



Greater Miami Valley EMS Council



2026-2028

Standing Orders

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Acknowledgement

Dear Region,

I hope these standing orders find you and your departments well. This live document serves as an approach to care in the pre-hospital setting. The efforts by this region, its providers, and those who volunteer their time, is what makes the delivery possible. The changes and progress made in this document are culminated via feedback from the regional surveys, changes in the state scope of practice, studies presented in progressive treatments, committee meetings with the hospital networks as well as agency leadership. We then assess relevance for the region and consistency for rural volunteer departments to busy urban departments, and interfacility transport is ensured. At times this progress can seem delayed. The goal is for the progression to be based off of science, well-established studies, and national review. This allows for consistent, well established patient care practices.

This region has provided a vast amount of welcomed input. These items were researched, discussed in public forum, and presented to the Regional Physicians Advisory Board for further discussion and approval. Of the numerous items that were presented, we are proud to say that over ninety-percent of the topics brought by the region were approved and are now part of this document. As a region we accomplished this.

The intention of this document is to provide a guided approach to patient care. It should be noted that not all situations and variables can be accounted for, nor does this document serve as a limitation to the decisions and chosen methods of each department's Medical Director when utilizing other treatments outside the GMVEMSC Drug bag. Online medical direction is always an acceptable approach when items are not addressed or don't align with the current situation. The structure of this document is supportive by design and not rooted in a punitive nature.

As a region we are successful and functional in EMS care. We cordially request continued input and feedback for the purposes of growth and progression. A deeply gracious "thank you" to everyone who takes part in medical care in the region and all those who have taken any of their time to serve the public in our region and be part of the continued growth.

At the posting of these standing orders, we have already started the process for improvement in the next two-year cycle. If any changes need to be made, please know that you have a voice regardless of your role in the system. Please continue to complete the regional surveys and take part in the bi-monthly Standing Orders and Regional Advisory Board Committee meetings.

Benjamin Valiton
Standing Orders Co-Chair



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Appendices

Appendix A	2026-2028 Protocol Changes
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Change Order Directives

COD 2026-1	No CODs Currently
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1000 Series

General Protocol

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Subject:

Introduction to Protocols

Effective:

June 1, 2021

Last modified:

May 15, 2026

1001.1 Introduction to Treatment Protocols

- a. Each protocol has been approved by the Greater Miami Valley EMS Council and the Regional Physician Advisory Board for Region 3 (as defined by the State Board of Emergency Medical, Fire and Transportation Services [EMFTS]).
- b. Each tab bears an effective date and a last modified date marking it as the latest version.
- c. A new addition to protocol would reflect a duplicate “Effective” and “Last Modified” date.
- d. When changes or revisions are made to a tab, only the “Last Modified” date will be changed.
- e. Each time changes or additions are made; they can be referred to by their specific line in the protocol. i.e. *A change was made to “1001.1.e”.*
- f. Each cycle, changes or additions will be listed in an addendum in the appendix.
- g. The date of release of the protocol will be in the lower right box of the title page.
- h. After release of a new version, all previous versions of the standing orders should be considered invalid.

1001.2 Printing, Retention, and Display

- a. All GMVEMSC Treatment Protocols are intended for color printing, and hard copy retention.
- b. These protocols are also intended for electronic display in Adobe Portable Document Format (PDF).
 - i. The PDF version includes links to the different tabs throughout the document.
 - ii. The GMVEMSC logo on most pages is a hyperlink back to the table of contents.
- c. Distribution is provided by means of the GMVEMSC official website.

1001.3 Application

- a. This protocol is for use by those individuals operating in and under the authority of the Greater Miami Valley EMS Council (GMVEMSC) Drug Bag Exchange Program and certified by the State of Ohio as an EMS provider.
- b. The provider must pass both the skills check-off and Computer Based Testing (CBT) for the current year.
- c. The GMVEMSC Treatment Protocols apply to the following certification levels:
 - i. Emergency Medical Responder (EMR)
 - ii. Emergency Medical Technician (EMT)
 - iii. Advanced Emergency Medical Technician (AEMT)
 - iv. Paramedic (PM)

1001.4 Stipulations

- a. The protocol is to be used in the field only.
- b. Communicate with the receiving facility as soon as practical:
 - i. When transporting unstable patients
 - ii. Transporting to hospitals that request contact for all patients delivered to their facility.
- c. No procedures, techniques, or drugs will be used without the proper equipment or beyond the training or capabilities of the prehospital personnel.
- d. Nothing in this protocol may be used without specific pre-approval of the Medical Director for the local department or agency.
- e. The protocol is to be utilized as clinically indicated. Not every standing order in a treatment protocol must be carried out on every patient treated under that treatment protocol.



Subject:

Introduction to Protocols

Effective:

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- f. Discretionary judgment is required and stepwise adherence to specific protocols may not be in the patient's best interest.
- g. At no time should treatment options exceed those authorized without direct consultation with the Medical Control Physician (MCP).

1001.5 Protocol Design

- a. The GMVEMSC protocols are organized around the General Patient Management Protocol which must be followed for all patients. This universally applicable protocol/flowchart allows the providers to integrate additional treatment protocols beyond general patient management as clinically necessary for specific patient care, emergency stabilization, and treatment.
 - i. As an example, while caring for a specific patient with chest pain, shortness of breath, and nausea the provider would:
 1. Follow the General Patient Management Protocol
 2. Integrate and follow the Chest Pain Protocol
 3. Integrate and follow the Respiratory Distress Protocol if indicated
 4. Integrate and follow the Cardiac Alert Protocol if indicated
 5. Integrate and follow the Abdominal Pain Protocol if indicated
 6. Refer to protocol for specific medication concentrations, dosages, and volumes.
 7. Complete the General Patient Management Protocol
- b. In most cases, a specific guideline will only be mentioned once within the protocol. All other circumstances where that guideline would be applicable will simply refer to the original guideline.
- c. Where applicable, a guideline mentioned in another section will have a hyperlink provided.
- d. Formatting
 - i. All attempts will be made to keep the protocol focused and specific.
 - ii. Extracurricular and enhancing information will be provided in an official study guide.
 - iii. All levels of providers will be addressed within a single protocol.
 - iv. Procedures and treatments marked with a diamond (♦) always require a physician's order.
 - v. Items enclosed in brackets ({ }) are at the option of the agency and their Medical Director.
 - vi. *Sections that apply only to adults are bulleted with an "A".*
 - vii. *All pediatric treatments will be in pink and bulleted with a "P".*
 - viii. There are also sections which apply to only Geriatric patients and are bulleted with a "G."

1001.6 Clinical Management Tables

- a. In addition to general statements, this protocol will utilize table-based algorithms where applicable.
- b. The table will demonstrate what care can be given at each provider level.
 - i. The level of certifications will be signified by the colored tabs to the right of each section.
- c. Even with a step-by-step algorithm in place, critical thinking is encouraged.
- d. While the table is sequential and listed by provider level, many elements in each section can be completed simultaneously.
- e. The following is an annotated example of a Clinical Management Table:



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Assessment			
Pediatric Considerations - This is where pediatric specific info might go. - Dosing and treatment will still be listed in the algorithm	Signs & Symptoms This is where S&S will go	Differential Diagnosis This is where differentials will go	
Treatment Algorithm			
- This will be where guidelines for all certification levels will go - Any EMR and above information will be listed in this box.			EMR
- Treatment directives for the EMT and above will be here. - If no EMT directives apply, then this box would read "No additional orders at this level".			EMT
- Treatment directives for the AEMT and above will be here. - If no AEMT specific directives apply, then this box would read "No additional orders at this level".			AEMT
- Treatment directives for the Paramedic will be listed here. - If no Paramedic specific directives apply, then this box would read "No additional orders at this level".			Paramedic
Consult			
- If requirements exist for any level to call for orders, that will be listed here. - If there is a guideline to call an alert, that will be listed here. - If there is a recommendation to call for MCP advice, that will be listed here. - If there is a request to call the receiving facility prior to arrival, that will be listed here.			
Clinical Pearls			
- Any important guidelines or clinical information will be added here. - This will not be a study guide nor a skill sheet. That information will be supplied in a separate format.			
END OF SECTION			

1001.7 Protocol Layout

- a. The protocol will be broken down into sections, each with a different series number or title. This version will be separated into the following:
 - i. 1000 Series – General Patient Management
 - ii. 2000 Series – Cardiac Protocol
 - iii. 3000 Series – Trauma Protocol
 - iv. 4000 Series – Medical Protocol
 - v. 5000 Series – Pediatric Considerations
 - vi. 6000 Series – Special Operations Protocol
 - vii. 7000 Series – Administration
 - viii. 8000 Series – EMS Drug Formulary
- b. Appendices
 - i. Any supporting materials not found in the standing orders can be added to the Appendix section.
 - ii. Each new version of standing orders will include a listing of all major changes, identifying the tab, section and specific edit, addition or deletion.
- c. Change Order Directives
 - i. This section will identify change orders introduced within the two-year life cycle of the protocol.
 - ii. Change orders will be introduced when a needed edit to the standing orders cannot wait until the next edition of the protocol. Previously, these came in the form of long-term just-in-time (JITSO) orders.



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- iii. Change Order Directives will be converted into permanent standing order changes at the end of the two-year cycle and the release of a new edition.
- iv. Each change order will be given a unique sequential number and include the specifics of the change.
- v. The change will also be reflected in the appropriate actual standing order.
- vi. When a change order is issued, a new Standing Orders will be released.
- vii. Each individual Change Order Directive will utilize the following format:

COD 20XX-#	Topic: This is where the title describing the subject of the change order will go		
Level of Providers Affected			
EMR	EMT	AEMT	Paramedic
Standing Order Tabs Affected			
XXXX.x.x – This will be the Tab, Sections and subsections of the change will be identified			
Change Orders			
1	This is where the specific addition, deletion, modification or edit of the change will be listed individually		
2	If there is more than one change to a specific order, they can be listed out on as many lines as needed		
Effective Date of Change: The effective date of the change will go here			

END OF SECTION



Subject: Communication with Hospital or Medical Control

Effective: June 1, 2021

Last modified: May 16, 2026

1002.1 Reasons to Contact the Hospital

- a. To notify the hospital when time is needed to prepare for patient arrival. Examples include:
 - i. Cardiac arrest
 - ii. Any of the defined alerts such as Cardiac Alert, Stroke Alert, Trauma Alert
 - iii. Pre-Arrival notification of behavioral patients (e.g. Dr. White, ACE, etc.)
 - iv. Indications of sepsis
 - v. Significant communicable disease
 - vi. Other serious patients that may require acute care
 - vii. Hazardous material exposures (*mandatory*)
 - viii. Bedbugs

1002.2 Reasons to Contact Medical Control

- a. To obtain orders for procedures or medications as indicated within the protocol.
- b. For field termination or DNR clarification.
- c. To obtain advice in a difficult situation or circumstance. Examples include:
 - i. Before a medication is given, even though protocol allows it to be used without permission.
 - ii. A situation where the patient has an unfamiliar condition.
 - iii. To discuss a destination decision.

1002.3 Call-in Procedures

- a. When contacting a hospital, make sure a clear picture is painted.
- b. When calling about a trauma patient, include:
 - i. MIVT – Mechanism, Injuries, Vital Signs and Time
 - ii. Estimated time of arrival (ETA)
 - iii. The components of the Glasgow Coma Score (GCS)
 - iv. Patient assessment findings which are relevant to the decision to transport to a Trauma Center.
- c. When calling to discuss a case with the Medical Control Physician, give the call taker a brief synopsis of the call and the reason for needing to speak with a physician:
 - i. After the summary, state your reason to need to speak to the Medical Control Physician.
 - ii. When calling with an Alert (Cardiac, Stroke, Trauma, etc.):
 - 1. Request to speak directly to the Medical Control Physician after the brief summary.
 - 2. Verbalize to the MCP (or in some cases the RN), "We recommend a _____ Alert."
 - 3. The MCP has the discretion to withhold the Alert and may decide not to activate it.

END OF SECTION

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Subject: Non-Initiation of Care

Effective: June 1, 2021

Last modified: Feb. 11, 2024

1003.1 General Guidelines

- a. This protocol may be applied by EMT, AEMT and Paramedic providers only. The EMR cannot determine that a patient is deceased.
- b. All patients (Adult, **Pediatric**, and Geriatric) may meet criteria for non-initiation of care.
- c. If care had begun and is readily apparent to the provider that the patient meets non-initiation of care criteria, **RESUSCITATION EFFORTS MAY CEASE.**

1003.2 Criteria for Non-Initiation of Care

- a. Resuscitation will not be initiated in the following circumstances:
 - i. Deep, penetrating, cranial injuries
 - ii. Massive truncal wounds
 - iii. DNR Order—present and valid (see [1004 Do Not Resuscitate](#))
 - iv. Frozen body
 - v. Rigor mortis, tissue decomposition, or severe dependent lividity
 - vi. Triage demands
 - vii. For patients in arrest resulting from **BLUNT OR PENETRATING TRAUMA** consider not initiating care for injuries obviously incompatible with life.
 1. Prolonged arrest (greater than 10 minutes)
 2. Consider possibility of MIXED MECHANISMS

1003.3 Exclusionary Conditions

- a. The following conditions will not meet non-initiation of care criteria:
 - i. Traumatic arrest in female patient with either:
 1. Known pregnancy greater than 24 weeks or
 2. Uterine fundus palpable at or above the umbilicus
 - ii. Possible medical etiology for traumatic cardiac arrest
 - iii. Arrest witnessed by EMS providers
 - iv. Lightning strike
 - v. Signs or symptoms of a hypothermic patient
 - vi. Focused blunt trauma to the chest, (commotio cordis)

1003.4 For an inquiry about organ donation, direct the call to Life Connection of Ohio at 1-800-535-9206.**END OF SECTION**

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Subject:

Do Not Resuscitate

Effective:

June 1, 2021

Last modified:

Jan. 19, 2025

1004.1 General Guidelines

- a. Per ORC [2133.01-2133.26](#), providers will consider and honor all valid Ohio Do Not Resuscitate orders.
- b. The two valid DNR orders are DNR: Comfort Care and DNR: Comfort Care Arrest.

1004.2 Do-Not-Resuscitate Orders Defined

- a. Do-Not-Resuscitate: Comfort Care Arrest (DNR-CCA)
 - i. Permits any GMVEMSC Protocol treatment until the patient goes into cardiac or respiratory arrest.
 - ii. Once the patient meets the above criteria, then only permitted DNR treatment is performed.
- b. Do-Not-Resuscitate: Comfort Care (DNR-CC)
 - i. Permits any medical treatment to diminish pain or discomfort
 - ii. No treatment should be used to postpone the patient's death.
 - iii. The order is initiated at the moment it is signed by the patient's physician.

1004.3 Permissible and Impermissible Treatments Once the DNR is Initiated

- a. The following treatments are permitted:
 - i. Conduct an initial assessment
 - ii. Perform basic medical care
 - iii. Clear airway of obstruction or suctioning
 - iv. If necessary, for comfort or to relieve distress, may administer oxygen, CPAP or BiPAP
 - v. If necessary, may obtain IV access for hydration or pain medication to relieve discomfort, but not to postpone death
 - vi. If possible, may contact other appropriate health care providers
- b. The following treatments are not permitted once an order is valid and effective:
 - i. Perform CPR
 - ii. Administer resuscitation medications with the intent of restarting the heart or breathing
 - iii. Insert an airway adjunct
 - iv. Defibrillation, cardioversion or initiate pacing
 - v. Initiate continuous cardiac monitoring

1004.4 Stipulations

- a. If more than one living will declaration or DNR exists, the most recent supersedes the previous.
- b. The authority of a DPOA-HC supersedes the DNR if the DPOA-HC previously consented to the DNR.
- c. The GMVEMSC protocol will recognize the following special situations as valid. If these scenarios present, then contact MCP and request to honor the DNR with physician permission.
 - i. Out-of-State DNR orders
 - ii. **Pediatric DNR orders**
- d. Blood glucose checks and treatment of [4008 Diabetic Emergencies – Hypoglycemia/Hyperglycemia](#), is acceptable even with a valid DNR.
- e. While [1005 General Patient Management](#) requires continuous cardiac monitoring when administering pain medications, this focused protocol supersedes that requirement in valid DNR patients.
- f. In situations where there are questions about the documents, try to keep the patient's intent in mind.
- g. If there is any confusion on scene, ♦ Call MCP for clarification

END OF SECTION

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Subject:

General Patient Management

Effective:

June 1, 2021

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June 6, 2026

1005.1 Guideline

- a. The General Patient Management protocol is to be applied to all patients.
- b. Once a primary impression and differential diagnosis is made, then the provider should look to specific treatment algorithms within these standing orders.
- c. Unless explicitly addressed, the GMVEMSC Protocol will not specify or endorse any type or brand of medical equipment. It is the responsibility of agency leadership and their medical direction to determine the appropriate equipment needed for medical care and insure that personnel are trained on it's use.

1005.2 Basic Patient Care

- a. The emphasis in patient care should ensure airway protection, oxygenation, and adequate ventilation without causing harm.
- b. Injury reduction strategies may include noninvasive ventilation when appropriate, titration of oxygen in certain settings, and being cautious not to over ventilate.
- c. Tailor treatment to the overall clinical picture.
- d. With the exception of suspected acute cerebral herniation, the rate and depth of ventilation in the prehospital setting should not be guided by the EtCO₂ reading alone.
- e. For the patient with cerebral herniation, ventilate the patient at approximately 20 times per minute to obtain an EtCO₂ of 30 mmHg.
- f. "Permissive hypercapnia" in most cases is appropriate, particularly in those with chronic lung disease who may chronically retain CO₂.
- g. It is recommended to listen to the chest to ensure that adequate exhalation is occurring during manual ventilation.

1005.3 EMT Assisting the Advanced Provider

- a. Per Ohio Revised Code, the EMT is permitted to assist the advanced provider with skills that are outside of the EMT's scope of practice.
- b. The EMT is only allowed to prepare ALS equipment under the direct supervision of the AEMT or Paramedic.
- c. The skills that an EMT may set up for and assist with are:
 - i. Endotracheal intubation
 - ii. Intravenous access
 - iii. IV fluid administration
 - iv. Saline locks
 - v. Placement of 4 and/or {12/15 Lead EKG} for cardiac monitoring
 - vi. Accessing the GMVEMSC Drug Bag to locate drugs and/or to assemble pre-jects.

1005.4 General Patient Management

Assessment		
Pediatric Considerations • Pediatric patients are defined as patients less than 16 years old • A Pediatric reference guide or length-based resuscitation tape may be used to reference pediatric equipment recommendations. • Pedi-Wheel may be used as a reference for pediatric vital signs. • Unless otherwise specified, the maximum dose for pediatric medication administration is the adult dose.	Signs & Symptoms • None	Differential Diagnosis • None
Treatment Algorithm		



Subject: General Patient Management

Effective: June 1, 2021

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- Scene/Crew Safety/PPE; with appropriate equipment/medications to patient side.
- Initial Assessment/Physical Exam
- Follow basic life support and airway algorithms as indicated based on current AHA guidelines.
- An unresponsive patient with gasping breaths and poor color should get supplemental oxygen via BVM
- Obtain chief complaint, OPQRST, SAMPLE history, and other pertinent information.
- Vital Signs
 - Blood Pressure (EMR are limited to obtaining manual blood pressures)
 - Pulse, rate and quality
 - Respirations; Rate, quality, and work-of-breathing
 - Assess every 5 to 15 minutes per patient condition
 - Temperature as needed
- Utilize monitoring devices, pulse oximeter, CO-oximetry, capnography, etc. as appropriate and approved by medical direction.

EMR

- Perform blood glucose check.
- Where indicated, the EMT may obtain a {12 Lead EKG} for the purpose of transmission.
- The EMT may assist the advanced provider with:
 - {12 /15 Lead EKG} application assisting a Paramedic who is present
- Set up an IV administration kit in the presence of an AEMT or Paramedic

EMT

- Utilize cardiac monitor as appropriate.
- Where indicated, the AEMT may obtain a {12 Lead EKG} for the purpose of transmission.
- The AEMT may apply a {12/15 Lead EKG} when assisting a Paramedic who is present.
- Start IV crystalloid solutions or saline lock as appropriate.
- **IV Therapy:** Follow [4016 Shock Protocol](#).
 - Lactated Ringers (pH and electrolyte balanced fluids) are preferred
 - For medical emergencies, head trauma, cardiac issues with stable BP, etc.: Use **TKO** rate.
 - Shock (not related to penetrating trauma):
 - Run **IV fluid** wide-open
 - Use macro-drip or blood tubing
 - Decrease fluid rate if SBP greater than 100
- **P IV fluid 20 ml/kg using macro-drip tubing. Titrate to maintain adequate perfusion.**
- Use of IO devices for both adults and **pediatrics** is limited to patients who are unresponsive or hemodynamically unstable, and only when less invasive means are not available or are ineffective (e.g. **Narcan** IN and **Versed** IN).
- Provide continuous cardiac monitoring, EtCO₂ and pulse oximetry (if available) for all patients with fentanyl, ketamine, morphine or midazolam if not already doing so.
- ♦ If a patient with an existing IV pump experiences an allergic reaction, consider discontinuing the pump.

AEMT

- Use of an {IV pump} is optional for any agency with approval from their Medical Director.
- Existing central venous catheters, dialysis catheters, fistulas, or grafts may be utilized for infusion of IV fluids and medication if the patient is hemodynamically unstable. These may also be used when the patient is deteriorating rapidly.

Paramedic

Consult

- Do not stop the flow of medication in an established medication pump except under direct orders from Medical Control.
- If a patient with an existing IV pump experiences an allergic reaction, call the MCP for an order to discontinue the pump.

Clinical Pearls

- If a patient was discharged from a hospital in the last 24 hours, it is recommended to return to the same facility or at the very least, the same network of hospitals.
- If the patient is experiencing complications from a recent surgery, if possible, transport the patient back to the facility where the surgery was performed. If that is not practical, then try to transport to the same network.
- If possible, bring medications or a list of the medications to the hospital; include the dose and frequency of administration.
- Crystalloid fluids include Lactated Ringers, Normosol, Plasmalyte, or Normal Saline in that order. Their pH is closer to neutral.
 - Medical emergencies, head trauma, cardiac problems with stable BP: Use TKO rate.
- IV medication administration: **Slow IV = over 2 minutes**, unless otherwise specified.
- Any medication given intravenous can also be administered intraosseous.
- Maintain normothermia.

END OF SECTION



Subject: Patient Abuse and Neglect

Effective: June 1, 2021

Last modified: Dec. 7, 2024

1006.1 Guideline

- a. EMS providers MUST, by law, report all alleged or suspected **pediatric** and adult abuse/neglect.
- b. Ohio Revised Code requires providers to report incidents of **pediatric** and adult abuse/neglect to:
 - A Their county's adult protective services agency (for patients over 60 years old)
 - P Their county's public children services agency**
 - iii. Or for both adults and **pediatrics**; Law enforcement
 - iv. For adult patients see ORC [5101.63](#) and for **pediatric patients** see ORC [2151.421](#)
- c. Simply notifying hospital personnel does not meet mandated EMS reporting responsibilities.
- d. Hospitals have copies of the EMS Social Services Referral Form, supplied by GDAHA, for documenting cases of abuse/neglect.
- e. Use of this form can help providers in providing information needed to their reporting agency, as well as provide for a continuum of care with hospital social services departments.
- f. Document on the Patient Care Report, all efforts that EMS made to report the suspected abuse; include name of agency notified, method used, and name of person contacted.

1006.2 Pediatric Abuse and Neglect**P Report all alleged or suspected child abuse or neglect to the appropriate agency.**

Pediatric Public Social Services Agencies			
County	Phone	After Hours Phone	Fax
Butler	513-887-4055	513-868-0888	513-887-4260
Champaign	937-484-1500	Contact County SO: 937-484-6092	937-484-1506
Clark	937-327-1700	937-324-8687	937-327-1910
Darke	937-548-7129	937-548-2020	937-548-8723
Greene	937-562-6600	937-372-4357	937-562-6650
Miami	937-335-4103	Contact County SO: 937-440-3965	937-339-7533
Montgomery	937-224-5437	937-224-5437 (same as daytime)	937-276-6597
Preble	937-456-1135	937-456-1135 (same as daytime)	937-456-6086
Shelby	937-498-4981	Contact County SO : 937-498-1111	937-498-1492
Warren	513-695-1558	513-695-1600	513-695-1800

1006.3 Adult Abuse or Neglect**A Report all alleged or suspected abuse or neglect to the appropriate agency.**

Adult Public Social Services Agencies			
County	Phone	After Hours Phone	Fax
Butler	513-887-4081	Contact County SO: 513-785-1000	513-785-5969
Champaign	937- 484-1500	Contact County SO: 937-484-6092	937-484-1506
Clark	937-327-1700	937-324-8687	937-327-1910
Darke	937-548-7129	937-548-2020	937-548-4928
Greene	937-562-6315	Contact County SO: 937-562-4800	937-562-6177
Miami	937-440-3471	Contact County SO: 937-440-3965	937-335-2225
Montgomery	937-225-4906	Contact County SO: 937-225-4357	937-496-7464
Preble	937-456-1135	937-456-1135 (same as daytime)	937-456-6086
Shelby	937-498-4981	Contact County SO: 937-498-1111	937-498-1492
Warren	513-695-1420	513-425-1423	513-695-2940

END OF SECTION

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Subject: Basic Airway Maintenance

Effective: June 1, 2021

Last modified: July 1, 2026

1007.1 Clinical Management

Assessment																				
Pediatric Considerations Repeated and prolonged suctioning could cause hypoxia and bradycardia <table><tr><th colspan="4">Respirations by Age</th></tr><tr><td>Up to 1 year</td><td>30-60</td><td>7-9 years</td><td>16-24</td></tr><tr><td>1-3 years</td><td>20-40</td><td>10-14 years</td><td>16-20</td></tr><tr><td>4-6 years</td><td>20-30</td><td>15+ years</td><td>12 -20</td></tr></table>		Respirations by Age				Up to 1 year	30-60	7-9 years	16-24	1-3 years	20-40	10-14 years	16-20	4-6 years	20-30	15+ years	12 -20	Signs & Symptoms - Respiratory difficulty or distress - Poor SpO₂ or EtCO₂ - Mechanism of Injury or Nature of Illness that would require O₂ therapy - Impending airway issues - Adventitious respiratory sounds		Differential Diagnosis None
Respirations by Age																				
Up to 1 year	30-60	7-9 years	16-24																	
1-3 years	20-40	10-14 years	16-20																	
4-6 years	20-30	15+ years	12 -20																	
Treatment Algorithm																				
- EtCO₂ monitors can be used on all patients with or without adequate perfusion, and with or without artificial airways. - Administer Oxygen as needed. Use the following rates as guidelines: 2 LPM by nasal cannula (NC) for patient with COPD, or as prescribed. 4-6 LPM by nasal cannula (NC) for other patients. 12-15 LPM by non-rebreather mask (NRM) for any patients with increased respiratory rates or effort (including COPD). - Ventilate patients who are symptomatic with an insufficient respiratory rate, depth or effort. P Patient less than 2 years old showing respiratory distress with nasal congestion, cough, rales, rhonchi or wheezing - without previous history of wheezing, reactive airway disease, breathing treatments: P Nasopharyngeal suctioning in both nares (3-5 seconds) with an appropriate device P If distress continues, repeat nasopharyngeal suctioning for 3-5 seconds P For patients less than 6 years old showing respiratory distress with agitation, upper airway noise, stridor, and/or “barky cough,”: P Lower temperature of ambulance as much as possible. P Deliver oxygen as the patient tolerates. P Often these symptoms resolve with less intervention. P Consider keeping distance from the patient.				EMR																
- Consider patient airway anatomy for the appropriate selection of the airway adjunct. - If indicated, suction the tracheostomy. P If patient has history of reactive airway disease with prescribed breathing treatments then treat with 4003 Asthma protocol. - Consider the need for a supraglottic or dual lumen rescue airway. - The EMT may only place a rescue airway in a pulseless, apneic patient. - For guidelines to placement of rescue airways, see 1008 Advanced Airway Management ♦ Oxygen flow rate for nebulized medications should be 8-10 LPM. ♦ Nebulized medication may be administered while ventilating with a BVM. Preferably use two oxygen sources.				EMT																
- Consider the need for intubation. - The AEMT may only intubate if patient is apneic. - If routine ventilation procedures are unsuccessful, try to visualize obstruction with laryngoscope. - If a foreign body is seen, attempt to remove it using suction or Magill forceps.				AEMT																
- When deciding whether to intubate, consider the following: - Insufficient respiratory rates, less than 10 or greater than 29, that are not rapidly controlled by other measures - Irregular respiratory rhythm - Abnormal breath sounds - Inadequate chest expansion and respiratory depth - Excessive effort to breathe - Use of accessory muscles - Nasal flaring - Pallor or cyanosis - Cardiac dysrhythmias					Paramedic															
Consult																				
The EMT needs MCP ordered to administer nebulized medications.																				
Clinical Pearls																				
- COPD patients in severe respiratory distress or with chest pain need the same O₂ devices and flow rates as any other patient in such condition. - CPAP and {BiPAP} are to be considered forms of “non-invasive ventilation”, or aggressive ventilatory support that avoids or eliminates the need for intubation.																				
END OF SECTION																				

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Subject: Advanced Airway Management

Effective: June 1, 2021

Last modified: May 16, 2026

1008.1 Clinical Management

Assessment			
Pediatric Considerations None	Signs & Symptoms - Patient unable to manage their own airway - Patient in cardiac arrest - Patient in respiratory arrest (AEMT & Paramedic) - Rapidly collapsing airway	Differential Diagnosis None	
Treatment Algorithm			
Advanced Airway Management is not an EMR skill			EMR
- The EMT may only place a rescue airway in a pulseless, apneic patient - If approved, “rescue airways” such as the Supraglottic Airways or Dual Lumen Airways are appropriate airway devices for both adult and pediatric patients. - Confirm correct placement of advanced airways by at least 5 methods, see protocol 1009 Advanced Airway Confirmation Devices - Reassess advanced airway placement every time the patient is moved.			EMT
- An AEMT may only intubate if patient is apneic. - Consider patient airway anatomy and condition for proper advanced airway device selection. - If a total of two attempts with an ET tube are not successful, move to a rescue airway. P Supraglottic airway is recommended as the <u>primary airway</u> except in extreme cases such as airway edema. - Always secure the ET tube in place, preferably with a commercial tube-securing device. - A cervical collar is effective in maintaining patient’s head in a neutral position during the intubation process. - If there are indications of tension pneumothorax and the patient is hemodynamically unstable: - Decompress the chest with a 14-gauge or larger, 3 ¼” angiocath - Location options include: - Fourth or fifth intercostal space in the mid-axillary line - Second or third intercostal space in the mid-clavicular line P less than 8 years old, site choice will only be the second or third intercostal space at the mid-clavicular line			AEMT
- Approved advanced airways satisfy the “rescue airway” component for 1010 {Sedate-to-Intubate or RSI}. - If a conscious patient requires intubation, consider the following: A Apply Lidocaine Jelly to the ET tube. A Lidocaine 100 mg IN (half dose per nostril) or nebulized with 8-10 LPM O₂. P Lidocaine 1.5 mg/kg nebulized with 8-10 LPM O₂ or IN. Maximum dose is 100 mg. - If the patient resists the tube after confirmed intubation: A SBP is greater than 100, consider Midazolam 2.5 mg slow IV. A SBP less than 100, consider Ketamine 100 mg slow IV. G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses. P SBP is age/weight appropriate consider Midazolam 0.1 mg/kg (max dose 2.5 mg), slow IV. A {If a patient needs intubation but is combative, agitated, or has jaws clenched, use 1010 {Sedate to Intubate or RSI} procedures if approved to do so by Medical Direction.} - Whenever all reasonable attempts to provide an adequate airway by less invasive means have failed due to a total airway occlusion and you are unable to ventilate: A Perform a needle cricothyrotomy or surgical airway utilizing an approved method. P Patient must be 8 years old or greater for a surgical airway. {With defined and specific indications, paramedics who have received appropriate training and Medical Director’s approval may perform a finger thoracostomy in accordance to their specific protocols}			Paramedic
Consult			
None			
Clinical Pearls			
- For the EMT, AEMT and Paramedic, Dual Lumen Airways, King Airway or Laryngeal Mask Airways (LMA), are acceptable airway devices. - For the AEMT and Paramedic, {Lighted Stylet Intubation} or {Camera Assisted Intubation} may be utilized. - For the Paramedic, Nebulized Lidocaine can be administered simultaneously with Albuterol and Ipratropium. - If feasible, wait one to two minutes before intubation			
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Subject: Advanced Airway Confirmation Devices

Effective: June 1, 2021

Last modified: May 15, 2026

1009.1 General Guidelines

- a. Confirm correct placement of advanced airways with waveform capnography and at least 4 other methods as listed below.
- b. Reassess advanced airway placement every time the patient is moved.

1009.2 Confirmation Methods

Assessment		
Pediatric Considerations	Signs & Symptoms	Differential Diagnosis
None	Inserted advanced airway	None
Treatment Algorithm		
Advanced Airway Management is not an EMR skill		
- Advance airway device confirmations (Utilize at least 5 methods after airway insertion): - Continuous EtCO₂ detection is mandatory for advanced airway confirmation - Auscultate the epigastrium, the lungs at the anterior chest, the lungs at the mid-axillary areas, and then the epigastrium again for ventilation sounds. - Observe rise and fall of the chest with each breath - Look for condensation in the tube of the advanced airway - Look at patient's appearance - If signs of cerebral herniation are present, hyperventilate at 20 ventilations per minute to an EtCO₂ value of 30 mmHg.		
- For ETT depth placement and measurement: A Keep the oral endotracheal tube at a minimum depth of tube size x 3 at the gum line/teeth. P Proper endotracheal tube depth can also be calculated by tube size x 3 at the gum line/teeth. - Avoid placing the ETT too deeply and the possibility of a right or left main stem bronchus intubation. - If confirmations indicated right or left main stem intubation, retract tube slightly till bilateral sounds are heard. - Do not confuse right main stem intubation for a pneumothorax. - Reassess endotracheal tube placement with at a minimum, EtCO₂ and lung sounds after every patient movement		
No additional orders at this level		
Consult		
None		
Clinical Pearls		
- Intravenous sodium bicarbonate will produce more carbon dioxide and affect EtCO₂ values. - End tidal capnography should be maintained through transfer to the hospital		

1009.3 Confirmation Devices

- a. These devices can help recognize esophageal intubation, but cannot identify bronchial placement.
- b. Maintain EtCO₂ devices until patient care is transferred to the receiving ED staff.
- c. Electronic End Tidal CO₂ (EtCO₂) Monitors (Capnography)
 - i. Continuous waveform capnography is a required confirmation device.
 - ii. EtCO₂ should be used on **EVERY** advanced airway
- d. End Tidal CO₂ Detector (EtCO₂) – Colorimetric
 - i. In cardiac arrest, if there is no color change, use other confirmation methods.
 - ii. Secretions, emesis, etc. can ruin the device.
 - iii. Large amounts of carbonated beverage in the stomach can give a false positive.
 - iv. The device can be used for no more than two hours.
 - v. Follow manufacturer's recommendations for weight restrictions.



Subject: Advanced Airway Confirmation Devices

Effective: June 1, 2021

Last modified: May 15, 2026

1009.4 D.O.P.E Advanced Airway Troubleshooting

- a. In the event of a deteriorating condition or presence of symptoms of acute hypoxia in the patient with an advanced airway, consider the D.O.P.E mnemonic:
- Displacement of the equipment – is the tube or advanced airway dislodged, too deep or out of the airway entirely? Check markers and verify positioning.
 - Obstruction (of the tube) – Is the advanced airway blocked by secretions (mucus plug), blood, or kinking?
 - Pneumothorax – Has a tension pneumothorax developed? Check for lung sounds and asymmetry.
 - Equipment Failure – Is there a failure in the oxygen source, ventilator tubing, or connections? Disconnect and switch to manual bagging (BVM) to verify.

END OF SECTION



Subject: {Sedate to Intubate or RSI}

Effective: June 1, 2021

Last modified: Feb. 19, 2025

1010.1 General Guidelines

- a. Sedate to Intubate and Rapid Sequence Intubation are optional skills in the GMVEMSC protocol.
- b. These skills are to be performed by the Paramedic only.
- c. This standing order applies to agencies whose personnel have received the appropriate training and Medical Director's approval only.
- d. Under no circumstances is RSI to be used as "behavioral control" or restraint in patients with otherwise intact airways.
- e. Some Medical Directors may recommend RSI as a primary airway control procedure.
- f. While this protocol recommends Succinylcholine as a short-term paralytic, a Medical Director may choose to use a different medication. Should a different paralytic be used, the Medical Director will be responsible to establish dosing and training.
- g. Inclusion criteria:
 - i. The patient must be an adult (16 years old or older)
 - ii. The patient cannot have suffered a paralyzing injury more than one week and less than 6 months ago (this guideline is specific to Succinylcholine)

1010.2 Clinical Management

Assessment			
Pediatric Considerations This protocol does not apply to pediatric patients.	Signs & Symptoms - Decreased LOC - Ineffective or absent breathing - Patient unable to maintain their own airway - Respiratory failure or inevitable loss of airway	Differential Diagnosis - Cardiac arrest - Anaphylaxis - Esophageal obstruction	
Treatment Algorithm			
Neither Sedate-to-Intubate nor Rapid Sequence Intubation are EMR skills			
Neither Sedate-to-Intubate nor Rapid Sequence Intubation are EMT skills			EMT
Neither Sedate-to-Intubate nor Rapid Sequence Intubation are EMT skills			A EMT
{Sedate-to-Intubate}: {Pre-oxygenate the patient with O₂ via BVM at 15 lpm} {Complete an airway assessment. Remove dentures or dental appliances. {If the paramedics doubt that they will be able to successfully intubate, the procedure should be avoided} } {Must have cardiac monitor, IV and pulse oximetry in place} {Sedate the patient}: {Administer Etomidate 0.3 mg/kg IV (maximum dose 40 mg)} OR {Ketamine 100 mg IV, may repeat 100 mg IV if patient not sufficiently sedated by first dose} (Recommended in hemodynamically unstable patients) OR {Midazolam 5 mg slow IV (in patients who are normotensive), may repeat up to 10 mg if patient not sufficiently sedated by first dose} <u>DO NOT</u> reduce Ketamine or Midazolam doses by half for patients greater than 69 y/o. Instead, give full doses to all patients to achieve sedation. - If stopping at {Sedate-to-Intubate}, then intubate the patient. {Rapid Sequence Intubation}: {Sedate the patient as outlined above} {Paralyze the patient with Succinylcholine 200 mg IV} {Once paralyzed, intubate the patient, and maintain continuous waveform capnography} {Maintain sedation}: {Midazolam 5-10 mg IV} {If hypotensive, then Ketamine 100-200 mg IV} - Maintain continuous waveform capnography after intubation.			

Paramedic



Subject: {Sedate to Intubate or RSI}

Effective: June 1, 2021

Last modified: Feb. 19, 2025

Consult

- Paramedics may seek guidance or approval from medical control prior to initiating the protocol; however, this is not required.

Clinical Pearls

- Paralytics or sedation do not change poor airway anatomy.
- The most important decision may be when NOT to paralyze the patient or intubate them.
- Succinylcholine paralyzes the muscles but does not affect LOC. ALWAYS SEDATE THE PATIENT.
- Tachycardia may be a sign that the patient is paralyzed but not adequately sedated.
- No more than 2 intubation attempts.
- If you can still ventilate the patient with a BLS airway, a cricothyroidotomy is not necessary.

1010.3 RSI Educational Recommendations

- Rapid Sequence Intubation should not be available to all paramedics in the system.
- Only those paramedics willing to undergo additional initial training and continuing training should be allowed to perform it.
- In initial training, the paramedic should demonstrate proficiency during the following practical evaluations:
 - 2 endotracheal intubations on airway simulators
 - 3 endotracheal intubations on airway simulator with C-spine immobilization
 - 5 surgical cricothyrotomies on simulators using surgical technique or an approved device
 - 4 intubations using the eschmann stylet (gum bougie) on airway simulators (*optional*)
 - 5 insertions of a rescue airway on airway simulators
- Once a quarter, the paramedic should demonstrate proficiency during the following practical evaluations:
 - 1 endotracheal intubation on airway simulators
 - 2 endotracheal intubations on airway simulator with C-spine immobilization
 - 1 surgical cricothyrotomy on airway simulator
 - 1 intubation using the eschmann stylet (gum bougie) on airway simulators (*optional*)
 - 1 insertion of rescue airway on airway simulators
- Any of the above evaluations could be credited if the procedure is performed under direct supervision by the Medical Director, Supervisor or Training Officer the field or a clinical setting.

END OF SECTION



Subject: Tracheostomy and Laryngectomy Care

Effective: June 1, 2021

Last modified: Dec. 8, 2021

1011.1 General Guidelines

- a. Consult the patient's caregiver for assistance. They are typically trained to manage these airways.
- b. Find out why they have an artificial airway (cancer, stroke, ventilator dependent, etc.)
- c. Ask if there have been any prior difficulties (reinserting, plugging, etc).
- d. Find out when the airway was first placed (newer airways may be more difficult to replace).
- e. For assessing failed tracheostomies and laryngectomies, consider:
 - i. D - displaced, dislodged or damaged
 - ii. O - obstructed (mucus, food, blood, secretions)
 - iii. P - pulmonary problems
 - iv. E - equipment failure (bent tubing, ventilator malfunction, depleted oxygen supply)
- f. Look for subcutaneous air in the neck as it might indicate a false passage of tube.

1011.2 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms Patient with tracheostomy or laryngectomy tube with signs of respiratory distress or failure	Differential Diagnosis None
Treatment Algorithm		
- Place the patient on high-flow O₂ - Assess pulse oximetry - Assess EtCO₂ if available - Assess and frequently reassess the artificial airway for easily reversible causes of distress (D.O.P.E.) - Administer high-flow oxygen over the stoma or tube. - Consider assisting ventilations using a bag-valve-mask attached to the device end. - BVM typically will only attach over the inner cannula - If there is no inner cannula, an endotracheal tube adapter (BVM end of ETT) a half size larger than the trach tube may be inserted into the outer cannula. - Consider infant BVM to stoma ventilation if the tracheostomy or laryngectomy tube has been removed.		EMR
- Pre-oxygenate when possible for 30-60 seconds prior to suctioning - Suction the tracheostomy tube if: - Unable to ventilate with BVM. - Coarse upper airway sounds are heard. - If respiratory distress continues despite BVM ventilation. - If the airway tube has an inner cannula, remove it prior to suctioning. - Use the patient's suctioning supplies or a catheter that is no more than 1/2 the tube diameter. (typical size is 10 fr) - DO NOT force the suction catheter into the tracheostomy tube. - Determine the proper suction catheter depth by measuring the length of the obturator or inner cannula and advancing slightly beyond this measure. - If no obturator is available: A Insert the suction catheter 2-3 inches into the tube. P Use the patient's pinky finger as an approximate length to insert the suction tubing. - Consider inserting 2 - 3 mL of saline or nebulized saline to help loosen thick or hard secretions. - Suction on the way out, for no more than 10 seconds, rotating the catheter as you go. - If respiratory distress continues, consider likely cause and reference appropriate protocol.		EMT
- Place patient on cardiac monitor. - If measures have not succeeded in improving respiratory status, consider replacing the airway tube as defined in 1011.3 - If no replacement tube is available, insert an ETT as a replacement.		AEMT
If all other means fail, including tube replacement, consider attempting oral tracheal intubation.		Paramedic
Consult		
None		



Subject: Tracheostomy and Laryngectomy Care

Effective: June 1, 2021

Last modified: Dec. 8, 2021

Clinical Pearls

- Patients with laryngectomy airways have the larynx removed, completely separating oral- and nasal- pharynx from the trachea and lungs.
 - These patients are sometimes referred to as neck breathers.
- Established stomas are less likely to close off.
 - Closed off stomas require surgical techniques to replace the tube and replacement should be avoided in the field.
- Often the cuff is deflated allowing the patient to have more air movement past the vocal cords thus enabling speech.
- There may also be speaking valve (a one-way valve allowing air in – not out) attached to the outside end of the tracheal tube.
- Tube replacement is a clean procedure (mask, splash protection, and clean gloves). Keep the patient's airway as clean as possible.

1011.3 Artificial Airway Tube Replacement (AEMT & Paramedic)**a. Necessary Equipment:**

- i. Replacement tracheostomy tube or laryngectomy tube (from the patient or care giver).
 1. If patient is pediatric, there is a one size smaller tracheostomy tube in the GoBag that should always be with the patient.
- ii. If no replacement tracheostomy tube is available, use an ETT of similar internal diameter
- iii. If possible, water-based lubricant jelly.

b. Procedure:

- i. Apply high-flow O₂, pulse oximetry, EtCO₂, and cardiac monitor.
- ii. Place patient semi-recumbent with slight neck extension (consider a roll under the neck).
- iii. Keep the head midline (may need additional personnel to maintain head position).
- iv. For adults, consider use of a bougie when removing the old tube. (this is not a pediatric practice)
- v. Lubricate the new tracheostomy tube or replacement ETT.
- vi. Deflate the old tracheostomy tube's balloon and remove during exhalation by gently pulling and rotating towards the patient's feet.
- vii. Remove the stoma dressing, then wipe area clean with only saline or medically packaged water.
- viii. Using the replacement tracheostomy tube's obturator or (in adults only) the bougie, gently advance the replacement tracheostomy tube in a fluid fashion, using the natural curvature of the tube until the flange is flush against the neck.
- ix. If present, remove the obturator and insert the hollow internal cannula
 1. Internal cannulas are not part of the most commonly used tracheostomy tubes for pediatric patients).
 2. If possible, use a non-fenestrated (no window) inner cannula.
 - a. Note: A fenestrated inner cannula will allow air leak through the glottis; potentially allowing air to enter the stomach and not allowing PEEP (positive end-expiratory pressure) to be achieved.
- x. If using an ETT as a replacement:
 1. Insert a bougie (adults only) into the stoma directed downward.
 2. Slowly advance the lubricated ETT into the stoma.
 3. Only advance the ETT a few centimeters into the stoma (as deep as the trach tube).
 4. Consider shortening the ETT by cutting the tube AFTER the takeoff for the pilot balloon.
- xi. Inflate the cuff of the replacement tracheostomy tube or ETT with the minimum amount of air to stop any audible leak at the stoma.
- xii. Place clean gauze around the stoma to absorb mucous.
 1. Never cut this gauze.



Subject: Tracheostomy and Laryngectomy Care

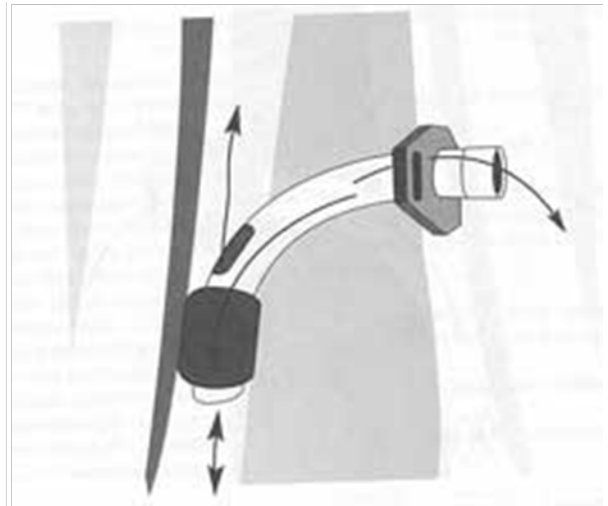
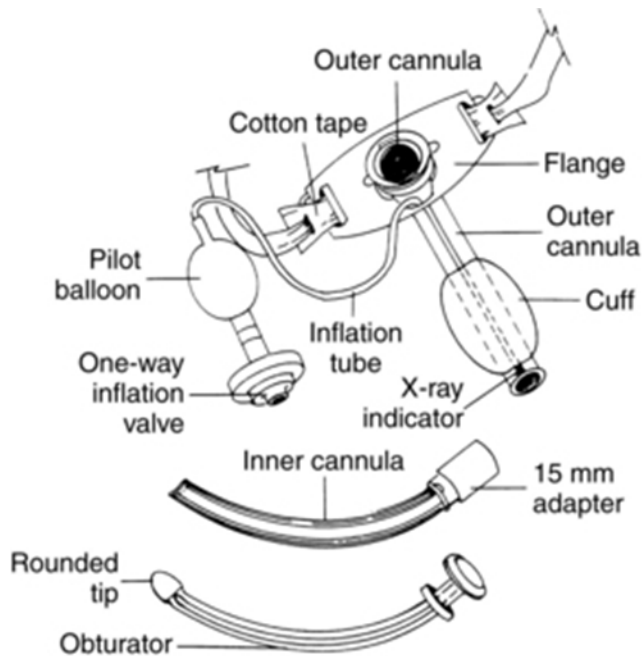
Effective: June 1, 2021

Last modified: Dec. 8, 2021

2. Fold it to size, to avoid creating small particulates of lint that could enter the airway.
- xiii. Secure the device to the patient's neck.

c. Emergency Procedures

- i. If the airway has been surgically altered and the glottis is hard to recognize, consider pushing on the chest to force air into the pharynx.
- ii. Where air bubbles are seen, insert bougie (in adults) and/or insert the ETT into the opening.



