (pt last known normal) Check blood glucose (if glucometer available) Transport as soon as possible (scene time should be ≤10 mins) ALS Consider advanced airway if GCS ≤8 or need for airway protection Cardiac monitor, consider 12-lead EKG (do not delay transport to perform 12-lead EKG) Obtain blood draw if requested by stroke receiving center IV/IO NS TKO (may bolus up to 1000 mL) • Transport to closest Transport to closest Are both the following stroke receiving center appropriate hospital present? Advise of "Stroke Alert" & Onset of symptoms ≤24 hrs Contact base/modified ·NΟ YEStime pt. last known normal base hospital for (including wake-up stroke*) Provide pt. identifying destination consultation if • ≤45 minute transport time to information if requested a stroke receiving center necessary by stroke receiving center

*Wake-up stroke definition: Pt awakens with stroke symptoms that were not present prior to falling asleep



Sierra – Sacramento Valley EMS Agency Treatment Protocol T-4 Hemorrhage Effective: 12/01/2024 Approval: Troy M. Falck, MD – Medical Director Approval: John Poland – Executive Director Next Review: 10/2027 Approved Commercial Tourniquet Devices: - Emergency and Military Tourniquet - Mechanical Advantage Tourniquet - Combat Application Tourniquet - SAM XT Extremity Tourniquet - Special Ops. Tactical Tourniquet - RECON Medical Tourniquet **Tourniquet Utilization Notes:** • Tourniquets applied by lay rescuers or other responders shall be evaluated for appropriateness and may be adjusted or removed if necessary - improvised tourniquets should be removed by prehospital personnel. • If application is indicated and appropriate, a commercial tourniquet should not be loosened or removed by prehospital personnel unless time to definitive care will be greatly delayed (>2 hrs). **Approved Hemostatic Agents:** - HemCon ChitoGauze XR2 PRO - QuikClot EMS 4x4 & Combat Gauze - HemCon ChitoGauze XR PRO - HemCon ChitoGauze OTC - HemCon Bandage PRO - HemCon OneStop Bandage BLS Assess V/S, including SpO₂ • O2 at appropriate rate if hypoxemic (SpO2 <94%) Attempt to control bleeding with direct pressure Uncontrolled Monitor & reassess Hemorrhage? YES Non-compressible or Extremity, area Non-extremity, suspected internal compressible bleeding amenable to tourniquet hemorrhage Consider hemostatic Pressure dressing agent application • Consider hemostatic • Apply tourniquet agent application proximal to bleeding if necessary • Apply 2nd tourniquet proximal to 1st for continued bleeding Continued Hemorrhage? YES Evaluate for TXA administration (page 2)

Hemorrhage

Tranexamic Acid (TXA) Administration

TXA Administration Notes:



• For post-partum hemorrhage, refer to Childbirth Protocol (OB-G1).





SEE PAGE 2 FOR ALS TREATMENT OF WHEEZING OR SUSPECTED CROUP/EPIGLOTTITIS





Pediatric General Medical Treatment

M-6P

Approval: Troy M. Falck, MD - Medical Director

Effective: 12/1/2024

Approval: John Poland – Executive Director

Next Review: 07/2027

GENERAL PEDIATRIC TREATMENT PRINCIPLES

- The purpose of this protocol is to provide standing order assessment/treatment modalities for pediatric pt complaints not addressed in other S-SV EMS treatment protocols including Nausea/Vomiting (Page 2), Brief Resolved Unexplained Event BRUE (Page 3) & Suspected Shock/Sepsis (Page 4).
- The Neonatal Resuscitation Protocol (C-1N) shall be used for pts during the first 28 days of life.
- Pediatric protocols shall be utilized for pts >28 days up to and including 14 years old.
- Applicable adult protocols may be utilized when there is not a pediatric protocol applicable to the pt's complaint/condition. Prehospital personnel shall consult with the base/modified base hospital for additional direction, if needed, when there is no standing order treatment protocol applicable to the pt's condition.
- A parent/reliable family member reported weight, length-based pediatric resuscitation tape or Handtevy shall be utilized for determining sizes of equipment and defibrillation/cardioversion joule settings. Once weight has been determined, medication dosing shall be based on S-SV EMS pediatric protocols.

