BATTLE AND NON-BATTLE INJURY DOCUMENTATION: THE RESUSCITATION RECORD

Original Release/Approval		1 Jun 2008	Note: This CPG requires an annual review.		
Reviewed: Sep 2012		Approved:	18 Sep 2012		
Supersedes: Use of Trauma		a Flow Sheets 1 D	ec 2008		
Minor Changes (<i>or</i>)		Changes are	e substantial and	d require a thorough reading of this CPG (or)	
Significant Changes		PI monitoring pl prior Trauma Flo		i-service approved Resuscitation Record replaces and Role 3.	

- **1. Goal**. Obtain complete trauma documentation, including evacuation documentation, on all trauma patients from Role 2 and Role 3 within the CENTCOM AOR.
- 2. Background. The role of trauma documentation within the Joint Theater Trauma System for trauma performance improvement has continuously increased since the Joint Theater Trauma Registry (JTTR) was initiated in 2004. This progression is not unlike the first civilian trauma registries and standardized trauma flow sheets that were developed in the late 1980s. JTTR data acquisition and processing has improved greatly, partly because of the continuing advances (i.e., development of a standardized Resuscitation Record, formerly trauma flow sheet, initiation of Oracle-based registry database, and Level II Access trauma database) that offer new approaches and maximize computer technologies and the deployment of trauma coordinators to Role 3 sites. Data collection that allows theater-wide comparison is important for the continuous learning process and to improve outcomes, standard of care development, analysis of differences in the mechanisms of injury, rescue systems, and approved treatment guidelines.

Although Resuscitation Record documentation can incorporate information from numerous sources (nursing flow sheets, monitors, MEDEVAC run-sheets, I-stat print outs, etc.); if the history taking, physical examination, or decision making is not documented by the trauma team leader, it did not occur. Therefore, good documentation on the Resuscitation Record is most important for care of the individual patient and the system-wide delivery of trauma/critical care to all injured patients within the CENTCOM AOR. It is easy to forget or only capture limited data on the Resuscitation Record when trauma patients spend very little time in the ED prior to heading to the OR. However, it is imperative to document the thought process and to take the time to complete the Resuscitation Record when time permits, even if completed the next day.

Although trauma documentation requirements are well known, it is noted that this is an area in need of improvement. Although not exhaustive, the following are documentation performance improvement areas that repeatedly surface which need careful attention:

- a. Complete set of initial vital signs, including temperature and respiration rate
- b. GCS total score and individual Motor, Verbal and Eye opening scores
- c. Total IV volume (blood, colloid and crystalloid) infused in the ED, even if fluid administration continues after transport
- d. Disposition: Place and time

- e. Arrival time
- f. Mechanism of Injury
- g. Labs transferred to trauma flow sheet (especially HCT, INR, and BE)
- h. Lethal Triad Indicators (Hypothermia, Acidosis, Coagulopathy)
- **3.** Indications for Initiation and Completion of Resuscitation Record. A Resuscitation Record should be initiated on *ALL* patients (battle/non-battle injury coalition forces, ANA, ANP, LN, contractors, etc.) triaged as Immediate. In addition, Resuscitation Record should be completed on all patients seen within the first 72 hours following injury, including but not limited to the following injury causes:
 - a. Building Collapse
 - b. Bullet/GSM/Firearm
 - c. Burn
 - d. EFP
 - e. Fall
 - f. Fire/Flame
 - g. IED
 - h. Inhalation Injury
 - i. Mine
 - j. Mortar/Rocket/Artillery Shell
 - k. Multi-Frag
 - l. MVC
 - m. Sports
 - n. UXO
 - o. Other
 - p. All trauma admissions to any/all Role 3 facilities in the continuum

It is the intent of this guideline that the broadest definition of trauma be used. This should include the majority of patients with single or multi-system injury seen in the emergency department or admitted directly to the ICU and is to be used as the primary method of initial documentation.