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Subject: Intraosseous Infusion

Effective: June 1, 2021

Last modified: May 15, 2026

1012.1 General Guidelines

- a. Use of IO devices is limited to patients who are unresponsive or hemodynamically unstable; and then, only when less invasive means are ineffective or not available.
- b. In patients with acceptable perfusion, and all other routes of access have failed, then consider an intraosseous access of in the following acceptable locations, in no particular order:
 - i. The proximal humeral head (must be greater than 12 years old)
 - ii. The distal femur
 - iii. The proximal tibia
- c. For an adult in cardiac arrest, the preferable order of vascular access is:
 - i. External jugular (EJ) vein IV
 - ii. Antecubital (AC) vein IV
 - iii. Proximal humeral head or distal femur IO (if the distal femur is used, better access should be sought as the cardiac arrest scenario progresses)
- d. For equipment sizing, follow manufacturers recommendations.

1012.2 Intraosseous Equipment Sizing

- a. The longer yellow (55 or 45 mm) needle should be used for proximal humeral IOs in adults.
- b. For the distal femur or proximal tibia in adults, use the properly sized needle.

P For pediatrics, access the distal femur or proximal tibia:

P Use the (25 mm or blue) IO needle for 3-45 kg.

P Use the (15 mm or pink) IO needle for 0-3 kg.

1012.3 Clinical Management

Assessment		
Pediatric Considerations Consider weight for IO selection	Signs & Symptoms Hemodynamically unstable patient needing vascular access with no IV	Differential Diagnosis None
Treatment Algorithm		
IO Insertion is not an EMR skill		EMR
IO Insertion is not an EMT skill		EMT
- After IO confirmation, IV pressure bags may facilitate infusion. - For the pain associated with infusion: A Lidocaine 2% 1.5 mg/kg via IO up to 100 mg. P Lidocaine 2% 0.5 mg/kg via IO (max 100 mg)		AEMT
No additional orders at this level		Paramedic
Consult		
None		
Clinical Pearls		
For the AEMT and the Paramedic, all approved medications and fluids that can be administered IV can be administered in the IO		
END OF SECTION		

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Subject:

Alternate Vascular Access

Effective:

June 1, 2021

Last modified:

Dec. 8, 2020

1013.1 General Guidelines

- a. This guideline is not for EMR, EMT or AEMT. Only Paramedics may utilize alternative vascular routes.

1013.2 Central Vascular Access Devices (CVAD)

- a. Patients who require long-term intravascular therapy may have Central Vascular Access Devices (CVAD).
- b. CVADs may be used for IV access if the patient is hemodynamically unstable or in arrest.
 - i. Central catheter: Catheter placed through chest wall into the internal jugular or subclavian vein.
 - 1. Central catheters can be single or multilumen.
 - 2. Distal portion of catheter has two access ports, either of which may be used for access.
 - ii. PICC Line: Catheter placed in arm.
 - 1. Distal portion of catheter is external with access port.
 - 2. Do not force fluids or drugs through the device or failure could result in an embolism.
 - 3. PICC line diameter creates significant resistance to fluid flow making it difficult to infuse large quantities of fluids.
 - 4. Dextrose 10% (D10) by PICC is preferable to IM Glucagon.
 - iii. Subcutaneously Implanted Port: Device surgically placed under the skin on the chest.
 - 1. No external access. PARAMEDICS ARE NOT PERMITTED TO ACCESS THIS DEVICE.
- c. Complications of CVADs
 - i. Infection: Thorough cleaning of the port must be done three times during the procedure:
 - 1. Before attaching each syringe
 - 2. Before attaching the IV tubing.
 - ii. Air Embolism: The catheter must be clamped before attaching or removing the syringes.
 - iii. Heparin Bolus: These catheters remain in place without fluids continually flowing through them. To prevent blood clot formation, a bolus of Heparin or other anticlotting agents will be in the catheter. Remove 5 ml of blood to ensure that the Heparin is not systemically administered to the patient.
 - iv. Catheter Damage:
 - 1. Use a 10 ml syringe or larger when drawing off the blood. Smaller syringes create too much pressure.
 - 2. After verifying blood return, flush catheter with 10 ml of NS with a 10 ml or larger syringe utilizing a pulsating technique.
 - 3. Administer medications slowly to avoid creating too much pressure. Do not use catheter if unable to get blood return.
 - v. DO NOT USE A PRESSURE INFUSION DEVICE ON CVADs.

1013.3 Internal Dialysis Fistula

- a. An artificial passage between an artery and a vein used to gain access for hemodialysis.
- b. Usually located in the inner aspect of the patient's forearm or bicep.
- c. A bulge under the skin that should be visible or easily palpated.
- d. In cardiac arrest or with a profoundly unstable patient, a dialysis fistula may be used to administer IV fluids or medication.
 - i. Use aseptic technique.
 - ii. Be careful not to puncture back wall of vessel.
 - iii. Use IV pressure bag.
 - iv. Blood may still back-up into tubing.
 - v. Control bleeding with direct pressure.
 - vi. Dialysis patients are usually on anticoagulants.

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Subject: Pain Management

Effective: June 1, 2021

Last modified: Feb. 27, 2025

1014.1 General Guidelines

- This protocol is for management of acute moderate to severe pain, including pain from suspected cardiac events, trauma (including thermal and chemical burns), crush syndrome, frostbite, fractures, dislocations, sprains, and abdominal pain (including unilateral flank pain).
- It is not for the treatment of exacerbations of chronic pain.
- Prehospital pain management reduces time to pain relief, avoids exacerbation of pain during movement, is compassionate, and is good medical care.
- Ketamine** is not to be administered to patients with suspected cardiac chest pain

1014.2 Clinical Management

Assessment					
Pediatric Considerations • Fentanyl is <u>not</u> to be administered to anyone less than 2 years old • If unable to obtain a blood pressure, look for evidence adequate perfusion (skin color, capillary refill, and mental status) prior to Fentanyl administration. • To account for medication remaining in the needle and syringe, add an additional 0.1 ml Fentanyl for pediatric intranasal doses. • Ketamine is <u>not</u> to be administered for pain to anyone less than 16 years old • Fentanyl IN, is the first choice for pediatrics	Signs & Symptoms • Severity of pain (pain scale) • Quality (sharp, dull, etc.) • Radiation of pain • Pain upon movement • Increased pain upon palpation	Differential Diagnosis • Chronic pain			
Treatment Algorithm					
• Use ice packs, position of comfort, and splinting to reduce pain. • Provide oxygen as indicated.		EM		Paramedic	
• No additional orders at this level.		EMT			
• For an alert patient with moderate to severe pain, give Fentanyl for relief. ◦ Ketamine should be considered a second line medication for the management of pain ◦ If Fentanyl dosing does not relieve pain or if the patient refuses Fentanyl, then administer Ketamine • ♦ Call for orders if you feel narcotics are needed for pain from a chronic condition. G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses. A If SBP is greater than 100, then Fentanyl 50-100 mcg IV A May repeat Fentanyl 50-100 mcg IV after 5 minutes. A If no IV, Fentanyl 50-100 mcg IN, SQ, or IM A May repeat Fentanyl 50-100 mcg IN, SQ or IM after 10 minutes. P If SBP is normal for patient's age (80 + 2 times age) or evidence of adequate perfusion then Fentanyl 1 mcg/kg IN, max 100 mcg P May repeat Fentanyl 1 mcg/kg IN, max 100 mcg after 10 minutes P If unable to administer IN, Fentanyl 1 mcg/kg IV, max 100 mcg P May repeat Fentanyl 1 mcg/kg IV, max 100 mcg after 5 minutes P As a last resort, Fentanyl 1 mcg/kg SQ or IM, max 100 mcg P May repeat Fentanyl 1 mcg/kg SQ or IM, max 100 mcg after 10 minutes A Ketamine 25 mg IV A May repeat Ketamine 25 mg IV after 5 minutes A If no IV, Ketamine 25 mg IN or 50 mg IM A May repeat Ketamine 25 mg IN or 50 mg IM after 10 minutes			AEMT		
• No additional orders at this level					
Consult					
P MCP contact required before administration of Fentanyl for pediatric patients with <u>abdominal pain</u>.					
Clinical Pearls					
• Always consider the weight of your patient when dosing pain meds, especially for the elderly. • Document patient's reported pain during initial patient contact, during treatment, and after any intervention.					
END OF SECTION					

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2000 Series

Cardiac Protocol

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Subject: Resuscitation Guidelines

Effective: June 1, 2021

Last modified: May 26, 2026

2001.1 General Guidelines

- a. A patient's BEST CHANCE for resuscitation is at the scene with high quality CPR and code management.
- b. Paramedics are expected to provide ALS resuscitative care at the scene.
- c. All providers should prioritize on-scene resuscitation efforts, regardless of age, unless clinically indicated.
- d. These standards apply to all patient populations, including both adults and **pediatrics**.

2001.2 Resuscitation and Field Termination

FIELD TERMINATION DOES NOT APPLY TO PEDIATRIC PATIENTS	- Pulseless and apneic - Does not meet Non-initiation of Care Guideline	Meets Non-initiation of Care Guideline
The EMR will continue resuscitation until the patient is handed off to a higher-level provider.		
- The EMT will continue resuscitation until the patient is handed off to a higher-level provider. - If no higher-level provider is available, then transport. - Patient with return of spontaneous circulation (ROSC) should be transported to an interventional facility if transport time is less than 30 minutes. ♦ If no ALS equipment is available at the scene, and transport time to a medical facility will exceed 20 minutes, field termination may be considered.		
- Patients will require prolonged resuscitation efforts if: - They have PEA with a rate greater than 40 per minute - They have an upward trending or persistent EtCO₂ greater than or equal to 20 mmHg, refractory to VF or VT. - If arrest due to profound hypothermia, then rapidly transport to a Trauma Center. ♦ Following all appropriate efforts, field termination requires MCP approval, and may only be considered when the following criteria are met: 18 years or older - In asystole or PEA, with rates less than 40 - Not be in arrest due to hypothermia - Have an advanced airway in place - Have vascular access in place - There are no signs of neurological function such as reactive pupils, response to pain or spontaneous movement		
- The following should be transported to a cardiac interventional facility if transport time is less than 30 minutes: - A documented STEMI and a witnessed cardiac arrest - A return of spontaneous circulation (ROSC)		
A When the AEMT or Paramedic contacts MCP directly to receive consent for field termination, they must provide the following information: - The duration of the resuscitation - How long the patient may have been in arrest prior to EMS arrival - Whether it was a witnessed or unwitnessed event - The current EtCO₂ - The presenting rhythm		
- There are situations where resuscitation may take 30 minutes or more. - Research has shown that CPR quality diminishes while being transported. - Consider aeromedical transport for transports greater than 30 minutes if the patient has ROSC. - In pseudo PEA, the patient may not be in true cardiac arrest, but simply not have palpable pulses due to profound shock.		
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Subject: Cardiac Arrest - BLS

Effective: June 1, 2021

Last modified: May 20, 2026

2002.1 This protocol has adopted the 2025 American Heart Association CPR Guidelines

	ADULTS	CHILDREN	INFANTS	NEWBORNS
CPR Order	CAB: Compression, Airway, Breathing			
Compression to Breaths Ratio <u>Without</u> Advanced Airway	1 or 2 Rescuers 30:2	1 Rescuer - 30:2 2+ Rescuers - 15:2		3:1
Compression to Breaths Ratio <u>With</u> Advanced Airway	Continuous compressions at a rate of 100-120 /min. Give 1 breath every 6 seconds.	Continuous compressions at a rate of 100-120 /min. Give 1 breath every 2-3 seconds.		40-60 breaths/min
Compression Rate	100 to 120 per minute			120 per minute
Compression Notes	Minimize interruptions in chest compressions. Limit interruptions to less than 10 seconds			
Compression Depth	At Least 2 Inches	1/3 Depth of Chest (About 2")	1/3 Depth of Chest (About 1 ½ ")	1/3 Depth of Chest
Rescue Breathing	1 breath every 5-6 seconds (10-12 breaths/min)	1 breath every 2-3 seconds (20-30 breaths/min)		40-60 breaths/min

2002.2 Basic Life Support

Assessment				
Pediatric Considerations If available, use age-appropriate AEDs or pads	Signs & Symptoms - Unresponsive - Pulseless and apneic		Differential Diagnosis - Signs of irreversible death - Other causes of unresponsiveness	
Treatment Algorithm				
- If witnessed or unwitnessed arrest, initiate quality CPR for 1-2 minutes - If available, initiate mechanical CPR using an approved device - Attach and use AED after at least 2 minutes of CPR - Utilize AED as it is programmed. (Even if it is not to AHA guidelines) - Repeat cycles of defibrillation and CPR for 2 minutes			EMR	AEMT
- Patient should be transported as appropriate - Patient with return of spontaneous circulation (ROSC) should be transported to an interventional facility if transport time is less than 30 minutes. - Obtain and transmit {12 Lead EKG} if patient has ROSC			EMT	
No additional orders at this level				
- Paramedics are expected to provide resuscitative care for all age groups at the scene. - P Pediatric cardiac arrest patients should receive at least a first dose of epinephrine and adequate ventilation (BVM or advanced airway) prior to transport, unless clinically indicated. - Cardiac arrests should not be transported unless: - When indicated, at least one defibrillation has been delivered. - Return of spontaneous circulation (ROSC) - The airway cannot be secured - Vascular access is not established - MCP declines to authorize Field Termination				
Consult				
No consult required unless applying Field Termination Guideline				
Clinical Pearls				
- Use jaw-thrust method to open airway on trauma patients - Allow the chest to fully recoil after each compression - Change person compressing chest every 2 minutes - Resume CPR beginning with compressions after each defibrillation - Minimize interruptions to compressions before and after each shock to less than 10 seconds - For pregnant patients in cardiac arrest; consider need for manual uterine displacement				
In all cardiac arrests, consider the ACLS treatable causes (Hs & Ts) to your level of certification:				
EMR - Hypoxia - Hypothermia	EMT Toxins	AEMT - Hypovolemia - Hydrogen Ion - Tension pneumothorax	Paramedic - Tamponade, Cardiac - Thrombosis (Coronary, Pulmonary)	
END OF SECTION				

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Subject: Cardiac Arrest: Asystole or PEA

Effective: June 1, 2021

Last modified: Oct. 10, 2021

2003.1 General Guidelines

- In all cardiac arrest patients, apply the [2002 Cardiac Arrest: Basic Life Support](#) protocol.
- Apply the appropriate guideline after rhythm interpretation.
- The rhythms may change and will require flexibility to move between the different protocols.
- If ROSC, then follow [2001 Resuscitation Guidelines](#)

2003.2 Clinical Management

Assessment		
Pediatric Considerations Pediatric dosing should never exceed adult doses	Signs & Symptoms - Unresponsive - Pulseless and apneic - Either: - No electrical activity on cardiac monitor - Electrical activity on monitor with no pulse present	Differential Diagnosis - Ventricular Fibrillation - Pulseless Ventricular Tachycardia - Other causes of unresponsiveness - Device (lead) error - Signs of irreversible death
Treatment Algorithm		
- If witnessed or unwitnessed arrest, initiate quality CPR for up to 2 minutes. - Follow 2002 Cardiac Arrest -BLS protocol - Apply the Automatic External Defibrillator (AED) and check for a shockable rhythm. - If no defibrillation is indicated, continuous CPR		EMR
Obtain and transmit {12 Lead EKG } if patient has ROSC		EMT
- Consider possible causes - Consider Field Termination as identified in 2001 Resuscitation Guidelines		AEMT
A Epinephrine (1:10,000) 1 mg, IV or IO, repeat every 3-5 minutes. P Epinephrine (1:10,000) 0.01 mg/kg, IV or IO, repeat every 3-5 minutes. The Paramedic may consider Field Termination after administering Epinephrine		Paramedic
Consult		
- No consult required unless applying Field Termination Guideline. - The AEMT or paramedic may consult MCP to field terminate. - Contact ED to request a Cardiac Alert if applicable.		
Clinical Pearls		
Contact receiving hospital prior to arrival with a cardiac arrest patient		
END OF SECTION		

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Subject: Cardiovascular Emergencies –
Renal Failure/Dialysis

Effective: June 1, 2021

Last modified: Dec. 24, 2024

2004.1 General Guidelines

- This protocol is for cardiac patients who receive renal dialysis treatment and is only to be administered by Paramedics.
- Dialysis patients who are bradycardic or experience cardiac arrest should be given both calcium (chloride or gluconate) and sodium bicarbonate.

2004.2 Clinical Management

Assessment				
Pediatric Considerations Pediatric dosing should never exceed adult doses	Signs & Symptoms - Cardiac arrest - Bradycardia - Confirmed history of renal dialysis		Differential Diagnosis None	
Treatment Algorithm				
No additional orders at this level			EMR	Paramedic
No additional orders at this level			EMT	
No additional orders at this level			AEMT	
- For renal dialysis patients in arrest: - A Calcium Chloride 10% 1 g IV - P Calcium Chloride 10%, 20 mg/kg (0.2 ml/kg) IV (max dose 500 mg) - A Sodium Bicarbonate 100 mEq IV - P Sodium Bicarbonate 1 mEq/kg IV ♦ For a renal dialysis patient presenting with a wide complex bradycardia: - A Calcium Chloride 10% 1 g IV. - P Calcium Chloride 10%, 20 mg/kg (0.2 ml/kg) IV (max dose 500 mg) - A Sodium Bicarbonate 100 mEq IV - P Sodium Bicarbonate 1 mEq/kg IV				
Consult				
In the treatment of hyperkalemia (wide complex bradycardia)				
Clinical Pearls				
- It is critical that these drugs not be given together, as they will precipitate. - Flush well between these medications.				
END OF SECTION				

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Subject: Cardiac Arrest:
V-Fib or Pulseless V-Tach

Effective:
June 1, 2021

Last modified:
July 1, 2026

2005.1 General Guidelines

- In all cardiac arrest patients, apply the [2002 Cardiac Arrest: Basic Life Support](#) protocol.
- Apply the appropriate guideline after rhythm interpretation.
- The rhythms may change and will require flexibility to move between the different protocols.
- If ROSC, then follow [2001 Resuscitation Guidelines](#)

2005.2 Clinical Management

Assessment		
Pediatric Considerations Pediatric dosing should never exceed adult doses	Signs & Symptoms - Unresponsive - Pulseless and apneic - Ventricular fibrillation or ventricular tachycardia on cardiac monitor or AED	Differential Diagnosis - Asystole - Artifact/Device failure - Signs of irreversible death - Other causes of unresponsiveness
Treatment Algorithm		
- If witnessed or unwitnessed arrest, initiate quality CPR for 1-2 minutes and proceed to first defibrillation - Follow Basic Life Support protocol - Defibrillate as indicated by the Automatic External Defibrillator (AED)		EMR
Obtain and transmit {12 Lead EKG} if patient has ROSC		EMT
- Defibrillate as required based on EKG interpretation - A {For refractory V-Fib/PVT after three shocks, consider Vector Change Defibrillation for subsequent shocks} - Consider possible causes		AEMT
- Alternate between CPR/Defibrillation/Medication Administration - A Epinephrine 1 mg 1:10,000, IV or IO, repeat every 3-5 minutes - P Epinephrine (1:10,000) 0.01 mg/kg, IV or IO, repeat every 3-5 minutes - After third defibrillation: - A Amiodarone 300 mg, IV or IO - P Amiodarone 5 mg/kg IV or IO (max first dose 300 mg) - If Amiodarone is not available, use Lidocaine - A Lidocaine 150 mg, IV or IO - P Lidocaine 1.0 mg/kg IV or IO (max first dose 100 mg) - A With evidence of pulseless polymorphic ventricular tachycardia (torsades de pointes), Magnesium Sulfate 2 gm infused with macro-drip tubing over 10 minutes. - A {After three traditional defibrillations and at least one antiarrhythmic medication, consider Double Sequential Defibrillation for subsequent shocks} - After sixth defibrillation: - A Amiodarone 150 mg, IV or IO - P Amiodarone 5 mg/kg IV or IO (max first dose 150 mg) - If Amiodarone is not available, use Lidocaine - A Lidocaine 75 mg, IV or IO - P Lidocaine 1.0 mg/kg IV or IO (max first dose 75 mg) - If patient converts with ROSC from a ventricular arrhythmia and no anti-arrhythmic has been given, then: - A Amiodarone 150 mg in 250 ml NS, IV over 10 minutes using 60 drop/ml tubing - Do not infuse unless SBP is greater than 100 - Consider IV fluid 500 ml to increase SBP to 100 or higher prior to infusion		



Subject: Cardiac Arrest:
V-Fib or Pulseless V-Tach

Effective:
June 1, 2021

Last modified:
July 1, 2026

Consult

- No consult required unless applying Field Termination Guideline.
- The AEMT or paramedic may consult MCP to field terminate.
- Contact ED to request a Cardiac Alert if applicable.

Clinical Pearls

- For initial and subsequent defibrillations, follow manufacturer recommendation for energy settings
- Pediatric defibrillation settings will start at 2 J/kg (or biphasic equivalent) and increase by 2 J/kg (or biphasic equivalent) each shock.
- Maximum pediatric shock will be 10 J/kg (or biphasic equivalent)
- Resume chest compressions immediately following each defibrillation, without performing pulse check, for 1-2 minutes
- Contact receiving hospital prior to arrival with a cardiac arrest patient

2005.3 Alternate Defibrillation Techniques

- Vector Change and Double Sequential Defibrillation are optional skills in the GMVEMSC Protocol
- Providers SHOULD NOT apply these techniques without the explicit consent of their Medical Director.
- These procedures are approved for adult patients only.
- Vector Change Defibrillation (for Advanced EMTs and Paramedics)
 - This technique is for refractory ventricular fibrillation/pulseless ventricular tachycardia.
 - Refractory V-Fib/PVT is defined as NOT CONVERTED by three standard defibrillations.
 - The AEMT or Paramedic will place a second set of defib pads in an anterior-posterior position.
 - There should be minimal interruption in CPR when placing the second set of pads.
 - Subsequent defibrillations will be through the anterior-posterior placed pads.
 - NOT indicated for Recurrent V-Fib/PVT which is defined as V-Fib/PVT that reoccurs episodically after successful conversion with intervening episodes of organized electrical activity.
- Double Sequential Defibrillation (for Paramedics)
 - This technique is for refractory V-Fib/PVT following three standard defibrillations and a least one round of an antiarrhythmic agent (amiodarone or lidocaine).
 - This requires the presence of two manual biphasic defibrillators.
 - One set of pads will be placed in the anterior-apical (traditional) position and one set will be placed in the anterior-posterior position.
 - With both sets of pads in place and both machines charged to maximum energy level, the discharge of the monitors should be as simultaneous as possible.
 - Repeat as indicated. All subsequent defibrillations should be double sequential.
 - CAUTION: Every agency considering applying this procedure needs to consult with the manufacturer of their cardiac monitor for advice. This technique is considered "off-label".
 - Agency Administrators should be made aware of the risk to monitors if the DSD technique is utilized.
- Neither Vector Change nor Double Sequential Defibrillation is indicated in Recurrent V-Fib/PVT, which is defined as V-Fib/PVT that reoccurs episodically after successful conversion with intervening episodes of organized electrical activity.

END OF SECTION



Subject: AICD Activations

Effective: June 1, 2021

Last modified: May 17, 2023

2006.1 General Guidelines

- a. A patient experiencing repeated AICD (Automatic Implantable Cardioverter-Defibrillator) activations should receive sedation or pain management from the AEMT or Paramedic.

2006.2 Clinical Management

Assessment			
Pediatric Considerations None	Signs & Symptoms - AICD in place and firing - Sudden pain - Muscle spasms	Differential Diagnosis None	
Treatment Algorithm			
- Monitor and be prepared to provide BLS care. - Be prepared to defibrillate in the event of AICD failure.			EMR
- Monitor and transport as indicated. - Consider calling for ALS care.			EMT
- Be prepared to defibrillate in the event of AICD failure. Midazolam 2.5 mg slow IV for sedation. - Consider 1014 Pain Management Protocol. - For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses.			AEMT
Be prepared to manually cardiovert or defibrillate in the event of AICD failure.			Paramedic
Consult			
None			
Clinical Pearls			
None			
END OF SECTION			

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Subject:

Ventricular Assist Devices

Effective:

June 1, 2021

Last modified:

Dec. 7, 2024

2007.1 General Guidelines

- a. It is important to recognize the patient with a ventricular assist device (VAD).
- b. Routinely, your agency will be advised when a VAD patient is in your community.
- c. Otherwise, these patients could be travelling through, or visiting in your jurisdiction.
- d. The patient or family members are generally knowledgeable about the VAD and how to troubleshoot it.

2007.2 Assessing the VAD Patient

- a. Skin color and mental status are the best indicators of stability in the VAD patient.
- b. A pulse is usually not palpable in the VAD patient. Nearly all VADs are continuous flow devices.
- c. If the device is a pulsatile flow device, a pulse should be palpable.
- d. Blood pressure may or may not be obtainable and auscultated readings are usually unreliable.
 - i. In a continuous flow device, mean arterial blood pressure (MAP) can be obtained by auscultating with a {Doppler}.
 - ii. The first sound heard during auscultation reflects the MAP.
 - iii. The MAP displayed by an automated non-invasive measurement may also be used.
 - iv. A normal and expected MAP in a patient with a VAD is 65 – 90 mmHg.
 - v. If the device is a pulsatile flow device, a blood pressure should be measurable.
- e. Pulse oximetry readings seem to be accurate, despite the manufacturer stating otherwise.
- f. Quantitative waveform capnography should be accurate and can be reflective of cardiac output
- g. An EtCO₂ of less than 30 mmHg can be indicative of low perfusion secondary to poor pump function.
- h. {ECG 12/15 lead} as usual, no interference from the VAD is expected
- i. Temperature should be measured as infection and sepsis are common.

2007.3 Transporting the VAD Patient

- a. Patients with or without a VAD problem should be transported to the nearest appropriate Hospital ED.
- b. Do NOT delay ground transportation waiting to speak with the patient's VAD Coordinator.
- c. Always bring the patients resource bag with you. It should contain:
 - i. Spare batteries and a battery charging unit
 - ii. Spare control unit
 - iii. Contact information for the VAD Coordinator.
 - iv. Directions for equipment and alarm troubleshooting.
- d. Always bring spare batteries for the VAD with the patient, even if it is not a VAD related problem.
- e. If the transport is going to be prolonged or it is expected that the patient will be away for a while, try to bring the VAD base power unit with you.
 - i. Alternately, you can ask the patient's family/caregiver to bring it to the hospital.
- f. There may be a need to bring it with the patient and plug it into an inverter for power.

2007.4 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - VAD equipment - VAD vests or battery packs	Differential Diagnosis None



Subject: Ventricular Assist Devices

Effective: June 1, 2021

Last modified: Dec. 7, 2024

Treatment Algorithm

- Determine if you have a patient with a VAD problem, or a patient with a VAD that has a medical/trauma problem.
- If there is no indication of possible VAD malfunction or failure, exit to appropriate protocols.
- Assess the VAD:
 - Auscultate over the VAD pump location (Should be just to the left of the epigastrium, immediately below the heart)
 - If the pump is functioning, a low hum should be audible.
 - Do not assume that the pump is functioning just because the control unit does not indicate a problem.
 - Palpate the control unit.
 - A hot control unit indicates the pump may be working harder than it should be
 - This often indicates a pump problem such as a thrombosis.
 - Look at the alarms on the control panel
 - Trouble with the VAD will usually be identified by an alarm.
 - The patient will usually have a resource guide to direct alarm troubleshooting.
 - Ask if the device is a continuous or pulsatile flow device.
 - Ask if the patient can receive electrical therapy.
 - Ask if chest compressions can be performed in the event of pump failure.
- Inquire about DNR status.
- If there is indication of possible device malfunction or failure:
 - Attempt to restart VAD if previously off for less than 5 minutes.
 - If VAD off longer than 5 minutes, then:
 - Locate the patients "Emergency Contact Card"/VAD ID Card
 - Contact the VAD coordinator.
 - Discuss the plan with caregivers.
- If a VAD patient is unresponsive and pulseless with a non-functioning VAD and has previously indicated a desire for resuscitative efforts, begin chest compressions.
 - AVOID THE USE OF MECHANICAL CPR DEVICES
 - Defibrillation pads should be placed anterior/posterior
 - Ensure that all troubleshooting efforts (reconnecting wires, changing batteries, replacing the control unit) have failed prior to starting chest compressions.
- Follow BLS protocol.

EMR

- Transport urgently.
- No additional directives at this level.

EMT

- No additional directives at this level.

AEMT

- Only symptomatic dysrhythmias not at the patient's baseline should be treated.
- If indicated, place electrical therapy/defibrillation pads away from VAD site and AICD.
- VAD patients may receive ACLS interventions.

Paramedic

Consult

- None

Clinical Pearls

- Utilize the patient and family as a resource.
- Always contact the VAD Coordinator if there is a VAD related problem or question.
- Common complications in VAD patients include stroke (incidence up to 25%), bleeding, dysrhythmias, and infection.
- The most common causes of death in VAD patients are sepsis and stroke. Consider this with a VAD patient showing altered mental status.
- VAD patients are preload dependent. Consider that a fluid bolus can often reverse hypoperfusion.

END OF SECTION



Subject: Suspected Cardiac Chest Pain

Effective: June 1, 2021

Last modified: Feb. 13, 2023

2008.1 General Guidelines

- a. Unstable cardiac patients are hypotensive, or have chest pain with poor skin color or diaphoresis.

2008.2 Clinical Management

Assessment					
Pediatric Considerations - Chest pain in the pediatrics is rarely related to a cardiac event. - Assessment for other causes (e.g., muscle pain, respiratory difficulties, injury) should be completed to determine the source of pain. - Apply supplemental oxygen and transport. - THE REST OF CHEST PAIN ALGORITHM DOES NOT APPLY TO PEDS.	Signs & Symptoms - Chest pain - Shortness of breath - Syncope - Pallor, Diaphoresis - Radiation of pain - Weakness - Nausea - Vomiting	Differential Diagnosis - Pericarditis - Pulmonary embolism - Asthma/COPD - Pneumothorax - Aortic dissection or aneurysm - GE reflux or hiatal hernia - Chest trauma - Esophageal spasm			
Treatment Algorithm					
- Arrange for rapid ALS transport. - Apply O₂ as appropriate. - Oxygen saturations less than 94%, should be given oxygen via NC and titrated to 94%. - Oxygen saturations 94% or higher, should not get any oxygen. - Do not withhold oxygen from a patient with SOB or respiratory distress.			EMR	AEMT	Paramedic
◆ Give Aspirin (ASA) 324 mg (chewed) to every patient greater than 25 y/o with symptoms of Acute Coronary Syndrome (ACS). ◆ Administer Nitroglycerin 0.4 mg SL, every 5 minutes, for pain, to a total of three pills with vital signs between doses. - SBP must be greater than 100. - Patient must be greater than 25 y/o. - Prior to moving patient, acquire a supine {12 Lead EKG} on all patients with ACS symptoms. {Transmit 12 Lead EKG} with two identifiers to MCP. - The MCP shall be contacted after at least the initial {12 Lead EKG transmission} is completed. - Consult MCP for appropriate destination. - Consider and transmit repeat {12 Lead EKGs} during transport.			EMT		
◆ Must obtain MCP permission to administer Aspirin (ASA) to patients 25 y/o or younger - The AEMT must also transmit the {12 Lead EKGs} - Administer Nitroglycerin 0.4 mg SL, every 5 minutes, for pain, to a total of three pills with vital signs between doses. - Prior to Nitroglycerin administration, establish vascular access for patients who have not previously had Nitroglycerin. - Consider 1014 Pain Management Protocol, provided SBP greater than 100 after first dose of nitroglycerin. DO NOT WAIT UNTIL 3 NITROGLYCERIN TABLETS ARE GIVEN BEFORE CONSIDERING FENTANYL. IV fluid, up to 500 ml, may be administered to a patient with SBP less than 100 without pulmonary edema.					
- Treat cardiogenic shock with or without pulmonary edema as identified in 4016 Shock. - If evidence of STEMI, transport to an interventional cardiac catheterization lab. - The Paramedic should only transmit a {12/15 Lead EKG} that meets Cardiac Alert criteria, or that is questionable.					
Consult					
- Without consultation, the Suspected Cardiac Chest Pain protocol only applies to patients greater than 25 years old with ACS symptoms. - Contact MCP for further advice with pediatric chest pain as needed. - For the EMT, the following requires MCP orders: - Aspirin administration - Nitroglycerin administration - Accessing the GMVEMSC Drug Bag					
Clinical Pearls					
- No significant change in patient condition in the field should be expected from the administration of Aspirin. - Patient must chew Aspirin. - Aspirin is contraindicated in third trimester of pregnancy. - Do not administer Nitroglycerin (NTG) if the patient has taken Viagra, Cialis, Levitra, Revatio, or similar medications within the last 24 hours.					
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Subject: Cardiac Alert Program

Effective: June 1, 2021

Last modified: Jan. 9, 2023

2009.1 General Guidelines

- The intent of the Program is to decrease the "Door to Balloon" time for pre-hospital AMI Patients.
- Providers will make early notification to the receiving facility and speak directly with the Physician.
- The Physician may activate a Cardiac Alert, based on provider impression and {12 Lead EKG} interpretations.

2009.2 Inclusionary Criteria

- Patients presenting with anginal-type chest pain or an equivalent anginal event may be candidates.
- Evidence of an AMI (greater than 1mm ST elevation in 2 contiguous leads) on a diagnostic {12-lead EKG}.

2009.3 Exclusionary Criteria

- Patient with a Left Bundle Branch Block (QRS greater than 120 milliseconds).
- Patients with a pacemaker rhythm.

2009.4 Clinical Management

Assessment		
Pediatric Considerations Consider differential diagnosis	Signs & Symptoms - Chest pain - Difficulty breathing - Syncope - Anginal equivalents	Differential Diagnosis - None in the presence of ACS symptoms - Chest trauma - Pulmonary issues - Cardiac Alert imitators on 12 Lead EKG
Treatment Algorithm		
No additional orders at this level.		
- Acquire and transmit the {12 Lead EKG} in any suspected AMI or in cardiac arrest with ROSC. - Contact the receiving hospital for further orders and/or destination directives. - Acquire serial {12 Lead EKGs} enroute to the hospital. - The recommendation is to repeat {12 Lead EKGs} every 5 minutes or with any change in condition/presentation) - Consider applying defibrillation pads to confirmed myocardial infarction patients. - Transport patients with ST Elevation MI (STEMI) or ROSC after cardiac arrest to an Interventional facility.		
Consider aggressive fluid administration of up to 500 ml to manage cardiogenic shock.		
- Reassess lungs frequently. - In significant AMIs, patients may develop cardiac conduction disorders (PVCs, BBB and 2° or 3° blocks). - If patient develops significant bradycardia, then utilize 2010 Bradycardia - Monitor blood pressure and administer Nitroglycerin or Fentanyl cautiously.		
A If patient is still hypotensive after other therapy, begin Norepinephrine by adding 4 mg to 250 ml of IV fluids. Infuse starting at 30 drops per minute (max. 45 drops) with 60 drop tubing and titrate to effect. Increase by 5 drops/minute every 5 minutes.		
Consult		
- The EMT and AEMT should contact the MCP after {12 Lead EKG} transmissions for further orders. - The Paramedic is expected to read and interpret the {12-lead EKG}. - Do <u>not</u> rely solely on the computer interpretation or expect the physician to interpret the transmitted {12 Lead EKG} for you.		
Clinical Pearls		
- An interventional facility is a hospital that provides Percutaneous Cardiac Interventions 24 hours a day. - To determine the regional interventional facilities, see 7013 Hospital Capabilities Chart. - Rerouting at interventional facilities does not apply to Cardiac Alerts. - Consider air medical transport if the interventional facility is over 30 minutes away. - Exceptions to transporting to an interventional facility include: - It is medically necessary to transport the patient to the closest hospital for stabilization. - It is unsafe to transport the patient directly due to adverse weather/ground conditions or excessive transport time. - Transporting the patient to would cause a critical shortage of local EMS resources. - Patient requests transport to a different facility, despite EMS education of patient.		

END OF SECTION

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Subject: Bradycardia

Effective: June 1, 2021

Last modified: Feb. 11, 2024

2010.1 General Guidelines

- Bradycardia is any rate less than 60 bpm.
- Non-symptomatic bradycardia may be a normal finding in otherwise healthy individuals.
- Assess the patient and determine medical history.
- Treat unexplained or symptomatic bradycardia

2010.2 Clinical Management

Assessment			
Pediatric Considerations - With adequate perfusion, monitor vital signs, and apply oxygen if needed. - Hypoxia in pediatric patients will produce bradycardia.	Signs & Symptoms - Heart rate less than 60 bpm - Syncope - Unstable bradycardia - Hypotension - Altered mental status - Unresolved chest pain - Poor skin color - Diaphoresis	Differential Diagnosis - Acute myocardial infarction - Hypoxia - Hypothermia - Elevated ICP (Stroke or Trauma) - Spinal cord lesion - Sick sinus syndrome - Athletic patients	
Treatment Algorithm			
- Administer oxygen as indicated. - Call for transport immediately. - For adequate perfusion, observe and monitor vital signs.			EMR
- Obtain {12 Lead EKG}, transmit and call receiving facility. - Transport immediately unless ALS intercept is less than 5 minutes. P For Pediatric patients less than 2 years old: P Look for signs and symptoms of shock or hypoperfusion P Secure the airway and ventilate with BVM at 1 breath every 3-4 seconds P If heart rate and perfusion do not increase within 30 to 60 seconds, then perform CPR.			EMT
No additional orders at this level.			AEMT
- Obtain and interpret {12/15 Lead EKG} - Wide complex bradycardia patients should spark consideration of treatment of hyperkalemia. A ♦ Administer both Calcium Chloride 10% 1 g (Calcium Chloride or Gluconate) <u>and</u> Sodium Bicarbonate 100 mEq. - Flush well between these medications. It is critical that these drugs not be given together, as they will precipitate. - With evidence of poor perfusion in adults and pediatrics: A Consider Atropine 1 mg IV, up to total of 3 mg. - If treatments are ineffective begin pacing: A If time permits, Ketamine 25 mg IV (preferred method) <u>or</u> Midazolam 2.5 mg slow IV prior to pacing. A <u>DO NOT</u> reduce Ketamine or Midazolam doses by half for patients greater than 69 y/o A Set at 70 BPM, 20 mA and increase until mechanical capture is obtained. P Epinephrine (1:10,000) 0.01 mg/kg, IV, repeat every 5 minutes. - If AV block: P Consider Atropine 0.02 mg/kg IV (minimum dose 0.1 mg, maximum single dose 0.5 mg) P May repeat dose every 5 minutes. Max total dose of 1 mg. P Consider pacing: P Pediatric electrodes should be used on patients less than 15 kg. P Consider Midazolam 0.1 mg/kg (max dose 2 mg) slow IV prior to pacing. P Start with 5 mA increasing as needed to 200 mA at a rate of 80 bpm until capture.			Paramedic
Consult			
The paramedic will consult for administration of Calcium Chloride 10% (or Gluconate) and Sodium Bicarbonate.			
Clinical Pearls			
None			
END OF SECTION			

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Subject: Tachycardia

Effective: June 1, 2021

Last modified: May 16, 2026

2011.1 General Guidelines

- Tachycardia is any heart rate greater than 100 bpm.
- Assess the patient and determine medical history.
- Treat unexplained or symptomatic tachycardia

2011.2 Clinical Management

Assessment				
Pediatric Considerations With adequate perfusion, monitor vital signs, and apply oxygen if needed.	Signs & Symptoms - Heart rate greater than 100 bpm - Dizziness - Chest pain - Shortness of breath - Unstable tachycardia - Hypotension - Altered mental status thought to be due to tachycardic rhythms	Differential Diagnosis - Myocardial infarction - Electrolyte imbalance - Exertion/pain/emotional stress - Fever - Hypoxia - Hypovolemia or anemia - Drug overdose - Hyperthyroidism - Pulmonary embolus		
Treatment Algorithm				
- Administer oxygen as indicated. - Call for transport immediately.			EMR	
- Obtain {12 Lead ECG}, transmit and call receiving facility. - Transport immediately unless ALS intercept is less than 5 minutes.			EMT	
No additional orders at this level.			AEMT	
- Obtain and interpret {12/15 Lead ECG} <u>Stable:</u> - Narrow Complex - Regular - A Vagal maneuvers - A Adenosine 6 mg rapid IVP, saline flush - A May repeat Adenosine 12 mg rapid IVP x 2, saline flush - Atrial Fibrillation with a Rapid Ventricular Rate (RVR) - A Transport for further evaluation - Wide Complex – Regular or Irregular - A Amiodarone 150 mg in 250 ml NS, IV over 10 minutes using 60 drop/ml tubing. - A If Amiodarone not available, use Lidocaine - A Lidocaine 150 mg IV/IO <u>Unstable:</u> - A Consider administration of a sedative/analgesic prior to cardioversion - A Ketamine 25 mg IV (preferred method) <u>or</u> Midazolam 2.5 mg slow IV - A <u>DO NOT</u> reduce Ketamine or Midazolam doses by half for patients greater than 69 y/o - A Cardioversion: 100, 200, 300, 360 J for monophasic or biphasic equivalent - A ♦ If Atrial Fibrillation with a Rapid Ventricular Rate (RVR): Cardioversion at 200 J for monophasic or biphasic equivalent <u>With evidence of polymorphic ventricular tachycardia (torsades de pointes):</u> - A If stable: Magnesium Sulfate 2 gm infused with macro-drip tubing over 10 minutes - A If unstable: Unsynchronized Cardioversion: 100, 200, 300, 360 J for monophasic or biphasic equivalent				



Subject: Tachycardia

Effective: June 1, 2021

Last modified: May 16, 2026

- Stable Pediatrics:
 - P Vagal maneuvers (blowing through a straw or oxygen tubing, etc.)
- Unstable Pediatrics:
 - P **Adenosine 0.1 mg/kg rapid IVP** (max dose 6 mg), saline flush.
 - P If no response, **Adenosine 0.2 mg/kg rapid IVP** (max dose 12 mg), saline flush. Repeat x 1.
 - o Consider cardioversion.
 - P If time permits, **Midazolam 0.1 mg/kg slow IV** (max dose 2.5 mg).
 - P **Cardioversion 1 J/kg**
 - P If no response, repeat cardioversion at 2 J/kg

Paramedic

Consult

- Paramedics must consult with MCP for orders to cardiovert unstable patients presenting Atrial Fibrillation with a Rapid Ventricular Rate (RVR)

Clinical Pearls

- Rule out or treat compensatory and physiological causes prior to treating tachycardia in isolation
- Paramedics should **not** cardiovert:
 - o Patients without hemodynamic changes
 - o Patients whose hemodynamic changes have other apparent causes.
- Hemodynamically unstable patients will typically present with:
 - o Hypotension
 - o Altered mental status
 - o Diaphoresis or cool/clammy skin
- If patient has history of Paroxysmal Supraventricular Tachycardia (PSVT) and advises it takes 12 mg of Adenosine, then skip the 6 mg dose.

END OF SECTION



3000 Series

Trauma Protocol

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Subject: General Trauma Management

Effective: June 1, 2021

Last modified: Feb. 11, 2024

3001.1 General Guidelines

- Minor trauma patients may be transported to non-trauma centers.
- Major trauma patients are to be transported as soon as possible to the nearest appropriate facility.
- Scene size-up, with rapid assessment and recognition of major trauma/multiple system trauma and effective evaluation of the mechanism of injury are essential to the subsequent treatment.
- If patient meets criteria as defined in [3018 Trauma Transport Guidelines](#), then call "Trauma Alert".
- If transporting by helicopter, ensure a copy of the patient care report gets to the receiving facility.

3001.2 Clinical Management

Assessment			
Pediatric Considerations • May not exhibit typically • Injuries may not present the same as in adults • Will present decompensated shock late	Signs & Symptoms • Traumatic injuries ○ DCAP-BTLS	Differential Diagnosis • Medical complaints with S/S that mimic traumatic injuries	
Treatment Algorithm			
• The <i>only</i> procedures that should take precedence to transport of major trauma patients are: ○ Airway management ○ Stabilization of neck/back or obvious femur and pelvic fractures on a backboard ○ Exsanguinating hemorrhage control ○ Extrication • Maintain patient’s body temperature. • Take a manual BP on all trauma patients. • Repeat vitals on trauma patients every 5 minutes.		EMR	
		• On-scene time should be limited to 10 minutes or less, except when there are extenuating circumstances • Report Mechanism of Injury, Injuries, Vital signs, Treatment (MIVT), GCS with components, and ETA to the receiving facility	
• IVs should be established en route to the hospital unless the patient is trapped, transport is otherwise delayed, or patient has no life-threatening injuries, and transport prior to analgesia would be extremely painful. - A Start the IV with a large bore catheter and macro drip tubing. - A Administer up to a 1000 ml IV fluid bolus - P Administer 20 ml/kg of IV fluid - A IV flow rates are as follows: • Keep open rate for major head trauma with adequate perfusion • IV wide open if the patient has inadequate perfusion (including head trauma) utilizing {IV Pressure Infusion Pump or Bag} or similar equipment if available • Titrate all IV flow rates to maintain SBP ~ 100 • For penetrating trauma to the chest and abdomen: ○ If a radial pulse is present and the patient is conscious and mentating, load and go. ○ If no radial pulse, infuse IV fluid in 250 ml boluses until radial pulse is present and then stop fluid. • Consider 1014 Pain Management Protocol.		AEMT	
• No additional orders at this level			Paramedic
Consult			
• Use of on-line MCP for medical direction in the field for difficult cases is encouraged. • Pre-arrival notification of the receiving facility is essential! • Keep the receiving hospital informed on the patient’s condition, significant changes should be reported.			
Clinical Pearls			
• Hypothermia is a significant and frequent problem in shock for major trauma patients. • Surgical emergencies with increased fluid administration cause dilution, lower body temperatures and increase coagulopathies, all of which increase mortality. ○ To address this, allow for “permissive hypotension,” • This means that IV fluids are not administered to these patients unless there is loss of radial pulse.			
END OF SECTION			

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Subject: Major Trauma

Effective: June 1, 2021

Last modified: May 16, 2026

3002.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms Significant injuries or life threats	Differential Diagnosis None
Treatment Algorithm		
- Patients meeting criteria for transport to a Trauma Center are considered "Load and Go." - Consider the trauma care mnemonics M.A.R.C.H. to establish treatment priorities: Massive Hemorrhage: Assess for and address significant bleeding Airway: Assess for and ensure a patent airway using adjuncts and positioning Respirations: Assess respiratory rate, quality and effort and oxygenate the patient with 100% O₂ Circulation: Double check all bleeding interventions and perform a head-to-toe sweep for additional bleeding Hypothermia: Prophylactically treat patient for hypothermia - Other trauma care considerations: - Open pneumothorax: cover wound with an occlusive dressing, tape down three sides. - Tension pneumothorax: lift one side of any occlusive dressing. - Flail chest: stabilize immediately with a gloved hand, then immobilize with a bulky dressing or towels taped to the chest		EMR
No additional orders at this level.		EMT
- Tension pneumothorax: - Use caution not to confuse right main stem intubation for a pneumothorax. - Perform needle decompression as indicated - Decompress the chest with a 14-gauge or larger, 3 ¼" angiocath - Location options include: - Fourth or fifth intercostal space in the mid-axillary line - Second or third intercostal space in the mid-clavicular line (use nipple line as a guide) P In patients less than 8 years old, decompression site choice will only be the second or third intercostal space at the mid-clavicular line		AEMT
{With defined and specific indications, paramedics who have received appropriate training and Medical Director's approval may perform a finger thoracostomy in accordance to their specific protocols}		Paramedic
Consult		
Contact Medical Control and advise them of patient condition with MIVT, ETA, and GCS components.		
Clinical Pearls		
For pregnant patient in arrest consider need for manual uterine displacement.		
END OF SECTION		

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Subject: Glasgow Coma Score

Effective: June 1, 2021

Last modified: Dec. 28, 2024

3003.1 General Guideline

- a. When assessing the level of consciousness, use the appropriate Glasgow Coma Score.
- b. All patients should have at least one recorded and reported GCS.

3003.2 The Glasgow Coma Score

	LESS THAN 2 YEARS OLD		ADULT & PEDIATRIC OVER 2 YEARS OLD	
EYES	SPONTANEOUSLY	4	SPONTANEOUSLY	4
	TO VOICE	3	TO VOICE	3
	TO PAIN	2	TO PAIN	2
	NO RESPONSE	1	NO RESPONSE	1
VERBAL	COOS, BABBLER	5	ORIENTED	5
	IRRITABLE CRY, CONSOLABLE	4	CONFUSED	4
	CRIES TO PAIN	3	INAPPROPRIATE WORDS	3
	MOANS TO PAIN	2	GRUNTS, GARBLED SPEECH	2
	NO RESPONSE	1	NO RESPONSE	1
MOTOR	NORMAL MOVEMENTS	6	OBEYS COMMANDS	6
	WITHDRAWS TO TOUCH	5	LOCALIZES PAIN	5
	WITHDRAWS TO PAIN	4	WITHDRAWS TO PAIN	4
	FLEXION (DECORTICATE)	3	FLEXION (DECORTICATE)	3
	EXTENSION (DECEREBRATE)	2	EXTENSION (DECEREBRATE)	2
	NO RESPONSE	1	NO RESPONSE	1

END OF SECTION

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Subject: Trauma Arrest

Effective: June 1, 2021

Last modified: Dec. 7, 2024

3004.1 General Guidelines

- a. Traumatic cardiac arrest care will follow the same algorithm as other cardiac arrest scenarios.
- b. Mechanical CPR devices are contraindicated in traumatic arrest where there is:
 - i. Injury or mechanism of injury to the neck
 - ii. Injury or mechanism of injury to the thoracic cavity (anterior or posterior)
 - iii. Injury or mechanism of injury to the abdominal cavity
 - iv. Minor injuries to these areas or the extremities do not apply.
- c. If appropriate, providers may consider termination of resuscitation.