NORMAL VITAL SIGNS & HYPOTENSION DEFINITION FOR NEONATAL & PEDIATRIC PATIENTS

Age	Normal Pulse Rate	Normal Resp. Rate	Normal SBP	Hypotension
≤28 days	100 - 205	30 - 50	60 - 80	SBP <60
29 days -12 months	90 - 180	30 - 50	70 - 100	SBP <70
1-2 years	80 - 140	24 - 40	80 - 110	SBP <70 + age x2
3-5 years	65 - 120	20 - 30	90 - 110	SBP <70 + age x2
6-9 years	60 - 120	20 - 30	100 - 120	SBP <70 + age x2
10-14 years	50 - 100	12 - 20	100 - 120	SBP <90

PEDIATRIC PROTOCOLS PROCEDURE/MEDICATION TREATMENT AGE RESTRICTIONS

- **<28 days old:** Base/modified base hospital order required to administer a fluid bolus (C-1N)
- <3 years old: Needle cricothyrotomy is not allowed (PR-3 & R-3P)
- <4 years old: Base/modified base hospital order required to administer the following medications:
 - Zofran/Ondansetron for nausea/vomiting (M-6P)
 - Analgesic medications for pain management (M-8P)
 - Midazolam for severe anxiety/combative symptoms (M-11P)
 - PO acetaminophen for febrile symptoms (N-2P & M-6P)
- <8 years old: CPAP is not allowed (R-3P)
- <15 years old: Base/modified base hospital order required to utilize the following procedures/medications:
 - Transcutaneous pacing for bradycardia (C-3P)
- Synchronized cardioversion for tachycardia (C-4P)
- Adenosine for tachycardia (C-4P)





Pediatric General Medical Treatment

Brief Resolved Unexplained Event (BRUE)















Multiple Patient Incidents

Approval: Troy M. Falck, MD – Medical Director

Approval: John Poland – Executive Director

Effective: 12/01/2024

G-1

DEFINITIONS

Control Facility (CF): An acute care hospital or EMS dispatch center responsible for situation status reporting and patient dispersal during a MCI or URVI.

EMS Surge Incident: An incident that does not overwhelm prehospital resources but has the potential to overwhelm hospital resources with multiple patients.

Unified Response to Violent Incident (URVI): An evolving event, primarily managed by law enforcement (LE), involving the use of force or violence on a group of people (e.g. mass shooting, bombing, riots, etc.). These incidents present a significantly higher threat of injury or loss of life to first responders, victims, and the public.

Multiple Casualty Incident (MCI): An incident that requires more prehospital and/or hospital resources to adequately manage patients than those available during a routine response. A MCI is categorized by the following levels:

LEVEL 1 MCI: Approximately 5-14 patients, expected duration ≤1 hour

LEVEL 2 MCI: Approximately 15-49 patients, expected duration ≥1 hour

LEVEL 3 MCI: 50+ patients, expected duration ≥1 hour

EMS SURGE ALERT	MCI ALERT
When:	When:
 Three (3) or more ground or air transport resources are requested to respond to an incident; or Three (3) or more patients are identified after arrival at the scene of an incident; or Multiple patients are released at scene who may arrive at a hospital by private vehicle. A URVI. 	 An incident that requires more EMS system resources to manage patients than those available during a routine response; or The number of patients from a single incident overwhelms the CF or closest appropriate receiving hospital. Who:
Who:	 Dispatch center, prehospital resources, or CF.
 Dispatch center or first dispatched ground transport resource. 	Why:
 Why: To provide early notification to the CF for situation status reporting and hospital polling. 	 To provide early notification for situation status reporting, hospital polling and initiation of the Regional MCI Plan.





