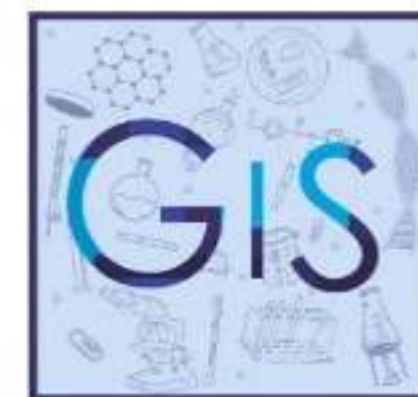




Search for this author in: [Google Scholar](#)

- 1-Jhan Sebastián Saavedra-Torres
- 2-Luisa Fernanda Zúñiga-Cerón
- 3-Luisa Fernanda Mahecha Virgúez
- 4-Alicia Andrea Ortega Narváez
- 5-María Virginia Pinzón Fernández
- 6-Nelson Adolfo López Garzón



# INFORMATION CIRCULAR: IMMUNE SYSTEM IN SPACE.



## Conflicts of interests

None stated by the authors

## Financing

None stated by the authors.

**While** it has been shown that decades of astronauts and cosmonauts suffer from immune disorders both during and after spaceflight, the underlying causes are still poorly understood, due in part to the fact that there are so many variables to consider when investigating the human immune system in a complex environment.

Immune system alterations have been previously characterized as essentially a postflight phenomenon, in animal studies or using terrestrial modeled microgravity or in-flight cell culture. The immune system is a complex weaving of biological structures and processes.

The immune response is affected by the spatial radiation, either solar or cosmic, which can not be contained entirely by the structure of the ship, stress, neuroendocrine response, changes in the pattern of sleep and exposure to the contaminants themselves Inside the ship. Studies have shown in microgravity there are immune system modifications. This may create an environment where, in some crew members, rashes, unusual allergies and latent virus reactivation may present themselves.

Invertebrates have become popular models for studying human disease because they are cheap, highly amenable to experimental manipulation, and have innate immune systems with a high genetic similarity to humans. Fruit flies (*Drosophila melanogaster*) have been shown to experience a dramatic shift in immune gene expression following spaceflight, but are still able to fight off infections when exposed to bacteria.

Extended exposure to radiation and microgravity in space has been linked to astronauts developing chronic diseases upon returning to Earth.

Data generated early in NASA's Integrated Immune study indicated that the distribution of immune cells in the blood of crew members aboard the space station is relatively unchanged during flight. However, they also revealed that some cell function is significantly lower than normal, or depressed, and some cell activity is heightened. In a sense, the immune systems of crew members are confused.

Researchers are also finding latent viruses are reactivating, but do not cause sickness in crew members. Evidence of viral 'shedding', virus DNA present in otherwise healthy individuals, has been found in crew member blood, urine and saliva samples. This can happen anytime the immune system is weakened in microgravity or even in stressful situations on Earth.

The reactivation of latent herpesviruses, including Epstein-Barr virus (EBV), cytomegalovirus (CMV), and varicella zoster virus (VZV), has been well documented in astronauts participating in short-duration spaceflight.

The reactivation of latent herpesviruses in astronauts was recently found to correlate with altered immunity. Should immune alterations persist for the duration of a human Mars mission, it could elevate specific clinical risks to crewmembers including infectious disease, allergies and hypersensitivities, autoimmunity, altered wound healing, and the consequences of persistent latent herpes virus reactivation.

