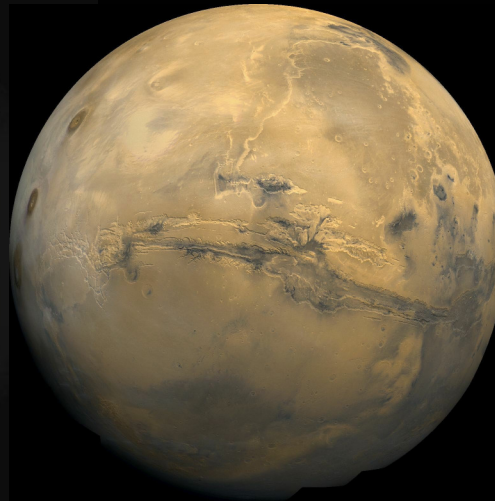
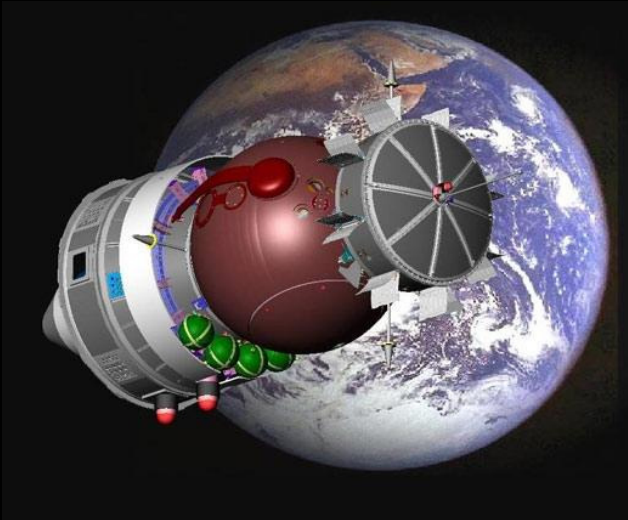


10 Tage im All:

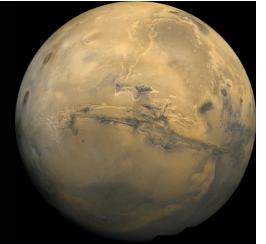
Alpine Flechten und die Wahrscheinlichkeit der Marsumwelt zu widerstehen



DLR-Status-Seminar Strahlen-/Astrobiologie 13.06.2008

Jean-Pierre de Vera, S. Ott

Earth/Mars-Characteristics



PARAMETER	ERDE	MARS
Mittlerer Sonnenabstand:	149,6 Mio. km	227,9 Mio. km
Mittlere Umlaufzeit um die Sonne:	365,25 Tage	686,98 Tage
Rotationsperiode:	Ca. 23 h 56 min	Ca. 24 h 37 min
Äquatorialdurchmesser:	12.756 km	6794 km
Mittlere Oberflächentemperatur:	+22° Grad Celsius	-55° Grad Celsius
Mittlere Sonneneinstrahlung:	1,37 kW/m ²	0,6 kW/m ²
Mittlerer Atmosphärendruck an der Oberfläche:	1013 mbar	6-10 mbar
Zusammensetzung der Atmosphäre:	78% Stickstoff 21% Sauerstoff 0,93% Argon 0,035% Kohlendioxid	0,3% Sauerstoff 2-3% Stickstoff 1-2% Argon 95% Kohlendioxid
Monde	Luna	Phobos und Deimos

Important
for
simulation

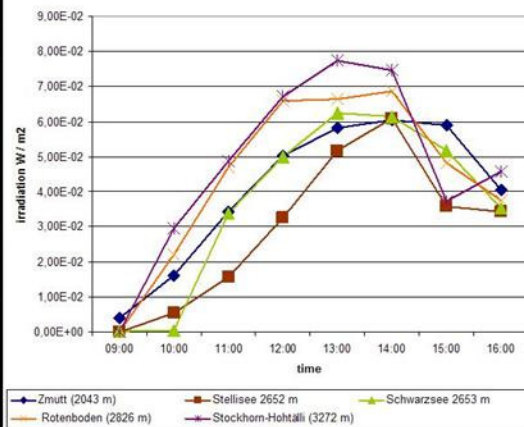
HABITAT CHARACTERISATION OF THE LICHEN SAMPLES HAS BEEN PERFORMED IN SWITZERLAND AND SPAIN

- **Radiation profile (UV, InfraRed and PAR)**
- **Temperature profile (air, soil/rock)**
- **Humidity profile (air, soil/lichen)**
- **Ozone profile**
- **altitude profile (between 2000 m and 3200 m)**
- **pressure**
- **Aerosol profile**
- **photosynthetic activity profile**
- **cartographic overview by helicopter (thanks to Air Zermatt)**

Some examples ...

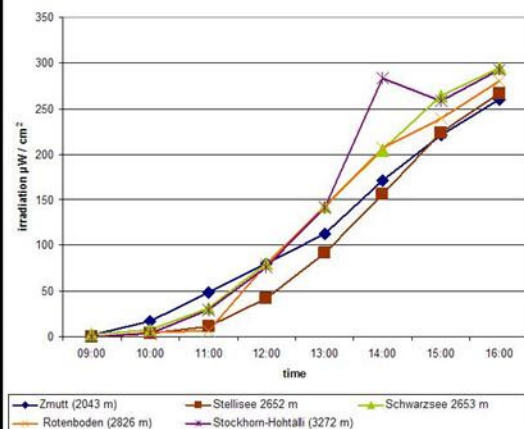
UV-irradiation profile

Irradiation of UV B (305 nm) at different habitats (record time between 26.08 and 03.09.2007)



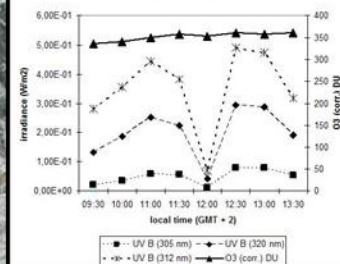
UV-dosis/day profile

Erythema radiation UV B daily dose augmentation (record time between 26.08 and 03.09.2007)



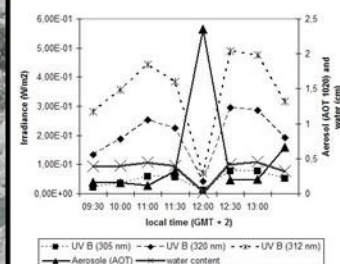
Ozone

Changing UV B irradiance in relation to Ozone concentration



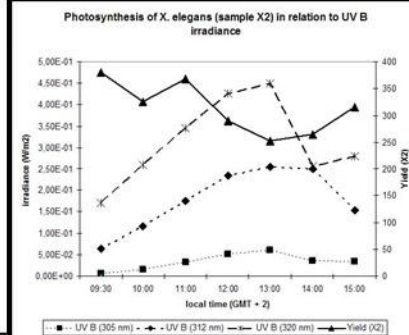
Aerosol and water conc.

Changing UV B irradiance in relation to aerosole and water concentration

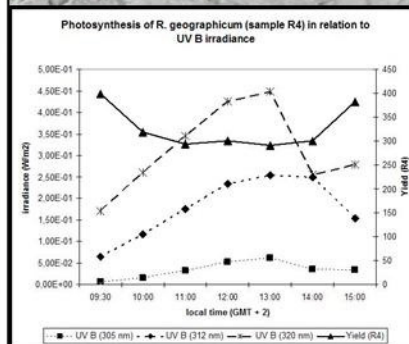


Photosynthesis profile of both lichens in relation to UV radiation

X. elegans

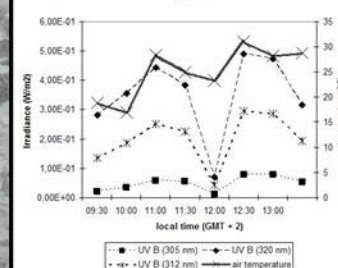


R. geographicum



Changing rock and air temperatures

Changing UV B irradiance and changing rock and air temperature



ETC ...

Habitat area characterization of *Rhizocarpon geographicum* and *Xanthoria elegans* by an overview out of the helicopter

Rotenboden (2826 m)



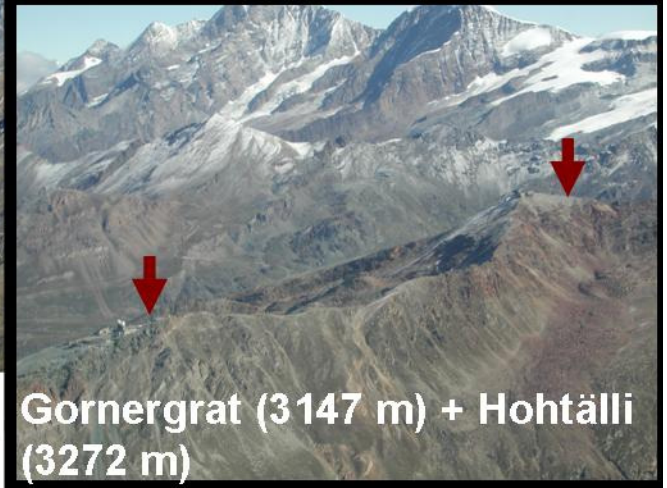
Trockener Steg (2948 m)

Schwarzsee (2653 m)

Furgg (2316 m)



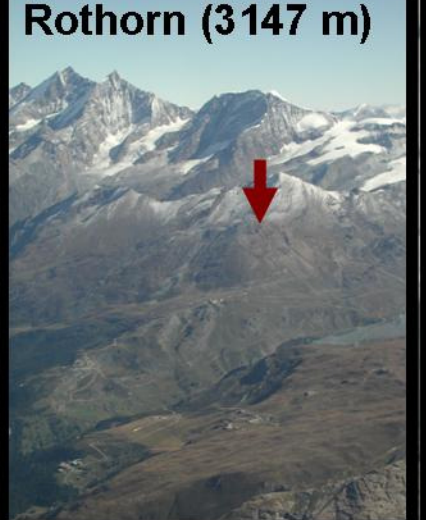
Gornergrat (3147 m) + Hohtälli (3272 m)



Zmutt (2043 m)



Rothorn (3147 m)



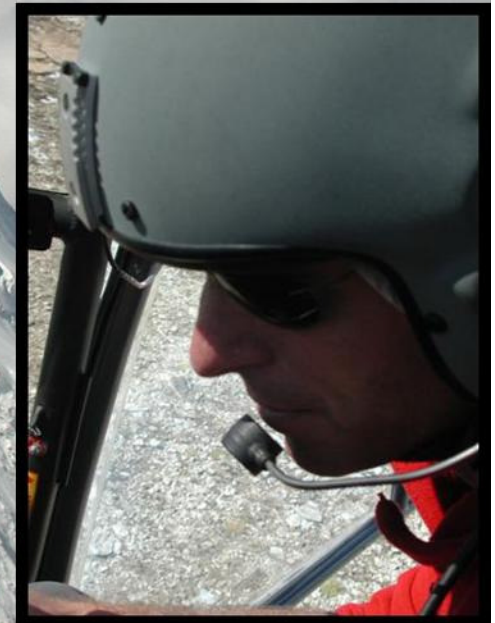
Stellisee (2652 m)



Habitat area characterization by an overview out of the helicopter

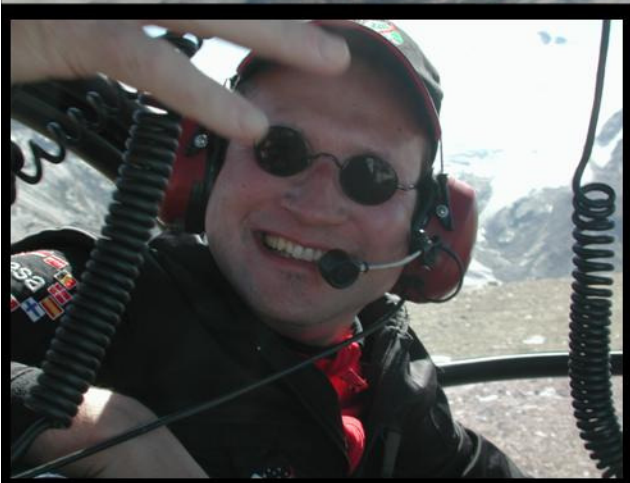
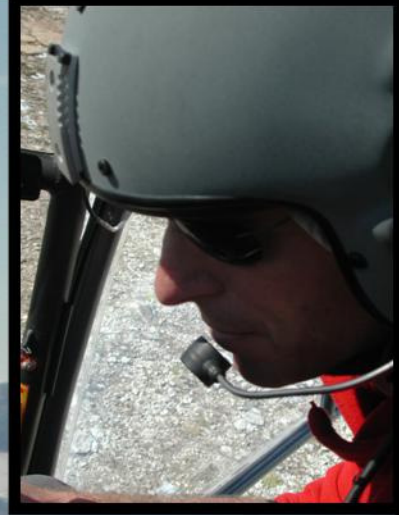


Publicity for ESA



Conclusion

- The habitats are mostly on exposed high altitudes with high UV radiation income
- The soil humidity is 2/3 of the year between 20 and 47 %
- Water availability by ice, snow-melting, fog and rain (but half of the year under snow)
- Some valleys and mountain peaks are more dominated by *Rhizocarpon geographicum* and others by *Xanthoria elegans*
- Soil composition varies between calciferous rocks, serpentinite and gneiss

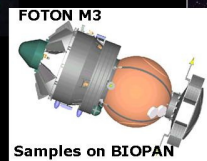
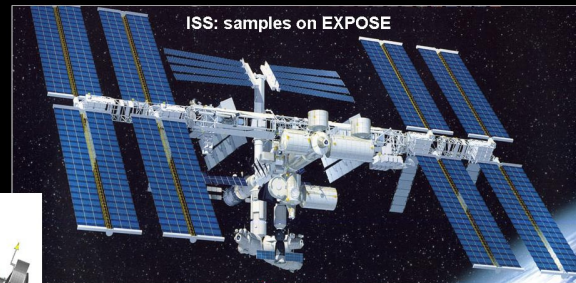


Lithopanspermia

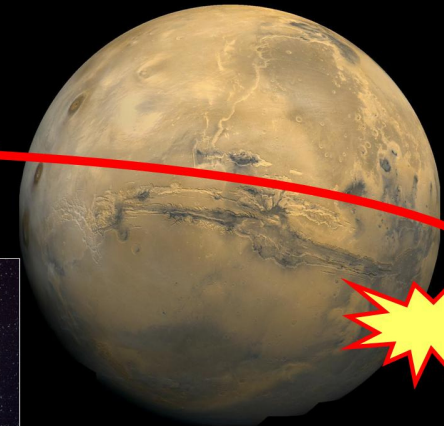
After space simulation: real space exposure
(long duration experiments)



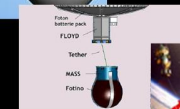
Travel through space



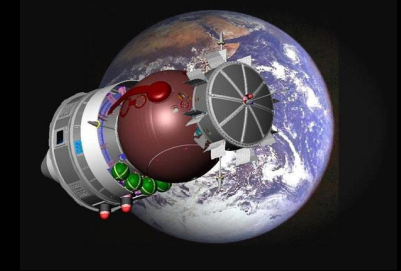
Impact
and
ejection



Reentry
process



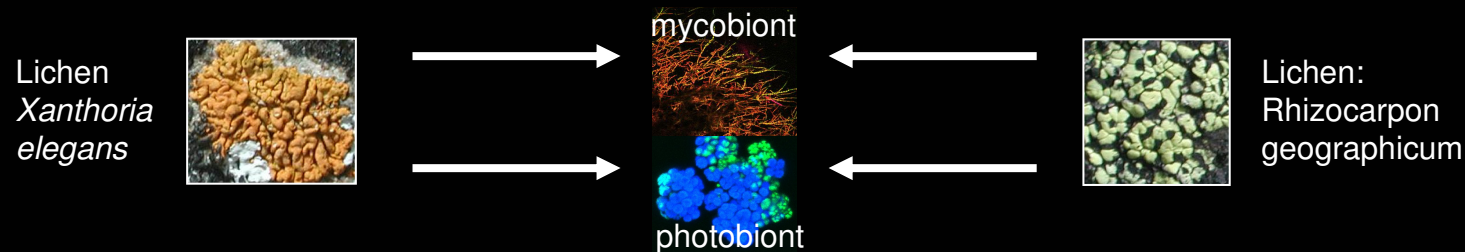
BIOPAN 6 / Lithopanspermia



Science objectives

Viability, resistance and adaptation mechanisms of lichens and their symbionts (mycobiont / photobiont) / maintenance or conservation of photosynthetically active cells in space.

