Task Book Report Generated on: 04/23/2024

| E:1 V | EV 2014 | OT 1 T 4 TT 1 4 T | EV 01/07/2015 |
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| Fiscal Year: | FY 2014 | Task Last Updated: | FY 01/0//2015 |
| PI Name: | Bodmer, Rolf Ph.D. | | |
| Project Title: | The Effects of Microgravity on Cardiac Function, Structure and Gene Expression using the Drosophila Model | | |
| Division Name: | Space Biology | | |
| Program/Discipline: | SPACE BIOLOGY | | |
| Program/Discipline Element/Subdiscipline: | SPACE BIOLOGYCellular and molecular biology | | |
| Joint Agency Name: | | TechPort: | No |
| Human Research Program Elements: | None | | |
| Human Research Program Risks: | None | | |
| Space Biology Element: | (1) Animal Biology: Invertebrate | | |
| Space Biology Cross-Element Discipline: | Reproductive Biology Developmental Biology Musculoskeletal Biology | | |
| Space Biology Special Category: | (1) Translational (Countermeasure) Potential | | |
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| City: | La Jolla | State: | CA |
| Zip Code: | 92037-1005 | Congressional District: | 49 |
| Comments: | | | |
| Project Type: | FLIGHT | Solicitation / Funding Source: | 2012 Space Biology NNH12ZTT001N |
| Start Date: | 09/01/2013 | End Date: | 10/31/2015 |
| No. of Post Docs: | | No. of PhD Degrees: | 0 |
| No. of PhD Candidates: | 1 | No. of Master' Degrees: | |
| No. of Master's Candidates: | | No. of Bachelor's Degrees: | |
| No. of Bachelor's Candidates: | | Monitoring Center: | NASA ARC |
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| Flight Program: | ISS | | |
| Flight Assignment: | ISS NOTE: Extended to 10/31/2015 per NSSC information (Ed., 9/15/15) | | |
| Key Personnel Changes/Previous PI: | | | |
| COI Name (Institution): | Bhattacharya, Sharmila Ph.D. (NASA Ames Research Center) Ocorr, Karen Ph.D. (Burnham Institute for Medical Research) | | |
| Grant/Contract No.: | NNX13AN38G | | |
| Performance Goal No.: | | | |
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The detrimental effects of spaceflight on the cardiovascular system are well known. It is believed that these effects may lead to clinically significant risks to astronauts on long duration space missions as well as to the success of these missions themselves. Current studies are limited primarily to human studies and rodent experiments. However, these model systems and human studies have significant limitations that may be addressed by using the well-established Drosophila model. Drosophila have previously been successfully launched into space and a ground-based Drosophila model for cardiac disease and function has been developed. However, the genetically versatile Drosophila model has yet to be used for studying the effects of spaceflight on the cardiovascular system. We are currently preparing flies for a scheduled launch in Sept. 2015 and analyzing data from a prelimnary space flown test of our experimental system. In this proposal we propose to fly groups of Drosophila aboard the International Space Station for approximately 30 days, along with identical on-board 1-g controls as well as ground controls. The Drosophila will require minimal astronaut intervention involving changing feeding trays on 1 or 2 occasions. The samples will be retrieved post-flight **Task Description:** and analyzed using established methods. Heart function, including measurements of diastolic and systolic intervals, heart rate, heart diameters, contractility, and arrhythmias will be recorded. Microscopic and immunohistochemical evaluations of heart morphology will also be carried out. We will also conduct intracellular membrane potential recordings of the heart. Finally, we will analyze mRNA expression with a microarray. The ultimate goal of this research is to obtain data while validating the Drosophila model for studying the effects of spaceflight on cardiac disease and function. The development of such a model would be a potentially significant advancement in the study and understanding of how spaceflight affects the cardiovascular system, and may ultimately lead to countermeasures to prevent them. **Rationale for HRP Directed Research:** Information about cardiac muscle function in microgravity is also expected to provide insights on genetic and molecular changes that occur with muscle atrophy on Earth. For example, we expect to identify basic molecular alterations that are Research Impact/Earth Benefits: associated with muscle atrophy that occurs during prolonged bed rest or muscle disuse in muscular dystrophies. We have sent a preliminary set of flies to the International Space Station (ISS) and have received back live flies that were born in space. We are currently analyzing the data from these flies and performing RNA sequencing of muscle and nervous tissue. This preliminary run has and will continue to provide important information (concerning numbers of flies we need and which genes to test further) that will allow us to maximize the data output from the next scheduled flight in Task Progress: September. We are also refining our abilities to extract data from the limited numbers of flies that return from the ISS. We have also been testing the new Fruit Fly Lab 1 flight hardware that will be used on our next mission to see if it will provide similar results as the initial mission that used only a Nanoracks box.

Description: (Last Updated: 06/23/2023)

Bibliography Type: