

### **2022 MELISSA CONFERENCE** 8-9-10 NOVEMBER 2022



OLE project: a hydroponic greenhouse demonstrator for fresh food production in space

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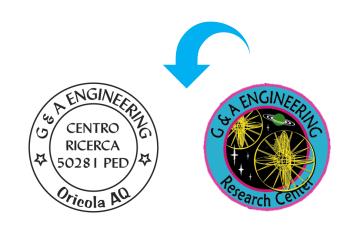
## **COMPANY HISTORY**



**1996** 

**G & A ENGINEERING** 

Foundation 2 DEPT





PRIVATE RESEARCH CENTER
For Microelectronic for Space Application

INNOVATIVE SME
Defence and Aerospace





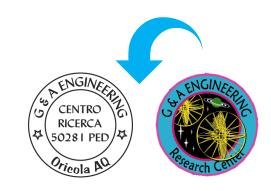
### **COMPANY HISTORY**



# May 2022







PRIVATE RESEARCH CENTER For Microelectronic for **Space Application** 



**INDUSTRY** Defence, Aerospace, Hydroponic, Automotive, Medical



## THE COMPANY



**18.000** SQM

**5**WORKSHOPS



ISO 9001
ISO 14001
BUREAU VERITAS
Certification





## THE COMPANY



A personnel focused company having a mix of skills, high tech internal technologies and industrial processes

**14** 

**2.5** 

**70%** 

**M€**TURNOVER













2

Research Projects
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An entire LOB dedicated to create a line of hydroponic products.

- ✓ RobotLamp, the best performing LED lamps on the market
- ✓ RobotFarm, the hydroponic greenhouse in the size of a washing machine.
- ✓ Hydrowall, the hydroponic vertical farm in the size of a planter
- ✓ MIG, the vertical farm in container ISO20 for extreme applications
- ✓ **SOLE**, the hydroponic vertical farm designed to cultivate in Space
- A Patent for Electronic Cultivation Recipe for CEA systems

FARMING HAS NEVER BEEN EASIER



## THE « SOLE » PROJECT





A plant greenhouse demonstrator for the soilless cultivation of plants, based on solid state artificial lighting (LED).

The project studied the best light recipes for duration, intensity and spectral quality, optimized to achieve the different phases of plant growth.







The goal is to use this demonstrator to support research for future manned space missions, to carefully evaluate the resources needed to produce adequate quantities of fresh food, reducing the astronaut's operating times.



### **BACKGROUND**





Future human long-term space exploration demands for fresh food production during missions, independently on periodic supplies from Earth.

The possibility to grow plants in space positively impacts both astronaut diet and psychological wellbeing.

It has been estimated that a facility of a few cubic meters is sufficient to supply the diet of 4 to 6 crew members with key vitamins and fresh bioactive substances.

Soilless cultivation is the best suited for production of high quality food in space, where resources are limiting factors.

In this context, the project SOLE aimed to realize a hydroponic cultivation module based on LED lighting systems for growing plants, specifically microgreens, for space applications.



## SOLE DEMONSTRATOR





SOLE is a hydroponic vertical farm built in a closed environment, fully controlled and the light radiation for photosynthesis (PAR) is provided by Proprietary LED lamps.

The cultures are arranged on 5 trays that can be lighted independently, while climate and irrigation parameters are in common.