The role of symbiosis and the likelihood of Lithopanspermia of an eukaryotic system



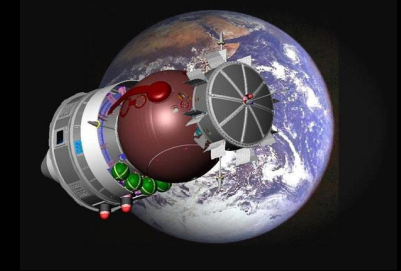
Expected science results

- Preservation of life processes (e.g. growth and germination capacity) due to resistance and adaptation mechanisms
- determination of the most resistant symbiont in the lichen community
- detection of the capacity of photosynthesis preservation after space exposure (Panspermia)
- the influence of space parameters on the DNA on an eukaryotic system / micro-ecology
- the influence of space environment on physiological processes

Space-relevance

-- the likelihood of Lithopanspermia (natural transfer of eukaryotes, evolution aspects)

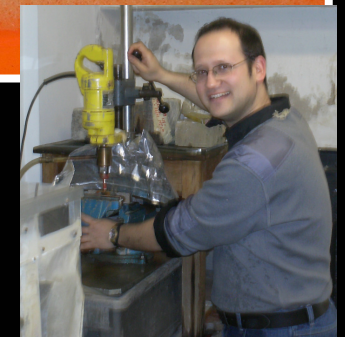
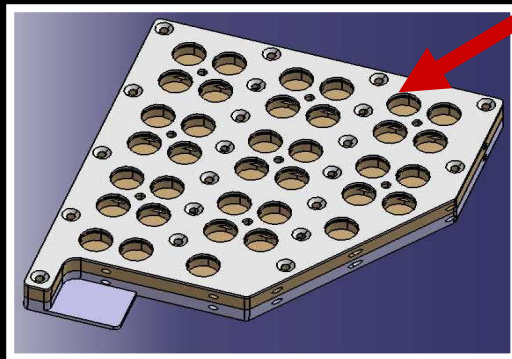
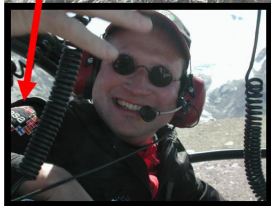
Samples: fruiting bodies and thalli of *Rhizocarpon geographicum* and *Xanthoria elegans*



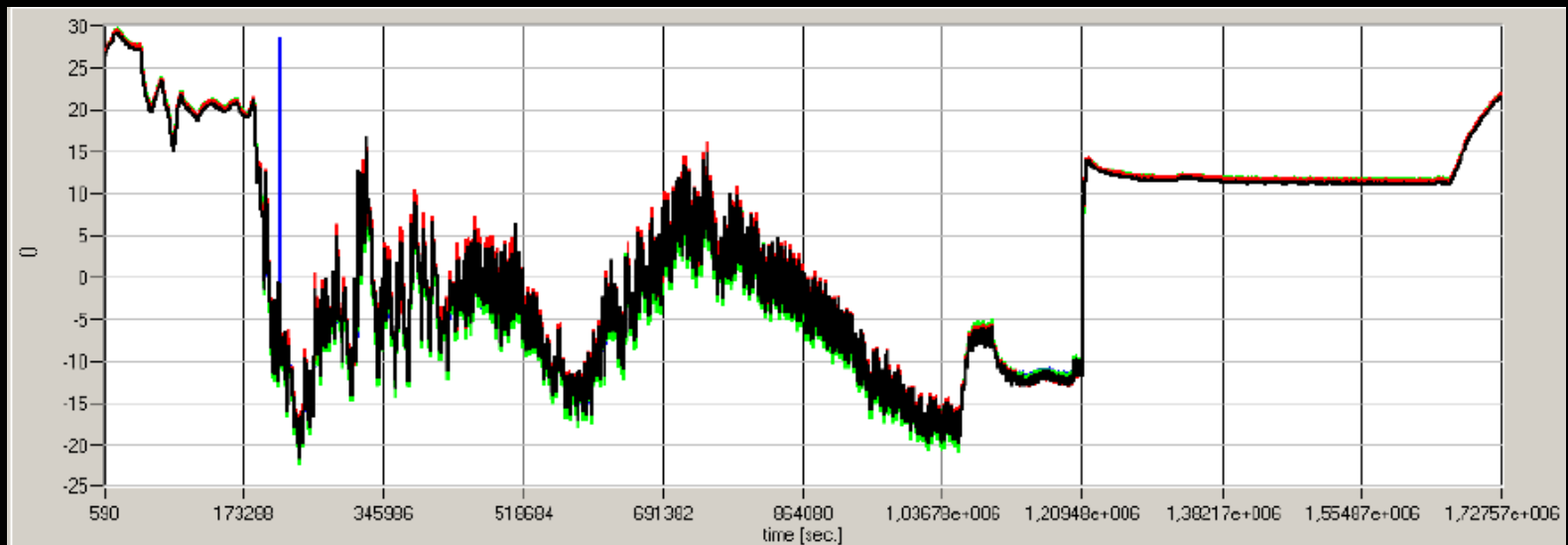
Habitat area characterization by an overview out of the helicopter



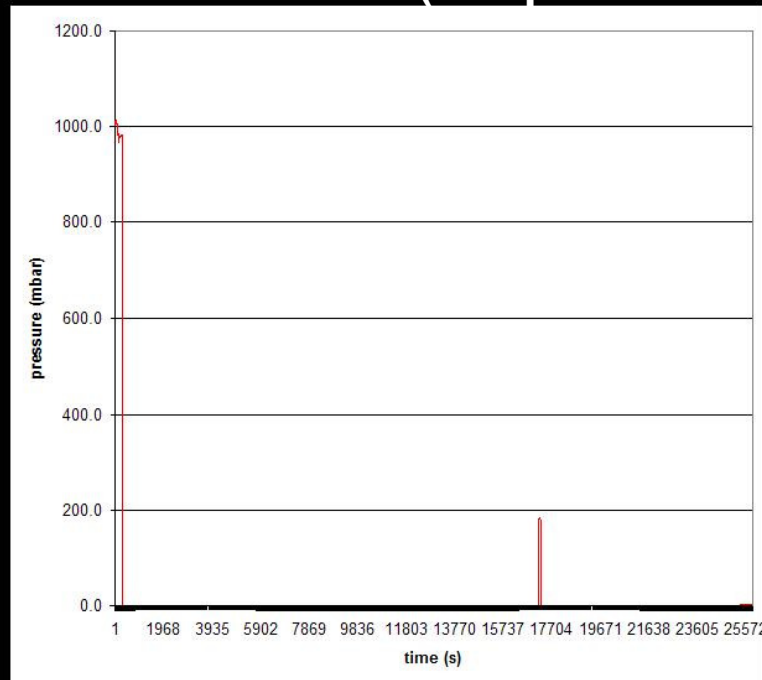
Publicity for ESA



Temperature on BIOPAN



Pressure on BIOPAN (Endpressure ~ 0.4 mbar)

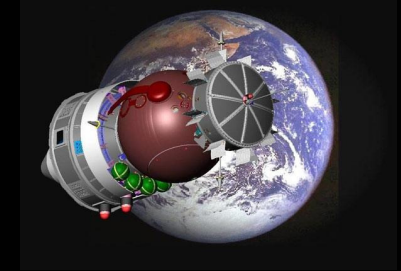


Dose on BIOPAN (SCh = solar constant hour)

mission	Biopan lid open	Biopan sensors		R3D-B sensors	
		total dose	daily dose	total dose	daily dose
Biopan-6	240 h = 10.0 d	24 SCh	2.4 SCh	36.3 SCh	3.63 SCh
		11.8 kJ cm ⁻²	1.18 kJ cm ⁻²	17.9 kJ cm ⁻²	1.79 kJ cm ⁻²

MEHODS:

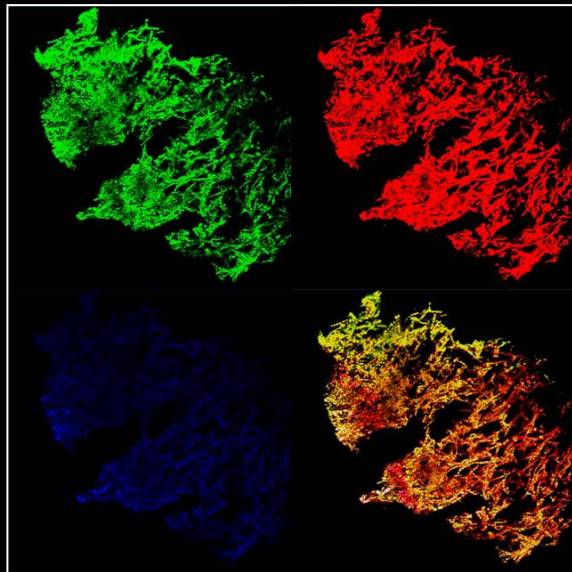
vitality check



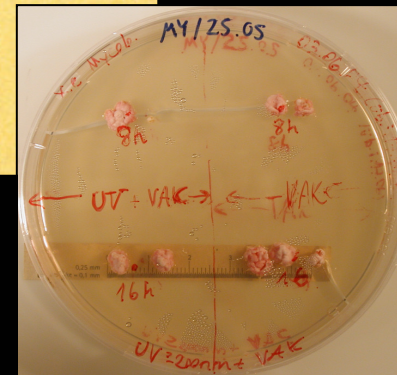
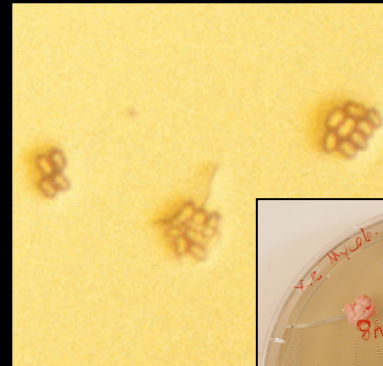
Physiology /
LIVE/DEAD

FUN I +
SYTOX-
green

Image-
Tool /
Statistics

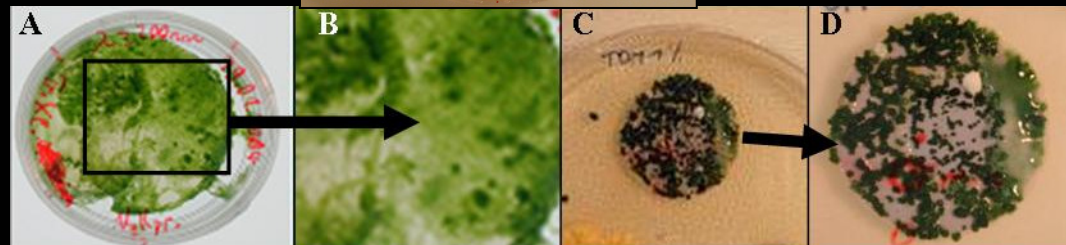
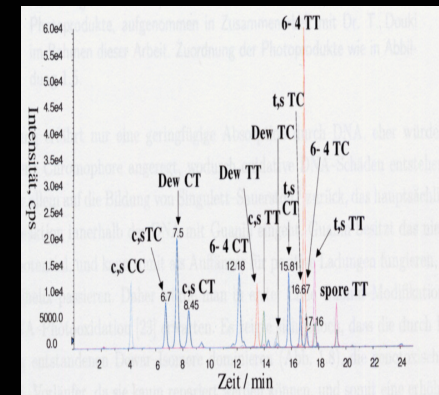


Growth capacity of
symbionts /
germination capacity of
ascospores



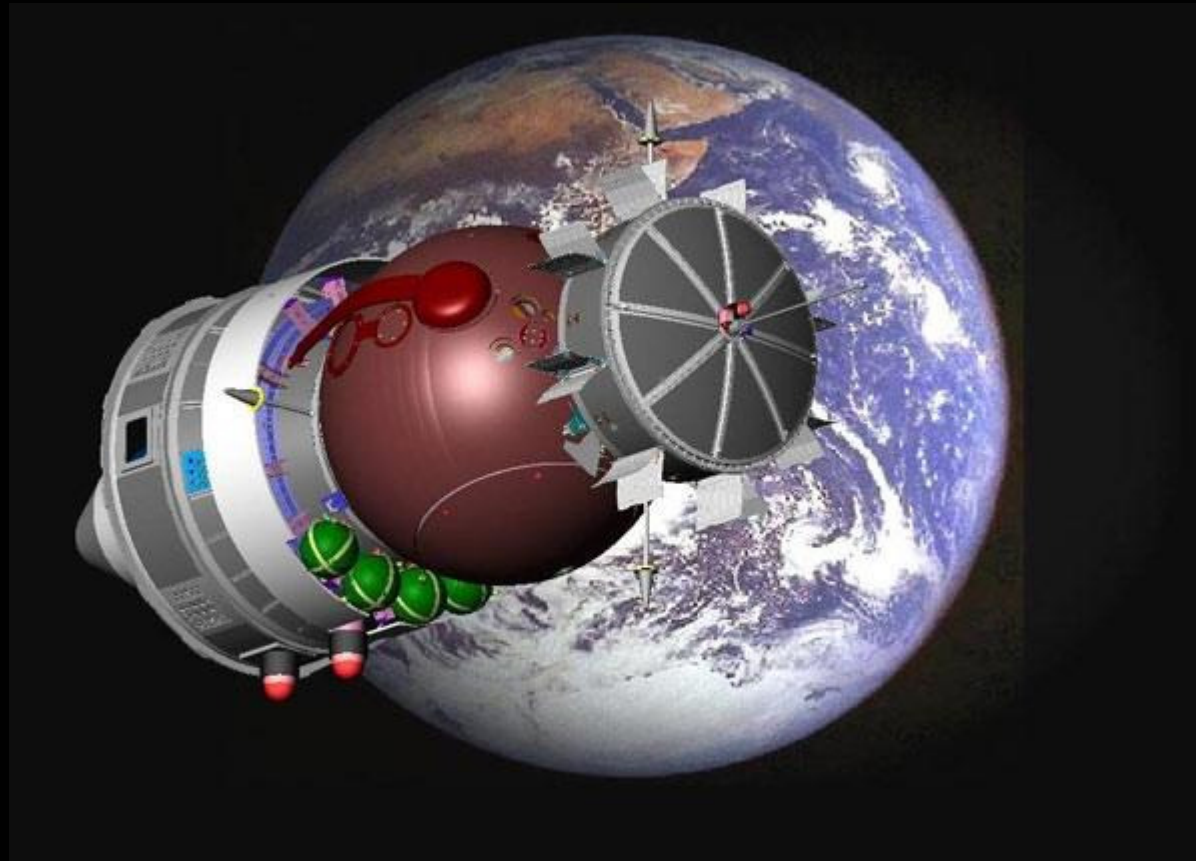
mutagenity

T-T-
photoproducts in
the DNA after UV
radiation



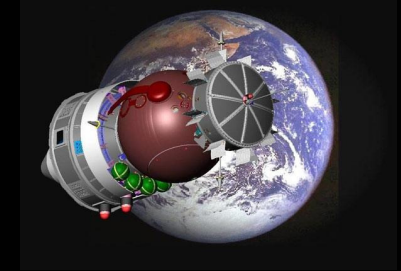
Results

LIVE / DEAD analysis by CLSM

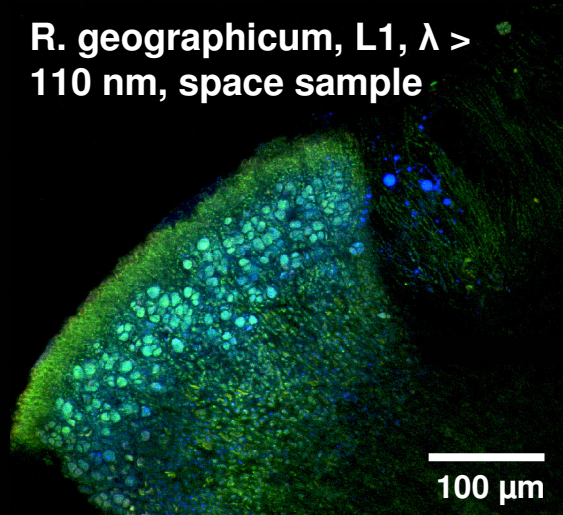


RESULTS:

CLSM Analysis by the use of LIVE/DEAD dye



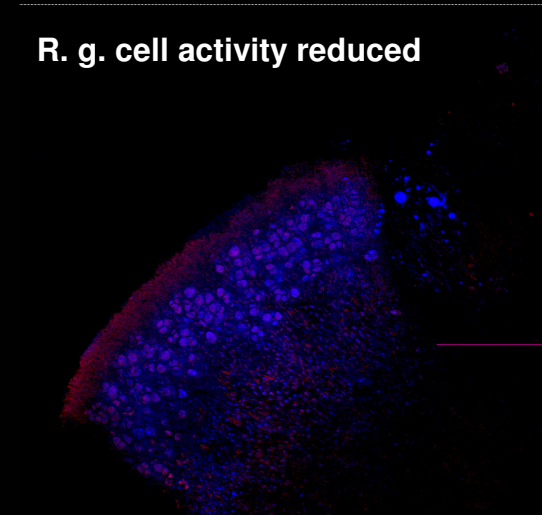
R. geographicum, L1, $\lambda > 110$ nm, space sample



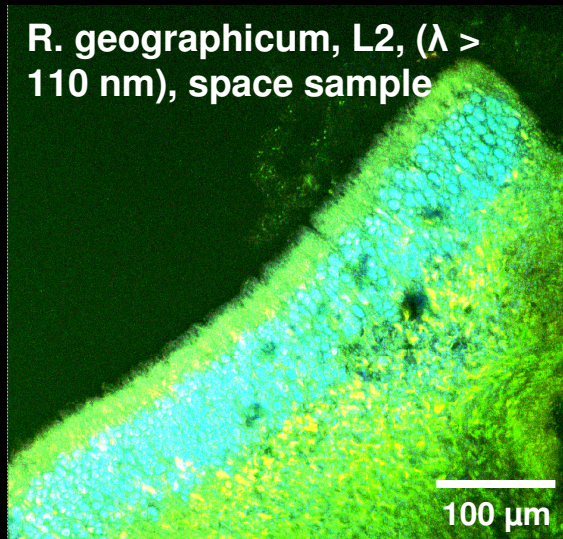
30 min later



R. g. cell activity reduced



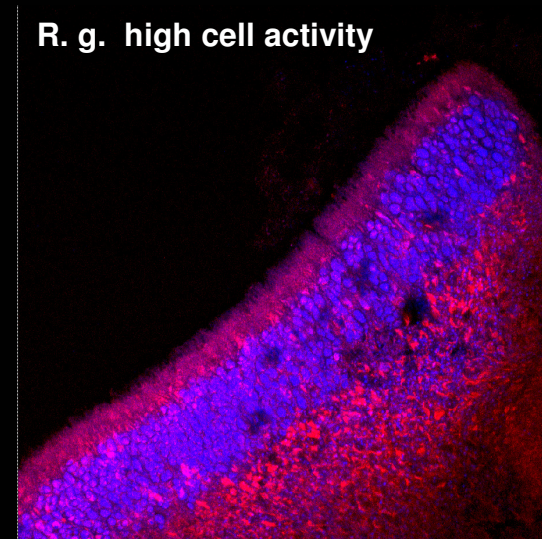
R. geographicum, L2, ($\lambda > 110$ nm), space sample



