Footprints and Fingerprints: Microbiome on the Moon

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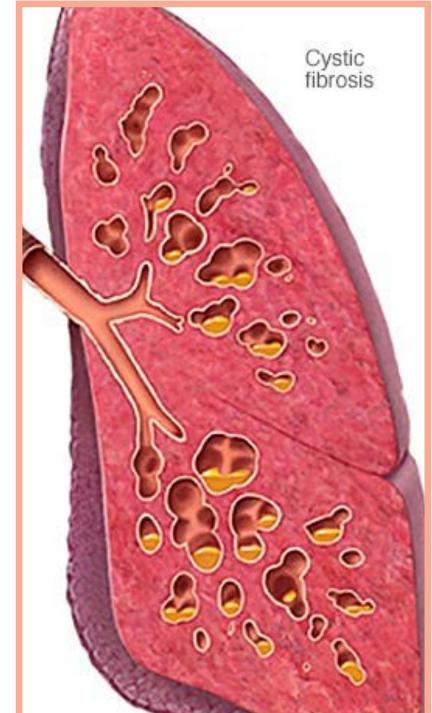
Micah Hamady¹, Jeffrey J Walker², J Kirk Harris³, Nicholas J Gold² & Rob Knight⁴

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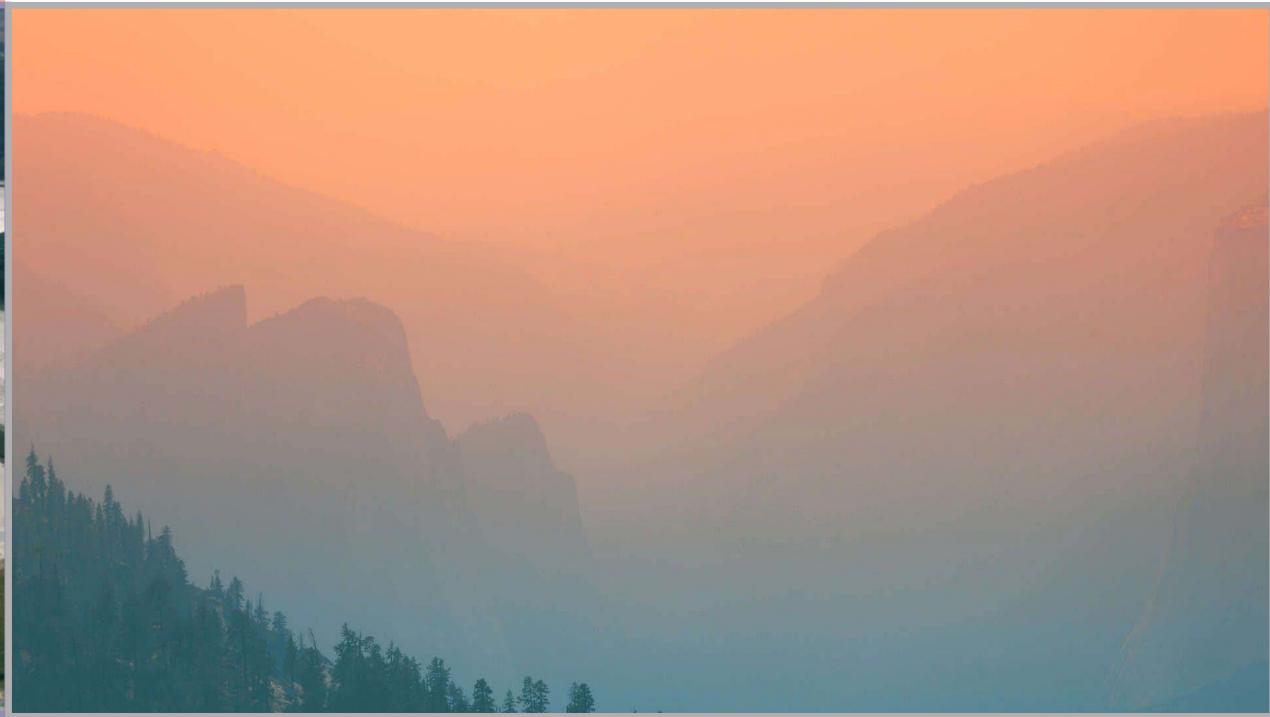




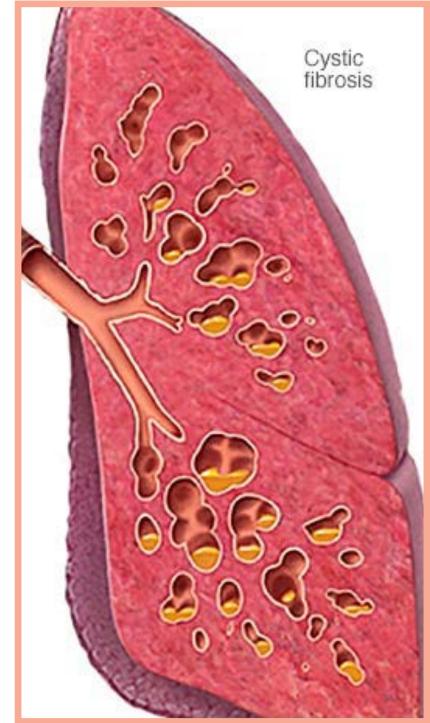
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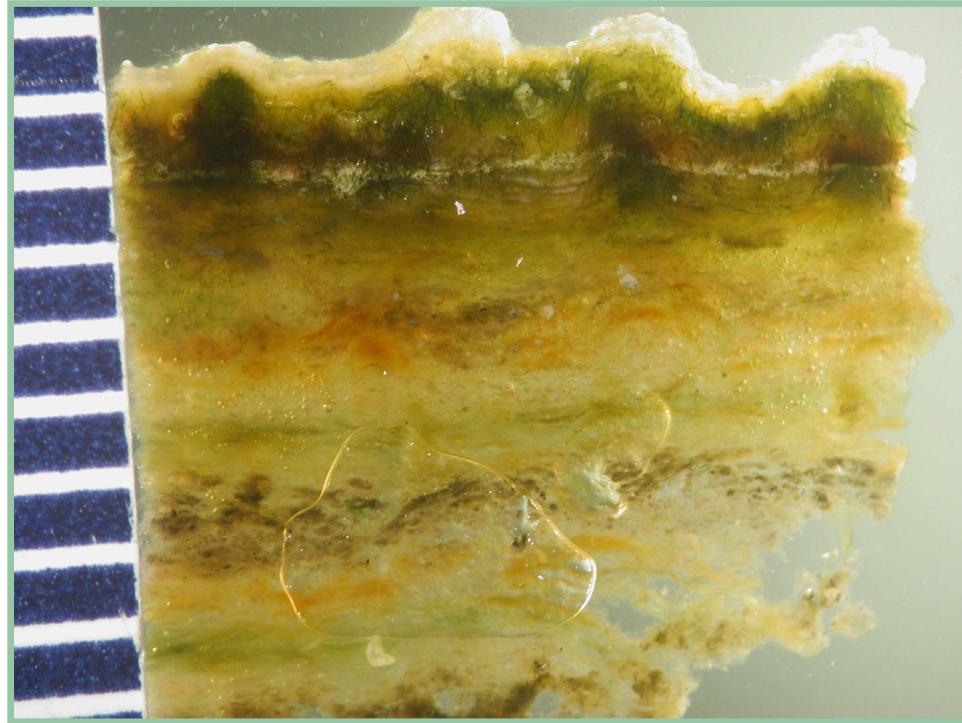




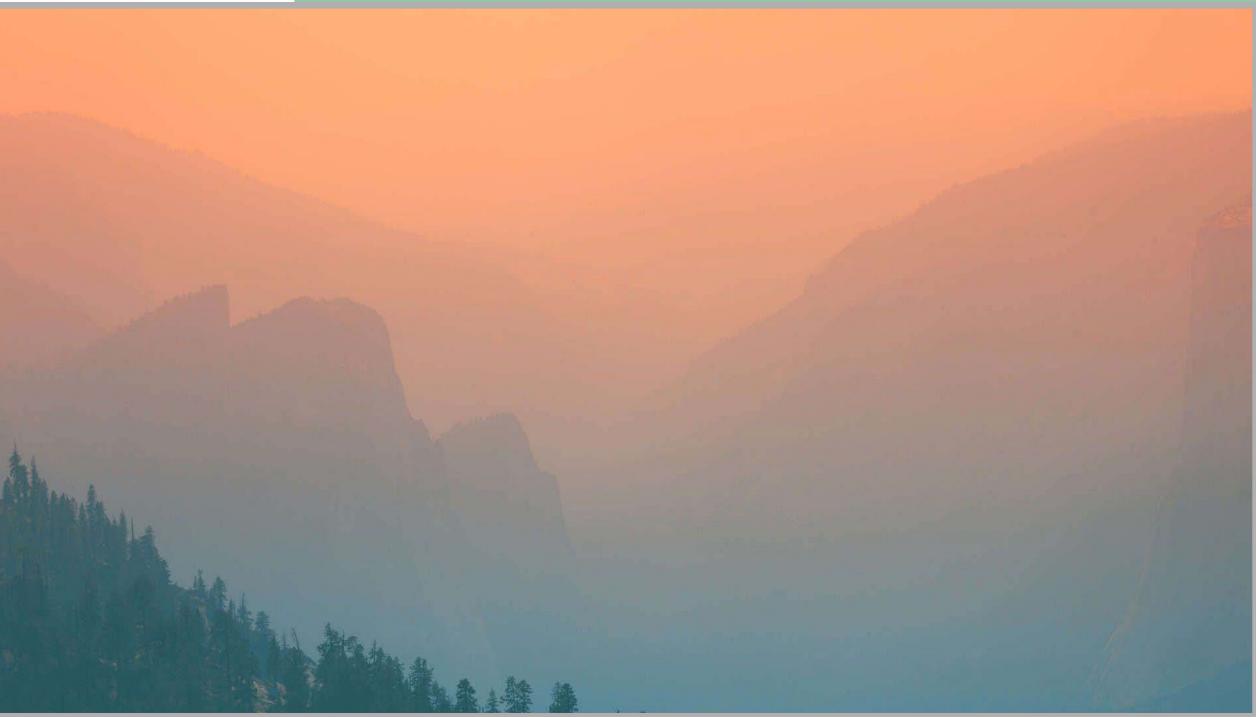


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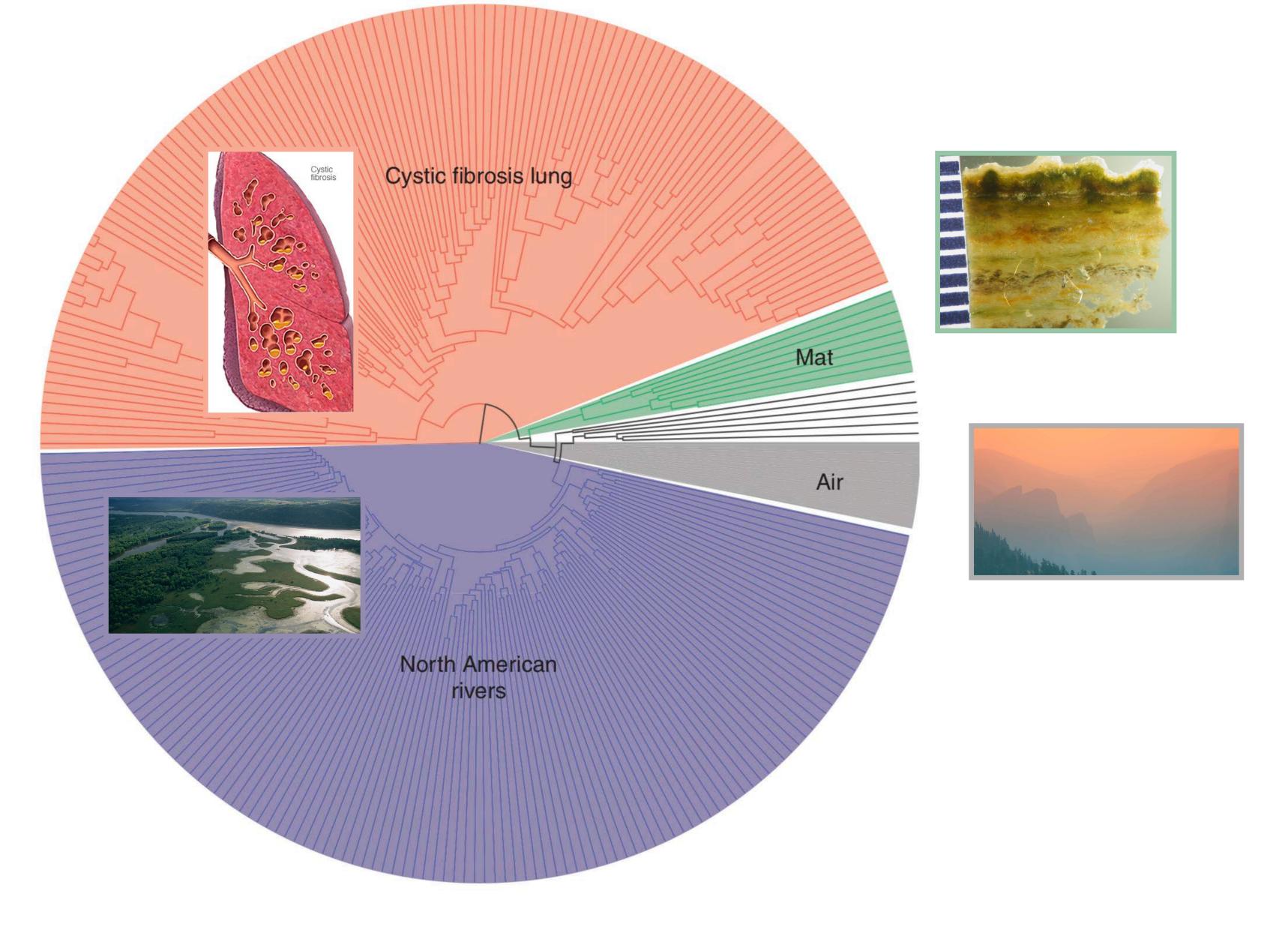


Figure 2 | UniFrac clustering by community was essentially perfect with sequences from pyrosequencing. Samples from cystic fibrosis lung, Guerrero Negro microbial mat, air and North American rivers cluster by environment type.

