

GERMS IN SPACE

A COMPARATIVE ANALYSIS OF GERM GROWTH ACTIVITY ON EARTH AND IN SPACE

By Christopher Future Astronaut – Space Explorer Science Project Assignment 2021

HELLO & WELCOME







INTRODUCTION

On planet Earth, we are accustomed to having gravity which is a force that keeps everything from floating around.

In space, there is very little gravity. On the International Space Station we call it microgravity.

Gravity affects things very differently. It is important to understand how it affects germs in space.





PURPOSE

The purpose of my research is to understand how germs grow in low gravity environments.

I also intend to understand the different types of germs present on the ISS compared to the germs that I will collect in my experiment here on Earth.

I hope to learn about the challenges and concerns with germ behavior on Earth so that I can better understand its behavior in space. This is very important as humanity prepares to establish a permanent presence in space and to explore beyond. We need to understand how humans will grow food in space and how germs will impact human health and performance in low gravity environments.

Through my research, I hope to develop solutions that will benefit life on Earth and in space.





HYPOTHESIS

PROBLEM STATEMENT

How does microgravity environments affect the growth of germs (bacteria, fungi, etc) in space?

Like with most biological forms of life, a few common parameters that are ideal for bacterial growth are gravity, moisture, light, temperature and nutrients.

In space (on the ISS), the only thing that will be different is the low gravity environment.

HYPOTHESIS

I hypothesize that the germs in space will not grow as much as germs found on earth. Their growth patterns, rates and bacterial colonies will differ.

I also hypothesize that different types of germs may be found on the ISS compared to here on Earth. Some of the germs found in space may thrive on the ISS because the low gravity environment is what they prefer.



MATERIALS

- 13 Petri Dishes
- 15g Nutrient Powder (Agar)
- 1 Roll of Transparent Tape
- 6 Disposable Aprons
- 12 Swabs
- 20 Pairs of Disposable Gloves
- 13 Zip Top Bags
- Exploration Guide
- 1 Gallon of Distilled Water
- 1 Laboratory Beaker

- **1 Laboratory Beaker**
- 1 Microwave Safe Container
- 1 Stirring Utensil
- Silicon Pot Holder
- Thermometer
- Magnifying Glass
- Microscope
- 1 Black Marker











PROCEDURES

- It was important to practice aseptic techniques to ensure that my experiment was not contaminated.
- I followed the sterilization procedures for my equipment provided with a previous space-related ISS experiment.
- I iterated the procedures as needed.



PROCEDURES

- I cleaned my 13 Petri dishes and corresponding covers with a 10% bleach solution for around 5 minutes.
- I also cleaned the beaker using this method.
- I rinsed my petri dishes and caps 5 times in distilled water.
- With supervision, I prepared my agar solution with a hot plate, distilled water and a beaker.
- I then placed 10ml of agar growth medium in each test tube after it cooled off.
- I collected my germ samples from around my home with the cotton swabs and smeared on agar.
- I labeled each sample with the tape and covered the petri dishes with lids.
- I recorded my observational data inside my experiment log for 15 days.
- I monitored the external conditions (temperature, light, moisture & nutrients).



EXPERIMENT

For 15 days I observed and recorded data on the germ growth rate, behavior and characteristics. Additionally, I plotted the data within Excel to analyze it.







EXPERIMENT PREPARATION

BEAKER	AGAR AMOUNT	WATER AMOUNT	NUMBER OF PETRI DISHES FOR PLATING	HEATING TIME
1500ml	15g	600ml	13	90-120 seconds

- I used an agar called Standard Method Agar provided by DreamUp to make my agar.
- Using safety equipment and with supervision, I mixed and heated my distilled water and agar making sure that the agar was dissolved completely.
- The media appeared cloudy initially but after boiling it and occasionally stirring it, it cleared up.





EXPERIMENT PREPARATION

BEAKER	AGAR AMOUNT	WATER AMOUNT	NUMBER OF PETRI DISHES FOR PLATING	HEATING TIME
1500ml	15g	600ml	13	90-120 seconds

- After I prepared my agar solution using distilled water, I let it cool for 20 minutes then I checked the agar temperature.
- At 40C, I distributed the agar evenly in my 13 petri dishes.
- I immediately placed the lid on the petri dishes after filling them.
- I allowed the prepared petri dishes to cool more so that the agar media would solidify.
- I placed them upside down so that condensation would not form on the media.