30 min later



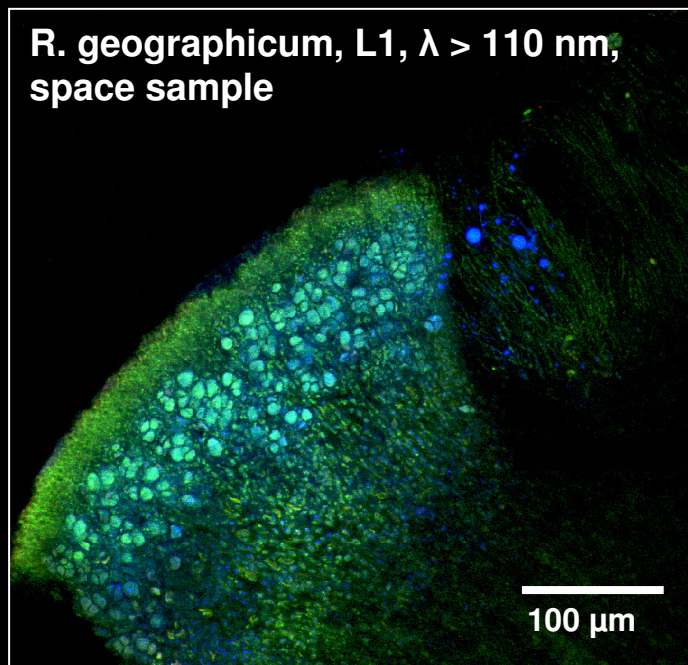
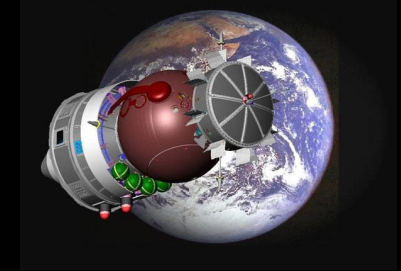
R. g. high cell activity



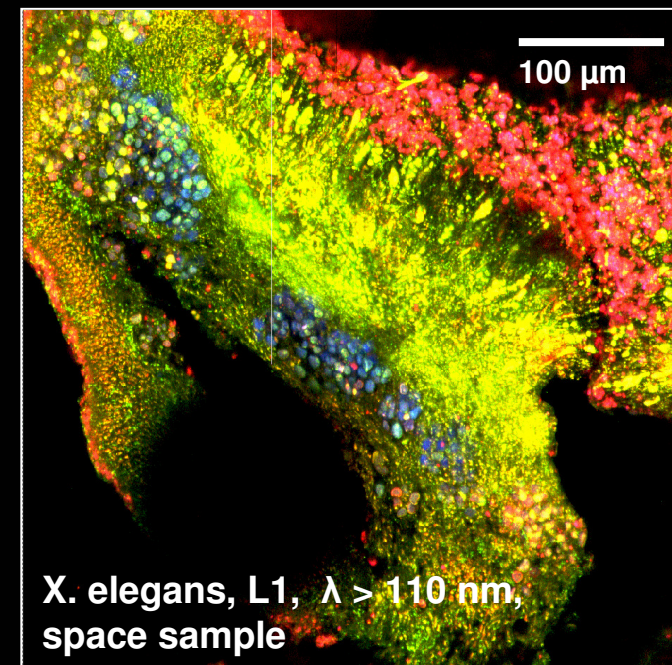
RESULTS:

CLSM Analysis by the use of LIVE/DEAD dye

Comparison between *R. geographicum* and *X. elegans*



Photobiont cells of *R.g.* are more vital than those of *X.e.*

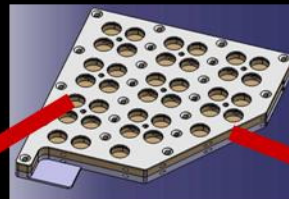


X. elegans, L1, $\lambda > 110$ nm, space sample

Mycobiont cells of *X.e.* are more vital than those of *R.g.*

RESULTS: samples from space

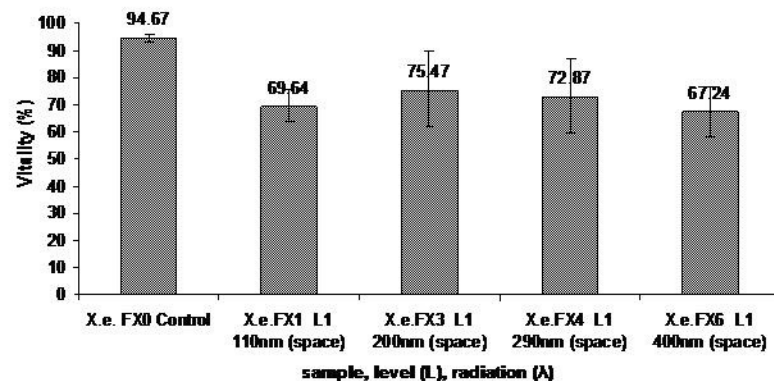
Quantified results after Analysis of CLSM images by the use of Image J



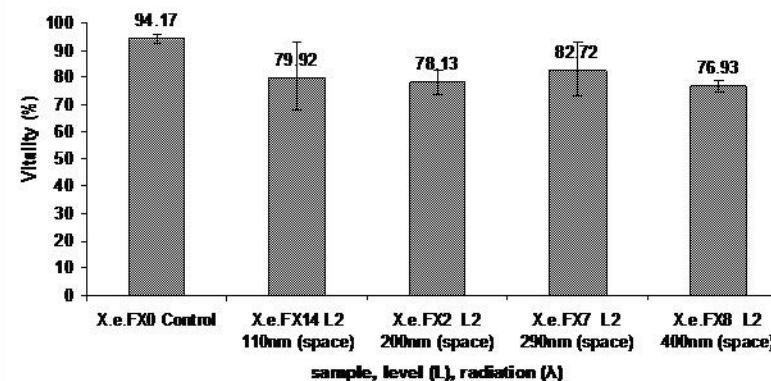
L 1

L 2

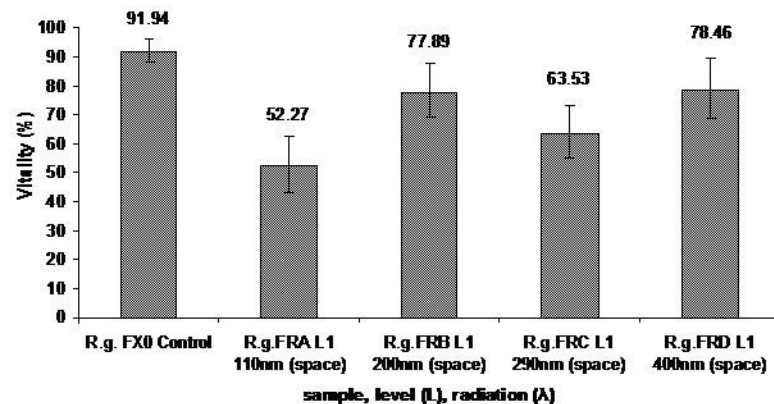
Vitality of thalli and fruiting bodies of *X. elegans* after space exposure and CLSM analysis with LIVE/DEAD dye FUN I on level 1



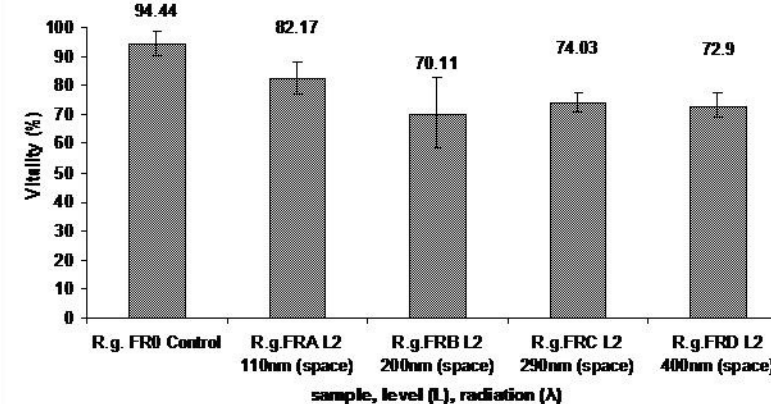
Vitality of thalli and fruiting bodies of *X. elegans* after space exposure and CLSM analysis with LIVE/DEAD dye FUN I on level 2



Vitality of thalli and fruiting bodies of *R. geographicum* after space exposure and CLSM analysis with LIVE/DEAD dye FUN I on level 1

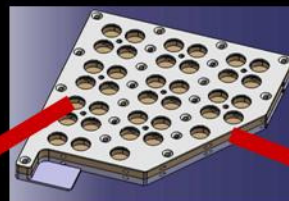
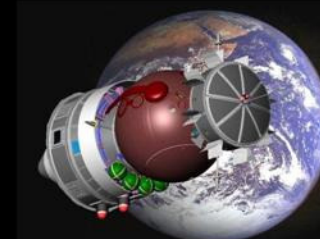


Vitality of thalli and fruiting bodies of *R. geographicum* after space exposure and CLSM analysis with LIVE/DEAD dye FUN I on level 2



RESULTS: simulation tests

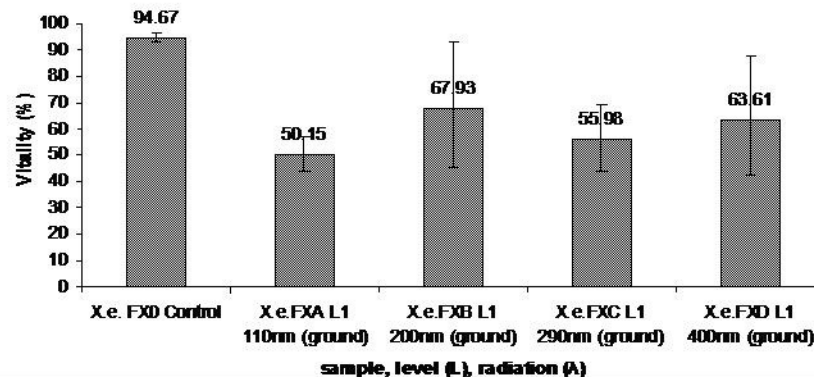
Quantified results after Analysis of CLSM images by the use of Image J



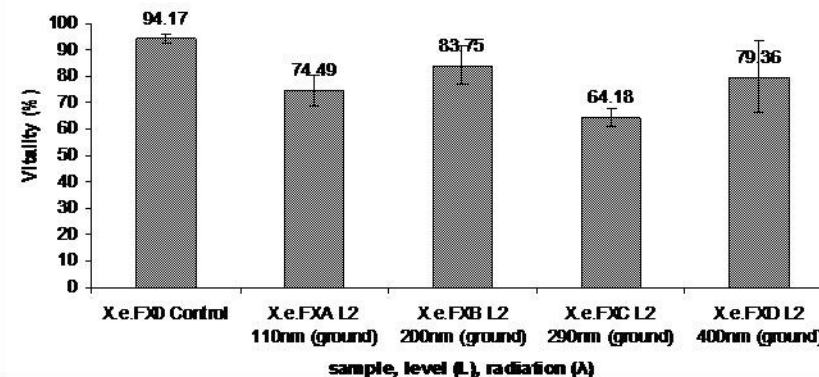
L 1

L 2

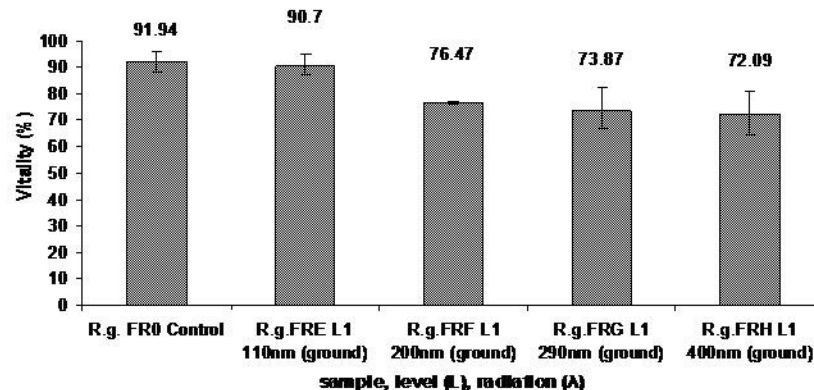
Vitality of thalli and fruiting bodies of *X. elegans* after ground simulation tests and CLSM analysis with LIVE/DEAD dye FUN 1 on level 1



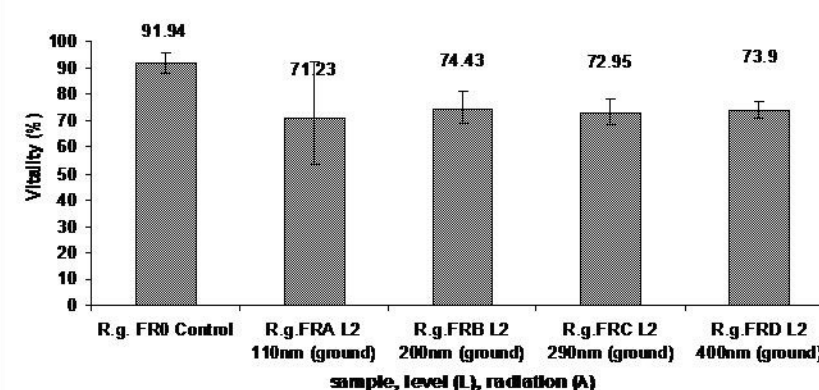
Vitality of thalli and fruiting bodies of *X. elegans* after ground simulation tests and CLSM analysis with LIVE/DEAD dye FUN 1 on level 2



Vitality of thalli and fruiting bodies of *R. geographicum* after ground simulation tests and CLSM analysis with LIVE/DEAD dye FUN 1 on level 1

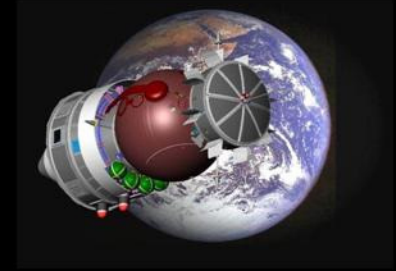


Vitality of thalli and fruiting bodies of *R. geographicum* after ground simulation tests and CLSM analysis with LIVE/DEAD dye FUN 1 on level 2

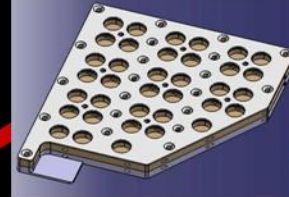


RESULTS: summary

Quantified results after Analysis of CLSM images by the use of Image J

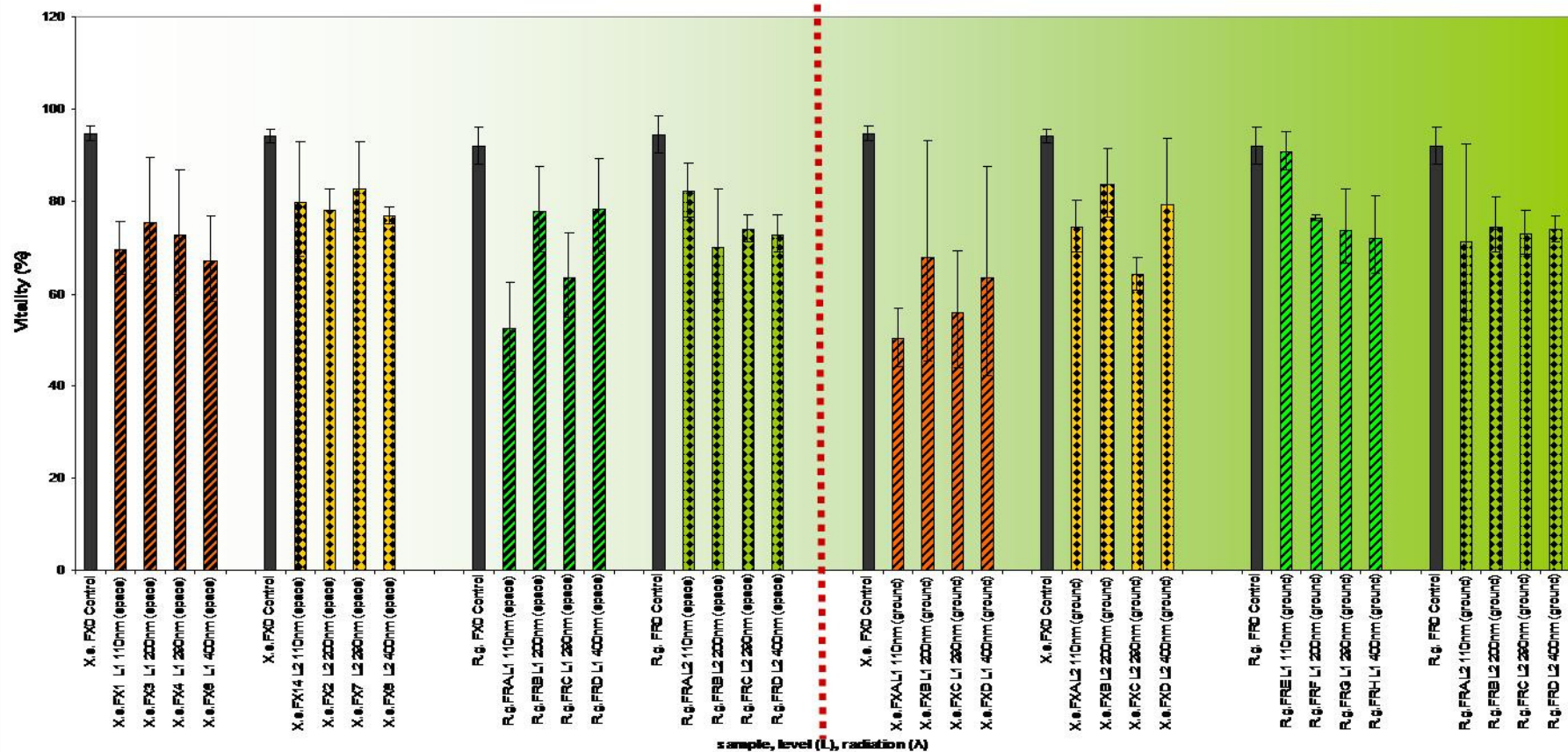


Space exposure

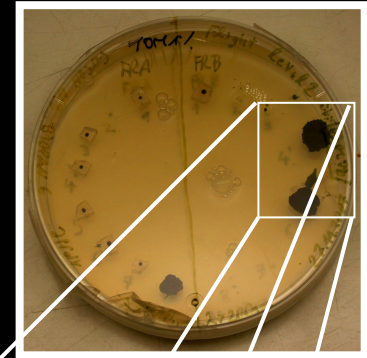
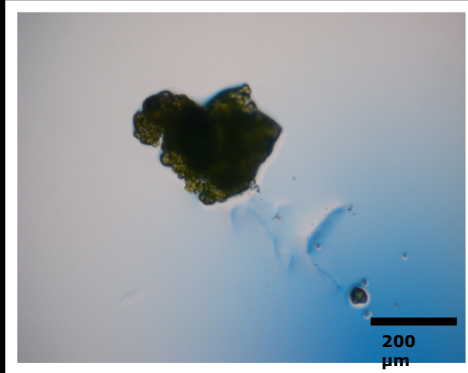