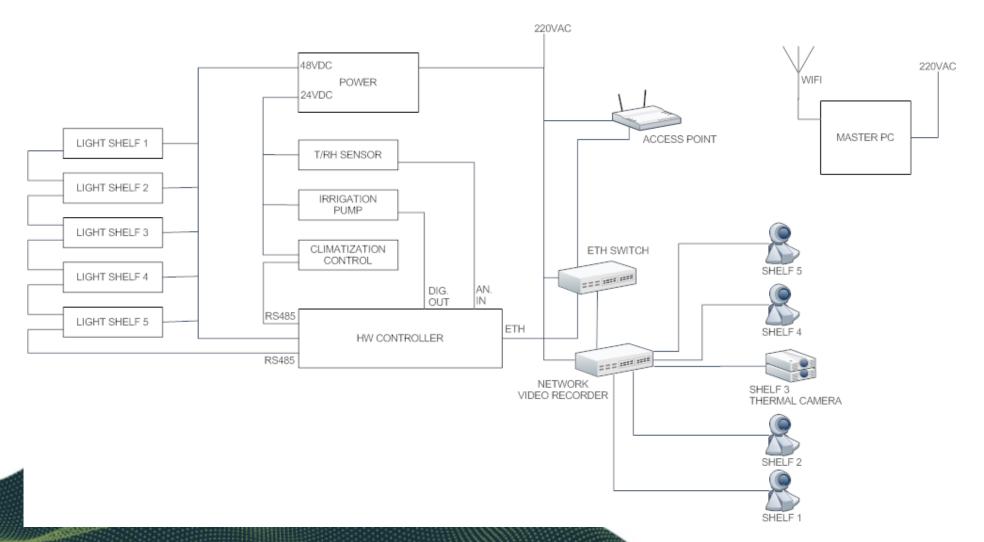
The command and control system is Proprietary and is divided into MASTER and HW CONTROLLER units. The MASTER consists of a PC external to the device, while the HW CONTROLLER consists of a Proprietary board with microprocessor. The two units communicate with a MODBUS protocol over Ethernet through the Wi-Fi network of SOLE. The MASTER has the task of providing the HMI interface, managing the automatic execution of cultivation recipes processes, allowing maintenance operations and storing system operating data, the HW CONTROLLER has the responsibility of managing the hardware by applying the commands from the MASTER.



## **BLOCK DIAGRAM**









## **DEMONSTRATOR DESIGN**





The SOLE demonstrator has been designed for the automatic hydroponic cultivation of microgreens, babyleaf and dwarf species.

It consists of a modular cabinet implemented with cultivation trays, light, HVAC, aeraulic and hydraulic systems.

The cabinet was realized in aluminium alloy, which ensure excellent mechanical properties and low weight. It was insulated to minimize energy dispersal and external environmental influence and was equipped with temperature, pressure and relative humidity sensors, else than real-time video monitoring system.

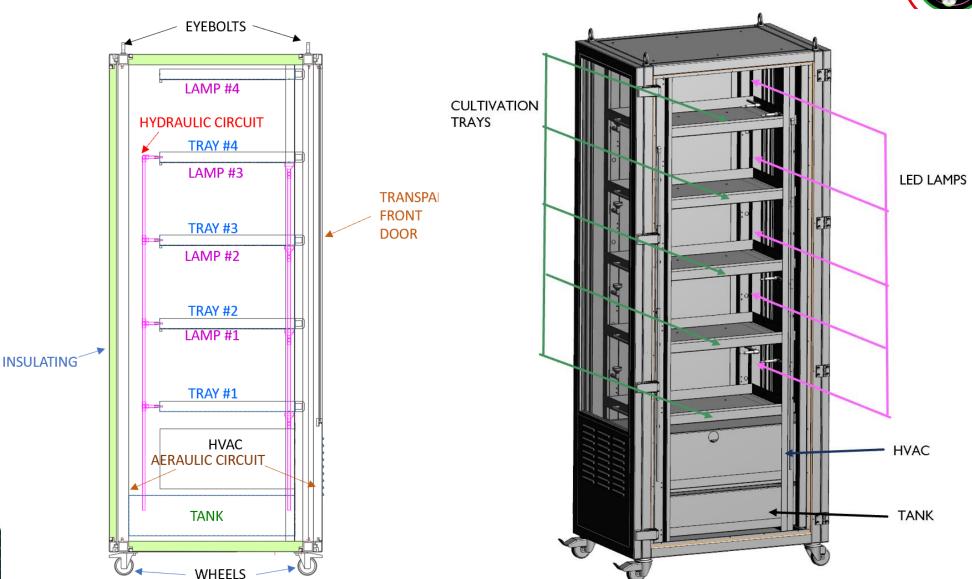
The demonstrator can be operated and controlled remotely using a LAN connection and a simple user-friendly HMI.