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\$4,000,000

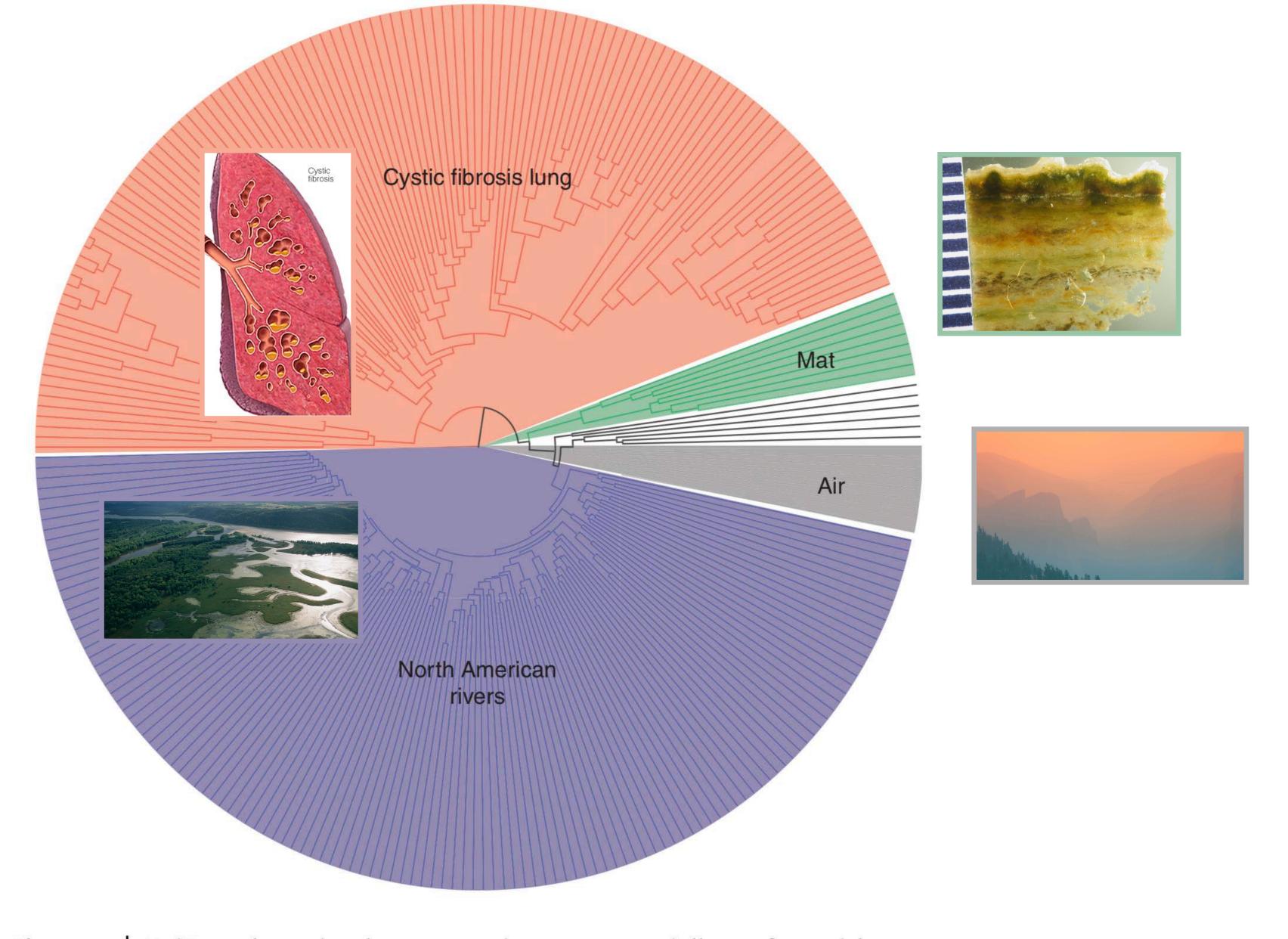


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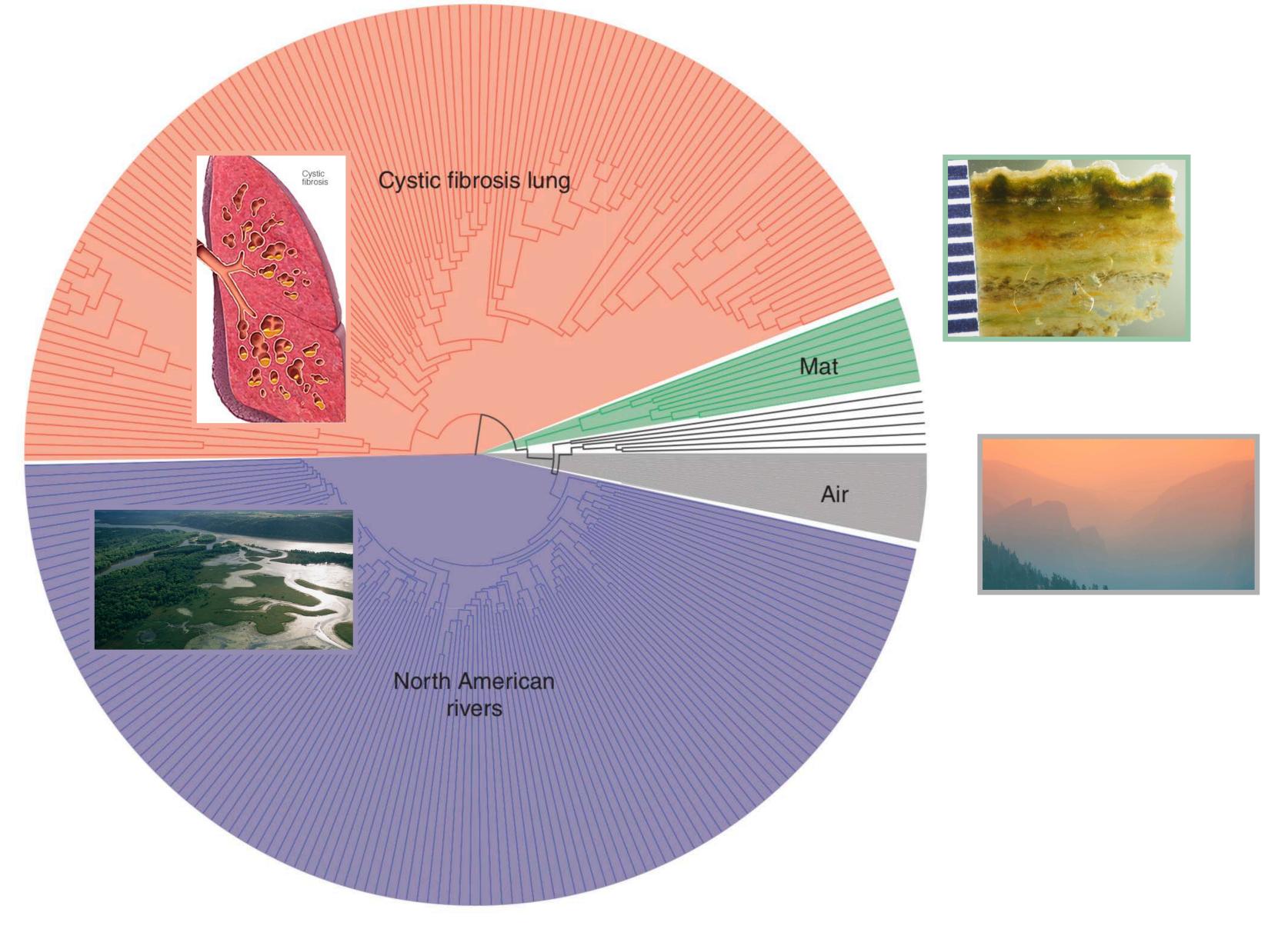


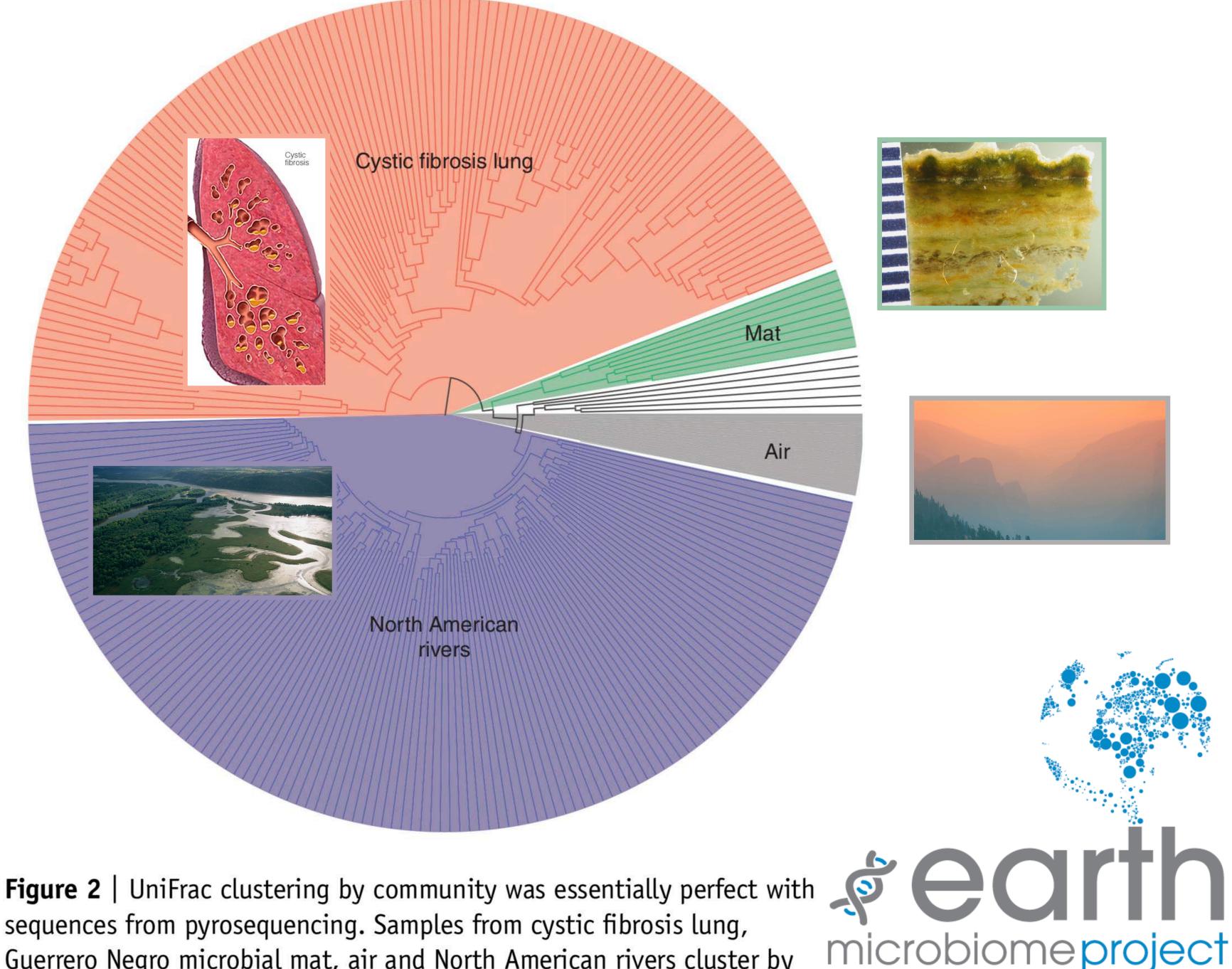
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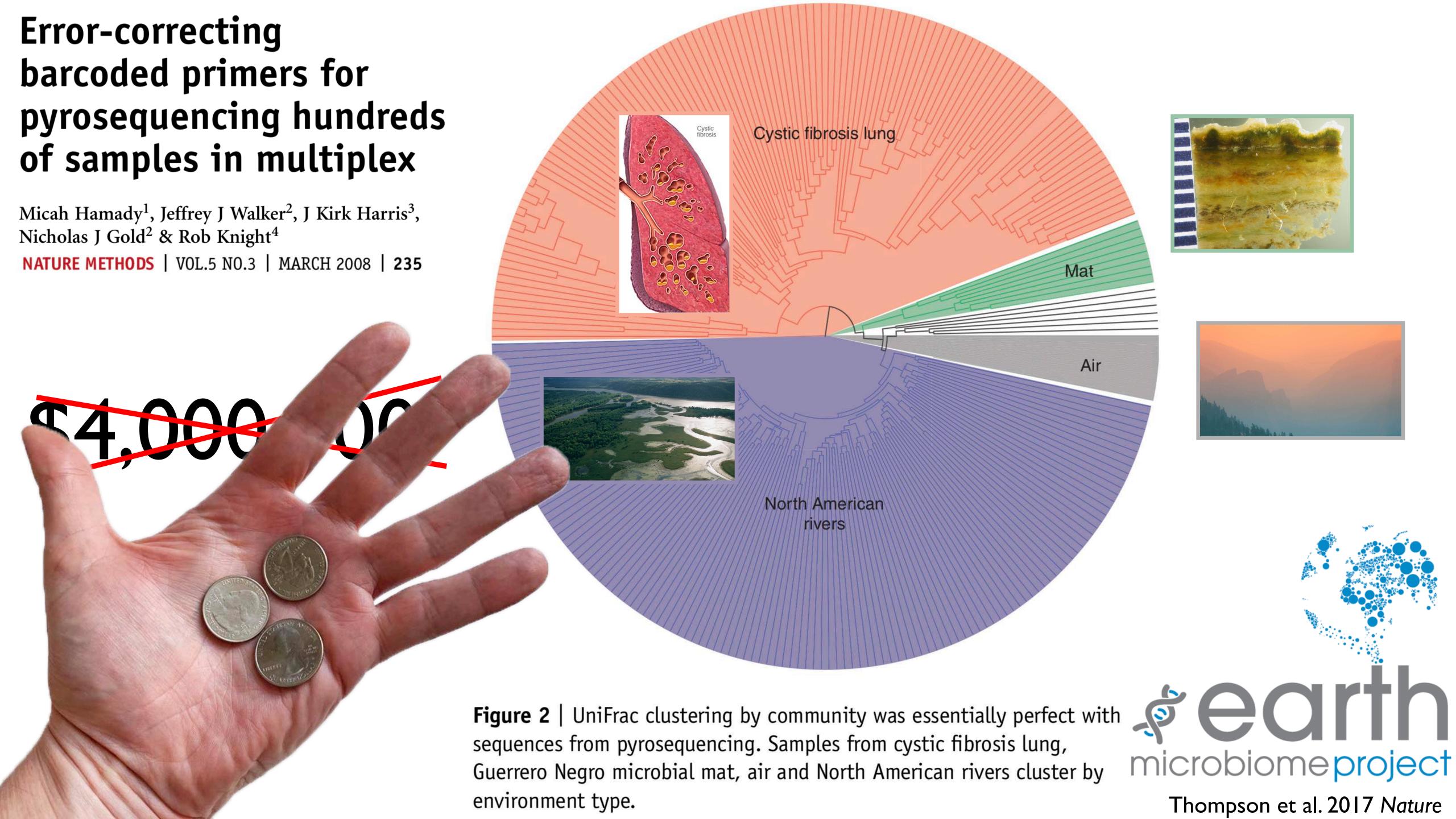
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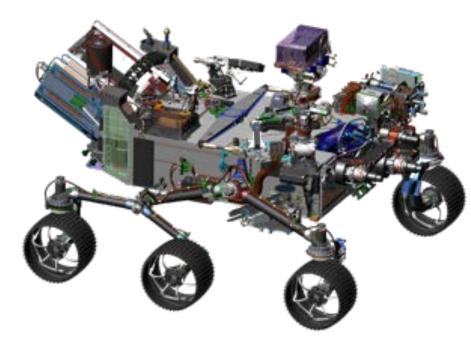
Thompson et al. 2017 Nature

