REPORT DOCUMENTATION PAGE		Form Approved OMB No. 0704-0188
The public reporting burden for this collection of information is estimated to average 1 hour per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing the burden, to the Department of Defense, Executive Service Directorate (0704-0188). Respondents should be aware that notwithstanding any other provision of law, no person shall be subject to any penalty for failing to comply with a collection of information if it does not display a currently valid OMB control number. PLEASE DO NOT RETURN YOUR FORM TO THE ABOVE ORGANIZATION.		
1. REPORT DATE (DD-MM-YYYY) 2. REPORT TYPE Poster		3. DATES COVERED (From - To) 12/04/2017
4. TITLE AND SUBTITLE Comparison of an Endotracheal Cardiac Output Monitor to a Pulmonary Artery Catheter	5a. CONTRACT NUMBER tery 5b. GRANT NUMBER	
	5c. PRO	GRAM ELEMENT NUMBER
6. AUTHOR(S) Maj Maddry, Joseph	5d. PROJECT NUMBER	
5e. TASK NU		KNUMBER
	5f. WOR	K UNIT NUMBER
7. PERFORMING ORGANIZATION NAME(S) AND ADDRESS(ES) 59th Clinical Research Division 1100 Willford Hall Loop, Bldg 4430 UDSA Lockland TV 78226 0008		8. PERFORMING ORGANIZATION REPORT NUMBER 17510
JBSA-Lackland, TX 78236-9908 210-292-7141		
 9. SPONSORING/MONITORING AGENCY NAME(S) AND ADDRESS(ES) 59th Clinical Research Division 1100 Willford Hall Loop, Bldg 4430 		10. SPONSOR/MONITOR'S ACRONYM(S)
JBSA-Lackland, TX 78236-9908 210-292-7141		11. SPONSOR/MONITOR'S REPORT NUMBER(S)
12. DISTRIBUTION/AVAILABILITY STATEMENT Approved for public release. Distribution is unlimited.		
13. SUPPLEMENTARY NOTES Science and Technology Leadership Conference, Falls Church, Virginia, December 4, 2017		
14. ABSTRACT		
15. SUBJECT TERMS		
16. SECURITY CLASSIFICATION OF: 17. LIMITATION OF a. REPORT b. ABSTRACT c. THIS PAGE ABSTRACT OF PAGES	Clarice	IE OF RESPONSIBLE PERSON Longoria
	19b. TELI	EPHONE NUMBER (Include area code) 210-292-7141 Standard Form 298 (Rev. 8/98)

Approved for public release. Distribution is unlimited.

Comparison of an Endotracheal Cardiac Output Monitor to a Pulmonary Artery Catheter

N Garrett¹, JK Maddry^{2,3}, VS Bebarta⁴, M Reilly⁵, S Boudreau², M Castaneda², K Canellis², A Arana⁶ ¹Geneva Foundation, San Antonio, TX, ²Dept of Emergency Medicine, San Antonio Military Medical Center, San Antonio, TX, ³CREST Program, Wilford Hall Ambulatory Surgical Center, Lackland AFB, TX ⁴University of Colorado-Denver, ⁵Audie L. Murphy VA Hospital, ⁶USAF En route Care Research Center



Background

In combat, initial resuscitation and life saving measures are initiated by securing a patent airway and administering fluid therapy. While methods of fluid resuscitation remain controversial, maintenance of a patent airway and hemodynamic stability as indicated by invasive monitoring can influence the overall outcome of an injured individual. A patent airway may be maintained via an endotracheal tube. The use of invasive monitoring, however, is complicated by several factors including the extent and type of injuries suffered by the patient. It is imperative for those wounded in battle that we explore potential technologies that can aid in the management of effective fluid resuscitation while considering the limitations presented by remote locations and limited resources.

Methods

Power analysis (G*Power 3.1) suggested 8 animals would be sufficient for comparisons. After induction of anesthesia, instrumentation, and stabilization in experiment 1 (hemorrhage), animals were exsanguinated to produce Type III hemorrhadic conditions. Cardiac output (CO) values were collected from the PAC and the ECOM over a 3 hour period. In experiment 2 (hypothermia), swine were cooled to a temperature of 33°C using the Stryker Gaymar TP700 cooling device and CO values recorded from both instruments. The protocol was approved by the Wilford Hall Ambulatory Surgical Center's Institutional Animal Care and Use Committee (FWH 20140100A).

Results Continued



Objective

The purpose of this study was to explore the accuracy and precision of a FDA approved device, the CONMED endotracheal cardiac output monitor (ECOM) [™] apparatus, by comparing it to the Edwards Vigilance II monitor (Edwards LifeSciences, Irvine, CA) pulmonary artery catheter (PAC) under hypothermic and hemorrhagic conditions.

Results

Using GraphPad Prism® to conduct non-linear fit analyses comparing the slopes of the curves for ECOM versus PAC, we found that the curves from the ECOM data were significantly different from the PAC data curves under both conditions, but more pronounced differences were found under hemorrhagic conditions.

Animal model

Controlled hemorrhage model

Conclusions

Although the ECOM apparatus simplifies data acquisition while limiting potential complications associated with the PAC, the ECOM does not appear to reliably reproduce CO values acquired from a traditional PAC under hemorrhagic or hypothermic conditions.