Simulation tests

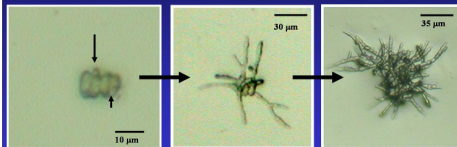
Vitality of thalli and fruiting bodies of *R. geographicum* and *X. elegans* after space / simulation exposure and CLSM analysis with FUN I



Germination / Growth capacity



FOTON M3 / BIOPAN: > 110 nm and vacuum, level 1 (sample FX1, on MY agar with amp.)
Growth capacity of ascospores of *Xanthoria elegans*

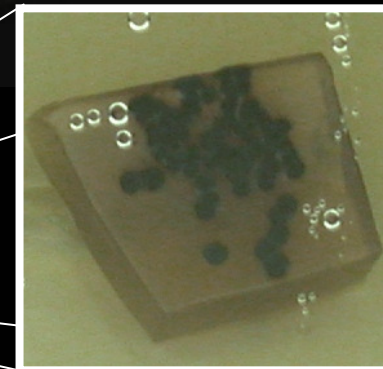
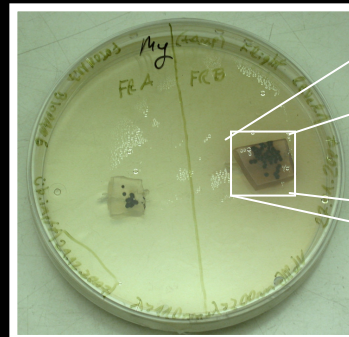
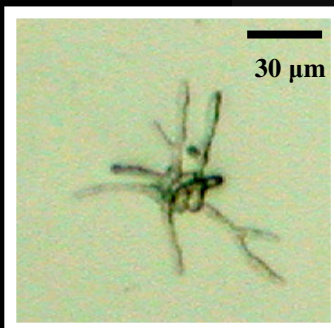


2 days after sporulation:
first germination process
occurs

10 - 12 days after
sporulation: first
branching of the young
mycelia

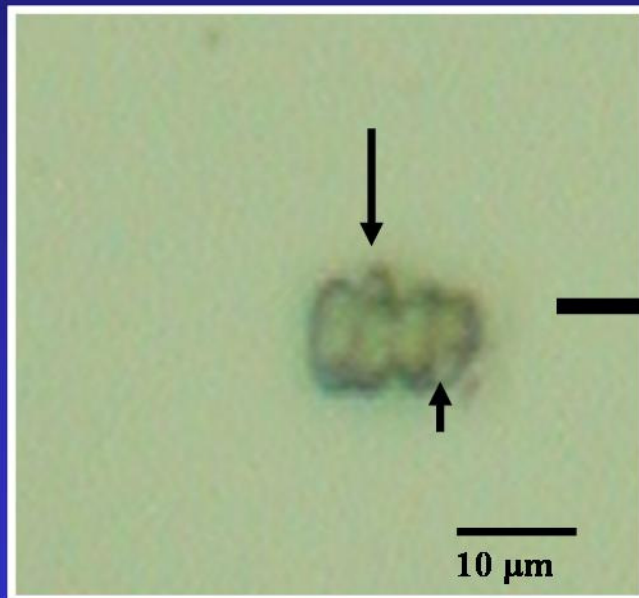
22 days after sporulation:
multiple branching

JPP4V

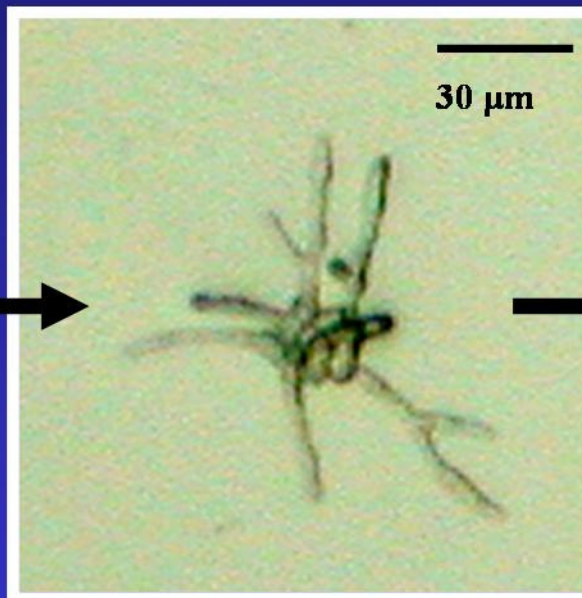


**FOTON M3 / BIOPAN: > 110 nm and vacuum, level 1 (sample FX1,
on MY agar with amp.)**

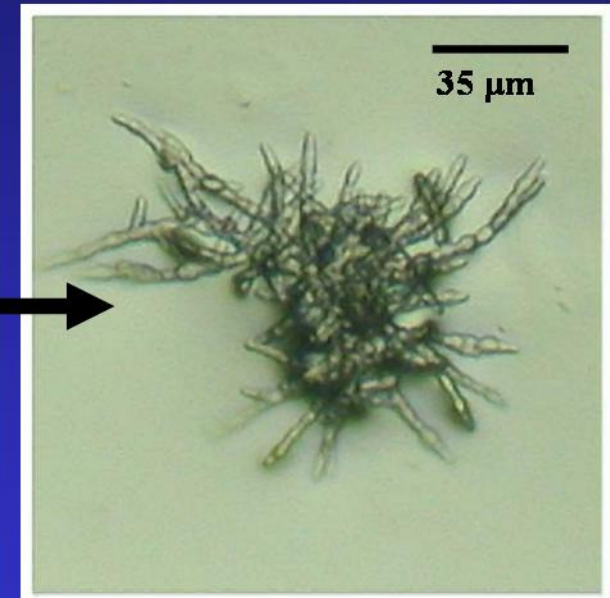
Growth capacity of ascospores of *Xanthoria elegans*



**2 days after sporulation:
first germination process
occures**



**10 - 12 days after
sporulation: first
branching of the young
mycelia**



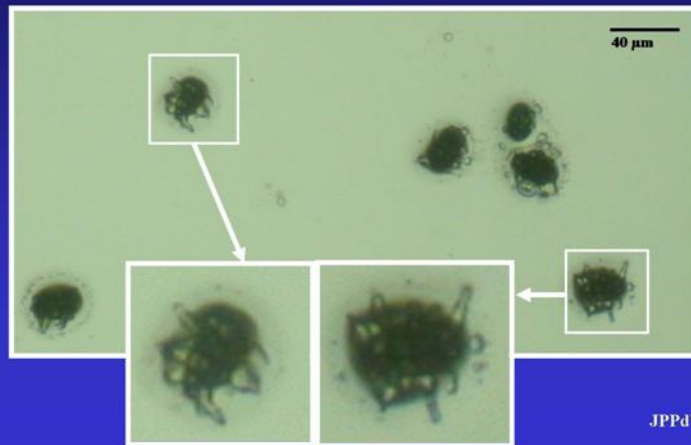
**22 days after sporulation:
multiple branching**

JPPdV

Germination and Growth capacity of ascospores

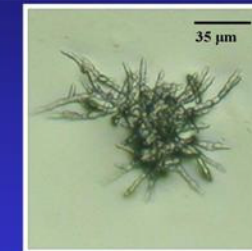
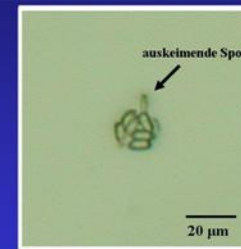


FOTON M3 / BIOPAN: > 290 nm and vacuum, level 1 (sample FR, on MY agar with amp.)
Germination capacity of ascospores of *Rhizocarpon geographicum*; 3 days after sporulation

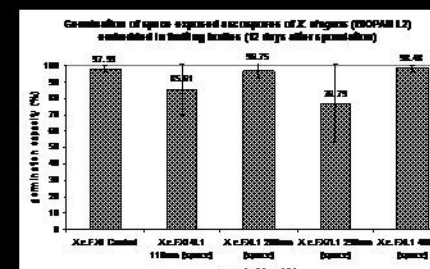
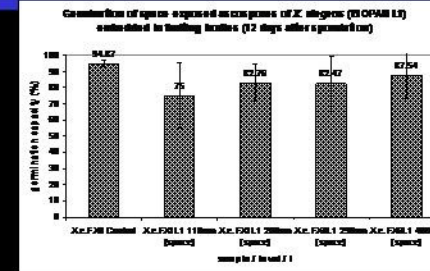
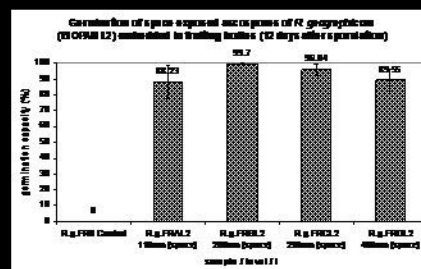
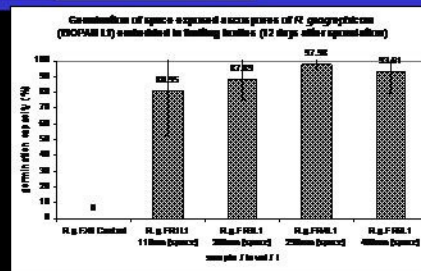
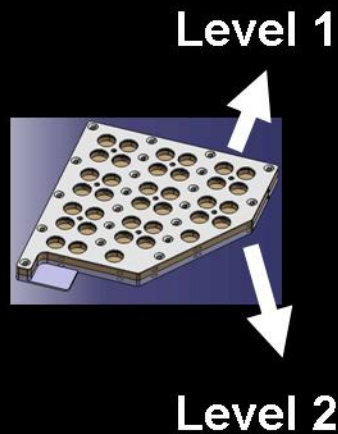


JPPdV

FOTON M3 / BIOPAN: >110 nm and vacuum, level 1 (sample FX on MY agar with amp.). Germination capacity of ascospores of *Xanthoria elegans*, 2 days after sporulation and growth of mycelia after 22 days



JPPdV

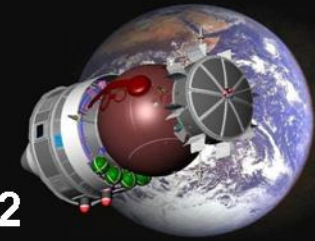


Germination and Growth capacity of ascospores

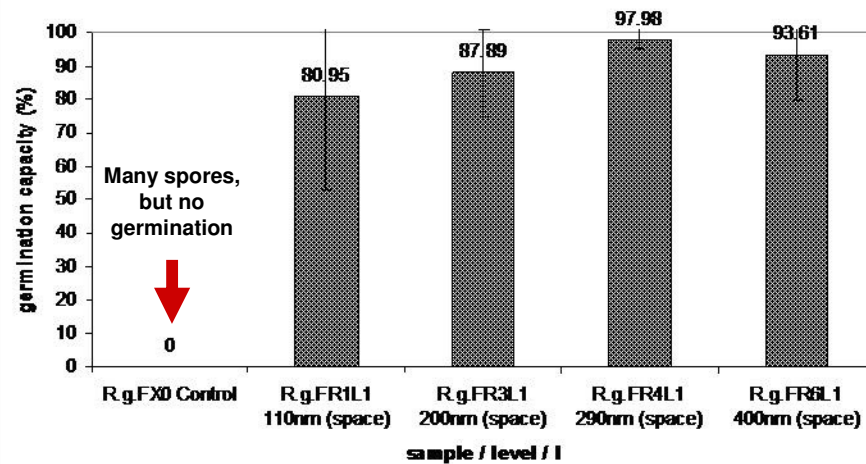
Level 1



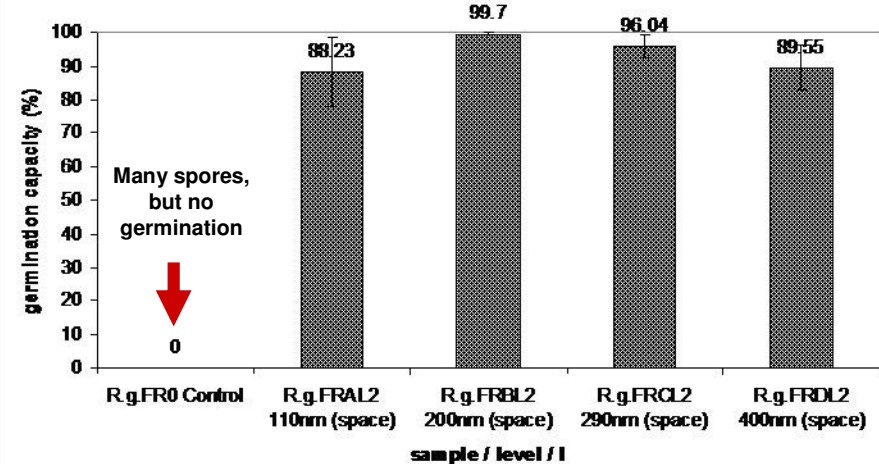
Level 2



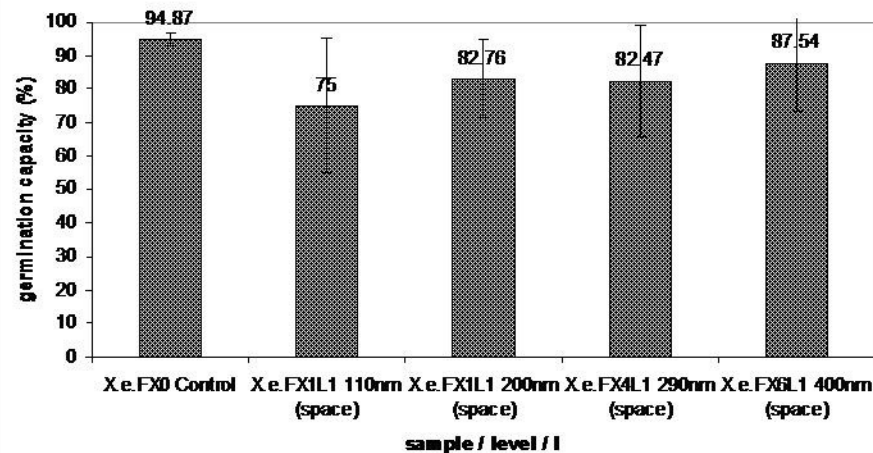
Germination of space exposed ascospores of *R. geographicum* (BIOPAN L1) embedded in fruiting bodies (12 days after sporulation)



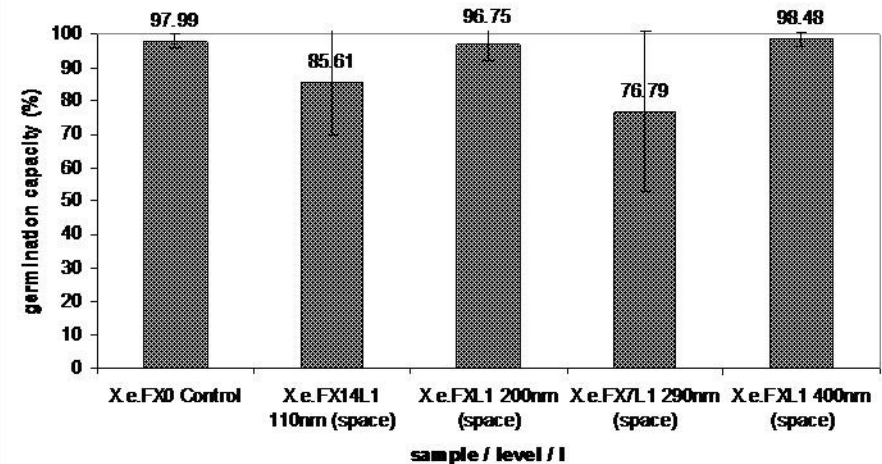
Germination of space exposed ascospores of *R. geographicum* (BIOPAN L2) embedded in fruiting bodies (12 days after sporulation)



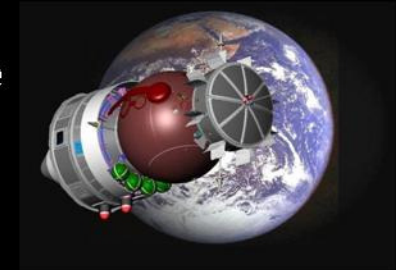
Germination of space exposed ascospores of *X. elegans* (BIOPAN L1) embedded in fruiting bodies (12 days after sporulation)



Germination of space exposed ascospores of *X. elegans* (BIOPAN L2) embedded in fruiting bodies (12 days after sporulation)