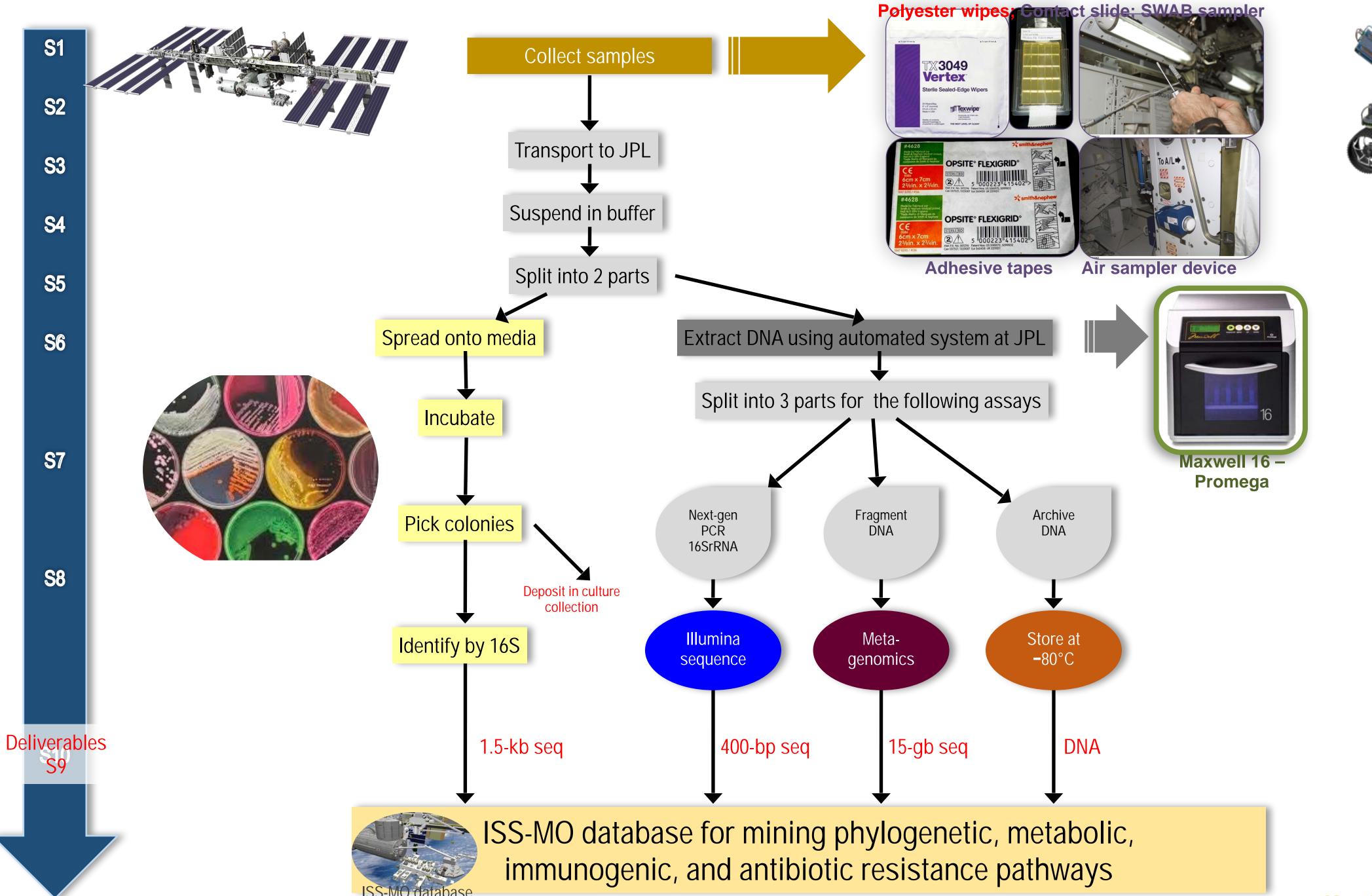


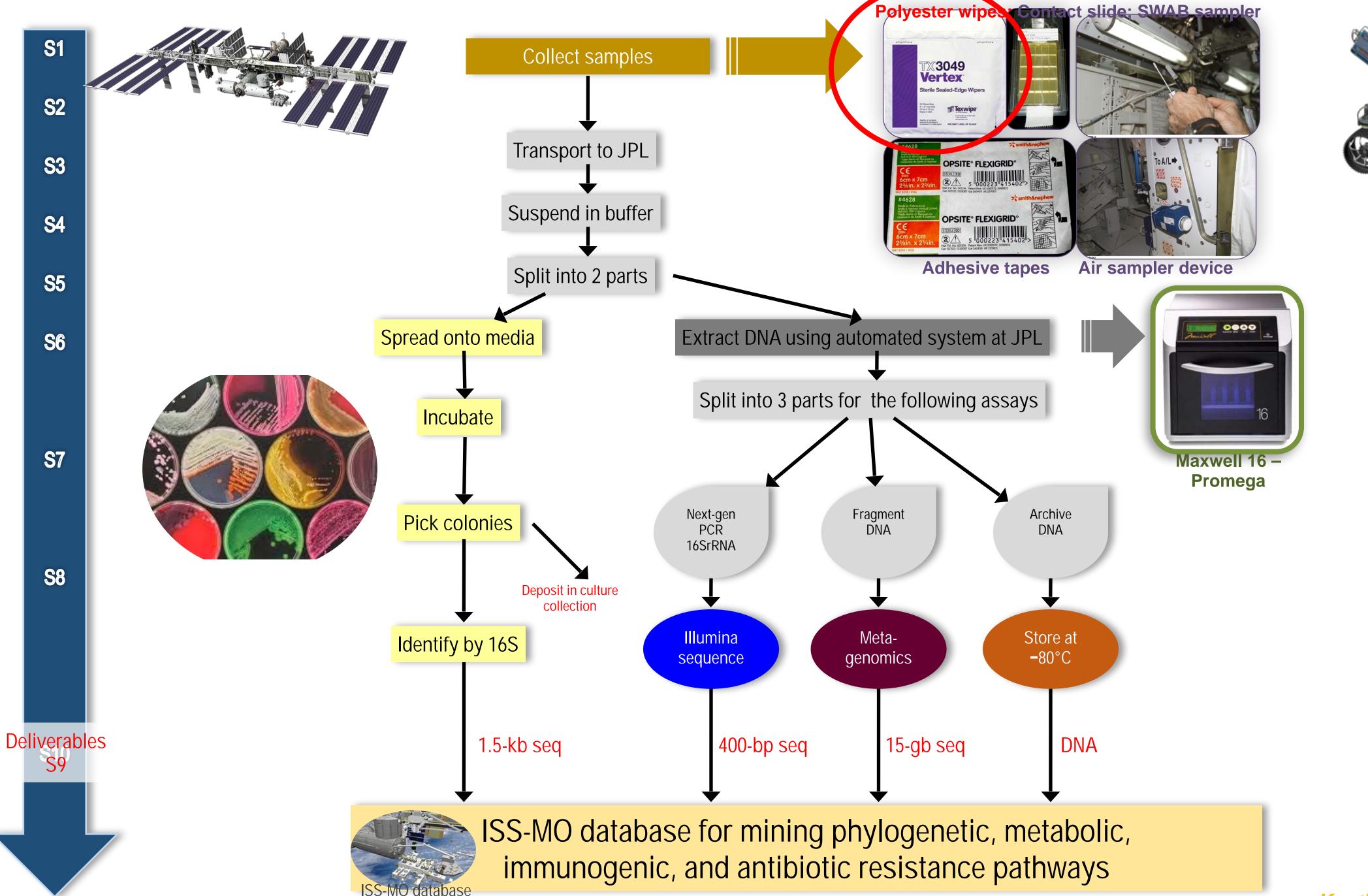
What does the microbiome have to do with lunar exploration?

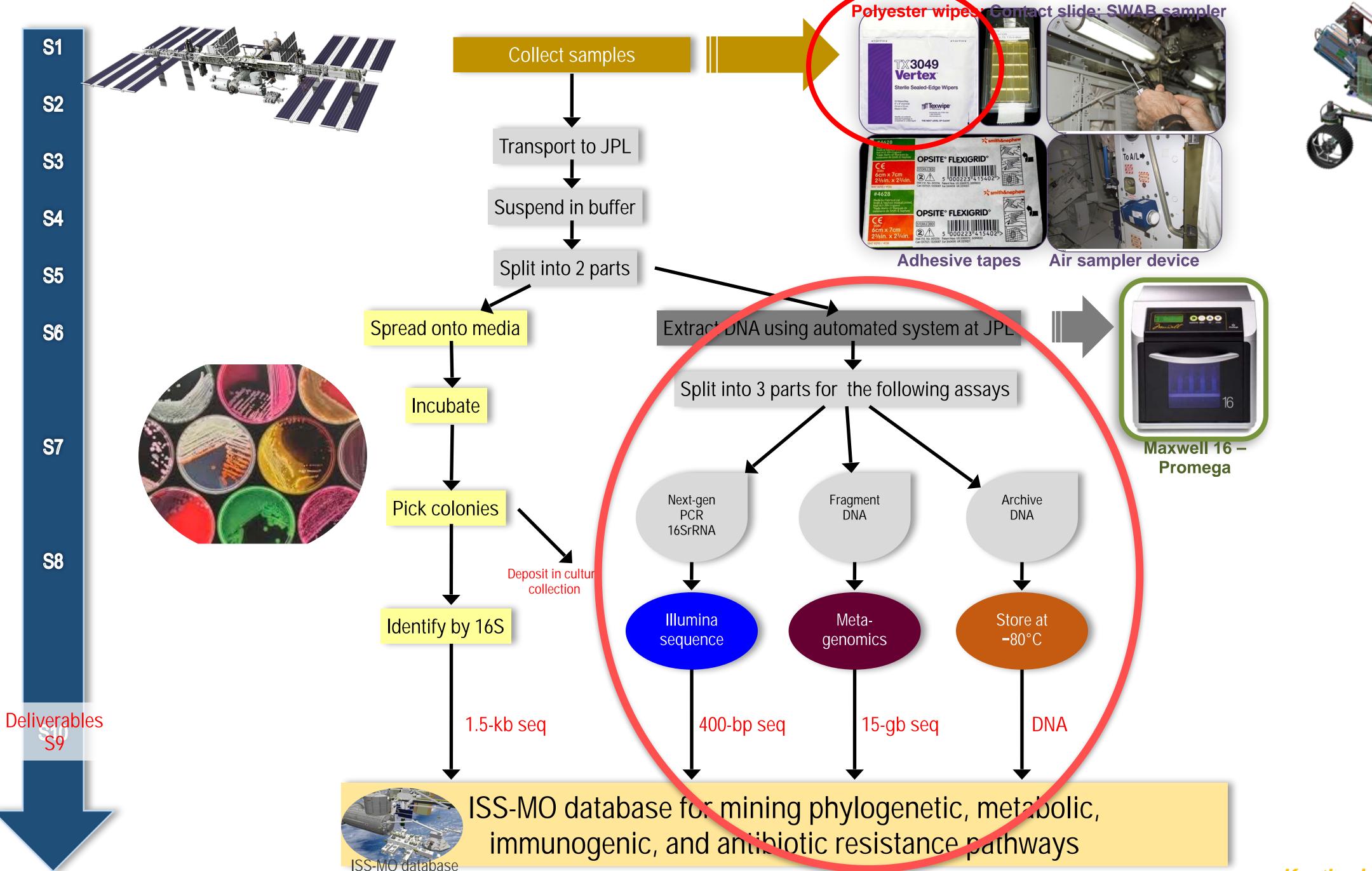
1. Planetary protection: let's not find life that we put there



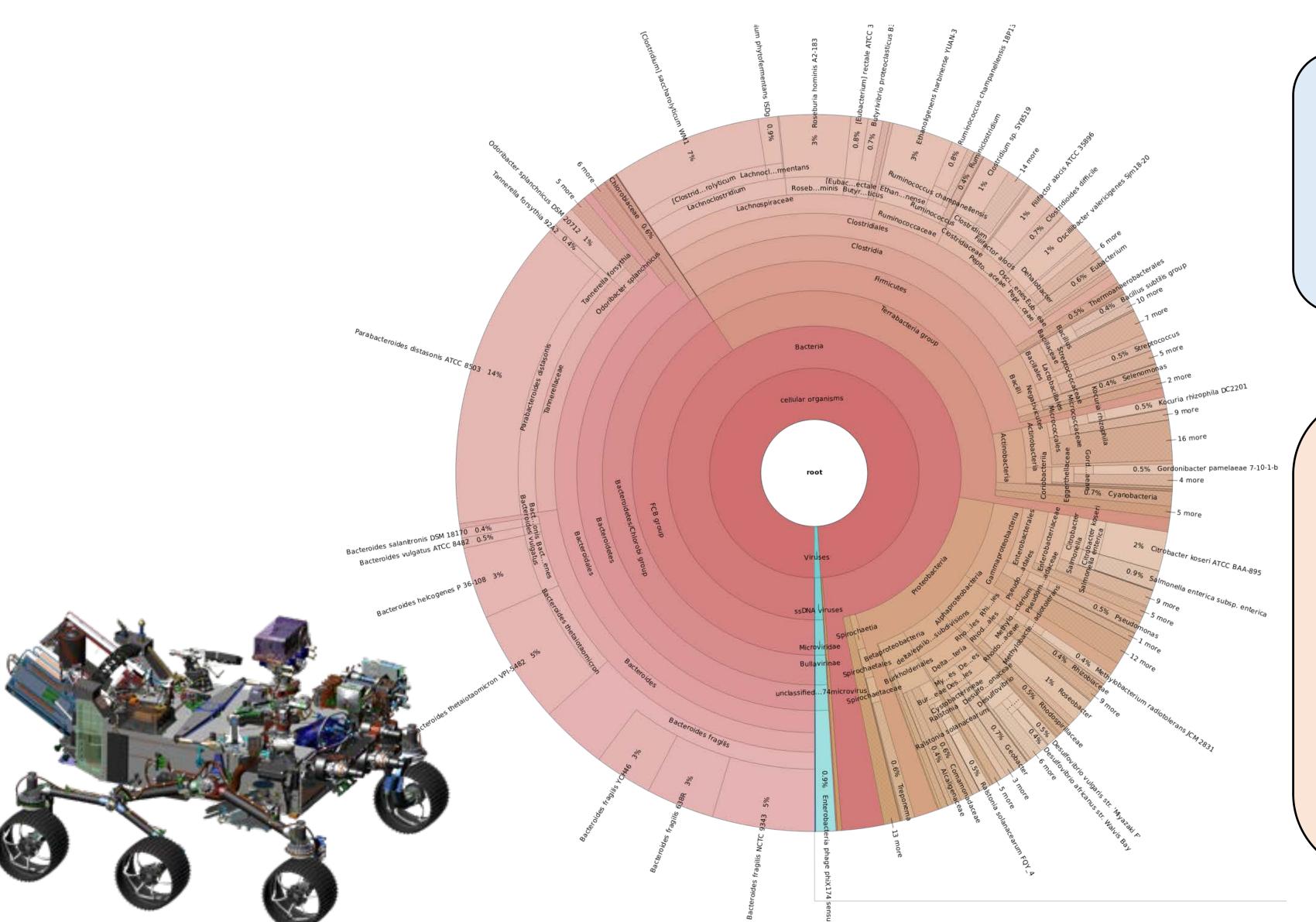








Mars 2020 JPL SAF associated microbiome (a shotgun metagenome approach)



A bird eye view of SAF Microbiome. Red – Brown portion represents Bacteria while blue represent Fungi, Archaea and Viruses.

Bacteria: 5194

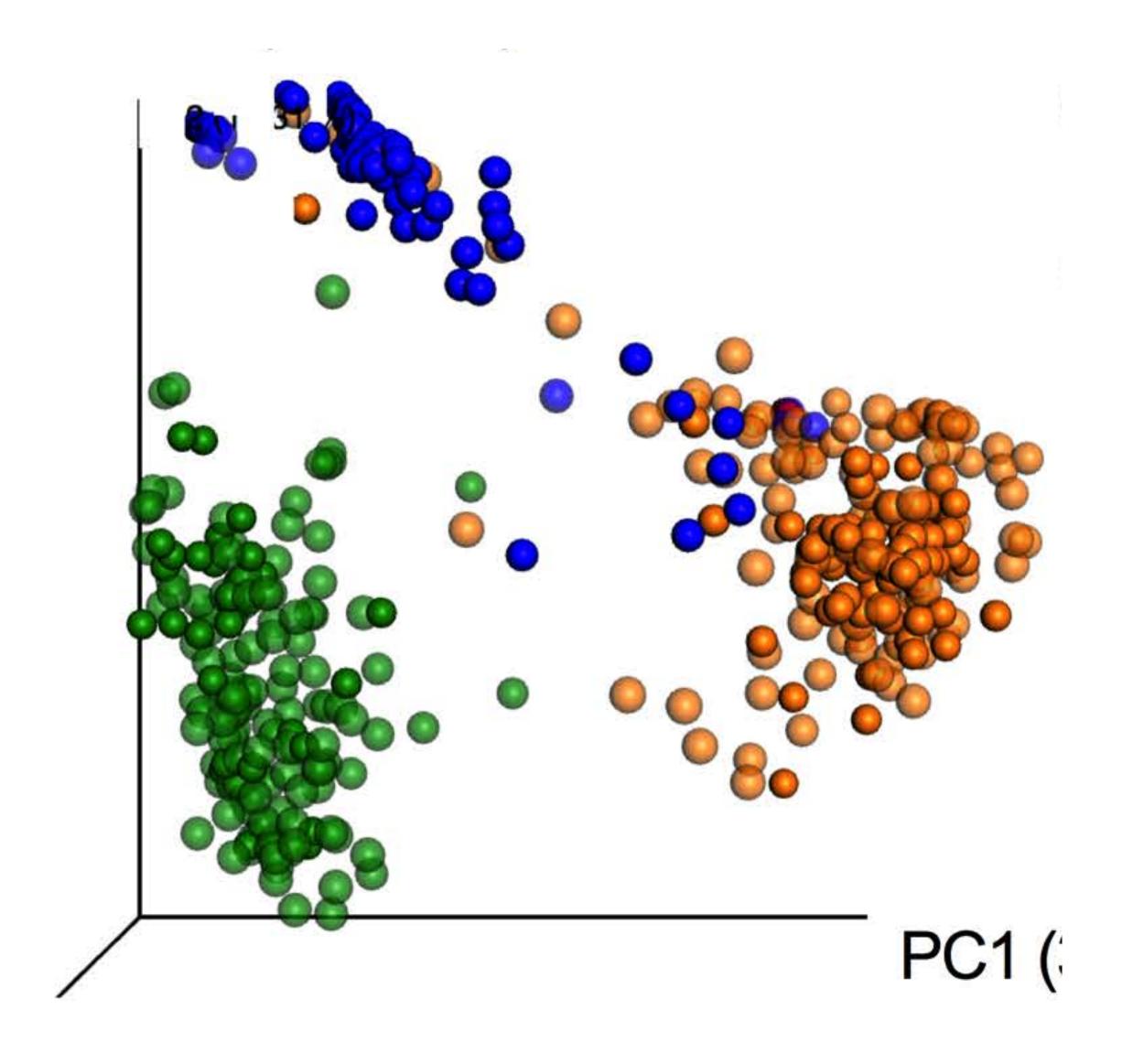
Fungi: 282

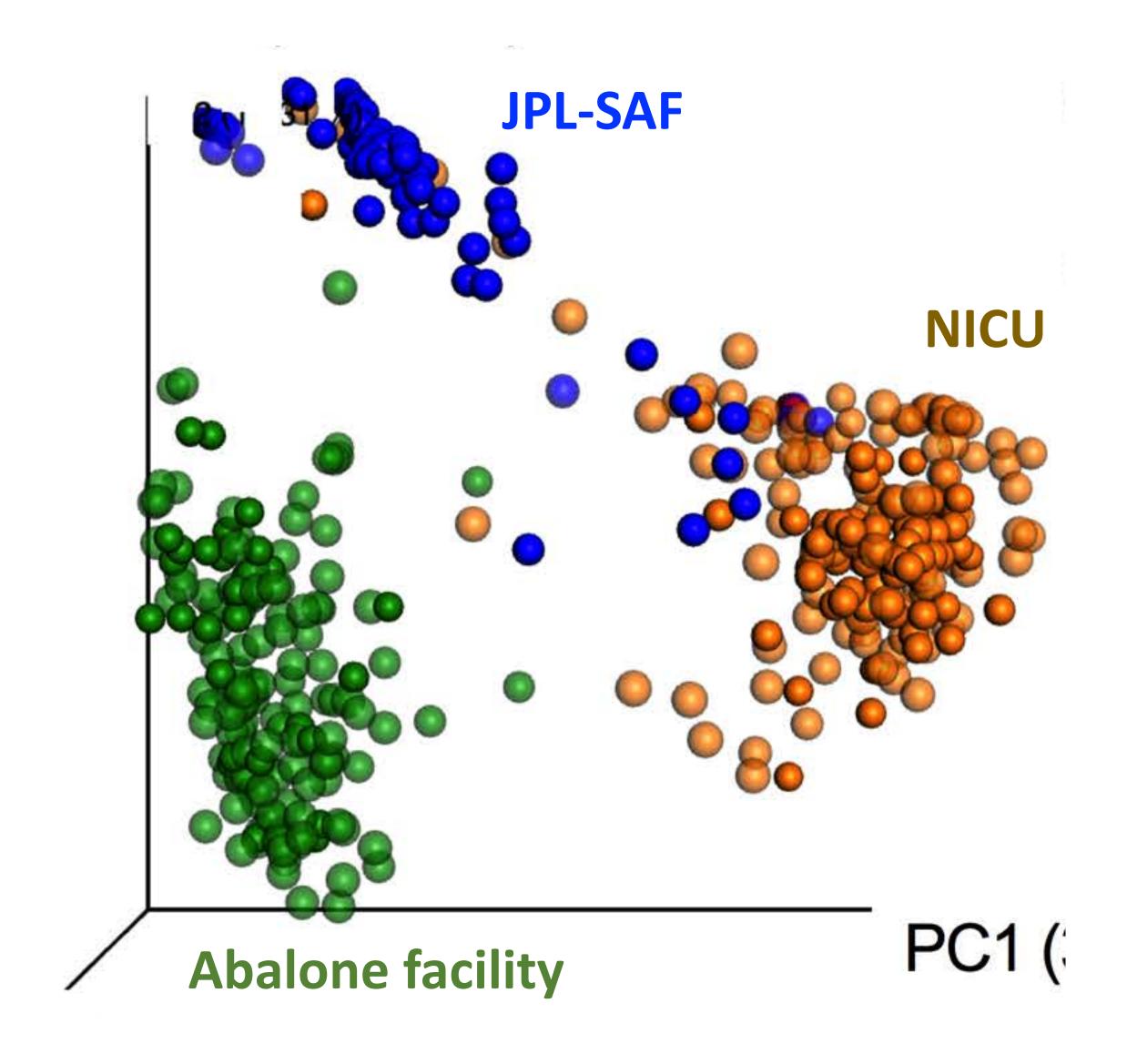
Archaea: 38

Viruses: 63

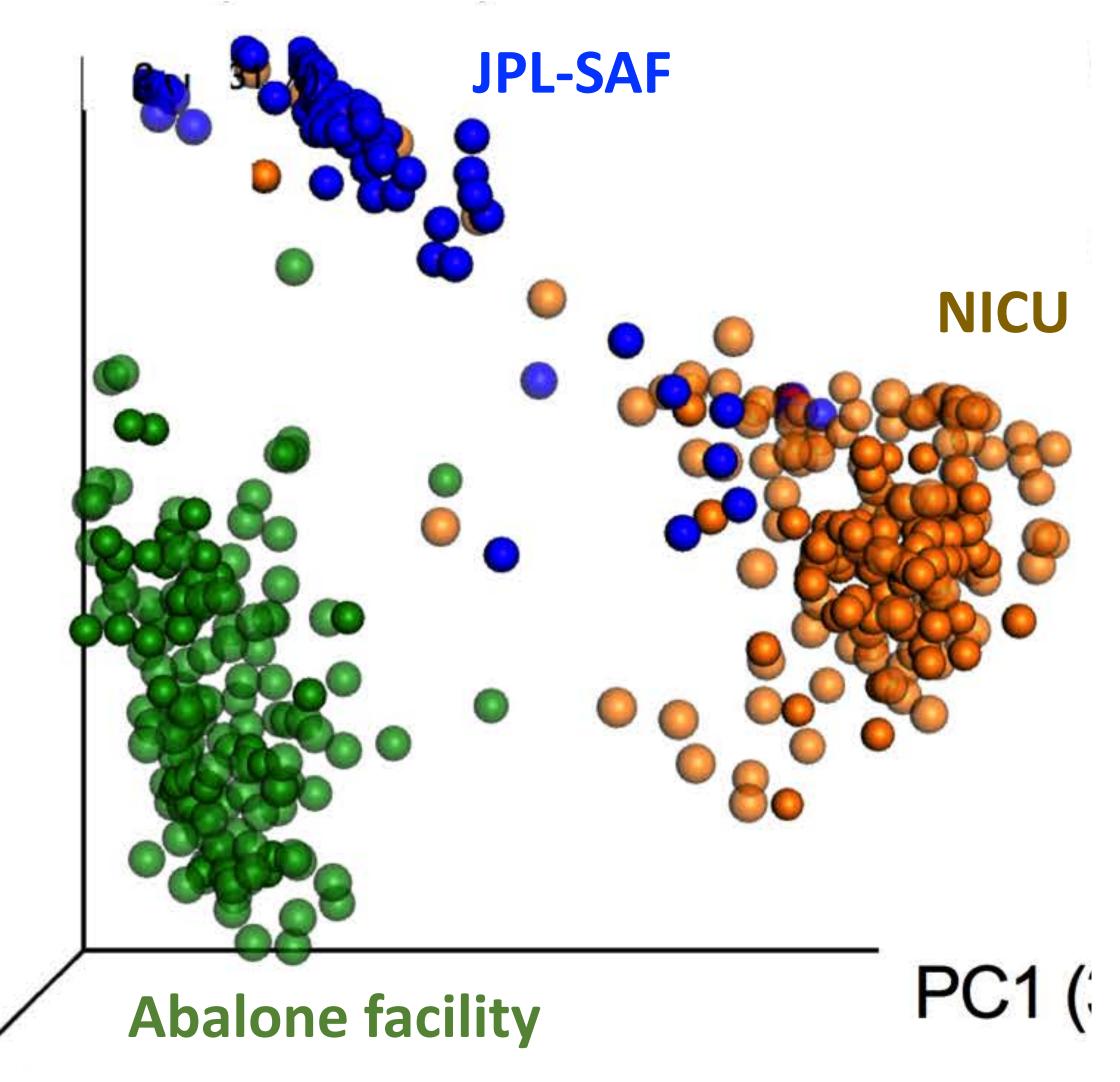
PP relevant Group

Firmicutes: 891

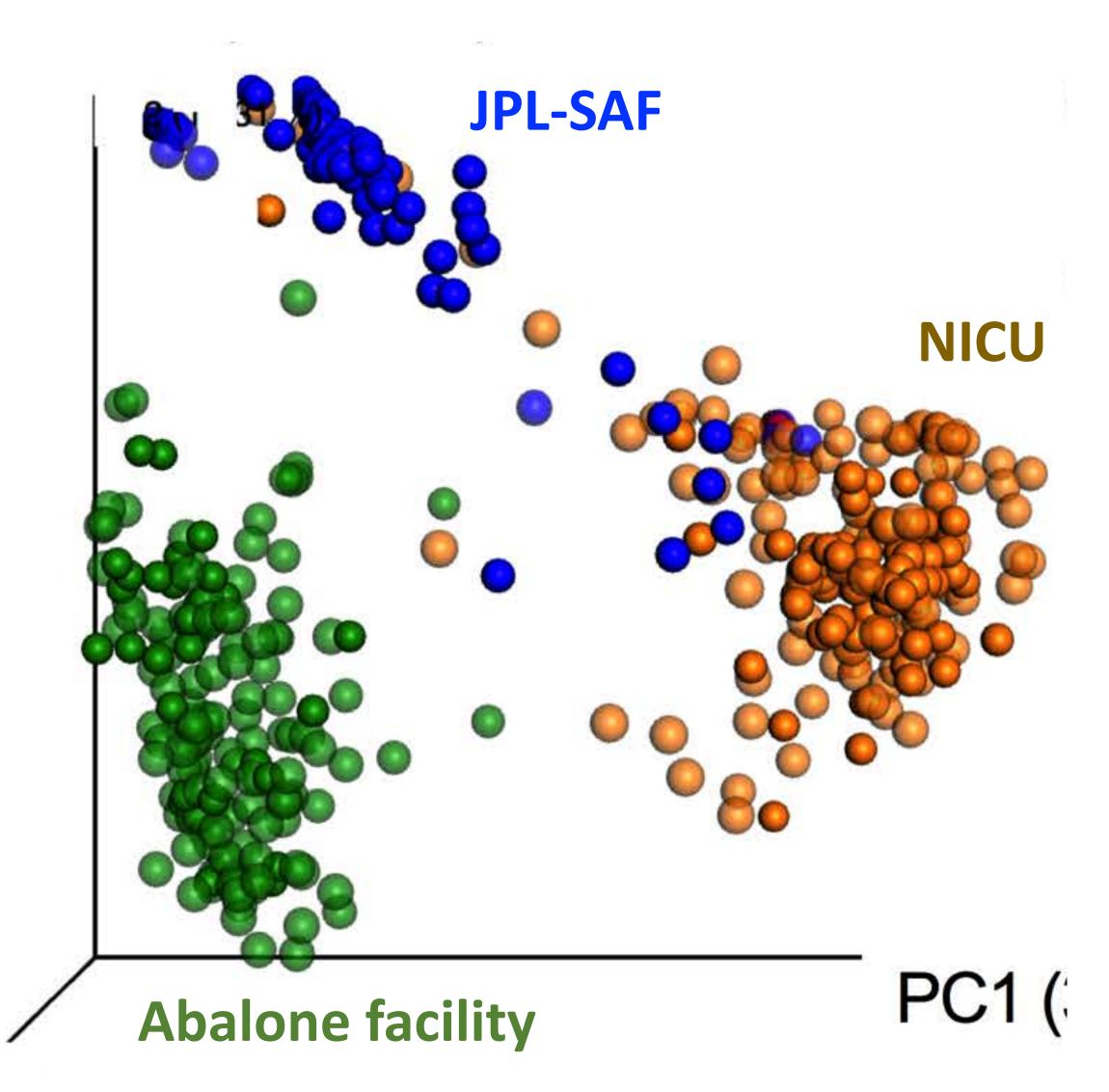


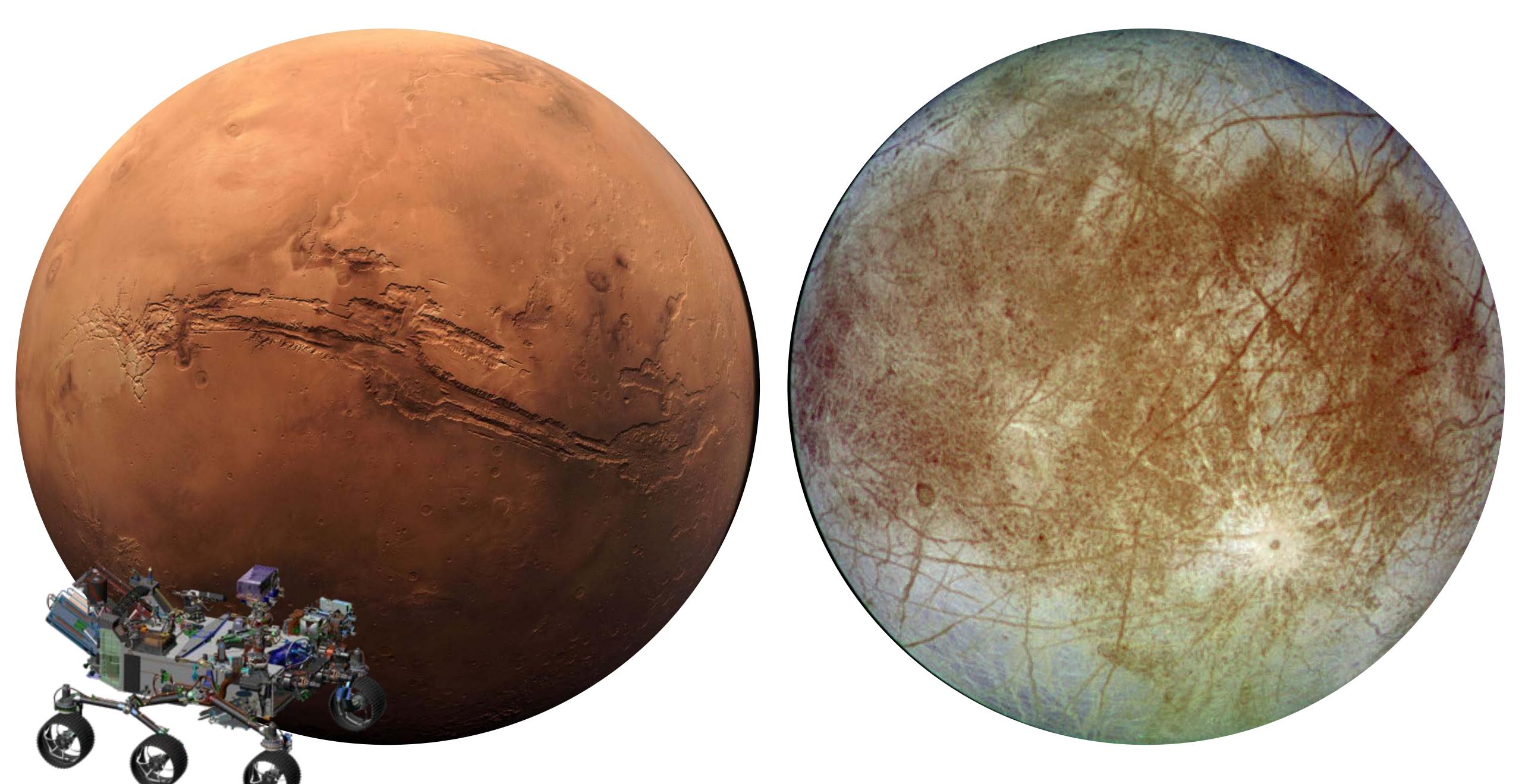


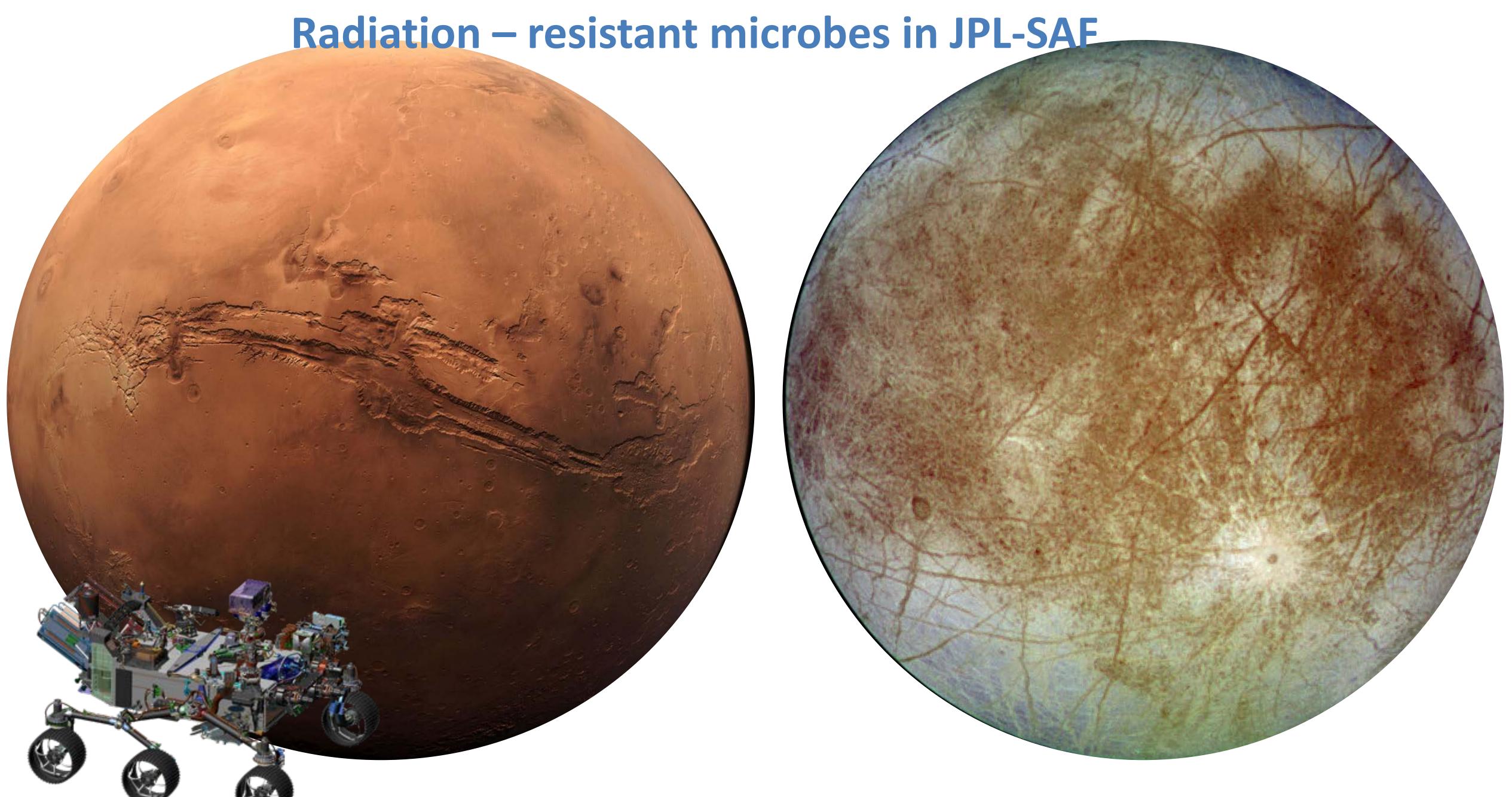
 Among the built environments of neonatal ICU and JPL-SAF cleanrooms, the alpha diversity measured by microbial taxon richness was highest in the NICU than the JPL-SAF cleanroom floors.

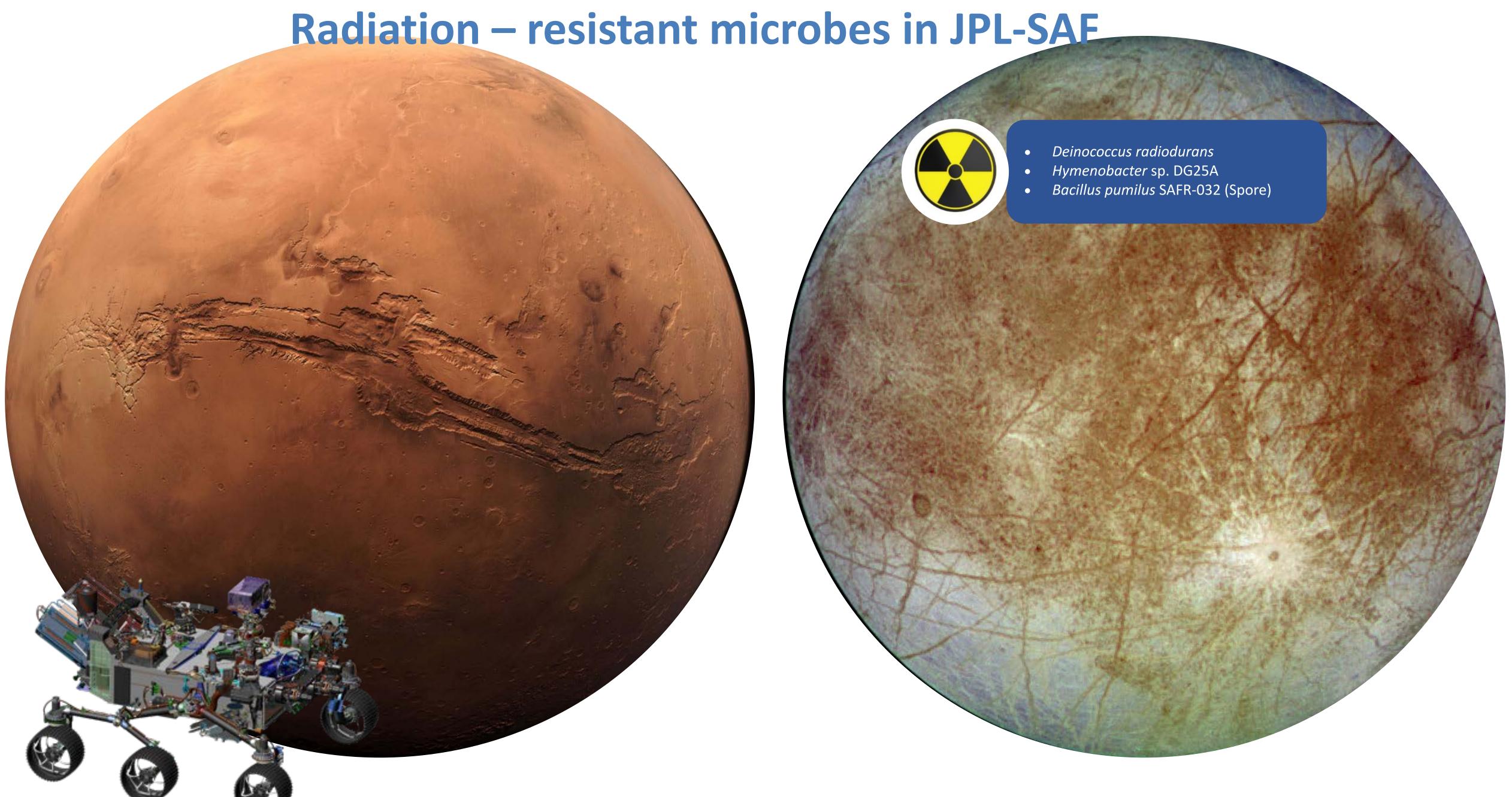


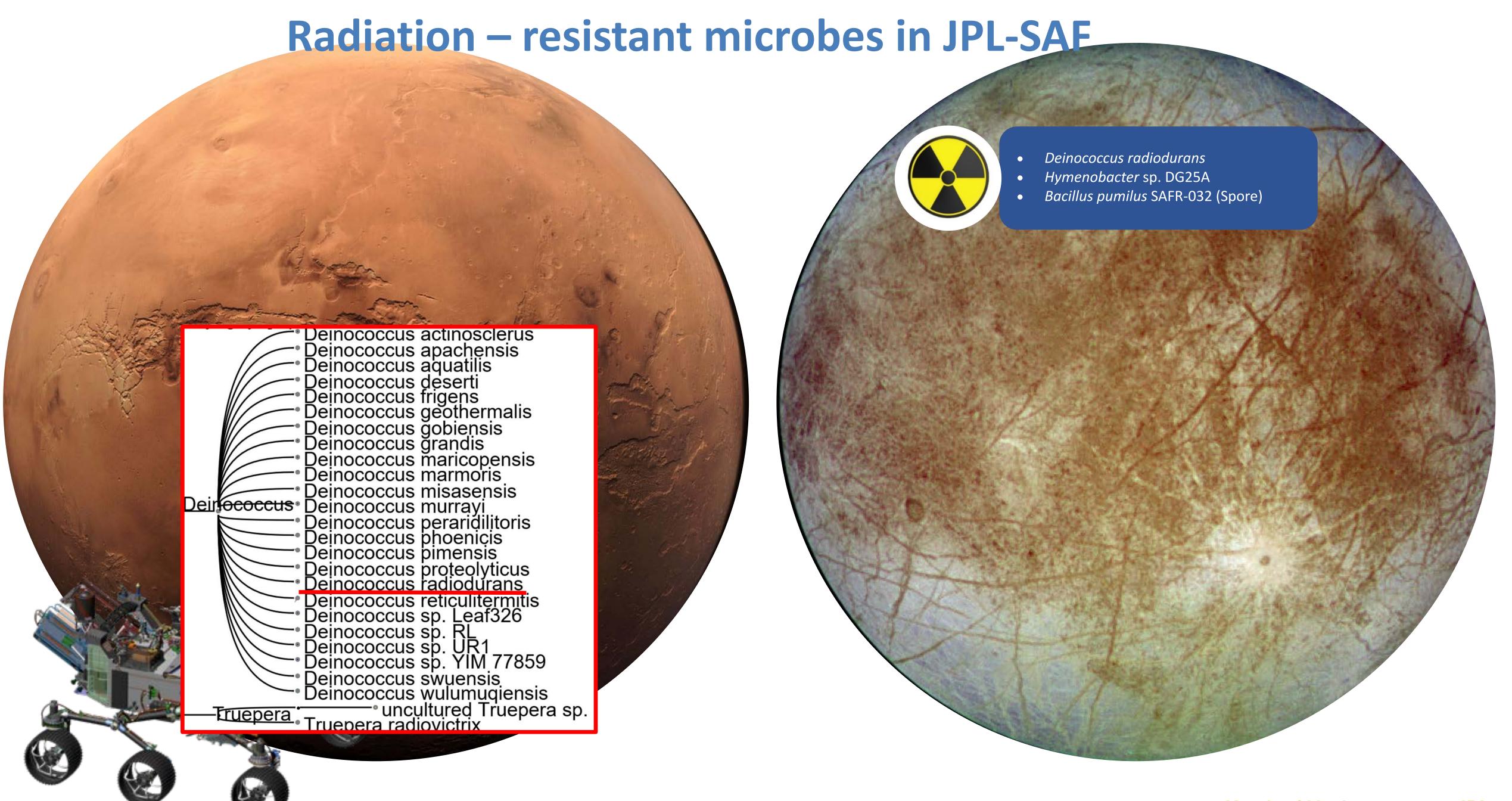
- Among the built environments of neonatal ICU and JPL-SAF cleanrooms, the alpha diversity measured by microbial taxon richness was highest in the NICU than the JPL-SAF cleanroom floors.
- Human exposure was associated with microbial taxon richness, although some of the details differed among the environments.
 - Klebsiella and Staphylococcus with NICU
 - Acinetobacter baumannii, Acinetobacter equi, and Acinetobacter johnsonii in JPL-SAF floors

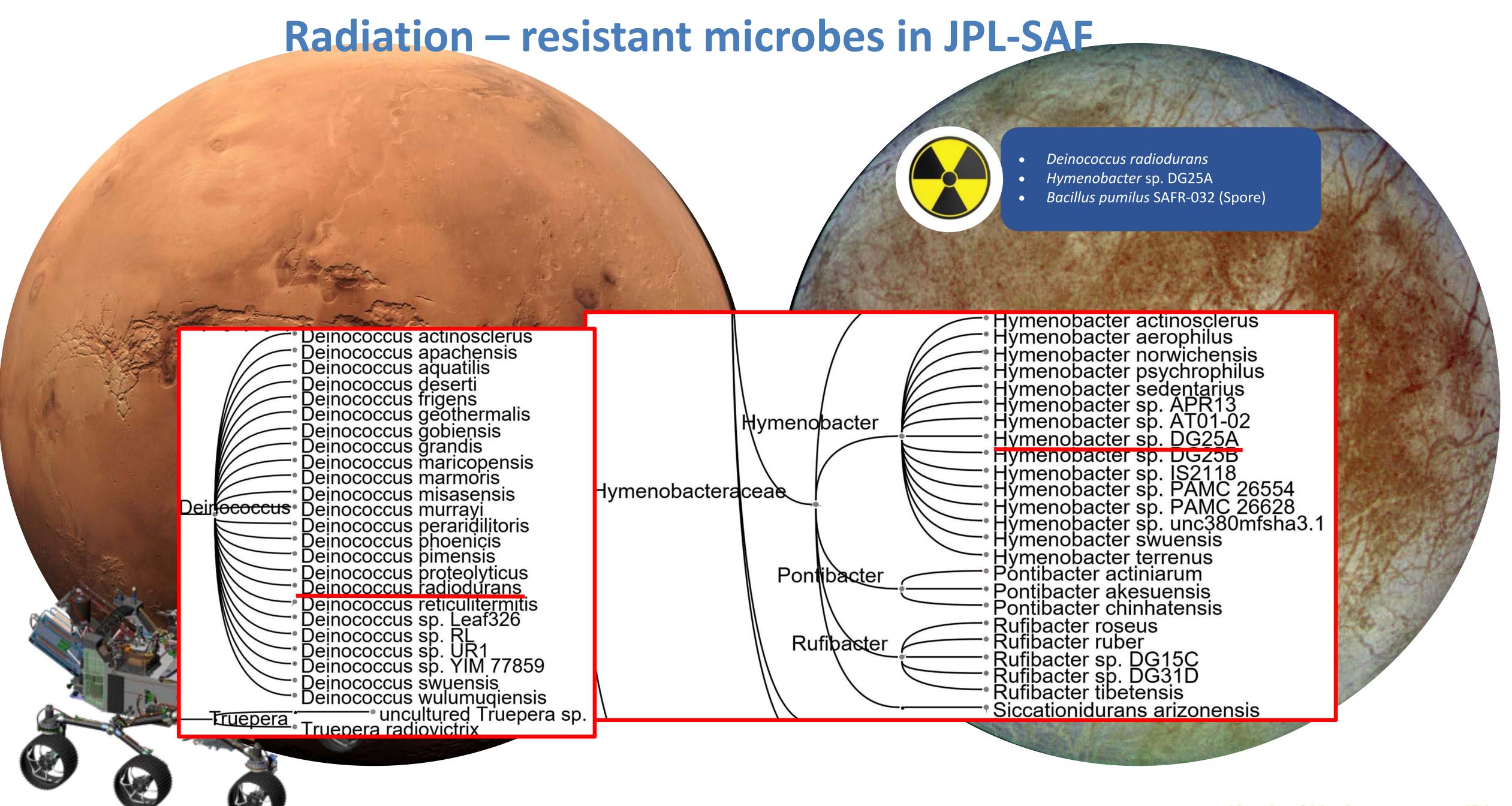




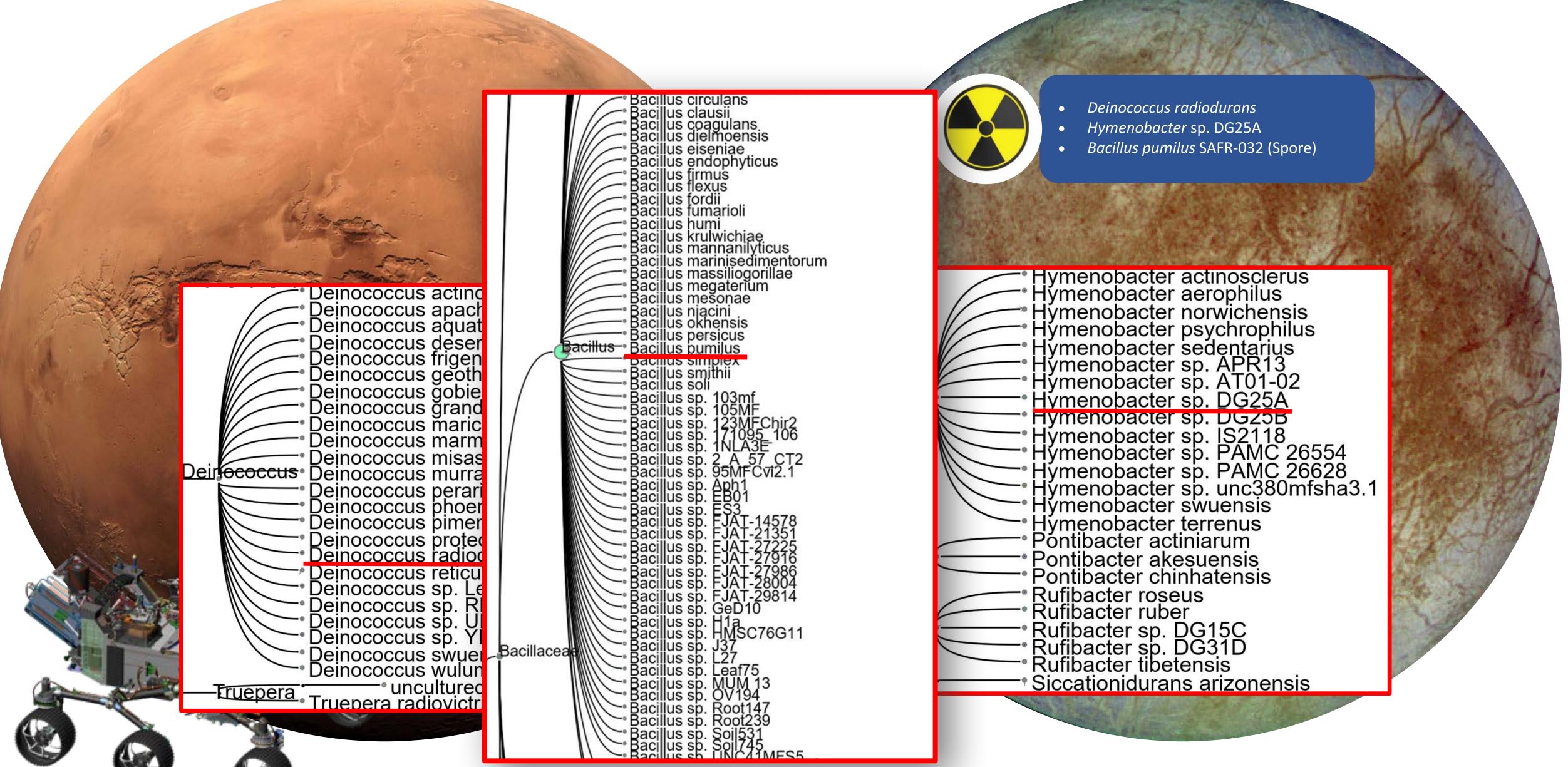








Radiation - resistant microbes in JPL-SAE



2. Creating a healthy built environment on the moon

Sloan Program on Microbiology of the Built Environment

microBEnet is funded by a grant from the Alfred P. Sloan Foundation as part of their program on the Microbiology of the Built Environment.

microbe.net

A conversation with Paula Olsiewski, program director, Alfred P. Sloan Foundation program on the Microbiology of the Built Environment

What is the Foundation's program on the Microbiology of the Built Environment?