- 4. Performance Improvement (PI) Monitoring.
 - a. Intent (Expected Outcomes).
 - 1) All patients in a US lead Role 2 or Role 3 facility have a Trauma Resuscitation Record complete and in the patient's record.
 - 2) Trauma Resuscitation Record Part I Nursing Flow Sheet has complete and accurate documentation from the primary survey in sections 3.1, 3.1, and 3.3.

- 3) Trauma Resuscitation Record has complete and accurate documentation in the patient identification section, i.e. patient name, patient ID/SSN, facility, nurse and provider.
- 4) Trauma Resuscitation Record Part II Physician H & P has complete and accurate documentation in sections 1.3, 1.5 and 6.3.
- b. Performance/Adherence Measures.
 - 1) All trauma patients triaged as immediate or with injuries sustained from one of the causes listed in 2i had the Trauma Resuscitation Record completed and their record.
 - 2) The Trauma Resuscitation Record was completed by the provider and the nurse on every patient expected to be admitted to a Role 3 or actually admitted to a Role 3 facility.
- c. Data Source.
 - 1) Patient Record
 - 2) Joint Theater Trauma Registry (JTTR)
- d. System Reporting & Frequency.

The above constitutes the minimum criteria for PI monitoring of this CPG. System reporting will be performed biannually; additional PI monitoring and system reporting may be performed as needed.

The system review and data analysis will be performed by the Joint Theater Trauma System (JTTS) Director, JTTS Program Manager, and the Joint Trauma System (JTS) Performance Improvement Branch.

5. Responsibilities.

- a. It is the trauma team leader's responsibility to ensure the Resuscitation Record Part II, Physician H&P is complete at Role 2 and Role 3.
- b. It is the responsibility of the nurse assigned to the trauma bay/patient to ensure the Resuscitation Record Part I, Nursing Flow Sheet is completed at Role 3.
- c. A member of the trauma team that is receiving report (CCATT, medevac, ground ambulance) should request a copy of the transport run-sheet and ensure it is included in the patient's record. All times on the Resuscitation Record should be local 24-hour military format (hhmm).

Approved by CENTCOM JTTS Director, JTS Director and CENTCOM SG

Opinions, interpretations, conclusions, and recommendations are those of the authors and are not necessarily endorsed by the Services or DoD.