EXPERIMENT PROCEDURES



- I collected my samples from various locations using a sterile cotton swab.
- I streaked my samples on the agar media.
- I labeled and sealed my petri dishes with clear tape to secure the lids.
- I placed each dish in a sealed clear transparent bag and kept them upside down.
- Samples were kept at room temperature around 73 degrees F.



EXPERIMENT SETUP

EXPERIMENT SAMPLES



KITCHEN SINK

13



EXPERIMENT SETUP





DATA COLLECTION

During this experiment I collected a lot of data for current and future analysis.





DATA COLLECTION METHODOLOGY

I collected both **QUANTITATIVE & QUALITATIVE** data.

- **A OBSERVATION OF MORPHOLOGY & OTHER FACTORS**
- **B** GENERAL MORPHOLOGY ANALYSIS
- **C COMPARATIVE ANALYSIS EARTH vs SPACE**
- **D** GROWTH RATE ANALYSIS



DATA COLLECTION /

OBSERVATION LOG (DAYS 1-15)





DATA COLLECTION B

COLONY MORPHOLOGY GENERAL OBSERVATIONS FOR EACH SAMPLE

SAMPLE 13

COLONY	MORPHOLC	GY		
FORM		ELEVATION		MARGIN
PUNTIFORM		FLAT		ENTIRE
CIRCULAR		RAISED	Х	ULDINATE
FILAMENTOUS		CONVEX	Х	FILAMENTOUS
IRREGULAR	X	PULVINATE		LOBATE
RHYZIOD		UMBINATE		EROSE
SPINDLE				CURLED

I ALSO RECORDED DETAILS ABOUT MY OBSERVATIONS ON THIS CHART ALSO FOR EACH SAMPLE WITH A FOCUS ON THE COLONY MORPHOLOGY

MORPHOLOGY CLASSIFICATION CHART





DATA COLLECTION

COLONY MORPHOLOGY GENERAL OBSERVATIONS FOR EACH SAMPLE

COLONY MC	ORPHOLOGY	SAMP	LE 1	COLONY MORPHOLOGY SAMPLE 2			COLONY MORPHOLOGY				SAMPLE 3			
FORM	ELEVATION	MARGIN		FORM	ELEVATIO	N	MARGIN		FORM		ELEVATION		MARGIN	
PUNTIFORM X	FLAT	× ENTIRE	x	PUNTIFORM	FLAT		ENTIRE	Х	PUNTIFORM		FLAT		ENTIRE	Х
CIRCULAR x	RAISED	ULDINATE		CIRCULAR >	X RAISED	Х	ULDINATE		CIRCULAR	х	RAISED	x	ULDINATE	
FILAMENTOUS	CONVEX	FILAMENTOL	s	FILAMENTOUS	CONVEX		FILAMENTOUS		FILAMENTOUS		CONVEX		FILAMENTOUS	
IRREGULAR	PULVINATE	LOBATE		IRREGULAR	PULVINAT	E	LOBATE		IRREGULAR		PULVINATE		LOBATE	
RHYZIOD	UMBINATE	EROSE		RHYZIOD	UMBINAT	E	EROSE		RHYZIOD		UMBINATE		EROSE	
SPINDLE		CURLED		SPINDLE			CURLED		SPINDLE				CURLED	
COLONY N	MORPHOLOGY	SAM	PLE 4	COLONY MORPHOLOGY SAMPLE 5			PLE 5	COLONY MORPHOLOGY S				SAM	PLE 6	
FORM	ELEVATIO	MARGIN		FORM	ELEVATIO	ON	MARGIN		FORM		ELEVATION		MARGIN	
PUNTIFORM	FLAT	X ENTIRE		PUNTIFORM	FLAT	х	ENTIRE		PUNTIFORM	x	FLAT		ENTIRE	
CIRCULAR	RAISED	ULDINATE	х	CIRCULAR	RAISED		ULDINATE		CIRCULAR	х	RAISED	xx	ULDINATE	Х
FILAMENTOUS	CONVEX	FILAMENTO	JS	FILAMENTOUS	CONVEX		FILAMENTOUS	5	FILAMENTOUS	6	CONVEX		FILAMENTOUS	
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RHYZIOD		EROSE		RHYZIOD	UMBINA	TE	EROSE	Х	RHYZIOD		UMBINATE		EROSE	Х
SPINDLE		CURLED		SPINDLE			CURLED		SPINDLE				CURLED	
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ANALYSIS A & B