The Alfred P. Sloan Foundation is funding an emerging field of scientific inquiry — the microbiology of the built environment. Science knows a lot about microbiology in the natural environment, but we know next to nothing about the complex microbial ecosystems found in the built environment.

The Sloan Foundation is funding basic science to better understand the trillions of life forms invisible to the naked eye in our homes and workplaces, hospitals and schools — any form of built environment — office buildings, airplanes, subways systems, you name it.

Why study the built environment?

People spend more than 90% of their time indoors, where we breathe in and come in contact with trillions of microorganisms – life forms invisible to the naked eye. These tiny creatures live inside and have probably been there from the beginning of construction. But they also come in through windows and mechanical systems such as heating and air-conditioning systems, and are brought inside by humans and other creatures, which are constantly shedding microbes.

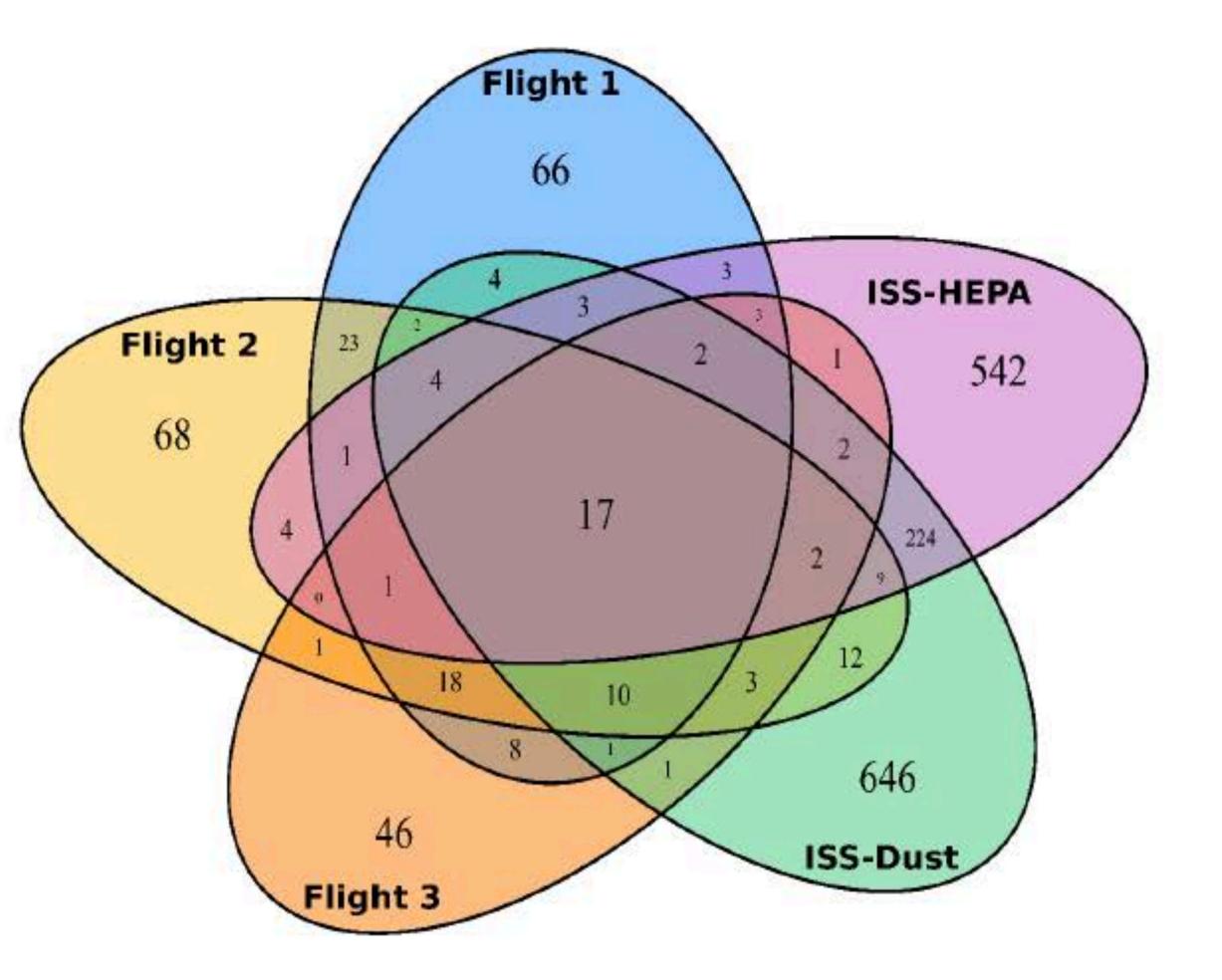
Our homes and workplaces, hospitals and schools are quite literally complex ecosystems filled with a variety of microbes. Given the amount of time we spend indoors it's important to understand what is living in these environments, how these microorganisms interact, what the potential implications are for human health, both positive and negative. We may also learn things that could influence building construction practices and inform other industrial processes.

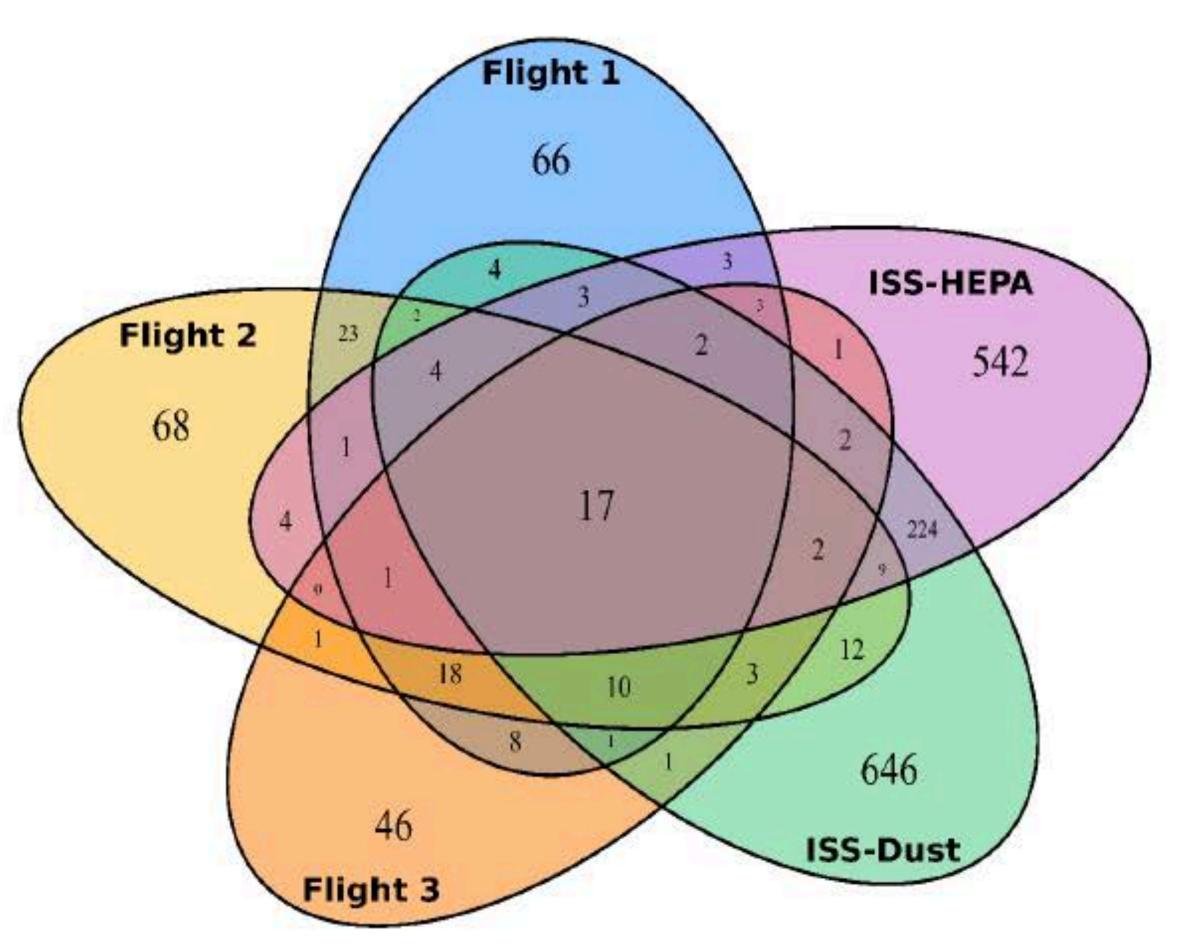
The National Academies of SCIENCES • ENGINEERING • MEDICINE