3004.2 Termination of Resuscitation

- a. Emergency medical responders (EMRs) may **not** terminate a trauma cardiac arrest.
- b. The criteria for termination of resuscitation in arrest from blunt or penetrating trauma is:
 - i. No immediately reversible cause can be determined after rapid primary survey and treatment.
 - ii. No signs of life after BLS (e.g. respiratory effort, purposeful movement, reactive pupils, etc.)
 - iii. Sustained EtCO₂ of below 10 mmHg
 - iv. If no ALS equipment is available at the scene and transport will exceed 20 minutes.
- c. Continue care and transport if patient arrests **after** in the care of EMS.

3004.3 Clinical Management

Assessment		
Pediatric Considerations If the pediatric patient does <u>not</u> meet non-initiation criteria, then <u>begin</u> resuscitation.	Signs & Symptoms - Cardiac arrest with traumatic injury or significant mechanism of injury - Unresponsive, pulseless, and apneic - Excessive hemorrhage	Differential Diagnosis - Signs of irreversible death - Other causes of unresponsiveness - Meets 1003 Non-initiation of Care Protocol
Treatment Algorithm		
- Initiate basic life support as defined in 2002 Cardiac Arrest – BLS - Internal/External hemorrhage control (e.g., tourniquets, pelvic binders, etc.)		EMR
- Consider the possibility of both medical and traumatic causes (mixed mechanisms). - Initiate a Rapid Primary Survey for reversible causes. TREATMENT OF REVERSIBLE CAUSES SHOULD BE A PRIORITY. - Cardiac monitoring/defibrillations via AED. ♦ Consider Termination of Resuscitation. (AEMT and Paramedic will continue through the algorithm before termination).		EMT
- Continue treating any organized rhythm with rate greater than 40 because of the potential of pseudo-PEA. - Secure airway and confirm with continuous EtCO₂. - Bilateral needle decompression as indicated (ex. high airway resistance, chest trauma, subcutaneous air). - Fourth or fifth intercostal space in the mid-axillary line - Second or third intercostal space in the mid-clavicular line (use nipple line as a guide) P In patients less than 8 years old, decompression site choice will be limited to the 2nd or 3rd intercostal space at the mid-clavicular line - Repeat needle decompression as indicated (continued high airway pressure). - Administer rapid IV fluid administration: A Administer up to 1000 ml IV fluid P Administer 20 ml/kg of IV fluid - If ROSC is achieved, transport immediately.		AEMT
No additional orders at this level		Paramedic

Subject: **Trauma Arrest**Effective: **June 1, 2021**Last modified: **Dec. 7, 2024****Consult**

- Contact MCP for Field Termination
- Be ready to provide the following information:
 - Duration of resuscitation
 - How long the patient was in arrest prior to EMS arrival
 - Witnessed or unwitnessed cardiac arrest
 - Capnography values
- Presenting rhythm (for AEMT and Paramedic)

Clinical Pearls

- For pregnant patient in arrest consider need for manual uterine displacement.

END OF SECTION



Subject: Burns and Smoke Inhalation

Effective: June 1, 2021

Last modified: May 16, 2029

3005.1 General Guidelines

- a. It is strongly recommended that at dispatch, agencies immediately call for the nearest available cyanide antidote cache whenever any of the following occur:
 - i. Dispatched on a report of a person trapped with exposure to fire or smoke in an enclosed area.
 - ii. Dispatched on a report of an incident involving cyanide.
 - iii. Report of a Mayday or firefighter down with exposure to fire or smoke in an enclosed area.
- b. Estimate and report total Body Surface Area (BSA) involved using universally accepted methods.
 - i. BSA estimates should include only full and partial thickness burns.
- c. Inhalation injuries with an unsecured airway should be transported to the nearest facility.
- d. Chemical burns are hazardous material situations and must be grossly decontaminated at the scene.

3005.2 Specific Care for Burns

- a. Radiation burns:
 - i. If there is radioactive material on the patient, then they must be decontaminated.
 1. Consider contacting a Hazardous Materials Team for assistance with decontamination.
 2. Contact the hospital prior to arrival like with any other hazardous materials case.
 - ii. Treat critical medical conditions first.
- b. Treat injuries like thermal burns once the area is decontaminated

3005.3 Clinical Management

Assessment		
Pediatric Considerations - Pediatric burn patients should be transported to Pediatric Burns Centers whenever practical. - Critical pediatric patients can go to the closest facility.	Signs & Symptoms - Burns, pain, swelling - Hypotension/shock - Airway compromise/distress - Singed facial or nasal hair - Hoarseness/wheezing	Differential Diagnosis - Superficial burns - Partial thickness burns - Full thickness burns - Chemical, Thermal, Electrical, Radiation burns
Treatment Algorithm		
- Stop the burning and minimize contamination. - Assess for respiratory distress, stridor, hoarseness, sooty sputum, singed eyebrows, and nares, or burns of the face or airway. - If available, use {CO oximeter}. - For inhalation burns: Administer high flow oxygen via non-rebreather mask. - Keep patient warm. - Superficial, partial thickness and full thickness burns should be covered with clean, dry sheets or dressings. - Do <u>not</u> apply ice or ice packs to burns, if ice was applied prior to arrival, then remove. - Remove clothing and jewelry from injured parts. Do not remove items which have adhered to the skin.		EMR
- If available deliver {humidified} oxygen. ♦ For inhalation burns: If no humidifier is available, administer Saline 3 ml via nebulizer. Repeat as needed.		EMT
- Apply cardiac monitor, especially if patient has suffered a lightning strike or electrical burn. - Provide endotracheal intubation if apneic. - Administer fluids to maintain perfusion, do not overhydrate. Fluids should be a balanced electrolyte solution when available. - IV access can be acquired in areas with burnt tissue if necessary and before intraosseous needle access. - Consider 1014 Pain Management Protocol.		AEMT

Subject: **Burns and Smoke Inhalation**Effective: **June 1, 2021**Last modified: **May 16, 2026**

- Early intubation as indicated. Do not wait for complete airway obstruction or respiratory arrest.
- For known or suspected cyanide poisoning, use [3014 Cyanide Poisoning and Antidotes](#)

Paramedic

Consult

- EMTs must call for orders to administer nebulized saline in the absence of humidified oxygen when treating inhalation burns.

Clinical Pearls

- Patients with severe burns should be transported to a Burn Center unless ETA greater than 30 minutes.
- BP may be taken over damaged tissue if no other site is accessible.

END OF SECTION



Subject: Carbon Monoxide Poisoning

Effective: June 1, 2021

Last modified: Jan. 19, 2025

3006.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Malaise, fatigue, drowsiness - Flu like symptoms - Headache - Dyspnea - Nausea/vomiting - Diarrhea - Abdominal pain - Syncope - Seizures	Differential Diagnosis - Flu/Severe cold - Chronic fatigue - Myocardial infarction - Diabetic crisis - Altitude sickness - Ingested toxins - Hypothyroidism
Treatment Algorithm		
- Remove patients from the environment. - Provide high flow O₂ to all suspected carbon monoxide poisonings. - Pulse oximeter will give false readings and should not be utilized. - If not already on scene, consider requesting an apparatus with equipment to test for CO in the atmosphere {CO oximeter}		EMR
Contact MCP to discuss transport considerations.		EMT
No additional orders at this level.		AEMT
No additional orders at this level.		Paramedic
Consult		
Look to Medical Control for guidance on transport destination.		
Clinical Pearls		
- When determining destination, consider possible hyperbaric oxygen treatment for the following patients with suspected CO exposure: - Underlying cardiovascular disease or symptoms such as chest pain or shortness of breath - Greater than 60 years of age - Obvious neurological symptoms, such as any interval of unconsciousness, loss of time, inability to perform simple motor tasks, or loss of memory - Smoke inhalation victims - Pregnancy		
END OF SECTION		

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Subject: Crush Syndrome Trauma

Effective: June 1, 2021

Last modified: Dec. 23, 2023

3007.1 Clinical Management

Assessment			
Pediatric Considerations No pediatric medication doses should exceed total adult doses.	Signs & Symptoms - Patient entrapped - Patient under a heavy load and crushed - Hypotension - Hypothermia - Abnormal ECG findings - Pain - Anxiety	Differential Diagnosis None	
Treatment Algorithm			
◆ Contact MCP immediately and prior to relieving the load. - Prepare for the patient to decompensate when extricated. - Monitor and reassess			EMR
{12 Lead ECG} as soon as feasible.			EMT
A 1 liter IV fluid bolus IV. Then 500 ml/hour IV P IV fluid, 20 ml/kg IV - Follow 1014 Pain Management protocol - If hypotensive and the patient has been entrapped for more than 1 hour: A Give additional IV fluid, 1 liter IV. P Give additional IV fluid, 20 ml/kg IV. ◆ Consider sedation: A Ketamine 250 mg IM, may repeat after 10 minutes G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses P Ketamine 5 mg/kg IM, max dose of 250 mg - Monitor for fluid overload			AEMT
<u>Normal ECG and hemodynamically stable, immediately prior to extrication:</u> A Sodium Bicarbonate 100 mEq IV P Sodium Bicarbonate 1mEq/kg IV <u>OR</u> <u>Abnormal ECG and hemodynamically unstable:</u> - If after release, hyperkalemia causes wide bizarre EKG complexes with: - Peaked T waves with a QRS greater than or equal to 0.12 seconds - QT ≥ 0.46 seconds - Loss of P wave - Bundle Branch Blocks - Premature ventricular contractions - Bradycardia ◆ Consider Calcium Chloride, 1 gm, flush line well before Sodium Bicarbonate Albuterol 10 mg nebulized A Sodium Bicarbonate 100 mEq IV P Sodium Bicarbonate 1mEq/kg IV			
Consult			
- Contact MCP immediately and prior to relieving the load. - MCP orders needed for sedation. - The paramedic must call MCP for orders to give Calcium Chloride to the unstable patient.			
Clinical Pearls			
- Consider the potential for multiple system trauma - Consider the potential for hypo or hyperthermia			
END OF SECTION			

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Subject: Cyanide Poisoning & Antidotes

Effective: June 1, 2021

Last modified: Mar. 9, 2025

3008.1 General Guidelines

- a. Cyanide antidotes are located in multiple caches in each of the counties throughout the region, and are available by contacting 937-333-USAR (8727).
- b. The cache agency closest to your incident will be dispatched, which will respond with both a Cyanokit and 3 doses of Sodium Thiosulfate, to provide for the potential of multiple patients.

3008.2 Indications to Call for the Cache

- a. It is strongly recommended that agencies immediately call for the nearest available cyanide antidote cache at the time of dispatch whenever any of the following occur:
 - i. Report of a person trapped with exposure to fire or smoke in an enclosed area.
 - ii. Report of an incident involving cyanide.
 - iii. Report of a Mayday or firefighter down with exposure to fire or smoke in an enclosed area.

3008.3 General Treatment

- a. Treatment of cyanide poisoning must include immediate attention to airway patency, adequacy of oxygenation and hydration, cardiovascular support, and management of any seizure activity.

3008.4 Clinical Management

Assessment				
Pediatric Considerations - For pediatric administration of Hydroxocobalamin (Cyanokit): - Mix 200 ml NaCl in 5 g vial (concentration is 25 mg/ml) 70 mg x patient weight in kg = total dose administered over 15 minutes. - Divide doses in half for repeat administration - See dosing chart at end of this tab for calculating pediatric doses		Signs & Symptoms - Known or strongly suspected cyanide exposure - Altered mental status - Seizures - Shock - Difficulty breathing		Differential Diagnosis None
Treatment Algorithm				
- Provide 100% O₂ via non-rebreather mask. - If unconscious, provide 100% O₂ by BVM				EMR
Consider CPAP for suspected smoke inhalation.				EMT
- Intubate if patient is apneic - Establish one IV in each arm if possible. - It is critical to control any seizure activity, as defined in 4014 Seizures				AEMT
- If available consider {BiPAP} for suspected smoke inhalation. ♦ Hydroxocobalamin (Cyanokit): ♦ Administer 5 grams via slow IV infusion over 15 minutes at a rate of 15 ml/min., using supplied 20 ml/min infusion set ♦ May repeat 5 grams via slow IV infusion over 15 min to 2 hours, depending on clinical response. ♦ Administer 70 mg/kg slow IV over 15 minutes; max dose of 5000 mg (5 grams), using supplied 20 ml/min infusion set ♦ May repeat 35 mg/kg slow IV; max dose 2500 mg (2.5 grams), depending on clinical response. or ♦ Sodium Thiosulfate: ♦ If greater than 25 kg: Administer 12.5 grams (50 ml) 25% solution slow IV. ♦ If less than 25 kg: Administer 412.5 mg/kg (1.65 ml/kg) 25% solution, slow IV (max dose 12.5 g (50 ml)).				
Consult				
- Orders for cyanide antidotes are not needed in cardiac arrest. ♦ Contact MCP to administer both Hydroxocobalamin (Cyanokit) and Sodium Thiosulfate to the same patient.				



Subject: Cyanide Poisoning & Antidotes

Effective: June 1, 2021

Last modified: Mar. 9, 2025

Clinical Pearls

- If a patient is in arrest, administer Hydroxocobalamin as quickly as possible.
- Only CAB, defibrillation, intubation, and epinephrine should precede use of the cyanide antidotes.
- Hydroxocobalamin is incompatible with numerous drugs including Diazepam.
- Whenever possible establish two IV lines in a different vein or limb, one for standard protocol drugs and one for cyanide antidotes.
- While IV infusion is the preferred method of cyanide antidote administration, in extreme cases the medications could be given via IO.
- If administering cyanide antidotes via IO, a traditional drip set may not be effective and measures may need to be taken to slowly push the medication in.

3008.5 Pediatric Hydroxocobalamin Dosing Chart

Weight (kg)	5	10	15	20	25	30	35	40	50	60	>70
Dose (mg)	350	700	1050	1400	1750	2100	2450	2800	3500	4200	5000
Amount needed for 70mg/kg	14 ml	28 ml	42 ml	56 ml	70 ml	84 ml	98 ml	112 ml	140 ml	168 ml	200 ml

3008.6 Pediatric Sodium Thiosulfate Dosing Chart

Weight (kg)	5	10	15	20	25	30
Dose (Gm)	2.06 g	4.125 g	6.18 g	8.25 g	10.3 g	12.5 g
Amount needed for 70mg/kg	8.25 ml	16.5 ml	24.75 ml	33 ml	41.25 ml	50 ml

END OF SECTION



Subject: Drowning

Effective: June 1, 2021

Last modified: Dec. 11, 2024

3009.1 Clinical Management

Assessment						
Pediatric Considerations None	Signs & Symptoms - History of submersion - Period of unconsciousness - Decreased or absent vital signs - Vomiting - Coughing	Differential Diagnosis - Trauma - Pre-existing medical problem - Barotrauma (diving) - Decompression sickness				
Treatment Algorithm						
- Consider Spinal Motion Restriction - Consider possibility of hypothermia. If present follow 3016 Hypothermia - Evaluate neurological status.			EMR	EMT	AEMT	Paramedic
- Drowning patients should be transported to a Trauma Center. - If patients are in cardiac arrest or respiratory failure/arrest, consider transport to the closest appropriate hospital						
Establish vascular access						
No additional orders at this level						
Consult						
None						
Clinical Pearls						
All submersion victims should be transported due to potential for worsening over the subsequent few hours.						
END OF SECTION						

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Subject: Extremity Injuries

Effective: June 1, 2021

Last modified: Dec. 8, 2021

3010.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Deformities - Inflammation - Pain upon movement - Immobility - Paresthesia	Differential Diagnosis None
Treatment Algorithm		
- For open fractures, control bleeding with direct pressure and cover with dry, sterile dressing. - If practical, consider elevating the limb. - Apply appropriate splinting device. - If the extremity is severely angulated and pulses are absent, apply gentle traction in an attempt to bring the limb back into a natural anatomic position. If resistance is encountered, splint the extremity in the angulated position. - Apply cold pack to reduce swelling.		EMR
No additional orders at this level		EMT
Consider 1014 Pain Management Protocol		AEMT
No additional orders at this level		Paramedic
Consult		
None		
Clinical Pearls		
- Document distal motor, sensation and circulation before and after splinting, or spinal motion restriction. - Open wounds should be covered with a sterile dressing before splinting. - Immobilize above and below the injury. - The patient who requires a load and go approach can be adequately immobilized by careful packaging on the long spine board. Do additional splinting enroute to the hospital as time and the patient's condition permit.		
END OF SECTION		

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Subject: Eye Injuries

Effective: June 1, 2021

Last modified: May 16, 2026

3011.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Irritation to eye - Visual disturbances or loss of vision - Obvious penetrating injury - Burns - Nausea	Differential Diagnosis - Hypertension - Contact lens issue
Treatment Algorithm		
- If possible, contact lenses should be removed. Contacts should be transported with patient. - Use nasal cannula with IV tubing for irrigation. - Chemical Burns: - Irrigate immediately with IV fluid or water for a minimum of 30 minutes or until patient transport is completed. - Determine chemical involved. Bring Safety Data Sheets, if available. - Major Eye Trauma: - Do not irrigate if there is penetrating trauma to the eye. - Cover affected eye(s) with a rigid cover. - Do not use a pressure or absorbent dressing on or near any eye that may have ruptured or have any penetrating trauma. - The patient should be transported with head elevated at least 30°.		EMR
No additional orders at this level.		EMT
No additional orders at this level.		AEMT
- Prior to irrigation with IV fluid or for significant eye pain, Tetracaine 2 drops in the affected eye. - Do not irrigate or use Tetracaine if penetrating trauma to the eye is present. - Use {Morgan Lens} or nasal cannula with IV tubing for irrigation.		Paramedic
Consult		
None		
Clinical Pearls		
None		
END OF SECTION		

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Subject: Frostbite

Effective: June 1, 2021

Last modified: Dec. 8, 2020

3012.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Cold, clammy skin - Shivering - Mental status changes - Extremity pain or sensory abnormality - Bradycardia - Hypotension or shock	Differential Diagnosis - Head Injury - Spinal cord injury
Treatment Algorithm		
- Protect injured areas. - Remove clothing and jewelry from injured parts. - Do not attempt to thaw injured part with local heat. - Maintain core temperature.		EMR
Severe frostbite injuries should be transported to a Burn Center.		EMT
- Establish vascular access and consider {warmed} fluids. - Consider 1014 Pain Management Protocol.		AEMT
No additional orders at this level		Paramedic
Consult		
None		
Clinical Pearls		
None		
END OF SECTION		

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Subject: Head Injury

Effective: June 1, 2021

Last modified: Dec. 7, 2024

3013.1 Clinical Management

Assessment		
Pediatric Considerations Assess the fontanelles in younger patients	Signs & Symptoms - Visible head trauma - Altered LOC - Cushing's Triad or similar V/S - Ataxic Respirations - Increased B/P - Bradycardia - Pupillary changes - Posturing	Differential Diagnosis - Alcohol/Acidosis - Epilepsy/Endocrine - Infection - Overdose/Oxygen Deficiency - Uremia - Tumor - Insulin - Psychogenic/Poison - Stroke/Shock
Treatment Algorithm		
- Evaluate level of consciousness, pupillary size and reaction. - Establish Glasgow Coma Score and reassess frequently. - Ventilate at 20 breaths per minute when signs of cerebral herniation are present: {Ventilate to maintain EtCO₂ readings of 30 mmHg (30 torr)}. - Never ventilate at less than 8 per minute. P Ventilate at a rate of ten faster than normal respiratory rate when the signs of cerebral herniation are present.		EMR
No additional orders at this level		EMT
No additional orders at this level		AEMT
No additional orders at this level		Paramedic
Consult		
None		
Clinical Pearls		
- Signs of cerebral herniation can include: - Decreased mental status - Dilated and/or unresponsive pupils - Bradycardia - Hypertension - Posturing - Hypoventilation increases the level of CO₂ in the brain, causing cerebral vasodilatation and increased swelling. - Hyperventilation decreases the level of CO₂ and causes cerebral vasoconstriction, hypoxia, and ischemia. - Both hyperventilation and hypoventilation could cause cerebral hypoxia and increased mortality. {Make every effort to maintain an EtCO₂ of 30 mmHg}		
END OF SECTION		

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Subject: Heat Exposure

Effective: June 1, 2021

Last modified: Oct. 10, 2021

3014.1 Clinical Management

Assessment		
Pediatric Considerations - May not exhibit typically - Do not thermoregulate well	Signs & Symptoms - History of heat exposure - Cramping - Hot or flushed skin - Excessive sweating - Nausea/vomiting - Mental status changes	Differential Diagnosis - Thyroid storm - Excited delirium - Malignant hyperthermia - Alcohol - Epilepsy - Insulin - Trauma - Infection - Psychosis - Stroke
Treatment Algorithm		
- Move patient to a cool environment - Remove patient's clothing - Continuously apply water to the skin to cool the patient, use fan for evaporation if available - Apply cold packs to underarms and groin area - Cold water submersion is an acceptable method for cooling heat stroke patients. You may encounter patients in cooling body bags. The goal is to lower temperature to less than 102.5°F - If conscious and not vomiting or extremely nauseous, provide oral fluids - Be prepared for seizures - Consider other medical conditions (e.g., overdose, hypoglycemia, CVA) and treat accordingly		EMR
Hyperthermia patients should be transported to a Trauma Center		EMT
- If hypotensive or mental status changes: A IV fluid 500 ml IV P IV fluid 20 ml/kg IV (max 500) - May repeat both adult and pediatric fluid bolus one time ♦ Additional IV fluid, if indicated - Consider other medical conditions (e.g., overdose, hypoglycemia, CVA) and treat accordingly		AEMT
No additional orders at this level		Paramedic
Consult		
For additional (more than 2) fluid challenges in adults		
Clinical Pearls		
- Geriatric patients, pediatric patients, patients with a history of spinal injury, and diabetics are most likely to suffer heat-related illnesses - Other contributory factors may include heart medications, diuretics, cold medications, and psychiatric medications - Heat exposure can occur due to increased environmental temperatures, prolonged exercise or a combination of both - Environments with temperatures above 90°F and humidity over 60% present the most risk		
END OF SECTION		

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Subject: Hemorrhage Control

Effective: June 1, 2021

Last modified: June 6, 2026

3015.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Significant bleeding - Shock-like symptoms	Differential Diagnosis None
Treatment Algorithm		
- Control of life-threatening external hemorrhage takes priority over any other treatment. - Constant, direct pressure is the primary method of bleeding control. - If direct pressure fails to control bleeding from extremities, use a tourniquet. {Commercial tourniquets such as the CAT or SOFTT are recommended} - Only use wide, flat materials such as cravats or BP cuffs as improvised tourniquets - Place a tourniquet as proximal as possible to the torso on the femur or humerus - Tighten the tourniquet until the bleeding stops - If bleeding persists, place another tourniquet abutted to the first tourniquet - Document time and location - Be sure that the ER staff is aware of the tourniquet {For life-threatening hemorrhage that can't be controlled by tourniquets, consider hemostatic dressings}. - Combat Gauze, or ChitoFlex PRO are examples - These can be used on the chest or abdomen - Place in direct contact with the source of bleeding and apply a pressure dressing or use Kerlix - DO NOT USE GRANULAR AGENTS - Wound Packing may be performed by providers at any level, as long as they have received proper training - This procedure is <u>not</u> to be used on open wounds to the head, chest or abdomen - Use sterile gauze or approved hemostatic products - Gauze should be placed as deeply in the wound as possible using a gloved digit and continuous pressure - Excessive force is not necessary and may be harmful. - Apply a pressure dressing and manual direct pressure over the packed wound for at least 3 minutes - Do not remove wound packing once placed in the cavity - Notify the ED staff of the use of wound packing on arrival at the destination - Treat for hypovolemic shock as indicated.		EMR
No additional orders at this level		EMT
- For known or suspected hemorrhage secondary to trauma: A Tranexamic Acid (TXA) 2 gram IV/IO over 1-2 minutes P In patients 8 y/o or greater, Tranexamic Acid (TXA) 15/mg/kg (max 1 gram) IV/IO over 1-2 minutes A If unsure if TXA is indicated, contact Medical Control for advice. - For adults with significant bleeding affecting the airway, including post-tonsillectomy: A Tranexamic Acid (TXA) 500 mg nebulized with an additional 3 ml normal saline, O₂ flowing at 8-10 LPM A ♦ If significant bleeding continues after nebulized Tranexamic Acid (TXA), consider administering the remainder IV/IO - In pediatric patients with uncontrolled bleeding from a recent tonsillectomy, consider Tranexamic Acid (TXA) nebulized: P Less than 25 kg: 250 mg, nebulized with an additional 3 ml normal saline, O₂ flowing at 8-10 LPM P 25 kg or greater: 500 mg, nebulized with an additional 3 ml normal saline, O₂ flowing at 8-10 LPM		AEMT
{Agencies with a Prehospital Whole Blood Program who have received appropriate training and Medical Director's approval may administer blood in accordance to their specific protocols} {The delivery of whole blood should not significantly delay transport}		Paramedic



Subject: Hemorrhage Control

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Consult

- If the AEMT or Paramedic needs assistance with a decision to administer **Tranexamic Acid (TXA)**
- Severe post-partum hemorrhage with **Tranexamic Acid (TXA)** as mentioned in [4007 Childbirth with Complications](#), call MCP for orders
- If uncontrolled bleeding in the adult airway after **500 mg Tranexamic Acid (TXA)** nebulized, contact MCP for orders to administer the remainder 1.5 grams IV/IO

Clinical Pearls

- For **Tranexamic Acid (TXA)**, time since injury must be less than three hours.
 - Greatest benefit will be to patients within 1 hour of injury
- Note and report the time of injury and the time of TXA administration

END OF SECTION



Subject: Hypothermia

Effective: June 1, 2021

Last modified: Oct. 11, 2021

3016.1 Clinical Management

Assessment						
Pediatric Considerations None	Signs & Symptoms - Cold, clammy skin - Shivering - Mental status changes - Extremity pain or sensory abnormality - Bradycardia - Hypotension or shock	Differential Diagnosis - Sepsis - Hypoglycemia - Stroke - Head Injury - Spinal cord injury				
Treatment Algorithm						
- Move patient to warm environment, remove all wet clothing, dry the patient, and cover with blankets. - Avoid any rough movement that may cause cardiac dysrhythmias or cardiac arrest. - It may be beneficial to consider spinal motion restriction measures. - Assess neurological status. - Oxygenate the patient with 100% O₂. - If patient goes into cardiac arrest: - CPR continuously - In severe hypothermia (less than 86°F [30°C]), limit defibrillation attempts to <u>one</u> except on orders from MCP. - If body temperature is (more than 86°F [30°C]), follow normal arrest protocols.			EMR	EMT	AEMT	Paramedic
- If available, provide {warmed and humidified} 100% O₂. - Hypothermic patients should be transported to a Trauma Center. - Resuscitative efforts should be continued while in transit, even if there is no response.						
- Use the least invasive means possible to secure airway. - Intubate if necessary, as gently as possible. - Establish vascular access and consider {warmed} fluids.						
Treat bradycardia only if patient is hypotensive.						
Consult						
- Consult with MCP for cardiac arrest management of the severely hypothermic patient. - All levels should consult with MCP for orders to administer second and subsequent defibrillations. - Paramedics must consult with MCP for orders to administer cardiac arrest medications.						
Clinical Pearls						
- It may be necessary to assess pulse and respirations for up to 45 seconds to confirm arrest. - Do not initiate CPR if there is any pulse present, no matter how slow.						
END OF SECTION						

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Subject:

Spinal Motion Restriction

Effective:

June 1, 2021

Last modified:

Jan. 5, 2024

3017.1 General Guidelines

- a. Studies indicate traditional spinal restriction has risks and may even cause harm in some cases.
- b. Spinal Motion Restrictions allows for an assessment-based management of the injured patient.
- c. Spinal precautions should always be taken when dealing with at risk patients.
- d. This protocol does not indicate that providers do not immobilize the spine; it simply provides a different means of restriction in selected patients.
- e. These guidelines apply to providers at all certification levels.

3017.2 Blunt Trauma Patients – Full Immobilization

- A All patients with clinical indications of a spinal injury and/or with altered levels of consciousness must be immobilized with both a C-collar and a spinal restriction device. (e.g., spine board, KED, vacuum splint).
- P Pediatric trauma patients less than 3 years of age with a GCS of less than 15 must be immobilized with both a C-collar and a spinal restriction device.

3017.3 Blunt Trauma Patients – SMR

- a. Other alert trauma patients, including all those listed below, should have a c-collar placed and moved with caution in-line as a unit to the cot. They would not need a backboard:
 - i. Patients with neck pain
 - ii. Patients with midline neck or spinal tenderness
 - iii. Patients with pain upon motion of the neck
 - iv. Cases with high risk mechanism (high speed MVC, fall greater than 10 feet, axial loading injury)

3017.4 Penetrating Trauma

- a. Patients with penetrating trauma do not need immobilization with either a cervical collar or backboard.
- b. Delays in transport are to be minimized and place the patient at greater risk.

3017.5 Airway or Ventilatory Management

- a. Patients who are immobilized and require airway and or ventilatory interventions (including intubation) may have the cervical collar removed during the intervention.
 - i. In-line stabilization should be maintained while the intervention is performed.
- b. The cervical collar should be reapplied after the intervention is either accomplished or abandoned.

3017.6 Equipment Issues

- a. In an emergency situation with equipment intensive sports such as football, hockey and lacrosse, the protective equipment shall be removed prior to transport to an emergency facility.
- b. Helmets of any kind that prevent either effective SMR or airway management should be removed.

3017.7 Other Considerations

- a. Patients greater than 69 y/o should be considered “high risk” patients for spinal injury and require closer assessment. With these patients, lean towards applying a cervical collar.
- b. If the patient meets the standards for a Trauma Alert Activation, consider a cervical collar at a minimum.
- c. Patients who do not tolerate any level of restriction should have that restriction adjusted to the point of removal if necessary based on clinical response.
 - i. Examples include shortness of breath, anxiety, and body habitus
 - ii. They should be transported in the manner of restriction that they can tolerate.
- d. Spinal restriction of the purpose of patient movement
 - i. Spinal restriction devices may be utilized for movement from a site of injury to the cot.
 - ii. Patients who do not require restriction should be removed from the device prior to transport.



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3017.8 Clinical Management

Full Spinal Motion Restriction

- Patients with GCS less than 15 including confusion and intoxication
- Patients with altered LOC
- Patients with neurologic deficits including paralysis
- Patients with clinical indications of a spinal injury
- Patients less than 3 y/o with GCS less than 15

C-Collar and Move In-line to Cot

- Patients that have a GCS of 15 and present with:
 - Neck pain
 - Midline neck tenderness
 - Pain on motion of the neck
 - High risk Mechanism of Injury (MOI)

SMR Is Not Required

- Penetrating trauma
- Patients that do not fall into the other two conditions

EXCEPTIONS

- Patients who require airway or ventilatory intervention may have the collar removed with inline stabilization during the intervention.
- Patients who do not tolerate restriction should have it adjusted to the point of removal if necessary.

END OF SECTION



Subject: Trauma Triage Guidelines

Effective: June 1, 2021

Last modified: Jan. 6, 2024

3018.1 Interpretation of Trauma Triage Guidelines

- a. This guideline meets the requirement of OAC 4765-14, defining Trauma Triage Guidelines for the region
- b. This guideline can separately provide direction as to when a provider should call a "Trauma Alert"
- c. Not all patients who meet Trauma Triage Criteria may need a trauma alert.

3018.2 State of Ohio Trauma Triage Age Considerations

- a. For the purposes of trauma guidelines, the criteria for patient age are:

- P Less than 16 years old will be pediatric patients**

- A 16 years old to 69 years old will be adult patients**

- G Greater than 69 years old will be geriatric patients**

3018.3 Trauma Center or Facility Capabilities:

- a. Level I and II Trauma Centers can care for the same trauma patients.
- b. Level III Trauma Centers offer services, based on individual hospital resources that provide for initial assessment, resuscitation, stabilization, and treatment of the trauma patient.
- c. In some areas of the region a Level III Trauma Center is the only trauma facility within 30 minutes ground transport time. This hospital may act as the primary receiving facility for the critically injured patient.
- d. In areas where the trauma patient is closer to a Level III Trauma Center, but a Level I or Level II Trauma Center is still within 30 minutes, the EMS Provider should decide whether the patient would benefit more from an immediate evaluation, stabilization, and treatment at the Level III Trauma Center, or from direct transport to a Level I or Level II Trauma Center.
- e. In areas of the region where there are no Trauma Centers within 30 minutes ground transport time, the acute care hospital may act as the primary receiving facility for critically injured trauma patients, or EMS Provider may arrange for air medical transport from the scene.
- P If a pediatric patient meets the trauma triage guidelines, transport to a Pediatric Trauma Center.**
- P Pediatric patients should be transported in an appropriately sized child restraint system.**
- h. If transportation time is greater than 30 minutes, transport to the nearest acute care hospital, or EMS providers may arrange for air medical transport from the scene.
- i. All pregnant trauma patients should be rapidly transported to the nearest Adult Trauma Center with labor and delivery capabilities, unless transport time is greater than 30 minutes.

3018.4 Air Medical Transportation:

- a. Prolonged delays at the scene waiting for air medical transport should be avoided.
- b. Cardiac arrest is not appropriate for air transport.
- c. In the rural environment, direct transfer of trauma patients by air medical transport may be appropriate.

3018.5 Exceptions to Transportation Guidelines:

- a. It is medically necessary to transport the victim to another hospital for initial assessment and stabilization before transfer to a Trauma Center.
- b. It is unsafe to transport the victim directly to a Trauma Center due to adverse weather or ground conditions or excessive transport time.
- c. Transporting the victim to a Trauma Center would cause a shortage of local EMS resources.
- d. No Trauma Center is able to receive and provide trauma care to the victim without undue delay.



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Trauma Triage Guidelines

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- e. Before transport begins, the patient requests to be taken to a particular hospital even if it is not a Trauma Center.
 - i. If the patient is a minor or otherwise considered incapable of making medical decisions, an adult relative or other legal representative may make this request.

3018.6 Trauma Criteria

a. Anatomical Criteria:

- i. Penetrating trauma to head, neck, torso
- ii. Significant, penetrating trauma to extremities proximal to elbow or knee with evidence of neurovascular compromise.
- iii. Injuries to the head, neck, or torso where the following physical findings are present:
 - 1. Visible crush injuries
 - 2. Abdominal injury with tenderness, distention, or seat belt sign
 - 3. Evidence of pelvic fracture
 - 4. Flail chest
- iv. Injuries to extremities where the following physical findings are present:
 - 1. Amputation proximal to wrist or ankle
 - 2. Visible crush injuries
 - 3. Fractures of two or more proximal long bones
 - G** One proximal long bone fracture in MVC only
 - 4. Evidence of neurovascular compromise
- v. Signs and symptoms of spinal cord injury
- vi. 2nd or 3rd degree burns greater than 10% total body surface area (BSA) or other significant burns involving the face, feet, hands, genitals, or airway
 - G** Injury sustained in two or more body regions
- viii. Open skull fracture

Meets Above Criteria = Transport to Trauma Center	Does Not Meet Above Criteria = Continue Assessment
Call Trauma Alert if patient presentation indicates	Assess for Physiologic Criteria

b. Physiological Criteria:

i. Adult Physiological Criteria

- A** GCS less than or equal to 13
- A** Loss of consciousness greater than five minutes
- A** Deterioration in level of consciousness at the scene or during transport
- A** Failure to localize pain
- A** Respirations less than 10 or greater than 29
- A** Needs ventilatory support
- A** Requires relief of tension pneumothorax
- A** Pulse greater than 120 in combination with evidence of hemorrhagic shock
- A** SBP less than 90 or absent radial pulse with carotid pulse present



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ii. Pediatric Physiological Criteria:

- P GCS less than or equal to 13
- P Loss of consciousness greater than five minutes
- P Deterioration in level of consciousness at the scene or during transport
- P Failure to localize pain
- P Evidence of poor perfusion (e.g., weak distal pulse, pallor, cyanosis, delayed capillary refill, tachycardia)
- P Evidence of respiratory distress or failure (e.g., stridor, grunting, retractions, cyanosis, nasal flaring, hoarseness, or difficulty speaking)
- P Respiratory rate less than 20 per minute in infants less than 1 year old

iii. Geriatric Physiological Criteria:

- G GCS less than or equal to 13
 - a. GCS less than or equal to 14 with evidence of Traumatic Brain Injury
- G Loss of consciousness greater than five minutes
- G Deterioration in level of consciousness at the scene or during transport
- G Failure to localize pain
- G Respirations less than 10 or greater than 29
- G Needs ventilatory support
- G Requires relief of tension pneumothorax
- G Pulse greater than 120 in combination with evidence of hemorrhagic shock
- G SBP less than 100 or absent radial pulse with carotid pulse present

Meets Physiological Criteria = Transport to Trauma Center**Call Trauma Alert if patient presentation indicates****Does Not Meet Above Criteria = Continue Assessment**

Look at Special Considerations

c. Special Considerations:

- i. Vehicle telemetry provides data consistent with high risk of injury
- ii. On scene fatality in the same vehicle
- G Pedestrian struck by a motor vehicle
- G Falls from any height, including standing falls, with evidence of traumatic brain injury

Special Considerations = Transport to Trauma Center**Call Trauma Alert if patient presentation indicates****Does Not Meet Above Criteria = Consider MOI**

Transport to most appropriate hospital

d. Mechanism of Injury:

- i. Auto-pedestrian/auto-bicycle injury with significant (faster than 5 mph) impact
- ii. Ejection from motor vehicle or unrestrained rollover
- iii. Extrication time longer than 20 minutes
- iv. Fall of more than 20 feet
 - P Fall greater than 3 times child's height
- v. High-speed auto crash
 - 1. Estimated speed faster than 40 mph



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2. Intrusion into passenger compartment of more than 12 inches
3. Major auto deformity of more than 20 inches
- vi. Open motor vehicle crashes faster than 20 mph or with separation of rider from vehicle
- vii. Pedestrian thrown or run over

e. Special Situations:

- i. Pre-existing cardiac or respiratory disease
- ii. Diabetes, cirrhosis, morbid obesity, seizure disorder
- iii. Patient with bleeding disorder or on anticoagulants or anti-platelets
- iv. Immuno-suppressed patients (renal dialysis, transplant, cancer, HIV)
- v. Congenital disorder

MOI or Special Considerations = Consider Trauma Center	No Significant MOI or Special Considerations
No need to call Trauma Alert if no significant symptoms	Transport to most appropriate hospital

END OF SECTION



Subject: SALT Triage System

Effective: June 1, 2021

Last modified: Dec. 7, 2024

3019.1 General Guidelines

- a. SALT stands for Sort, Assess, Life-Saving Intervention, and Treatment/Transport.
- b. Developed by the Centers for Disease Control and Prevention to address limitations in other systems.
- c. The CDC has proposed SALT as the national standard for Mass Casualty Incident (MCI) triage.
- d. SALT has the advantage of being the **fastest mass casualty triage system**.
- e. Notify hospitals of any MCI, especially a MCI involving contaminated patients.
 - i. Consider use of the [3020 Regional Hospital Notification System \(RHNS\)](#)

3019.2 Primary and Secondary Triage Prior to Transport

- a. Initial Triage:
 - i. Use triage ribbons (color-coded strips), not treatment tags, during initial triage.
 1. Treatment tags slow the process and should be used later, in the treatment areas.
 2. Treatment tags do need to be used at some point as they are sometimes the only documentation of EMS assessments and treatments.
 - ii. Tie the triage ribbon to an upper extremity in a **VISIBLE** location (on the right wrist, if possible).
 - iii. SALT Triage Levels:
 1. **RED – Immediate**
 2. **YELLOW – Delayed**
 3. **GREEN – Minimal**
 4. **GRAY – Expectant** (The patient is unlikely to survive given the current resources)
 5. **BLACK – Dead** (black & white zebra stripe for easier visibility in low light)
 6. **ORANGE and Polka Dot** - used in addition to one of the above ribbons to indicate victim has been contaminated with a hazardous material.
 - iv. Move as quickly and safely as possible; making quick decisions.
 - v. Victims will be re-triaged, probably multiple times. Revise the triage category as often as indicated.
 - vi. Over-triage can be as harmful as under-triage. If everyone is tagged red, those who are truly red will receive delayed treatment, delayed transport, and delayed definitive care.
 - vii. Treatment and transport should NOT be delayed especially for critical patients. Get the reds out.
 - viii. If there are extensive delays in the field, consider requesting orders for palliative care, e.g., pain medications if time and resources allow.
- b. Secondary Triage:
 - i. Reassess (i.e. secondary triage) as often as practical, including when the patient is moved to the Casualty Collection Point (CCP) or Treatment Area, and on all victims prior to transport.
 1. Also reassess patients when their condition or resources available change.
 - ii. Apply Treatment Tags after patients enter the CCP, or in the Transport Area (by the Transport Officer/Group) if the patient is being directly removed without going to the Treatment Area.
 - iii. Crews can also fill in pertinent and available information on the Tag during transport.
 - iv. Use the patient's ribbon to tie on the treatment tag
 1. Use treatment tags with individual barcodes consistent with this Standing Order and Juvare EMTrack, the patient tracking system used throughout Ohio.
 - v. **Orange & Polka-dot** ribbons (indicating contaminated patients) are removed after decontamination.
 1. Each contaminated patient initially receives two ribbons: one with the triage category (**Red, Yellow, Green, Gray, or Black**), and the second, the **Orange & Polka-dot** ribbon indicating contamination.
 2. EMS is responsible for performing primary decontamination prior to transport. However,



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the hospital must be made aware of both contamination and the decontamination procedures taken.

3. Make sure to decontaminate under the ribbons.
4. After decontamination, remove the **Orange & Polka-dot** ribbon.
5. Mark treatment tags for contaminated patients with two check marks on the orange strip:
 - a. Mark both the “dirty” and “decontaminated” boxes.
 - b. This indicates to the hospital personnel that the patient has had field decontamination, but may still be somewhat “dirty”.

c. Transport:

- i. Treatment Area or Transport Group personnel determine priority for transport.
- ii. Distribution of patients among various hospitals is one of EMS’ most crucial tasks.
- iii. **Do not overload any hospital**, regardless of transport distance to other hospitals.
 1. Consider use of Juvare EMResource from the scene to monitor hospital triage capabilities according to **RED**, **YELLOW** and **GREEN** patient categories.
- iv. In an MCI, transport trauma patients to non-Trauma Centers as necessary.
 1. All hospitals will accept and stabilize trauma patients during MCIs.
 2. Consider transporting minor (**GREEN**) patients to satellite EDs to relieve pressure on Trauma Centers and other hospitals.
- v. When assigning patient allocation, consider the likelihood that the closest hospitals may be overwhelmed by patients who were not transported by EMS.
- vi. In large scenarios, consider activation of the Forward Movement of Patients Plan as defined in [3021 Crisis Standards of Care in Massive Events](#).

3019.3 Sort, Assess, Life-Saving Intervention, Treatment/Transport Process

a. Sort

- i. Global Sorting: Action 1
 1. Action: “Everyone who can hear me please move to [designated area] and we will help you” (use loudspeaker if available)
 2. Goal: Group ambulatory patients using voice commands
 3. Result: Those who follow commands are *last* priority for individual assessment (Green)
 4. Assign someone to keep them together and notify Incident Command or EMS Group/Branch of number of patients and their location.
 5. Do not forget these victims.
 6. Someone must re-triage them as soon as possible.
 7. In smaller incidents, such as a motor vehicle crash with few victims that you do not want to move on their own, skip Action 1, and go to Global Sorting Action 2
- ii. Global Sorting: Action 2
 1. Action: “If you need help, wave. We will be there to help as soon as possible”
 2. Goal: Identify non-ambulatory patients who can follow commands or make purposeful movements
 3. Result: Those who follow this command are second priority for individual assessment
- iii. Global Sorting: Result
 1. Casualties are now prioritized for individual assessment
 - a. Priority 1: Still, and those with obvious life threat
 - b. Priority 2: Waving or purposeful movements
 - c. Priority 3: Walking



Subject: SALT Triage System

Effective: June 1, 2021

Last modified: Dec. 7, 2024

- iv. Begin assessing all non-ambulatory victims where they lie, performing Life Saving Interventions (LSIs) as needed, within your scope of practice, using the equipment is readily available.

b. Assess

i. Is the patient breathing?

1. If not, open the airway. In children, consider giving two rescue breaths.
2. If the patient is still not breathing, triage them to **BLACK** (dead).
3. Do not move patients triaged **BLACK** except to gain access to a living patient.
4. If patient is breathing, conduct next assessment.

ii. Assess for the following:

1. Can the patient follow commands or make purposeful movements?
2. Does the patient have a peripheral pulse?
3. Is the patient not in respiratory distress?
4. Is hemorrhaging under control?

iii. Grading the Assessment

1. If the answer to any of those questions is no (bad) and the patient **IS** likely to survive given current resources, tag them as **RED (Immediate)**.
2. If the answer to any of those questions is no (bad) and the patient is **NOT** likely to survive given current resources, tag them as **GRAY (Expectant)**.
3. If the answer to all of those questions is yes but injuries are not minor and require care, tag patient as **YELLOW (Delayed)**.
 - a. **YELLOWs** have serious injuries and need care, though not as urgently as **REDs**.
 - b. On secondary triage, some **YELLOWs** will need higher priority transport than others.
4. If the answers to all of those questions is yes and the injuries are minor, tag patient as **GREEN (Minimal)**.

Two mnemonics to remember the four assessment questions

C – follows Commands

R – No Respiratory distress

A – No (uncontrolled) Arterial bleeding

P – Peripheral Pulse Present

Think of the questions in terms of “bad” or “good”

If the answer to any of the questions is “bad” then the patient is tagged either **RED** (Immediate) or **GRAY** (Expectant)

c. Life Saving Interventions

i. Only correct life-threatening problems during triage.

1. Control major hemorrhage
2. Open airway (if child, consider giving two rescue breaths)
3. Needle chest decompression
4. Auto injector antidotes
5. See [3019.5 Special Situations](#)

ci. Treatment/Transport

i. Transport/treatment priority is typically given (in order) to

1. **RED (Immediate)**
2. **YELLOW (Delayed)**
3. **GREEN (Minimal)**
4. **GRAY (Expectant)** patients should be treated and transported as resources allow.



Subject: SALT Triage System

Effective: June 1, 2021

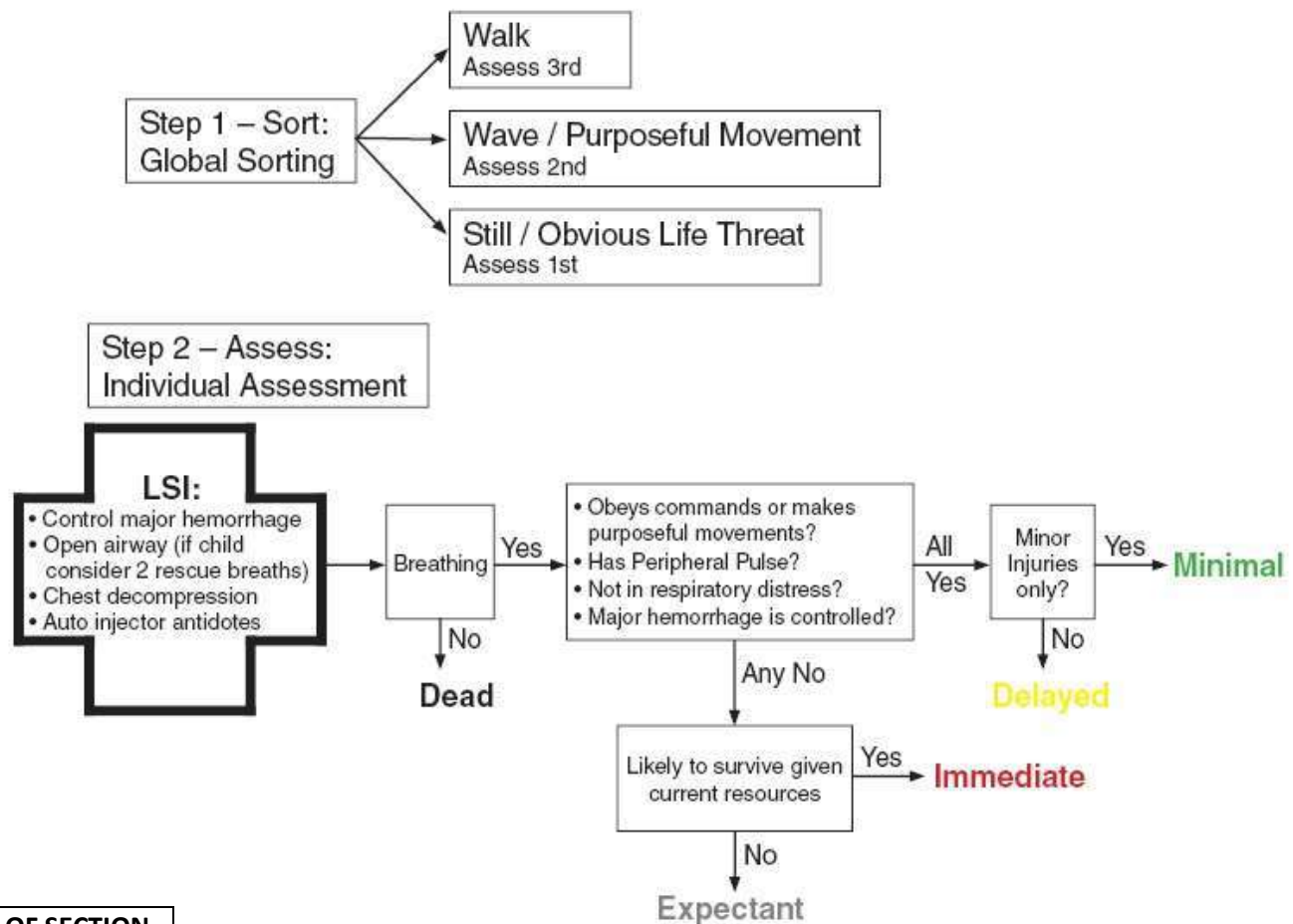
Last modified: Dec. 7, 2024

3019.4 General Considerations

- Patients must be reassessed periodically, including when moved to the CCP, or when their condition or resources change.
- Even after applying treatment tags, the main indicator of patient condition is the triage ribbon.
- Continue to use the same tag, even if the condition changes repeatedly, changing the ribbon to indicate the patient's current condition.
- If the patient's condition or the triage priority changes, indicate that on the tag.