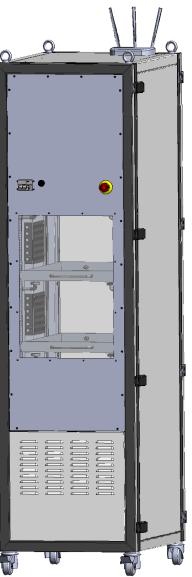


## SOLE DEMONSTRATOR



















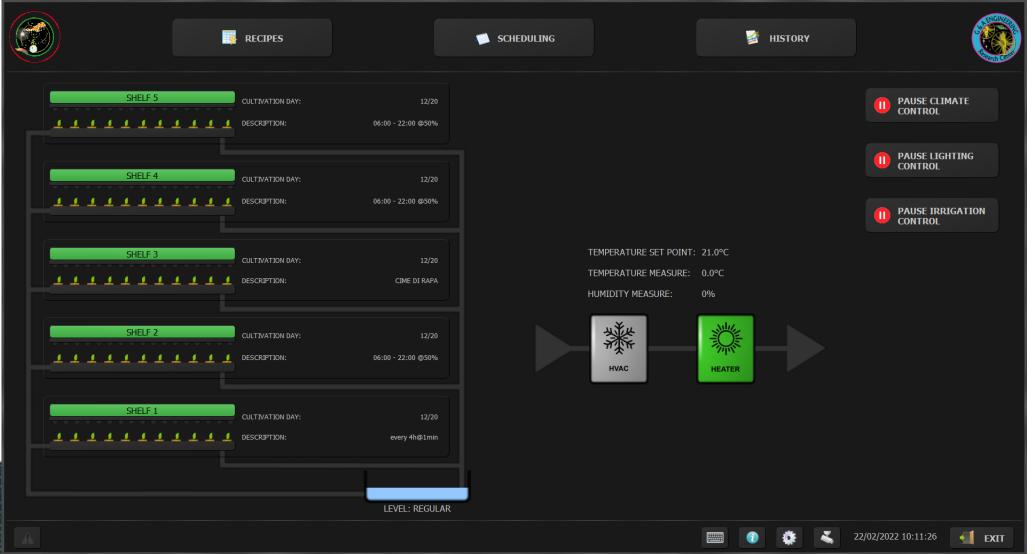




## **SOLE HMI**





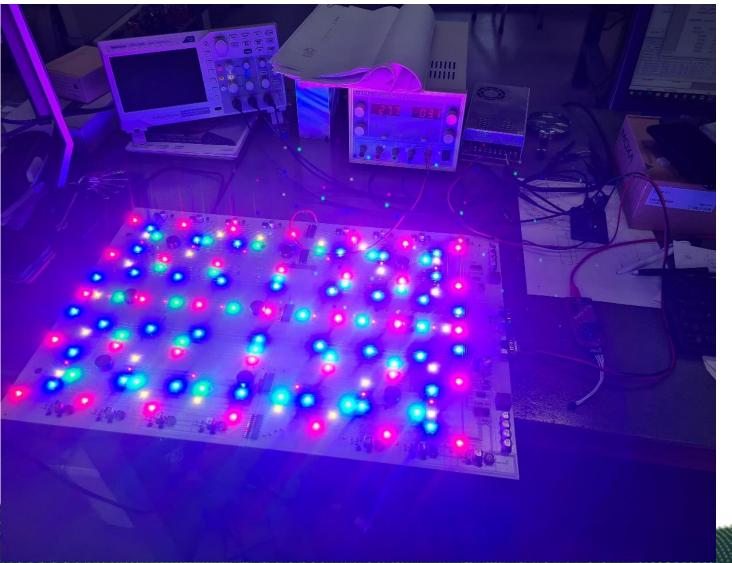




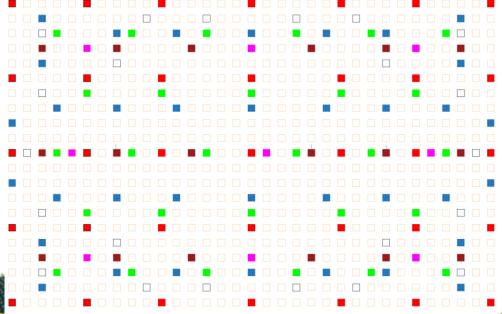
## SOLE LED LIGHTS







COLOR	WAVELENGHT	DOMINANT		
Deep Blue	439 – 461 nm	451 nm		
Far Red	720 – 740 nm	730 nm		
Hyper Red	635 – 666 nm	660 nm		
True Green	513 – 545 nm	521 nm		
UV-A	380 – 440 nm	395 nm		
White	3000K	N/A		





## **CULTIVATION CONDITIONS**





Temperature:

Relative Humidity:

Irrigation Cultivation Recipe:

Recipe Cultivation Lights:

+21°C to +25°C, adjustable

40% to 60%, adjustable

1.30 min every 4 hours, adjustable

18 h Light, 6 h Dark

150 μmoles/m<sup>2</sup>·s

Red 40% - Green 20% - Blu 40%



## **CULTIVATION CONDITIONS**





**Nutritive Solution:** 

**Cultivation Surface:** 

**Seed Germination:** 

**Custom Prepared** 

pH 7

 $68 \times 43 \text{ cm} = 2.924 \text{ cm}^2$ 

48h @ dark, +25°C, 90% RH

In this phase the imbibition of the

seed takes place.