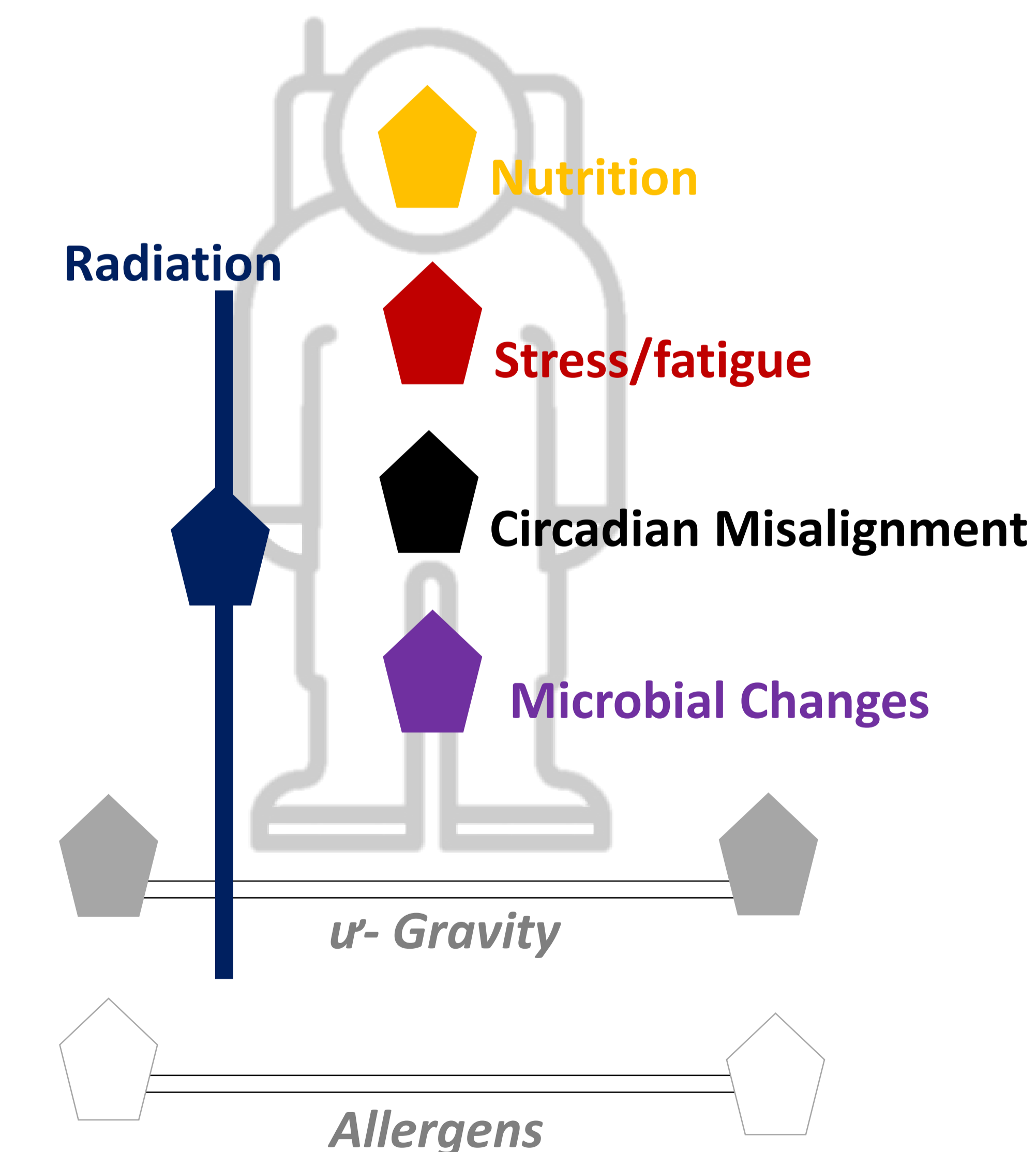


Expedition 30 crew members Dan Burbank and Andre Kuiper perform blood draws for the Integrated Immune investigation. Data collected during the Functional Immune study will build upon findings from this investigation.  
**Credits: NASA**



Astronaut Doug Wheelock collecting saliva sample during spaceflight operations. Image - **Credits: NASA**

*Risk of adverse event of health due to Immune response is altered in space travel*



**The** Gram-negative pathogen *Serratia marcescens* has been shown to potentially cause significant infections in humans and in insect models on Earth. Our recent findings also showed that *S. marcescens* shows an increase in virulence after a short period of growth in the spaceflight environment, which raises initiatives to find the correlation between space environment and the increased virulence. Because we know that the health of astronauts is immunocompromised in space, it is possible that the combination of increased bacterial virulence and the weakened immune system will cause astronauts to be more susceptible to chronic diseases in extended spaceflight.

Additional spaceflight experiments have also provided greater detailed information by investigating specific niches aboard spacecraft or using alternative methodologies beyond the culture-based isolation historically used.

Generally, the data indicate that the potable water, air, and surfaces to which the crew are exposed are free of obligate pathogens; however, opportunistic pathogens such as *P. aeruginosa*, *Stenotrophomonas maltophilia*, and *S. aureus* are not uncommon.

The primary post-infection countermeasure during spaceflight is the use of antibiotics; however, several spaceflight experiments have provided evidence suggesting alterations in antibiotic resistance when microorganisms are cultured during spaceflight.

Investigations of immunity immediately after spaceflight may be confounded by the high-G reentry and stressors of readaptation to terrestrial gravity following prolonged deconditioning. Some studies have indicated that the function of various immune subpopulations may be depressed either following or during short-duration spaceflight.

