

This SOP represents *practice maximums*.

CATEGORY		EMR	EMT	AEMT		Р
Analgesics		EIVIR	EIVI I	ACIVII		Р
Allaigesics	Oral analganian					
	Oral analgesics			•		
	Acetaminophen		•		•	•
	Nonsteroidal anti-inflammatory		•	•	•	•
	Parenteral non-opioid analgesics					
	Acetaminophen, ketorolac			•	•	•
	Opiates			•	•	•
	Dissociative analgesics					
	Ketamine 0.5 mg/kg or less IV/IN/IM					•
Anesthetics/Sedatives						
	Topical/Otic/Occular		•	•	•	•
	Inhaled-self administered		•	•	•	•
	Local (infiltration, intraosseus)			•		•
	General - initiate					
	General - maintain infusion intubated patient					
	Sedation for the violent/aggressive patient				•	•
	Benzodiazepine/antipsychotic combination				•	•
	Benzodiazepine/antipsychotic combination				•	•
	Ketamine greater than 0.5 mg/kg IV/IM					_
	Antipsychotics				•	•
	Benzodiazepines (for sedation)				•	•
Anticonvulsants	Includes benzodiazepines and anti-epileptic drugs			•		•
Glucose Altering Agents						
	Glucose elevating agents		•	•	•	•
	Glucose lowering agents					
	Insulin SQ/IV/infusion				•	•
Antidotes						
7 11.11.401.00	Anticholinergic Antagonists				•	•
	Antionolinergic Antagonists					
	Anticholenesterase Antagonists	•	•	•	•	•
	Anticholeriesterase Antagonists					
	Danas diagonia - Antononista					
	Benzodiazepine Antagonists					
	N. C. A. C. C.					
	Narcotic Antagonists	•	•	•	•	•
	Nondepolarizing Muscle Relaxant Antagonist					
	Beta/Calcium Channel Blocker Antidote				•	•
	Tricyclic Antidepressant Overdose				•	•
	,					
	I.					

"Investigational medications and procedures which have been reviewed and approved by an Institutional Review Board (IRB) will be considered to be approved by the Medical Direction Committee solely within the context of the approved study. Investigators involved in IRB approved research are asked to present their study plans to the MDC for informational purposes so that the committee can maintain an awareness of on-going pre-hospital research in the Commonwealth. Those who desire to conduct non-IRB reviewed pilot projects, demonstration projects, or research are asked to present those proposals to the MDC prior to their implementation for review and approval by MDC."



This SOP represents *practice maximums*.

CATEGORY		EMR	EMT	AEMT	ı	Р	
	Cyanide Antidote - hydroxycobalamin			•	•	•	Added per MDC 4/2023
	Cholinesterase Reactivator	•	•	•	•	•	
	Onomicotorado i todotivator						
	Combination antidotes, e.g. DuoDote	•	•	•	•	•	
	Combination antidotes, e.g. DuoDote						
				_			
Antihistamines & Combina	tions		•	•	•	•	
Biologicals							
-	Vaccines						
	Vaccines all ages			•	•	•	
	Vaccines to age < 18 years				•	•	
	Antibiotics		•	•		•	
	Artiblotics						
Blood/Blood products							
	Initiate						
	Maintain				•	•	
Blood Modifiers							
Dioca incamere	Anticoagulants				•	•	
	Anticoagularits						
	A C 1 (1 (_	_		
	Antiplatelet agents		•	•	•	•	
	Hemostatic agents		•	•	•	•	
	Thrombolytics					•	
	,						
	Anti-fibrinolytics (eg tranexamic acid)			•	•	•	
	Anti-libiliolytics (eg tranexamic acid)			_			
Cardiovascular Agents							
	Alpha adrenergic blockers				•	•	
	Adrenergic stimulants				•	•	
	Antiarrhythmics				•	•	
	7 and army annios						
	Beta adrenergic blockers				•	•	
	Deta aurenergic blockers				•	•	
	Calcium channel blockers				•	•	
	Diuretics				•	•	
	Inotropic agents				•	•	
	instropio agonto						

"Investigational medications and procedures which have been reviewed and approved by an Institutional Review Board (IRB) will be considered to be approved by the Medical Direction Committee solely within the context of the approved study. Investigators involved in IRB approved research are asked to present their study plans to the MDC for informational purposes so that the committee can maintain an awareness of on-going pre-hospital research in the Commonwealth. Those who desire to conduct non-IRB reviewed pilot projects, demonstration projects, or research are asked to present those proposals to the MDC prior to their implementation for review and approval by the MDC."