APPENDIX A Resuscitation Record—Part I Nursing Flow Sheet, page 1 of 5
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		SUSCITATIC art I, Nursing			
1. PATIENT INF 1.1TRAUMA TEAM DAT	ORMATION A	1.4 MODE OF ARRIVAL	1.6 INJURY	1.9 PATIENT CATEGORY	1.10 INJURY CAUSE
	illed Arrived Name	Walked/Carried CASEVAC - Air CASEVAC - Ground MEDEVAC - Air Mission #	CLASSIFICATION Battle Non-Battle Unknown	USA USAF USMC USN	Building Collapse Bullet/GSW/Firearm Bum EFP Fall
Lab/Blood Bank Radiology Pharmacy Consult (i.e., Ortho)		MEDEVAC - Ground Mission # CCATT Ship EVAC	Intrace CATEGORY Immediate Delayed Minimal Expectant	USCG USPHS Civilian - Local Civilian - Other	Fire/Flame IED Inhalation Injury Mine Mortar/Rocket/
1.2 ARRIVAL Date Time of Arrival Time of Injury Date of Injury Transit Time minutes	1.3 EVAC FROM 1st Responder Forward Resuscitative Care Theater Hospital Location	AE Other Other Slunt Burn Penetrating		Contractor EPW NATO - Coalition Non-NATO - Coalition Other	Artillery Shell Multi-Frag MVC Sports UXO Other
2. CARE DONE	PRIOR TO ARRIVAL			1	
Effective? Y	Lower Extremities: Type: CAT SOFTT Other Time On Off 3 R How 1 4 many? 2 3 L How 1 3 4 Effective? Y 3 4 L How 1 3 4 Effective? Y 3 4 L How 1 3 4 2 4	VITALS HE GCS CC Eye _/4 Verbal _/5 Motor _/6 Total _/15 4 P 3 RR 3 BP 4 BP	MORRHAGE WITCOL EASURES Celox ChitoFlex Combat Gauze Direct Pressure	ARMING L Blanket P Body Bag In HPMK C Space Blanket T Other N PREHOSPITALMEDS P C C C C C C C C C C C C C	.6.9REHOSPITAL NTERVENTIONS rehospital Airway ntubated
3. PRIMARY SU 3.1VITALS P RR BP O2Sat Pain Scale (0 - 10) 3.2AIRWAY Patent Stridor	RVEY 3.3 HYPO / HYPERTHERMIACONTE MEASURES Arrival Temp Date Route Oral Avillary Temperature Control Procedure: Bair Hugger Warming Bl Fluid Warmer Cooling Blar Other	C Unlabored C Labored Rectal Retraction Absent Chest Symmetry:	Rales	L Warm L Pink L Dry L Dry L Heart Sounds: Clear Capillary Refill: dline < 2 Second	Cool Hot Pale Cyanotic Moist Diaphoretic Muffled s (normal)
Drooling Obstructed Oral/Nasal Airway BVM Intubated Combi Tube Other	3.4 CPR IN ED Y N Start Time	Alert - Obeys C	Commands erbal Stimuli ainful Stimuli	- In Archite	tric Broselow Tape Color:
PATIENT IDENTIFIC		Indiant Barand #	First	MI	Rank
Patient ID/SSN Fadlity Name	BRNN	ledical Record #	DOB MOS/AF	SC/NEC Deploye	Gender M F
Nurse Name	- solid t	Nurse Sig	en nette statue et		
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*		100/00110-00500302	CITATION	RECOR	D	
4. SECONDARY	SURVEY	Part I,	Nursing Fl	ow Snee	<u>r</u>	
4.1 HEAD / NECK ENT Drainage: Nasal (Color) Ear (Color)		PHEART / THORACIC ythm NSR Tachy/Brady V-fib / V-tach PEA Asystole	4.3 AB DOMINAL/GU Open Wound Flat Obese Distended Tender Non-Tender		EXTREMITIES formities Puls RUE LUE RLE	es Present Motor Sensory Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N Y N
CSF (Halo Test)		Other Strong W = Weak = Strong W = Weak = Doppler A = Absent rotidRL	Rebound Tende Guarding Rigid Unable to Asses Pelvic Binder		ALLERGIES	
Brisk Slugg ish	Y N Br Brisk Ra Sluggish	moralRL achialRL dialRL dalRL	Blood at Meatus/Vagina FAST + desorbe - Equivocal Last Meal @		None Current Me	eds: (List med, dose, & route)
Procedure	Time	Size/Type	Site		Performed By	Results
0 ₂ TherapyLpm	0.4300	Nasal Cannula	Nasal Airway	8		
ET Intubation (Put ad ditional changes in Remarks)	Time	Teeth	_ cm	Dral Nasal		ETCO ₂ Change BBS Post Intubation
C-Collar Placed Time _	0	-Collar Removed Time				
Chest Tube #1	Time	2 2	4	L 🗌 R		Air Blood (cd)
Chest Tube #2	Time	 	_			Air Blood (cc)
Needle Decompression	Time		_ 🗆	L 🗌 R		Air Blood (cc)
Thoracotomy	Time		1 million (1997)	L 🔲 R Iamshell		
Toumiquet	Time	Types	Site			
Eye Shield	Time			Both		
A-line	Time	in ai			10	
Gastric Tub e	Time	<u>1</u>	_ 0	Dra I Nasal		Verified Y N Suction Y N
Urinary	Time	Amount Color FoleySize		Meatus Suprapubic		Heme Dip - / - + Results cc
Other Procedure	Time	Describe				
Other Procedure	Time	Describe				
Hemorrhage Control Measures	Celox ChitoFlex	Combat Gauze	Field Dressing	QuikClo		Unknown Other
PATIENTIDENTIFICA	TION Name: L	ast	First		MI	Patient ID/55N
BRN Facility	ocation	Nurse Name			Nurse Signatur	e
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3			TATION ursing Flo				
4. SECONDARY SU	JRVEY, continue						
4.8 INTUBATION MECH/VEN	T 4.9 ABGs / VBGs						
Time	20 20 STD	Time Fig	D2 pH	pCO2	pO2 BE	HCO3	SAT
MODE:		G	10. 3255			201	
FIO2:		G			5	6-13-	
RATE:		G					
PEEP:	ABG or VE	G	1000	्रांस	<u> </u>	15115	19.07
TV:	ABG or VE	G	11111	2030	8	1217	ta in
4.10 INTRAVENOUS ACCES	S AND FLUIDS		4.11	BLOOD PRODUCT	IS		
Bate 	Site IVE Type /	Amount Up Amount	Un Stop Un 	it# Type	<u>Start</u>	<u>Stop Volum</u>	e Initials
4.12 MEDICATIONS	Total Amount Infused:		4.13 VITAL SIGNS			Pain Sc	ale) Other (ICF
2 160 2 160 2 160		1	1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 				
						43.02	
14 LABS	15 CT Iype Jime	4.17 DISPOSITION	Date:		Host Nation Facility Name:	Coalition	CASE
CPC	Head	Admit	Time:		Charles Services of Adaptive	Priority	
Children (C-Spine		υ 🗆 κw		nsport Vehicle		Urgent
Steer 1	Chest	RID E S. M. D. O	uarters 🔲 Profile	and the second s	12-14 21	ng - 🗌 Med Tech 📃	Critical Care
	Abd				Fixed Win	T 33-33 38-3	CCATT
	Pelvis	RTD Mode of Tran			d: 🗌 Medical	Non-Medical	2005
	Pan Scan*		ory W/C	Evac Mo		Ambulatory	
EG .	Select Pan Scan only if all of	4.18 DEATH INFORM					in spirie boure
1100	ne ablove requested	Time of Death	Mortuary	Affairs Notified?		me to Morgue	
	.16 X-RAY	Death Remarks		. 3	6.97	25	
T&S	<u>Type Time</u> C-Spine	4.19 REMARKS					
T&C x	Chest						
LIA	Abd	-					
11000	Pelvis						
(Constant)	 Ext						
Spedfy Other:							
season dedication of the WA							
PATIENTIDENTIFICATI			First		MI	Patient ID/SSN	
BRN FacilityLoo	cation N	lurse Name		N	lurse Signature		
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Resuscitation Record—Part I Nursing Flow Sheet, page 3 of 5