Determination Of Death

Next Review: 07/2017

Approval: Troy M. Falck, MD – Medical Director	Effective: 12/01/2024

Approval: John Poland – Executive Director

General Procedures/Considerations:

- CPR need not be initiated and may be discontinued for pts who meet Obvious Death or Probable Death criteria as contained in this protocol, at the time of initial assessment.
- A valid Do Not Resuscitate (DNR) should be honored for any pt with absent respirations, pulses and neurological response, regardless of the cause of death (e.g. terminal illness, trauma).
- Hypothermia, drug and/or alcohol overdose can mask neurological reflexes. If any doubt exists about contributing environmental factors (e.g. cold water submersion) and no valid DNR exists, initiate resuscitation and treat according to applicable S-SV EMS protocol.
- In the event of a declared MCI, death may be determined in accordance with START/JUMP START criteria.
- For all pts treated under this protocol, the following must be assessed/confirmed (as possible):
 - Absent respirations: look, listen (auscultate), and feel for respirations for a minimum of 30 secs.
 - Absent pulses: palpate both the carotid and apical pulses for a minimum of 30 secs.
 - Absent neurological response: check pupil response with a light and check for response to painful stimuli.
- If the base/modified base hospital physician directs EMS personnel to stop resuscitation efforts once transport has begun, the ambulance shall reduce transport code and continue transport to the original destination hospital.
- If determination of death is made at rendezvous location with HEMS aircraft, the body shall not be moved from the ambulance and an immediate request for law enforcement shall be made.
- If there is any objection/disagreement by family members or EMS personnel to terminating or withholding
 resuscitation for pts who have a valid DNR or meet probable death criteria, BLS measures (including
 defibrillation) shall continue or begin immediately and EMS personnel shall contact the base/modified
 base hospital for further direction.

Instructions for EMS Personnel Upon Determination of Death:

- If not already on scene, request law enforcement
- Minimize contact with the body and scene to protect potential crime scene evidence
- Appropriate EMS personnel shall remain on scene until released by law enforcement
- Provide law enforcement with the following minimum information:
 - Unit ID
 - Name and certification/license # of EMS provider determining death
 - Patient demographics and known, pertinent medical history
 - Determination of death date and time
- At a minimum, the PCR must include the following:
 - Time of determination of death
 - Six-second cardiac monitor strip of two (2) leads for pts meeting probable death criteria

See page 2 for Determination of Death Assessment Criteria



Sierra – Sacramento Valley EMS Agency Treatment Protocol

DNR, POLST & End Of Life Option Act

Approval: Troy M. Falck, MD – Medical Director

Effective: 12/01/2024 Next Review: 07/2027

G-3

Approval: John Poland – Executive Director

DEFINITIONS

Advance Health Care Directive (AHCD) – A document that allows an individual to provide healthcare instructions &/or appoint an agent to make healthcare decisions when they are unable or prefer to have someone speak for them.

Agent or Attorney-In-Fact – An individual designated in a power of attorney for health care to make a health care decision for the pt, regardless of whether the person is known as an agent or attorney-in-fact, or by some other term.

Aid-in-Dying Drug – A drug prescribed by a physician for a qualified individual, which the qualified individual may choose to self-administer to bring about their death.

Do Not Resuscitate (DNR) – A request to withhold interventions to restore cardiac activity & respirations (no chest compressions, defibrillation, assisted ventilation, advanced airways, or cardiotonic medications).

DNR Wrist or Neck Medallion – A MedicAlert® or other approved wrist or neck medallion, engraved with the words "Do Not Resuscitate", and a patient ID number.

Durable Power of Attorney for Health Care (DPAHC) – A document that allows an individual to appoint an agent/attorney-in-fact to make health care decisions if they become incapacitated. The DPAHC must be immediately available and the agent/attorney-in-fact must be physically present. Decisions made by the agent/attorney-in-fact must be within the limits set by the DPAHC, if any.

EMSA/CMA Prehospital DNR Form – A form developed by the California Emergency Medical Services Authority (EMSA) and California Medical Association (CMA) for the purpose of instructing EMS personnel to forgo resuscitation attempts in the event of a pt's cardiopulmonary arrest in the out of hospital setting. The form must be signed and dated by a physician and pt/representative to be valid.

End of Life Option Act – A law authorizing an adult, 18 years or older, who meets certain qualifications and who has been determined by their attending physician to be suffering from a terminal disease, to request an aid-in-dying drug prescribed for the purpose of ending their life in a humane and dignified manner.