This SOP represents practice maximums.

CATEGORY		EMR	EMT	AEMT			Р	
	Vasodilatory Agents		•	•	•		•	
	, 5							
	Vasopressors							
	Includes infusions and push dose pressors				•		•	
	Epinephrine IV/IO for cardiac arrest			•	•		•	Epinephrine at the AEMT level added by MDC 7/7/2022
	Epinephrine IM for allergic reaction		•	•	•		•	
	Epinephrine administration systems for allergic reaction			_			_	
	(See note below)		•	•	•		•	
Central Nervous System	Antipsychotic						•	
Central Nervous System	Antipsychotic				_			
Dietary Supplements/Electro	olyty Vitamine							
Dietary Supplements/Liectiv	oryte vitainins							
	Minerale start at a health save facility		Coo cooti	on: Intraven	oue Eli	uido		
	Minerals - start at a health care facility Salts - start at a health care facility		see secu	on, muaven	ous Fit	lius		
	Sails - Start at a nearth care facility							
	Electrolytes Solutions - started at a health care facility							
	Hypertonic Saline				•		•	
	Hypertonic Saline				_		•	
Coo								
Gas	Ovugan	•	•	•	•		•	
	Oxygen Heliox						•	
	пенох				_			
Gastrointestinal								
Gastronnestinai	Antacids							
	OTC			•	•		•	
	OTC				_			
	Antidiarrheals		•	•	•		•	
	Antidiamicals		_					
	Antiemetics		•	•	•		•	
	EMT SL/PO route only		_		_			
	H2 blockers							
	PO		•	•	•		•	
	IV				•		Ť	
Hormones	Corticosteroids, Mineralocorticoids			•	•		•	
	Other Hormones							
	pitocin, octreotide, prostaglandins						•	
								EMT may transport patient with IV fluids not requiring titration or
								adjustment, and without additives including electrolytes (e.g.
Intravenous Fluids	isotonic		•	•	•		•	potassium, magnesium)
* See note below)	hypotonic		•	•	•		•	, , , , , , , , , , , , , , , , , , ,

"Investigational medications and procedures which have been reviewed and approved by an Institutional Review Board (IRB) will be considered to be approved by the Medical Direction Committee solely within the context of the approved resist involved in IRB approved research are asked to present their study plans to the MDC for informational purposes so that the committee can maintain an awareness of on-going pre-hospital research in the Commowealth. Those who desire to conduct non-IRB reviewed pilot projects, demonstration projects, or research are asked to present those proposals to the MDC prior to their implementation for review and approval by MDC."



This SOP represents *practice maximums*.

CATEGORY		EMR	EMT	AEMT		Р
	hypertonic			,	•	
	M = Maintenance I = Initiate					
	Crystalloid, +/- Dextrose/Lactate		М	I/M	I/M	I/M
	with Multi=vitamins		М	M	М	М
	with Thiamine		М	М	М	М
Neuromuscular Blockers						•
Respiratory	Anticholinergics		•	•	•	•
	Sympathomimetics					
	Beta agonists		•	•	•	•
	Epinephrine (nebulized)			•	•	•
Dosage and Concentration Ca	alculation			•	•	•
M = Maintenance						
I = Initiate						
	Note: EMT's may administer medications within their					
	scope of practice in addition to providing assistance in					
	administration of those medications. EMT's may access					
	a drug kit to access those medications.					
	Note: Med-Math skills including dosage calculations and					
	measurement of medication to be administered are					
	outside EMT scope of practice. EMT's may draw					
	epinephrine from vials or ampules for the treatment of					
	acute allergic reactions using devices/systems using					
	syringes with mechanical limiters or color-coded or other					
	clearly marked indicators to facilitate accurate dose					
	measurement.					
	EMTs may transport patients with IV fluids not requiring					
	titration or adjustment, and without additives including					
	electrolytes (e.g. potassium, magnesium)					

"Investigational medications and procedures which have been reviewed and approved by an Institutional Review Board (IRB) will be considered to be approved by the Medical Direction Committee solely within the context of the approved study. Investigators involved in IRB approved research are asked to present their study plans to the MDC for informational purposes so that the committee can maintain an awareness of on-going pre-hospital research in the Commonwealth. Those who desire to conduct non-IRB reviewed pilot projects, demonstration projects, or research are asked to present those proposals to the MDC prior to their implementation for review and approval by the MDC."