I USED THIS CHART AS A GUIDELINE TO CLASSIFY MY FINDINGS.

Colony 4 Colony 3 Colony 1 Colony 2 Colony 7 Colony 5 Colony 6 Colony 8 Colony 9 Colony 10 Colony 11 Colony 13 Colony 12

ABOVE IS AN EXAMPLE OF DIFFERENT TYPES OF COLONIES.

DATA COLLECTION C

COMPARATIVE DATA FOR MICROBIAL BACTERIA GROWTH ON EARTH AND IN SPACE

GERMS IN SPACE	COMPARISON CHART	
		DATE
	My Microbial Growth on Earth	Microbial Growth in Space
What do you first notice about the microbial growth?	Early on, I noticed that the growth seemed slow .	The growth occurs very fast in space.
How large is the microbial growth in general?	It depended on the bacteria present in the petri dishes. Several of them rapidly covered	In space, the bacteria forms biofilms which is a protective mode of growth. This allows them to survive in hostile environments and to colonize in new areas.
What colors are the microbes?	The colors vary	Colors vry.
Describe the shape of the colonies	The colony growth patterns varied based on the samples collected.	The colonies stick together forming biofilms.
What do you notice about the arrangement of the microbes?	Many of the microbes grew in circular patterns then they spreaded out.	The colonies stick together forming biofilms and groups.
How does your microbial growth compare with the microbial growth in space?	My microbes grew slower than the microbes in space.	According to research conducted on the ISS, 8 areas were sampled for bacteria. The data showed that there were no significant differences in the amount of bacteria found. They also found some bacteria that was genetically different than any know bacteria on Earth.
What do you expect to change? Do you expect any changes?	I expected to see steady growth over time.	I expected to see the bacteria grow slower in space.
What else do you notice?	Some of the samples were 100% covered in bacteria in a very short period of time. The colony took over and likely consumed all of the nutrients. Nothing else grew within the dish.	Most of the bacteria found on the ISS resembled bacteria found on animal skin surfaces so this means the bacteria comes from humans.
	No Data	Staph and Entero bacteria was found on the ISS.





Observation based findings in comparison from observation data collected from a similar experiment conducted on the International Space Station (ISS).



ISS Experiment Data provided by DreamUp & found through a literature review.





ANALYSIS D

I collected quantitative data by using a grid to gather data to calculate percent coverage.

There were 100 squares that each petri dish covered.

I counted the number of squares with bacterial growth.

Number of Squares Filled With Bacteria

I estimated the growth rates based on the samples and the experiment days.





DATA COLLECTION D

(Sample Number*)													
(Day)	S1	S2	S3	S4	S5	S6	S7	S8	S 9	S10	S11	S12	S13
1	0	0	0	0	0	0	0	0	0	0	0	0	0
2	6	15	5	50	100	1	100	100	0	0	1	1	5
3	6	17	11	60	100	9	100	100	8	80	1	1	20
4	15	20	13	70	100	10	100	100	9	85	100	95	25
5	55	25	30	75	100	20	100	100	25	95	100	95	100
6	<mark>60</mark>	30	40	75	100	30	100	100	30	95	100	95	100
7	70	55	50	79	100	35	100	100	45	95	100	95	100
8	80	70	<mark>6</mark> 0	80	100	40	100	100	55	96	100	95	100
9	85	70	75	90	100	50	100	100	<mark>6</mark> 0	98	100	95	100
10	90	70	75	90	100	50	100	100	75	98	100	95	100
11	95	75	80	90	100	55	100	100	80	98	100	95	100
12	100	75	80	90	100	55	100	100	80	98	100	95	100
13	100	75	80	95	100	60	100	100	80	98	100	95	100
14	100	75	80	95	100	60	100	100	85	98	100	95	100
15	100	75	80	95	100	60	100	100	85	98	100	95	100



Notice that many of my samples grew rapidly and filled all 100 squares within the first 5 days.