The seed swells and metabolic

processes are activated.



## CLEAN, SANITIZATION & SOWING



Before starting the cultivation it is necessary to carry out a cleaning and sanitizing cycle in order to guarantee the absence of pathogens for cultivation.

The procedure used is summarized below:

- Sterilize the seed in a 2% v/v hydrogen peroxide aqueous solution for 10 minutes
- Rinse in demi-water three times
- Dry the seeds in clan sterile paper
- Clean all cultivation trays with a 2% v/v hydrogen peroxide aqueous solution
- Cut out the Greenfelt mats to match the grow trays dimension
- Sterilize the Greenfelt by spaying hydrogen peroxide 2% v/v for 15 min before sowing
- Sterilize the tank with a 2% v/v hydrogen peroxide aqueous solution
- Prepare the nutrient solution to fill the tank
- Place the Greenfelt mats on the cultivation trays and carry out a watering cycle with the nutrient solution
- Sow











**TURNIP TOPS** 









TRAY DIVISION					
LETTUCE	BROCCOLI				
CABBAGE	RADISH				











TRAY DIVISION				
MUSTARD	BEANS			
LENTILS	ROCKET			









TRAY DIVISION					
CUCUMBER	CHARD				
ROCKET	VALERIAN				









TRAY DIVISION					
CHICKPEAS	ENDIVE				
CHICORY	RED CABBAGE				









TRAY DIVISION						
KOHLRABI	CLOVER					
TURNIP TOP	BRUSSELS SPROUT					









TRAY DIVISION					
RED CABBAGE	RADISH				
BROCCOLI	ROCKET				



## **EXPERIMENTAL TRIALS**





SOLE cultivation experiments were conducted in **ENEA**'s facilities. (SI)





Morphometric (hypocotyl height, cotyledons area, fresh and dry weight), fluorometric (Multiplex index analysis SFR\_R, FLAV, ANTH) and hyperspectral analysis (Specim IQ camera + Evince sw) were conducted on red and green microgreen species: Radish (Raphanus sativus) and Mustard (Brassica juncea).

#### Microgreens hydroponic cultivation

3 replicas/light recipe 20 samples/replica

#### 4 Genotypes

Red and Green Mustard Red and Green Radish

#### 3 Light spectra

 $150 \, \mu mol/m^2 s$ 12h photoperiod L1 60% R, 20% G, 20% B L2 40% R, 20% G, 40% B L3 20% R, 20% G, 60% B



### Non-destructive analysis

## Fluorometric analysis

Multiplex (SFR\_G, FLAV, ANTH RG)

## **Hyperspectral image**

analysis

### Destructive analysis

### **Morphological parameters**

(Plant height, Cotyledon area)

### Biomass production

(wet and dry weight)



## **EXPERIMENTAL TRIALS**





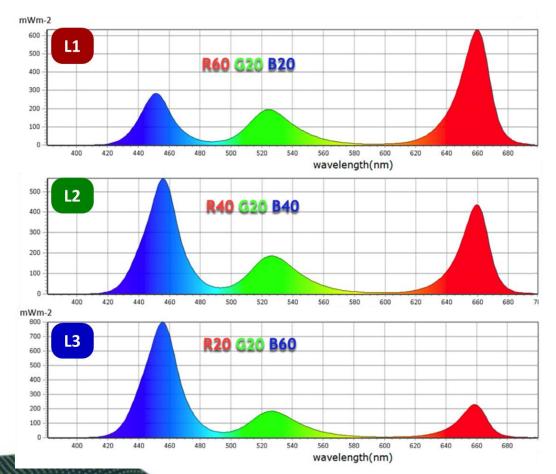




Plants were cultivated with three different light recipes to evaluate the best spectra in terms of primary and secondary metabolism development.