			ON RECORD sician H&P	
1. HISTORY & PHYSICAL - IN				
1.1 ARRIVAL	1.2 TRIAGE CATEGORY	1.4 INJURY DESC	IPTION	
Date	Immediate Delayed Minimal Expectant	(AB)rasion (AMP)utation (AV)ulsion (BL)eeding (B)urn %TBSA		Pulses Present S= Strong W= Weak D= Doppler
1.3 CHIEF COMPLAINT, HISTORY AND		(C)repitus (C)repitus (D)eformity (DG)Degloving (E)cchymosis (FX)Fracture (F)oreign Body (GSW)Gun Sho (H)ematoma (LAC)eration (PW)Puncture V (SS)Seatbelt Sig (SW)Stab Wour (P)ain (PP)Peppering	found n d	
	4		ANTERIOR POSTERIOR	
1.5 HISTORY AND PHYSICAL Head & Neck:			1.6 PRE / INITIAL PROCEDURES / DIAGNOSTICS Pre / Initial Cric C-Collar/ Time Rem ICP Monitor Cantholysis&Canthol Ventric Tympanic Membrane Eye Shield R	otomy R L es Rupture R L Blood R L
Chest:			Needle Decompression R L Pericardial Output Air Blood (cc) Pericardial Pericardial	describe: - / 🔲 + diocentesis
Abdomen/Back and Spine:			DPL GrossBlood: - / + describe Log Roll Time Back Exam WNL ABNL describe Rectal Exam WNL Weak/Absent Tone G	
Pelvis: Stable	Unstable 🗌 Binde	r	Prostate	
Upper Extremities:			Closed Reduction EXT Fixation Kound Washout Splint	Toumiquet R # L #
Lower Extremities:			Closed Reduction EXT Fixation Wound Washout Splint	Toumiquet R # L #
Interventions Prior to Arrival:				oc Site oc Site oc Site
1.7 PUPILS / VISION Brisk R L Hand Motion Sluggish R L Light Percept NR R L Light Percept Size Right mm Left mm		% Use the Burn Flow	RUE + + +	ROM + / - + / - + / - + / - + / -
PATIENT IDENTIFICATION Na	me: Last		First MI	Rank
Patient ID/SSN	BRN Medic	al Record #	DOB Age	Gender M F
Facility Name	Facility Location		Physician Signature	
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Resuscitation Record—Part II Physician H &P, page 4 of 5