Physician's Orders for Life Sustaining Treatment (POLST) – A physician order form that addresses a patient's wishes about a specific set of medical issues related to end-of-life care. The form must be signed and dated by a physician and pt/representative to be valid.

VALID DNR ORDERS/FORMS

- EMSA/CMA Prehospital DNR form
- POLST form
- DNR wrist or neck medallion
- DNR order in the medical record of a licensed healthcare facility signed by a physician (or an RN verifying a valid verbal physician order on a physician order sheet), or an electronic physician's order
- Verbal DNR order given by the patient's physician
- An AHCD or DPAHC with the agent/attorney-in-fact physically present and stating the pt refuses resuscitative measures

DNR orders do not expire and photocopies/electronic physician's orders are considered valid

DNR, POLST & End Of Life Option Act

• All pts shall receive an immediate assessment/evaluation by EMS personnel.

- A copy of applicable DNR orders/forms shall be attached to the EMS patient care report (PCR) when available. - If DNR orders/forms are not available, document the method of DNR verification in the PCR.
 - If DNR bracelet or neck medallion present, document the medallion number in the PCR.
- If applicable, document the name/contact information of any agent, attorney-in-fact or other pt representative.
- If pt is transported by EMS, DNR orders/forms shall be taken with the pt to the receiving facility.
- Pts with a POLST form indicating "Comfort-Focused Treatment", are typically only transported to a hospital if their comfort needs cannot be met in their current location/setting. These pts who have no signs of pain or respiratory distress, & who have sufficient family/caretaker support present, may be released at scene by EMS personnel & not transported to the hospital, unless transportation is requested by the patient/legal representative.
- EMS personnel shall contact the base/modified base hospital for consultation for any questions or concerns regarding EMS treatment/transport of a patient with a POLST form.
- Provide supportive care to family members/caregivers as appropriate.





12-Lead EKG



Approval: Troy M. Falck, MD – Medical Director

Effective: 12/01/2024

Approval: John Poland – Executive Director

Next Review: 07/2027

INDICATIONS 12-lead EKG procedures shall be performed on pts who present with one or more of the following: • Sign/symptoms suggestive of acute coronary syndrome (ACS) such as: - Non-traumatic chest or upper abdominal discomfort - Syncope/near-syncope - Acute generalized weakness - Dyspnea Cardiac dysrhythmias on 4-lead EKG ROSC following cardiac arrest **PRE-PROCEDURE** Assess vital signs including SpO₂ Administer O₂ as indicated by clinical condition PROCEDURE Prepare EKG monitor and connect 12-lead cables Utilize packaged electrodes designed for single pt use (not bulk) • Prep skin as necessary (e.g. wiping with 4x4 gauze, shaving) • Enter, at a minimum, pt's age, gender, and last name/first initial into the cardiac monitor • Apply chest leads using the landmarks indicated on the diagram While acquiring the 12-lead EKG: - Position pt away from 60hz RF noise (light switches, smartphones, LED lights, etc.) - Position pt supine, or semi-fowler with their arms at their side and legs uncrossed - Instruct pt to breath normally and remain still - Don't converse with or touch pt during acquisition Interpret the EKG findings • If isoelectric line has significant artifact or machine reads "poor data quality" (or equivalent), attempt to reacquire a clean 12-lead EKG if pt condition allows **POST-PROCEDURE** 12-lead EKG's meeting STEMI criteria shall be transmitted to the appropriate facility (closest hospital or STEMI Receiving Center depending on incident specific circumstances) as soon as possible if transmission capabilities are available • For pts with suspected ACS, serial 12-lead EKGs should be obtained if the pt's clinical status changes or if EKG changes are noted on the cardiac monitor, and every 15 minutes if transport times are long

 Copies of 12-lead EKGs shall be provided to the receiving hospital physician upon EMS arrival, left at the receiving hospital at time of pt delivery, and attached to the EMS pt care report (PCR)



Pleural Decompression

Approval: Troy M. Falck, MD – Medical Director

Approval: John Poland – Executive Director

Effective: 12/01/2024

Next Review: 07/2027

INDICATIONS

- Suspected tension pneumothorax with absent or diminished breath sounds & one or both of the following:
 - Combined hypotension (SBP <90) and SpO₂ <94%
- Penetrating injury to the thorax
- Traumatic cardiac arrest if chest or multi-system trauma is suspected