3019.5 Special Considerations

- SALT is a clinical guideline, not an absolute.
- Every MCI is extraordinary - use your clinical judgement
- A patient who is **GRAY** (Expectant) initially can become **RED (Immediate)** as soon as resources are available.
- MCIs with patients suffering traumatic (aka, compression) asphyxia who are not breathing initially, may start breathing after just a few ventilations.
 - Common to crowding situations and crowd surges, it is worth attempting a few ventilations during the LSI step, even in adults.
- In MCIs due to lightning strikes, the pathology can be very complex.
- Consider attempting ventilation or defibrillation, depending on resources and the conditions of other victims.

3019.6 SALT Triage Flow Chart

END OF SECTION



Subject: Regional Hospital Notification
System (RHNS)

Effective:
June 1, 2021

Last modified:
June 16, 2024

3020.1 General Guidelines

- a. The purpose of the Regional Hospital Notification System (RHNS) is to provide one number for EMS, hospitals, and EMAs to call that will make rapid, simultaneous notifications in a Mass Casualty Incident or Event (MCI/MCE), or other major emergency.
- b. The system can be used when an incident could involve a significant number of the region's hospitals.
- c. RHNS is critical. Activation should be a **high priority** to alert hospitals and regional coordinators. Early warning will start processes essential to handle major events.

3020.2 RHNS Activation

- a. To activate the RHNS, call 937-333-USAR (8727).
- b. The agency calling must ask for a Dispatch Supervisor, request a "Regional Hospital Notification", and use the phrase "Mass Casualty Page Hospitals". Then the agency calling will provide the following:
 - i. Name of agency
 - ii. Nature of emergency
 - iii. Location of emergency
 - iv. General statement on severity, such as approximate number of victims
 - v. Any other information to be conveyed
- c. The Montgomery County Regional Dispatch Center (RDC) will immediately put out a computerized message to the RHNS Group with the information provided.

END OF SECTION

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Subject: Crisis Standards of Care in
Massive Events

Effective: June 1, 2021

Last modified: June 16, 2024

3021.1 General Guidelines

- a. Some incidents are so large as to require extraordinary EMS procedures. Those scenarios are sometimes referred to as Mass Casualty Events (MCEs), instead of Mass Casualty Incidents (MCIs).
- b. These EMS procedures should be utilized in very large emergency scenarios, or when the duration is extended.
- c. In the event of an MCE, especially one lasting days or longer, Greater Miami Valley EMS Council, with the approval of the Regional Physicians Advisory Board (RPAB), may promulgate "Just in Time Standing Orders" (JITSO).
- d. With approval from Ohio Department of Public Safety, these orders might include triage standards for transport to other healthcare facilities and other crisis standards of care; possibly exceeding the standard scope of practice for EMS.
- e. Full information on the process can be found in the Dayton MMRS Regional MCI Plan Template

3021.2 Alternate Transports

- a. In some circumstances, EMS may be authorized to triage selected patients for transport to other healthcare facilities, including:
 - i. Urgent Care Centers
 - ii. Acute Care Center (ACC)
 - iii. Neighborhood Emergency Help Center (NEHC)
 - iv. Disaster Medical Assistance Team (DMAT)

3021.3 Forward Movement of Patients

- a. Planned by Dayton MMRS
- b. The intent is to relieve the burden on local hospitals by transporting patients, possibly directly from the scene, to more distant hospitals.

3021.4 Functional Needs Shelter Triage

- a. A regional protocol for Functional Needs Shelter Triage has been added to the Optional Standing Orders Manual and is also available at gmvemsc.org on the Training Materials page.
- b. Will help determine whether individuals with functional needs can be safely sheltered in a Red Cross Shelter during a disaster
- c. This Shelter Triage Protocol is a pre-approved Just-In-Time Standing Order (JITSO), authorized by the RPAB for use by an EMS agency assisting the Red Cross with shelter triage.
- d. It is intended to be printed and given to paramedics, nurses, and other healthcare personnel at the time of a shelter operation.
- e. At the option of local department chiefs and medical directors, the same protocol can be used during a disaster to determine patients who would be more appropriate for transport to Red Cross Shelters than to hospitals.
- f. In those cases, EMS should, if possible, contact the shelter before transporting.
- g. If locations or contact information for shelters is not known, contact the County EMA or the Red Cross.
- h. When transporting these non-emergency patients to shelters, it is critical that the patients bring their medications and medical equipment with them.

END OF SECTION

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4000 Series

Medical Protocol

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Subject: Abdominal Pain

Effective: June 1, 2021

Last modified: May 16, 2026

4001.1 General Guidelines

- Ensure an abdominal exam which includes inspection, auscultation and palpation is performed and documented on every patient with abdominal pain.
- Assess all abdominal pain patients for trauma, pregnancy, illness, or potential ingestion.

4001.2 Clinical Management

Assessment					
Pediatric Considerations None	Signs & Symptoms - Pain (location/migration) - Tenderness (point, palpation, rebound) - Nausea and/or vomiting - Diarrhea - Dysuria - Constipation - Vaginal bleeding/discharge - Pregnancy	Differential Diagnosis - Hepatitis - Peptic ulcer disease/gastritis - Gallbladder - Pancreatitis - Abdominal aneurysm - Appendicitis - Pelvic (PID, ovarian cyst, ectopic pregnancy) - Diverticulitis - Gastroenteritis - Bladder/prostate disorders - Kidney stone - Myocardial infarction - Pneumonia - Pulmonary embolus			
Treatment Algorithm					
- Place patient in position of comfort. - Give nothing by mouth.				EMR	Paramedic
- For nausea or active vomiting: - A Consider Ondansetron 4 mg PO dissolving tablet - P Consider Ondansetron 4 mg PO dissolving tablet if patient is 12 y/o or older				EMT	
- For nausea or active vomiting: - A Ondansetron 4 mg slow IV/IO. - P Ondansetron 0.1 mg/kg IV/IO (max 4 mg) if patient is 5 y/o or older and weight is more than or equal to 30 kg. - For pain relief: - A Including with unilateral flank pain, consider 1014 Pain Management Protocol. - P ♦ For pain relief, call MCP for orders.				AEMT	
A For nausea or if no IV access established, Ondansetron 4 mg PO (dissolving tablet) or consider administering 4 mg (2 ml) of the IV form PO by spraying it into the patient's mouth.					
Consult					
The AEMT and Paramedic need MCP orders when providing abdominal pain relief to pediatric patients.					
Clinical Pearls					
The Paramedic can administer the IV form of Ondansetron orally to adults by spraying it into the patient's mouth.					
END OF SECTION					

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Subject: Allergic Reaction/Anaphylaxis

Effective: June 1, 2021

Last modified: May 16, 2026

4002.1 General Guidelines

- Epinephrine is the mainstay of anaphylaxis in allergic reaction treatment.
- Epinephrine is particularly important in cases of any airway edema, hypotension, or when multiple body systems are involved.
- Advanced age is not a contraindication to epinephrine.

4002.2 Clinical Management

Assessment			
Pediatric Considerations - Epinephrine is dosed based on weight, not age. - While the protocol lists those patients under 15 kg as pediatric, it is understood that patients equal to or greater than 30 kg will get both the Adult EpiPen and the EpiPen Jr. or Epinephrine 0.5 mg, no matter what their age.	Signs & Symptoms - Itching - Hoarseness or stridor - Wheezing - Respiratory distress - Altered level of consciousness - Cyanosis - Pulmonary edema - Facial/airway edema - Urticaria/hives	Differential Diagnosis - Rash only - Shock (vascular effect) - Angioedema - Aspiration/airway obstruction - Vasovagal event - Asthma	
Treatment Algorithm			
- Provide O₂ as needed. - If allergic reaction: - If equal to or greater than 30 kg, give both Adult EpiPen and EpiPen Jr. (adult and pediatric) P If less than 15 kg, EpiPen Jr. P If equal to or greater than 15 kg and less than 30 kg, Adult EpiPen - If applicable, apply ice pack. ♦ If symptoms persist, may repeat Epinephrine (adult and pediatric) in 10 minutes. - Call for transport.		EMR	
- If patient develops wheezing, assist them with their prescribed metered dose inhaler or Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized with O₂ flowing at 8-10 LPM. - May repeat Albuterol 2.5 mg nebulized two times {If allergic reaction and an absence of Epi-pens in the drug bag, EMTs are permitted to administer Epinephrine IM via a syringe} {The EMT may only perform this skill after authorization and training from their Medical Director} - If equal to or greater than 30 kg, Epinephrine (1:1,000) 0.5 IM, (adult and pediatric) P {If equal to or greater than 15 kg and less than 30 kg, Epinephrine (1:1,000) 0.3 mg IM} P {If less than 15 kg, Epinephrine (1:1,000) 0.15 mg IM} ♦ {May repeat Epinephrine (1:1,000) 0.5 mg IM after 10 minutes (adult and pediatric)} P ♦ {May repeat Epinephrine (1:1,000) (at weight appropriate dose) after 10 minutes}		EMT	
- If an allergic reaction: - If equal to or greater than 30 kg, give both Adult EpiPen and EpiPen Jr or Epi (1:1,000) 0.5 mg IM (adult and pediatric) P If equal to or greater than 15 kg and less than 30 kg, Adult EpiPen or Epinephrine (1:1,000) 0.3 mg IM P If less than 15 kg, EpiPen Jr or Epinephrine (1:1,000) 0.15 mg IM - May repeat Epinephrine (1:1,000) 0.5 mg IM after 10 minutes (adult and pediatric) P May repeat Epinephrine (1:1,000) (at weight appropriate dose) after 10 minutes - If apneic, intubate, possibly with smaller than normal ET tube. - If patient intubated, Albuterol 2.5 mg by nebulizer into the ETT. If Ipratropium not given before intubation, add to first Albuterol. - If hypotensive: A IV fluid to maintain adequate BP. P IV fluid 20 ml/kg IV to maintain adequate BP. A Diphenhydramine 50 mg IM or IV P Diphenhydramine 1 mg/kg IM or IV (max dose 50 mg).			AEMT



Subject: Allergic Reaction/Anaphylaxis

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- If patient deteriorating or unresponsive, consider early intubation, possibly with a smaller than normal size endotracheal tube
- If a conscious patient requires intubation:
 - A **Lidocaine 100 mg IN** half dose per nostril or added to nebulizer with breathing treatment.
 - P **Lidocaine 1.5 mg/kg nebulized with O₂ 8-10 LPM or IN. Maximum dose is 100 mg.**
- If patient remains hypotensive after IV fluid, **Epinephrine (1:10,000) 0.1 mg, slow IV**, every 3 minutes up to 0.5 mg.
- **Solu-Medrol 125 mg IV**
- P **Solu-Medrol 2 mg/kg IV, max dose 125 mg.**

Paramedic

Consult

- The EMR and EMT need MCP orders to administer repeat epinephrine.
- EMT needs MCP orders to administer breathing treatments.

Clinical Pearls

- The EMT may only perform Epinephrine 1:1,000 draws and injections after authorization and training from their Medical Director
- No significant change in patient condition in the field should be expected from the administration of Solu-Medrol.
- Solu-Medrol will be given to all patients treated within the allergic reaction or anaphylaxis protocol only after all other applicable first-line medications have been delivered.

END OF SECTION



Subject: Asthma/Emphysema/COPD

Effective: June 1, 2021

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4003.1 Clinical Management

Assessment			
Pediatric Considerations • Younger patients may exhibit nasal flaring • Epinephrine is dosed based on weight, not age. • While the protocol lists those patients under 15 kg as pediatric, it is understood that patients equal to or greater than 30 kg will get Epinephrine 0.5 mg IM, no matter what their age.	Signs & Symptoms • Shortness of breath • Pursed lip breathing • Increased respiratory rate and effort • Wheezing, rhonchi • Accessory muscle use • Cough • Tachycardia • Tripod position	Differential Diagnosis • Anaphylaxis • Aspiration • Pleural effusion • Pneumonia • Pulmonary embolus • Pneumothorax • Cardiac event (AMI or CHF) • Pericardial tamponade • Hyperventilation • Inhaled toxins	
Treatment Algorithm			
• Provide O₂ as needed. • Call for transport.		EMR	
• If patient develops wheezing, assist them with taking their prescribed metered dose inhaler or ○ Consider Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized with O₂ flowing at 8-10 LPM ○ May repeat Albuterol 2.5 mg nebulized two times • For any patient who is bronchial constricted: CPAP ○ If indicated, consider administration of nebulized Albuterol 2.5 mg and Ipratropium 0.5 mg, in line with CPAP • Transport unless ALS intercept is less than 5 minutes.		EMT	
• If patient intubated, Albuterol 2.5 mg by nebulizer into the ETT. If Ipratropium not given before intubation, add to first Albuterol. • After intubation of an asthma patient, limit rate of ventilation to avoid auto-PEEP and hypotension, provided that you can adequately oxygenate the patient at below rate: - A 8-10 breaths per minute for adults - P 10-15 breaths per minute for pediatric patients • Consider needle decompression in the presence of auto-PEEP or hyperinflation: ○ If the patient is in cardiac arrest, perform bilateral needle decompression ○ If unilateral or bilateral diminished breath sounds and the patient is hemodynamically unstable, consider decompression of only the affected sides ○ Decompression sites: ▪ Fourth or fifth intercostal space in the mid-axillary line ▪ Second or third intercostal space in the mid-clavicular line (use nipple line as a guide) - P In patients less than 8 years old, decompression site choice will be limited to the 2nd or 3rd intercostal space at the mid-clavicular line • Asthmatics in severe distress (NOT for emphysema or COPD): ○ If equal to or greater than 30 kg, give both Adult EpiPen and EpiPen Jr or Epi (1:1,000) 0.5 mg IM (adult and pediatric) - P If equal to or greater than 15 kg and less than 30 kg, Adult EpiPen or Epinephrine (1:1,000) 0.3 mg IM - P If less than 15 kg, EpiPen Jr or Epinephrine (1:1,000) 0.15 mg IM ○ May repeat Epinephrine (1:1,000) 0.5 mg IM after 10 minutes (adult and pediatric) - P May repeat Epinephrine (1:1,000) (at weight appropriate dose) after 10 minutes		AE/MT	
• If a conscious patient requires intubation: - A Lidocaine 100 mg IN half dose per nostril or added to nebulizer with breathing treatment. - P Lidocaine 1.5 mg/kg nebulized with O₂ 8-10 LPM or IN. Maximum dose is 100 mg. • For any persistent bronchial constriction consider CPAP or {Bi-PAP} - A Solu-Medrol 125 mg IV - P Solu-Medrol 2 mg/kg IV, max dose 125 mg. • Bronchoconstriction due to asthma refractory to Albuterol, Ipratropium, and Epinephrine: - A ♦ Magnesium Sulfate 2 gm (half bag) infused with macro-drip tubing over 10 minutes.			Paramedic

Subject: **Asthma/Emphysema/COPD**Effective: **June 1, 2021**Last modified: **May 16, 2026****Consult**

- The Paramedic needs MCP orders to administer Magnesium Sulfate to the asthmatic patient.

Clinical Pearls

- A patient who has received a breathing treatment should be transported for evaluation.
- No significant change in patient condition in the field should be expected from the administration of Solu-Medrol.

END OF SECTION



Subject: Behavioral Emergencies

Effective: June 1, 2021

Last modified: May 16, 2026

4004.1 General Guidelines

- Per Ohio Revised Code, EMS providers may not “pink slip” an individual even if they are threatening harm to themselves or others.
- Only a police officer, crisis worker, health officer, psychiatrist or licensed physician can administer an involuntary admission form (“pink slip”) for a patient.
- Each EMS department, in consultation with its medical director and local law enforcement, should have a procedure to deal with these types of situations.

4004.2 Precautions

- Consider staging until law enforcement has made the scene safe.
- Have law enforcement search patient for weapons.
- Consider possible medical causes for patient’s condition:

Anemia
Hypoxia
Hypoglycemia
Stroke
Dysrhythmias
Hypertension

Toxicological Emergency
Pulmonary embolism
Hemorrhage
Metabolic disorders
Seizures and postictal states
Shock

Infection (especially meningitis/encephalitis)
Electrolyte imbalance
Myocardial ischemia or infarction
Head trauma or intracranial pressure
Drug or alcohol intoxication, side effects
Drug withdrawal

4004.3 Clinical Management

Assessment		
Pediatric Considerations For the purpose of transporting to appropriate behavioral health treatment facilities, patients are considered “pediatric” if they are less than 18 years old.	Signs & Symptoms - Anxiety, agitation, confusion - Affect change, hallucinations - Delusional thoughts, bizarre behavior - Violent or combative - Expression of suicidal/homicidal ideations	Differential Diagnosis - Medical causes listed above - Other altered mental status issues - Alcohol intoxication/Substance abuse - Medication effect/overdose - Withdrawal symptoms - Depression - Bipolar (manic-depressive) - Schizophrenia - Anxiety disorders
Treatment Algorithm		
- Determine patient capacity and consent. - Take actions to prevent imminent harm to the patient or others, if it is safe to do so. - Do not judge, just treat. - Consider possible medical causes for patient’s condition		EMR
- If patient is unwilling to go to a facility, consider whether they are a candidate for a “pink slip” - Transport all patients who are not making rational decisions and who are a threat to themselves or others for medical evaluation. - A If possible, transport a mental health patient to the facility where the individual has been previously treated. - A In all other cases, patients greater than or equal to 18 years old should be transported to the closest ED. - P Patients less than 18 years old, with mental health issues should be transported to a facility with pediatric mental health capabilities. (see 4004.5 c. Dayton Children’s Hospital for specific instructions)		EMT
No additional orders at this level		AEMT
Severe agitation is a medical emergency, and should be treated. See 4005 Behavioral Emergencies - Combative Patient/Emergency Sedation		Paramedic
Consult		
If no one is available or willing to issue the “pink slip”, contact MCP for advice on how to proceed.		



Subject: Behavioral Emergencies

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Clinical Pearls

- Consider that a patient may be incapable to make medical decisions if they are:
 - Suicidal
 - Confused
 - Severely developmentally or mentally disabled
 - Intoxicated
 - Injured/ill with an altered mental status
 - Physically/verbally hostile
 - Unconscious
- When obtaining medical history, determine:
 - Suicidal or violent history
 - Previous psychiatric hospitalization, when and where
 - Location where patient receives mental health care
 - Medications
 - Recreational drugs/alcohol: amount, names
- Exceptions to the outlined transport recommendations include:
 - It is medically necessary to transport the patient to the closest hospital for stabilization.
 - It is unsafe to transport the patient to the preferred/recommended facility due to adverse weather or ground conditions or excessive transport time.
 - Transporting the patient to the preferred/recommended facility would cause a critical shortage of local EMS resources.
 - Patient requests transport to a different facility.

4004.4 Transport Guidelines

- a. Adults can be transported to any facility. Considerations should be given to the capabilities of the receiving facility, combativeness of the patient and length of transport time.
- b. Pediatrics
 - i. Transporting pediatric patients to Dayton Children's Hospital should be a priority.
 - ii. For excessive transport times, consider transport to a closer facility which can facilitate transfer after an initial assessment, stabilization, and a consult with DCH.
 - iii. While GMVEMSC Standing Orders consider a pediatric patient to be less than 16 years old, DCH will accept behavioral patients up to the age of 17 years old.
 - iv. Pregnant patients of any age should be transported to an adult facility.

4004.5 Pre-Arrival Notification of Behavioral Patients

- a. Premier Health Facilities
 - i. Acute Crisis Evaluation (ACE) is used to manage behavioral patients in an acute crisis.
 - ii. Activating an "ACE Alert":
 1. When contacting the receiving facility, state that you are requesting an "ACE Alert"
 2. You do NOT need to speak to Med-Control to activate the alert
 3. Proceed with report and ETA
 - iii. All behavioral health patients must go to an ED for evaluation; there is no direct EMS admission to inpatient behavioral health
- b. Kettering Health Network Facilities
 - i. KHN uses the code word "Dr. White" to activate a response when bringing in a patient who needs management for mental health or behavioral disturbances.



Subject:

Behavioral Emergencies

Effective:

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- ii. To notify KHN facilities of an arrival patient with behavioral needs:
 - 1. When contacting the receiving facility, state that you are requesting “Dr. White”
 - 2. You do NOT need to speak to Med-Control to activate the alert
 - 3. Proceed with report and ETA
 - iii. No direct transport to Kettering Behavioral Health Center. Patient must go through the ED.
- c. Dayton Children’s Hospital (Main Campus only)
- i. Acute Crisis Evaluation (ACE) is used to manage behavioral patients in an acute crisis.
 - ii. Activating an “ACE Alert”:
 - 1. When contacting the receiving facility, state that you are requesting an “ACE Alert”
 - 2. You do NOT need to speak to Med-Control to activate the alert
 - 3. Proceed with report and ETA
 - iii. Transport destination:
 - 1. **Between 0800 hrs. and 0100 hrs., transport to** the DCH Mathile Center for Mental Health and Wellness.
 - 2. **For all other hours (0101 hrs. to 0759 hrs.) transport to** the Main Campus Emergency Department.
 - iv. Transport Exceptions (Other reasons to transport to the Main Campus Emergency Department)
 - 1. **Pediatric patients in need of medical care (with the exception of minor cuts, minor stitching, and orthopedic injuries)**
 - 2. **Intoxication**
 - 3. **Sexual Assault**
 - 4. **Strangulation**
 - 5. **Patient received Ketamine**
 - 6. **Pregnancy**
 - v. Mathile Center for Mental Health and Wellness arrival instructions:
 - 1. Enter the facility’s site, continue past the gates, and arrive at the EMS Arrival Bays.
 - 2. Present to the sally port where Public Safety will escort crews into the Intake Center.

END OF SECTION

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Subject: Combative Patients/Emergency Sedation

Effective: June 1, 2021

Last modified: May 25, 2025

4005.1 General Guidelines

- Restrained patients must **not** be transported in a prone position with hands & feet behind their back.
- Restrained patients must **not** be sandwiched between backboards or other items.
- Always maintain the ability to remove restraints if the patient vomits or develops respiratory distress.

4005.2 Combative Patients

- Identified as irrational behavior, examples include aggression, violence, and/or paranoia in the patient.
- This state can result from causes including, but not limited to:
 - Stimulant intoxication
 - Psychiatric illness
 - Hypoglycemia
 - Other medical illnesses.
- The combative patient often becomes significantly hyperthermic and/or hypoxic, even after the episode has subsided.

4005.3 Collaboration with Law Enforcement

- Providers should get as much information as possible from LE as to the reason for the encounter, observed behaviors, medical history, use of force and/or less-than-lethal weapons, etc.
- Decisions to use pharmacological interventions will be made solely by the EMS provider based on their own assessment and not at the behest of the law enforcement officers on scene.
- Consideration should be given to transitioning the patient from police restraints to medical restraints prior to transport.
- If restraint devices require a key, that key should be transported **with** the patient whenever possible.
 - Provider agencies should work with their law enforcement partners beforehand to establish SOPs to address this concern.

4005.4 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Patient out of control and dangerous to self or others. - Restraint required for patient control without causing harm - Combative or violent patient	Differential Diagnosis - Alcohol intoxication - Substance abuse - Medication effect/overdose - Withdrawal symptoms - Mental health history - Medical causes listed in 4004
Treatment Algorithm		
- Explain the need for restraint to the patient. - Recheck often a restrained patient's ability to breathe and distal circulation.		EMR
No additional orders at this level		EMT



Subject: Combative Patients/Emergency Sedation

Effective: June 1, 2021

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A Ketamine 250 mg IM (in anterolateral thigh) or **Ketamine 100 mg slow IV**.

A No change after 10 minutes with IM dose or 5 minutes with IV dose, consider additional medication:

- DO NOT ADMINISTER KETAMINE AND MIDAZOLAM SIMULTANEOUSLY.
 - Give the administered sedative time to work before moving on to a secondary medication and dosing.
- A Ketamine 250 mg IM** (in opposite anterolateral thigh) or repeat **Ketamine 100 mg IV**.

AND/OR:

A Midazolam 10 mg IN (5 mg in each nostril), or **Midazolam 2.5 mg slow IV**, or **Midazolam 5 mg IM**.

A If necessary, repeat Midazolam doses:

- A** Repeat **Midazolam 5 mg IN** (2.5 mg in each nostril) after 10 minutes.
- A** or repeat **Midazolam 2.5 mg slow IV** after 5 minutes.
- A** or repeat **Midazolam 5 mg IM** after 10 minutes.

P If the patient is age 8 or greater, consider **Ketamine 1 mg/kg slow IV** (max dose 100 mg) or **Ketamine 5 mg/kg IM** (max dose 250).

or

P **Midazolam 0.2 mg/kg IN** (max IN dose 10 mg) or **Midazolam 0.1 mg/kg slow IV** (max IV dose 2.5 mg) or **Midazolam 0.2 mg/kg IM** (max IM dose 4 mg)

P ♦ Call MCP for additional Ketamine or Midazolam.

A If an excited delirium patient goes into arrest: ♦ Consider **Sodium Bicarbonate 100 mEq IV**

Consult

- MCP needed for pediatric repeat medications and (for the paramedic) Sodium Bicarbonate in cardiac arrest.

Clinical Pearls

- Consider advising the receiving facility of the combative patient as outlined in [4004 Behavioral Emergencies](#)
 - Premier Health Facilities: ACE Alert
 - Kettering Health Facilities: "Dr. White"
 - Dayton Children's Hospital: ACE Alert
 - For other hospitals in the region, check to see if they have a code word for notification
- Any patients who are restrained or sedated should be constantly monitored for an effective airway, adequate breathing, and circulation.
- Patients who have been sedated with Ketamine can be deeply unconscious and present with hypersalivation. Management should include use of and nasopharyngeal airway, proper positioning, and persistent suctioning to maintain a clear airway.
- Avoid transporting combative patients to satellite EDs if safe to do so.

END OF SECTION

AEMT

Paramedic



Subject: Childbirth

Effective: June 1, 2021

Last modified: Jan. 18, 2025

4006.1 General Guidelines

- a. Obtain history of patient condition and pregnancy, including:
 - i. Contraction duration and interval
 - ii. Gestation age should be expressed in weeks whenever possible
 - iii. Due date
 - iv. First day of last menstrual period
 - v. Number of pregnancies and number of live births (gravida/para)
 - vi. Presence or absence of prenatal care
 - vii. Possibility of multiple births
 - viii. Any possible complications
 - ix. Any drug use by the mother
- b. The patient should be transported to a hospital with obstetrical capabilities
 - i. Unless delivery is imminent, (the baby is crowning during a contraction).
 - ii. **ABSOLUTELY NO PREGNANT PATIENTS TO DAYTON or CINCINNATI CHILDREN'S HOSPITALS.**
- c. Visualize the perineal area only when contractions are less than five minutes apart.
- d. Run reports must be completed for each patient. The newborn is a separate patient from the mother.

4006.2 Clinical Management

Assessment		
Pediatric Considerations Manage childbirth scenarios with patients under 16 years old in the same manner as an adult.	Signs & Symptoms - Spasmodic pain - Vaginal discharge or bleeding - Lengthening and narrowing contractions - Urge to push - Crowning	Differential Diagnosis - Abnormal presentations (foot, and, buttocks) - Prolapsed cord - Placenta previa - Abruptio placenta
Treatment Algorithm		
- The EMR may only assist with emergency childbirth management - Apply gentle pressure on the baby's head with a flat hand to prevent an explosive delivery. - Place a gloved hand inside the birth canal only in the case of: - Breech delivery with entrapped head - Prolapsed umbilical cord limiting fetal circulation - Keep the newborn warm with blankets and (if available) a head cover. - Cut the umbilical cord and then place the baby to suckle at the mother's breast. - Obtain one, five and ten minute APGAR scores if time and patient condition permit. (see table below) - Establish an IV for patients in active labor. - No additional orders at this level.		
Consult		
None		
Clinical Pearls		
- When transporting potentially complicated deliveries or emergent childbirths, consider transporting to the closest L & D capable facility - Changes in fundal (upper part of the uterus) height during pregnancy: Above the symphysis pubis = Greater than 12 to 16 weeks gestation		

APGAR Score	0	1	2
Appearance	Full body cyanosis	Cyanosis at the extremities	No cyanosis present
Pulse	Absent	Slow (less than 100)	Greater than 100
Grimace	Flaccid	Grimace with stimulation	Cough or sneeze with stimulation
Activity	Absent	Some flexion of extremities	Active motion
Respiratory Effort	Absent	Slow or irregular	Good, vigorous cry

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Subject: Childbirth with Complications

Effective: June 1, 2021

Last modified: Feb. 21, 2025

4007.1 General Guidelines

- a. With all complicated childbirth scenarios, evaluate the need for rapid transport to a birthing center or possibly, the nearest hospital.
- b. These guidelines apply to all levels of certification, except where noted.
- c. In all complicated childbirth scenarios, place the mother on oxygen by non-rebreather mask.

4007.2 Clinical Management

- a. Cord around Baby's Neck:
 - i. As baby's head passes out of the vaginal opening, feel for the cord.
 - ii. Initially try to slip cord over baby's head.
 - iii. If too tight, clamp cord in two places and cut between clamps.
- b. Breech Delivery:
 - i. When an appendage or buttocks first becomes visible, position patient to discourage delivery, coach patient to avoid pushing and transport patient immediately.
 - ii. If the delivery is in progress, take care to support the baby's body.
 - iii. If the head is caught in the birth canal:
 1. Apply gentle pressure above the pubis symphysis as the mother pushes.
 2. If the head will not deliver, you must create an airway for the baby.
 3. Support the body and insert two fingers into the birth canal, forming a "V" around the mouth and nose.
- c. Prolapsed Cord:
 - i. When the umbilical cord is exposed prior to delivery, check cord for pulse.
 - ii. Transport immediately with hips elevated and a moist dressing around cord.
 - iii. Insert two fingers into the birth canal to displace the presenting part away from cord, distribute pressure evenly if occiput presents.
 - iv. Do not attempt to reinsert cord.
- d. Excessive Bleeding:
 - i. Treat for shock.
 - ii. Post-delivery, massage uterus firmly and put baby to mother's breast.
 - iii. ♦ The AEMT and Paramedic may consider the use of **Tranexamic Acid (TXA)** 2 grams IV/IO over **1-2 minutes** as outlined in [3015 Hemorrhage Control](#)

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Subject: Diabetic Emergencies –
Hypoglycemia/Hyperglycemia

Effective: June 1, 2021

Last modified: July 1, 2026

4008.1 General Guidelines

- Hypoglycemia is defined as a blood glucose level less than 60 mg/dL, or there is strong suspicion of hypoglycemia despite glucometer readings
- Hyperglycemia is defined as a blood glucose level at or above 250 mg/dL. EMS will treat patients with a BGL greater than 400 mg/dL or when the glucometer reads “High”

4008.2 Clinical Management

Assessment			
Pediatric Considerations None	Signs & Symptoms (Hypo) - Altered LOC - Dizziness - Irritability - Diaphoresis - Seizures - Hunger - Confusion - Acute onset	Signs & Symptoms (Hyper) - Altered LOC - Malaise - Hypotension - Dehydration - Polydipsia - Muscle cramps - Nausea - Delayed onset	Differential Diagnosis - Alcohol related issues - Toxic overdose - Trauma - Seizure - Syncope - CNS disorder - Stroke or TIA - Pre-existing condition
Treatment Algorithm			
- Provide basic care. - Call for transport.			EMR
- Obtain blood sample via finger stick and measure blood glucose level - If Hypoglycemic: - Administer 1 tube of Oral Glucose. - Maintain normothermia. Unconscious diabetics are often hypothermic. - In a diabetic patient with an insulin pump and blood glucose less than 60 mg/dL, treat the hypoglycemia. - If Hyperglycemic: - Monitor and transport			EMT
- If Hypoglycemic: A Administer Dextrose 10% (D10), 250 ml at wide open rate, (250 ml = 25 g of Dextrose) P Administer Dextrose 10% (D10) 5 ml/kg, maximum single dose of 250 ml. P For newborn, Dextrose 10% (D10) 2 ml/kg if BGL is less than 40 mg/dL. Dextrose 10% (D10) may be repeated in ten minutes if blood sugar remains less than 60 mg/dL. - If Hyperglycemic: A If BGL reads over 400 mg/dL or “High” on glucometer, administer 500 ml fluid IV – wide open. P Do <u>not</u> administer fluid to a hyperglycemic pediatric patient, unless otherwise indicated.			AEMT
A If an excited delirium patient goes into arrest: ♦ Consider Sodium Bicarbonate 100 mEq IV			Paramedic
Consult			
None			
Clinical Pearls			
- Oral glucose is indicated for any conscious but disoriented patient with BGL less than 60 mg/dL, or in a strong suspicion of hypoglycemia despite BGL readings. - Oral glucose may be administered carefully under the tongue or between the gum and cheek of an unresponsive patient who then must be placed in the lateral recumbent position to promote drainage of secretions away from the airway. - If the AEMT or Paramedic cannot obtain an IV in a hypoglycemic patient, an intraosseous (IO) access is an option. Transport is recommended. - When documenting the administration of Dextrose 10% (D10), do so in terms of milliliters. - Insulin Pumps - For a diabetic patient with an insulin pump who is hypoglycemic, treat the hypoglycemia. Do not disconnect or turn off pump. - Take extra tubing and medication reservoir or vials to the receiving facility for patients with insulin pumps.			
END OF SECTION			

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Subject: Diabetic Emergencies –
Refusal of Transport

Effective: June 1, 2021

Last modified: Dec. 7, 2024

4009.1 General Guidelines

- a. EMTs and above may allow for diabetic patients to refuse transport after treatment.
- b. EMRs should call for transport or a provider of a higher-level certification.

4009.2 Refusal Procedures

- a. Patients 18 years of age or older may be permitted to refuse. Follow these guidelines:
 - i. Repeat physical examination and vital signs.
 - ii. Patient must be Alert & Oriented x 3.
 - iii. Warn the patient that there is a significant risk of going back into a hypoglycemic state, especially if on oral hypoglycemics.
 - iv. Advise the patient to eat something substantial immediately.
 - v. Advise the patient to contact their family physician as soon as possible to minimize future episodes.
 - vi. Advise the patient to stay with someone.
 - vii. Follow normal patient refusal procedures.
- b. If the diabetic patient is under 18, but a parent or guardian is present, then the responsible adult may refuse patient transportation under the same guidelines as listed above in 4009.2.a.

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Subject: Extrapyramidal (Dystonic) Reactions

Effective: June 1, 2021

Last modified: Dec. 8, 2021

4010.1 General Guidelines

- a. A patient who is currently on a phenothiazine (e.g., Phenergan, Thorazine, Compazine) or a butyrophenone (e.g., Haldol, Droperidol) and exhibiting signs of acute muscle spasm or motor restlessness may be suffering from an Extrapyramidal Reaction.
- b. Extrapyramidal reactions can occur with ingestion of recreational drugs
- c. Physical examination findings may include any of the following:
 - i. Oculogyric crisis (spasmodic deviation of eyes in all directions generally fixed upward.)
 - ii. Buccolingual crisis (protrusion of tongue with slurred speech)
 - iii. Trismus (closing of the jaw due to spasm of the muscles also called lockjaw.)
 - iv. Difficulty in speaking
 - v. Facial grimacing
 - vi. Torticollis crisis (stiff neck with deviation of the head with the chin pointing to the other side)
 - vii. Opisthotonus (extreme back arching)
 - viii. Tortipelvic crisis—Involves hip, pelvis, and abdominal wall muscles, causes difficulty walking.
 - ix. Mental status is unaffected.
 - x. Vital signs are usually normal.
 - xi. Remaining physical examination findings are normal.

4010.2 Clinical Management

Assessment							
Pediatric Considerations None	Signs & Symptoms As listed above	Differential Diagnosis - Alcohol intoxication - Toxin/substance abuse - Medication effect - Withdrawal syndromes - Anxiety disorders - Mental health history					
Treatment Algorithm							
- Provide basic care. - Call for transport.				EMR	EMT	AEMT	Paramedic
If blood glucose less than 60, or there is strong suspicion of hypoglycemia despite glucometer readings, then follow 4008 Diabetic Emergencies - Hypoglycemia protocol							
Initiate IV fluid to maintain adequate BP. A ♦ Diphenhydramine 50 mg IV or IM P ♦ Diphenhydramine 1 mg/kg IV or IM (max dose 50 mg)							
Paramedics do not need a MCP order to administer Diphenhydramine.							
Consult							
The AEMT needs orders for Diphenhydramine							
Clinical Pearls							
None							
END OF SECTION							

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Subject:

Obstetrical Emergencies

Effective:

June 1, 2021

Last modified:

May 15, 2026

4011.1 General Guidelines

- a. Consider the possibility of ectopic pregnancy in females of child-bearing age.
- b. Ask for first day of last menstrual period.
- c. Gestational age should be expressed in weeks whenever possible.
- d. Aggressively treat for hypovolemic shock (do not rely on standard vital sign parameters).
- e. Give psychological support to patient and family.
- f. Be sure to take all expelled tissue with you to the hospital.

4011.2 Transport Decisions

- a. Transport to Maternity Department if:
 - i. Pregnant patients, 20 weeks, or greater gestation with obstetric complaints.
 - ii. If unsure of time of gestation, then consider transport to a maternity department.
 - iii. Patient experiencing obstetric related issues up to 12 weeks postpartum.
- b. Transport to Emergency Department if:
 - i. Pregnant patients with minor trauma or medical (non-obstetric) complaints
 - ii. Pregnant patients less than 20 weeks gestation
- c. Pregnant trauma patients should be rapidly transported to the Emergency Department at an Adult Trauma Center with labor and delivery capabilities.
- d. Positional transport considerations:
 - i. Prepare for postural hypotension caused by fetus pressure on venous return.
 - ii. Passively or actively move the fetus off the vena cava by doing either/or:
 1. Place in left lateral recumbent position or place a pillow under the right hip/lower back.
 2. Apply continuous manual displacement of the uterus towards the patient's left side.

4011.3 Clinical Management

- a. Causes of cardiac arrest in pregnant patients can include:
 - i. Pulmonary embolism
 - ii. Trauma
 - iii. Hemorrhage
 - iv. Congenital or acquired cardiac disease.
- b. Load and go to the closest hospital and follow all cardiac arrest protocols enroute.

4011.4 Aspirin and the Pregnant Patient

- a. Aspirin is contraindicated in third trimester due to possibility of pre and post-partum hemorrhaging and protentional fetal harm.

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Subject: Overdose/Poisonings

Effective: June 1, 2021

Last modified: Mar. 9, 2025

4012.1 General Guidelines

- a. ♦ EMS personnel should contact and confer with MCP for direction on suspected poisonings.
- b. Poison Control is intended for use by the general public.
- c. If possible, provide receiving facility all available information about the substance:
 - i. Safety data sheets (SDS)
 - ii. The container (if it is safe to do so)
 - iii. The label or an image of the label and warning information if it is unsafe or impractical to transport the actual substance container
- d. See Clinical Pearls for recommendations on treatment for button battery ingestion.

4012.2 Clinical Management

Assessment				
Pediatric Considerations - Most pediatric patients with respiratory depression do not have narcotic overdose. They are either septic or have respiratory failure. - Treatment for button battery ingestion is for patients greater to or equal to 12 months of age.	Signs & Symptoms - Mental status changes - Hypo/hypertension - Decreased respiratory rate - Tachycardia or bradycardia - Cardiac dysrhythmias - Seizures	Differential Diagnosis - Respiratory depression - Insecticides (organophosphates) - Solvents, cleaning agents - Cardiac medications - Stimulants - Depressants		
Treatment Algorithm				
- If respirations are impaired or there is suspicion of narcotic overdose: - A Administer Naloxone, up to 4 mg IN (half dose per nostril) - A May repeat Naloxone doses in 2 minutes - P Naloxone: - Less than or equal to 20 kg then 0.1 mg/kg IN, (max dose 2 mg), may repeat x one - Greater than 20 kg 2 mg, IN, may repeat as needed - Titrate Naloxone to adequate respirations. - Consider patient restraint before administration of Naloxone.			EMR	
No additional orders at this level			EMT	
- If patient has a pulse, Naloxone should be administered before inserting an ETT. - When given IV or IN, the onset of action for Naloxone is approximately 2 minutes. - If respirations are impaired or there is suspicion of narcotic overdose: - A Administer Naloxone, up to 4 mg IN, 2mg IV or 4 mg IM - A May repeat Naloxone doses in 2 minutes. - A Consider repeat IV dosing if no or inadequate response is noted - P Administer Naloxone: - Less than or equal to 20 kg then 0.1 mg/kg IN, IV, IM (max dose 2 mg), may repeat x one - Greater than 20 kg 2 mg, IN, IV, IM, may repeat as needed - Naloxone slow IV is preferred, but it may be given IN or IM before IV is established. - Titrate to adequate respirations. - If using IN route and respirations don't improve after 2 minutes, establish IV and administer IV dose. <u>Stimulant Overdose</u> (cocaine, methamphetamines, amphetamines, crack cocaine) with chest pain: - A Nitroglycerin 0.4 mg SL, if SBP >100, every 5 minutes to a total of three doses with vital signs between doses - A Midazolam 10 mg, IN (5 mg in each nostril) <i>or</i> 2.5 mg slow IV, <i>or</i> 5 mg IM - A Repeat Midazolam 5 mg IN (2.5 mg in each nostril) <i>or</i> 2.5 mg slow IV <i>or</i> 5 mg IM for unrelieved chest pain - G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses.				AEMT



Subject: Overdose/Poisonings

Effective: June 1, 2021

Last modified: Mar. 9, 2025

- Tricyclic Antidepressant Overdose may be evidenced by bradycardia, tachycardia, hypotension and prolongation of the QRS complex. Risk of rapid deterioration or sudden onset V Fib is high.

A ♦ **Sodium Bicarbonate 100 mEq, slow IV**

P ♦ **Sodium Bicarbonate 1 mEq/kg slow IV**

A ♦ Repeat **Sodium Bicarbonate 50 mEq, slow IV** for persistent QRS prolongation

P ♦ Repeat **Sodium Bicarbonate 0.5 mEq/kg slow IV** for persistent QRS prolongation

- Calcium Channel Blocker Overdose:

A ♦ **Calcium Chloride, 1 Gm slow IV**

P ♦ **Calcium Chloride, 0.2 ml/kg (20 mg/kg) slow IV (max dose 500 mg)**

Paramedic

Consult

- For guidance on suspected poisonings contact MCP.
- Calcium Channel Blocker, Beta Blocker and Tricyclic antidotes in this protocol are by MCP order only.

Clinical Pearls

- Consider other causes of altered mental status such as hypoglycemia, head trauma, sepsis, and stroke.
- When Naloxone is given intranasal (IN), the onset of action is approximately 2 minutes.
- Naloxone is not felt to be effective in the reversal of cardiac arrest from opioid overdose.
 - Airway control, ventilation, and quality CPR are still the mainstay of treatment in cardiac arrest management.
- Naloxone is not indicated in the management of newborns as it can cause withdrawal and secondary issues.
- Ondansetron (Zofran) is NOT to be given prophylactically with Naloxone.
- For ingestion of lithium button batteries - Though not in EMS scope of practice, management of ingested button batteries includes administering pure honey to coat the battery and delay tissue damage.
 - If there is honey at the scene and the patient or patient's caregivers wish to do so, they may administer the treatment.
 - The patient should consume approximately 10 ml every 10 minutes, and may do so up to 6 times.
 - **This is contraindicated in patients less than 12 months old.**
 - This should be documented as "self or family administered", not administered by the provider.
- Tricyclic Antidepressant Examples:
 - Amitriptyline (Elavil, Endep, Etrafon, Limbitrol)
 - Nortriptyline (Pamelor, Aventyl)
 - Amoxapine (Asendin)
 - Clomipramine (Anafranil)
 - Desipramine (Norpramine)
 - Doxepin (Sinequan)
 - Imipramine (Tofranil)
 - Protriptyline (Vivactil)
 - Trimipramine (Surmontil)
- Calcium Channel Blocker examples:
 - Amlodipine (Norvasc)
 - Diltiazem (Cardizem, Dilacor)
 - Felodipine (Plendil)
 - Isradipine (Dynacirc)
 - Nifedipine (Procardia, Adalat)
 - Verapamil (Calan, Isoptin, Verelan)
- Beta Blocker examples
 - Acebutolol (Sectral)
 - Atenolol (Tenormin)
 - Carvedilol (Coreg)
 - Corzide, Inderide, Lopressor, HCT, Tenoretic, Timolide, Ziac
 - Labetalol (Normodyne, Trandate)
 - Metoprolol (Topral, Lopressor)
 - Nadolol (Corgard)
 - Pindolol (Viskin)
 - Propranolol (Inderal)
 - Sotalol (Betapace)
 - Timolol (Blocadren)

END OF SECTION



Subject: Respiratory Distress/Pulmonary Edema

Effective: June 1, 2021

Last modified: May 16, 2026

4013.1 General Guidelines

Assessment				
Pediatric Considerations None	Signs & Symptoms - Cyanosis - Clammy skin - Presence/Absence of fever - Coughing - Wheezing - Labored breathing - Diaphoresis - Pitting edema - Bilateral lower lobe rales - Tachypnea - Apprehension - Jugular vein distension (JVD) - Inability to talk.	Differential Diagnosis - Myocardial infarction - Congestive heart failure - Asthma - Anaphylaxis - Aspiration - Chronic obstructive pulmonary disease - Pleural effusion - Pneumonia - Pulmonary embolus - Pericardial tamponade		
		Treatment Algorithm		
- Evaluate breath sounds. - Obtain pulse oximetry reading. - Obtain capnography reading. - Provide high flow O₂. - Call for transport.		EMR		Paramedic
{Obtain and transmit 12 Lead EKG} A If Pulmonary Edema, then Continuous Positive Pressure Airway (CPAP)		EMT		
- If Pulmonary Edema: A CPAP use is encouraged prior to the initiation of drug therapy. A If patient has SBP greater than 100, Nitroglycerin 0.4 mg SL up to 3, 1 every 5 minutes.		AEMT		
- Cardiac monitoring - If Pulmonary Edema: - CPAP or {Bi-PAP} use is encouraged prior to the initiation of drug therapy. - Consider need for possible early endotracheal intubation.				
Consult				
None				
Clinical Pearls				
- Evaluate breath sounds: <u>Clear</u>: treat cause (e.g. MI, pulmonary embolism, metabolic disturbance, and hyperventilation). <u>Wheezes</u>: treat cause (e.g. pulmonary edema, FBAO, asthma, allergic reaction). <u>Rales</u>: treat cause (e.g. pulmonary edema, pneumonia). - Diminished or absent: <u>Unilateral</u>: treat cause (e.g., pneumothorax, hemothorax, pneumonia, surgically removed lung). <u>Bilateral</u>: treat cause (e.g., respiratory failure, COPD, asthma). - Pneumonia may look like CHF with pulmonary edema. However, the pneumonia patient is often dehydrated and has an elevated temperature. - Where indicated, consider administering Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized simultaneously inline with CPAP				
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Subject: Seizures

Effective: June 1, 2021

Last modified: May 16, 2026

4014.1 Clinical Management

Assessment			
Pediatric Considerations None	Signs & Symptoms - Decreased mental status - Sleepiness - Incontinence - Observed seizure activity - Evidence of trauma	Differential Diagnosis - Head trauma - Tumor - Metabolic, hepatic or renal failure - Hypoxia - Electrolyte abnormality - Drugs, medications - Infection/fever - Alcohol withdrawal - Eclampsia - Stroke/TIA - Hyperthermia - Psychogenic Non-epileptic Seizures	
Treatment Algorithm			
- BVM and nasopharyngeal airway during seizure as needed. - Maintain normothermia. - Obtain Pulse Oximeter and {Capnography} reading.			EMR
- If glucose less than 60, or there is strong suspicion of hypoglycemia despite glucometer readings, then follow 4008 Hypoglycemia/Hyperglycemia. - Place patient in the recovery position during assessment and transport.			EMT
- Cardiac monitor - For actively seizing adult patients: G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses A Midazolam 10 mg IN (5 mg in each nostril), <i>or</i> Midazolam 2.5 mg slow IV, <i>or</i> Midazolam 5 mg IM - If still seizing, repeat Midazolam doses: A Repeat Midazolam 5 mg IN (2.5 mg in each nostril) after 10 minutes. A <i>Or</i> repeat Midazolam 2.5 mg slow IV after 5 minutes. A <i>Or</i> repeat Midazolam 5 mg IM after 10 minutes. - In Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam: A Repeat any of the Midazolam doses/routes above A <i>Or</i> consider Ketamine 100 mg IV/IO or (if no IV/IO), then 250 mg IM - For actively seizing pediatric patients: P Midazolam 0.2 mg/kg IN (max IN dose 10 mg) <i>or</i> Midazolam 0.1 mg/kg slow IV (max IV dose 2.5 mg) <i>or</i> Midazolam 0.2 mg/kg IM (max IM dose 5 mg) - If still seizing, repeat Midazolam doses: P Repeat Midazolam 0.2 mg/kg IN (max IN dose 5 mg) after 10 minutes P <i>Or</i> repeat Midazolam 0.1 mg/kg slow IV (max IV dose 2.5 mg) after 5 minutes P <i>Or</i> repeat Midazolam 0.2 mg/kg IM (max IM dose 5 mg) after 10 minutes - In Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam: P Repeat any of the Pediatric Midazolam doses/routes above			AEMT



Subject: Seizures

Effective: June 1, 2021

Last modified: May 16, 2026

A If seizing pregnant patient with gestation greater than 20 weeks or postpartum less than 12 weeks, administer **Magnesium Sulfate 4 grams** (whole bag) infused with macro-drip tubing over **20 min**.