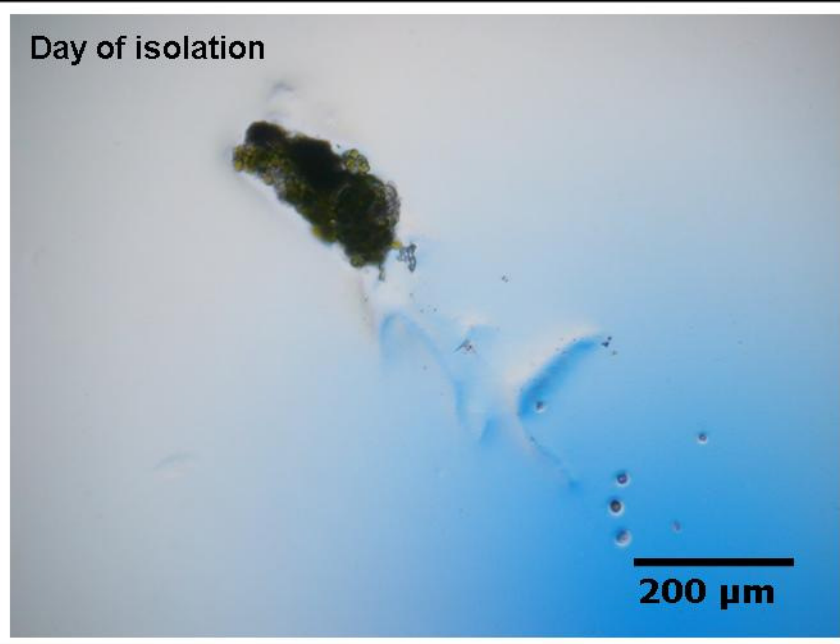


Growth capacity of photobiont cells after space exposure

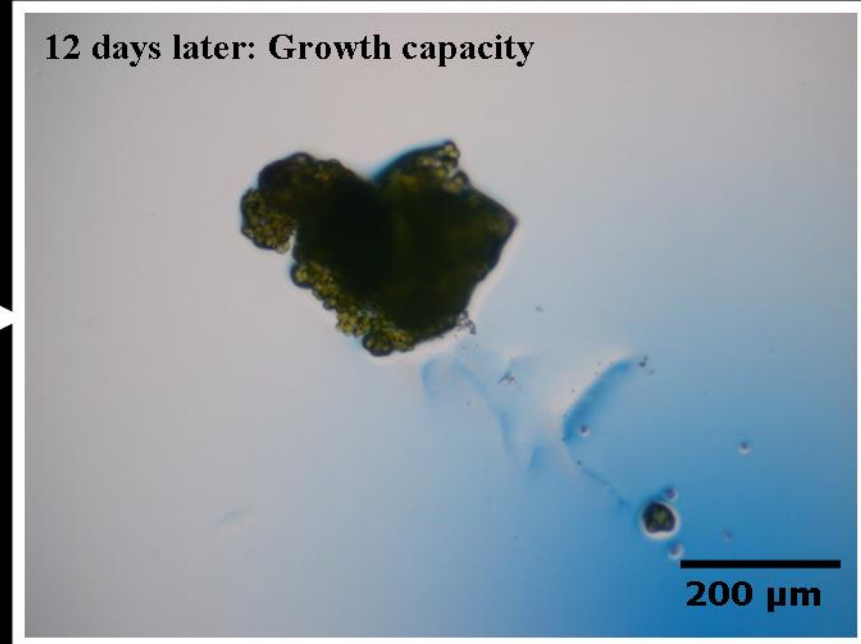


Photobiont of *Rhizocarpon geographicum* (>110 nm, vacuum, Level 1)

Day of isolation



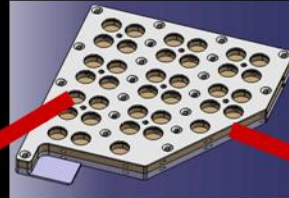
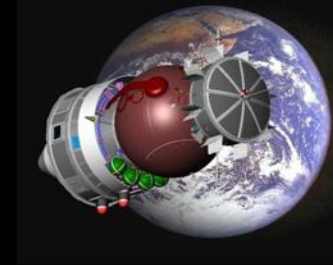
12 days later: Growth capacity



The work is further going on...

RESULTS:

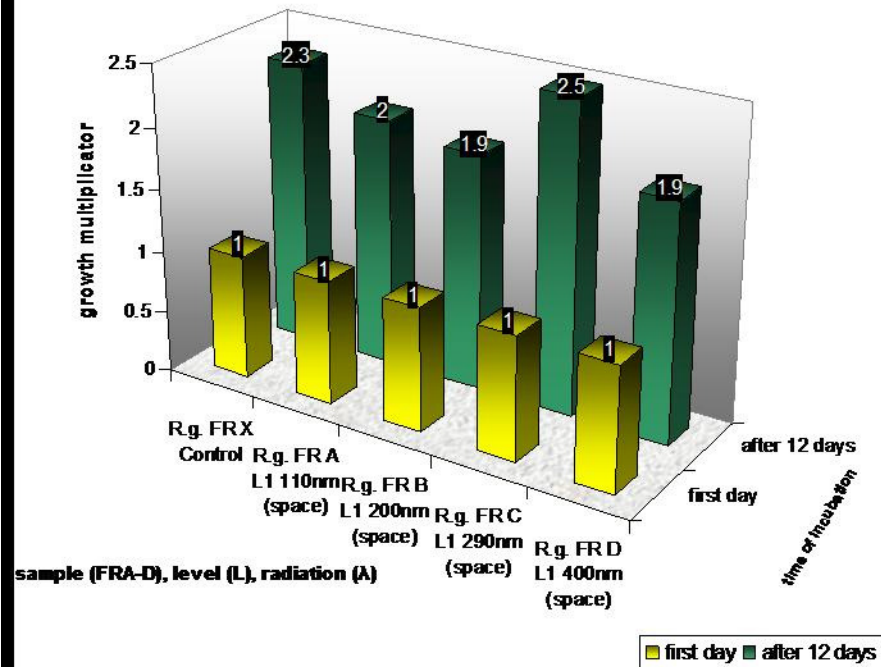
Growth potential of photobiont cells of *R. geographicum*



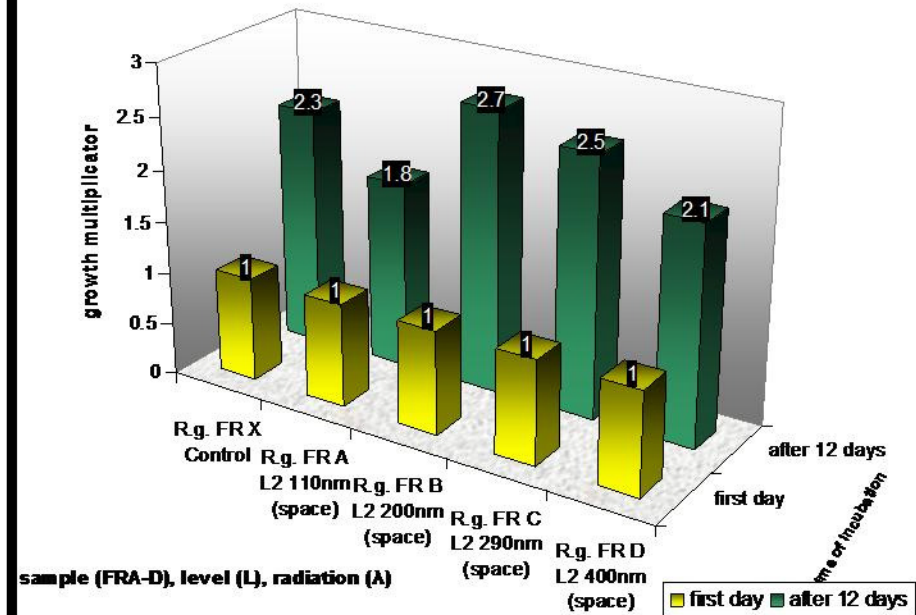
L 1

L 2

Growth capacity of photobiont cells of the exposed lichen
R. geographicum (BIOPAN, L1, space)

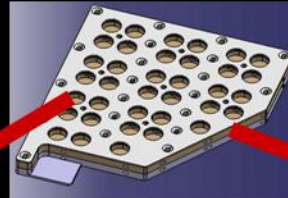
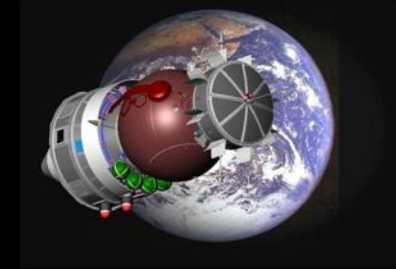


Growth capacity of photobiont cells of the exposed lichen
R. geographicum (BIOPAN, L2, space)



RESULTS:

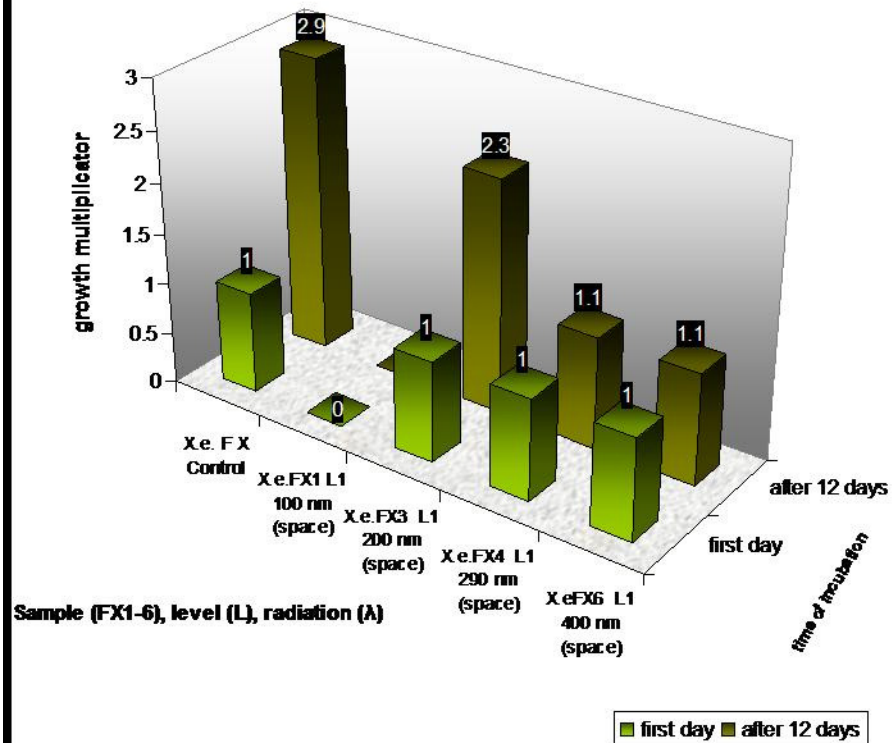
Growth potential of photobiont cells of *X. elegans*



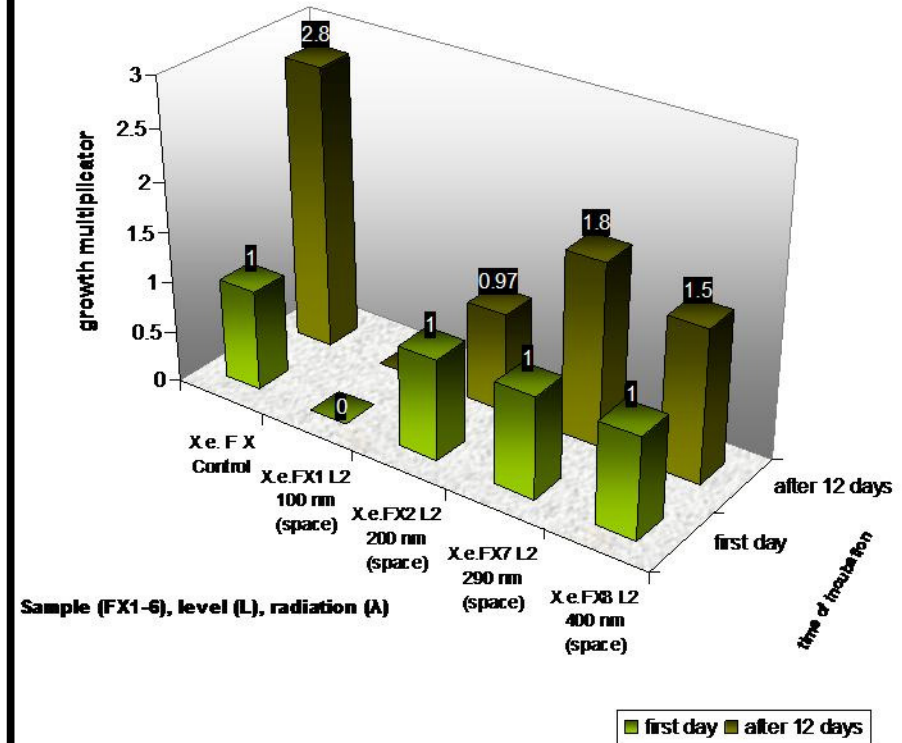
L 1

L 2

Growth capacity of photobiont cells of the exposed lichen *X. elegans*
(BIOPAN, L1, space)



Growth capacity of photobiont cells of the exposed lichen *X. elegans*
(BIOPAN, L2, space)



Preliminary CONCLUSIONS



- In space the mycobiont of *X. elegans* is more resistant than the mycobiont of *R. geographicum*
 - In space the photobiont of *R. geographicum* is more resistant than the photobiont of *X. elegans*
 - The germination capacity of ascospores of both lichens after space exposure is nearly the same compared to the control
 - In case of *R. geographicum* it can be postulated: “Germination induction by vacuum”
- 
- 10 days under Mars conditions are probably without damaging effects on the investigated lichens

Photosynthesis under Martian conditions



**Das Feuchtemesslabor HUMILAB
(Quelle: DLR/Berlin)**



**Die Experimentierkammer (Quelle:
DLR/Berlin)**



Das Gasmischsystem (Quelle: DLR/Berlin)



Fig. 1: The MARS-simulation humidity-lab "HUMILAB" (DLR/Berlin)



Fig. 2: The Mars-simulation chamber (DLR/Berlin)



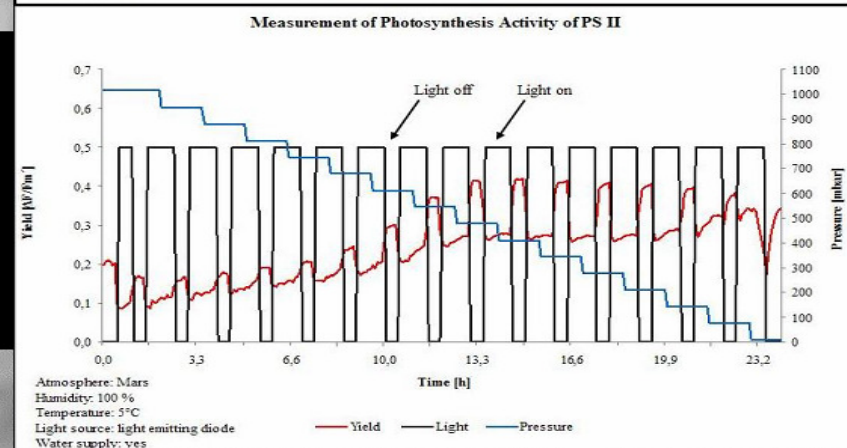
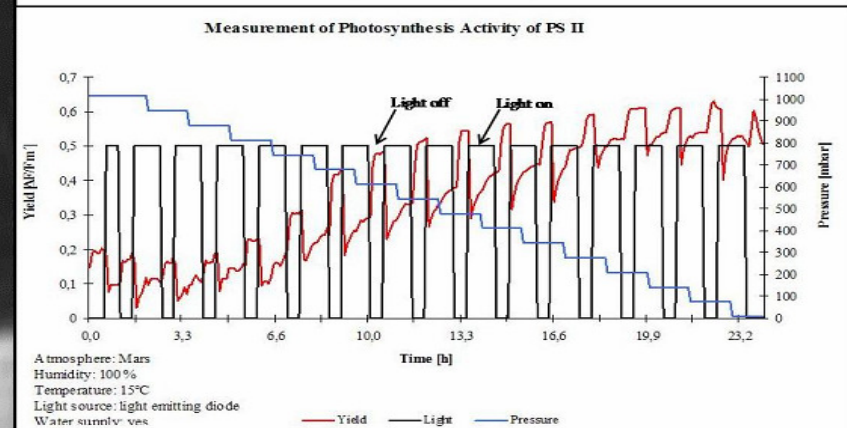
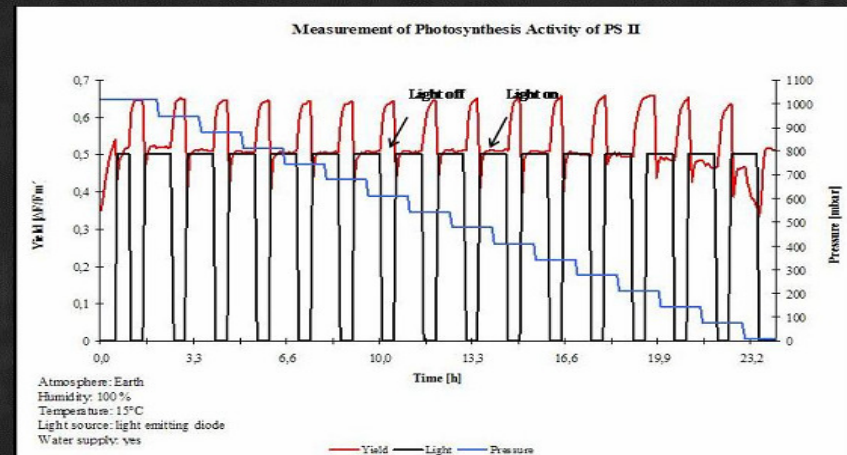
Fig. 3: The gas-mixing system for creation of a Martian atmosphere (DLR/Berlin)

Fig. 4: Photosynthetic activity in Earth atmosphere with decreasing pressures down to 6 mbar (Mars like pressure = blue line). The photosynthetic activity is represented by the red line (Yield value). No remarkable influence occurs.