CONSENSUS STUDY REPORT

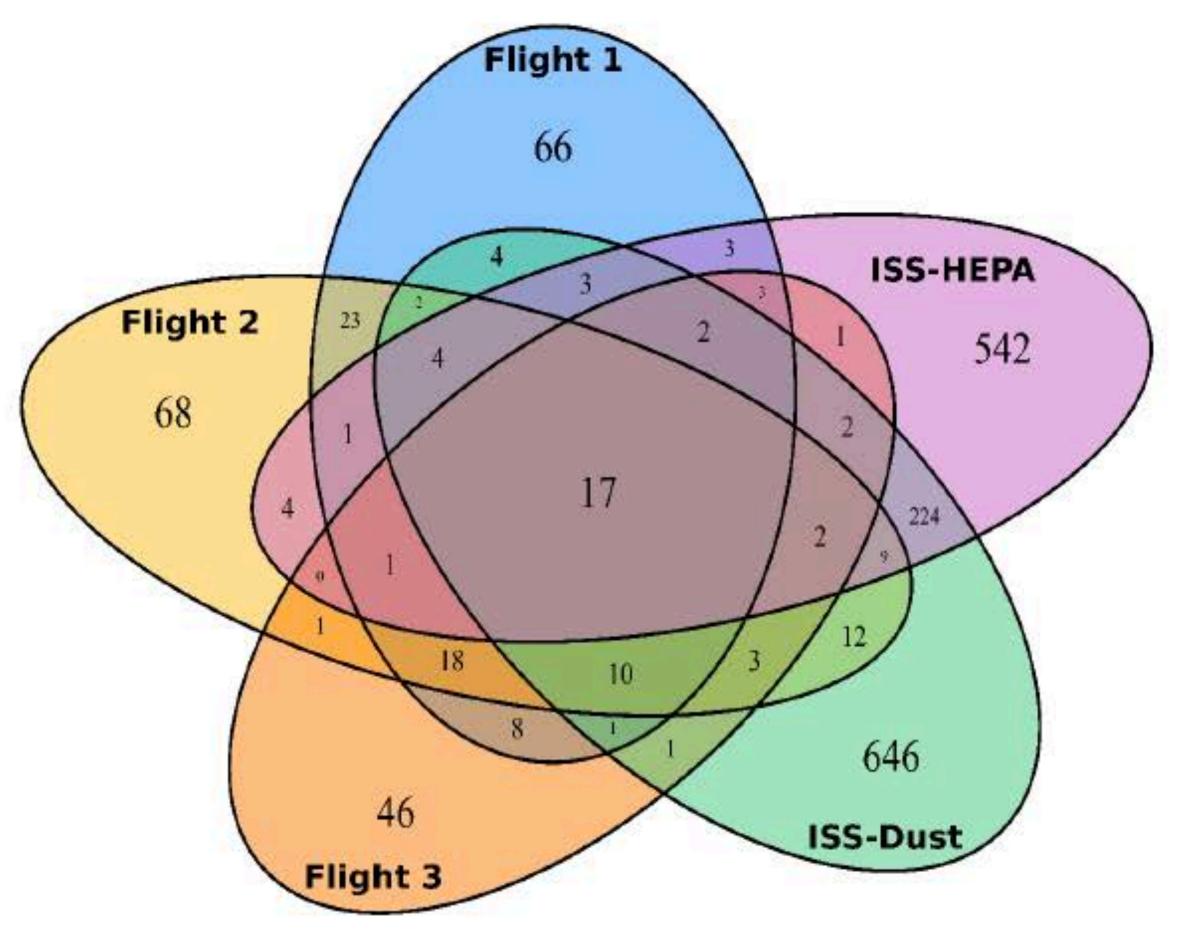
Microbiomes of the Built Environment

HUMAN HEALTH, AND BUILDINGS

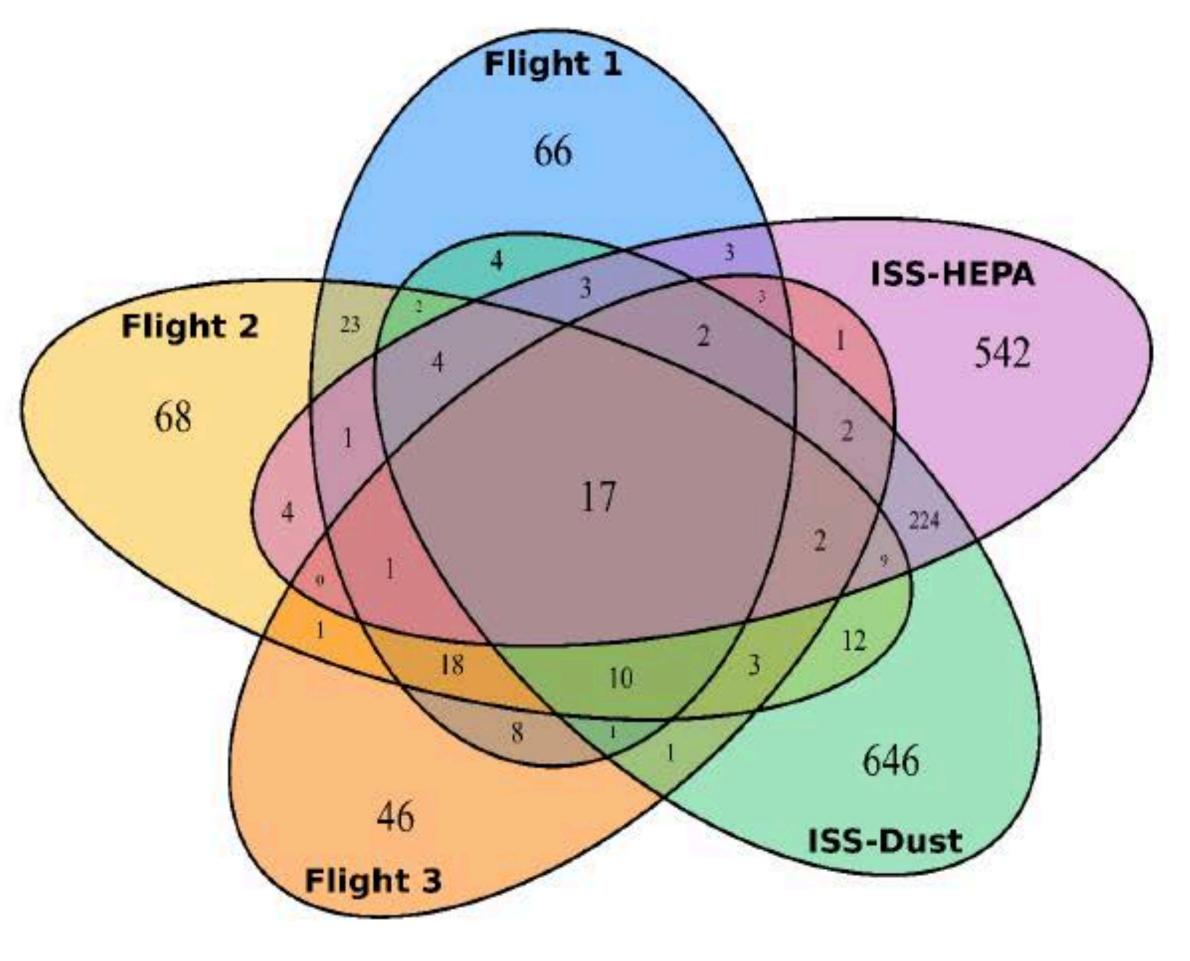




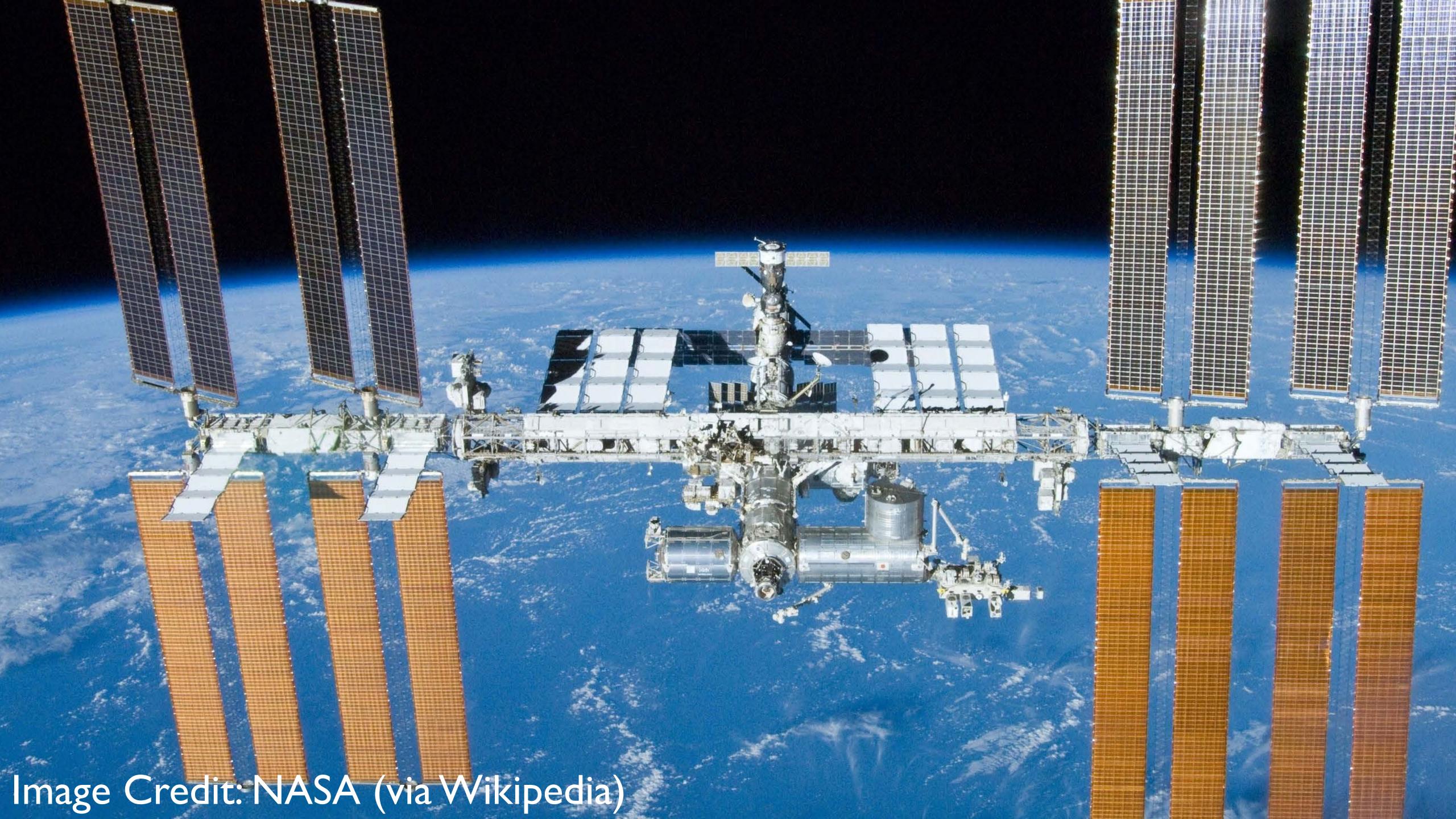
• The ISS microbiome is unique to a particular subsystem but also some microbial species are common (core microbiome; 17 species).

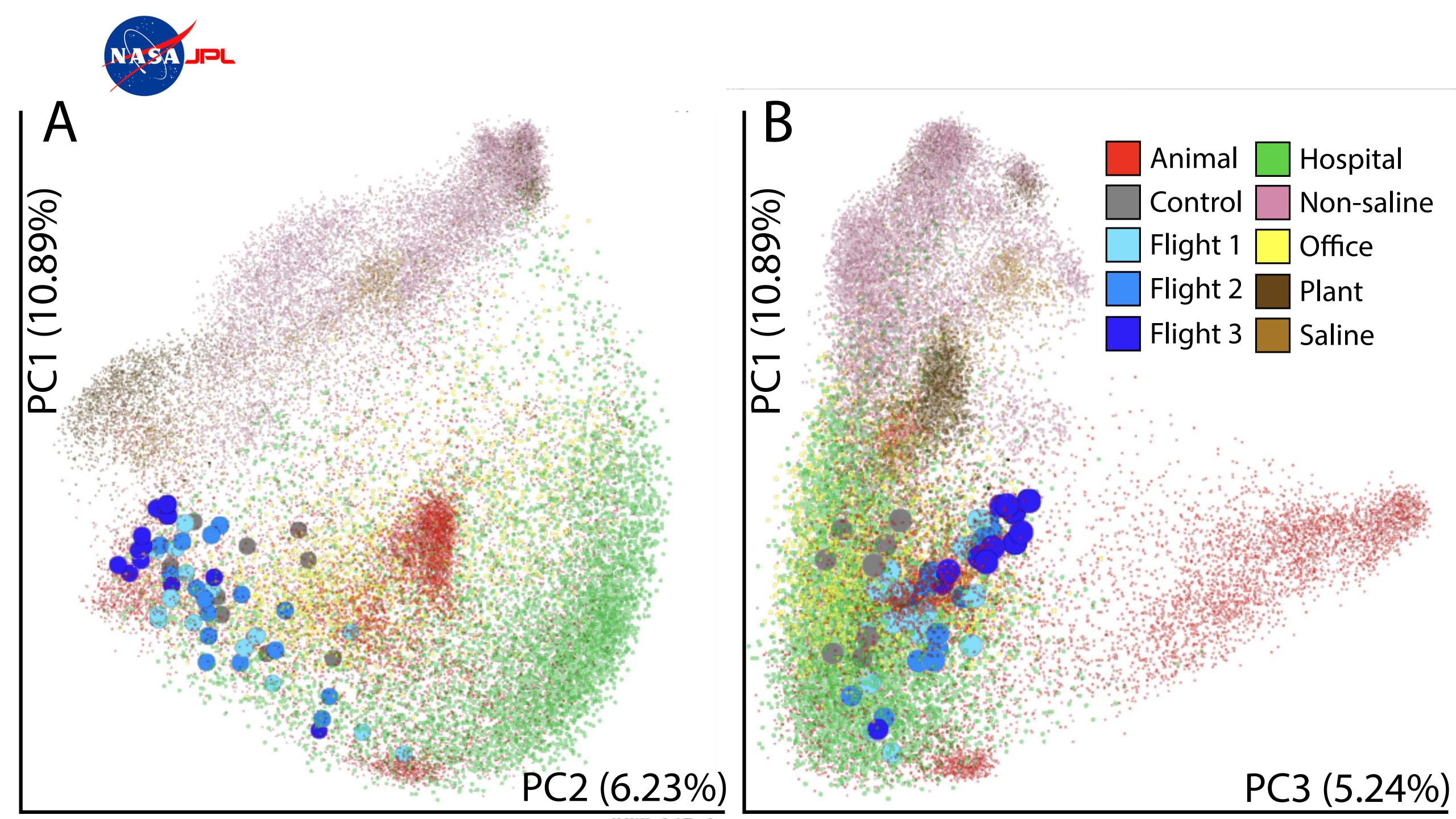


- The ISS microbiome is unique to a particular subsystem but also some microbial species are common (core microbiome; 17 species).
- Dominant ISS core microbiome:
 - Fungi (P. brasilianum, P. chrysogenum, P. digitatum, P. expansum, P. freii, P. griseofulvum, P. roqueforti, P. rubens, A. calidoustus, and A. niger).
 - Bacteria (*E. cloacae, E. coli, P. ananatis, S. enterica, S. aureus*, and *S. epidermidis*).



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 - Bacteria (*E. cloacae, E. coli, P. ananatis, S. enterica, S. aureus*, and *S. epidermidis*).
- Compare with other microbiome studies:
 - The ISS microbiome represents a "minimal core" human model in which all the human subjects shared few microbial species, large overlaps were found in subsets but a very little was common between all the sets.





Growing up on an Amish farm protects children against asthma by reprogramming immune cells

August 3, 2016

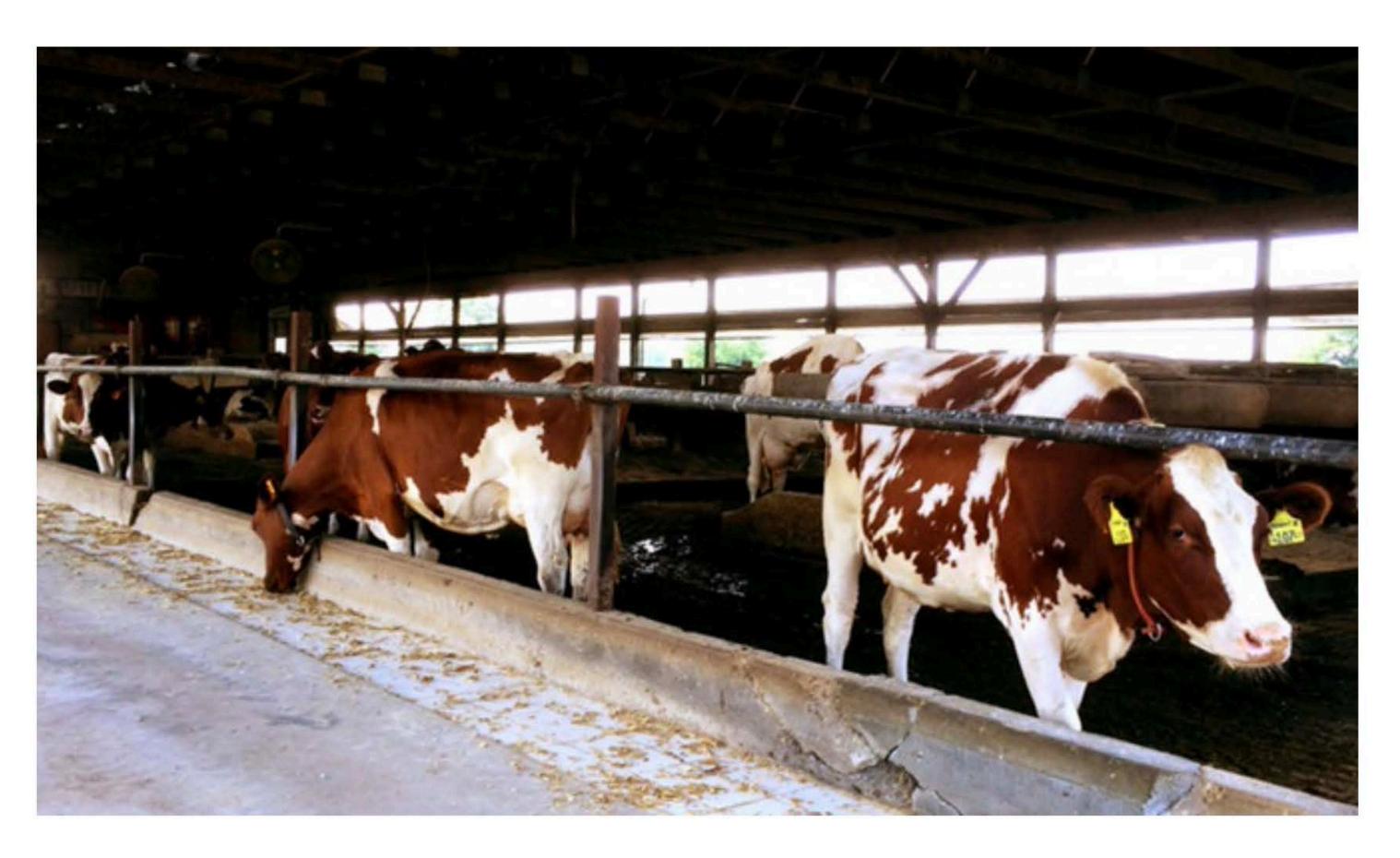
Written By John Easton

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By probing the differences between two farming communities -- the Amish of Indiana and the Hutterites of South Dakota -- an interdisciplinary team of researchers found that specific aspects of the Amish environment are associated with changes to immune cells that appear to protect children from developing asthma.

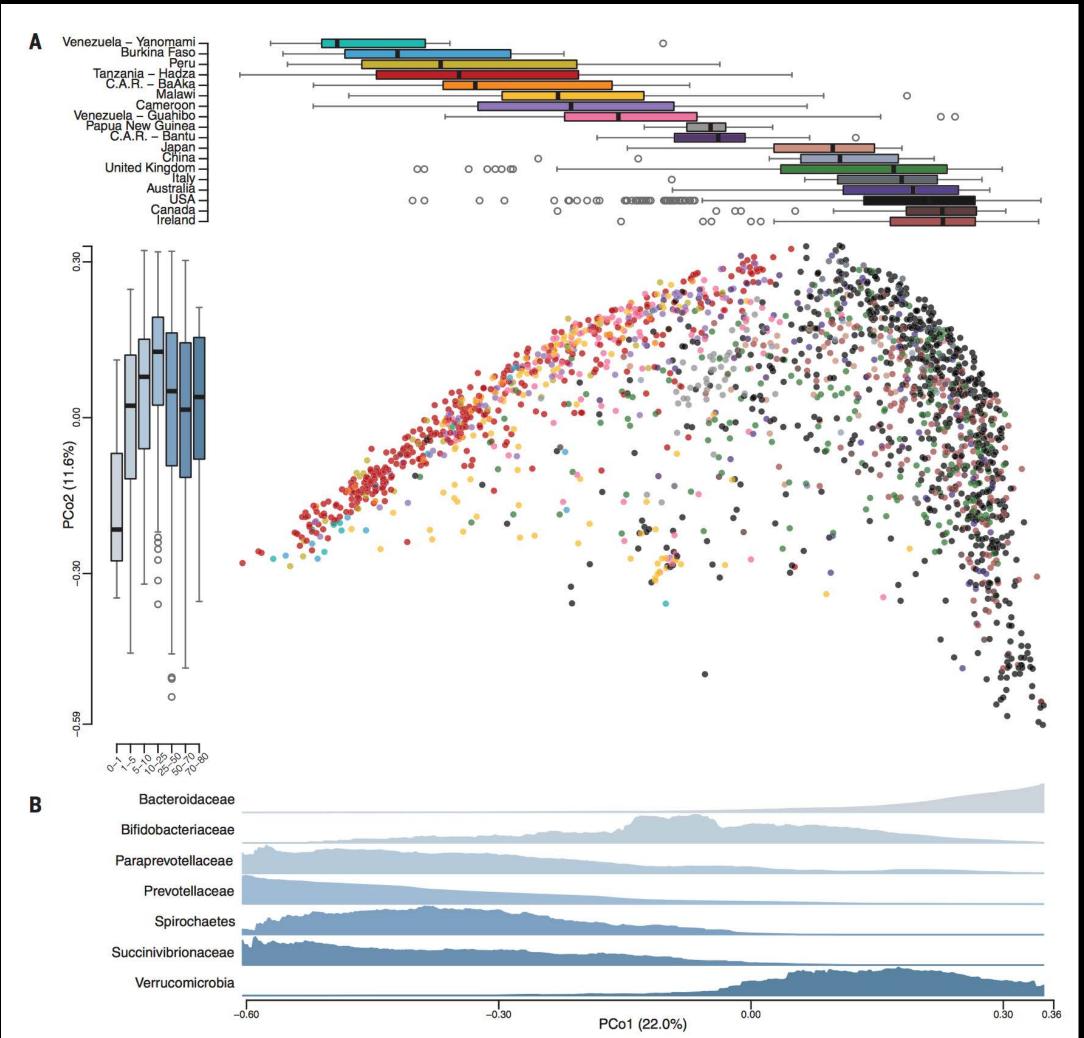
3. Precision medicine and nutrition for lunar explorers





Seasonal cycling in the gut microbiome of the Hadza hunter-gatherers of Tanzania

Samuel A. Smits, ^{1*} Jeff Leach, ^{2,3*} Erica D. Sonnenburg, ¹
Carlos G. Gonzalez, ⁴ Joshua S. Lichtman, ⁴ Gregor Reid, ⁵ Rob Knight, ⁶
Alphaxard Manjurano, ⁷ John Changalucha, ⁷ Joshua E. Elias, ⁴
Maria Gloria Dominguez-Bello, ⁸ Justin L. Sonnenburg ¹†



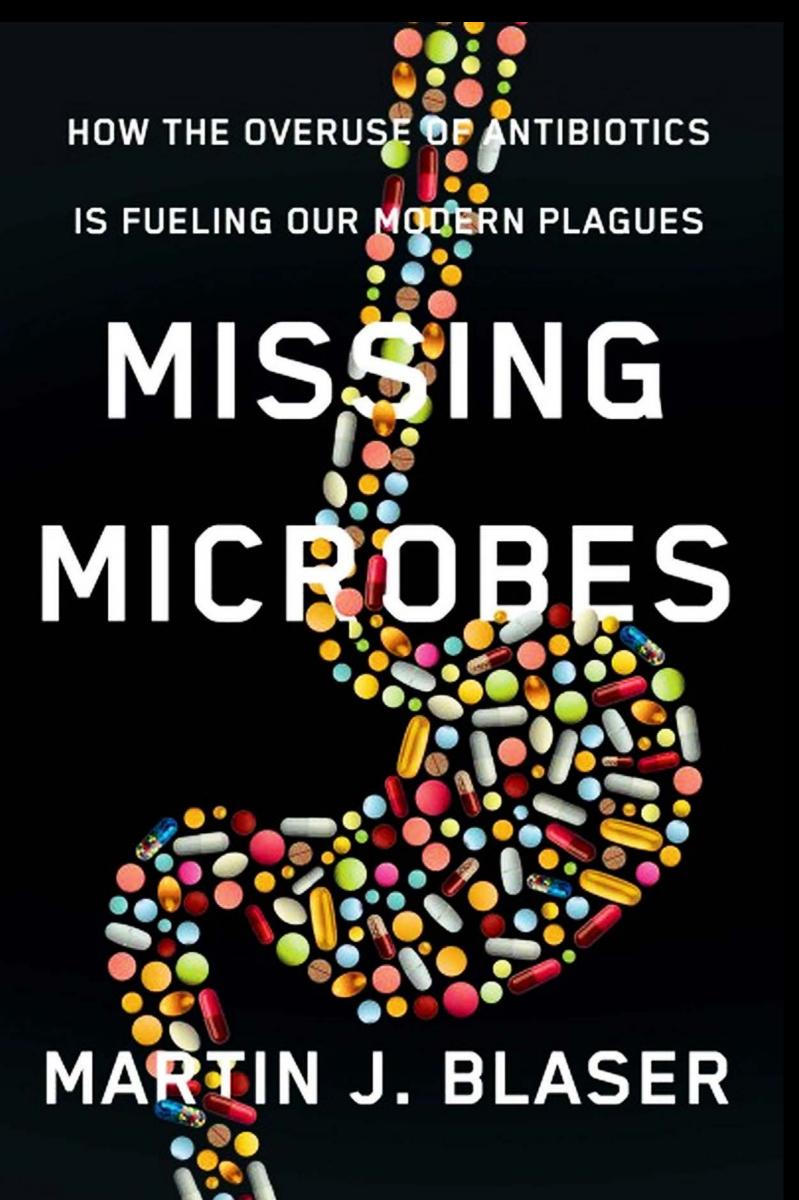




A global microbial biodiversity crisis?