PRE-PROCEDURE

- Assess respiratory status, manage airway & assist ventilations as appropriate
- Administer high flow O₂ & monitor SpO₂
- Assess & continually monitor vital signs

PROCEDURE

- Identify & prep the site approved sites in preferred order:
 - **A** Mid-clavicular line in the 2^{nd} intercostal space **B** - Mid-axillary line in the 4^{th} or 5^{th} intercostal space
 - **B** Mid-axillary line in the 4th or 5th intercostal space above the nipple line
 - **C** Anterior axillary line in the 5th intercostal space above the nipple line
- Capnospot® Pneumothorax Decompression Indicator Procedure:
- Use a minimum 14g x 3.25" catheter specifically designed for needle decompression
- Attach Capnospot[®] Decompression Indicator to the catheter prior to insertion
- Insert needle with syringe attached at a 90⁰ angle, just over the superior border of the rib, & advance until air is freely aspirated or a "pop" is felt, then advance only the catheter until the hub rests against the skin



- Observe for color change from blue to yellow within 10 secs to confirm catheter placement. Color change may not be reliable in patients with an open pneumothorax. Observe for clinical indicators of successful placement.
- Simplified Pneumothorax Emergency Air Release (SPEAR®) Procedure:
- Insert in accordance with manufacturer's directions for use
- Adequately secure catheter
- If an initial attempt at 1 approved site is unsuccessful, consider utilizing an alternate approved site
- 2 attempts allowed on affected side(s) without base/modified base hospital contact

POST-PROCEDURE

- Reassess breath sounds
- Administer high flow O₂ & monitor SpO₂
- Continuous cardiac & EtCO₂ monitoring
- Assess & document vital signs every 3-5 mins (if possible)
- Monitor Capnospot[®] (if used) & breath sounds for signs of development of tension pneumothorax



Venous Blood Draws

Approval: Troy M. Falck, MD – Medical Director

Effective: 12/01/2024

Approval: John Poland – Executive Director

Next Review: 07/2027

INDICATIONS

- Paramedics or AEMTs may perform blood draws on pts with a medical complaint, when there is an agreement to do so in place between the EMS provider agency & the receiving hospital
- Paramedics may perform chemical testing blood draws at the direction of law enforcement (LE) under the following parameters:
- Fire department/district employees are not allowed to perform chemical testing blood draws
- Personnel must be authorized to perform chemical testing blood draws by their employer
- Medical treatment & emergency calls take precedence over chemical testing blood draw requests

PRE-PROCEDURE

Assess for & provide medical treatment as indicated/appropriate

MEDICAL BLOOD DRAW PROCEDURE

- Select appropriate equipment & site:
- If drawing blood from an IV catheter, attach blood draw adapter to the IV catheter hub & draw blood sample prior to IV fluid administration
- If no IV has been established, or if IV fluids have been administered, prep site with an appropriate disinfectant agent, place tourniquet 3 4 inches above collections site & perform venipuncture
- Insert the blood tubes in the following order (releasing tourniquet when blood starts to flow):
 Blue, Red, Green, Purple
- Apply slight pressure to the site with a gauze pad & secure with tape
- Gently invert each tube a few times (do not shake or mix vigorously)
- Label samples as follows:
- Patient name & date of birth
- Date & time of blood draw
- EMS unit number
- Place labeled tubes in a specimen collection bag & turn over to appropriate hospital staff
- Adequately document medical blood draws on the PCR

CHEMICAL TESTING BLOOD DRAW PROCEDURE

- Suspects shall be in LE custody and shall consent to the blood draw if the suspect refuses or is unable to consent, the paramedic shall stop the procedure immediately
- Paramedics shall not draw blood on a struggling or restrained suspect
- Blood draw kits shall be supplied by the requesting LE agency
- Alcohol or other volatile organic disinfectant shall not be used to clean the skin at the draw site a suitable aqueous disinfectant (normally included in the LE supplied blood draw kit) shall be utilized
- The arresting officer must be present when the blood draw is performed & the blood sample is the property of the arresting officer
- In addition to routine incident information, the paramedic shall document the following on the PCR:
 Blood draw kit number
- Requesting officer's name & badge number
- Suspect/Pt's consent for the procedure
- Skin prep used and site of blood draw(s)



Vascular Access

Approval: Troy M. Falck, MD – Medical Director

Approval: John Poland – Executive Director

Effective: 12/01/2024

INDICATIONS

• Vascular access may be established by authorized EMS personnel when there is a current or anticipated need to administer intravenous medications/fluids.