P If seizing pregnant or postpartum patient is less than 16 years old but meets criteria above, then administer **Magnesium Sulfate 4 grams** (whole bag) infused with macro-drip tubing over **20 min**

Paramedic

Consult

- None

Clinical Pearls

- When obtaining history be sure to include the following:
 - Description of seizures, areas of body involved, and duration
 - Other known medical history (e.g., head injury, diabetes, drugs, alcohol, stroke, heart disease, recent fever or illness, possible toxicological agents)
- Refractory Status Epilepticus (RSE) is defined as seizures that continue despite treatment with first-line medications. Post-medication signs and symptoms could include:
 - Generalized Tonic/Clonic seizures lasting longer than 5 minutes
 - Focal motor seizures lasting longer than 10 minutes
 - Absences lasting longer than 10 minutes
 - Recurrent seizures without a lucid interval

END OF SECTION



Subject: Sepsis

Effective: June 1, 2021

Last modified: Feb. 18, 2024

4015.1 General Guidelines

- Severe sepsis is characterized by poor perfusion, leading to a buildup of serum lactate and resulting metabolic acidosis.
- To compensate for metabolic acidosis, patients increase their minute ventilation.
- This increased respiratory rate “blows off” carbon dioxide and lowers EtCO₂.
- EtCO₂ levels decline in the setting of both poor perfusion and metabolic acidosis.
- Poor tissue perfusion decreases the amount of blood flow to the alveoli of the lungs, reducing the amount of carbon dioxide that can be exhaled
- Sepsis is often associated with a high mortality rate. The key to improve patient outcomes in septic shock is early recognition, fluid resuscitation, O₂ therapy and rapid transport.

4015.2 Clinical Management

Assessment				
Pediatric Considerations None	Signs & Symptoms - Known or suspected infection - EtCO₂ less than 32 or greater than 47 with 2 or more of the following criteria: - Respiratory rate greater than or equal to 22 - Altered mental status (GCS less than 13) - Temperature over 100.4 (38 C) or under 96.8 (36 C) - Heart rate greater than 90 - Systolic BP less than 100 or Mean Arterial Pressure (MAP) below 65			Differential Diagnosis - Fever - Flu-like symptoms
	Treatment Algorithm			
- Administer oxygen - Call for transport immediately.				EMR
If possible, obtain blood sample via finger stick and measure blood glucose level				EMT
- Administer a bolus of 1 liter of IV fluid. ♦ For additional fluid administration.				AEMT
A Consider Norepinephrine by adding 4 mg to 250 ml of IV fluids. Infuse starting at 30 drops per minute (max 45 drops) with 60 drop tubing and titrate to effect. Increase by 5 drops/minute every 5 minutes.				Paramedic
Consult				
Consult with MCP to give more than 1 liter of fluids.				
Clinical Pearls				
- Mean Arterial Pressure (MAP) can also predict the organ perfusion pressure. - MAP = (SBP + 2 X DBP) / 3 and is normally 70 – 110 mm/hg. - Patients may be in septic shock with a normal blood pressure. CAUTION: Be especially suspicious of sepsis in geriatric patients with altered mental status - Consider calling the receiving facility ahead to advise ED staff of potentially septic patients.				
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Subject: Shock

Effective: June 1, 2021

Last modified: Mar. 20, 2024

4016.1 General Guidelines

- Shock is inadequate tissue perfusion.
- Be proactive in treatment of shock. Do not wait for symptoms to present.
- Management of shock should include trying to find and correct the underlying cause (if possible).

4016.2 Clinical Management

Assessment		
Pediatric Considerations - Pediatric patients will compensate longer than adults. - Apparent signs and symptoms of shock can indicate a critical patient.	Signs & Symptoms - Restlessness, confusion - Weakness and dizziness - Tachycardia - Tachypnea - Hypotension - Decreased mentation - Pale, cool, clammy skin	Differential Diagnosis - Hypovolemia - Cardiogenic - Septic - Neurogenic - Anaphylactic - Pulmonary emboli - Tension pneumothorax - Medications or overdose - Vasovagal hypotension
Treatment Algorithm		
- Call for transport immediately. - Provide O₂ as appropriate - Keep patient warm. - Control external bleeding and treat for hypovolemic shock as indicated.		EMR
Transport immediately unless ALS intercept is less than 5 minutes.		EMT
- Only give fluids for specific signs and symptoms of shock and not to every trauma patient. - For persistent shock, establish additional vascular access. <u>Non-traumatic shock without Pulmonary Edema:</u> <i>Patient does not have JVD, edema, or rales.</i> - A IV fluid 500 ml IV. Maintain adequate perfusion. - P IV fluid 20 ml/kg IV. - P Titrate to maintain adequate perfusion. - A Additional IV fluid 500 ml IV, if needed. - P ♦ Additional IV fluid 20 ml/kg IV, if needed. <u>Non-traumatic shock with Pulmonary Edema:</u> <i>Patient may have JVD, edema, or rales present.</i> - A Consider IV fluid 250 ml IV. <u>Exsanguinating Hemorrhage:</u> - A IV fluid to maintain approximately 100 SBP or a radial pulse. Do not allow blood pressure to get too high. - P IV fluid 20 ml/kg IV. May repeat x 2. Titrate to maintain adequate perfusion.		AEMT
- For non-traumatic shock: - A Treat arrhythmias as indicated. - A If SBP remains less than 100, begin Norepinephrine by adding 4 mg to 250 ml of IV fluids. Infuse starting at 30 drops per minute (max 45 drops) with 60 drop tubing and titrate to effect. Increase by 5 drops/minute every 5 minutes.		Paramedic
Consult		
None		
Clinical Pearls		
Perform manual BP on all patients presenting with signs and symptoms of shock.		
END OF SECTION		

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Subject: Stroke

Effective: June 1, 2021

Last modified: Jan. 21, 2024

4017.1 General Guidelines

- a. If one or more signs of the Cincinnati Prehospital Stroke Scale (CPSS) are abnormal, and less than 24 hours since patient was last seen normal, call a "Stroke Alert", and transport to the closest appropriate Stroke Center.
- b. In addition to the CPSS, providers should screen patients for possible large vessel occlusions (LVO) before making transport destination decisions.
- c. If greater than 24 hours since last known well, consider transport to a Comprehensive or Thrombectomy Capable facility.
- d. When reporting last known well, state actual clock time. Do not say, "20 minutes ago."
- e. With such a diverse group of agencies and receiving hospital capabilities covered by this protocol, all agencies should discuss "best practices" for assessment, management, and transport of possible strokes with their Medical Directors. With approval, agencies may deviate from this guideline in the following manners:
 - i. Agencies may use alternative stroke screening scales (RACE, MEND, LAPSS, etc.) for evaluation of possible CVAs. All screening tools should include the routine assessments found in the CPSS and methods to screen for large vessel occlusions.
 - ii. Agencies may make transport destination decisions based on their proximity to stroke management facilities and the capabilities of those hospitals.
 - iii. All modifications to this protocol should be made in the form of a supplemental guideline specifically approved and signed off by the medical director.

4017.2 Clinical Management

Assessment						
Pediatric Considerations None	Signs & Symptoms - Facial drooping - Arm drift or weakness - Slurred or difficult speech - Aphasia (expressive or receptive) - Pupillary changes (in hemorrhagic strokes) - Gaze deviation/abnormal eye movement (indicative of large vessel occlusions)	Differential Diagnosis - Seizure - Subdural hematoma - Brain tumor - Syncope - Toxic or metabolic disorders (e.g., hypoglycemia) - Migraine headaches				
Treatment Algorithm						
- Perform a Cincinnati Pre-hospital Stroke Scale {or alternative approved by Medical Direction} assessment. - A patient in respiratory distress with pale, moist skin and altered mental status should get oxygen via NRB mask. - Be prepared to assist ventilations with OPA/NPA and Bag-valve-mask. - If signs of cerebral herniation are present, ventilate at the following rates: - A Approximately 20 times per minute. P Ventilate at a rate of ten faster than normal respiratory rate if the signs of cerebral herniation are present. {If numeric EtCO₂ readings are available, ventilate at a rate to maintain readings at approximately 30 mmHg (30 torr)} - Never ventilate at less than 8 per minute. - A patient with indications of stroke with a SpO₂ less than 94%, should be given oxygen via NC and titrated to 94%. - A patient with indications of stroke with a SpO₂ greater than 94%, should not get any oxygen.			EMR			



Subject: Stroke

Effective: June 1, 2021

Last modified: Jan. 21, 2024

- The presence of a single abnormal finding in the CPSS {or alternative screening approved by Medical Direction} should dictate a stroke alert and transport to the closest stroke center (unless contraindicated by greater than 24-hour onset, presence of LVO indicators or thrombolytic considerations).
- Perform a Large Vessel Occlusion (LVO) screening looking for:
 - Difficulty in balance or gait
 - Eye deviation – eyes may only move to one side, or be forced to one side
 - Visual disturbances – field of view cut, double vision, new onset blindness
 - Aphasia – expressive (inability to speak or paraphasic errors) or receptive (not understanding or following commands)
 - Denial/Neglect – can a patient feel you touch both of their arms and do they recognize their own hand?
- Consider the following contradictions to thrombolytics:
 - Neurosurgery, head trauma or stroke in the last 3 months
 - Major surgery or serious non-head trauma in the previous 14 days
 - History of gastrointestinal or urinary tract hemorrhage within 21 days
 - Current (within the last 48 hours) use of anticoagulants. Examples include:
 - Warfarin (Coumadin, Jantoven)
 - Edoxaban (Savaysa)
 - Apixiban (Eliquis)
 - Rivaroxaban (Xarelto)
 - Abigatran (Pradaxa)
 - Lovenox injections
- Patients with onset greater than 24 hours, clinical findings indicative of LVO or with contraindications to thrombolytics consider transport to a Comprehensive Stroke Center or Thrombectomy Capable Facility.
 - Patients with signs or symptoms that strongly indicate a possible hemorrhagic stroke should not be transported to a Thrombectomy Capable Facility
- Transport the patient with the bed flat, if able to tolerate. If showing signs of increased ICP, do not lay patient flat.
- If glucose is less than 60, or there is strong suspicion of hypoglycemia despite glucometer readings, then follow [4008 Diabetic Emergencies - Hypoglycemia](#) protocol.

EMT

AEMT

Paramedic

- No additional orders at this level

- No additional orders at this level

Consult

- Contact MCP for Stroke Alerts or for advice regarding transport destination, if not clear.

Clinical Pearls

- Cincinnati Prehospital Stroke Scale: (normal or abnormal)
 - Facial Droop (*patient shows teeth or smiles*).
 - Arm Drift (*patient closes eyes and holds both arms straight out for about 10 seconds*).
 - Abnormal Speech (*have patient say "You can't teach an old dog new tricks." or any other phrase*).
- Possible indicators of a large vessel occlusion (LVO):
 - The presence of abnormal findings in all three categories of the Cincinnati Prehospital Stroke Test increase the possibility of LVO
 - Visual neglect, gaze deviation, or abnormal eye movement are key clinical findings
 - New onset loss of balance or coordination may indicate a possible LVO stroke
- Arrange for transport a historian with patient both to provide patient history and for permission to treat.

4017.3 Stroke Centers

- Telemedicine Stroke Center:** Also known as drip and ship, has thrombolytic capabilities and immediate access to a Neurologist via telemedicine.
- Primary Stroke Center:** Facility with capability to administer thrombolytics and has an ICU.
- Comprehensive Stroke Centers or Thrombectomy Capable:** Facilities with 24/7 endovascular capabilities.
 - Miami Valley Hospital (Comprehensive)
 - Kettering (Comprehensive)
 - Mercy Health – Springfield (Thrombectomy Capable)

END OF SECTION



5000 Series

Pediatric Protocol

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Subject: Brief Resolved Unexplained Event

Effective: June 1, 2021

Last modified: Dec. 27, 2024

5001.1 General Guidelines

- a. A Brief Resolved Unexplained Event (BRUE) involves any infant under 1 year of age reported by a bystander as sudden, brief (less than 1 minute), unexplained, and completely resolved upon EMS arrival that includes one or more of the following:
 - i. Breathing change (absent, decreased, or irregular)
 - ii. Color change (central cyanosis or pallor)
 - iii. Change in muscle tone (Increase or decrease in muscle tone)
 - iv. Altered level of responsiveness (including irritability)
- b. Children who experience a BRUE event often have a normal exam on assessment.
A cause will be difficult to determine in most BRUE cases.

5001.2 Important Information to Gather

- a. Document the symptoms of the event given by the observer:
 - i. Was the child apneic, cyanotic, or limp during event?
 - ii. Infant's color, respirations, and muscle tone
 - iii. Was seizure-like activity noted?
 - iv. Was any resuscitation attempted or did event resolve spontaneously?
 - v. How long did the event last?
- b. Obtain past pertinent medical history:
 - i. Recent trauma, infection (e.g., fever, cough)
 - ii. History of gastroesophageal reflux (GERD)
 - iii. History of congenital heart disease
 - iv. History of seizures
 - v. Medication history
 - vi. Birth defects

5001.3 Clinical Management

- a. Support airway, breathing, circulation.
- b. Keep warm.
- c. Head-to-Toe exam for trauma, bruising, or skin lesions.
- d. Check anterior fontanel: is it bulging, flat or sunken?
- e. Pupillary exam.
- f. Respiratory exam for rate, pattern, work of breathing and lung sounds.
- g. Cardiovascular exam symmetry of brachial and femoral pulses.
- h. Neuro exam for level of consciousness.
- i. Observe for repetition of reported occurrences.
- j. The patient should be transported to the hospital for further assessment.

5001.4 Management and Transport of Febrile Pediatric Patients

- a. Transport all infants less than 2 months of age with a history or reported temperature of greater than 38.0 C (100.4 F) or less than 35.6 C (96.0 F).

END OF SECTION

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Subject: Newborn Care and Resuscitation

Effective: June 1, 2021

Last modified: Mar. 4, 2021

5002.1 General Guidelines

- a. Maintain airway. Place in the sniffing position (1" towel under shoulders).
- b. If drying and suctioning has not provided enough tactile stimulation, flick the infant's feet or rub the infant's back.
- c. Suction only infants in distress, until airway is clear of all secretions. Bulb suctioning is preferred.
- d. If meconium staining is present:
 - i. Newborn is vigorous, with strong respirations, good muscle tone, and heart rate greater than 100 BPM; monitor the patient and maintain a patent airway.
 - ii. Newborn is depressed, has poor respiratory effort, decreased muscle tone, or heart rate less than 100 BPM; clear the airway by suctioning before taking other resuscitative steps.
- e. Avoid direct application of cool oxygen to infant's facial area as may cause respiratory depression due to a strong mammalian dive reflex present immediately after birth.
- f. If stimulation does not improve the infant's breathing, then BVM assist may be necessary.

5002.2 Viable Fetus

- a. If the fetus is greater than 23 weeks gestation, follow normal resuscitative procedures.
- b. A fetus is viable if:
 - i. Eyelids not fused
 - ii. If measurable or known, must be greater 500 grams.

5002.3 Clinical Management

Assessment					
Pediatric Considerations Nothing additional	Signs & Symptoms - Respiratory distress - Central cyanosis - Altered level of consciousness - Bradycardia	Differential Diagnosis - Peripheral cyanosis (normal) - Infection - Maternal medication effect - Hypothermia, hypoglycemia, hypovolemia			
Treatment Algorithm					
- After delivery of the infant; - P Assess the airway and breathing. - P Warm, dry and stimulate - P Position head lower than body. - P Ventilate with BVM at 40-60/minute to increase HR (if less than 100) or for apnea or persistent central cyanosis. - P If heart rate is less than 60 bpm begin CPR. - P Compress at 120/min. - P Compression to Ventilation ratio of 3:1				EMR	Paramedic
Obtain APGAR scores at 1, 5 and 10 minutes post-delivery.				EMT	
- P If hypovolemic, IV fluid 10 ml/kg over 5-10 minutes. - P NEWBORN: D10 (2 ml/kg) if blood glucose less than 40.				AEMT	
- If heart rate remains less than 60 bpm after CPR: - P Epinephrine 1:10,000, 0.01 mg/kg IV - P If no response, repeat Epinephrine 1:10,000, 0.01 mg/kg IV, every 3-5 minutes.					
Consult					
Contact MCP for instructions and guidance when attempting to determine the viability of a fetus.					
Clinical Pearls					
- Use length-based resuscitation tape on all neonatal resuscitations. - Mechanical suction may be used on infants only if the suction pressure does not exceed 100 mmHg or 136 cmH₂O.					
END OF SECTION					

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Subject: Pediatric Assessment Triangle

Effective: June 1, 2021

Last modified: Dec. 8, 2020

5003.1 General Guidelines

- a. The Pediatric Assessment Triangle establishes a level of severity, assists in determining urgency for life support measures, and identifies key physiological problems using observational & listening skills.
- b. This assessment tool can be utilized by providers of all certification levels.

5003.2 Appearance

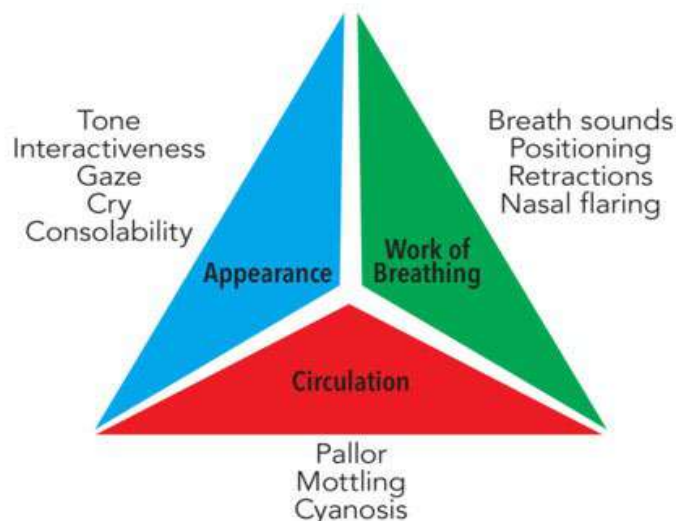
- a. Appearance reflects adequacy of: oxygenation ventilation, brain perfusion, CNS function.
 - i. The mnemonic used for pediatric assessment of appearance is: TICLS.
 1. Tone- Moves spontaneously, sits or stands (age appropriate)
 2. Interaction- Alert, interacts with environment
 3. Consolability- Stops crying with comfort measures (holding, warmth, distraction)
 4. Look/gaze – Makes eye contact with clinician, tracks objects
 5. Speech/cry – Uses age-appropriate speech or crying

5003.3 Work of Breathing

- a. WOB is a more accurate indicator of oxygenation and ventilation than respiratory rate or breath sounds.
- b. Assess for effort in breathing, accessory muscle use, and depth of breathing.
- c. Capillary refill is an accurate predictor of pediatric oxygenation.
- d. Under work of breathing, the patient should fall into one of four categories:
 - i. Normal Breathing
 - ii. Respiratory difficulty
 - iii. Respiratory failure
 - iv. Respiratory arrest

5003.4 Circulation

- a. Circulation reflects adequacy of cardiac output and perfusion of vital organs (core perfusion).
- b. Cyanosis reflects decreased oxygen levels in arterial blood, vasoconstriction, and respiratory failure.
- c. Mottling of the skin indicates hypoxemia, vasoconstriction, and respiratory failure.

5003.5 The Pediatric Assessment Triangle**END OF SECTION**

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Subject: Safe Harbor

Effective: June 1, 2021

Last modified: Dec. 8, 2020

5004.1 General Guidelines

- a. Safe Harbor is for the voluntary separation of newborn infant.
- b. It is designed to allow desperate parents to separate from their babies to hospitals, EMS, or law enforcement agencies, confidentially.

5004.2 Clinical Management

- a. Stipulations of separation:
 - i. Infant can be no older than be 30 days old.
 - ii. Infant can have no signs of abuse or neglect
- b. History which should be obtained:
 - i. Date and time of birth
 - ii. Any pertinent family medical history
 - iii. Information regarding prenatal care
 - iv. Information concerning the birth.
- c. Information should be obtained in a manner, which will not lead to the revealing of the identity of the parents.
- d. Information collected should be based on patient (infant) care needs and assure confidentiality.
- e. Transport the infant to the hospital.

END OF SECTION

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6000 Series

Special Operations Protocol

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Subject: General Management for Haz Mat

Effective: June 1, 2021

Last modified: Dec. 8, 2020

6001.1 General Guidelines

- a. This section will provide the responders with direction toward the management and mitigation of Hazardous Material events.
- b. The initial goal of any hazardous materials release is to isolate and identify.

6001.2 Initial Actions

- a. Personnel safety:
 - i. Consider potential for secondary devices
 - ii. Don appropriate PPE
 - iii. Stage personnel & equipment
- b. Call for additional resources. (Haz Mat Teams, Decon crews, Law Enforcement, etc.)
- c. Field decontamination:
 - i. Remove all contaminated clothing
 - ii. Thoroughly wash the patient with {Dawn} dishwashing detergents
 - iii. Pay special attention to skin folds and other areas where simple irrigation may not remove it
 - iv. If a patient has been contaminated with any fuel, irrigate well
- d. Contact Medical Control and the hospital immediately to allow time for their set-up of decontamination equipment.
 - i. Provide the following information:
 - 1. Estimated number of confirmed or potential adult and pediatric patients
 - 2. Signs and symptoms exhibited by the patients
 - 3. Name and identification information of the contaminant if known, or as much information as possible
 - 4. Form of the contaminant (liquid, gas, etc.) if known
 - 5. Routes of exposure of the patients (percutaneous, inhalation, ingestion, etc.) if known
 - 6. Additional anticipated decontamination needs if necessary.
 - ii. Obtain permission from hospital upon arrival before entering with a potentially contaminated patient or crew.
- e. ♦ In the event of an MCI involving cyanide or nerve agents, request an “Antidote free” order, allowing you to treat all of the patients on the scene with the appropriate antidote, rather than calling for patient orders individually.
- f. Do not transport a patient until gross decontamination is completed.
- g. Decontaminate EMS vehicles prior to leaving hospital.

END OF SECTION

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Subject: Antidote Resources

Effective: June 1, 2021

Last modified: Mar. 9, 2025

6002.1 Antidote Options

- a. {EMS Departments are authorized to stockpile **Atropine**, **2-PAM**, auto-injectors, and antidote delivery supplies at their own expense}
- b. Dayton MMRS Caches
 - i. Dayton MMRS stores additional supplies of cyanide antidotes in each county in Ohio Homeland Security Region 3.
 - ii. To obtain Dayton MMRS antidotes: call 937-333-USAR (8727).
 - iii. The closest department with an antidote cache will respond as a mutual aid request.
- c. CHEMPACK Resources:
 - i. Store of antidotes to treat about 500 victims of a nerve agent or organophosphate incident.
 - ii. EMS CHEMPACK contents:
 1. **Atropine**—blocks effects of excess acetylcholine
 - a. **0.5 mg AtroPen** auto-injectors (for patients less than 20 kgs)
 - b. **1.0 mg AtroPen** auto-injectors (for patients 20-40 kgs)
 - c. Multi-dose vials
 2. **Pralidoxime Chloride (2-PAM)**—reduces levels of acetylcholine
 - a. **600 mg** auto-injectors
 - b. Multi-dose vials
 3. **Midazolam** or **Diazepam (Valium)**—treats seizures.
 - a. Carpu-jects
 - b. Multi-dose vials
 4. Both EMS and Hospital CHEMPACKs contain the same three drugs.
 - iii. Hospital CHEMPACK contents
 1. More multi-dose vials for more precise dosing of children and long-term patients.
 2. Hospitals have the option to keep the materials for use at their hospital.
 3. If a hospital opens its CHEMPACK, it must notify OSP Central Dispatch.
 - iv. CHEMPACK Limitations
 1. Only useful against nerve agents or organophosphate
 2. Only to be utilized when other resources are inadequate for number of victims.
 3. CHEMPACKs opened contrary to guidelines will not be replaced by CDC.



Subject: Antidote Resources

Effective: June 1, 2021

Last modified: Mar. 9, 2025

v. CHEMPACK procurement:

1. ♦ Obtain MCP approval
2. ♦ Contact OSP Central Dispatch 866-599-LERP (5377) and request a CHEMPACK.
3. You must indicate that the scenario meets both of the following criteria:
 - a. The agent has been identified, or patients are exhibiting signs and symptoms of organophosphate/nerve agent exposure.

AND

 - b. The need for antidotes is greater than the available resources.
4. OSP Central Dispatch will:
 - a. Notify closest CHEMPACK hospital
 - b. Dispatch Troopers to deliver the CHEMPACK to the MCI's staging area.
 - c. Troopers will expect EMS to sign a form indicating receipt.

END OF SECTION



Subject: Hazardous Drug Exposure

Effective: June 1, 2021

Last modified: Oct. 10, 2021

6003.1 Identification or Recognition of a Hazardous Drug Situation

- a. Hazardous drug situations include:
 - i. Patients who have just had IV chemotherapy at the clinic or hospital
 - 1. Body fluids could have traces of hazardous drugs for up to 48 hours.
 - ii. Patients taking oral chemotherapy drugs.
 - iii. Patients who have continuous IV chemotherapy at home.
- b. Potential routes of exposure include:
 - i. Absorption through skin or mucous membranes
 - ii. Accidental injection by needle stick or contaminated sharps
 - iii. Inhalation of drug aerosols, dust, or droplets
 - iv. Ingestion through contaminated food, tobacco products, beverage, etc.
- c. Don PPE listed below whenever there is a risk of hazardous drug being released into the environment.
 - i. When handling leakage from tubing, syringe, and connection sites.
 - ii. When disposing of hazardous drugs or items contaminated by hazardous drugs.
 - iii. When handling the body fluids of a patient who received hazardous drugs in the past 48 hours.
 - iv. When cleaning hazardous drug spills

6003.2 Guidelines for Personal Protective Equipment:

- a. Gloves: two sets of nitrile gloves are recommended. Change gloves every 30 minutes.
- b. Disposable, non-permeable gowns
- c. NIOSH-approved respirator masks
- d. Eye and face protection: wear a face shield whenever there is a possibility of splashing.

6003.3 Procedures:

- a. Wipe up liquids with an absorbent pad or spill-control pillow.
- b. If necessary, consult with the appropriate Haz-Mat team.
- c. Dispose hazardous drugs or contaminated materials per MSDS or Haz Mat Team direction.
- d. Report and document spills as required.
- e. For accidental skin exposure: Remove contaminated garments, place in leak-proof plastic bag, and immediately wash contaminated skin with soap and water. Rinse thoroughly.
- f. For accidental eye exposure: immediately flush eye with saline solution or water for at least 30 minutes or until patient transport is completed.

6003.4 Identification or Clarification

- a. For more information about a hazardous drug or handling procedures, contact:
 - i. The homecare agency that is supplying the infusion.
 - ii. The physician who ordered the infusion.
 - iii. A hospital pharmacy, if necessary (there should be a label on the IV bag with the drug's name, concentration, and dosage).

END OF SECTION

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Subject: Hydrofluoric Acid Exposure

Effective: June 1, 2021

Last modified: Oct. 10, 2021

6004.1 Clinical Management

Assessment			
Pediatric Considerations None	Signs & Symptoms - Breathing difficulty - Abdominal pain - Chest pain - Burns (with blisters) - Stridor (if inhaled)	Differential Diagnosis Chemical burns	
Treatment Algorithm			
- Ensure the safety of all responders. - Begin decontamination and irrigate the chemical burn with water as quickly as possible. - Flush affected eyes and skin with copious amounts of water or IV Fluids for a minimum of 30 minutes. - Continue flush until patient transport is completed. - If ingested, do not induce vomiting. Dilute with water or milk. - Monitor for cardiac arrest.			EMR
{Perform a 12 Lead EKG and transmit it to the hospital}			EMT
- Intubate if apneic. - Consider 1014 Pain Management Protocol			AEMT
- When feasible, use {Epsom Salt solution} as an additional irrigating solution for affected skin. - A {Epsom Salt Solution} is not for eyes or mucous membranes. - A Getting water on the burn is more urgent than the use of Epsom salt. - A Do not delay irrigation or decontamination. - A If available, use {Epsom Salt solution} on the skin for at least 30 minutes. - If ingested, in addition to water or milk, give {3-4 ounces of magnesium-containing antacid (i.e., Maalox or Mylanta)}. - Intubate if unconscious or at first sign of pulmonary edema or respiratory distress. - Perform a 12-lead EKG and monitor for prolonged QT interval. - Apply {magnesium-containing antacid (Maalox or Mylanta)} topically to burned areas. - A Omit if topical agents have already been applied prior to arrival. ♦ If patient with HF exposure experiences tetany or cardiac arrest, administer Calcium Chloride 10% 1 g (10 ml) 10%, IV. - A Calcium Chloride 10% should be considered a first line drug in cardiac arrest associated with Hydrofluoric Acid. - A Only ABCs, defibrillation, intubation and Epinephrine should precede its administration. ♦ If patient was exposed to high concentration HF (greater than 40%), discuss prophylactic Calcium Chloride 10% 400 mg (4 ml), slow IV with MCP.			
Consult			
The paramedic should contact MCP for administration of Calcium Chloride 10%			
Clinical Pearls			
Death due to Hydrofluoric Acid has been reported from burns involving less than 3% body surface area.			
END OF SECTION			

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Subject: Organophosphate or Nerve Agent Exposure

Effective: June 1, 2021

Last modified: Mar. 9, 2025

6005.1 Clinical Management

Assessment		
Pediatric Considerations None	Signs & Symptoms - Salivation - Lacrimation - Urination - Defecation - Gastrointestinal Issues - Emesis - Miosis - Muscle Twitching	Differential Diagnosis None with a recent history of exposure to nerve agents
Treatment Algorithm		
- Administer oxygen ♦ Administer Atropine by DuoDote every 5 minutes, as available until the lungs are clear to auscultation. ♦ DuoDotes can be given to adult and pediatric over 40 kgs patients.		EMR
No additional orders at this level.		EMT
♦ Treat seizures with Midazolam or Diazepam		AEMT
G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses		
♦ Administer Atropine every 5 minutes (up to a total of three doses), as available until lungs are clear to auscultation. Atropine may be given IV, IM, IO or by AtroPen auto-injector for children, or by DuoDote. A ♦ Adults and children greater than 40 kgs, give DuoDote, or Atropine 2 mg, IV, IM. P ♦ Children 20 – 40 kg, give 1.0 mg Atropine, or the 1.0 mg Atropen auto-injector. P ♦ Children less than 20 kg, give 0.5 mg Atropine, or the 0.5 mg Atropen auto-injector. A ♦ Follow Atropine with 2-PAM (Pralidoxime) 600 mg IM. (each DuoDote contains 600 mg of 2-PAM) P ♦ Infants and young children should receive Pralidoxime, 25-50 mg/kg IV or IM, if available. - Treat seizures with Midazolam or Diazepam		Paramedic
Consult		
Contact MCP for administration of medications listed above.		
Clinical Pearls		
- Treat any case of known or suspected Organophosphate or Carbamate (e.g., insecticides such as Parathion or Malathion); or nerve agent (e.g., Tabun, Sarin, Soman, VX) exposure. - Mild to moderate cases should be treated with one or two doses of DuoDote. - Severe cases will generally require repeating every 5 minutes up to 3 doses. - Organophosphate poisonings may require more Atropine (3 DuoDotes). - Atropine in these circumstances is <u>not</u> for bradycardia, which may or may not be present. - Procedures for DuoDotes, pediatric AtroPens are the same as administering an Epi-Pen. - Primary endpoints for treatment are diminished airway secretions (lungs are clear to auscultation), hypoxia improves, airway resistance decreases, and dyspnea improves		
END OF SECTION		

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Subject: Other Hazardous Materials

Effective: June 1, 2021

Last modified: Mar. 15, 2023

6006.1 General Guidelines

- a. These guidelines are for the management of specific materials.
- b. Unless otherwise noted, these orders apply to all certification levels.

6006.2 Specific Materials

- a. Biological materials
 - i. ♦ {For the possibility of a bioterrorist attack, agencies may store their own supply of Ciprofloxacin (Cipro) or Doxycycline.}
 - ii. They can also provide prophylaxis against Anthrax, Cholera, and some protection against Plague.
- b. Pepper Spray
 - i. {**Sudecon Wipes**} can assist in the decontamination of patients or public safety personnel who have been sprayed with Pepper Spray.

END OF SECTION

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7000 Series

Administrative

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Subject: **Drug Bag Exchange Program:
General Operating Guidelines**

Effective: **June 1, 2021**

Last modified: **July 1, 2026**

7001.1 Drug Bag Exchange Committee Make-up

- a. Co-Chairpersons:
 - i. One Hospital EMS coordinator
 - ii. One Hospital pharmacy representative or one Greater Miami Valley EMS Council member
- b. Members:
 - i. EMS Coordinator from each participating hospital
 - ii. Pharmacy representative from each participating hospital
 - iii. Any interested GMVEMSC (Greater Miami Valley EMS Council) member
- c. Meetings
 - i. One scheduled meeting per year
 - ii. Unscheduled as needed to discuss problem areas

7001.2 General Operating Guidelines

- a. In order to participate in the GMVEMSC Drug Bag program, an agency must have the capability to communicate with Medical Control at participating hospitals.
- b. There are two types of drug bags: ALS/BLS and BLS only.
 - i. The ALS/BLS drug bag is a navy, standard issue drug bag with 5 outside compartment.
 - ii. The BLS only bag is red “fanny-pack” style bag.
 - iii. Each bag is labeled with a metal tag reflecting the assigned bag number.
 - iv. Bags may have additional tags from time to time with specific instructions or inventory changes.
- c. All drug bags, both ALS/BLS and BLS, are the property of the GMVEMSC
- d. GMVEMSC drug bags are only for use by EMS providers located or stationed within GMVEMSC’s region.
- e. Agencies may not use GMVEMSC drug bags for runs originating from stations outside of or responding to an address outside of GMVEMSC’s region (except in case of mutual aid responses to those areas).
- f. Except in extreme circumstances, a GMVEMSC drug bag should not be used on multiple runs.
- g. There is an initiation fee for each new bag that EMS agencies add to the program.
- h. There is an annual maintenance fee for each ALS/BLS bag and BLS bag.
- i. For replacement of lost or stolen drug bags, see [7005 Drug Bag Exchange Program: Lost or Stolen Drug Bag Policy](#).
- j. To maintain the integrity of the drug bag contents, pharmacy departments’ seal each compartment of stocked drug bags with a blue plastic device. The seal should only be broken for administration of prehospital emergency medical treatment by approved EMS personnel. After prehospital emergency medical treatment use, the drug bag should be cleaned and re-sealed with the red plastic device contained inside each drug bag compartment.
- k. The following actions may be taken for any agency found to be in non-compliance with the Drug Bag Exchange Program Operating Guideline regarding maintaining custody of, opening and resealing the drug bag:
 - i. Notification of the Fire Chief, EMS Administrator, or Private Ambulance Administrator.
 - ii. The governing agency, e.g., city council, trustees, EMFTS for private ambulance service, will be notified that action is being initiated for the Fire, EMS and Private ambulance service.



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- iii. After the third strike (see [7001.5](#)), removal of all drug bags from all locations of said Fire, EMS and Private ambulance service.
- iv. Written notification to the following that the said service is in violation of the operating policy of the Drug Bag Exchange Program:
 - 1. Medical Director
 - 2. Regional Physician Advisory Board
 - 3. Ohio State Board of Pharmacy
 - 4. Ohio Division of EMS
 - 5. All hospitals participating in the drug bag exchange program
- l. GMVEMS Council maintains an information database for all EMS personnel authorized to participate in the Drug Bag Exchange Program.
- m. Rosters with certification expiration dates for EMS providers are available via an online database for review and updates.

7001.3 Participation Requirements

- a. Active membership in the GMVEMS Council.
- b. Each agency in GMVEMSC must understand that Council typically communicates with departments and agencies via email, and that some of those messages concern changes to Standing Orders, pharmaceuticals in our Drug Bags, or other critical issues. Council maintains two lists of emails:
 - i. The GMVEMSC Listserve
 - ii. A distribution list of Agency Contacts
- c. As such, to participate in the Drug Bag Program, each agency must provide a minimum of one functioning email contact for each of those lists (may be the same person or different). Council desires to communicate as freely and effectively as possible, and agencies may provide as many as they like for each list, but must have at least one person who can reliably receive messages. Since in rare cases, these messages may be urgent, we encourage use of the “three-deep” rule: provide Council with three (or more) emails for each list.
- d. Additional Requirements for Drug Bag Program
 - i. Upon release of the current Implementation Guide, the Department Chief/Agency Head will sign and submit the Attestation Form accessible at this link to the GMVEMSC Attestation & Compliance: <https://forms.office.com/r/UhMMApn34c>.
 - ii. This Form, which is also in the Implementation Guide, is to be completed annually by April 1 to ensure understanding and adherence with the testing processes for each year. The form MUST be completed by using the link above.
 - iii. The copy of the license needs to go to Council by March 31 of each calendar year that the agencies’ drug license is renewed. This is required, as the Pharmacy at each hospital needs the license on file in order to exchange drug bags with your department.
 - iv. Complete drug bag updates when scheduled. This is essential. The Pharmacy Board has made it very clear that updates must be completed on time.
 - v. Signed agreement to abide by the GMVEMS Council Operating Guidelines for the Drug Bag Exchange Program (see [7007 Drug Bag Exchange Program: Agency Agreement Letter](#))



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- e. **No department which participates in the Drug Bag Exchange Program shall possess a DEA License.**
- f. Area hospital participation according to Council guidelines. (See [7006 Drug Bag Exchange Program: Hospital Participation Policy](#))
- g. Document medical advisor approval for the use of the GMVEMS Council Operating Protocols with a signed, notarized letter, which is attached to the drug license renewal application form with a copy submitted to Council. Notarized letter is not required for renewal unless medications are added or there is a change in Medical Director from previous year.
- h. Agreement to complete the GMVEMSC annual skills and annual written test between in the timeline established by Council (see Non-Compliance Procedures).
- i. **Maintain all drugs at all times in a clean, temperature-controlled environment per Rule 4729-33-03 of the Ohio State Pharmacy Board Administrative Code.**
- j. The rules can be seen at: <https://codes.ohio.gov/ohio-administrative-code/rule-4729:3-3-03>
- k. The ideal temperature span is 59-86 degrees Fahrenheit.
- l. In order to utilize an ALS/BLS or BLS drug bag in the pre-hospital emergency setting, the following equipment must be available, unless otherwise noted:
 - i. BLS Provider:
 1. Oxygen
 2. Pulse Oximetry
 3. Extraglottic Airways
 4. CPAP administration and management
 5. Oral Glucose
 6. Glucometry
 7. Ice Packs
 8. Suction (manual is acceptable)
 9. AED
 - ii. ALS Provider:
 1. Oxygen
 2. EtCO₂ detection, monitoring and waveform for intubated patients
 3. 12 or 15 Lead acquisition, transmission and interpretation
 4. Mucosal Atomizer Device (MAD)
 5. IO and device
 6. IV pressure infuser
 7. Suction (manual is acceptable)
 8. Monitor or defibrillator or AED & intubation equipment
- m. Departments are required to have a tracking system that tracks all drug bag exchanges.

7001.4 General Non-Compliance Procedures

- a. Each agency and their Medical Director(s) will be notified if the annual written test and skills check-off has not been completed within the prescribed time period.
- b. The Ohio State Board of Pharmacy will be notified that a department or individual members of a department have not completed the annual written test and skills check-off within the prescribed time period.



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- c. Hospital EMS coordinators and pharmacy departments will receive a list of departments or individuals within a department that are not in compliance with the operating guidelines.
- d. At the end of the testing season, if a department does not have 100% of their personnel completing both skill and written tests (or explanations for individuals not in compliance) noted in the Standing Orders database, then appropriate action, up to and including the removal of department from the Drug Bag program, may be taken by the chair of the drug bag committee.
- e. If copy of drug license(s) is not received by due date, GMVEMS Council will notify the agencies' medical director. GMVEMS Council reserves the right to initiate the non-compliance action process for any Fire/EMS/Private Ambulance service that does not provide documentation for drug license(s) renewal.

7001.5 Three Strike Policy

- a. An agency may be issued a strike for failure to comply with the participation requirements and or general operating guidelines set forth by council and the State Board of Pharmacy.
- b. In the event that a violation has occurred that reaches the level of issuing a strike notice, then the agency or agencies that have incurred the infraction will be notified by certified mail from the Greater Miami Valley EMS Council.
- c. The infraction will remain for a minimum of 1 year and will be reviewed by the Drug Bag Co-Chairs and the President and President-Elect at the end of the 1-year period to determine if the strike can be removed from that agency.
- d. An agency issued a strike has the ability to appeal the infraction if they are able to show proof that the infraction did not occur. This must be supported by proper documentation (i.e. at time of infraction was not able to show a drug bag exchange log but was able to produce one that can be verified). The strike can be removed.
- e. If an agency reaches three strikes, then the agency, their Medical Director as well as the State Board of Pharmacy will be notified of that departments removal from the Drug Bag Program.

7001.6 Levels of Participation

- a. Paramedic Level
 - i. A Paramedic can access any of the compartments within the bag to obtain medications.
- b. AEMT Level
 - i. A side compartment of the ALS/BLS bag will be labeled "Intermediate"
 - ii. The AEMT can access the Airway, BLS, Intermediate and Naloxone compartments to obtain medications per their protocol.
 - iii. The AEMT cannot access the Center Inside Compartment
- c. EMT Level
 - i. The BLS Only fanny-pack style bag will carry:
 1. Albuterol
 2. Atrovent
 3. Baby Aspirin
 4. Nitrostat
 5. Epi-pen and Epi-pen, Jr. (1 each)
 - ii. The RED BLS Pouch on an ALS/BLS bag will carry the following medications ONLY:



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1. Nitrostat
 2. Baby Aspirin
 3. Epinephrine 1:1,000 ampule
- iii. The EMT can only access following to treat their patient per protocol:
1. The BLS Only fanny-pack (if available)
 2. The Airway Pouch of the ALS/BLS bag
 3. The BLS Pouch of the ALS/BLS bag
 4. The Naloxone Pouch of the ALS/BLS bag
 5. The EMT cannot access the Intermediate and Center Compartments

END OF SECTION

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Subject: Drug Bag Exchange Program:
Wasted Drug Procedure

Effective: June 1, 2021

Last modified: May 16, 2026

7002.1 Guideline

- a. Most hospitals require the use of the GMVEMSC approved Controlled Drug Usage Form in addition to documentation of medications given on the run sheet.
- b. This GMVEMSC approved form must be filled out for any controlled drug use, even if there is no wastage.
- c. This information shall be on both the original EMS agency form and the hospital copy for reference.
- d. Every crew transporting a patient will provide a completed run sheet to the hospital within 3 hours.
- e. In cases where communication methods are not functioning, run sheets need to be delivered by alternative communications or by a hard copy delivered to the hospital within 24 hours of patient transfer.

7002.2 Procedure

- a. Fentanyl, Ketamine, Morphine, Versed and Valium are all controlled drugs.
 - i. If a controlled medication from a GMVEMSC Drug Bag is only partially administered, the paramedic or AEMT must account for the all the unused portions.
 - ii. When wasting controlled medications at the end of a call, providers will waste the drug directly into a secure sharps container in the ambulance.
 1. Wasting controlled medications into a hospital sharps container is not permitted.
 2. The substance must be drawn out of the vial and discharged into the sharps container.
 - iii. Some private organizations may have different rules for waste of controlled medications that are approved by the Ohio Board of Pharmacy and may use those procedures.
 1. If an agency varies from the procedures listed in 7002.2.ii, they must provide their policy in writing to the GMVEMSC.
 - iv. Agencies that need to waste a controlled medication not from the GMVEMSC Drug Bag, at their home facilities, will do so in a similar manner and document the procedure.
 1. Discharging the medication into an absorbent material or a commercial drug waste device are acceptable options as well.
 2. If this practice is employed, the discharge vessel must be in a secured area.
 3. Any drug wasting practices employed by agencies at their own facilities must be written into a policy and approved by medical direction.
- b. To ensure the medications are properly accounted for, all paramedics and AEMTs will document:
 - i. The drug name
 - ii. The amount used
 - iii. The amount wasted (if all the medication was administered, then list "none")
 - iv. If an amount was wasted, specifically where did that waste occur (i.e. into sharps container)
 - v. The signature of a second witness if there is wastage.
 1. The second witness can be a member of the EMS crew.
 2. Often hospital employees are not permitted to witness or sign for drug wastage.

END OF SECTION

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Subject: Drug Bag Exchange Program:
Exchange Process

Effective: June 1, 2021

Last modified: Dec. 12, 2022

7003.1 Guideline

- a. Each department is assigned to a "home" hospital.
- b. The assigned hospital is the central resource for initial fulfillment of medications for the drug bags and wholesale exchanges, replacement, or additions as required by revisions to the protocols.
- c. Drug bags can be exchanged at any participating hospital or within the same department.
- d. ALS/BLS bags may be exchanged one-for-one with another ALS/BLS bag.
- e. BLS bags may be exchanged one-for-one with another BLS bag.
- f. It is not permissible to exchange drug bags between two different Fire/EMS Agencies.
- g. The primary care provider is responsible for the inventory of the drug bag prior to sealing it.
- h. If two departments have accessed a drug bag, they should jointly seal the drug bag.
- i. Each hospital designates a specific location for the exchange of drug bags.
- j. EMS personnel are **required** to complete the Sign In and Out log when exchanging a drug bag.
- k. Each agency is responsible to track drug bag exchanges within their own organization (i.e. documentation, internal log, tracking software, etc.)
- l. Once sealed, any provider can exchange the drug bag.
- m. Unless the patient was removed to a non-participating drug bag exchange hospital or the patient was a non-removal, the drug bag must be exchanged at the time of patient delivery to the hospital.
- n. In the exceptions listed above, the drug bag will be exchanged at a participating hospital within 8 hours.
- o. Drug Bag Exchange after field termination will be at the facility from where the order was given, unless that hospital is not part of the Drug Bag Exchange Program.

7003.2 Drug Bag Blue Seals

- a. Blue seals:
 - i. Blue seals are used by the pharmacy that inventories and restocks the ALS/BLS drug bags.
 - ii. The blue seals will have a hospital sticker attached to the seal that identifies the hospital and pharmacist that inventoried the bag and the expiration date of the next drug to expire.
 - iii. The inner compartment of the ALS bag and Intermediate will be sealed with a blue seal and will have the expiration date noted.
 - iv. The blue seal will be looped through the proximal portion of the zipper tab (not the outermost portion of the zipper tab).
 - v. EMS should verify the blue seal is intact and has an expiration date before accepting the bag.
 - vi. When a provider opens a drug bag compartment, they should keep the blue seal in their possession until they have verified the contents are accounted for.
 - vii. Once they have verified the contents, they should place the blue seal in the compartment, unless there is a discrepancy and then seal the compartment with RED tag.
 - viii. EMS MUST PLACE THE BLUE SEAL IN THE COMPARTMENT!
- b. Red Seals:
 - i. Red seals identify ALS/BLS bags as being used.
 - ii. EMS providers are required to inventory each opened pouch, discard any used sharps and clean any contaminants from bag used and then take red seal from the inside compartment (supplied by pharmacy when restocking the ALS/BLS bag) and seal the used compartment.
 - iii. The red seal will be looped through the proximal portion of the zipper tab (not the outermost portion of the zipper tab).

END OF SECTION

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Subject: Drug Bag Exchange Program:
Drug Bag Discrepancies

Effective: June 1, 2021

Last modified: May 15, 2026

7004.1 General Guidelines

- a. **EMS providers are required to inventory each opened pouch prior to applying the red seal.**
- b. All discrepancies (missing meds, wrong med or dose, altered or tampered meds, drug bag number discrepancy, etc.) that are identified shall be reported to GMVEMSC using the Drug Bag Discrepancy Report (Addendum E).
- c. If at any time, an EMS provider encounters a discrepancy they will:
 - i. Notify their EMS Officer of the discrepancy.
 - ii. If the discrepancy was discovered after opening the bag, retain the blue seal and the hospital sticker that was attached to the drug bag in question.
 - iii. If the EMS provider is at the hospital, he/she will log the bag in using the normal procedure at that hospital while retaining the blue seal.
 - iv. He/she will advise the pharmacist or EMS Coordinator of the discrepancy and that they will be initiating the Discrepancy form as described below (pharmacist may request a copy of the Discrepancy form).
 - v. The EMS Officer may contact the EMS Coordinator if assistance is needed.
- d. In the event an expired medication is discovered, complete a drug usage form identifying the expired medication and exchange the bag at a participating hospital.