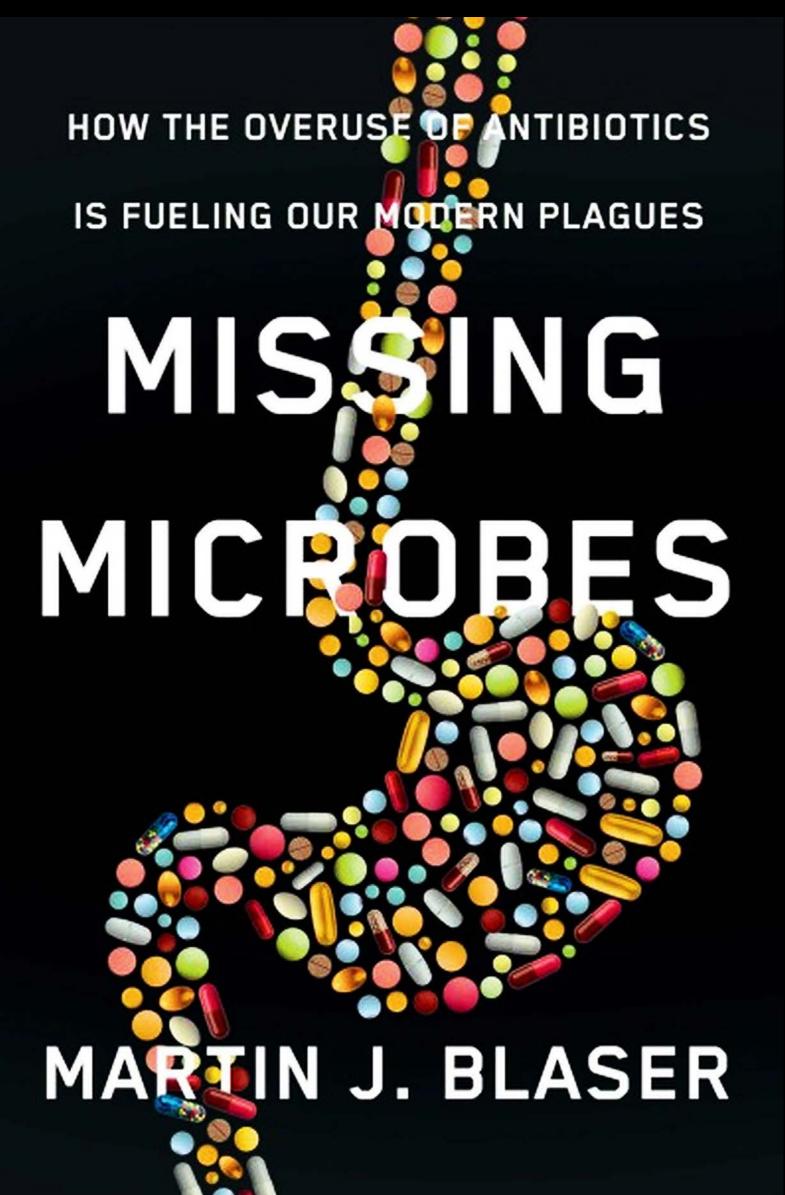


A global microbial biodiversity crisis?





A global microbial biodiversity crisis?



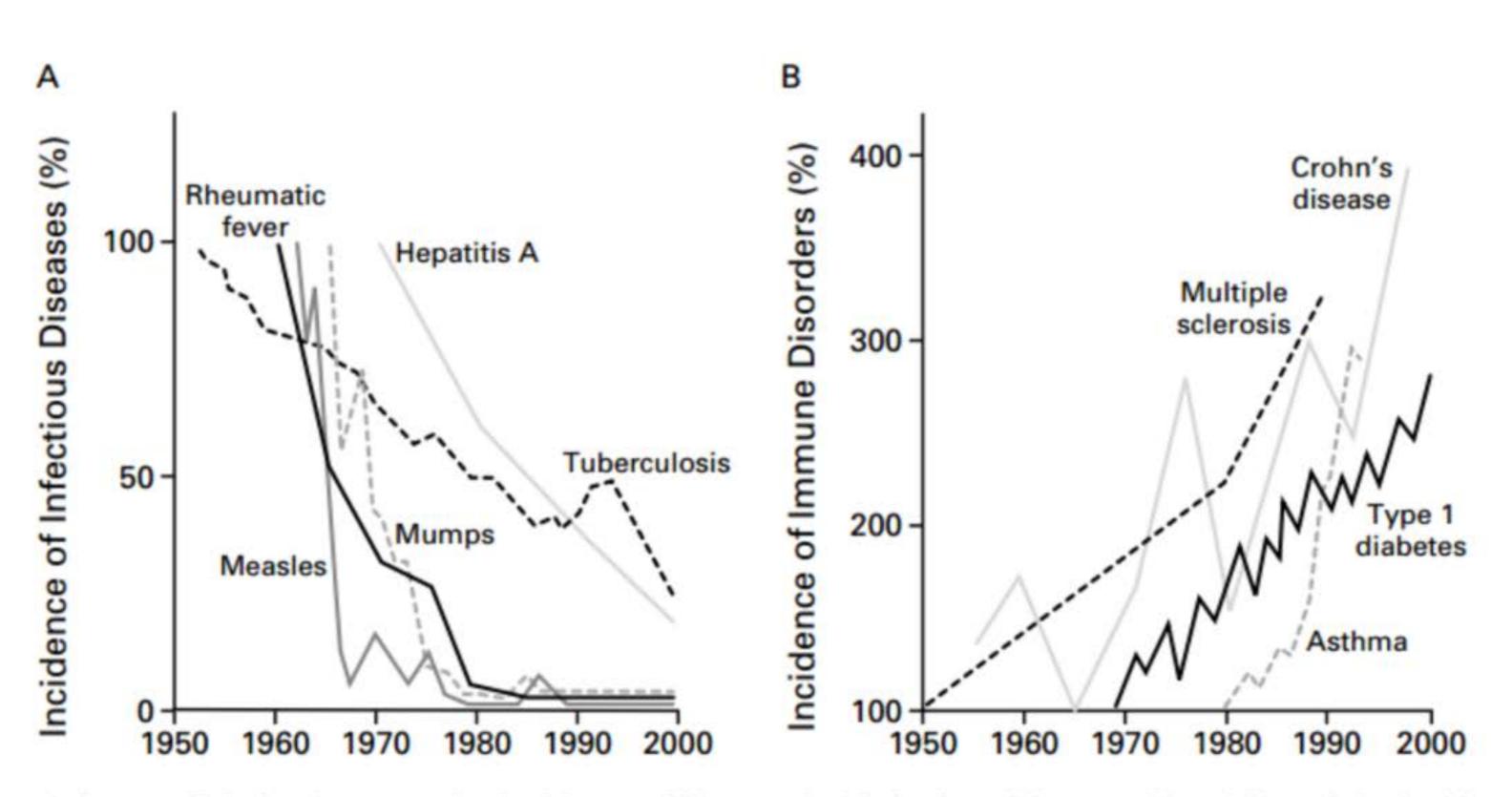


Figure 1. Inverse Relation between the Incidence of Prototypical Infectious Diseases (Panel A) and the Incidence of Immune Disorders (Panel B) from 1950 to 2000.

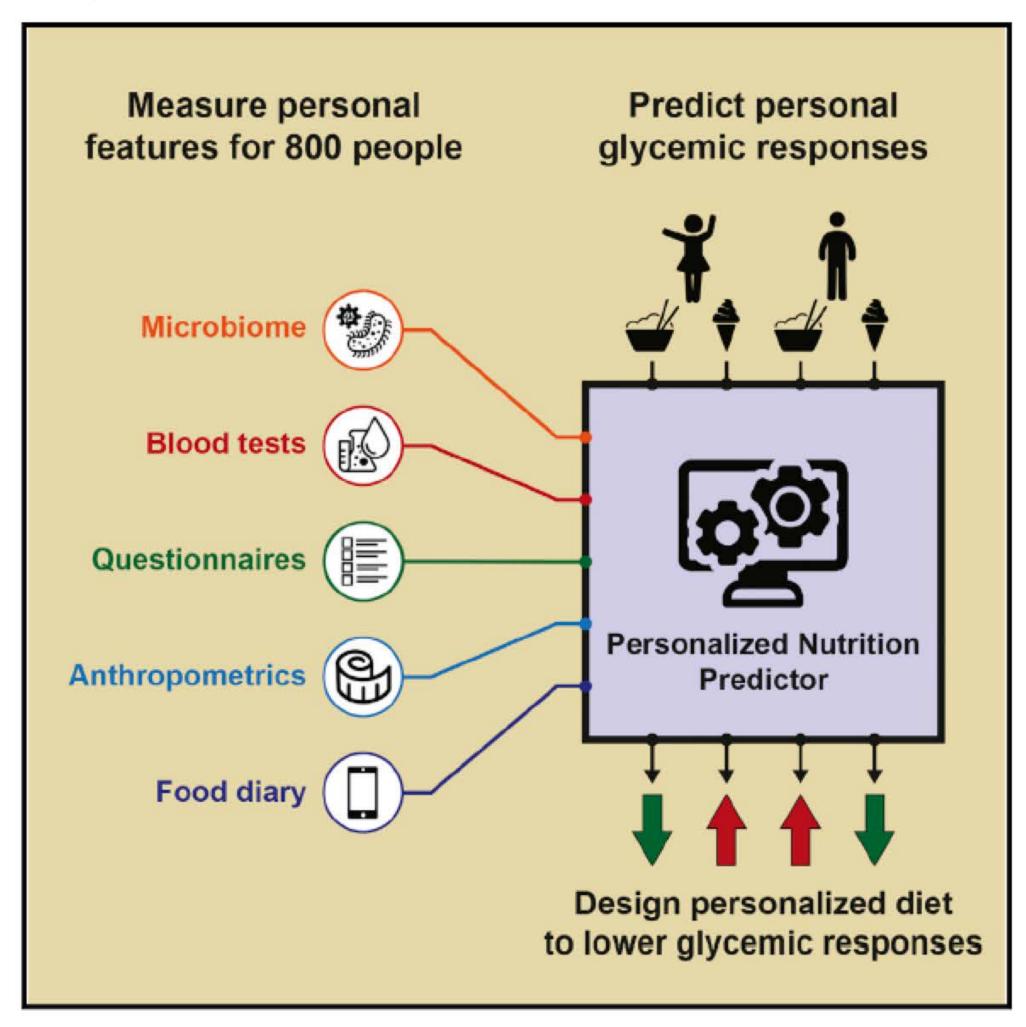
In Panel A, data concerning infectious diseases are derived from reports of the Centers for Disease Control and Prevention, except for the data on hepatitis A, which are derived from Joussemet et al.¹² In Panel B, data on immune disorders are derived from Swarbrick et al.,¹⁰ Dubois et al.,¹³ Tuomilehto et al.,¹⁴ and Pugliatti et al.¹⁵

Bach (2002) N Engl J Med, Vol. 347, 911-920



Personalized Nutrition by Prediction of Glycemic Responses

Graphical Abstract



Authors

David Zeevi, Tal Korem, Niv Zmora, ..., Zamir Halpern, Eran Elinav, Eran Segal

Correspondence

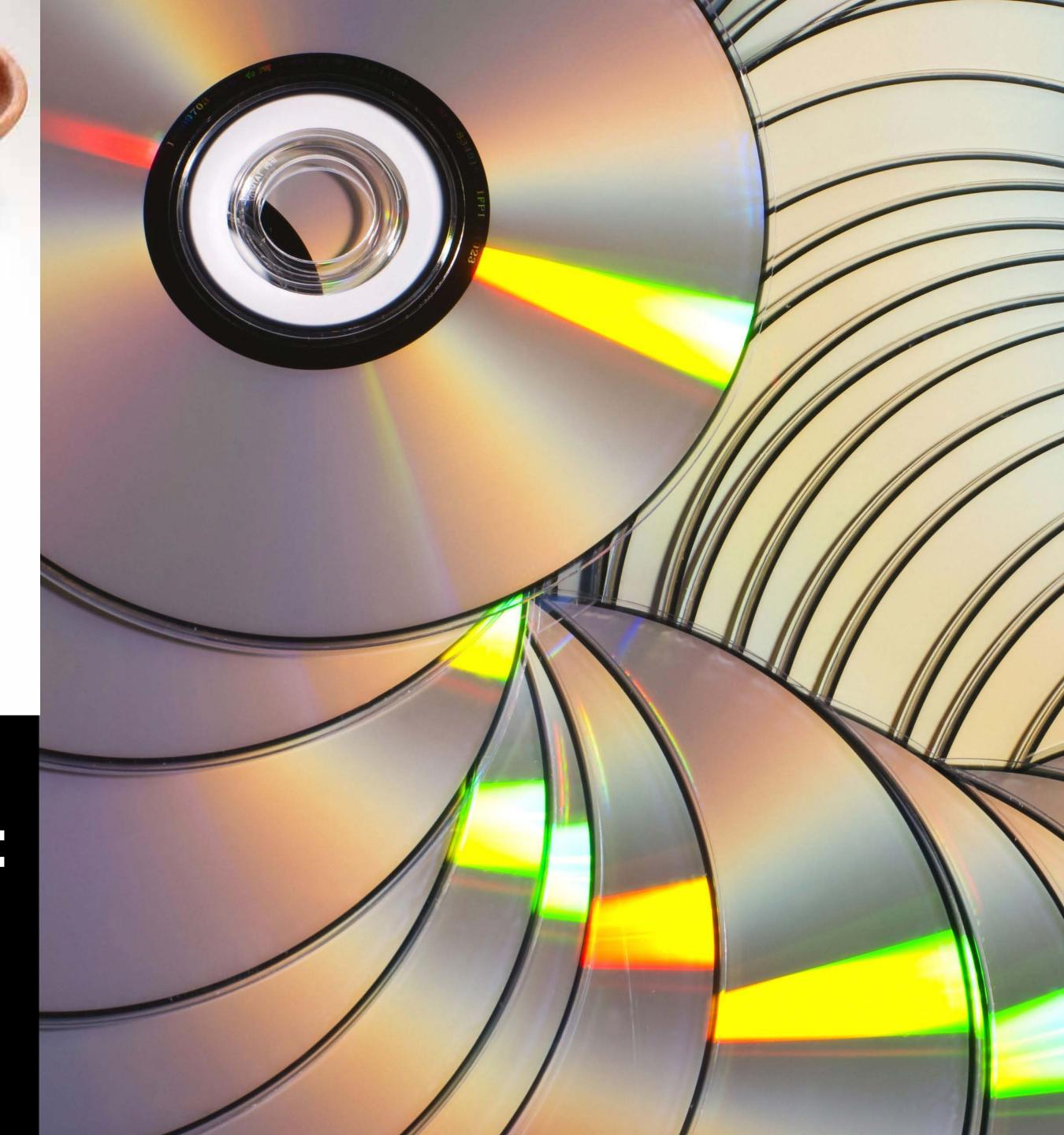
eran.elinav@weizmann.ac.il (E.E.), eran.segal@weizmann.ac.il (E.S.)

In Brief

People eating identical meals present high variability in post-meal blood glucose response. Personalized diets created with the help of an accurate predictor of blood glucose response that integrates parameters such as dietary habits, physical activity, and gut microbiota may successfully lower post-meal blood glucose and its long-term metabolic consequences.



l teaspoon stool =
l ton of DVDs



Questions?

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