DATA COLLECTION C













These charts represent the sample growth over a 15 - day time period.



DATA COLLECTION C









These charts represent the sample growth over a 15 - day time period.





INDEPENDENT VARIBLES

- Light Duration & Intensity
- Moisture
- Temperature
- Nutrients

- DEPENDANT VARIBLES
- Rate of Growth
- Size of bacteria
- Shape of Bacteria
- Shape of Bacterial Colony





- I conducted a literature review of several germ growth articles relating to experiments conducted in space.
- I conducted a few informational interviews with NASA experts that have been involved in space germ growth experiments.
- I used actual ISS experimental data via DreamUp for my comparative analysis.



Interviews & Expert Discussions

INTERVIEW 1 ASTRONAUT DR. SIAN PROCTOR DATE: 11 October 2021

I had a discussion with Astronaut Dr. Sian Proctor about her recent trip to Space with SpaceX in September 2021.

We discussed her biological research conducted in space regarding germs and how after a period of time some germs have been known to swap from individual to individual.

Individual swabs were collected for analysis and study upon the crew's return to Earth.









Interviews & Expert Discussions

INTERVIEW 2 ANONOMOUS NASA SCIENTIST DATE: 24 October 2021

We discussed data collection methods and data analysis methods.

We discussed how bacteria react in space.



RESULTS & FINDINGS

I assessed my findings and determined that my hypothesis for conducting a comparative analysis with my data and with data from a similar experiment conducted on the ISS was incorrect.

I assumed that the bacteria would grow slower in space. Findings suggest that bacteria and fungi grow faster in space. They both thrive in low gravity environments and often form groups. They form biofilms which is a protective mode of growth. This allows them to survive in hostile environments and to colonize in new and extreme areas. It is important to note that some types of opportunistic microbes were found on the ISS which could pose risks to humans.

Most of the bacteria found on the ISS resembled bacteria found on animal skin surfaces so this means the bacteria more than likely comes from the astronauts on the ISS. This aligns with some of the things that I learned during my informational interview with Dr. Sian Proctor who had recently returned from space.

Its important to note that bacteria were also found outside of the ISS. Specific research conducted determined that they can survive in the harsh space environment for at least 3 years.

The bacteria found in the locations that I sampled on Earth appeared to have all types of bacterial growth from both animals and plants. This finding differed from the ISS findings however this was not surprising.

My quantitative data was surprising as several of my samples grew extremely fast. One day there was no growth. The following day, several of my samples were 100 percent covered with growth. It appeared that one specific type of colony consumed all of the nutrients.

RESULTS & FINDINGS

- My hypothesis was incorrect.
- Germs grow much faster in space.
- In space, germs go into a survival mode and form groups and biofilms.
- My samples consisted of animal and plant cells.
- Bacteria found on the ISS are animal cells coming from humans.
- My quantitative data showed exponential bacterial growth.

APPLICATIONS

- There are many applications for my research.
- Astronauts who travel into deep space will need to be able to understand the impacts of bacteria and what disinfectants and potential medicines that they may need to bring with them.

- Bacteria in space may be able to assist with plant growth in low nutrient environments (Earth and in space).
- The discoveries that we make in space, the ones that I hope to make, will benefit life in space and on earth.

CONCLUSION

KEY POINTS

- It is important to understand the behaviors bacteria and fungi in the space environment.
- This is important for human health, performance and for plant growth. It is also important to understand these behaviors so that we do not contaminate or expose ourselves to harmful environments that we visit while exploring space.
- This type of research is important also for extreme environments on Earth. As we understand bacterial behavior more in space, hopefully we can come up with solutions to grow food for populations on Earth that need it.
- I intend to carry out research in the future to understand extremophile bacteria and how it can benefit life on Earth.

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