7004.2 Discrepancies Involving Controlled Drugs or Potential Tampering:

- a. When an issue arises concerning any of the following, a collaborative effort between the EMS organization or provider and the Hospital EMS Coordinator or Pharmacist shall be made in an attempt to resolve the issue:
 - i. A controlled drug (Fentanyl, Ketamine, Valium, Versed, or Morphine)
 - ii. A stolen, missing or lost bag
 - iii. Any medication that appears to have been altered or tampered with.
- b. If the issue cannot be resolved, the following steps shall be taken:
 - i. If the discrepancy was discovered by the EMS organization/provider, the person designated by the organization/provider shall comply with the requirements of OAC 4729-9-15 and GMVEMSC requirements as indicated below.
 - ii. If the discrepancy was discovered by the hospital, the person designated by the hospital shall comply with the requirements of OAC 4729-9-15 and GMVEMSC requirements as indicated below.
- c. Required reporting for unresolved issues involving Controlled Drug or potential/suspected tampering or lost or stolen drug bags pursuant to Federal and State Laws and GMVEMSC Protocol include:
 - i. If you have knowledge of or suspect a discrepancy is due to a theft, contact your State of Ohio Board of Pharmacy agent immediately. Advise them you want to report a theft or drug discrepancy. They will connect you with the appropriate person. (OAC 4729-9-15)
 - ii. Notify the Drug Bag Exchange Committee Chairs immediately.
 - iii. File a report with the appropriate law enforcement authorities (ORC 2921.22).
 - iv. Notify the Drug Enforcement Agency within 24 hours of discovery using DEA Form 106
 - v. DEA Form 106: <https://www.deadiversion.usdoj.gov/webforms/app106Login.jsp>.
 - vi. A 30-day extension may be requested in writing from the DEA. (CFR 1301.76(b)).



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Drug Bag Discrepancies

Effective: June 1, 2021

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- vii. Submit a completed GMVEMSC Drug Bag Discrepancy Report located at Addendum #E, with appropriate supporting documentation, to the GMVEMSC.
- d. "Dangerous drug" means any of the following:
 - i. Any drug to which either of the following applies:
 - 1. Under the "Federal Food, Drug, and Cosmetic Act," 52 Stat. 1040 (1938), 21 U.S.C.A. 301, as amended, the drug is required to bear a label containing the legend "Caution: Federal law prohibits dispensing without prescription" or "Caution: Federal law restricts this drug to use by or on the order of a licensed veterinarian" or any similar restrictive statement, or the drug may be dispensed only upon a prescription;
 - 2. Under Chapter 3715 or 3719 of the Revised Code, the drug may be dispensed only upon a prescription.
 - ii. Any drug that contains a schedule V controlled substance and that is exempt from Chapter 3719. of the Revised Code or to which that chapter does not apply;
 - iii. Any drug intended for administration by injection into the human body other than through a natural orifice of the human body;
 - iv. Any drug that is a biological product, as defined in section [3715.01](#) of the Revised Code.

7004.3 Discrepancies Not Involving Controlled Drugs or Potential Tampering:

- a. Examples may include:
 - i. Non-controlled drugs that were not in the bag
 - ii. Wrong number of medications or doses
 - iii. Wrong drug concentration
 - iv. Expired medications found
 - v. No expiration date on tag
 - vi. Medications improperly labeled
 - vii. Empty vials or packages left in bag. DO NOT PUT ANY USED VIALS BACK IN DRUG BAG
 - viii. Unsealed medications
 - ix. Wrong medication administered
 - x. Unsealed pouch discovered
 - xi. Bag logged out with red seal (used bag)
- b. If discovered by EMS, the EMS Officer will initiate the Discrepancy form. They shall provide a copy of the form and the Blue Seal to the Hospital EMS Coordinator.
- c. If the Hospital discovers the discrepancy, the EMS Coordinator will initiate the Discrepancy Form and submit to GMVEMSC.
- d. If the EMS Coordinator is able to determine which EMS agency/hospital is responsible for the discrepancy, the agency or hospital will be notified and will receive a copy of the Discrepancy Form and the Blue Seal if applicable.

7004.4 Follow Up Procedures

- a. The GMVEMSC will:
 - i. Maintain a record of all discrepancies that occur.



Subject: **Drug Bag Exchange Program:
Drug Bag Discrepancies**

Effective: **June 1, 2021**

Last modified: **May 15, 2026**

- ii. Follow up with the agencies involved as needed.
 - iii. Advise the Drug Bag Chairperson of any and all discrepancies and action taken.
- b. The Drug Bag Committee Chairperson will:
 - i. Report at the Drug Bag Committee meetings for discussion and resolutions to all discrepancies encountered.
 - ii. Assist the Council and or affected departments with any issues or questions that may result.

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Subject: **Drug Bag Exchange Program:
Lost or Stolen Drug Bag Policy**

Effective: **June 1, 2021**

Last modified: **Mar. 1, 2022**

7005.1 Purpose

- a. To provide a uniform mechanism for the investigation and reporting of lost or stolen drug bags.

7005.2 Notification

- a. Upon discovery of a missing GMVEMSC drug bag, agencies will notify or cause to be notified the GMVEMSC Drug Bag Committee Chair(s).
- b. A responsible party at the agency will initiate the Drug Bag Discrepancy Form and follow instructions for reporting lost or stolen drug bags. Completed paperwork and reports will be submitted to GMVEMSC.
- c. The agency representative or the GMVEMSC Drug Bag Committee Chair (s) will notify the State of Ohio Board of Pharmacy (SOBP) at 614-466-4143
 - i. The Drug Bag Chair(s) may elect to notify the SOBP for the agency or advise the agency to contact them individually.
 - ii. Either way contact with the SOBP must be coordinated and accomplished

7005.3 Investigation

- a. The EMS agency shall develop and implement an internal search mechanism for lost drug bags.
- b. The internal search mechanism should include:
 - i. Determine if drug bag was left at the scene.
 - ii. Determine if drug bag was not exchanged on last run.
 - iii. Determine if drug bag is in the wrong vehicle.
- c. The GMVEMSC will seek the assistance of the GMVEMSC Drug Bag Chair(s) to check with all hospitals to determine if the bag might be in inventory or be alerted if it shows up at one of the hospitals.
- d. The GMVEMSC will contact the hospital EMS Coordinator with whom the EMS Department is assigned to work out a drug bag replacement.
 - i. Drug bag replacement will only occur after all paperwork is submitted
 - ii. The GMVEMSC will assess a fee for replacement bag to be paid for by the receiving agency.

END OF SECTION

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Subject: **Drug Bag Exchange Program:
Hospital Participation Policy**

Effective: **June 1, 2021**

Last modified: **Dec. 8, 2020**

7006.1 Purpose

- a. To assure uniformity of hospital pharmacy participation in the Drug Bag Exchange Program.

7006.2 The Hospital Shall:

- a. Purchase (at cost), fill, and maintain a supply of drug bags sufficient to meeting the needs of an average day, plus a few extra to meet peak demands for drug bag replacement.
- b. Accept responsibility for filling new drug bags for departments or vehicles as assigned by GMVEMS Council, at hospital expense.
- c. Assign one licensed pharmacist and an EMS coordinator to attend and participate in the Standing Orders and Drug Bag Exchange Program Committees.
- d. Agree to pay annual dues and any fees assessed by GMVEMS Council that are approved by the Drug Bag Exchange Program Committee and the GMVEMS Council that pertain to the Drug Bag Exchange Program.

7006.3 The Greater Miami Valley EMS Council shall:

- a. Maintain a current State Drug Licenses for all participants in the Drug Bag Exchange Program.
- b. Furnish hospital pharmacy with a current listing of all departmental personnel authorized to access the GMVEMSC drug bags and copy of the protocol.
- c. Assign departments to hospitals in both a geographic and otherwise equitable fashion.

END OF SECTION

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Subject: Drug Bag Exchange Program:
New Agency Member Policy

Effective: June 1, 2021

Last modified: Dec. 8, 2020

7007.1 Purpose

- a. To establish the procedures required to provide new agency members with an ALS or BLS drug bag from the GMVEMSC Drug Bag Exchange Program.

7007.2 Procedure:

- a. Those agencies who have applied for membership and require a GMVEMSC Drug Bag to license their units may request a GMVEMSC Drug Bag be available 24 hours prior to the Ohio Medical Transportation Board (OMTB) inspection date.
- b. In order to receive a drug bag, the EMS agency shall:
 - i. Have applied for a GMVEMSC membership.
 1. Have been given a provisional membership by the GMVEMSC Executive Committee if the inspection is before regularly scheduled Council meeting.
 - ii. Provide a copy of their State Pharmacy License.
 - iii. Check off all agency personnel on Standing Orders and data entered in the GMVEMSC data base.
 - iv. Have the Medical Director submit a notarized letter to the State Pharmacy Board with License application stating they approve their department to use the GMVEMSC protocols.
 1. Medical Directors have the right to limit their personnel from using certain medications or procedures within the scope of the GMVEMSC protocols.
 2. Medical Directors may elect to change or add medications or procedures to the protocol.
 3. The Medical Director must include those protocols in addendum to the GMVEMSC, be responsible for the training and documentation of training in of their protocol as well as purchasing and maintaining those drugs that are not included in the standard inventory of the GMVEMSC ALS or BLS drug bag.
 - v. The agency has 72 hours to show proof of a temporary permit from the date of inspection to the GMVEMS Council office.

7007.3 Agreement Letter

- a. In order to participate in the GMVEMS Council Drug Bag Exchange program, the agency will provide the agreement letter that follows to the Greater Miami Valley EMS Council.
- b. A similar example of the agencies' choosing may also be used.

**Subject:** Drug Bag Exchange Program:
New Agency Member Policy**Effective:** June 1, 2021**Last modified:** Dec. 8, 2020

**Greater Miami Valley EMS Council
Drug Bag Exchange Program
Agency Agreement Letter**

Please type or print legibly

DEPARTMENT/SERVICE: _____**CONTACT PERSON:** _____**TELEPHONE:** _____**FAX:** _____

This department/service agrees to abide by the GMVEMS Council Drug Bag Exchange Program and Standing Orders.

SIGNATURE: _____

Fire Chief, EMS Administrator, or Private Ambulance Administrator

DATE: _____

Return to:
GMVEMSC
124 E. Third St.
Dayton OH 45402

END OF SECTION



Subject: **Drug Bag Exchange Program:
GMVEMSC Drug Bag Discrepancy Report**

Effective: **June 1, 2021**

Last modified: **Dec. 8, 2020**

7008.1 General Guideline

- a. If at any time an EMS provider encounters a discrepancy in the GMVEMS Council Drug Bag they are using, they will notify their agencies' EMS Officer (or their supervisor if an EMS Officer does not exist).
- b. If the EMS provider is at a hospital that participates in the GMVEMS Council Drug Bag Exchange Program, they will log the bag in using the normal procedure at that hospital.
- c. If the discrepancy was discovered after opening the bag, retain the blue seal and the hospital sticker that was attached to the drug bag in question. The seals (or photo copies of the seals should be attached to the **GMVEMSC Drug Bag Discrepancy Report**.
- d. They will advise the pharmacist or EMS Coordinator of the discrepancy and that they will be initiating the **GMVEMSC Drug Bag Discrepancy Report** provided on the opposite page.
- e. Examples of the **GMVEMSC Drug Bag Discrepancy Report** should be available at all hospitals. They will often be found in the EMS rooms.
- f. The **GMVEMSC Drug Bag Discrepancy Report** will be completed in triplicate with a copy going to the GMVEMS Council, the receiving pharmacy and the EMS agency reporting.
- g. The pharmacist may request a copy of the **GMVEMSC Drug Bag Discrepancy Report**.

END OF SECTION

GMVEMSC Drug Bag Discrepancy Report

If at any time an EMS provider encounters a discrepancy they will notify their EMS Officer of the discrepancy. If the discrepancy was discovered after opening the bag, retain the blue seal and the hospital sticker that was attached to the drug bag in question. If the EMS provider is at the hospital, they will log the bag in using the normal procedure at that hospital. They will advise the pharmacist or EMS Coordinator of the discrepancy and that they will be initiating the Discrepancy form as described below (pharmacist may request a copy of the Discrepancy form).

Date of report: _____ Bag Number: _____ Date Discrepancy discovered: _____

Discovered by: _____ Hospital/EMS Dept making discovery: _____

Have blue Hospital seal? YES/NO If yes - Attach seal to report

Tracking:

Date bag was logged out: _____ from (hospital) _____ To (EMS agency) _____

Date Bag turned in: _____ to (hospital) _____

Description of the discrepancy: (Attach addendum if additional space needed)

Describe efforts to resolve the discrepancy: (Attach addendum if additional space needed)

Was the discrepancy satisfactorily resolved? _____ If not, what steps are to be taken:

Who will be responsible for any required reporting: _____

Reporting requirements:

Was a police report filed? _____

Date: _____ By whom? _____

Was a DEA report filed? _____

Date: _____ By whom? _____

Was the Stat Pharmacy Board notified? _____

Date: _____ By whom? _____

Required documents submitted to GMVEMSC By: _____ **Date:** _____

For Drug Bag committee use:

Wrong medication stocked		Bag logged out with red seal	
Expired medication found		Empty vials/packages found	
Wrong dose packaged		Open pouch found	
Missing medications		Unsealed bottles found	
Wrong number packaged		Medication found in wrong compartment	
No expiration date on tag		Wrong medication administered	
Atrovent/Albuterol not labeled		Lost or stolen bag	
Damaged medications		Other:	
Other:			

GMVEMSC – White

Pharmacy - Yellow

EMS Department - Blue



Subject: **Drug Bag Exchange Program:
Report of Theft or Loss of Dangerous
Drugs, Controlled Substances and Drug
Documents**

Effective:
June 1, 2021

Last modified:
Dec. 8, 2020

7009.1 OAC 4729-9-15

(A) Each prescriber, terminal distributor of dangerous drugs, or wholesale distributor of dangerous drugs shall notify the following upon discovery of the theft or significant loss of any dangerous drug or controlled substance, including drugs in transit that were either shipped from or to the prescriber, terminal distributor of dangerous drugs, or wholesale distributor of dangerous drugs:

- (1) The state board of pharmacy, by telephone immediately upon discovery of the theft or significant loss;
- (2) If a controlled substance, the drug enforcement administration (DEA) pursuant to section 1301.76(b), Code of Federal Regulations;
- (3) Law enforcement authorities pursuant to section 2921.22 of the Revised Code.

(B) Controlled substance thefts must also be reported by using the Federal DEA Report form whether or not the controlled substances are subsequently recovered and/or the responsible parties are identified and action taken against them. A copy of the federal form regarding such theft or loss shall be filed with the State Board of Pharmacy within thirty days following the discovery of such theft or loss.

- (1) An exemption may be obtained upon sufficient cause if the federal form cannot be filed within 30 days.
- (2) A request for a waiver of the thirty-day limit must be requested in writing.

(C) Each prescriber, terminal distributor of dangerous drugs, or wholesale distributor of dangerous drugs immediately upon discovery of any theft or loss of:

- (1) Uncompleted prescription blank(s) used for writing a prescription, written prescription order(s) not yet dispensed, and original prescription order(s) that have been dispensed, shall notify the state board of pharmacy and law enforcement authorities.
- (2) Official written order form(s) as defined in division (Q) of section 3719.01 of the Revised Code shall notify the state board of pharmacy and law enforcement authorities, and the drug enforcement administration (DEA) pursuant to section 1305.12(b), Code of Federal Regulations.

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Subject:

Ambulance Restocking Policy

Effective:

June 1, 2021

Last modified:

Dec. 8, 2020

7010.1 History

- a. The member hospitals of Greater Dayton Hospital Association (GDAHA) have supported Emergency Medical Services agencies in the region for decades.
- b. In 1998, GDAHA received permission (Advisory Opinion No. 98.7) from the Department of Health & Human Services to continue to exchange drugs (GMVEMSC Drug Bag Exchange Program) and supplies with EMS agencies and avoid violating the anti-kickback (safe harbor) statute of the Social Security Act.
- c. The hospitals named in the advisory are in the eight (8) county West Central Region: Champaign, Clark, Darke, Greene, Miami, Montgomery, Preble and Shelby.
- d. In December 2001, the Centers for Medicare and Medicaid Services issued an Ambulance Final Rule on Ambulance Restocking Safe Harbor.
 - i. Elements of the Safe Harbor include:
 1. Billing and claim submission
 2. Documentation
 3. Not tied to referrals
 4. Compliance with other laws

7010.2 EMS Supply Exchange Program:

- a. EMS agencies and personnel should understand the benefits of the EMS Supply exchange program, as offered by GDAHA members participating in this program.
- b. Hospitals are not required to participate in this restocking program.
- c. EMS agencies and personnel must adhere to the agreement, particularly the areas highlighted below:
 - i. For all transports to member hospitals, the EMS agencies will provide the receiving hospital with copies of the written records **describing each of the medical supplies and/or medications utilized by or for the patient during the transport**. In most cases, this should be done immediately after patient transfer.
 - ii. Participating hospitals will restock EMS agency ambulances, at no charge to the EMS agency, with the medical supplies and/or medications which were **utilized by or for the patient during the transport to the receiving hospital**.
- d. **Hospitals will not restock items used on patients delivered to another hospital.**
 - i. Restocking an ambulance at a participating hospital for items used on a patient delivered to a hospital not participating in the agreement will jeopardize this program.
 - ii. It is the responsibility of the EMS agencies to restock items used on patients delivered to a hospital that is not a participant in the Agreement.
- e. **Participating hospitals will restock drug bags.**
- f. Hospitals will not provide medical supplies to a new ambulance, or an old ambulance being returned to service.
 - i. These ambulances must be stocked for the first time by the EMS agency.

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Subject: Diversion of Emergency Patients

Effective: June 1, 2021

Last modified: Jan. 5, 2024

**Greater Dayton Area Hospital Association, Greater Miami Valley Emergency Medical Services Council, and
Greater Montgomery County Fire Chiefs' Association Policy Statement for Temporary Diversion of
Emergency Patients**

7011.1 EMS and Dispatch Procedures:

- a. When situations exist that prevent the timely treatment of additional emergency cases or certain types of emergency patients, the designated hospital or satellite emergency department (ED) Official will report that they are on "Diversion of Emergency Patients," formerly referred to as rerouting.
- b. For patients impacted by the type of diversion specified, EMS should utilize hospitals in normal status. Transport to a hospital in diversion status may jeopardize patient care more than the delay in treatment caused by longer transport times except for patients that are in extreme life/limb threatening circumstances.
- c. When a patient and /or the patient's physician requests EMS to transport to a hospital which is on diversion, EMS have the responsibility to advise the patient and/or the physician that "due to diversion patient care may be jeopardized."

7011.2 Monitoring Emergency Department Status

- a. Anyone with a Juvare EmResources account can set up preferences to receive an alert when the hospital status changes.
 - i. Dispatch centers should set up Juvare EMResources preferences to receive an alert when the hospital status changes.
 - ii. Dispatch centers are encouraged to continuously monitor Juvare EMResources.
 - iii. Dispatch centers must notify EMS of hospital status changes.

7011.3 Diversion Categories:

- a. Hospitals communicate the following status information via Juvare EMResources:
 - i. **CLOSED:**
 1. The hospital or satellite ED has activated its disaster plan because of an internal emergency or other situation rendering the hospital or satellite ED unable to accept **any emergency patient. EMS will not transport any patient to a CLOSED Facility.**
 - ii. **DIVERSION OF CERTAIN TYPES OF PATIENTS:**
 1. Limited Divert/Operations:
 - a. Limited operations/ability to handle some types of traffic/special situation (examples include CT scanner downtime, no ICU beds available, specialty care limitations). Write the specific issue in the comment section.
 2. Divert/At Capacity:
 - a. Facility is at capacity and/or on diversion; ED is paused to inbound EMS traffic and the facility is not in a designated load balancing plan.



Subject: Diversion of Emergency Patients

Effective: June 1, 2021

Last modified: Jan. 5, 2024

7011.4 Hospital and Satellite ED Procedures:

- a. The hospital or satellite ED will:
 - i. Update the Juvare EMResources page with ED status and activity between 6 and 9 am daily and anytime the status or activity changes.
 - ii. Notify EMS Coordinators and appropriate dispatch centers. Hospitals and satellite EDs located in the southern Miami Valley region may also need to contact northern Cincinnati area hospitals or dispatch centers.
- b. Status Management - Changes/Updates
 - i. It is the responsibility of the **diverting** hospital or satellite ED to review and update their diversion status **hourly**, making changes as needed.
 - ii. When the status changes, including return to normal operations, notify EMS Coordinators and appropriate dispatch centers and update Juvare EMResources using the same notification protocols used to initiate the diversion procedure.

7011.5 Participating Hospitals (Additional hospitals added upon approval)**Atrium Medical Center (Middletown)**

1 Medical Center Dr, Middletown, OH 45005

Austin Boulevard Emergency Center

300 Austin West Blvd., Miamisburg, OH 45342

Dayton Children's Hospital

1 Children's Plaza, Dayton, OH 45404

Dayton Children's Hospital – South Campus

South Campus 3333 W. Tech Blvd, Miamisburg, OH 45342

Dayton-Springfield Emergency Center

1840 Springfield Road, Fairborn, OH 45324

Joint Township District Memorial Hospital

200 St. Clair Ave, St. Marys, OH 45885

Kettering Health Dayton

405 W Grand Ave, Dayton, OH 45405

Kettering Health Network Franklin Emergency Center

100 Kettering Way, Franklin, OH 45005

Kettering Health Greene Memorial

1141 N Monroe Dr, Xenia, OH 45385

Kettering Health Hamilton

630 Eaton Ave, Hamilton, OH 45013

Kettering Health Network Huber Emergency Center

8701 Troy Pike, Huber Heights, OH 45424

Kettering Health Main Campus

3535 Southern Blvd, Kettering, OH 45429

Kettering Health Miamisburg

4000 Miamisburg Centerville Rd, Miamisburg, OH 45342

Kettering Health Middletown Emergency Center

6147 W. State Route 122 Middletown, OH, 45005

Kettering Health Preble Emergency Center

450-B Washington-Jackson Rd, Eaton, OH 45320

Kettering Health Springfield

2300 N. Limestone St., Springfield OH 45503

Kettering Health Troy

600 W. Main St., Troy, OH 45373

Kettering Health Washington Township

1997 Miamisburg Centerville Rd, Dayton, OH 45459

Mercy Health – Springfield

100 Medical Center Drive, Springfield, OH 45504

Mercy Health Urbana Hospital

904 Scioto St, Urbana, OH 43078

Miami Valley Hospital

1 Wyoming St, Dayton, OH 45409

Miami Valley Hospital – Beaver Creek Emergency Center

2400 Lakeview Dr., Beaver Creek, OH 45431

Subject: **Diversion of Emergency Patients**Effective: **June 1, 2021**Last modified: **Jan. 5, 2024**

Miami Valley Hospital - Jamestown Emergency Center
4940 Cottonville Rd, Jamestown, OH 45335

Miami Valley Hospital North
9000 N Main St, Dayton, OH 45415

Miami Valley Hospital South
2400 Miami Valley Dr, Centerville, OH 45459

Soin Medical Center
3535 Pentagon Blvd, Beavercreek, OH 45431

Upper Valley Medical Center
3130 N Co Rd 25A, Troy, OH 45373

Dayton VA Medical Center
4100 West 3rd Street, Dayton, OH 45428

Wayne Healthcare
835 Sweitzer St, Greenville, OH 45331

Wilson Memorial Hospital
915 West Michigan Street, Sidney, OH 45365

WPAFB 88th Medical Center
4881 Sugar Maple Dr, Wright-Patterson AFB, OH 45433

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Subject: Hospital Capabilities Chart

Effective: June 1, 2021

Last modified: July 1, 2026

HOSPITAL	Trauma Center	Burn Center	24 hr Interventional Cardiac Cath	Stroke Telemedicine	Stroke Primary	Stroke Comprehensive	L & D
Atrium Medical Center (Middletown)	Adult 3		Y	Y	Y		Y
Austin Blvd. Emergency Center				Y			
Bethesda Arrow Springs				Y			
Bethesda Butler Hospital				Y	Y		
Christ Hospital Liberty				Y			Y
Dayton Children's Hospital	Pedi 1	Y					
Dayton Children's - South Campus							
Dayton-Springfield Emergency Center				Y			
Joint Township District Memorial Hosp.				Y			
Kettering Health Dayton	Adult 3		Y	Y	Y		
Kettering Health Franklin				Y			
Kettering Health Greene Memorial				Y			
Kettering Health Hamilton			Y	Y	Y		Y
Kettering Health Huber				Y			
Kettering Health Main Campus	Adult 2		Y	Y	Y	Y	Y
Kettering Health Miamisburg				Y	Y		
Kettering Health Middletown				Y			
Kettering Health Preble				Y			
Kettering Health Springfield				Y			
Kettering Health Troy				Y			
Kettering Health Washington Twp.				Y	Y		Y
McCullough-Hyde Hospital				Y			Y
Mercy Health – Kings Mill				Y			
Mercy Health - Springfield			Y	Y	Y	(Thrombectomy Capable)	Y
Mercy Health - Urbana Hospital				Y			
Miami Valley Hospital	Adult 1	Y	Y	Y	Y	Y	Y
Miami Valley – Beavercreek EC				Y			
Miami Valley - Jamestown EC				Y			
Miami Valley Hospital North				Y			
Miami Valley Hospital South			Y	Y	Y		
Reid Health	Adult 3		Y	Y	Y		Y
Soin Medical Center			Y	Y	Y		
Upper Valley Medical Center	Adult 3		Y	Y	Y		
Dayton VA Medical Center							
Wayne Health Care				Y			Y
West Chester Hospital	Adult 3		Y	Y	Y		Y
Wilson Memorial Hospital			Y	Y			Y
WPAFB 88 th Medical Center							

Notes: Comprehensive stroke centers have the capability of endovascular intervention 24/7. Primary stroke centers have CT and tPA capabilities and focus on evaluating patients for intravenous tPA. Telemedicine with tPA ready offers immediate access to a Neurologist.

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Subject:

Hospital Contact Information

Effective:

June 1, 2021

Last modified:

Feb. 28, 2025

Hospitals in **bold type** ask to be called for every patient

HOSPITAL	PHONE	FAX
Atrium Medical Center, Middletown Maternity	513-424-3924 513-974-8700	513-420-5133
Austin Boulevard Emergency Center	937-865-9663	937-641-2608
Bethesda Arrow Springs	513-282-7222	513-867-2581
Bethesda Butler Hospital	513-893-8222	513-893-8321
Christ Hospital Liberty	513-648-7874	513-648-7962
Cincinnati Children's Stat Line	513-636-8008	513-636-4050
Dayton Children's Hospital	937-641-4444	937-641-5301
Dayton Children's Hospital South	937-641-5642	937-641-4880
Dayton-Springfield Emergency Center	937-523-8792	937-523-8788
Joint Township District Memorial Hospital	419-394-7333	419-394-1902
Kettering Health Dayton	937-723-3419	937-723-4609
Kettering Health Franklin Emergency Center	937-458-4728	937-458-4737
Kettering Health Greene Memorial	937-372-2297	937-352-3501
Kettering Health Hamilton	513-867-2144	513-867-2581
Kettering Health Huber	937-558-3301	937-558-3349
Kettering Health Main Campus	937-395-8080	937-395-8347
Kettering Health Miamisburg	937-384-8766	937-384-8729
Kettering Health Middletown	513-261-3415	513-261-3419
Kettering Health Preble	937-456-8328	937-456-8377
Kettering Health Springfield	937-504-8306	937-504-8309
Kettering Health Troy	937-980-7015	937-980-7019
Kettering Health Washington Township Maternity	937-435-1832 937-401-6850	937-401-6447 937-401-6861
McCullough-Hyde Hospital	513-524-5353	513-523-0144
Mercy Health – Kings Mill	513-637-9360	513-978-5010
Mercy Health - Springfield	937-523-1902	937-523-1950
Mercy Health Urbana Hospital	937-484-6160	937-484-6183
Miami Valley Hospital Maternity	937-208-2440 937-208-3677	937-641-2608 937-208-2651
Miami Valley – Beaver Creek Emergency Center	937-429-0708	937-641-2608
Miami Valley – Jamestown Emergency Center	937-374-5274	937-641-2608
Miami Valley North Hospital	937-540-1067	937-641-2608
Miami Valley South Hospital Maternity	937-438-2662 937-974-8700	937-641-2608
Regional Hospital Notification System	937-333-8727	
Reid Memorial Hospital	765-983-3161	765-983-3038
Soin Medical Center	937-702-4525	937-702-4509
Upper Valley Medical Center	937-440-9444	937-440-4346
Dayton VA Medical Center	937-262-2172	937-267-5364
Wayne Health Care	937-547-5777	937-569-6087
West Chester Hospital Maternity	513-298-7777 513-298-7777	513-298-8978
Wilson Memorial Hospital	937-498-5300	
WPAFB Medical Center	937-257-3295	937-656-1673

Hospitals in **bold type** ask to be called for every patient.

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Subject: Infectious Disease Exposure
Reporting Policy

Effective:
June 1, 2021

Last modified:
Jan. 31, 2021

7014.1 General Guidelines

- a. The purpose of this policy is to provide public safety personnel (including fire, EMS, and law enforcement) and hospitals with a set of standard guidelines and expectations for defining, responding to, and following up on an infection control exposure incident involving an emergency response provider.
- b. This guideline is a cooperative effort between the Greater Miami Valley EMS Council (GMVEMSC) and the Greater Dayton Area Hospital Association (GDAHA).

7014.2 Blood-borne Exposure

a. Definition of A Blood-borne Exposure

- i. An exposure incident that may place a public safety worker at risk for Hepatitis B Virus (HBV), Hepatitis C Virus (HCV), or Human Immunodeficiency Virus (HIV) infections or other blood borne pathogens that includes:
 1. A percutaneous injury (e.g., a needle stick or cut), or
 2. Contact of mucous membrane or non-intact skin (e.g., exposed skin that is chapped, abraded, or afflicted with dermatitis) with blood, tissue, or other body fluids that are potentially infectious.
- ii. What is NOT an exposure?
 1. A percutaneous injury with a clean or sterile needle or instrument.
 2. Intact skin splashed with potentially infectious blood, body fluid, or tissue.

b. Post Exposure Procedure

- i. An exposed public safety worker should take the following immediate “first aid” action steps:
 1. Immediately irrigate the involved area.
 2. Flush eyes with copious amounts of IV fluids, if indicated.
 3. Wash skin vigorously with soap and water.
 4. If soap and water is not available, rinse area with another available solution such as IV fluids or a water-based liquid.
 5. Waterless hand cleaners are not recommended for post-exposure gross decontamination, but can be used when other options are not available.
- ii. The Employee shall report the exposure incident to the receiving hospital and to their immediate supervisor.
- iii. Exposed employees are required to register as a patient at the same hospital as the source.
- iv. Once at the receiving hospital, the exposed employee should locate and complete the “Request for Information by Emergency Care Workers (RIECW)” form (see Appendix A).
- v. When completed, the form should be submitted to the nurse handling the exposed employee’s care in the Emergency Department (ED).
- vi. The EMS Coordinator for the receiving hospital can serve as a liaison between the organization and the hospital.
- vii. The department’s infection control officer (ICO) or designated supervisor should, upon receiving notification that there has been an exposure incident, notify the receiving hospital’s EMS Coordinator.



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- viii. *For the purpose of this policy the “department’s Infection Control Officer (ICO), designated supervisor, or designee” refers to the person responsible for reporting and coordinating an exposed employee’s incident within that Public Safety entity.*
- ix. Follow-up care/exam(s) will be provided to each employee involved when indicated. All follow-up care/exam(s) will be coordinated through your employer.

c. Testing The Source Patient

- i. A blood sample is required to determine whether a patient has HIV, HBV or HCV. Blood/Body Fluid (B/BF) testing of a source patient includes the following (MMWR, June 29, 2001):
 - 1. HIV antibody
 - 2. HBV surface antigen (HBsAg)
 - 3. HCV antibody
- ii. If the source patient is transported to a hospital:
 - 1. The ED obtains patient consent and the blood specimen for testing.
 - 2. In the event that the patient refuses to or cannot give consent (e.g., due to an altered level of conscious) a hospital’s “infection control committee... or other body of a health care facility performing a similar function” has the authority to obtain the HIV screening when there has been a significant exposure (Ohio Revised Code §3701.242).
- iii. If the source patient refuses transport to a hospital:
 - 1. If the patient refuses to give consent for blood sampling and refuses transport, the public safety worker must follow up with their ICO or designee.
 - 2. At this point it is a legal matter to obtain the source patient’s blood for testing (Ohio Revised Code §3701.247).
 - 3. Following a significant exposure in which the source patient refuses to provide a blood sample and refuses transport, the employee should seek immediate medical evaluation and counseling for their selves (MMWR, Sept. 30, 2013).
 - 4. In cases where the patient refuses transport, or in exposure incidents where the source patient is unknown, an exposed employee should follow the steps outlined in [7014.2e - Patients Not Transported to a Hospital](#).
 - 5. EDs or hospitals will not run source patient blood samples if the source patient is not a patient at their hospital.

ci. Source Patient (Transported to Hospital) Results

- i. Hospital-run HIV test results should be available within an hour (may be longer for “stand alone” or smaller EDs); HBV and HCV results may not be available for several days.
- ii. The exposed employee is expected to remain a patient in the ED until they have received the results of the rapid HIV test and any additional counseling from the attending physician.
- iii. The employee is expected to communicate his/her follow-up needs to your department’s ICO or designated supervisor.



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- iv. Written notification of positive test results shall be provided directly to the affected employee by the hospitals designated infection control point of contact within three (3) days after oral notification (Ohio Revised Code §3701.248).
- v. Confidentiality of the source patient and public safety worker information shall be maintained
- vi. Only information pertaining to source patient results will be released to the organization's ICO or designee and/or an employee who is still present in the ED as described above.
- vii. The department ICO or designee and the public safety worker shall not disclose any medical information publicly about the source patient.

e. Patients Not Transported to a Hospital by EMS

- i. Employees should notify their immediate supervisor, and their immediate supervisor should notify the organization's ICO or designee. Federal regulations dictate that, "following report of an exposure, the employer shall make immediately available to the exposed employee a confidential medical evaluation and follow-up" (OSHA 29 CFR, 1910.1030(f) (3)).
- ii. Exposed employee should be directed to any ED for treatment.
- iii. Employee shall locate, complete, and sign the Request for Information by Emergency Care Workers (RIECW) Form (Appendix A), which should be available, completed, and submitted to the nurse handling care in the ED.
- iv. If the public safety worker is aware that the patient went to an ED by other means, the employee's supervisor may call the ED charge nurse of the patient's destination and notify them of the exposure, with a request to obtain baseline testing of the source patient.
- v. The written Request for Notification of Test Results shall be faxed to the ED charge nurse as soon as possible by the employee or the department's ICO.

f. Prophylaxis for Blood/Body Fluid Exposed Public Safety Worker

- i. Post-exposure prophylaxis (PEP) treatment may be offered to the public safety worker by the ED or workplace health provider in accordance with current clinical guidelines and local PEP protocols. Additionally, the employee may wish to consult their personal physician.
 - 1. The decision to take PEP includes a risk-based assessment based on known or unknown source patient and type of exposure.
 - 2. Employees receiving PEP treatment should be followed up within 72 hours of starting treatment.
 - 3. The PEP treatment decision should consider laboratory results when available.
- ii. HIV prophylaxis:
 - 1. Decisions about chemoprophylaxis can be modified if additional information becomes available.
 - 2. Public safety workers must register as ED patients to receive HIV prophylaxis from the hospital.
 - 3. HIV PEP should be started as soon as possible.



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4. Consideration should be given by the ED for expert consultation and guidance on HIV PEP (e.g., infectious disease physician, MMWR, 2011) or the National Clinicians' Post Exposure Prophylaxis Hotline @ #888-448-4911).
 5. Counseling should be made available through the agency's employee assistance program (EAP) or by contractual agreements.
- iii. Hepatitis Prophylaxis
1. Hepatitis Prophylaxis is dependent on the public safety worker's vaccine status.
 2. A small percentage of immunized individual's protection from the vaccine declines over time, which may require Hepatitis B Immunoglobulin (HBIG) and additional doses of the Hepatitis B vaccine to protect against both the current exposure and future exposures.
 3. The results of the HBV Surface Antibody test will demonstrate the employee's immunity to HBV, but are not typically given in the ED as the results of the HBV Surface Antibody test are usually not available immediately.
 4. Employees must follow up with his/her organization's workplace health provider for related prophylaxis as soon as possible.
 5. There is no prophylaxis for HCV at this time. In cases of positive source HCV results, the employee should follow up with their workplace health provider for evaluation and care.
- g. Public Safety Worker Baseline Testing
- i. Baseline testing of the exposed public safety worker is the employee's choice.
 - ii. Agencies should maintain signed statements of employees who decline baseline testing/evaluation at the time of an exposure.
 - iii. Baseline testing is the term given to the set of initial laboratory tests that should be drawn on an exposed employee.
 - iv. This data may be used to compare future assessments in determining if an infectious disease was contracted.
 - v. Baseline testing is not emergent; however, evaluation for PEP as discussed above should be considered urgent and care sought immediately.
 - vi. In cases where PEP was determined not an appropriate emergency treatment, the public safety worker should seek follow up care as instructed.
 - vii. This follow up should be by the organization's workplace health provider. This follow up should optimally occur the next day and no later than seven days post exposure (MMWR, 2001).
 - viii. In cases where the source patient testing is negative but the public safety worker still wants further testing, the employee is encouraged to follow up with their private physician or your department's workplace health provider.
 - ix. Public safety worker baseline testing includes at minimum:
 1. HIV antibody
 2. Hepatitis B surface antibody
 3. Hepatitis C virus antibody



Subject: Infectious Disease Exposure
Reporting Policy

Effective: June 1, 2021

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- x. A positive Hepatitis and/or HIV test of the source patient should trigger viral load testing of the source patient.

7014.3 Respiratory Exposure

a. Definition of A Respiratory Exposure

- i. Respiratory exposure is defined as contamination with an infectious agent through the respiratory tract.
- ii. This occurs via one of two routes (*CDC, Rationale for Isolation Precautions in Hospitals, 1996*):
- iii. Via airborne infectious agents with small-particle residue [5 µm or smaller] of evaporated droplets containing microorganisms that remain suspended in the air for long periods of time (example is tuberculosis, rubella, and varicella virus).
- iv. Via droplet infectious agents which are propelled a short distance (less than three feet) through the air by coughing or sneezing: these droplets are acted upon rapidly by gravity (examples are meningitis, pertussis and influenza).
- v. Respiratory exposures may not be immediately known by the public safety worker, especially if the patient is not overtly symptomatic.

b. Immediate actions of the airborne-exposed public safety worker

- i. Don PPE as soon as possible at the scene or during transport if the patient is known to have a respiratory infection or is coughing or spraying secretions.
- ii. If secretions are splashed or coughed into the eyes or other mucous membranes, flush with copious amounts of IV fluids as soon as possible.
- iii. The public safety worker who suspects or is notified of respiratory exposure:
 - 1. Notify the department ICO that an exposure occurred
 - 2. Notify the ED charge nurse of the exposure upon delivery of the patient
 - 3. Complete the *Request for Notification of Test*.
 - 4. In these cases being checked in as an ED patient may or may not be necessary.
- iv. Upon receipt of the source patient's diagnosis, follow-up care and prophylaxis may be necessary for those exposed.
 - 1. At this point exposed employees may have to return to the receiving hospital and be checked in as a patient to receive care.
 - 2. In other situations follow-up care and prophylaxis may come from your department's workplace health provider.

c. Prophylaxis for The Airborne-Exposed Public Safety Worker

- i. If an exposed employee needs prophylaxis, prophylaxis should be coordinated thru the receiving (or notifying) hospital or when immediately available at the department's workplace health provider's clinic.



Subject: Infectious Disease Exposure
Reporting Policy

Effective: June 1, 2021

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d. Testing The Source Patient

- i. Source testing for respiratory exposures is done by the hospital based on patient symptoms.

e. Source Patient Results

- i. The hospital ICO or designee will notify the department ICO or designee of the infectious agent as soon as possible after symptoms of clinical presentation, or within 48 hours of a positive infectious agent determination.
- ii. Your organization's ICO, possibly after consulting with your department physician, will assess the potential exposure of the employee based on the interaction history with the source patient and the agent involved.
- iii. Confidentiality of source patient and the employee's information shall be maintained.
- iv. Only information pertaining to source patient results will be released to the department's ICO.

7014.4 Blood or Body Fluid & Airborne Exposures by Coroner's Cases

a. Exposure during resuscitation

- i. In cases where there is a public safety worker exposure during resuscitation efforts, it is recommended that crews transport the patient to the hospital where source testing can be performed, rather than follow field termination procedures.
- ii. However, in some incidents, exposure of a public safety worker may occur from a deceased victim who must remain at a scene for a period of time pending a coroner's investigation.

b. Immediate actions of the exposed provider:

- i. Decontaminate self as described in previous sections.
- ii. Notify the department ICO or designee that the exposure occurred.
- iii. At the direction of the department ICO or designee, seek treatment at an ED or at your organization's workplace health provider.
- iv. Consider prophylaxis based on the index of suspicion.

c. Actions of the ICO or designee:

- i. The Coroner or Coroner's Investigator shall be notified as soon as possible by the department's ICO or designee that an exposure has occurred.
- ii. A *Request for Information by Emergency Care Workers* form (Appendix A) shall be forward to the Coroner's Office as soon as possible after notification.

d. Testing the source patient:

- i. The Coroner shall make every effort to test a source patient by the next business day of being notified of the exposure.
- ii. In some cases, the Coroner may elect to send a specimen to an outside lab for testing. The public safety worker shall not wait for testing results from the Coroner to seek medical evaluation.



Subject: **Infectious Disease Exposure
Reporting Policy**

Effective: **June 1, 2021**

Last modified: **Jan. 31, 2021**

e. Source patients test results:

- i. The Coroner or Deputy Coroner shall notify the department ICO or designee of source patient test results as soon as possible.
- ii. Oral notification of source HIV status (positive or negative) shall be provided to the department ICO or designee within two days of test results, and written notification of positive test results shall be provided within three days after oral notification (ORC §3701.248).



Subject: Infectious Disease Exposure
Reporting Policy

Effective: June 1, 2021

Last modified: Jan. 31, 2021

Appendix A

REQUEST NO. 10349

REQUEST FOR INFORMATION
BY EMERGENCY CARE WORKERS

PLEASE PRINT - Use Blue or Black Ink - PRESS HARD

This form is for use by emergency care workers to request information on the presence of a contagious or infectious disease (if known) of a person, alive or dead, who has been treated, handled, or transported for medical care by an emergency care worker.

Before you can be provided with this information, you must believe that you have suffered significant exposure through contact with the person about whom you are requesting the information. A significant exposure means:

- (1) A percutaneous (break in skin or needle stick) or mucous membrane exposure (eyes, nose, mouth) to the blood, semen, vaginal secretions, or spinal, synovial (joint, bone, tendon), pleural (lung), peritoneal (abdomen), pericardial (heart), or amniotic fluid of another person; or
- (2) Exposure to a contagious or infectious disease.

You may expect to receive a reply to this request within 2 days after contagious or infectious disease testing results are known. This may be longer than 2 days after you submit your request. A written notification will follow. Your supervisor will also be informed.

Deposit top (white) copy in designated area or with charge nurse. Submit yellow copy to your agency or employer. Retain pink copy.

The requestor should follow his/her agency's or employer's exposure control plan for post-exposure follow up.

PLEASE PRINT CLEARLY

1. Your Name: _____
2. Your Home Address: _____
City/State/Zip: _____
3. Your telephone number: Home: _____ Work: _____ Pager: _____
4. Have you completed more than two (2) injections in Hepatitis B series. Yes _____ No _____
5. Employer or volunteer agency for whom you were administering health care when exposure occurred:
Employer or Agency: _____
Address: _____
City/State/Zip: _____ Phone: _____
6. Name of your supervisor at above listed place of employment or volunteer agency: _____
7. Regarding the exposure, what was
Name of Source Patient: _____
Date: _____ Time: _____
Place: _____
Manner of exposure:
_____ Dirty Needle Stick _____ Broken Skin Exposure
_____ Splash - Eye, Nose, Mouth _____ Unprotected Mouth to Mouth
Other: Describe the Incident (be specific) _____

This is to attest that the above statements are true and correct to the best of my knowledge and belief.

Your Signature: _____ Date: _____

ACKNOWLEDGEMENT

Name of Health Care Facility/Coroner: _____

Name of Person Receiving Request: _____

Signature of Person Receiving Request: _____

Received: Date _____ Time _____

White: Hospital/Coroner

Yellow: Agency/Employer

Pink: Requestor's Copy



Subject: Infectious Disease Exposure
Reporting Policy

Effective: June 1, 2021

Last modified: Jan. 31, 2021

Appendix B

RESPONSE TO EMERGENCY CARE WORKER REQUEST FOR MEDICAL INFORMATION

REQUEST NO. _____

THIS INFORMATION HAS BEEN DISCLOSED TO YOU FROM CONFIDENTIAL RECORDS PROTECTED FROM DISCLOSURE BY STATE LAW. YOU SHALL MAKE NO FURTHER DISCLOSURE OF THIS INFORMATION WITHOUT THE SPECIFIC, WRITTEN, AND INFORMED RELEASE OF THE INDIVIDUAL TO WHOM IT PERTAINS, OR AS OTHERWISE PERMITTED BY STATE LAW. A GENERAL AUTHORIZATION FOR THE RELEASE OF MEDICAL OR OTHER INFORMATION IS NOT SUFFICIENT FOR THE PURPOSE OF THE RELEASE OF HIV TEST RESULTS OR DIAGNOSES, DISCLOSED ON THIS FORM.

1. Date of oral report: _____ Person giving report: _____
Report given to worker ☐ Supervisor ☐ Supervisor's name _____
Written report will be given to worker and supervisor within 3 working days following oral notification of final results.

2. Date of written report: _____ Person sending report: _____
Report sent to worker ☐ supervisor ☐ Supervisor's name _____

3. Your request for information has been received.

a. _____ The request has been rejected because: _____

Presence of a contagious or infectious disease at this time is unknown due to:

- b. _____ No tests were performed. c. _____ The source person in question has refused HIV testing.
d. _____ Source patient discharged home. e. _____ No blood available
f. _____ Source patient discharged to health care facility/coroner's office/funeral home.

Address of facility/coroner's office/funeral home (if known): _____

g. The following tests were performed on source patient with **negative results**: _____

h. Testing on source person in question was **positive for**: _____

Comments: _____

4. Written and oral report included:

- ☐ Name of disease ☐ (Medical) precautions necessary to prevent transmission
☐ Signs & symptoms of disease ☐ Recommended prophylaxis (if any)
☐ Date of Exposure ☐ Suggested treatment
☐ Incubation period of disease ☐ Appropriate Counseling
☐ Mode of transmission

5. Sources of materials provided regarding disease: _____

6. It is expected that the emergency care worker will consult a physician in cases of true disease exposure. It is understood by provider of report and recipients that decisions related to prophylaxis, treatment, and counseling will be at the discretion of that physician.

