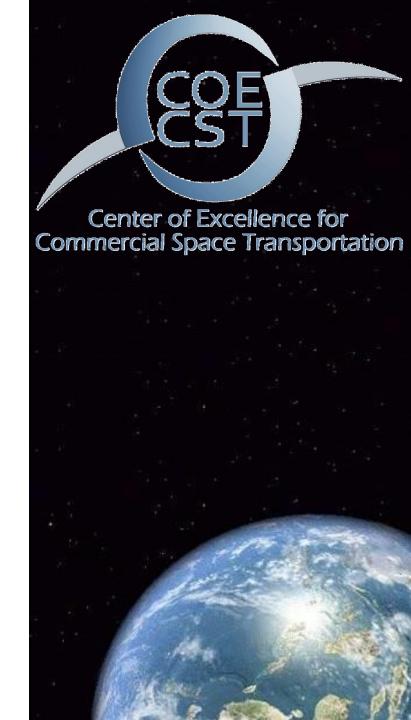
COE CST 3rd Annual Technical Meeting:

Task 255: Validation of Non-Invasive Biomedical Monitoring in Centrifuge-Simulated Suborbital Spaceflight Richard Jennings, MD, MS Tarah Castleberry, DO, MPH









October 30th, 2013

Disclaimers

- Will discuss off-label use of commerciallyavailable physiologic monitoring device, Equivital EQ01-1000 (Hidalgo Ltd., Cambridge, United Kingdom)
- Hidalgo Ltd provides technical expertise and materials to investigators for the purpose of research

Disclaimers

- NASTAR Center partners with UTMB to provide no-cost centrifuge time for research
 - 112 days
 - Value of contribution: 3.4 million
- No other financial relationships to disclose

Overview



- Team Members
- Purpose of Task
- Research Methodology
- Results
- Conclusions
- Next Steps
- Contact Information

Team Members



- PI: Richard Jennings, MD, MS; Tarah Castleberry, DO, MPH (UTMB Aerospace Medicine)
- Co-I: James Vanderploeg, MD, MPH (UTMB Aerospace Medicine)
- Co-I: Rebecca Blue, MD, MPH (UTMB Aerospace Medicine)
- Student: Alejandro Garbino, MD, PhD (Baylor College of Medicine)
- Industry Partner: Brienna Henwood (NASTAR Center)
- Program Manager: Ken Davidian (FAA)
- Technical Monitor: Henry Lampazzi

Purpose of Task



- Purpose:
 - Identify the utility of a commercial, noninvasive, biomedical monitoring device to support operational monitoring needs in a centrifuge-simulated suborbital spaceflight experience.

NASTAR Center





Study Hardware





http://cdn.medgadget.com/wp-content/uploads//2013/01/Equivital-belt-large.jpg

http://vivonoetics.com/wp-content/uploads/2012/04/SEM02-300x210.jpg

Research Methodology



- Physiological parameters, including:
 - Heart rate
 - Respiratory rate
 - Pulse Oximetry
 - Tri-axial acceleration
- Physiologic data were synchronized with standard electrocardiogram monitoring for validation

Research Methodology

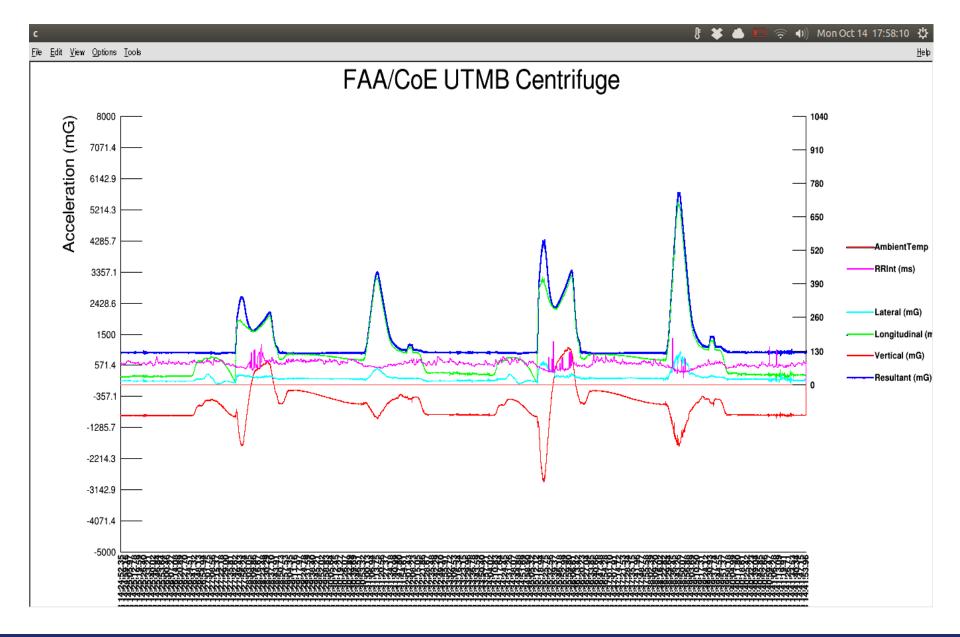


- Instrumented subjects underwent 7 centrifuge runs over two days
 - Day 1 consisted of two +Gz runs (peak=+3.5Gz) and two +Gx runs (peak=+6.0Gx)
 - Day 2 consisted of three runs approximating suborbital spaceflight (combined +Gx and +Gz).

Results



- The device performed well during the centrifuge profiles, providing hemodynamic data with little disruption of signal
- Accelerometer data were reliably synchronized with centrifuge acceleration profiles and served as excellent run-timing markers for hemodynamic data



Conclusion



- Despite the significant acceleration exposures, the monitoring system performed well and provided accurate and reliable hemodynamic monitoring of subjects
- Limitations of the device include difficulty in identifying altered electrocardiographic morphology due to the off-nominal electrode placement, cumbersome analysis techniques, and limited harness size to accommodate larger subjects.

Next Steps



- Complete training and evaluation using the NASTAR centrifuge
- Perform data analysis
- Publish results

Contact Information



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Task 256: Tolerance of Centrifuge-induced G-force by Disease State

Project At-A-Glance

- University: The University of Texas Medical Branch
- Principal Investigator: Tarah Castleberry, DO, MPH
- Student Researchers: Alejandro Garbino, MD, PhD

Relevance to Commercial Spaceflight Industry

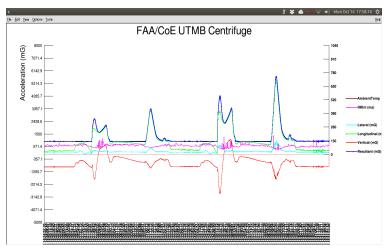
- Commercial spaceflight participants (SFPs) represent a population with potentially significant medical problems that may warrant in-flight medical monitoring
- Commercial SFPs may be hesitant to wear highly invasive, obtrusive monitoring equipment

Statement of Work

- Identify the utility of a commercial, non-invasive, biomedical monitoring device to support operational monitoring needs in a centrifuge-simulated suborbital spaceflight experience
- Volunteers wearing the monitoring device experienced Gforces simulating a commercial spaceflight.







Status

 Complete evaluation using the NASTAR centrifuge

Future Work

- Perform data analysis
- Publish results