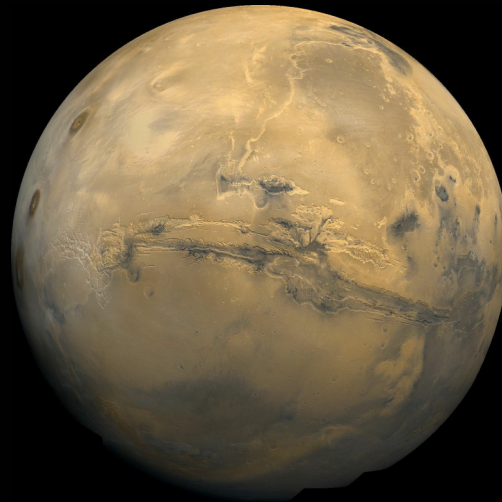
Fig. 5: The Photosynthetic activity in a Mars like atmosphere (95 % CO₂) with decreasing pressures down to 6 mbar at 15°C. The photosynthetic activity (Yield value) is increasing with decreasing pressure and reaches higher levels as at 5°C what is similar to Earth conditions.

Fig. 6: The Photosynthetic activity in a Mars like atmosphere (95 % CO₂) with decreasing pressures down to 6 mbar at 5°C. The photosynthetic activity (Yield value, red curve) is increasing with decreasing pressure.

RESULTS

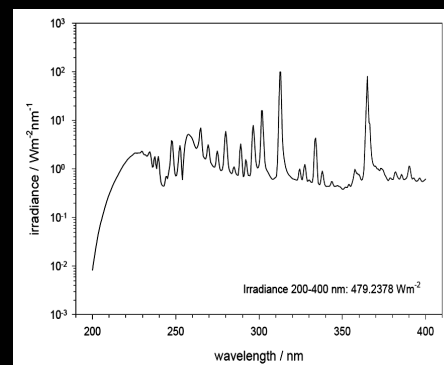
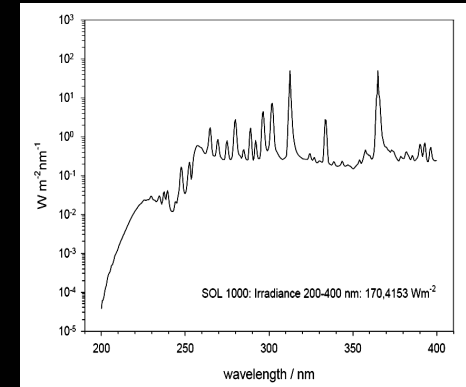
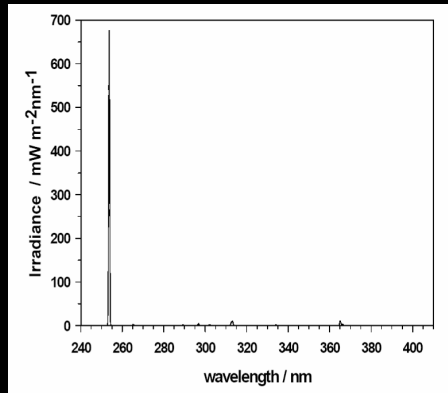


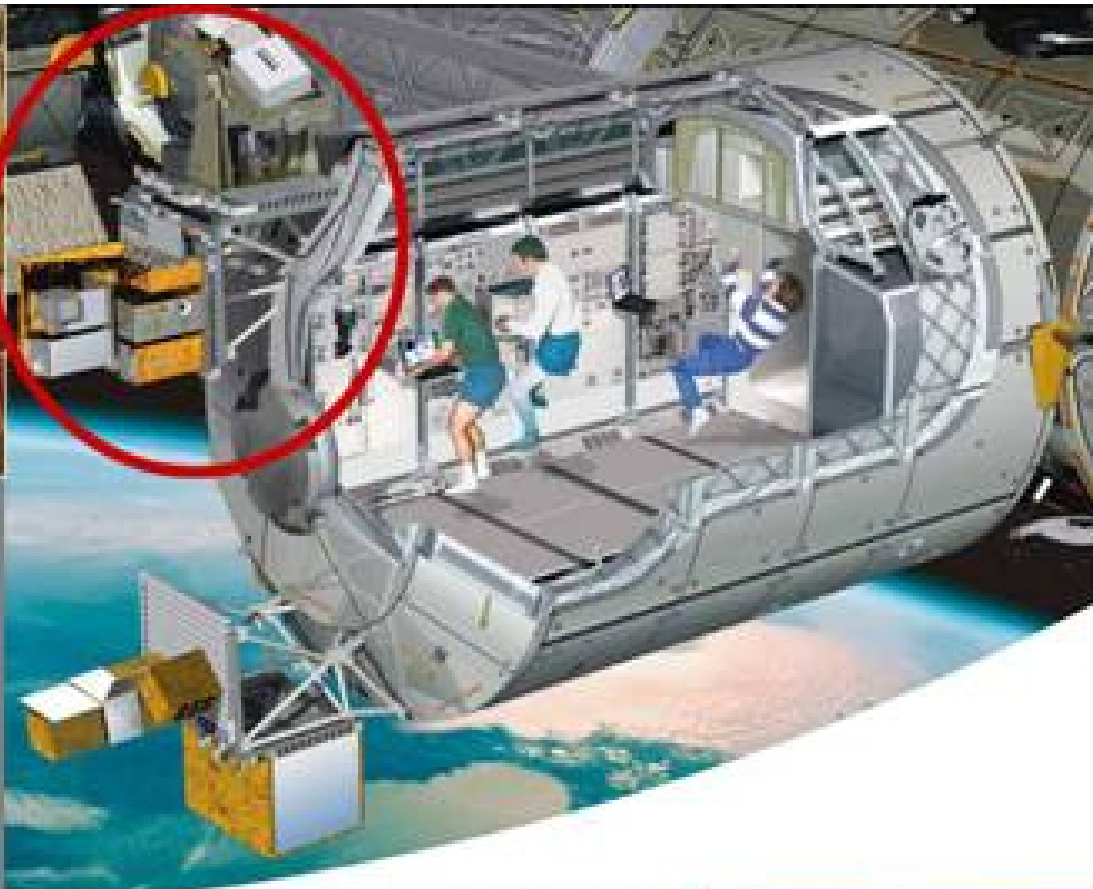
Low Pressure and the high CO₂ concentration seem not to be limiting factors for photosynthesis on Mars



Low temperatures and high UV radiation may disturb photosynthetic activity much more!!!

Preparation for EXPOSE on the ISS and Outlook





Expose-EuTEF: Flugmodell
Expose-EuTEF: Flight Model



Expose-EuTEF: Behälter für die Proben
Expose-EuTEF: Sample Containers

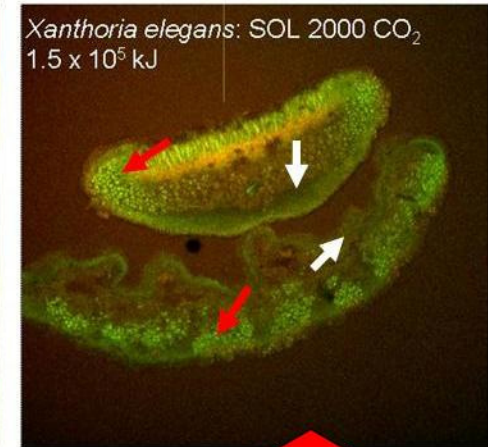
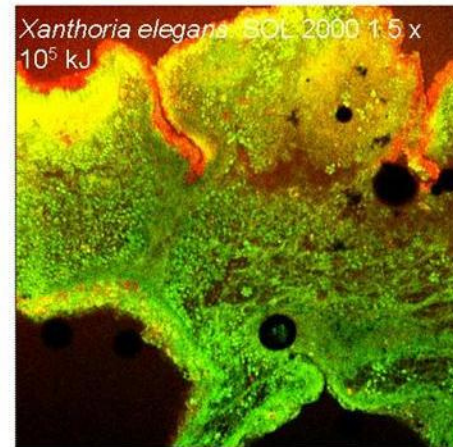
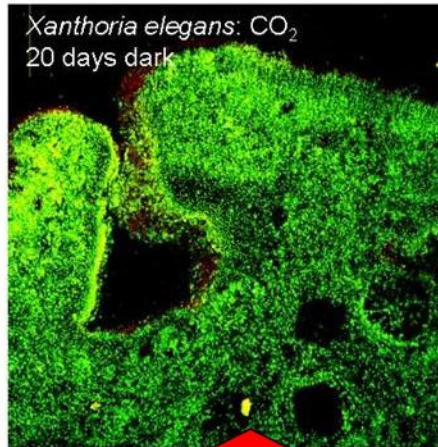
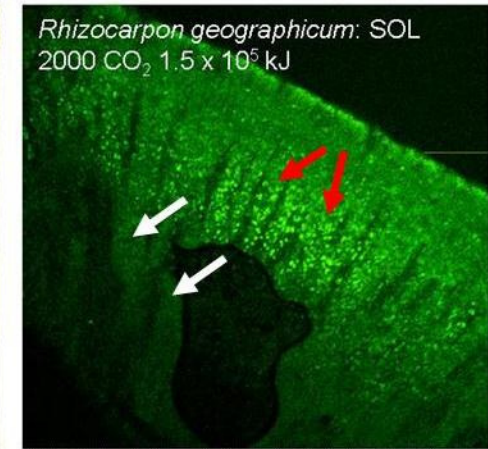
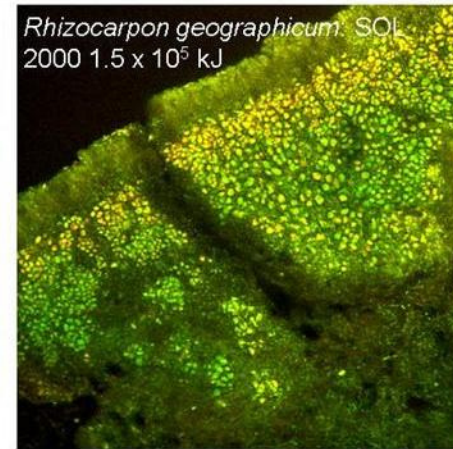
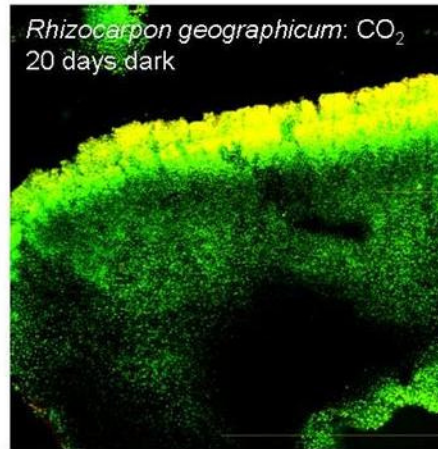
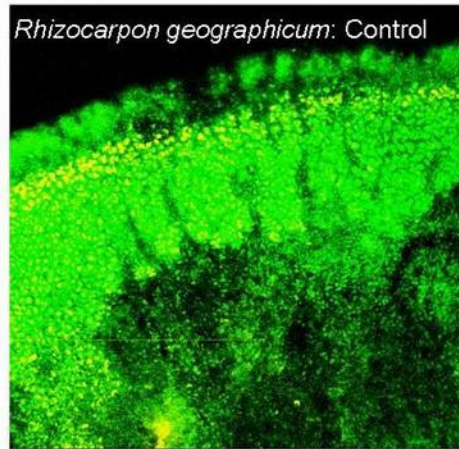


Proben

EVT: Selected and combined space simulation parameters

EVT	parameter	duration/exposition		no. samples
E1				
	Vacuum 10^{-5} Pa	1 h	1.3×10^{-5} Pa	3
		1 wk	2.3×10^{-6} Pa	3
	Temperature oscillation 50 cycles -20 °C to +20 °C, 1 atm air	2 wk		3
	UV-C irradiation monochromatic 254 nm, 1 atm air, 71.4 $\mu\text{W}/\text{cm}^2$	14 s	10 Jm^{-2}	3
		2 min 20 s	100 Jm^{-2}	3
		23 min 20 s	1000 Jm^{-2}	3
	UV irradiation polychromatic 200-400 nm, 1 atm air	3 s (SOL2000)	$1,44 \text{ kJm}^{-2}$	3
		52 min (SOL2000)	$1.5 \times 10^3 \text{ kJm}^{-2}$	3
		87 h (SOL2000)	$1.5 \times 10^5 \text{ kJm}^{-2}$	3
	total number of samples EVT-E1			27
E2	parameter	duration	exposition	no. samples
	Vacuum 10^{-5} Pa (Dark)	22 days		3
	Vacuum 10^{-5} Pa + UV irradiation polychromatic (200-400 nm)	22 days 244.5 h (SOL1000)	$1.5 \times 10^5 \text{ kJm}^{-2}$	3
	Mars atmosphere 600 Pa (Dark)	21 days		3
	Simulated CO ₂ Mars atmosphere 600 Pa + UV irradiation polychromatic (200-400 nm)	21 days 18 min (SOL2000) + 10 dd 3 h 40 min 48 s (SOL1000)	$1.5 \times 10^5 \text{ kJm}^{-2}$	3
	total number of samples EVT-E2			12
Control				
	Room temperature, dark, 1 atm air	2 months		3

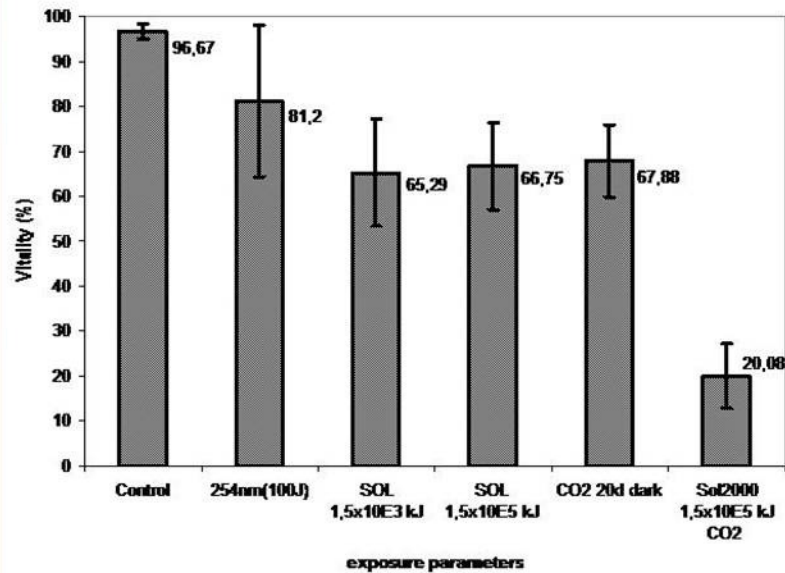
Results of EVT experiments



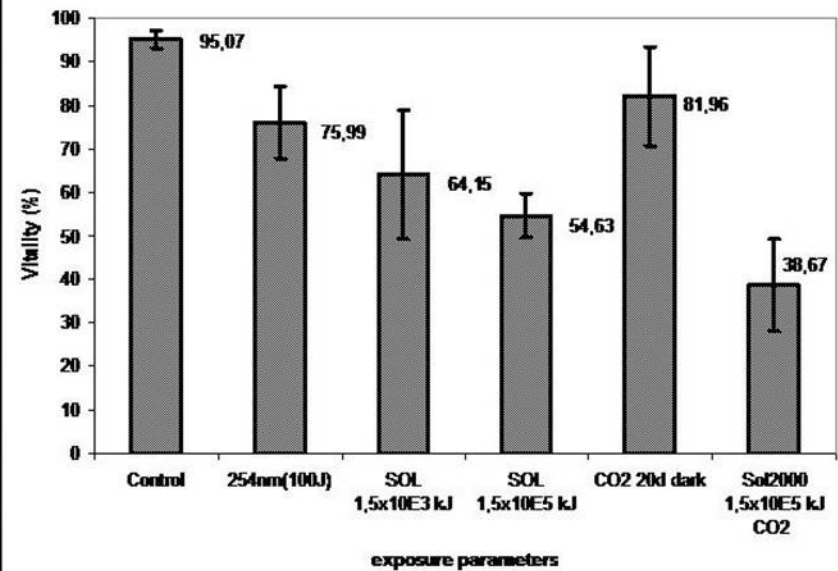
CO₂-Effects: More vitality reduction effects on the fungal part (mycobiont) in the lichen of *R. geographicum* than in the lichen of *X. elegans*. The alga (photobiont) is less influenced / green color = vital cells. Yellow, red and orange color = physiological activity

Radiation and CO₂ effects: in both lichens the mycobiont is more influenced (white arrow). The photobiont survived better (red arrow) due to protecting surrounding fungal hyphae and secondary lichen metabolites.