			ATION RECORD Physician H&P	
2. X-RAYS and CT				
21 CT OBTAINED 22X	RAYS OBTAINED	2.3 PENDING STUDIES	2.4 RESULTS (include TEG/Rotem results)	2.5 C-SPINE RESULTS
	-Spine Extremity			CT Scan Normal
	pine DRUE			CT Scan Abnormal
	Thest/Upright LUE			C-Spine cleared based on:
Contraction of the second second	Pelvis 🗌 RLE			Normal Exam, reliable Pt
Pan Scan*	13-25			Normal CT scan, normal exam C-Spine not cleared based on:
and the first				Neuro c/o, abnormal exam
only if all of the	3			Abnormal imaging
above requested Othe	<u> </u>			Unreliable Pt
3. LABORATORY RES	ULTS			
3.1 CBC	<u>.</u> <u>3.</u>	2 CHEMISTRY 7	3.ALFT	3.5 URINALYSIS
	1		Amylase Bili	SpGr Chem
			Alk Phos SGOT	Micro HCG
			LDH SGPT	pH Bact
3.3 PT / INR / PTT	7	7	Other	WBC RBC
		- <u></u>		
4. IMPRESSION		102		
2 3 6. PLAN 6.1 PLAN			5 6	
6.2 TRIAD INDICATORS U			FW	B Requested Yes No
Temp < 96F/36C Ye	s 🗌 No 🛛 INR>	1.4 Yes No	Base Deficit >5 Yes No Dar	nage Control 🗌 Yes 📄 No
6.3 DISPOSITION		CW Transfer	Date: Tin	ne:
7. DNBI/NBI CATEG	ORY			
Injury, Sports	🗌 Injury, Work/Train	ing 🔄 Surgical		
Injury, MVC	linjury, Other			
8. CAUSE OF DEATH	588 CON			
8.1 ANATOMIC			8.2 PHYSIOLOGIC	
Airway Neck	Abdomen D Fv	tremity 🗍 U / 🗍 L	MOF Sepsis	Total Body Disruption
Head Chest	(F 3) (F 3)	ther, Sped fy	CNS Hemori	12
		ina, speary	Other, Specify	inge 🗋 neonig
PATIENT IDENTIFICA	TION Name: Last		First A	AI Patient ID/SSN
-	Location	Physician Name	Physician 1	
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Resuscitation Record—Part II Physician H &P, page 4 of 5

APPENDIX B General Instructions for Resuscitation Record, Page 1 of 5

General Instructions for Resuscitation Record

Purpose: The Resuscitation Record is for documenting a trauma patient's injuries and related medical treatment and resuscitation care provided at DoD medical treatment facilities (MTFs). It is to be used at all DoD MTFs which have a surgical capability or emergency department (ED). A trauma patient is defined as a person who has an injury with the potential of requiring a surgical intervention. The form is comprised of two parts. Part I, Nursing Flow Sheet is completed by the nurse fulfilling the role as a scriber or the nurse providing bed side care. Part II, Physician H&P (History and Physical) is completed by the trauma physician providing care for the patient. The Resuscitation Record becomes part of the patient's permanent DoD medical record.

PART I, NURSING FLOW SHEET

General Instructions:

- To be completed by the nurse fulfilling the role as a scriber or the nurse providing bed side care.
- Time Zones: Record all time local 24 hour military format, hh:mm
 - A+ (plus sign) means positive test result; a (minus sign) means negative test result.

PATIENT IDENTIFICATION (at bottom of each page). As stated.