### 3 Light spectra

150 μmol/m²s 12h photoperiod L1 60% R, 20% G, 20% B L2 40% R, 20% G, 40% B L3 20% R, 20% G, 60% B





## HYPERSPECTRAL ANALYSIS





The images were acquired by means of a **Specim IQ** hyperspectral camera, with an acquisition interval

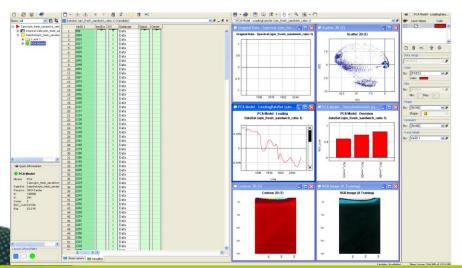
between 400-1000 nm.

The **Evince software** was used for the analysis of the images, the extraction and processing of the data present in the images, which allows to process, analyze and obtain results thanks to the integrated advanced statistical analysis package specific for hyperspectral images





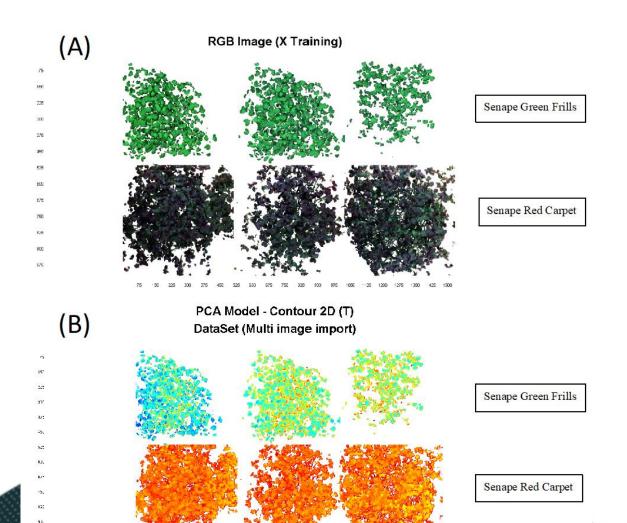






## HYPERSPECTRAL RESULTS



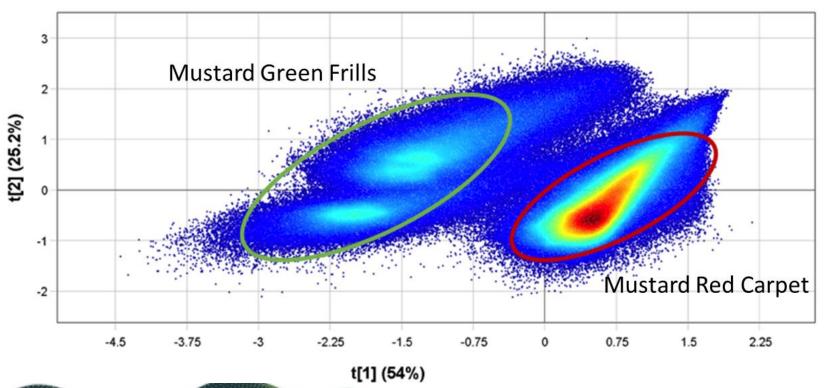


RGB image of the Green Frills and Red Carpet mustard samples used for display in RAW format (A) on which the PCA (Principal Component Analysis) analysis was conducted and of the responses due to the analyzed variables (B).



## HYPERSPECTRAL RESULTS

PCA Model - Scatter 2D (T)
DataSet (Multi image import)





2D Scatter plot from a PCA (Principal Component Analysis) of a hyperspectral image of two mustard cultivations (Green Frills vs. Red Carpet).

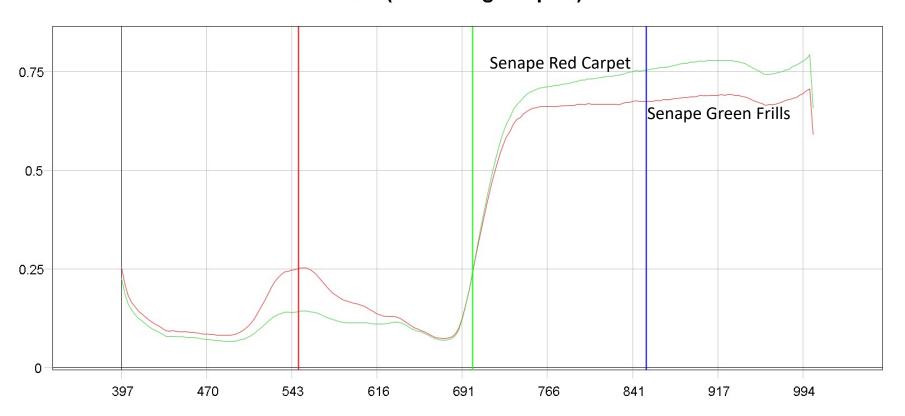
Colour density highlights the difference between the two cultivations.