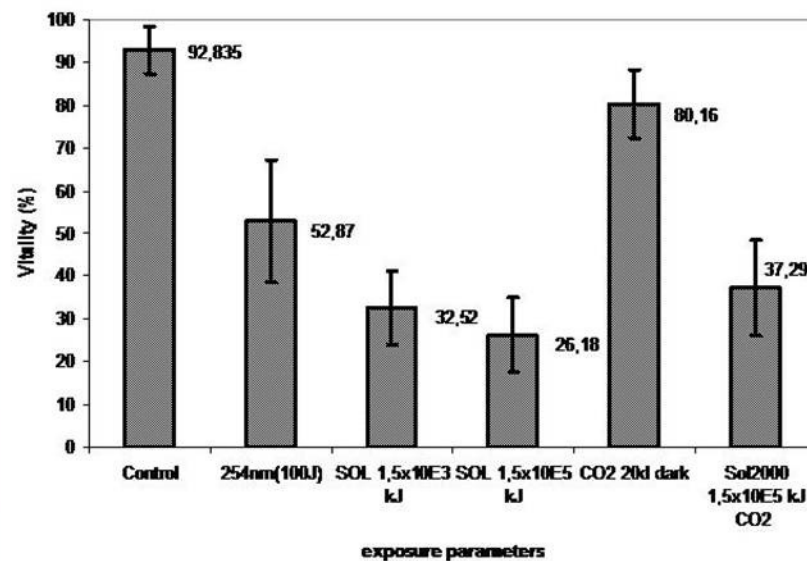
Viability of thalli and apothecia of *Rhizocarpon geographicum* after EVT 1 (space parameter exposure)



Viability of thalli and apothecia of *Xanthoria elegans* after EVT 1 (space parameter exposure)



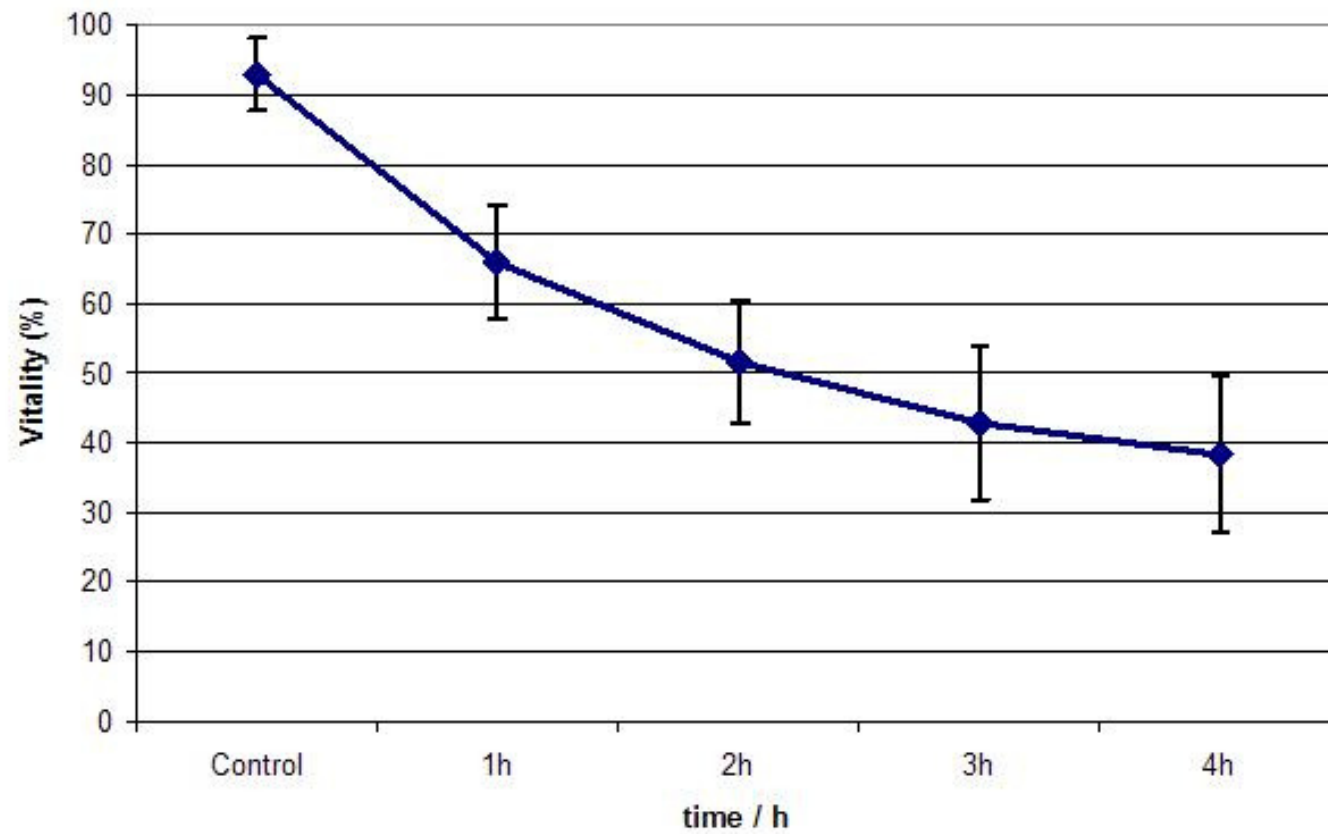
Viability of mycobiont of *Xanthoria elegans* after EVT 1 (space parameter exposure)



During the simulation tests no cultures of the mycobiont of *R. geographicum* were available

Temperature resistance

Vitality of *Xanthoria elegans* at $t = 80^{\circ}\text{C}$ with different exposure times

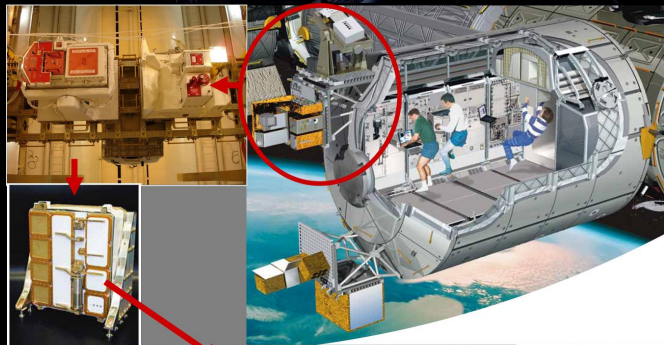


Additional check

DNA damage of the lichen *X. elegans* and its symbionts induced by monochromatic UV radiation

	Proben-Nr.	Probenart	λ (nm)	Parameter		Photoprodukte			DNA
				Dosis (kJm ⁻²)	t _e (h)	Cyclo-TT	(6-4) TT	Spore TT	Masse (μg)
photobiont	1	P ^C	0	0	0	0	0	0	0,019
	2	P	254	5,04	2	31,7	8,2	7,5	0,007
	3	P		10,08	4	23,9	11,7	10,6	0,011
	4	P		20,16	8	63,5	13,5	13,2	0,006
mycobiont	5	M ^C	0	0	0	0	0	0	0,007
	6	M	254	5,04	2	0	0	0	0,018
	7	M		10,08	4	0	0	0	0,012
	8	M		20,16	8	0	0	0	0,013
lichen	9	Xe ^C	0	0	0	0	0	0	0,054
	10	Xe	254	5,04	2	0	0,5	0	0,027
	11	Xe		10,08	4	2,3	1,2	2,6	0,054
	12	Xe		20,16	8	1,4	3,3	4,5	0,037
	19	XeApo	0	0	0	0	0	0	0,017
	20	XeApo	254	5,04	2	0	0	0	0,046
	21	XeApo		10,08	4	0	0	0	0,016

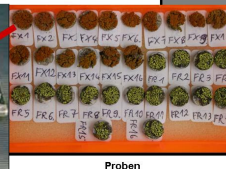
Other experiments are further going on On EXPOSE!!!



Expose-EUTEF: Flugmodell
Expose-EUTEF: Flight Model



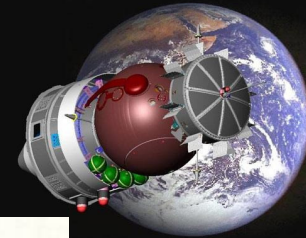
Expose-EUTEF: Behälter für die Proben
Expose-EUTEF: Sample Containers



Proben



PUBLICITY AND OUTREACH



B4 DÜSSELDORF

RHEINISCHE POST MITTWOCH 5. DEZEMBER 2007

Der Countdown läuft

Wenn am Donnerstag die **Raumfähre Atlantis** zur internationalen Raumstation ISS startet, drücken mindestens zwei Wissenschaftler der **Heinrich-Heine-Universität** fest die Daumen: **Sieglinde Ott** und **Jean-Pierre de Vera** vom Botanischen Institut schicken Proben mit ins All.

VON DIRKE KÖPP

Eigentlich könnten Sieglinde Ott und Jean-Pierre de Vera ganz ruhig sein: Die Crew für den Flug der Atlantis-Raumfähre am Donnerstag steht schon fest, und das „Go“ für den Start hat die Nasa auch bereits gegeben. „Aber wir sind trotzdem nervös“, gesteht Sieglinde Ott, Professorin am Botanischen Institut der Uni, und ihr Kollege, der Wissenschaftler Jean-Pierre de Vera, fügt hinzu: „Man fragt sich immer: Klappt alles?“ Das Interesse der Düsseldorfer Forscher an der Mission der Nasa ist nicht ganz uneigennützig: Wenn das Spaceshuttle Atlantis am Donnerstag, 16.11 Uhr amerikanischer Zeit, vom Welt-

„Flechten sind höhere Organismen als Bakterien. Dass sie im All überleben, ist ein gutes Zeichen“

raumbahnhof Cape Kennedy in Florida zur internationalen Raumstation ISS startet, haben die Astronauten auch „Geplack“ der Heine-Universität an Bord: Proben von Flechten, einem Symbiose-Organismus aus Alge und Pilz.

Schon im September hatte die Arbeitsgruppe um Ott und de Vera zusammen mit spanischen Forschern unter Leitung von Rosa de la Torre Noetzel zwei Kapseln mit Flechten ins All geschickt. Sie wollten wissen, wie die Organismen auf die weite Reise unter extremen Bedingungen reagieren. Tage lang kreisten Sojas und Flechten in knapp 300 Kilometern Höhe im All. Partner für das Millionenprojekt zur Grundlagenforschung war damals und ist diesmal das Deutsche Zentrum für Luft- und Raumfahrt in Köln (DLR).

Das aktuelle Experiment ist für die Wissenschaftler noch spektakulärer: Diesmal bleiben die Proben mehr als zwei Jahre im All und werden zudem außen an der Raumstation „angebracht“, beschreibt de Vera. Dafür müssen zwei Astronau-



Wenn die **Atlantis-Spaceshuttle** am Donnerstag gegen Raumstation ISS startet, hat es auch die Proben der Heine-Uni an Bord.

ten aussteigen und sie dort befestigen. Die Flechten sind in einer speziellen Box, dank derer sie zeitweilig vor der allzu intensiven Sonneneinstrahlung geschützt werden können. „Es ist wichtig, dass dieser Mechanismus funktioniert“, so Ott.

Mit ihrem Experiment wollen die Wissenschaftler zu denen außer den spanischen Kollegen auch Silvio Jorda in Vitoria (Italien) und Elke Gahbauer vom DLR zählen, beiraten jedoch, wie resistent die Flechten gegen die extrem lebensfeindlichen Umwelt des Weltraums sind und was sich daraus für die Forschung ableiten lässt. Etwa könne man das Wissen über die Struktur resistenter Flechten bei der Entwicklung von Strahlenschutz-Kleidung oder – aber erst in ferner Zukunft – wie Ott und de Vera betonen – in der Krebsforschung nutzen.

Erste Ergebnisse aus Untersuchungen der Flechten, die im September ins All waren, lassen hoffen: „Mehr als 90 Prozent der Sporen (so etwas wie Samen) keimten und wuchsen“, sagt de Vera. Auch habe man nachweisen können, dass der Stoffwechsel der Zellen intakt sei und die Algen Photosynthese betreiben. „Nun müssen wir die Proben weiter beobachten“, erklärt Ott. „Hören sie auf zu wachsen? Verhalten sie sich anders als Flechten, die nicht im All waren? Zudem waren die Proben verschiedenen



Die Wissenschaftler **Jean-Pierre de Vera** und **Sieglinde Ott** müssen sich rund zwei Jahre gedulden, bis die Flechten aus dem All zurück sind.



Mikroskopbilder eines Querschnitts der Flechte **Xanthoria elegans**. Links (vorher) sind lebende Zellen grün/rot gefärbt, rechts (nachher) sind sie noch lebendig.

INFO

Langzeitexperiment

Der Langzeitversuch mit den Flechten an der internationalen Raumstation ISS wird mehr als zwei Jahre dauern. Bei der erweiterten Weltraumuntersuchung mit Namen „LIFE“ (Lichen and Function Experiment) werden außer den Flechten auch im Gestein lebende Organismen wie **Cyanobakterien** (Blaualgen) und antarktische **Mikropilze** studiert.

Dosen von UV-Licht ausgesetzt – auch die Reaktion darauf prüfen wir.“ Insgesamt sind die Forscher sehr zufrieden: „Flechten sind höhere Organismen als Bakterien. Dass sie die Weltraumbedingungen überleben, ist ein gutes Zeichen.“

SERVICE

„Saufen ist uncool“: Schüler zeigen ab heute Plakate

(gök) Es ist eine besondere Ausstellung, die Bürgermeister Dirk Ebers heute um 11 Uhr zusammen mit Michael Baites, Leiter des Ursulinen-Gymnasiums, im Weiterbildungszentrum eröffnet: „Saufen ist uncool“ lautet der Titel der Schau, die die Schüler der Klasse 9e des St. Ursula-Gymnasiums konzipiert und umgesetzt haben. Zu sehen sind 14 Plakate, die sich mit Alkoholkonsum von Jugendlichen beschäftigen. Ausgangspunkt der Klasse waren Sprüche wie „Komusaufer“ und „Dicht ist Pflicht“, mit denen sie sich im Biologie-Unterricht auseinandergesetzt haben.

Bei der Umsetzung der Ergebnisse haben Kunstlehrerin Nicole Busch und Design-Professor Wilfried Korfmaier von der FH geholfen.

WBZ, Bertha-von-Suttner-Platz, Foyer. Zu sehen ist die Schau bis 19. Dezember.

Kneipp-Verein wandert im Dezember

(lani) An drei Tagen im Dezember lädt der Kneipp-Verein Düsseldorf zum Wandern ein. Heute wird ein Spaziergang am Rundwanderweg angeboten. Treffpunkt ist um 14.30 Uhr an der Haltestelle der Straßenbahnlinie 703 auf der Burgmühlstraße. Für Samstag, 8. Dezember, ist ein Ausflug zum Rosenhof geplant. Treffpunkt hierfür ist die Bushaltestelle der Linie 730 am Knappertbrück. Für den letzten Tag, Samstag, 15. Dezember, ist ab 13 Uhr eine Fahrt nach Lörick geplant.

Kontakt: Telefon 21.92.44 oder E-Mail: Kneipp-Verein.Düsseldorf@t-online.de.

Tipps für Umgang mit Demenzerkrankten

(RP) Zu einem praxisnahen Vortrag zum Umgang und zur Kommunikation mit demenziell Erkrankten lädt der Caritasverband für heute um 17.15 Uhr in den Caritas-Treffpunkt ein. Referent Andreas Kutsche möchte auch schwierige Themen und Situationen im Umgang mit Demenzerkrankten respektvoll, anschaulich und mit Humor darstellen.

Caritas-Treffpunkt Oststraße 64

Über Hiroshima und



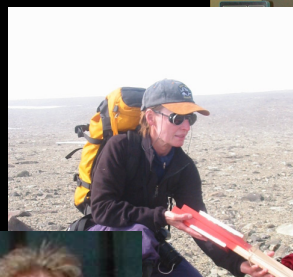
International Cooperation



J.P.de Vera; C. Cockell; C. Meyer; G. Horneck

R. Möller

S. Ott



P. Rettberg

U. Hornemann



P. Inderbilen (Assistent, Walliser Hof, Zermatt)



M. Melles

TNT-experts



Technicians of DLR/Cologne



S. Onofri



D. Stöffler

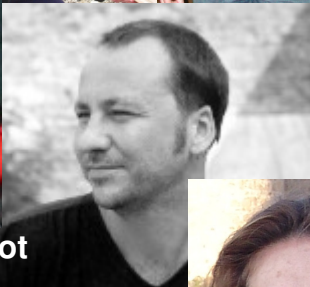


J. Fritz



R. de la Torre Noetzel

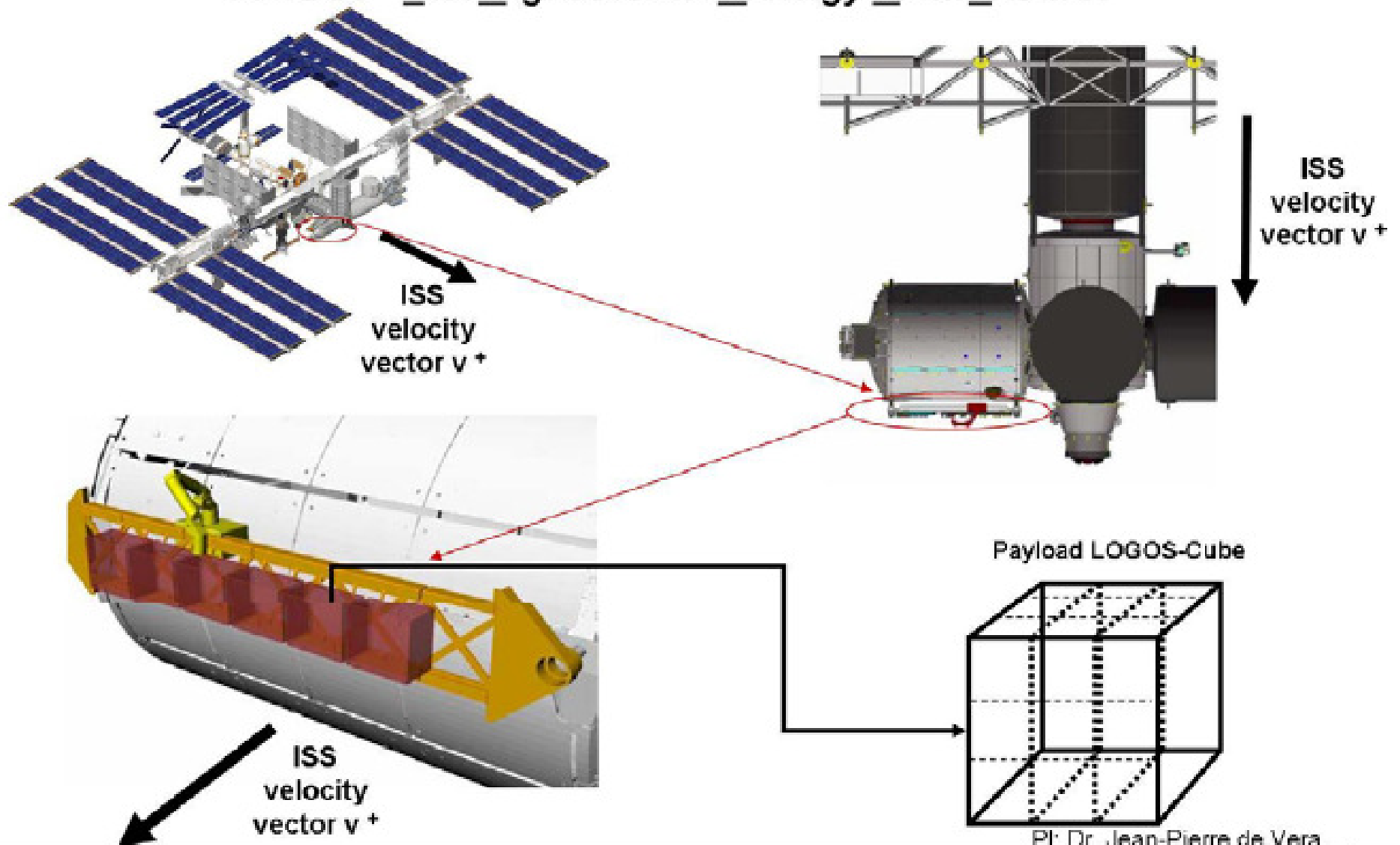
S. Dreesen, Pilot (Air Zermatt)