**Recommended Reading:** NASA (Laurie J. Abadie, Charles W. Lloyd, Mark J. Shelhamer, NASA Human Research Program) - see link: <https://www.nasa.gov/hrp/bodyinspace>

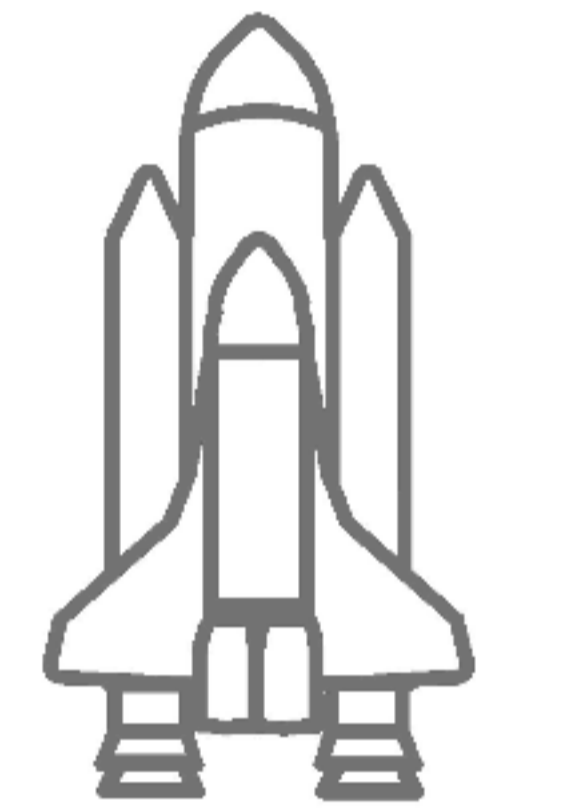
**REFERENCES**

1. Kacena MA, Todd P. Gentamicin: effect on *E. coli* in space. *Microgravity Sci Technol.* 1999; 12(3-4):135-7.
2. Crucian BE, Stowe RP, Mehta SK, Yetman DL, Leal MJ, Quiariarte HD et al. Immune status, latent viral reactivation, and stress during long-duration head-down bed rest. *Aviat Space Environ Med* 2009; 80: A37-A44.
3. Stowe RP, Sams CF, Pierson DL. Effects of mission duration on neuroimmune responses in astronauts. *Aviat Space Environ Med* 2003; 74: 1281-1284.
4. Juergensmeyer MA, Juergensmeyer EA, Guikema JA. Long-term exposure to spaceflight conditions affects bacterial response to antibiotics. *Microgravity Sci Technol.* 1999; 12(1):417.
5. Gmunder FK, Konstantinova I, Cogoli A, Lesnyak A, Bogomolov W, Grachov AV. Cellular immunity in cosmonauts during long duration spaceflight on board the orbital MIR station. *Aviat Space Environ Med* 1994; 65: 419-423.
6. Stowe RP, Sams CF, Pierson DL. Adrenocortical and immune responses following short- and long-duration spaceflight. *Aviat Space Environ Med* 2011; 82: 627-634.
7. Nash PV, Konstantinova IV, Fuchs BB, Rakhmilevich AL, Lesnyak AT, Mastro AM. Effect of spaceflight on lymphocyte proliferation and interleukin-2 production. *J Appl Physiol* 1992; 73: 1865-1905.
8. Mehta SK, Cohrs RJ, Forghani B, Zerbe G, Gilden DH, Pierson DL. Stress-induced subclinical reactivation of varicella zoster virus in astronauts. *J Med Virol* 2004; 72: 174-179.
9. Payne DA, Mehta SK, Tying SK, Stowe RP, Pierson DL. Incidence of Epstein-Barr virus in astronaut saliva during spaceflight. *Aviat Space Environ Med* 1999; 70: 1211-1213.

# INFORMATION CIRCULAR: IMMUNE SYSTEM IN SPACE.



NASA astronaut Kate Rubins removes samples from the Minus Eighty-Degree Laboratory Freezer for ISS (MELFI). Blood, saliva and urine samples will be stored in MELFI until they can be transported back to Earth for analysis.  
**Credits: NASA**



**Consult** reports and evidence of human research of NASA "Human Research Roadmap":

- <https://humanresearchroadmap.nasa.gov/explore/>
- <https://www.nasa.gov/content/study-reveals-immune-system-is-dazed-and-confused-during-spaceflight-u>
- [https://www.nasa.gov/mission\\_pages/station/research/news/functional-immune](https://www.nasa.gov/mission_pages/station/research/news/functional-immune)

**HUMAN EXPLORATION**  
NASA's Path to Mars

MISSION TYPE	MISSION DURATION	RETURN TO EARTH
<b>EARTH RELIANT</b>	MISSION: 6 TO 12 MONTHS	RETURN TO EARTH: HOURS
<b>PROVING GROUND</b>	MISSION: 1 TO 12 MONTHS	RETURN TO EARTH: DAYS
<b>MARS READY</b>	MISSION: 2 TO 3 YEARS	RETURN TO EARTH: MONTHS

Mastering fundamentals aboard the International Space Station

U.S. companies provide access to low-Earth orbit

Expanding capabilities by visiting an asteroid redirected to a lunar distant retrograde orbit

The next step: traveling beyond low-Earth orbit with the Space Launch System rocket and Orion spacecraft

Developing planetary independence by exploring Mars, its moons and other deep space destinations

www.nasa.gov

**Credits: NASA**