FACILITY NAME. Record your MTF unit identifier

FACILITY LOCATION. Record FOB, COB, or geographic site

BRN. Battle Roster Number

MOS. Military Occupational Specialty

AFSC. Air Force Specialty Code

NEC. Navy Enlisted Classification

1 PATIENT INFORMATION

- 1.1 TRAUMA TEAM DATA. As stated. Record all time local 24 hour military format, hh:mm
- 1.2 ARRIVAL. As stated.
- 1.3 EVAC FROM. Check all that apply. Location is the facility name.
- 1.4 MODE OF ARRIVAL. Check one. MEDEVAC Air includes DUSTOFF. If Other, describe the method by which the patient arrived, such as PJ or MERT, but not DUSTOFF.
- 1.5 INJURY TYPE. Check all that apply.
- 1.6 INJURY CLASSIFICATION. Check one.
- 1.7 TRIAGE CATEGORY. Check one.
 - Immediate Patients who require rapid, immediate intervention in order to preserve life and/or limb AND are likely to survive because of the intervention--damage control surgery (ex: respiratory obstruction, unstable casualty with chest or abdominal injuries, uncontrolled hemorrhage, hypovolemic shock, emergency amputation)
 - Delayed Patients who require surgery or other specific therapeutic intervention, but who will not be severely compromised if the intervention is delayed to a later time (ex: closed fx without neurovascular compromise, moderate burns of < 50% TBSA, large muscle wounds, intraabdominal and/or thoracic wounds)
 - Minimal Non-Urgent: Minor Injuries; patient can safely care for themselves or be helped by nonmedical personnel. (ex: Minor lacerations, abrasions, fractures of small bones, and minor burns). Can safely wait 12-24 hours or longer for care.
 - Expectant Patients whose injuries are so severe that even with the benefit of optimal medical resources, their survival would be unlikely (ex: massive open head injury with brain matter present, high spinal cord injuries, mutilating explosive wounds involving multiple anatomical sites and organs, second/third degree burns in excess of 60% TBSA, profound shock with multiple injuries and agonal respirations)
- 1.8 VALUABLES FOUND. Check one. Time correlates to checked item.

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	General Instructions for Resuscitation Record
1.9	PATIENT CATEGORY. Check one. If Other, describe the patient's classification as it relates to military, government or civilian organizations. USA. United States Army
	USAF. United States Air Force
	USMC, United States Marine Corp
	USN. United States Navy
	USCG. United States Coast Guard
	USPHS. United States Public Health Services
	Civilian – Local, Includes Host Nation.
	Civilian – Other, Includes Host Nation Police
	EPW. Enemy Prisoner of War
	NATO-Coalition. Joining military forces
	Non-NATO Coalition. Opposing military forces
	Other, Describe not otherwise specified category.
1.10	
1.10	EFP. Explosively Formed Projectile/Penetrator
	IED. Improvised Explosive Device
	Mortar/Rocket/Artillery Shell. Includes Indirect and Direct Fire
	MVC. Motor Vehicle Crash
	UXO. Unexploded Ordnance
2	CARE DONE PRIOR TO ARRIVAL
2.1	PREHOSPITAL TOURNIQUET. Check all that apply.
	SOFTT. Special Operations Forces Tactical Tourniquet
	CAT. Combat Application Tourniquet
	If Other. Describe the type of tourniquet.
	Effective. An effective tourniquet controls active hemorrhage. May be combined with a dressing.
2.2	PREHOSPITAL VITALS. As stated.
2.3	PREHOSPITAL HEMORRHAGE CONTROL MEASURES – Check all that apply. Celox. Granules, applicator or gauze. Stops bleeding by bonding with red blood cells and gelling with fluids to produce a sticky pseudo clot. This clot sticks to moist tissue to plug the bleeding site. Celox is made with chitosan, a natural polysaccharide.
	ChitoFlex. A stuffable wound dressing conducive to narrow wound tracks.
	Combat Gauze. Combat Gauze™ is a 3-inch x 4-yard roll of sterile gauze. The gauze is impregnated with kaolin, a material that causes the blood to clot.
	Direct Pressure. Pressure applied directly to a wound, usually with sterile, low-adherent gauze between the wound and source of bleeding.
	Field Dressing. A casualty's dressing applied to a wound to control hemorrhaging.
	HemCon. Bandage or patch that becomes sticky when in contact with blood, seals the wound and controls the bleeding. HemCon products are made from chitosan, a naturally occurring, bio-compatible polysaccharide.
	QuikClot. Emergency dressing, combat gauze, interventional bandage, QuikClot ACS+™, QuikClot 1st Response™. When QuikClot [®] comes into contact with blood in and around a wound, it takes in the smaller water molecules from the blood. The larger platelet and clotting factor molecules remain in the wound in a concentrated form. This promotes rapid natural clotting and prevents severe blood loss.
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includes non-palpable, but detected with Doppler. Absent means no pulse, non-palpable and not detected with Doppler.		Gene	ral Instructions for Resuscitation Record
 If Other, describe the not otherwise specified hemorrhage control measure. If PREHOSPITAL WARMING. Check all that apply. HPMK. Hypothermia Prevention and Management Kit. Check only if all three components were used: Hat/Hood, Activated Liner, and Outer Shell. If Other. Describe the not otherwise specified warming device. PREHOSPITAL INTERVENTIONS. As stated. PREHOSPITAL INTERVENTIONS. As stated. PRIMARY SURVEY 11 VITALS. As stated. For Pain Scale, enter level that patient indicates their pain to be. Zero indicate the least pain; 10 is the most severe pain. 21 AIRWAY. As stated. If Other, describe the not otherwise specified type of airway. 21 AIRWAY. As stated. 10 forber, describe the not otherwise specified type of airway. 23 AIRWAY. As stated. 24 AIRWAY. As stated. 25 BREATHING. As stated. 26 CIRCULATION. As stated. 27 OFR IN ED. As stated. 28 COLULATION. As stated. 29 COLULATION. As stated. 20 CIPVINE As stated. 21 Grey/Pink <u>3 - 7 Kg</u> 24 - 29 Kg 24 - 29 Kg 24 - 29 Kg 27 Orange <u>24 - 29 Kg</u> 24 - 29 Kg 27 Orange <u>24 - 29 Kg</u> 24 - 29 Kg 27 Orange <u>24 - 29 Kg</u> 28 EECONDARY SURVEY 11 HEAD/NECK ENT. As stated. 24 HEART / THORACIC. 21 HEAD/NECK ENT. As stated. 24 HEART / THORACIC. 27 Rhythm. As stated. In other, describe not otherwise specified rhythm. Pulses. Enter S, W, D, A as appropriate. Doppler includes non-palpable, but detected with Dopple Absent means no pulse, non-palpable and not detected with Doppler. 33 ABDOMINAL/GU. As stated. Unable to Assess includes TAC (Temporary Abdominal Closure). Last meal @. Enter date and time. 28 EXTREMITIES. Check all that apply. For Pulses Present (positive) enter S, W, D, or A. Doppler includes non-palpable. Biot detected with Doppler. 34 ABDO		None. Check if no hem	norrhage control measures.
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4.9 ABGs/VBGs. As stated.	1.7		
	1.8	INTUBATION MECH/\	/ENT. As stated.
	1.9	ABGs/VBGs. As stated	i.