Proposed facility / Payload:

LOGOS-CUBE and Experiment:

LOGOS – Life-Organics and Geology Orbit Science



Proposed facility / Payload:
LOGOS-CUBE and Experiment:

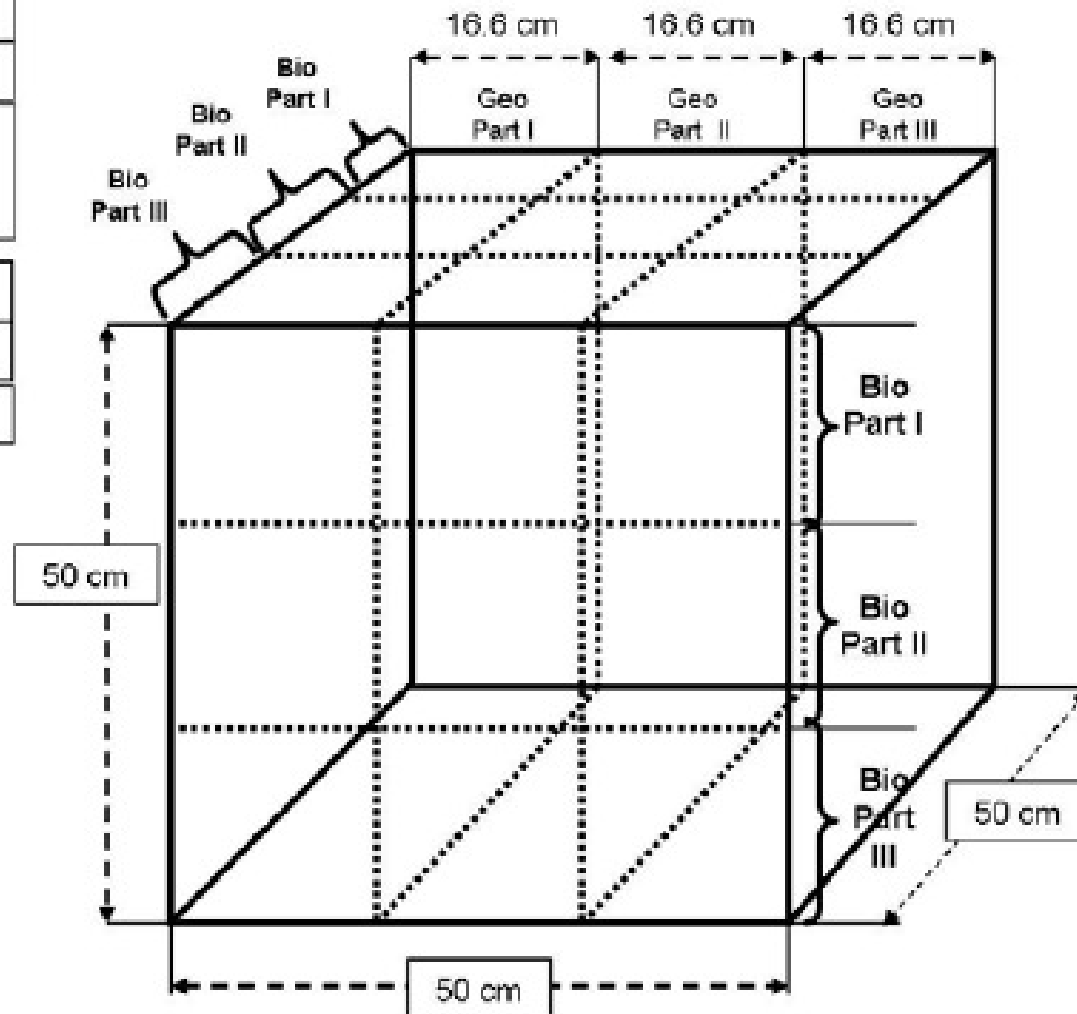
LOGOS – Life-Organics and Geology Orbit Science

Bio Part I: Epilithic organisms on rock
Bio Part II: Endolithic organisms in rocks
Bio Part III: Clean rock samples with integrated dosimeter, temp.instr., pressure instr., fluorescence instruments
Geo Part I: sandstone
Geo Part II: Dunit
Bio Part III: Gabbro

A sophisticated autonomic payload facility for scientific tests: space weathering on organisms and rock and degradation or resistance of organics produced by life forms. Results will be essential for definition of real life markers/bio signatures as reference to the search of life in the universe.

Realisation of a space performed bio signature data bank (BSDB)

**ISS
velocity
vector $v +$**

PI: Dr. Jean-Pierre de Vera,

Proposed facility / Payload:

LOGOS-CUBE and Experiment:

LOGOS – Life-Organics and Geology Orbit Science

Option: windows of cube may serve additionally as solar panel

Window I = solar panel I

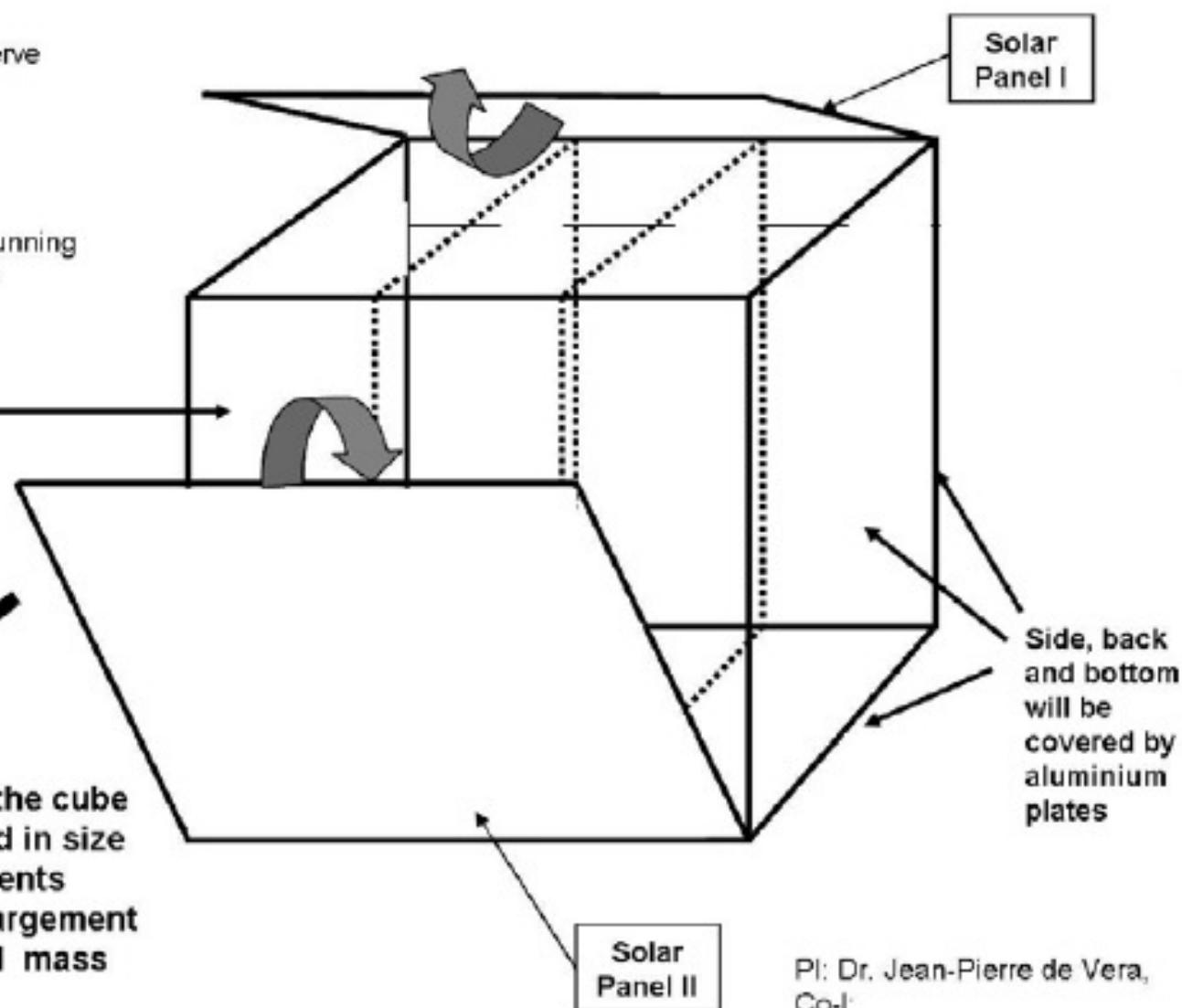
Window II = solar panel II

External energy source for the running measuring instruments and data transfer devices

Inside, bottom: logger, memory device, dark control samples

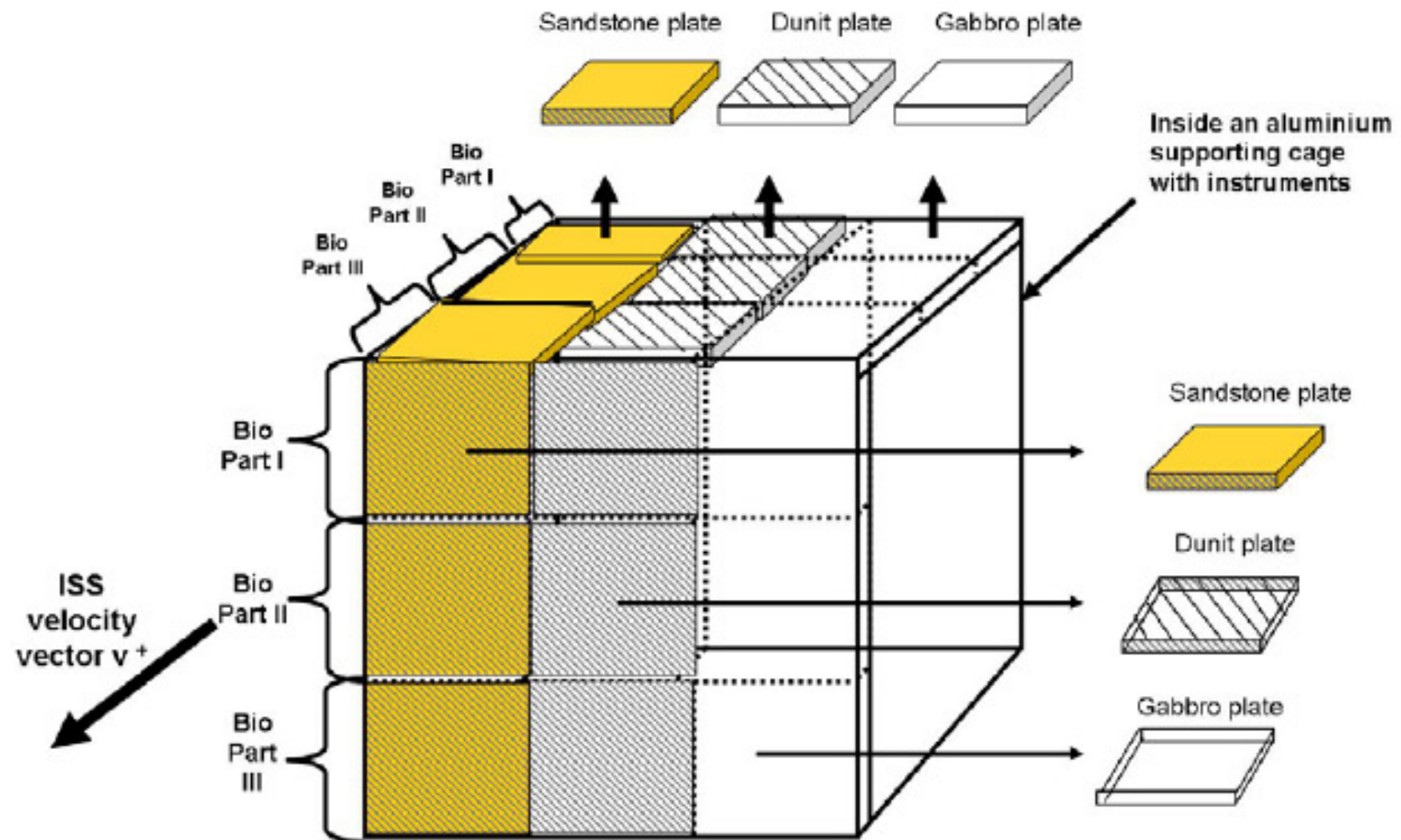
ISS
velocity
vector $v +$

Solar panels are optional and the cube has the flexibility to be adapted in size to different scientific requirements (flexibility for reduction or enlargement of the payload with maintained mass and low power requirements)



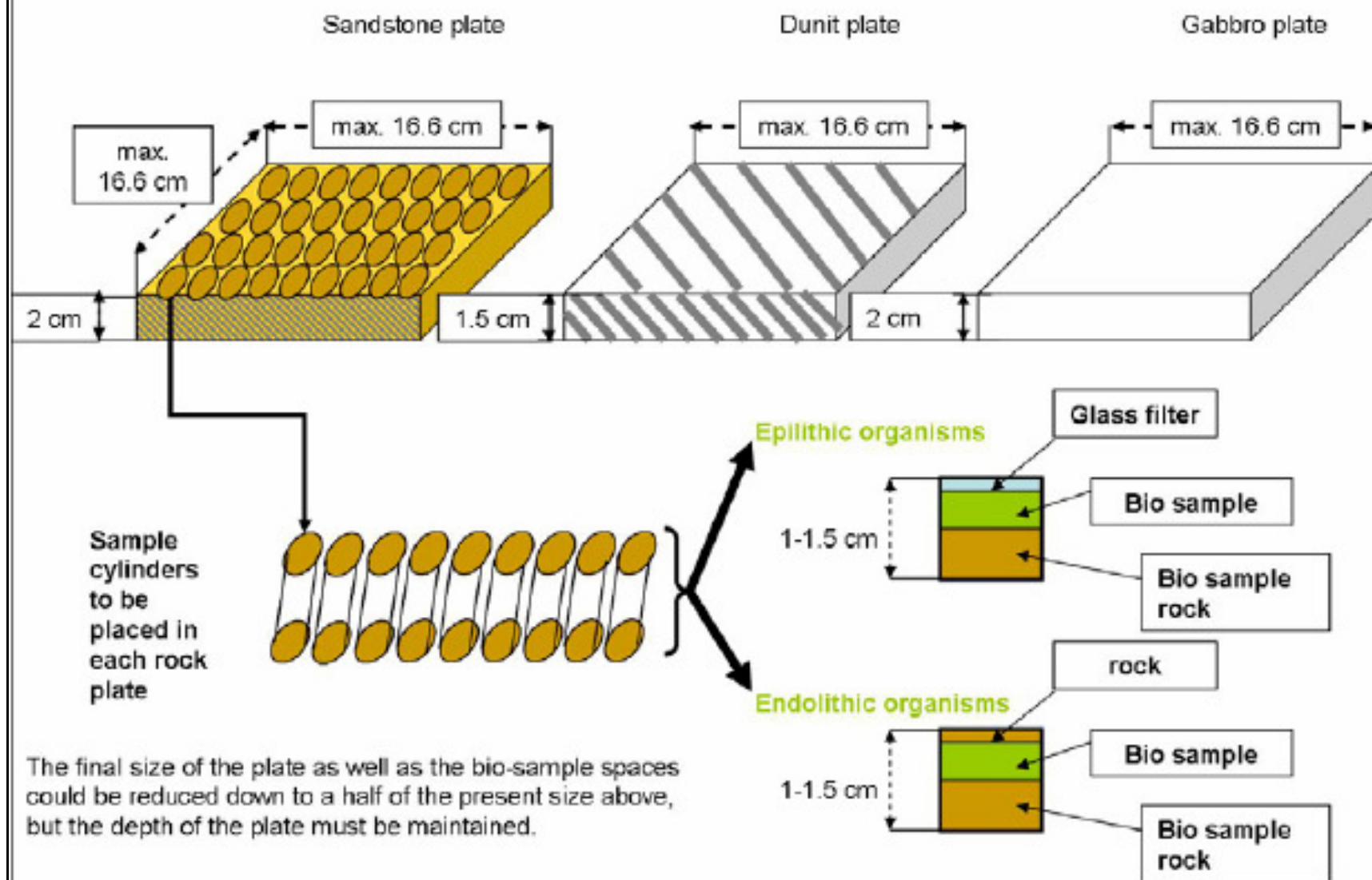
PI: Dr. Jean-Pierre de Vera,
Co-I:

Payload: LOGOS-CUBE Hardware Elements top and front surface



Experiment: LOGOS – Life-Organics and Geology Orbit Science

on Payload: LOGOS-CUBE-Hardware Plates for Biology and Geology



Proposed Bio-Samples

List of proposed organisms:

1. **Cyanobacteria (Chlorophyll)**
2. **Xanthoria elegans (Chlorophyll, Carotene, Parietin, Emodine, Chitin...)**
3. **Rhizocarpon geographicum (Chlorophyll, Melanin, Usnic acid, Chitin...)**
4. **Bacteria (*B. subtilis*)**
5. **Fungi (Melanin, Anthraquinones, Chitin, ...)**
6. **Seeds (flavonoids, plastids...)**

List of proposed isolates:

1. **Chlorophyll a**
2. **Chlorophyll b**
3. **Carotene**
4. **Anthraquinones as Parietin**
5. **Chitin**
6. **Flavonoids**
7. **Plastids**

The list could be enlarged, reduced or some of proposed isolates exchanged according to the requirements of each team member and ESA guide lines