ADDITIONAL DIRECTIONS/CONSIDERATIONS

- Do not delay transport to establish vascular access unless clinically necessary.
- Avoid establishing vascular access in an extremity with a functioning dialysis shunt unless no other vascular access is available/appropriate.
- Intraosseous (IO) access or external jugular (EJ) vein cannulation shall only be attempted if unable to establish peripheral vascular access & immediate medication/fluid administration is necessary.
- Preexisting Vascular Access Devices (PVADs) may be utilized for pts in extremis when no other vascular access is available/appropriate.
- Limit vascular access attempts to three (3) unless necessary for emergent treatment.
- Do not connect the primary IV tubing directly to the IV catheter. IV extension/saline lock tubing shall be utilized between the primary IV tubing and the IV catheter.

INTRAOSSEOUS (IO) ACCESS

Contraindications:

- Fracture/suspected vascular compromise in targeted bone or infection at area of insertion site.
- Excessive tissue or absence of adequate anatomical landmarks.
- Previous significant orthopedic procedure at site or IO access in targeted bone within past 48 hours.

Procedure:

- Prep selected site (see images) with a recognized antiseptic agent & wipe dry with a sterile gauze pad.
- Insert device per manufacturer specific instructions.
- Attach primed extension set to needle & secure needle per manufacturer instructions.
- For pts unresponsive to pain:
 - Rapid flush with 10 mL of normal saline.
- For pts responsive to pain:
 - Prime extension set with 2% lidocaine.
- Slowly administer 2% lidocaine over 120 sec.
 - Adult pts 40 mg.
 - Pediatric pts 0.5 mg/kg (max: 40 mg).
- Allow lidocaine to dwell in IO space 60 sec.
- Rapid flush with 10 mL of normal saline.
- Slowly administer a subsequent ½ dose of 2% lidocaine over 60 sec.
- Connect fluids to extension set infusion may need to be pressurized to achieve desired rate.
- Dress site and secure tubing.





Vascular Access

EXTERNAL JUGULAR (EJ) VEIN CANNULATION

Contraindications:

- Suspected coagulopathy (e.g. advanced liver disease, anti-coagulant medications)
- Suspected cervical spine injury
- Inability to tolerate supine position

Procedure:

- Place pt in Trendelenburg or supine position and elevate shoulders.
- Turn head 45° 60° to side opposite of intended venipuncture site.
- Palpate to assure no pulsatile quality to vessel.
- Prep site with recognized antiseptic agent & wipe dry with a sterile gauze pad.
- 'Tourniquet' vein by placing finger just above clavicle near midclavicular line.
- Stabilize skin over vein with thumb.
- Point needle toward shoulder in direction of vein & puncture vein midway between jaw & clavicle, over belly of sternocleidomastoid muscle.
- Maintain compression of vein at clavicle area until needle is withdrawn & IV tubing has been connected.
- Secure IV site.

PREEXISTING VASCULAR ACCESS DEVICE (PVAD) UTILIZATION

Contraindications:

• Subcutaneous access requiring special equipment & entry through the skin is not approved for use by EMS personnel

Procedure:

- Do not remove injection cap from catheter.
- Do not use a syringe smaller than 10 ml to prevent catheter damage from excess infusion pressure.
- Always expel air from syringe prior to administration.
- Follow all medications with 5 ml of saline to avoid clots.
- Do not inject medications or fluids if resistance is met when establishing patency.
- Do not allow IV fluids to run dry.
- Do not manipulate or remove an indwelling catheter under any circumstances.
- Should damage occur to the external catheter, clamp immediately between the skin exit site & the damaged area to prevent air embolism or blood loss.