**THIS RESPONSE PROVIDES ALL INFORMATION AVAILABLE AS OF THE DATE OF THIS WRITTEN RESPONSE.
ANY ADDITIONAL REQUEST WILL NEED TO BE SUBMITTED FOR ANY FUTURE INFORMATION REGARDING
THIS PATIENT.**

White: Requestor's Copy Yellow: Agency/Employer Pink: Hospital Infection Control Committee/Coroner

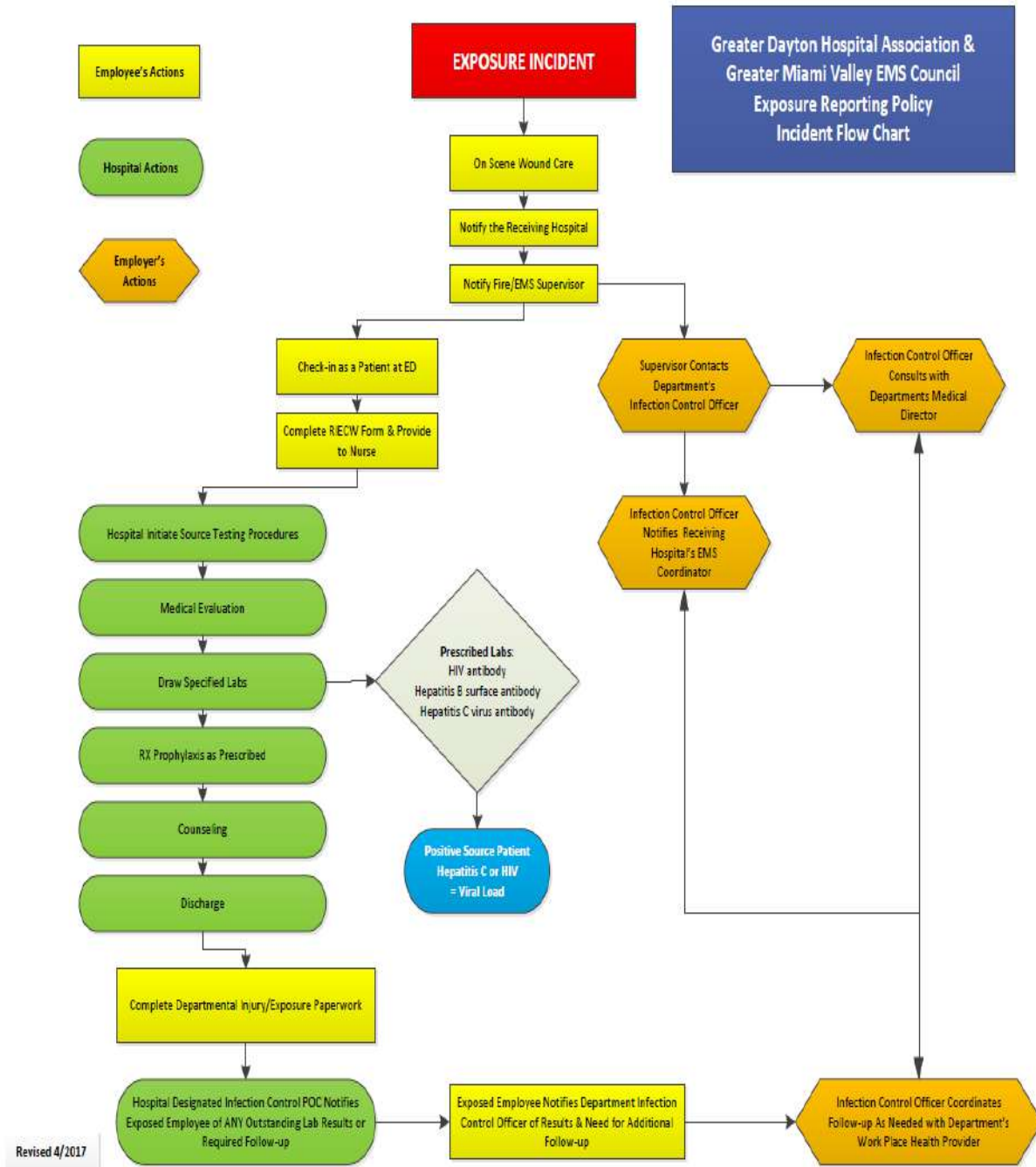


Subject: Infectious Disease Exposure Reporting Policy

Effective: June 1, 2021

Last modified: Jan. 31, 2021

Exposure Incident Flowchart



END OF SECTION



8000 Series

EMS Drug Formulary

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Subject: Adenosine (Adenocard)

Effective: June 1, 2021

Last modified: Oct. 10, 2021

EMR	EMT	AEMT	Paramedic
Packaging	6 mg (1 in drug bag) and 12 mg (2 in drug bag) prefilled syringes		
Indications	Stable Paroxysmal Supraventricular Tachycardia (PSVT)		
Adult Dosing	A 6 mg rapid IV as quickly as possible A If not successful, may repeat 12 mg rapid IV. A If not successful, may repeat 12 mg rapid IV. A All doses of Adenosine are followed by 20 ml bolus of IV fluid. A Go directly to 12 mg if patient with history of PSVT advises it takes 12 mg. May repeat once.		
Pediatric Dosing	P 0.1 mg/kg rapid IV followed by 10 ml rapid saline flush. Max single dose 6 mg. P If unsuccessful, 0.2 mg/kg rapid IV followed by 10 ml rapid saline flush. P Max single dose 12 mg. May repeat x one.		
Therapeutic Action	- Decreases electrical conduction through the AV node without causing negative inotropic effects - Acts directly on SA node to decrease chronotropic activity		
Contraindications	- Second or third degree AV block or sick sinus syndrome - Hypersensitivity to Adenosine		
Precautions And Side Effects	- Lightheadedness, - Paresthesia - Headache - Diaphoresis - Palpitations - Chest pain - Hypotension - Shortness of breath, - Transient periods of sinus bradycardia, sinus pause, or asystole - Ventricular ectopy - Nausea - Metallic taste. - May produce bronchoconstriction in patients with asthma and in patients with bronchopulmonary disease		
Medical Control	- Adult patient: No - Pediatric Patient: No		
Protocols	Cardiac Protocol 2011 - Tachycardia		
END OF SECTION			

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Subject: Albuterol (Proventil)

Effective: June 1, 2021

Last modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	2.5 mg in 3 ml plastic ampule (4 in drug bag)		
Indications	- For the EMT, AEMT and Paramedic: - Exacerbation of Asthma, Emphysema, or COPD - Bronchospasm in Asthma, COPD - Allergic reaction with wheezing - For the Paramedic only: - Hyperkalemia in the presence of Crush Syndrome Trauma		
Adult Dosing	A 2.5 mg (3 ml), nebulized with O₂ at 8-10 LPM. A Combine Ipratropium with first dose of Albuterol. A May repeat Albuterol up to 2 times for a total of 3 doses A Give all 4 doses for hyperkalemia A In Crush syndrome: administer 10 mg nebulized		
Pediatric Dosing	P 2.5 mg (3 ml), nebulized with O₂ at 8-10 LPM. P Combine Ipratropium with first dose of Albuterol. P May repeat Albuterol up to 2 times for a total of 3 doses P In Crush Syndrome: administer 10 mg nebulized		
Therapeutic Action	Bronchodilator		
Contraindications	- Prior hypersensitive reaction to Albuterol - Cardiac dysrhythmias associated with tachycardia.		
Precautions And Side Effects	- Once initiated, the patient should be removed by EMS. - Side Effects - Restlessness - Apprehension - Dizziness - Palpitations - Tachycardia - Dysrhythmias - May precipitate angina pectoris		
Medical Control	Adults: No Pediatrics: No		
Protocols	General Protocol 1008 – Advanced Airway Management Trauma Protocol 3007 – Crush Syndrome Trauma (Paramedic only) Medical Protocol 4002 – Allergic Reaction/Anaphylaxis Medical Protocol 4003 – Asthma/Emphysema/COPD		
END OF SECTION			

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Subject: Amiodarone (Cordarone)

Effective: June 1, 2021

Last modified: Oct. 10, 2021

EMR	EMT	AEMT	Paramedic
Packaging	150 mg in 3 ml vial, 50 mg/ml 3 vials in drug bag		
Indications	- Ventricular Fibrillation or Pulseless Ventricular Tachycardia - Stable Wide-Complex Tachycardia		
Adult Dosing	<u>Ventricular Fibrillation or Pulseless Ventricular Tachycardia</u> A 300 mg IV or IO. A May repeat with half the initial dose (150 mg IV or IO) no sooner than 10 minutes after first dose. <u>If patient converts with ROSC from a ventricular arrhythmia and no anti-arrhythmic has been given:</u> A 150 mg in 250 ml NS, IV wide open over 10 minutes using 60 gtt/ml tubing & 18 g angiocath <u>Stable Wide-Complex Tachycardia:</u> A 150 mg in 250 ml NS, IV wide open over 10 minutes using 60 gtt/ml tubing & 18 g angiocath		
Pediatric Dosing	<u>Ventricular Fibrillation or Pulseless Ventricular Tachycardia</u> P 5 mg/kg IV or IO (max first dose 300 mg). P May repeat 5 mg/kg IV or IO no sooner than 10 minutes after first dose. - Max repeat dose is 150 mg - Not indicated for stable wide complex tachycardia		
Therapeutic Action	Antidysrhythmic agent with multiple mechanisms of action		
Contraindications	- Pulmonary congestion - Cardiogenic shock - Hypotension (SBP less than 100) - Sensitivity to Amiodarone		
Precautions And Side Effects	- Continuous EKG monitoring is required. - Side Effects - Hypotension - Headache - Dizziness - Bradycardia - AV conduction abnormalities - Flushed skin - Abnormal salivation		
Medical Control	Adult patient: No Pediatric Patient: No		
Protocols	Cardiac Protocol 2005 – Cardiac Arrest: Ventricular Fib or Pulseless V-Tach Cardiac Protocol 2011 – Tachycardia		
END OF SECTION			

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Subject: Aspirin

Effective: June 1, 2021

Last Modified: Oct. 10, 2021

EMR		EMT	AEMT	Paramedic
Packaging	81mg tablets in a blister pack (4 tablets total)			
Indications	Given as soon as possible to the patient with AMI.			
Adult Dosing	A 324 mg chewed (Four 81 mg tablets)			
Pediatric Dosing	P Not applicable to pediatric patients			
Therapeutic Action	Anti-platelet			
Contraindications	- Hypersensitivity to salicylates - Active ulcer disease - Bleeding disorders - Third trimester pregnancy			
Precautions And Side Effects	- Suspected cardiac chest pain must be at least 25 years old. - Patient <u>must</u> chew the tablets - Side Effects - Stomach irritation - Heartburn or indigestion - Nausea or vomiting - Allergic reactions			
Medical Control	Adult patient: For AEMT and Paramedic: No, unless patient is 25 y/o or younger with AMI symptoms. For EMTs: Yes Pediatric Patient: Not applicable			
Protocols	Cardiac Protocol 2008 – Suspected Cardiac Chest Pain Medical Protocol 4011 – Obstetrical Emergencies			
END OF SECTION				

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Subject: Atropine

Effective: June 1, 2021

Last modified: Dec. 6, 2024

EMR	EMT	AEMT	Paramedic
Packaging	1mg in 10 ml prefilled syringe; (3 in drug bag) - In Haz Mat/WMD Security Bag: - Duodote: 2 mg auto-injector (along with 2-Pam 600 mg autoinjector) - In WMD Drug Caches and Chempacks: 2 mg, 1mg and 0.5 mg AtroPen auto-injectors; - Multidose vial 8 mg in 20 ml, 0.4 mg/ml		
Indications	- Symptomatic bradycardia - Organophosphate or Nerve Agent poisoning (regardless of cardiac rate)		
Adult Dosing	<u>Bradycardia:</u> A 1 mg IV up to 3 mg <u>Organophosphate or Nerve Gas poisoning:</u> A ♦ EMR, EMT, AEMT, Paramedic: 2 mg Duodote auto-injector. Paramedic only: 2 mg IV, IO or IM A No max dose, given every 5 min or until lungs are clear to auscultation.		
Pediatric Dosing	<u>Bradycardia:</u> P 0.02 mg/kg IV or IO every 5 min. P Minimum single dose of 0.1 mg, max single dose 0.5 mg P Maximum total dose 1 mg <u>Organophosphate or Nerve Gas poisoning:</u> P For EMR, EMT, AEMT or Paramedic: P ♦ Less than 20 kgs: 0.5 mg AtroPen auto-injector P ♦ 20 - 40 kgs: 1.0 mg AtroPen auto-injector P ♦ Greater than 40 kgs: 2.0 mg AtroPen auto-injector P Paramedic only: ♦ May give atropine doses listed IV or IM P No max dose, given every 5 minutes or until lungs are clear to auscultation.		
Therapeutic Action	Anticholinergic		
Contraindications	- None for severe organophosphate exposure. - Tachycardia - Hypersensitivity to atropine - Obstructive disease of GI tract - Obstructive neuropathy - Unstable cardiovascular status in acute hemorrhage with myocardial ischemia - Narrow angle glaucoma - Thyrotoxicosis		
Precautions And Side Effects	- EMR, EMT and AEMT can <u>only</u> administer the Duodote auto-injector to Organophosphate or Nerve Agent patients - Pupillary dilation rendering the pupils nonreactive. Pupil response may not be useful in monitoring CNS status. - Side Effects - Dysrhythmias, tachycardia, palpitations - Paradoxical bradycardia when pushed too slowly or when used at doses less than 0.5 mg - Headache or dizziness - Anticholinergic effects (dryness, photophobia, blurred vision, urinary retention, constipation) - Nausea and vomiting - Flushed, hot, dry skin - Allergic reactions.		
Medical Control	Adult patient: Bradycardia —No, Organophosphate Nerve Agent Poisoning—Yes Pediatric Patient: Bradycardia—No, Organophosphate Nerve Agent Poisoning—Yes		
Protocols	Cardiac Protocol 2010 – Bradycardia Special Operations Protocol 6002 – Antidote Resources Special Operations Protocol 6005 – Organophosphate or Nerve agent Exposure		
END OF SECTION			

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Subject: Calcium Chloride 10%

Effective: June 1, 2021

Last Modified: Oct. 10, 2021

EMR	EMT	AEMT	Paramedic
Packaging	1 gram in 10 ml vial, 100 mg/ml (1 in drug bag)		
Indications	- Renal dialysis patient in cardiac arrest or with ♦ bradycardia - Calcium Channel Blocker OD ♦ Hydrofluoric Acid exposure with tetany <u>or</u> cardiac arrest. - Tetany may present as: overactive neurological reflexes, spasms of the hands and feet, cramps, and laryngospasm. - May be given prophylactically, after exposure to high concentration (> 40%) Hydrofluoric Acid ♦ Adults with Crush Syndrome presenting with abnormal ECG or hemodynamic instability		
Adult Dosing	A 1 gm (10 ml) IV for: - Cardiac arrest in renal dialysis patients ♦ Calcium Channel Blocker OD ♦ Hydrofluoric Acid exposure with tetany or cardiac arrest A ♦ For prophylaxis in high concentration Hydrofluoric Acid exposure: 400 mg (4 ml) IV A ♦ Renal dialysis patient with bradycardia: 1 gm (10 ml) IV A ♦ Crush syndrome: 1 gm (10 ml) IV		
Pediatric Dosing	P 20 mg/kg IV (max dose 500 mg) for: - Cardiac arrest in renal dialysis patients ♦ Calcium Channel Blocker OD P ♦ Call in advance to treat crush syndrome or hydrofluoric acid exposures in pediatric patients		
Therapeutic Action	- Antagonizes cardiac toxicity in hyperkalemia associated with dialysis patients. - Reverses symptoms of Calcium Channel Blocker		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Do not administer with Sodium Bicarbonate because if mixed, a precipitate develops. - Flush tubing between drugs. - Side Effects: - Bradycardia (may cause asystole) - Hypotension - Metallic taste - Severe local necrosis and sloughing following IV infiltration - May produce vasospasm in coronary and cerebral arteries - Hypertension and bradycardia may occur with rapid administration.		
Medical Control	Adults: - Cardiac Arrest—No - Renal dialysis patient in bradycardia---Yes - Calcium Channel Blocker OD—Yes - Hydrofluoric Acid Exposure—Yes - Crush syndrome—Yes Pediatrics - Arrest—No - Calcium Channel Blocker OD—Yes - Hydrofluoric Acid Exposure—Yes - Crush syndrome--Yes		
Protocols	Cardiac Protocol 2004 – Cardiac Arrest - Renal Failure/Dialysis Cardiac Protocol 2010 – Bradycardia Trauma Protocol 3007 – Crush Syndrome Trauma Medical Protocol 4012 – Overdose or Poisoning Special Operations Protocol 6004 – Hydrofluoric Acid Exposure		
END OF SECTION			

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Subject: Calcium Gluconate

Effective: June 1, 2021

Last Modified: Oct. 10, 2021

EMR	EMT	AEMT	Paramedic
Packaging	1 gram in 10 ml vial, 100 mg/ml. Only in the drug bag in the event of Calcium Chloride 10% shortage		
Indications	- Renal dialysis patient in cardiac arrest or with ♦ bradycardia - Calcium Channel Blocker OD ♦ Hydrofluoric Acid exposure with tetany <u>or</u> cardiac arrest. - Tetany may present as: overactive neurological reflexes, spasms of the hands and feet, cramps, and laryngospasm. - May be given prophylactically, after exposure to high concentration (> 40%) Hydrofluoric Acid ♦ Adults with Crush Syndrome presenting with abnormal ECG or hemodynamic instability		
Adult Dosing	A 1 gm (10 ml) IV for: - Cardiac arrest in renal dialysis patients ♦ Calcium Channel Blocker OD ♦ Hydrofluoric Acid exposure with tetany or cardiac arrest A ♦ For prophylaxis in high concentration Hydrofluoric Acid exposure: 400 mg (4 ml) IV A ♦ Renal dialysis patient with bradycardia: 1 gm (10 ml) IV A ♦ Crush syndrome: 1 gm (10 ml) IV		
Pediatric Dosing	P 20 mg/kg IV (max dose 500 mg) for: - Cardiac arrest in renal dialysis patients ♦ Calcium Channel Blocker OD P ♦ Call in advance to treat crush syndrome or hydrofluoric acid exposures in pediatric patients		
Therapeutic Action	- Antagonizes cardiac toxicity in hyperkalemia associated with dialysis patients. - Reverses symptoms of Calcium Channel Blocker		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Do not administer with Sodium Bicarbonate because if mixed, a precipitate develops. - Flush tubing between drugs. - Side Effects: - Bradycardia (may cause asystole) - Hypotension - Metallic taste - Severe local necrosis and sloughing following IV infiltration - May produce vasospasm in coronary and cerebral arteries - Hypertension and bradycardia may occur with rapid administration.		
Medical Control	Adults: - Cardiac Arrest—No - Renal dialysis patient in bradycardia---Yes - Calcium Channel Blocker OD—Yes - Hydrofluoric Acid Exposure—Yes - Crush syndrome—Yes Pediatrics - Arrest—No - Calcium Channel Blocker OD—Yes - Hydrofluoric Acid Exposure—Yes - Crush syndrome--Yes		
Protocols	Cardiac Protocol 2004 – Cardiovascular Emergencies: Renal Failure/Dialysis Cardiac Protocol 2010 – Bradycardia Trauma Protocol 3007 – Crush Syndrome Trauma Medical Protocol 4012 – Overdose or Poisoning Special Operations Protocol 6004 – Hydrofluoric Acid Exposure		
END OF SECTION			

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Subject: Ciprofloxacin (Cipro)

Effective: June 1, 2021

Last Modified: Feb. 20, 2024

EMR	EMT	AEMT	Paramedic
Packaging	Tablets		
Indications	As prophylaxis against Anthrax, Cholera or Plague		
Adult Dosing	A ♦ 500 mg tablet by mouth, twice a day		
Pediatric Dosing	P ♦ Dosage will be specified at time of incident.		
Therapeutic Action	Antibiotic		
Contraindications	- Allergy to quinolones - Tendon pain or inflammation - Pediatrics - Pregnancy		
Precautions And Side Effects	- Side Effects - Atrial flutter - Hypotension - Premature Ventricular Contractions - QT prolongation - Torsade De Pointes - Tendon pain/inflammation		
Medical Control	- Adult: Yes - Pediatric: Yes		
Protocols	Special Operations Protocol 6006 – Other Hazardous Materials		
END OF SECTION			

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Subject: Dextrose 10% (D10)

Effective: June 1, 2021

Last Modified: Dec. 6, 2024

EMR	EMT	AEMT	Paramedic
Packaging	500 ml of D10W, contains 50 g Dextrose 1 bag of solution in drug bag		
Indications	- Diabetic with mental status changes - Evidence of hypoglycemia in cardiac arrest - Generalized hypothermia with or without arrest - Altered level of consciousness of unknown cause - Seizures with BGL of less than 60 mg/dl - No blood sugar monitor is available or suspicion of hypoglycemia despite glucometer readings.		
Adult Dosing	A 250 ml IV at wide open rate A May repeat in 10 minutes if patient fails to respond or BGL remains less than 60 mg/dl. A Maximum dose is 500 ml.		
Pediatric Dosing	- Pediatric patients: P 5 ml/kg P Maximum dose is 250 ml - Newborn patients: P 2 ml/kg if BGL is less than 40 mg/dl		
Therapeutic Action	Principal form of carbohydrate utilized by the body		
Contraindications	Known or suspected CVA in the absence of hypoglycemia		
Precautions And Side Effects	- May precipitate severe neurologic symptoms in thiamine deficient patients <u>Side Effects:</u> - Warmth - Pain - Hyperglycemia - Burning from medication infusion - Thrombophlebitis		
Medical Control	Adults: No Pediatrics: No		
Protocols	Medical Protocol 4008 – Diabetic Emergencies - Hypoglycemia Pediatric Considerations 5002 – Newborn Care and Resuscitation		
END OF SECTION			

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Subject: Dextrose 25% (D25) (JITSO)

Effective: June 1, 2021

Last Modified: Jan. 8, 2025

EMR	EMT	AEMT	Paramedic
Packaging	10 ml prefilled syringe of Dextrose 25%, contains 2.5 g Dextrose 1 in drug bag in the absence of Dextrose 10%		
Indications	- Diabetic with mental status changes - Evidence of hypoglycemia in cardiac arrest - Generalized hypothermia with or without arrest - Altered level of consciousness of unknown cause - Seizures with BGL of less than 60 mg/dl - No blood sugar monitor is available or suspicion of hypoglycemia despite glucometer readings.		
Adult Dosing	Not indicated in adult patients		
Pediatric Dosing	- For pediatric patients greater than 25 kg, see 8011 Dextrose 50% - Pediatric patients at or less than 25 kg: P 2 ml/kg of Dextrose 25% (D25) - Newborn patients: P 2 ml/kg of Dextrose 25% (D25) diluted in equal volume of saline solution P Only administer if BGL is less than 40 mg/dl ♦ If no Dextrose 25% (D25) found in the drug bag, contact Medical Control for advice		
Therapeutic Action	Principal form of carbohydrate utilized by the body		
Contraindications	Known or suspected CVA in the absence of hypoglycemia		
Precautions And Side Effects	- May precipitate severe neurologic symptoms in thiamine deficient patients <u>Side Effects:</u> - Warmth - Pain - Hyperglycemia - Burning from medication infusion - Thrombophlebitis		
Medical Control	Adults: Not indicated in adult patients Pediatrics: No, unless no Dextrose 25% (D25) found in the drug bag		
Protocols	Medical Protocol 4008 – Diabetic Emergencies - Hypoglycemia Pediatric Considerations 5002 – Newborn Care and Resuscitation		
END OF SECTION			

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Subject: Dextrose 50% (D50) (JITSO)

Effective: June 1, 2021

Last Modified: Jan. 8, 2025

EMR	EMT	AEMT	Paramedic
Packaging	50 ml prefilled syringe of Dextrose 50%, contains 25 g Dextrose 2 in drug bag in the absence of Dextrose 10%		
Indications	- Diabetic with mental status changes - Evidence of hypoglycemia in cardiac arrest - Generalized hypothermia with or without arrest - Altered level of consciousness of unknown cause - Seizures with BGL of less than 60 mg/dl - No blood sugar monitor is available or suspicion of hypoglycemia despite glucometer readings.		
Adult Dosing	A One 50 ml (25 gm) IVP May repeat in 10 minutes if patient fails to respond or BGL remains less than 60 mg/dl.		
Pediatric Dosing	- Pediatric patients greater than 25 kg: P 1 ml/kg of Dextrose 50% (D50) - For pediatric patients at or less than 25 kg, see 8010 Dextrose 25% or P 1 ml/kg of Dextrose 50% (D50) diluted in equal volume of IV solution (if no D25 present)		
Therapeutic Action	Principal form of carbohydrate utilized by the body		
Contraindications	Known or suspected CVA in the absence of hypoglycemia		
Precautions And Side Effects	- May precipitate severe neurologic symptoms in thiamine deficient patients <u>Side Effects:</u> - Warmth - Pain - Hyperglycemia - Burning from medication infusion - Thrombophlebitis		
Medical Control	Adults: No Pediatrics: No		
Protocols	Medical Protocol 4008 – Diabetic Emergencies - Hypoglycemia Pediatric Considerations 5002 – Newborn Care and Resuscitation		
END OF SECTION			

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Subject: Diazepam (Valium) (JITSO) & Carpuject

Effective: June 1, 2021

Last Modified: Jan. 16, 2025

EMR	EMT	AEMT	Paramedic
Packaging	• Vial for AEMT and Paramedic only ○ 10 mg in 2 ml vial (5 mg/1ml) ○ One vial present in the drug bag in the event of Midazolam shortage • WMD Drug Cache & CHEMPACK resource for all certification levels • Convulsive Antidote, Nerve Agent (CANA) 10 mg carpuject		
Indications	• Vial for AEMT and Paramedic only ○ Seizures - A Chest pain associated with stimulant overdose (adults only) • CANA carpujects for all certifications • Seizures associated with Organophosphate or Nerve Agent event		
Adult Dosing	• Vial for AEMT and Paramedic only - A Seizures: 5 mg slow IV; may repeat dose once. - A Cocaine or crack use: 5 mg slow IV, may repeat dose once. • CANA carpujects for all certifications - A 10 mg IM • For patients 70 y/o or older, reduce dosing for sedatives and analgesics to one half (½) of the adult doses		
Pediatric Dosing	• Vial for AEMT and Paramedic • Seizures: - P 0.2 mg/kg slow IV over 2 min. (maximum dose 5 mg IV) - or - P 0.5 mg/kg rectally, (maximum dose 10 mg rectally) - P May repeat 0.2 mg/kg slow IV over 2 min (maximum 5 mg) • CANA carpujects for all certifications - P 10 mg IM by auto-injector		
Therapeutic Action	• Treats alcohol withdrawal and grand mal seizure activity • Used to treat anxiety and stress.		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• Side Effects: ○ Hypotension ○ Reflex tachycardia (rare) ○ Respiratory depression ○ Ataxia ○ Psychomotor impairment ○ Confusion ○ Nausea ○ May cause local venous irritation		
Medical Control	• Vial for AEMT and Paramedic only ○ Adults: No ○ Pediatrics: No • CANA carpujects for all certifications ○ Adults: Yes ○ Pediatrics: Yes		
Protocols	• Trauma Protocol 3008 – Cyanide Poisoning • Medical Protocol 4012 – Overdose/Poisoning • Special Operations Protocol 6002 – Antidote Resources • Special Operations Protocol 6005 – Organophosphate or Nerve Agent Exposure		
END OF SECTION			

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Subject: Diphenhydramine (Benadryl)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	50 mg in 1ml vial		
Indications	- Allergic reaction or Anaphylaxis - In anaphylaxis, for the patient who goes into cardiac arrest if not previously given - Extrapyramidal reaction		
Adult Dosing	A 50 mg IM or slow IV		
Pediatric Dosing	P 1 mg/kg (max dose 50 mg) IM or slow IV		
Therapeutic Action	Prevents the physiologic actions of histamine by blocking histamine receptors		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Use cautiously in patients with CNS depression or lower respiratory diseases such as asthma. - Side Effects: - Dose related drowsiness - Sedation - Disturbed coordination - Hypotension - Palpitations, tachycardia or bradycardia - Thickening of bronchial secretions - Dry mouth and throat		
Medical Control	- Adults: No, for the Paramedic. Yes, for the AEMT when treating Extrapyramidal Reactions - Pediatrics: No, for the Paramedic. Yes, for the AEMT when treating Extrapyramidal Reactions		
Protocols	- Medical Protocol 4002 – Allergic Reactions/Anaphylaxis - Medical Protocol 4010 – Extrapyramidal (Dystonic) Reactions		
END OF SECTION			

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Subject: Dopamine (JITSO)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	- Premixed 250 ml bag (400 mg/250 ml) - Concentration: 1600 mcg/ml - Only present in the drug bag in the event of Norepinephrine shortage		
Indications	Shock with or without Pulmonary Edema		
Adult Dosing	A IV drip rate, 5 to 20 mcg/kg/min of 400 mg/250 ml; increase by increments of 5 mcg/kg/min .		
Pediatric Dosing	P IV drip rate, 5 to 20 mcg/kg/min of 400 mg/250 ml; start at 5 mcg/kg/min . P Titrate to maintain adequate perfusion		
Therapeutic Action	- Acts on alpha, beta and dopaminergic receptors in dose dependent fashion - Increases cardiac output in higher doses		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Correct hypovolemia prior to using Dopamine. - Infuse through large stable vein to avoid possibility of extravasation injury. - Side Effects: - Dose related tachydysrhythmias - Hypertension - Increased myocardial oxygen demand (ischemia)		
Medical Control	- Adults: No - Pediatrics: No		
Protocol	- As a replacement for Norepinephrine: - Cardiac Protocol 2009 – Cardiac Alert Program - Medical Protocol 4015 – Sepsis - Medical Protocol 4016 – Shock		
END OF SECTION			

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Subject: Doxycycline

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Tablets		
Indications	• As prophylaxis against Anthrax, Cholera or Plague		
Adult Dosing	A ♦ 100 mg tablet by mouth, twice a day		
Pediatric Dosing	P ♦ Dosage will be specified at time of incident		
Therapeutic Action	• Antibiotic		
Contraindications	• Pregnancy • Allergies to Tetracycline antibiotics		
Precautions And Side Effects	• <u>Side Effects:</u> ○ May make birth control pills less effective ○ Use with caution in patients with liver disease, kidney disease and asthma ○ Can cause headache, blurred vision and flu-like symptoms		
Medical Control	• Adults: Yes • Pediatrics: Yes		
Protocols	• Special Operations Protocol 6006 – Other Hazardous Materials		
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Subject: Duodote

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Auto-injector Atropine 2 mg and Pralidoxime Chloride (2-Pam) 600 mg • In WMD Drug Caches and CHEMPACKS		
Indications	• Organophosphate or Nerve Agent poisoning		
Adult Dosing	A ♦ Single auto-injector containing Atropine 2 mg and 2-Pam 600 mg		
Pediatric Dosing	P ♦ Single auto-injector containing Atropine 2 mg and 2-Pam 600 mg		
Therapeutic Action	• Anticholinergic as a result of WMD MCI; also reactivates cholinesterase.		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• Use with caution in myasthenia gravis, renal impairment, pregnancy, lactation or children. • Atropine causes pupillary dilation rendering the pupils nonreactive. Pupil response may not be useful in monitoring CNS status. • <u>Side Effects:</u> ○ Tachycardia ○ Paradoxical bradycardia when pushed too slowly or when used at doses less than 0.5 mg ○ Palpitations or dysrhythmias ○ Headache ○ Dizziness ○ Anticholinergic effects (dry mouth, nose, skin, photophobia, blurred vision, urinary retention, constipation) ○ Nausea & vomiting ○ Flushed, hot, dry skin ○ Allergic reactions		
Medical Control	• Adults: Yes • Pediatrics: Yes		
Protocols	• Special Operations Protocol 6005 – Organophosphate or Nerve Agent Exposure		
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Subject: Epinephrine Auto-Injector

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	- EpiPen auto-injector: 0.3 mg (one in BLS Only small drug bag) - EpiPen Jr. auto-injector: 0.15 mg (one in BLS Only small drug bag)		
Indications	- For the EMR, EMT, AEMT and Paramedic: - Anaphylaxis or allergic reaction - The EMR and the EMT cannot treat Asthma with Epinephrine		
Adult Dosing	- Asthma (AEMT and Paramedic) or Anaphylaxis (EMR, EMT, AEMT and Paramedic) - A If equal to or greater than 30 kg, give both Adult EpiPen 0.3 mg and EpiPen Jr 0.15 mg - A May repeat after 10 minutes		
Pediatric Dosing	- Asthma (AEMT and Paramedic) or Anaphylaxis (EMR, EMT, AEMT and Paramedic) - P If less than 15 kg, EpiPen Jr 0.15 mg - P If equal to or greater than 15 kg and less than 30 kg, Adult EpiPen 0.3 mg - P If greater than 30 kg, give both Adult EpiPen 0.3 mg and EpiPen Jr 0.15 mg - P May repeat after 10 minutes		
Therapeutic Action	- Directly stimulates alpha and beta adrenergic receptors in dose-related fashion - Causes bronchodilation, vasoconstriction, and increased cardiac output.		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- While the protocol lists those patients under 15 kg as pediatric, it is understood that patients equal to or greater than 30 kg will get both the Adult EpiPen and the EpiPen Jr., no matter what their age. - Epinephrine dosing for Asthma, Allergies and Anaphylaxis is based on weight, not age - Side Effects - Headache - Nausea - Restlessness - Weakness - Dysrhythmias, including ventricular tachycardia and ventricular fibrillation - Hypertension - Tachycardia - May increase myocardial oxygen demand or precipitation of angina pectoris - Syncope has occurred following epinephrine administration to asthmatic children.		
Medical Control	Adults: Initial dose administration at all levels and follow-up dosing for AEMT and Paramedics – No In allergies/anaphylaxis, repeat doses by EMR/EMTs - Yes Pediatrics: Initial dose administration at all levels and follow-up dosing for AEMT and Paramedics – No In allergies/anaphylaxis, repeat doses by EMR/EMTs - Yes		
Protocols	Medical Protocol 4002 – Allergic Reactions/Anaphylaxis Medical Protocol 4003 – Asthma/Emphysema/COPD		
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Subject: Epinephrine 1:1,000

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	1:1,000 – 1mg/ml ampule (One in BLS compartment)		
Indications	- For the EMT, AEMT and Paramedic: - Anaphylaxis or allergic reaction - For the AEMT and Paramedic: - Asthma in severe distress - The EMT cannot treat Asthma with Epinephrine {The EMT may only administer 1:1,000 IM after authorization and training from their medical director}		
Adult Dosing	- Asthma (AEMT or Paramedic) or anaphylaxis ({EMT}, AEMT and Paramedic) - A If equal to or greater than 30 kg, Epinephrine (1:1,000) 0.5 mg IM - A May repeat in 10 minutes		
Pediatric Dosing	- Asthma (AEMT and Paramedic) or Anaphylaxis ({EMT}, AEMT and Paramedic) - P If equal to or greater than 30 kg, Epi (1:1,000) 0.5 mg IM - P If equal to or greater than 15 kg and less than 30 kg, Epi (1:1,000) 0.3 mg IM - P If less than 15 kg, Epi (1:1,000) 0.15 mg IM - P May repeat Epi (1:1,000) (at weight appropriate dose) IM after 10 min.		
Therapeutic Action	- Directly stimulates alpha and beta adrenergic receptors in dose-related fashion - Causes bronchodilation, vasoconstriction, and increased cardiac output.		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- While the protocol lists those patients under 15 kg as pediatric, it is understood that patients equal to or greater than 30 kg will get 0.5 mg IM., no matter what their age. - Epinephrine dosing for Asthma, Allergies and Anaphylaxis is based on weight, not age. - Side Effects - Headache - Nausea - Restlessness - Weakness - Dysrhythmias, including ventricular tachycardia and ventricular fibrillation - Hypertension - Tachycardia - May increase myocardial oxygen demand or precipitation of angina pectoris - Syncope has occurred following epinephrine administration to asthmatic children.		
Medical Control	Adults: Initial dose administration at all levels and follow-up dosing for AEMT and Paramedics – No In allergies/anaphylaxis, repeat doses by EMR/EMTs - Yes Pediatrics: Initial dose administration at all levels and follow-up dosing for AEMT and Paramedics – No In allergies/anaphylaxis, repeat doses by EMR/EMTs - Yes		
Protocols	Medical Protocol 4002 – Allergic Reactions/Anaphylaxis Medical Protocol 4003 – Asthma/Emphysema/COPD		
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Subject: Epinephrine 1:10,000

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	1:10,000 – 1 mg/10ml prefilled syringes (six in drug bag)		
Indications	- For the Paramedic only: - Ventricular Fibrillation, Pulseless Ventricular Tachycardia, Asystole, and PEA in all patients - Bradycardia in pediatric patients		
Adult Dosing	Ventricular Fibrillation, Pulseless Ventricular Tachycardia, Asystole, and PEA (Paramedic) A 1 mg (1:10,000) IV, repeat every 3-5 minutes		
Pediatric Dosing	Ventricular Fibrillation, Pulseless Ventricular Tachycardia, Asystole, PEA, and Bradycardia P 0.01 mg/kg (1:10,000) IV; repeat every 3-5 minutes (max single dose 1 mg)		
Therapeutic Action	- Directly stimulates alpha and beta adrenergic receptors in dose-related fashion - Causes bronchodilation, vasoconstriction, and increased cardiac output.		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Side Effects - Headache - Nausea - Restlessness - Weakness - Dysrhythmias, including ventricular tachycardia and ventricular fibrillation - Hypertension - Tachycardia - May increase myocardial oxygen demand or precipitation of angina pectoris - Syncope has occurred following epinephrine administration to asthmatic children.		
Medical Control	- Adults: No - Pediatrics: No		
Protocols	Cardiac Protocol 2003 – Cardiac Arrest: Asystole or PEA Cardiac Protocol 2005 – Cardiac Arrest: V-Fib or Pulseless V-Tach Cardiac Protocol 2010 – Bradycardia Pediatric Considerations 5002 – Newborn Care and Resuscitation Special Operations Protocol 6004 – Hydrofluoric Acid Exposure		
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Subject: Etomidate

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	40 mg in 20 ml vial (2 mg/ml)		
Indications	To provide sedation prior to {Sedate-to-Intubate} or {Rapid Sequence Intubation} procedures		
Adult Dosing	A 0.3 mg/kg IV A May repeat within 2 minutes if patient resistant to intubation. A Average dose is 15 mg - 25 mg		
Pediatric Dosing	Not applicable to pediatric patients		
Therapeutic Action	- Short-acting, potent sedative - Hypnotic		
Contraindications	- Hypersensitivity - Not to be administered to pediatric patients		
Precautions And Side Effects	- Must be authorized for use by the agencies’ Medical Director - Side Effects: - Bradycardia - Respiratory depression or tachypnea - Sinus tachycardia - Hypotension - Nausea and vomiting		
Medical Control	- Adults: No - Pediatrics: Not applicable		
Protocols	General Protocol 1010 – {Sedate to Intubate and Rapid Sequence Intubation}		
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Subject: Fentanyl (Sublimaze)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	100 mcg/2 mL (50 mcg/ml) vial - One in drug bag		
Indications	- Suspected Cardiac Chest Pain - Pain associated with traumatic events - Extremity Fractures - Dislocations or Sprains - Frostbite - Abdominal Pain - Hydrofluoric Acid (Hf) exposure		
Adult Dosing	A 50-100 mcg slow IV, provided SBP is greater than 100. A May repeat 50-100 mcg slow IV, after 5 minutes provided SBP greater than 100. A If no IV, Fentanyl 50-100 mcg IN, SQ or IM A May repeat Fentanyl 50-100 mcg IN, SQ or IM after 10 minutes G Patient greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult dose		
Pediatric Dosing	P Fentanyl is <u>not</u> to be administered to anyone less than 2 years of age. P If unable to obtain a blood pressure, look for evidence adequate perfusion prior to administration. ♦ Contact MCP prior to treatment of abdominal pain - First choice treatment for pain: P 1 mcg/kg IN, max dose 100 mcg., provided age appropriate normal SBP (80 + 2x age in years) P Repeat 1 mcg/kg IN after 10 minutes, if an additional drug bag is available. - Second choice treatment for pain: P 1 mcg/kg, slow IV, max dose 100 mcg, P Repeat 1 mcg/kg, slow IV after 5 minutes, max dose 100 mcg P Maintain age appropriate blood pressure - If unable to obtain IV: SQ or IM for pediatric patients is a last resort P 1 mcg/kg SQ or IM, max dose 100 mcg P Repeat 1 mcg/kg SQ or IM, max dose 100 mcg, no sooner than 10 minutes after first dose.		
Therapeutic Action	- Provides analgesia - Reduces cardiac preload by increasing venous capacitance and decreasing afterload		
Contraindications	Hypersensitivity		
Precautions And Side Effects	- Chest wall rigidity ("wooden chest syndrome") may occur: - Prevents adequate chest wall excursion and ventilation. - Typically occurs with high doses (6-7 mcg/kg) or with rapid administration. - Reversible with naloxone. - Provide continuous cardiac monitoring, EtCO₂ and pulse oximetry with sedated patients. - Geriatric & debilitated patients require lower doses & are more prone to side effects. - Apnea - CNS depression - Bradycardia which may be transient. - Ensure adequate ventilation and oxygenation first. - Atropine only if bradycardia is symptomatic and hemodynamically significant. - For the Paramedic, follow bradycardia protocol.		
Medical Control	- Adults: No - Pediatrics: Yes, for abdominal pain		
Protocols	General Protocol 1014 – Pain Management Cardiac Protocol 2006 – AICD Activations Cardiac Protocol 2008 – Suspected Cardiac Chest Pain Cardiac Protocol 2009 – Cardiac Alert Program		
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Subject: Hydroxocobalamin (Cyanokit)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	1 vial, containing 5 g lyophilized Hydroxocobalamin dark red crystalline powder for injection. - After reconstitution with 200 ml fluid, the vial contains Hydroxocobalamin for injection, 25 mg/mL. - Available in caches located in each county in Homeland Security Region 3.		
Indications	- Known or strongly suspected cyanide intoxication - Smoke inhalation with suspected cyanide component. - Victim exposed to fire or smoke who presents with altered mental status, seizures, shock, or difficulty breathing. - To reconstitute follow package directions: - Place the vial in an upright position. - Add 200 mL of NS or LR to the vial using the transfer spike. Fill to the line. - Mix: The vial should be inverted or rocked, not shaken, for at least 1 min. before infusion. - Infuse Vial: Use vented intravenous tubing, hang and infuse over 15 minutes.		
Adult Dosing	A ♦ 5 gram vial via slow IV infusion over 15 minutes (Can be given IO as a last resort) A ♦ May repeat 5 grams IV via slow IV infusion over 15 minutes to 2 hours depending on clinical response		
Pediatric Dosing	P ♦ 70 mg/kg slow IV over 15 minutes; max dose of 5 grams (Can be given IO as a last resort) P ♦ May repeat a dose of 35 mg/kg IV; max dose 2.5 g, depending on severity of poisoning and clinical response.		
Therapeutic Action	Binds to cyanide molecules and is eliminated as waste		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Must not be used in conjunction with other Cyanide antidotes - May cause hypertension		
Medical Control	Adults: - In cardiac arrest—No - In patients not in arrest—Yes Pediatrics: - In cardiac arrest—No - In patients not in arrest—Yes		
Protocols	Trauma Protocol 3008 – Cyanide Poisoning & Antidotes		
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Subject: Ipratropium (Atrovent)

Effective: June 1, 2021

Last Modified: May 16, 2026

EMR		EMT	AEMT	Paramedic
Packaging	0.5 mg in 2.5 ml plastic ampule 1 in drug bag			
Indications	- Bronchospasm in Asthma, COPD, Emphysema - Allergic reaction/Anaphylaxis with wheezing			
Adult Dosing	A 0.5 mg (2.5 ml), nebulized with O₂ at 8-10 LPM A Combined with first dose of Albuterol			
Pediatric Dosing	P 0.5 mg (2.5 ml), nebulized with O₂ at 8-10 LPM P Combined with first dose of Albuterol			
Therapeutic Action	Causes bronchodilation by anticholinergic effect			
Contraindications	None in the emergency setting			
Precautions And Side Effects	- Once initiated, the patient should be removed by EMS. - Use with caution in patients with narrow-angle glaucoma and lactating mothers.			
Medical Control	- Adults: No - Pediatrics: No			
Protocols	Medical Protocol 1008 – Advanced Airway Management Medical Protocol 4002 – Allergic Reactions/Anaphylaxis Medical Protocol 4003 – Asthma/Emphysema/COPD			
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Subject: Ketamine (Ketalar)

Effective: June 1, 2021

Last Modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	500 mg/10 mL vial (50 mg/ml) - One in drug bag		
Indications	- For the AEMT and Paramedic - Chemical restraint for combative patient, including excited delirium - Pain control (should be considered a second line medication for the management of pain) - Seizing patient with Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam - For the Paramedic {Sedate-to-Intubate} or {RSI} - Conscious adult patient requiring pacing or cardioversion (preferred method)		
Adult Dosing	- For pain: - A 25 mg IV, may repeat 25 mg IV after 5 minutes. - A If unable to obtain IV: - A 25 mg IN or 50 mg IM, may repeat 25 mg IN or 50 mg IM after 10 minutes. - For combative patients: - A 250 mg IM anterolateral thigh. - or - A 100 mg slow IV - A If no change in 10 minutes for IM or 5 minutes for IV, repeat: - A 250 mg IM anterolateral thigh - or - A 100 mg slow IV - For the seizing patient with Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam: - A Consider 100 mg IV/IO or (if no IV/IO), then 250 mg IM - Do not reduce geriatric dosing to half dose when sedating for pacing and cardioversion - For the Paramedic performing {Sedate to Intubate} or {Rapid Sequence Intubation}: - A 100 mg slow IV, may repeat 100 mg IV after 5 minutes - Do not reduce geriatric dosing to half dose when attempting to achieve complete sedation - For the Paramedic preparing the conscious adult patient for pacing or cardioversion - A 25 mg IV - Do not reduce geriatric dosing to half dose when sedating for pacing and cardioversion - A For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses (Exceptions: {RSI or STI}, RSE seizures, combative patients, pacing or cardioversion)		
Pediatric Dosing	P Not to be administered for pain to any patient less than 16 y/o - Emergency sedation for combative patient, including excited delirium: P Limited to use in patients age 8 or greater. P 1 mg/kg slow IV (max dose 100 mg). - or P 5 mg/kg IM (maximum dose is two doses of no more than 250 mg or 500 mg total) P ♦ Call MCP for repeat doses		



Subject:

Ketamine (Ketalar)

Effective:

June 1, 2021

Last Modified:

Dec. 11, 2024

Therapeutic Action	- Ketamine is a Schedule III Phencyclidine (PCP) derivative that is rapid acting and produces a “dissociative” anesthesia in which the patient’s consciousness is detached from their nervous system. - Due to its “dissociative” properties, Ketamine is a potent analgesic. - May be given as an adjunct to narcotic pain medication, particularly in patients at risk for hypotension or respiratory depression.
Contraindications	- Suspected cardiac chest pain - Hypertensive crisis - When significant elevations in BP might prove harmful: - Acute Myocardial Infarction - Angina Pectoris - Aortic dissection
Precautions And Side Effects	- Emergence reaction may occur, when patient is awakening (hallucinations, delirium, confusion, etc.) - Provide continuous cardiac monitoring, EtCO₂ and pulse oximetry with sedated patients. - Management should include use of a nasopharyngeal airway, proper positioning and persistent suctioning to maintain a clear airway. - Geriatric & debilitated patients require lower doses & are more prone to side effects. - Catecholamine release (hypertension, tachycardia) - Hypersalivation (the ketamine drool) - Nausea, vomiting, particularly prevalent in pediatrics. - Minimal cardiac depression occasionally reported with high doses administered rapidly IV. - May transiently increase heart rate and blood pressure by central sympathetic stimulation. - May require administration of midazolam prior to wearing off.
Medical Control	Adults: No Pediatrics: - No - For repeat sedation doses - yes
Protocols	General Protocol 1008 – Advanced Airway Management General Protocol 1010 – {Sedate to Intubate and Rapid Sequence Intubation} General Protocol 1014 – Pain Management Cardiac Protocol 2010 – Bradycardia Cardiac Protocol 2011 – Tachycardia Trauma Protocol 3007 – Crush Syndrome Trauma Medical Protocol 4005 – Behavioral Emergencies - Combative Patients/Emergency Sedation
END OF SECTION	



Subject: Lactated Ringers

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Usually a 1000 ml flexible, non-latex plastic bag • Generally with a pH of 6.5. • Not in drug bags or caches		
Indications	• Solution for fluid and electrolyte replenishment • Hypovolemia • Flushing of wounds • Shock • Pulmonary edema with systolic BP over 100 mmHg • Sepsis		
Adult Dosing	A Non traumatic shock without pulmonary edema: 500 ml IV, may repeat up to two times if needed A Non traumatic shock with pulmonary edema: 250 ml IV • Sepsis: - A 1 L IV - A ♦ Additional IV fluid if indicated - A Penetrating trauma to chest or abdomen: enough fluid to obtain a radial pulse - A If BGL reads over 400 mg/dL or “High” on glucometer, administer 500 ml fluid IV – wide open. • Crush syndrome: - A Initial treatment: 1 L IV then 500 ml/hour IV - A If hypotensive and the patient has been trapped more than 1 hour, then additional 1 L IV • Heat exposure: - A 500 ml IV, may repeat one time - A ♦ Additional IV fluid, if indicated		
Pediatric Dosing	P 20 ml/kg IV bolus P In heat exposures, may repeat 20/ml/kg IV bolus P ♦ In shock, call for orders to administer additional fluid		
Therapeutic Action	• Used for hydration and management of hypotension		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• None		
Medical Control	• Adults: Yes, for additional fluid administrations in some circumstances • Pediatrics: Yes, for additional fluid administrations in some circumstances		
Protocols	• General Protocol 1005 – General Patient Management • Cardiac Protocol 2005 – Cardiac Arrest; V-Fib or Pulseless V-Tach • Cardiac Protocol 2008 – Suspected Cardiac Chest pain • Cardiac Protocol 2009 – Cardiac Alert Program • Trauma Protocol 3001 – General Trauma Management • Trauma Protocol 3004 – Trauma Arrest • Trauma Protocol 3007 – Crush Syndrome Trauma • Trauma Protocol 3014 – Heat Exposure • Medical Protocol 4002 – Allergic Reaction/Anaphylaxis • Medical Protocol 4008 – Diabetic Emergencies – Hypoglycemia/Hyperglycemia • Medical Protocol 4015 - Sepsis • Medical Protocol 4016 – Shock		
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Subject: Lidocaine 2%