General Instructions for Resuscitation Record, Page 4 of 5

	General Instructions for Resuscitation Record
4.10	INTRAVENOUS ACCESS AND FLUIDS. As stated.
4.11	BLOOD PRODUCTS. As stated. Initials. Legible initials of person who performed task.
4.12	MEDICATIONS. As stated. Initials. Legible initials of person who performed task.
4.13	VITAL SIGNS. As stated.
4.14	LABS. Enter time as stated.
4.15	CT. As stated.
4.16	X-RAY. As stated.
4.17	DISPOSITION. As stated.
4.18	DEATH INFORMATION. If death, as stated. Leave blank if patient is alive.
4.19	REMARKS. Enter additional information relevant to the patient's nursing care.
PAR	T II, PHYSICIAN H&P
Gene	eral Instructions:
	 To be completed by the trauma physician providing care for the patient. Time Zones: Record all time local 24 hour military format, hh:mm A+ (plus sign) means positive test result; a - (minus sign) means negative test result.
DATI	
PAII	ENT IDENTIFICATION (at bottom of each page). As stated.
	FACILITY NAME. Record your MTF unit identifier
	FACILITY LOCATION. Record FOB, COB, or geographic site
	BRN. Battle Roster Number
1	HISTORY & PHYSICAL – INJURY DESCRIPTION
1.1	ARRIVAL. As stated.
1.2	TRIAGE CATEGORY. Check one. Refer to 1.7 for definitions from Part I Nursing Flow Sheet.
1.3	CHIEF COMPLAINT, HISTORY AND PRESENTING ILLNESS. As stated.
1.4	INJURY DESCRIPTION. As stated. Doppler includes non-palpable, but detected with Doppler. Absent means no pulse, non-palpable and not detected with Doppler.
1.5	HISTORY AND PHYSICAL. As stated. Interventions Prior to Arrival is any intervention performed in a prehospital or transferring facility.
1.6	PRE / INITIAL PROCEDURES / DIAGNOSTICS. As stated. Pre means prior to arrival. Cntrl Line is Central Line.
1.7	PUPILS/VISION. As stated.
1.8	BURN. As stated. Describe the cause of burn.
<mark>1.9</mark>	EXTREMITIES. As stated.
2	X-RAYS AND CT
2.1	CT OBTAINED. As stated.
2.2	X-RAYS OBTAINED. As stated.
2.3	PENDING STUDIES. As stated.
2.4	RESULTS. Include TEG/Rotem results.
2.5	C-SPINE RESULTS. As stated.
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General Instructions for Resuscitation Record, Page 5 of 5

APPENDIX C

ADDITIONAL INFORMATION REGARDING OFF-LABEL USES IN CPGs

- **1. Purpose**. The purpose of this Appendix is to ensure an understanding of DoD policy and practice regarding inclusion in CPGs of "off-label" uses of U.S. Food and Drug Administration (FDA)–approved products. This applies to off-label uses with patients who are armed forces members.
- **2. Background**. Unapproved (i.e., "off-label") uses of FDA-approved products are extremely common in American medicine and are usually not subject to any special regulations. However, under Federal law, in some circumstances, unapproved uses of approved drugs are subject to FDA regulations governing "investigational new drugs." These circumstances include such uses as part of clinical trials, and in the military context, command required, unapproved uses. Some command requested unapproved uses may also be subject to special regulations.
- **3.** Additional Information Regarding Off-Label Uses in CPGs. The inclusion in CPGs of off-label uses is not a clinical trial, nor is it a command request or requirement. Further, it does not imply that the Military Health System requires that use by DoD health care practitioners or considers it to be the "standard of care." Rather, the inclusion in CPGs of off-label uses is to inform the clinical judgment of the responsible health care practitioner by providing information regarding potential risks and benefits of treatment alternatives. The decision is for the clinical judgment of the responsible health care practitioner within the practitioner-patient relationship.

4. Additional Procedures.

- a. Balanced Discussion. Consistent with this purpose, CPG discussions of off-label uses specifically state that they are uses not approved by the FDA. Further, such discussions are balanced in the presentation of appropriate clinical study data, including any such data that suggest caution in the use of the product and specifically including any FDA-issued warnings.
- b. Quality Assurance Monitoring. With respect to such off-label uses, DoD procedure is to maintain a regular system of quality assurance monitoring of outcomes and known potential adverse events. For this reason, the importance of accurate clinical records is underscored.
- c. Information to Patients. Good clinical practice includes the provision of appropriate information to patients. Each CPG discussing an unusual off-label use will address the issue of information to patients. When practicable, consideration will be given to including in an appendix an appropriate information sheet for distribution to patients, whether before or after use of the product. Information to patients should address in plain language: a) that the use is not approved by the FDA; b) the reasons why a DoD health care practitioner would decide to use the product for this purpose; and c) the potential risks associated with such use.