## HYPERSPECTRAL RESULTS



## Spectral (X Training) DataSet (Multi image import)



Lines Plot for the analysis of average spectral data, which allow to identify spectral bands discriminating between Mustard "Green Frills" and "Red Carpet"



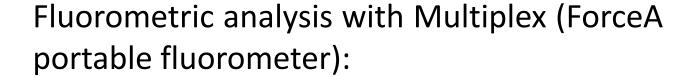
## FLUOROMETRIC ANALYSIS











- SFR\_R index related to leaves chlorophyll concentration
- FLAV index related to leaves flavonols concentration
- ANTH\_RG index related to leaves anthocyanins concentration

## FOICE A

#### CALCOLI DEL MULTIPLEX

- FLAV content
- Anthocyanins content

$$FLAV = \log \frac{FRF_R}{FRF_UV}$$

$$ANTH = \log \frac{FRF_R}{FRF_G}$$

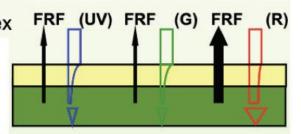
$$ANTH = \log \frac{5000}{FRF_R}$$

Chlorophyll content

$$SFR = \frac{FRF\_R}{RF\_R}$$

· NBI: Nitrogen Balance Index

$$NBI\_G = \frac{FRF\_UV}{RF\_G}$$





## FLUOROMETRIC RESULTS



NAataual	Green Frills				Red Carpet			
Mustard	ANOVA	L1	L2	L3	ANOVA	L1	L2	L3
Wet weight (g)	ns	0,0865	0,0800	0,0931	***	0,0390	0,0579	0,0222
Dry weight (g)	ns	0,0045	0,0049	0,0034	*	0,0028	0,0029	0,0013
Hypocotyl height (mm)	*	63,4	74,3	66,1	***	28,9	43,6	18,5
Cotyledons area (mm²)	**	125,0	169,7	104,9	***	112,3	151,5	73,3
SFR_G	ns	0,8577	0,8432	0,7425	**	0,7555	0,8305	0,7278
FLAV	ns	-0,0001	0,0240	0,0249	ns	-0,0005	-0,0211	-0,0259
ANTH_RG	ns	-0,0395	-0,0279	-0,0334	*	0,0527	0,0240	-0,0203

<u>Radish</u>	Green Daikon				Rioja Improved			
	ANOVA	L1	L2	L3	ANOVA	L1	L2	L3
Wet weight (g)	*	0,3226	0,3529	0,2621	***	0,1870	0,2118	0,1461
Dry weight (g)	*	0,0150	0,0174	0,0141	*	0,0092	0,0110	0,0079
Hypocotyl height (mm)	ns	74,4	68,3	57,6	*	61,6	54,2	48,4
Cotyledons area (mm²)	**	471,9	584,0	419,6	**	453,0	574,0	413,0
SFR_G	***	1,1250	1,0200	1,2320	ns	0,8404	0,8020	0,8889
FLAV	**	0,0210	0,0315	0,1071	**	0,1552	0,1133	0,1914
ANTH_RG	*	-0,0490	-0,0348	-0,0330	**	0,2330	0,2056	0,2557

Statistical analysis (one-way ANOVA) of the effect of LED light on morphometric and fluorometric parameters in red and green genotypes of mustard and radish. SFR G: clorophyll content

FLAV: Flavonols content

ANTH\_RG: antochyanins content



## **WORKING GROUP**









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## **CONCLUSIONS**



SOLE demonstrator allows to monitor and analyze data on plant growth and on the accumulation of bioactive metabolites in real time and remotely.

SOLE demonstrator can be easily adapted for space applications as growing plant on-board the International Space Station or in mini/micro satellites.

This biotechnological innovation ensures an improvement of the nutritional quality of the ready-to-eat product in future space missions.











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## THANK YOU.

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