Effective: June 1, 2021

Last Modified: Feb. 27, 2025

EMR	EMT	AEMT	Paramedic
Packaging	100 mg in 5 ml syringe (20 mg/ml) - Three in drug bag		
Indications	- For AEMT and Paramedic: - For pain caused by pressure of intraosseous fluid administration - For Paramedic: - Intubation on conscious patient - JITSO – Cardiac arrest: V-Fib/Pulseless V-Tach and Tachycardia, in the absence of Amiodarone		
Adult Dosing	- Pain associated with IO infusion (AEMT, Paramedic): A 1.5 mg/kg IO (maximum dose 100 mg) - Intubation on conscious patient (Paramedic): A 100 mg (5 ml) nebulized with 8-10 LPM O₂ or A 100 mg (5 ml) IN with 50 mg (2.5 ml) in each nostril - JITSO for Cardiac Arrest: V-Fib or Pulseless V-Tach (Paramedic): A 150 mg (7.5 ml) IV or IO A Repeat dose of 75 mg (3.75 ml) IV or IO - JITSO for Tachycardia (Paramedic) A 150 mg (7.5 ml) IV or IO		
Pediatric Dosing	- Pain associated with IO infusion (AEMT, Paramedic): P 0.5 mg/kg IO (maximum dose 100 mg) - Intubation on conscious patient (Paramedic): P 1.5 mg/kg nebulized with 8-10 LPM O₂ or IN (maximum dose 100 mg) - JITSO for Cardiac Arrest: V-Fib or Pulseless V-Tach (Paramedic): P 1 mg/kg IV or IO (maximum dose 100 mg) P Repeat dose of 1 mg/kg IV or IO (maximum dose 75 mg)		
Therapeutic Action	Decreases automaticity		
Contraindications	- Hypersensitivity - Second degree or third degree heart block, in absence of an artificial pacemaker		
Precautions And Side Effects	- Use extreme caution in patients with hepatic disease, heart failure, marked hypoxia, severe respiratory depression, hypovolemia or shock, incomplete heart block or bradycardia and atrial fib. - Side Effects: - Altered level of consciousness, confusion or lightheadedness - Cardiovascular collapse and/or hypotension - Bradycardia - Blurred vision - irritability - Muscle twitching and seizures with high doses		
Medical Control	- Adults: No - Pediatrics: No		
Protocols	General Protocol 1008 – Advanced Airway Management General Protocol 1012 – Intraosseous Infusion Cardiac Protocol 2003 – Cardiac Arrest: Asystole or PEA Cardiac Protocol 2005 – Cardiac Arrest: V-Fib or Pulseless V-Tach Cardiac Protocol 2011 – Tachycardia Medical Protocol 4002 – Allergic Reactions/Anaphylaxis Medical Protocol 4003 – Asthma/Emphysema/COPD		
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Subject: Lidocaine 2% Gel

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	2% gel in a tube - Not carried in drug bag		
Indications	Lubrication of airway adjunct on conscious patient		
Adult Dosing	A Apply to airway adjunct.		
Pediatric Dosing	P Apply to airway adjunct.		
Therapeutic Action	Suppresses stimulation of the upper airway activity such as, swallowing, gagging or coughing that can cause cardiovascular stimulation and elevation in intracranial pressure		
Contraindications	None		
Precautions And Side Effects	None		
Medical Control	- Adults: No - Pediatrics: No		
Protocols	General Protocol 1008 – Advanced Airway Management		
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Subject: Magnesium-Containing Antacid

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	- Varies by manufacturer or vendor - Not carried in drug bag - Examples include Maalox and Mylanta		
Indications	- Ingestion of Hydrofluoric Acid - Hydrofluoric Acid on skin		
Adult Dosing	- For Ingestion: - A Following dilution with water or milk, have patient drink 3-4 oz. Maalox or Mylanta. - For exposure: - A Following irrigation, apply topically to burned area unless topical agents is already applied		
Pediatric Dosing	P Same application as for adult patients		
Therapeutic Action	Neutralize acid and increases the pH		
Contraindications	None in the emergency setting.		
Precautions And Side Effects	- Use with caution in: - Neonates - Geriatric patients - Patients with renal impairment - Side Effects: - Hypercalcemia - Hypermagnesemia - Hypotension - Nausea & vomiting		
Medical Control	- Adults: No - Pediatrics: No		
Protocols	Special Operations Protocol 6004 – Hydrofluoric Acid Exposure		
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Subject: Magnesium Sulfate

Effective: June 1, 2025

Last Modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	4 grams pre-mixed in 100 ml IV bag (40 mg/ml) - May be a 4 gm vial and a 100 ml NaCl IV bag depending on availability - One in drug bag		
Indications	- Polymorphic ventricular tachycardia (torsades de pointes) in adult patient (with or without a pulse) - Seizing pregnant patient with gestation greater than 20 weeks - Seizing patient with less than 12 weeks postpartum - Asthmatic bronchoconstriction, refractory to Duo-Neb, and Epinephrine and Solu-Medrol		
Adult Dosing	A Polymorphic ventricular tachycardia: 2 gm (1/2 IV bag) infused with macro-drip tubing over 10 min.* A Seizing pregnant/postpartum: 4 gm (whole bag) with macro-drip tubing over 20 min.* A ♦ Asthma: 2 gm (1/2 IV bag) infused with macro-drip tubing over 10 min.* *when using either 10 gtt/ml or 15 gtt/ml tubing, a drip rate of 1 gtt/second will deliver 2 gm in 10 min. or 4 gm in 20 min. (approximately)		
Pediatric Dosing	P Seizing pregnant/postpartum: 4 gm (whole bag) with macro-drip tubing over 20 min.* *when using either 10 gtt/ml or 15 gtt/ml tubing, a drip rate of 1 gtt/second will deliver 2 gm in 10 min. or 4 gm in 20 min. (approximately)		
Therapeutic Action	- CNS depressant - Blocks peripheral neuromuscular transmission - Slows rate of sino-atrial node impulse formation in myocardium and prolongs conduction time - Bronchial smooth muscle relaxation		
Contraindications	- Hypersensitivity - Any heart block, except for 1st degree block - Administer with caution if SBP less than 90 mmHg - Acute myocardial infarction (AMI) - Abdominal pain		
Precautions And Side Effects	- Any patient receiving intravenous Magnesium Sulfate should have continuous cardiac monitoring and frequent monitoring of vital signs. - Can cause respiratory depression - Hypotension, bradycardia and conduction disturbances may occur if given rapidly - Toxicity is associated with CNS and neuromuscular depression		
Medical Control	Adult: Yes, when given to treat asthma refractory to primary medications Pediatric: No		
Protocols	Cardiac Protocol 2005 – Cardiac Arrest: V-Fib or Pulseless V-Tach Cardiac Protocol 2011 – Tachycardia Medical Protocol 4003 – Asthma/Emphysema/COPD Medical Protocol 4014 – Seizures		
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Subject: Methylprednisolone (Solu-medrol)

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• 125 mg in 2 ml • One in drug bag		
Indications	• Severe allergic reactions • Anaphylaxis • Asthma • COPD • Emphysema • Intended to augment standard therapy for anaphylaxis, allergic reaction, and to address airway edema and inflammation in asthma.		
Adult Dosing	A Solu-Medrol 125 mg IV A Given to patients in the Allergic reaction or Anaphylaxis protocol only after all other applicable first-line medications have been delivered.		
Pediatric Dosing	P Solu-Medrol 2 mg/kg IV, max dose 125 mg P Given to patients in the Allergic reaction or Anaphylaxis protocol only after all other applicable first-line medications have been delivered.		
Therapeutic Action	• Potent anti-inflammatory steroid • Accelerates detoxification of cyanide		
Contraindications	• None in emergency setting		
Precautions And Side Effects	• Intended for cases that are of a more urgent nature. • No significant change in patient condition in the field should be expected after administration. • Do not to initiate an IV only to administer this medication. • Side Effects: ○ Cardiac arrhythmias ○ Syncope		
Medical Control	• Adults: No • Pediatrics: No		
Protocols	• Medical Protocol 4002 – Allergic Reactions/Anaphylaxis • Medical Protocol 4003 – Asthma/Emphysema/COPD		
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Subject: Midazolam (Versed)

Effective: June 1, 2021

Last Modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	10 mg in 2 ml vial, (5 mg/ml) - Two in drug bag		
Indications	- For the AEMT and Paramedic: - Seizures - As chemical restraint for combative patient - Chest pain associated with stimulant overdose (adults only) - For the Paramedic: - Conscious patient requiring cardioversion - Conscious patient requiring pacing {Sedate-to-Intubate} or {RSI} in normotensive patients - After intubation, if patient is resisting and SBP is normal for age.		
Adult Dosing	- If seizures, chemical restraint, or chest pain in stimulant overdose (AEMT, Paramedic): A 10 mg IN (5 mg in each nostril) <u>or</u> 2.5 mg slow IV <u>or</u> 5 mg IM A Repeat 5 mg IN (after 10 min.) <u>or</u> 2.5 mg slow IV (after 5 min.) <u>or</u> 5 mg IM (after 10 min.) - If evidence of Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam (AEMT, Paramedic), then repeat at second dose levels/routes - If conscious patients requiring cardioversion/pacing or patient resisting ETT (Paramedic) A 2.5 mg slow IV A In {Sedate-to-intubate} or {RSI}, 5 mg slow IV (in patients who are normotensive), may repeat up to 10 mg IV (Paramedic) G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses (Exceptions: {RSI or sedate-to-intubate}, combative patients, pacing or cardioversion)		
Pediatric Dosing	P If seizures, or chemical restraint for combative patients (AEMT, Paramedic): P 0.2 mg/kg IN (maximum dose 10 mg) <u>or</u> P 0.1 mg/kg slow IV (maximum dose 2.5 mg) <u>or</u> P 0.2 mg/kg IM (maximum dose 5 mg) P In seizures, repeat same doses (maximum IN 5mg, maximum IV 2.5 mg, maximum IM 5 mg) P ♦ In chemical restraint, call MCP for repeat doses P If evidence of Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam (AEMT, Paramedic), <u>then repeat at second dose levels/routes</u> P If conscious patients requiring cardioversion/pacing or patient resisting ETT (Paramedic) P 0.1 mg/kg slow IV (maximum dose 2.5 mg)		
Therapeutic Action	Provides sedation		
Contraindications	Respiratory distress		
Precautions And Side Effects	- Use with caution with lactating mothers. - Geriatric & debilitated patients require lower doses & are more prone to side effects. - Can cause respiratory depression - Monitor respirations and ventilate if necessary. - The Paramedic should intubate as indicated, the AEMT should intubate if apneic. - Provide continuous cardiac monitoring, EtCO₂ and pulse oximetry with sedated patients.		
Medical Control	Adults: No Pediatrics: Yes, for repeat doses in Combative Patient/Emergency Sedation Protocol		



Subject: Midazolam (Versed)

Effective: June 1, 2021

Last Modified: May 16, 2026

Protocols

- [General Protocol 1008 – Advanced Airway Management](#)
- [General Protocol 1010 – {Sedate to Intubate and Rapid Sequence Intubation}](#)
- [Cardiac Protocol 2006 – AICD Activations](#)
- [Cardiac Protocol 2010 – Bradycardia](#)
- [Cardiac Protocol 2011 – Tachycardia](#)
- [Medical Protocol 4005 – Behavioral Emergencies - Combative Patients/Emergency Sedation](#)
- [Medical Protocol 4012 – Overdose/Poisoning](#)
- [Medical Protocol 4014 – Seizures](#)
- [Special Operations Protocol 6002 – Antidote Resources](#)
- [Special Operations Protocol 6005 – Organophosphate or Nerve Agent Exposure](#)

END OF SECTION



Subject: Morphine (JITSO)

Effective: June 1, 2021

Last Modified: Apr. 7, 2025

EMR	EMT	AEMT	Paramedic
Packaging	5 mg in 1ml vial - Two in drug bag in the absence of fentanyl		
Indications	Pain relief in suspected cardiac chest pain, trauma emergencies, extremity fractures, dislocations, sprains, frostbite, abdominal pain, Hydrofluoric Acid (HF) exposure		
Adult Dosing	A Up to 5 mg slow IV based on patient’s weight, provided SBP greater than 100. A May repeat up to 5 mg slow IV after 5 minutes A If unable to establish IV, Morphine 5 mg IM, may repeat after 10 minutes. G For patients greater than 69 y/o, reduce dosing for sedatives and analgesics to one half (½) of the adult doses		
Pediatric Dosing	- Pain relief in pediatric patients greater than 2 years old: - P 0.1 mg/kg slow IV (maximum dose 5 mg) provided appropriate SBP. - P ♦ May repeat 0.1 mg/kg, (maximum dose 5 mg) after 5 minutes - P If unable to establish IV, 0.1 mg/kg IM (maximum dose 5 mg) - P ♦ May repeat IM dose after 10 minutes		
Therapeutic Action	Provides analgesia, reduces cardiac preload by increasing venous capacitance and decreasing afterload		
Contraindications	- Hypersensitivity to narcotics - Hypotension - Head injury, increased intracranial pressure - Severe respiratory depression - Patients who have taken MAO inhibitors within 14 days		
Precautions And Side Effects	- Use with caution in the elderly, those with asthma, and in those susceptible to CNS depression. - Provide continuous cardiac monitoring, EtCO₂ and pulse oximetry with sedated patients. - Geriatric & debilitated patients require lower doses & are more prone to side effects. - Hypotension - Tachycardia, or bradycardia - May worsen bradycardia or heart block in inferior MI (vagotonic effect) - Palpitations - Syncope - Euphoria - Facial flushing - Respiratory depression - Bronchospasm - Dry mouth - Allergic reaction		
Medical Control	Adults: No ♦ Pediatrics: Yes, for repeat doses		
Protocols	General Protocol 1014 – Pain Management Cardiac Protocol 2006 – AICD Activations Cardiac Protocol 2008 – Suspected Cardiac Chest Pain Cardiac Protocol 2009 – Cardiac Alert Program		
END OF SECTION			

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Subject: Naloxone (Narcan)

Effective: June 1, 2021

Last Modified: Jan. 12, 2025

EMR	EMT	AEMT	Paramedic
Packaging	2 mg in 2 ml vial (1 mg/ml) - Four in drug bag		
Indications	- High index of suspicion of narcotic overdose - Respiratory depression - Suspicion of drug abuse in cardiac arrest		
Adult Dosing	(EMR or EMT): - A Up to 4 mg IN (half dose per nostril) (AEMT or Paramedic): - A Up to 4 mg IN (half dose per nostril) or 2 mg IV - A If no IV, up to 4 mg IM - A Titrate dosing to adequate respirations, repeat as needed		
Pediatric Dosing	(EMR, EMT): - P If 20 kg or less, then 0.1 mg/kg IN (maximum dose 2 mg) (half dose per nostril) - P If greater than 20 kg, then 2 mg IN, may repeat as needed (AEMT or Paramedic): - P If 20 kg or less, then 0.1 mg/kg IN (half dose per nostril), IV or IM (maximum dose 2 mg) - P If greater than 20 kg, then 2 mg IN (half dose per nostril) - P If using IN route and respirations don't improve after 2 mins., establish and administer via IV - P Titrate dosing to adequate respirations, repeat as needed.		
Therapeutic Action	A competitive narcotic antagonist		
Contraindications	- Hypersensitivity - Newborn patients		
Precautions And Side Effects	- Any intranasal administration should be given at a half dose in each nostril - Onset of action is two minutes, if no response two minutes after dosing, then give additional doses - For the AEMT or Paramedic: if the patient has a pulse, Naloxone should be given before intubation. - After administration, patient transport by EMS is encouraged, even if patient becomes responsive. - Use with caution in narcotic-dependent patients who may experience withdrawal syndrome (including neonates of narcotic-dependent mothers). - Caution should be exercised when administering to narcotic addicts (may precipitate withdrawal symptoms) - Side Effects: - Tachycardia - Hypertension - Dysrhythmias - Diaphoresis - Blurred vision - Nausea and vomiting - May not reverse hypotension		
Medical Control	- Adult: No - Pediatric: No		
Protocols	- General Protocol 1005 – General Patient Management - General Protocol 1012 – Intraosseous Infusion - Medical Protocol 4012 – Overdose/Poisoning		
END OF SECTION			

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Subject: Nitroglycerin (Nitrostat)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	- Dark brown glass bottle, 0.4 mg SL tablets - One bottle in drug bag		
Indications	- For the EMT, AEMT and Paramedic: - Cardiac related chest pain - For the AEMT and Paramedic: - Pulmonary edema with systolic BP over 100 mmHg - Stimulant overdose with chest pain		
Adult Dosing	A 0.4 mg SL every 5 min for continued chest pain up to a total of 3 tablets		
Pediatric Dosing	Not indicated in pediatric patients		
Therapeutic Action	Vasodilator which decreased preload and to a lesser extent, afterload		
Contraindications	- Hypersensitivity - Hypotension - Use of sexual enhancement drugs (Viagra, Cialis, Levitra) in last 24 hours - Taking Revatio (a pulmonary hypertension medication) - Head injury		
Precautions And Side Effects	- Use only on patients who are greater than 25 years old or have been prescribed Nitroglycerin - Side Effects: - Transient headache - Reflex tachycardia - Hypotension - Diaphoresis - Postural syncope - Nausea & vomiting		
Medical Control	- Adult: - For the EMT: Yes - For the AEMT and Paramedic: No - Pediatric: Not applicable		
Protocols	Cardiac Protocol 2008 – Suspected Cardiac Chest Pain Medical Protocol 4012 – Overdose/Poisoning Medical Protocol 4013 – Respiratory Distress/Pulmonary Edema		
END OF SECTION			

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Subject: Norepinephrine (Levophed)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic															
Packaging	4 mg in 4ml (1mg/ml) vial for dilution in 250 ml of IV fluids - One in drug bag																	
Indications	For blood pressure control in acute hypotensive states in the non-trauma patient.																	
Adult Dosing	A Add 4 mg to 250 ml of IV fluids. A Infuse starting at 30 drops per minute (max 45 drops) with 60 drop tubing and titrate to effect. A Increase by 5 drops every 5 minutes. <table><tr><th>gtts/min</th><th></th><th>mcg/min</th></tr><tr><td>30</td><td>=</td><td>8</td></tr><tr><td>35</td><td>=</td><td>9.35</td></tr><tr><td>40</td><td>=</td><td>10.7</td></tr><tr><td>45</td><td>=</td><td>12</td></tr></table>			gtts/min		mcg/min	30	=	8	35	=	9.35	40	=	10.7	45	=	12
gtts/min		mcg/min																
30	=	8																
35	=	9.35																
40	=	10.7																
45	=	12																
Pediatric Dosing	P ♦ Contact MCP for dosing and administration guidance.																	
Therapeutic Action	- Peripheral vasoconstrictor. - Positive inotrope (increases cardiac contractility) and chronotrope (increases heart rate).																	
Contraindications	- Should not be given to patients who are hypotensive from acute hemorrhage. - Do not use the solution if its color is pinkish or darker than slightly yellow or if it contains particles.																	
Precautions And Side Effects	- Protect the vial from light - This drug <u>must</u> be diluted before administration. - Administer in free-flowing IV and watch for infiltration. - Avoid hypertension. - If extravasation occurs, stop the infusion immediately as necrosis may occur. - Leave the catheter in place so that a reversal agent can be given through the infiltrated catheter.																	
Medical Control	- Adult: No. - Pediatric: Yes																	
Protocols	Cardiac Protocol 2009 – Cardiac Alert Program Medical Protocol 4015 – Sepsis Medical Protocol 4016 – Shock																	
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Subject: Normal Saline (Sodium Chloride Solution)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Usually a 1000 ml flexible, non-latex plastic bag • Generally with a pH of 6.5. • Not in drug bags or caches		
Indications	• Solution for fluid and electrolyte replenishment • Hypovolemia • Flushing of wounds • Shock • Pulmonary edema with systolic BP over 100 mmHg • Sepsis		
Adult Dosing	A Non traumatic shock without pulmonary edema: 500 ml IV, may repeat up to two times if needed A Non traumatic shock with pulmonary edema: 250 ml IV • Sepsis: - A 1 L IV - A ♦ Additional IV fluid if indicated - A Penetrating trauma to chest or abdomen: enough fluid to obtain a radial pulse - A If BGL reads over 400 mg/dL or “High” on glucometer, administer 500 ml fluid IV – wide open. • Crush syndrome: - A Initial treatment: 1 L IV then 500 ml/hour IV - A If hypotensive and the patient has been trapped more than 1 hour, then additional 1 L IV • Heat exposure: - A 500 ml IV, may repeat one time - A ♦ Additional IV fluid, if indicated		
Pediatric Dosing	P 20 ml/kg IV bolus P In heat exposures, may repeat 20/ml/kg IV bolus P ♦ In shock, call for orders to administer additional fluid		
Therapeutic Action	• Used for hydration and management of hypotension		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• None		
Medical Control	• Adults: Yes, for additional fluid administrations in some circumstances • Pediatrics: Yes, for additional fluid administrations in some circumstances		
Protocols	• General Protocol 1005 – General Patient Management • Cardiac Protocol 2005 – Cardiac Arrest; V-Fib or Pulseless V-Tach • Cardiac Protocol 2008 – Suspected Cardiac Chest pain • Cardiac Protocol 2009 – Cardiac Alert Program • Trauma Protocol 3001 – General Trauma Management • Trauma Protocol 3004 – Trauma Arrest • Trauma Protocol 3007 – Crush Syndrome Trauma • Trauma Protocol 3014 – Heat Exposure • Medical Protocol 4002 – Allergic Reaction/Anaphylaxis • Medical Protocol 4008 – Diabetic Emergencies – Hypoglycemia/Hyperglycemia • Medical Protocol 4015 - Sepsis • Medical Protocol 4016 – Shock		
END OF SECTION			

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Subject: Normosol-R

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Usually a 1000 ml flexible, non-latex plastic bag • Generally with a pH of 6.5. • Not in drug bags or caches		
Indications	• Solution for fluid and electrolyte replenishment • Hypovolemia • Flushing of wounds • Shock • Pulmonary edema with systolic BP over 100 mmHg • Sepsis		
Adult Dosing	A Non traumatic shock without pulmonary edema: 500 ml IV, may repeat up to two times if needed A Non traumatic shock with pulmonary edema: 250 ml IV • Sepsis: - A 1 L IV - A ♦ Additional IV fluid if indicated - A Penetrating trauma to chest or abdomen: enough fluid to obtain a radial pulse - A If BGL reads over 400 mg/dL or “High” on glucometer, administer 500 ml fluid IV – wide open. • Crush syndrome: - A Initial treatment: 1 L IV then 500 ml/hour IV - A If hypotensive and the patient has been trapped more than 1 hour, then additional 1 L IV • Heat exposure: - A 500 ml IV, may repeat one time - A ♦ Additional IV fluid, if indicated		
Pediatric Dosing	P 20 ml/kg IV bolus P In heat exposures, may repeat 20/ml/kg IV bolus P ♦ In shock, call for orders to administer additional fluid		
Therapeutic Action	• Used for hydration and management of hypotension		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• None		
Medical Control	• Adults: Yes, for additional fluid administrations in some circumstances • Pediatrics: Yes, for additional fluid administrations in some circumstances		
Protocols	• General Protocol 1005 – General Patient Management • Cardiac Protocol 2005 – Cardiac Arrest; V-Fib or Pulseless V-Tach • Cardiac Protocol 2008 – Suspected Cardiac Chest pain • Cardiac Protocol 2009 – Cardiac Alert Program • Trauma Protocol 3001 – General Trauma Management • Trauma Protocol 3004 – Trauma Arrest • Trauma Protocol 3007 – Crush Syndrome Trauma • Trauma Protocol 3014 – Heat Exposure • Medical Protocol 4002 – Allergic Reaction/Anaphylaxis • Medical Protocol 4008 – Diabetic Emergencies – Hypoglycemia/Hyperglycemia • Medical Protocol 4015 - Sepsis • Medical Protocol 4016 – Shock		
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Subject: Ondansetron (Zofran)

Effective: June 1, 2021

Last Modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	4 mg in 2 ml vial, (2 mg/ml) 1 vial in drug bag 4 mg tablet 1 tablet in drug bag		
Indications	For nausea or active vomiting		
Adult Dosing	- For the EMT, AEMT and Paramedic: A 4 mg tablet PO dissolving tablet - For the AEMT, and Paramedic: A 4 mg slow IV/IO, preferred route for active vomiting as patient may need hydration. - For the Paramedic: A If no IV/IM, 4 mg (2 ml) PO or the IV form discharged into the patient's mouth.		
Pediatric Dosing	- For the EMT, AEMT and the Paramedic: P 4 mg tablet PO dissolving tablet if patient 12 y/o or older - For the Paramedic: P 0.1 mg/kg IV/IM (max 4 mg) if patient 5 y/o or older and weight is 30 kg or more.		
Therapeutic Action	- Stimulation of 5-HT₃ receptors causes transmission of sensory signals to the vomiting center via vagal afferent fibers to induce vomiting. - By binding to 5-HT₃ receptors, Ondansetron blocks vomiting mediated by serotonin release.		
Contraindications	Known hypersensitivity to Ondansetron		
Precautions And Side Effects	- During pregnancy it should only be used where clearly needed. - Ondansetron (Zofran) is NOT to be given prophylactically with Naloxone. <u>Side effects:</u> - Constipation or diarrhea - Fever - Headache. - Sudden blindness of 2-3 minutes duration. (the speed of delivery may contribute to the blindness)		
Medical Control	Adults: No Pediatrics: No		
Protocols	Medical Protocol 4001 – Abdominal Pain		
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Subject: Oral Glucose

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Tube; concentration varies, check label • Not carried in drug bag		
Indications	• Hypoglycemia • Generalized hypothermia without arrest • Altered level of consciousness of unknown cause • Seizures with BGL of less than 60 mg/dl, no BGL monitor; or suspicion of hypoglycemia despite BGL reading • For the AEMT and Paramedic, no IV access		
Adult Dosing	A 1 tube A May be repeated in 10 minutes if BGL remains less than 60 mg/dl		
Pediatric Dosing	P 1 tube P May be repeated in 10 minutes if BGL remains less than 60 mg/dl		
Therapeutic Action	• Raise blood glucose concentration		
Contraindications	• Inability to control the airway		
Precautions And Side Effects	• Use caution when giving to unresponsive patients. • Hyperglycemia		
Medical Control	• Adults: No • Pediatrics: No		
Protocols	• Medical Protocol 4008 – Diabetic Emergencies - Hypoglycemia		
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Subject: Plasmalyte-A

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	• Usually a 1000 ml flexible, non-latex plastic bag • Generally with a pH of 7.4. • Not in drug bags or caches		
Indications	• Solution for fluid and electrolyte replenishment • Hypovolemia • Flushing of wounds • Shock • Pulmonary edema with systolic BP over 100 mmHg • Sepsis		
Adult Dosing	A Non traumatic shock without pulmonary edema: 500 ml IV, may repeat up to two times if needed A Non traumatic shock with pulmonary edema: 250 ml IV • Sepsis: - A 1 L IV - A ♦ Additional IV fluid if indicated - A Penetrating trauma to chest or abdomen: enough fluid to obtain a radial pulse - A If BGL reads over 400 mg/dL or “High” on glucometer, administer 500 ml fluid IV – wide open. • Crush syndrome: - A Initial treatment: 1 L IV then 500 ml/hour IV - A If hypotensive and the patient has been trapped more than 1 hour, then additional 1 L IV • Heat exposure: - A 500 ml IV, may repeat one time - A ♦ Additional IV fluid, if indicated		
Pediatric Dosing	P 20 ml/kg IV bolus P In heat exposures, may repeat 20/ml/kg IV bolus P ♦ In shock, call for orders to administer additional fluid		
Therapeutic Action	• Used for hydration and management of hypotension		
Contraindications	• None in the emergency setting		
Precautions And Side Effects	• None		
Medical Control	• Adults: Yes, for additional fluid administrations in some circumstances • Pediatrics: Yes, for additional fluid administrations in some circumstances		
Protocols	• General Protocol 1005 – General Patient Management • Cardiac Protocol 2005 – Cardiac Arrest; V-Fib or Pulseless V-Tach • Cardiac Protocol 2008 – Suspected Cardiac Chest pain • Cardiac Protocol 2009 – Cardiac Alert Program • Trauma Protocol 3001 – General Trauma Management • Trauma Protocol 3004 – Trauma Arrest • Trauma Protocol 3007 – Crush Syndrome Trauma • Trauma Protocol 3014 – Heat Exposure • Medical Protocol 4002 – Allergic Reaction/Anaphylaxis • Medical Protocol 4008 – Diabetic Emergencies – Hypoglycemia/Hyperglycemia • Medical Protocol 4015 - Sepsis • Medical Protocol 4016 – Shock		
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Subject: Pralidoxime (2-PAM)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	600 mg auto-injector		
Indications	- To be used following Atropine in organophosphate, or nerve agent poisoning. - Both for treatment of civilian patients at the scene, as well as for protection of public safety personnel who walk into scene & become unexpectedly contaminated.		
Adult Dosing	A ♦ 600 mg IM auto-injector		
Pediatric Dosing	P ♦ Patients greater than 20 kg: 600 mg IM auto-injector		
Therapeutic Action	- Reactivates cholinesterase after poisoning with anticholinesterase agents, (Organophosphate or Nerve Gas) - Reverses muscle paralysis after organophosphate poisoning		
Contraindications	Hypersensitivity		
Precautions And Side Effects	- Use with caution in myasthenia gravis, renal impairment, pregnancy, children. - Can spread to child through breast feeding		
Medical Control	- Adults: Yes - Pediatrics: Yes		
Protocols	Special Operations Protocol 6002 – Antidote Resources Special Operations Protocol 6005 – Organophosphate or Nerve Agent Exposure		
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Subject: Sodium Bicarbonate

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	50 mEq in 50 ml syringe (1 mEq/ml) - Two in drug bag		
Indications	- Not for routine arrests. Studies indicate no proven efficacy. - Renal dialysis patient in asystole or PEA cardiac arrest - Excited delirium patients that go into cardiac arrest - Known tricyclic overdose - Crush Syndrome		
Adult Dosing	<u>Cardiac Arrest:</u> - A In renal dialysis patient: 100 mEq IV - A ♦ Consider for the excited delirium patient who goes into arrest: 100 mEq IV <u>Tricyclic Antidepressant OD:</u> - A ♦ 100 mEq IV - A ♦ May repeat dose of 50 mEq IV for persistent or prolonged QRS <u>Crush Syndrome:</u> - A 100 mEq IV		
Pediatric Dosing	<u>Cardiac Arrest:</u> - P In renal dialysis patient: 1 mEq/kg IV <u>Tricyclic Antidepressant OD:</u> - P ♦ 1 mEq/kg IV - P ♦ May repeat dose of 0.5 mEq/kg IV for persistent or prolonged QRS <u>Crush Syndrome:</u> - P 1 mEq/kg IV		
Therapeutic Action	Buffers metabolic acidosis		
Contraindications	None in the emergency setting		
Precautions And Side Effects	- Metabolic alkalosis - Hypoxia - Rise in intracellular PCO₂ and increased tissue acidosis - Electrolyte imbalance (hypernatremia) - Seizures - Tissue sloughing at injection site		
Medical Control	Adults: - Renal Dialysis Arrest – No - Tricyclic OD – Yes - Excited Delirium Arrest - Yes Pediatrics: - Arrest – No - Tricyclic OD – Yes - Crush Syndrome - No		
Protocols	Cardiac Protocol 2004 – Cardiac Arrest - Renal Failure/Dialysis Cardiac Protocol 2010 – Bradycardia Trauma Protocol 3007 – Crush Syndrome Trauma Medical Protocol 4005 – Behavioral Emergencies - Combative Patients/Emergency Sedation Medical Protocol 4012 – Overdose/Poisoning		
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Subject: Sodium Nitrate (JITSO)

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	300 mg in 10 ml vial (30 mg/ml) - Available in caches located in each county in Homeland Security Region 3.		
Indications	Patients with known or suspected cyanide poisoning		
Adult Dosing	A ♦ 300 mg (10 ml) 3% solution slow IV		
Pediatric Dosing	Not indicated in pediatric patients		
Therapeutic Action	Oxidizes hemoglobin which then combines with cyanide to form an inactive compound		
Contraindications	Nitrite/nitrate allergy		
Precautions And Side Effects	Methemoglobinemia if given in excessive amounts		
Medical Control	- Adults: Yes - Pediatrics: Not applicable		
Protocols	Trauma Protocol 3008 – Cyanide Poisoning & Antidotes		
END OF SECTION			

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Subject: Sodium Thiosulfate

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	12.5 gm in 50 ml vial (250 mg/ml) - Available in caches located in each county in Homeland Security Region 3.		
Indications	- Conscious patient with known or suspected cyanide poisoning - Smoke inhalation with suspected cyanide component - Cardiac arrest from known or suspected cyanide poisoning or smoke inhalation		
Adult Dosing	A ♦ 12.5 gm (50 ml) 25% solution slow IV		
Pediatric Dosing	P ♦ Greater than 25 kg: 12.5 gm (50 ml) 25% solution slow IV P ♦ Less than 25 kg: 412.5 mg/kg (1.65 ml/kg) of 25% solution (max dose 12.5 g (50 ml))		
Therapeutic Action	Accelerates detoxification of cyanide		
Contraindications	None		
Precautions And Side Effects	Possible hypotension		
Medical Control	- Adults: - In cardiac arrest—No - In patients not in arrest—Yes - Pediatrics: - In cardiac arrest—No - In patients not in arrest—Yes		
Protocols	Trauma Protocol 3008 – Cyanide Poisoning & Antidotes		
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Subject: Tetracaine

Effective: June 1, 2021

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	0.5%/ml eye drop bottle (10 ml) - One in drug bag		
Indications	Prior to eye irrigation in cases of chemical injury to the eye and in other situations with significant eye pain without possibility of penetrating trauma to eye.		
Adult Dosing	A 2 drops in each affected eye		
Pediatric Dosing	P 2 drops in each affected eye		
Therapeutic Action	Provides rapid, brief, superficial anesthesia by inhibiting conduction of nerve impulses from sensory nerves		
Contraindications	- Hypersensitivity to Tetracaine - Open injury to eye		
Precautions And Side Effects	- May cause burning or stinging sensation or irritation - Can cause epithelial damage and systemic toxicity - Incompatible with mercury or silver salts often found in ophthalmic products		
Medical Control	- Adults: No - Pediatrics: No		
Protocols	Trauma Protocol 3011 – Eye Injuries		
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Subject: Tranexamic Acid (TXA)

Effective: June 1, 2025

Last Modified: May 16, 2026

EMR	EMT	AEMT	Paramedic
Packaging	1 gram in 10 ml vial (100 mg/ml) - Two in drug bag		
Indications	- Known or suspected hemorrhage secondary to trauma - Time since injury is known to be less than 3 hours ♦ Uncontrolled post-partum hemorrhage - Uncontrolled bleeding in the adult airway - Hemorrhage secondary to recent tonsillectomy in pediatric patients		
Adult Dosing	- In patients with known or suspected hemorrhage secondary to trauma: - A 2 gm IV/IO over 1-2 minutes - In patients with significant bleeding affecting the airway: - A 500 mg nebulized with an additional 3 ml normal saline, O₂ flowing at 8-10 LPM - A ♦ If significant bleeding continues after, consider administering the remainder IV/IO		
Pediatric Dosing	- In patients 8 y/o or greater in known or suspected hemorrhage secondary to trauma: - P 15/mg/kg (max 1 gram) IV/IO over 1-2 minutes - In pediatric patients with uncontrolled bleeding from a recent tonsillectomy, TXA nebulized: - P Less than 25 kg: 250 mg, nebulized with 3 ml normal saline, O₂ flowing at 8-10 LPM - P 25 kg or greater: 500 mg, nebulized with 3 ml normal saline, O₂ flowing at 8-10 LPM		
Therapeutic Action	- An anti-fibrinolytic and the synthetic equivalent of the amino acid lysine - Inhibits both plasminogen activation and plasmin activity - Prevents clot breakdown and reduces hemorrhage		
Contraindications	- Hypersensitivity - Pediatric patients, unless treating uncontrolled bleeding from a recent tonsillectomy - Time elapsed from initial injury is greater than 3 hours or otherwise unknown - Known or suspected pregnancy of greater than 20 weeks - Gastrointestinal bleeding		
Precautions And Side Effects	- Do not delay transport for administration of Tranexamic Acid (TXA) - It is important to accurately note the time of administration - Gastrointestinal disturbances may occur - Monitor patient for hypotension usually found in rapid infusions - Side effects ○ Nausea ○ Vomiting ○ Visual disturbances ○ Seizure		
Medical Control	Adult: ○ Uncontrolled hemorrhage – No ○ Bleeding in Airway (Nebulized) – No ○ Bleeding in Airway (IV/IO) secondary to nebulized – Yes ○ Post-partum hemorrhage - Yes • Pediatric: No		
Protocols	• Trauma Protocol 3015 – Hemorrhage Control • Medical Protocol 4006 – Childbirth with Complications		
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Subject: Vasopressin (JITSO)

Effective: June 1, 2025

Last Modified: Dec. 11, 2024

EMR	EMT	AEMT	Paramedic
Packaging	20 units in 1 ml vial, 20 units/ml - Usually 2 vials (20 ml) present - Not routinely present in the drug bag		
Indications	Adult patients in cardiac arrest		
Adult Dosing	A 40 units IV A Once IV is established, Vasopressin is permitted after either first or second dose of Epinephrine.		
Pediatric Dosing	Not indicated in pediatric patients		
Therapeutic Action	- Potent peripheral vasoconstrictor. - May be used as an alternative pressor to Epinephrine in the treatment of adult shock-refractory VF and PEA		
Contraindications	None in the adult cardiac arrest		
Precautions And Side Effects	May produce cardiac ischemia and angina		
Medical Control	- Adults: No - Pediatrics: Not applicable		
Protocols	Cardiac Protocol 2005 – Cardiac Arrest: V-Fib or Pulseless V-Tach		
END OF SECTION			

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Appendix A

2026-2028

Protocol Changes

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Subject: 2026-2028 Protocol Changes

Effective: Nov. 1, 2026

Last modified: June 6, 2026

Appendix A.1 General Guidelines

- a. All the important changes made to the 2025 GMVEMSC protocol are identified in this section.
- b. Any changes made since the June 1, 2024 release are included.
 - i. As there were mid-year changes to the 2024 Protocol, those are included and listed in green text
- c. Grammatical changes, formatting or clerical corrections are not mentioned.
- d. The different tabs are:
 - i. General Protocol Changes – includes any changes that effect the protocol as a whole or direct all of the different disciplines
 - ii. EMR – changes affecting the patient care from an EMR
 - iii. EMT – changes affecting the patient care from an EMT, including from EMR tabs
 - iv. AEMT – changes affecting the patient care from an EMT, including from EMR & EMT tabs
 - v. Paramedic - changes affecting the patient care from a Paramedic, including from all other tabs
 - vi. Drug Formulary – changes made to the 8000 series drug listings, affecting all levels
- e. It is recommended that each discipline review the changes to all the other levels as well as their own as some changes could affect their practice.

Appendix A.2 2026-2028 GMVEMSC Protocol Changes

General Protocol Changes		
Tab	Section	Change/Edit/Addition
N/A	N/A	Added a sub-section after Appendices to the protocol. This section will show mid-year changes when they occur
TOC	TOC	Added Section for Change Order Directives
Whole	Whole Document	At the Paramedic level, changed {12 Lead EKG} to {12/15 Lead EKG}
1001	1001.1.g	Added stipulation that all versions of the protocol will include a release date in the right corner of the title page
1001	1001.1.h	Added stipulation making all old versions of the protocol invalid on release of a new protocol
1001	1001.7.a & b	Added section to describe the layout of the standing orders
1001	1001.7.c	Described the look and purpose of "Change Order Directives", or mid-cycle SO changes
1002	1002.3.c	Added language to the call for orders procedure to include giving the call taker a brief synopsis of the call
1007	1007.1 Pearls	Added emphasis that CPAP and {BiPAP} are forms of "Non-invasive Ventilation"
2001	2001.1.c	Added emphasis that providers should prioritize resuscitation efforts over transport in most cardiac arrest cases
2001	2001.1.d	Added a clarifying statement that 2001.1.c applies to both adult and pediatric patients.
3005	3005.3 Pediatric	Added bullets to Pediatric Considerations , recommending pediatric burn patients go to pediatric burn centers .
4004	4004.5.c.iii	Changed admission times of DCH Mathile Center for Mental Health and Wellness to 0800-0100 hrs.
4011	4011.2.a.iii	Added: Transport to Maternity Dept. if patient experiencing obstetric related issues up to 12 weeks postpartum.
7001	7001.3.h	Changed the timeline for testing to "established by Council" to conform with new and future standards
7002	7002.1.e	Added that if normal methods of transmitting ePCRs is down, the hospital must get the report in 24 hours
7004	7004.3.b	Removed requirement to fax the drug bag discrepancy form, it can just be sent to the Hospital EMS Coordinator

Emergency Medical Responder		
Tab	Section	Change/Edit/Addition
3011	3011.1 EMR	Changed "Cover both eyes..." to "Cover affected eye(s) with a rigid cover."

Emergency Medical Technician		
Tab	Section	Change/Edit/Addition
1009	1009.4	Added the D.O.P.E trouble shooting mnemonic, for advanced airways (primarily ET tubes)



Subject:

2026-2028 Protocol Changes

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3011	3011.1 EMR	Changed "Cover both eyes..." to "Cover affected eye(s) with a rigid cover."
4001	4001.2 EMT	Added: Consider Ondansetron (Zofran) 4 mg PO dissolving tablet for nausea or active vomiting in adults
4001	4001.2 EMT	Added: Consider Ondansetron (Zofran) 4 mg PO dissolving tablet for nausea or active vomiting in pediatrics
4002	4002.2 EMT	Removed requirement for the EMT to contact MCP for nebulized Albuterol and Ipratropium
4003	4003.1 EMT	Removed requirement for the EMT to contact MCP for nebulized Albuterol and Ipratropium
4003	4003.1 EMT	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP
4013	4013.1 Pearls	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP

Advanced Emergency Medical Technician

Tab	Section	Change/Edit/Addition
1009	1009.2 AEMT	Changed recommendation for ETT depth to 21-23 cm mark to a minimum of 3x tube size for adults
1009	1009.2 AEMT	Added instructions to realign tube in the case of right or left bronchial intubation
1009	1009.2 AEMT	Added directions to reassess tube placement with EtCO2 and lung sounds after every patient movement
1009	1009.4	Added the D.O.P.E trouble shooting mnemonic, for advanced airways (primarily ET tubes)
1012	1012.1.c.iii	Added option for distal femur IO in adult arrests, emphasized looking for better access later
1012	1012.3 Pearls	Added reminder that all medications and fluids that can be given IV can be given IO
2005	2005.3.d.vi	Added stipulation to NOT do Vector Change Defibrillation for recurrent V-Fib/PVT, but only for refractory v-fib
3011	3011.1 EMR	Changed "Cover both eyes..." to "Cover affected eye(s) with a rigid cover."
3015	3015.1 AEMT	Added: In patients 8 y/o or greater, Tranexamic Acid (TXA) 15/mg/kg (max 1 gram) IV/IO over 1-2 minutes
3015	3015.1 AEMT	Added: Tranexamic Acid (TXA) 500 mg nebulized for adults with bleeding affecting the airway
3015	3015.1 AEMT	Added: If uncontrolled airway bleeding after nebulized TXA in adults, obtain MCP to administer the remainder IV/IO
3015	3015.1 Consult	Added language referencing IV/IO TXA for continued airway bleeding in adults requiring MCP orders
4001	4001.2 AEMT	Moved Ondansetron slow IV/IO for adults and pediatrics with nausea or active vomiting to AEMT level
4001	4001.2 AEMT	Change age/weight restrictions for pediatric IV/IO Ondansetron to 5 y/o or older and 30 kg or more
4002	4002.2 AEMT	Removed line stating no orders needed for the AEMT to administer nebulized meds, due to EMT change for orders
4003	4003.1 AEMT	Removed line stating no orders needed for the AEMT to administer nebulized meds, due to EMT change for orders
4003	4003.1 EMT	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP
4008	4008.2 Pearls	Added confirmation that if no IV access in a hypoglycemic patient, an IO is an appropriate option.
4013	4013.1 Pearls	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP
4014	4014.1 AEMT	Added dosing for Refractory Status Epilepticus (RSE) for adult and pediatric patients
4014	4014.1 Pearls	Added signs and symptoms of Refractory Status Epilepticus (RSE)

Paramedic

Tab	Section	Change/Edit/Addition
1008	1008.1 Paramedic	Removed Nasal Intubation to be in line with Ohio Scope of Practice
1008	1008.1 Paramedic	Added optional skill of {finger thoracostomy} with medical director's approval and specific training
1009	1009.2 AEMT	Changed recommendation for ETT depth to 21-23 cm mark to a minimum of 3x tube size for adults
1009	1009.2 AEMT	Add instructions to realign tube in adults in the case of right or left bronchial intubation
1009	1009.2 Paramedic	Removed Nasal Intubation to be in line with Ohio Scope of Practice
1009	1009.3	Removed reference to Beck Airway Airflow Monitor (BAAM) since nasal intubations are removed
1009	1009.4	Added the D.O.P.E trouble shooting mnemonic, for advanced airways (primarily ET tubes)
1012	1012.1.c.iii	Added option for distal femur IO in adult arrests, emphasized looking for better access later
1012	1012.3 Pearls	Added reminder that all medications and fluids that can be given IV can be given IO
2002	2002.2 Paramedic	Added emphasis that Paramedics are expected to provide on scene resuscitation efforts to all age groups
2002	2002.2 Paramedic	Added bullet that pediatric arrests should get at least a first dose of Epi and a functioning airway prior to transport
2002	2002.2 Paramedic	Added statement that the Paramedic should defibrillate at least once when indicated, prior to transport
2005	2005.3.d.vi	Added stipulation to NOT do Vector Change Defibrillation for recurrent V-Fib/PVT, but only for refractory v-fib
2005	2005.3.e.vii	Added statement that Administrators need to understand the risk of damage to monitors when using DSD
2011	2011.2 Paramedic	Added language to transport stable Atrial Fibrillation with a Rapid Ventricular Rate (RVR)
2011	2011.2 Paramedic	Added language to cardiovert unstable Atrial Fibrillation with a Rapid Ventricular Rate (RVR) with MCP order
2011	2011.2 Consult	Added instructions that cardioversion of unstable A-Fib with RVR is by MCP order only
2011	2011.2 Pearls	Added recommendation to rule out or treat compensatory or physiological causes of tachycardia
2011	2011.2 Pearls	Added descriptions of the hemodynamically unstable patient



Subject:

2026-2028 Protocol Changes

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3002	3002.1 Paramedic	Added optional skill of {finger thoracostomy} with medical director's approval and specific training
3011	3011.1 EMR	Changed "Cover both eyes..." to "Cover affected eye(s) with a rigid cover."
3015	3015.1 AEMT	Added: In patients 8 y/o or greater, Tranexamic Acid (TXA) 15/mg/kg (max 1 gram) IV/IO over 1-2 minutes
3015	3015.1 AEMT	Added: Tranexamic Acid (TXA) 500 mg nebulized for adults with bleeding affecting the airway
3015	3015.1 AEMT	Added: If uncontrolled airway bleeding after nebulized TXA, obtain MCP to administer the remainder IV/IO
3015	3015.1 Consult	Added language referencing IV/IO TXA for continued airway bleeding in adults requiring MCP orders
3015	3015.1 Paramedic	Added language for agencies that have a Medical Director approved whole blood delivery program
4001	4001.2 AEMT	Moved Ondansetron slow IV/IO for adults and pediatrics with nausea or active vomiting to AEMT level
4001	4001.2 AEMT	Change age/weight restrictions for pediatric IV/IO Ondansetron to 5 y/o or older and 30 kg or more
4003	4003.1 EMT	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP
4008	4008.2 Pearls	Added confirmation that if no IV access in a hypoglycemic patient, an IO is an appropriate option.
4013	4013.1 Pearls	Added recommendation to provide Albuterol 2.5 mg and Ipratropium 0.5 mg, nebulized inline with CPAP
4014	4014.1 AEMT	Added dosing for Refractory Status Epilepticus (RSE) for adult and pediatric patients
4014	4014.1 Pearls	Added signs and symptoms of Refractory Status Epilepticus (RSE)
4014	4014.1 Paramedic	Changed postpartum timeline indication for Magnesium Sulfate in seizures from less than 6 weeks to 12 weeks

Drug Formulary

Tab	Section	Change/Edit/Addition
8002	Medical Control	Removed requirement for EMTs to contact medical control
8023	Medical Control	Removed requirement for EMTs to contact medical control
8024	Indications	Added: Seizing patient with Refractory Status Epilepticus (RSE) untreated by two doses of Midazolam
8024	Adult Dosing	Added dosing of 100 mg IV/IO or (if no IV/IO), then 250 mg IM for treatment of RSE
8024	Adult Dosing	Added reminder to not reduce Ketamine in half when treating RSE in geriatric patients
8029	Indications	Changed indication in postpartum seizures from less than 6 weeks to less than 12 weeks
8031	Adult Dosing	Added third dose for adults with evidence of Refractory Status Epilepticus untreated by two previous doses
8031	Pediatric Dosing	Added third dose for pediatrics with evidence of Refractory Status Epilepticus untreated by two previous doses
8038	Adult Dosing	Added EMT as capable of administering 4 mg tablet PO
8038	Adult Dosing	Moved slow IV/IO route for adults and pediatrics to AEMT level
8038	Pediatric Dosing	Removed pediatric requirement that the patient be 40 kg or more for 4 mg PO dissolving tablet
8038	Pediatric Dosing	Changed age/weight restrictions to for IV/IO to 5 y/o or older and 30 kg or more
8046	Indications	Added uncontrolled bleeding in adult airway
8046	Adult Dosing	Added 500 mg TXA nebulized for uncontrolled bleeding in adult airway, followed by IV/IO with MCP orders
8046	Pediatric Dosing	Added: In patients 8 y/o or greater, Tranexamic Acid (TXA) 15/mg/kg (max 1 gram) IV/IO over 1-2 minutes
8046	Medical Control	Changed adult medical control indicators to align with new indications to administer TXA

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Change Order Directives

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