



**CATRIN**

Czech Advanced  
Technology and Research  
Institute

# Electrochemical biosensors as an emerging data collection tool for plant phenotyping and agriculture

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**Photoelectrochemistry**  
 $2\text{H}_2\text{O} \xrightarrow{h\nu} 2\text{H}_2 + \text{O}_2$

**Magnetic nanostructures**

**Nanomaterials in biomedicine**

**Environmentally active complexes**

**Environmental nanomaterials**

**Carbon nanomaterials**

**Xe**

**Molecular Nanostructures on Surfaces**



**Laboratory of experimental medicine**

**Laboratory of genome integrity**

**Innovative Chemistry**

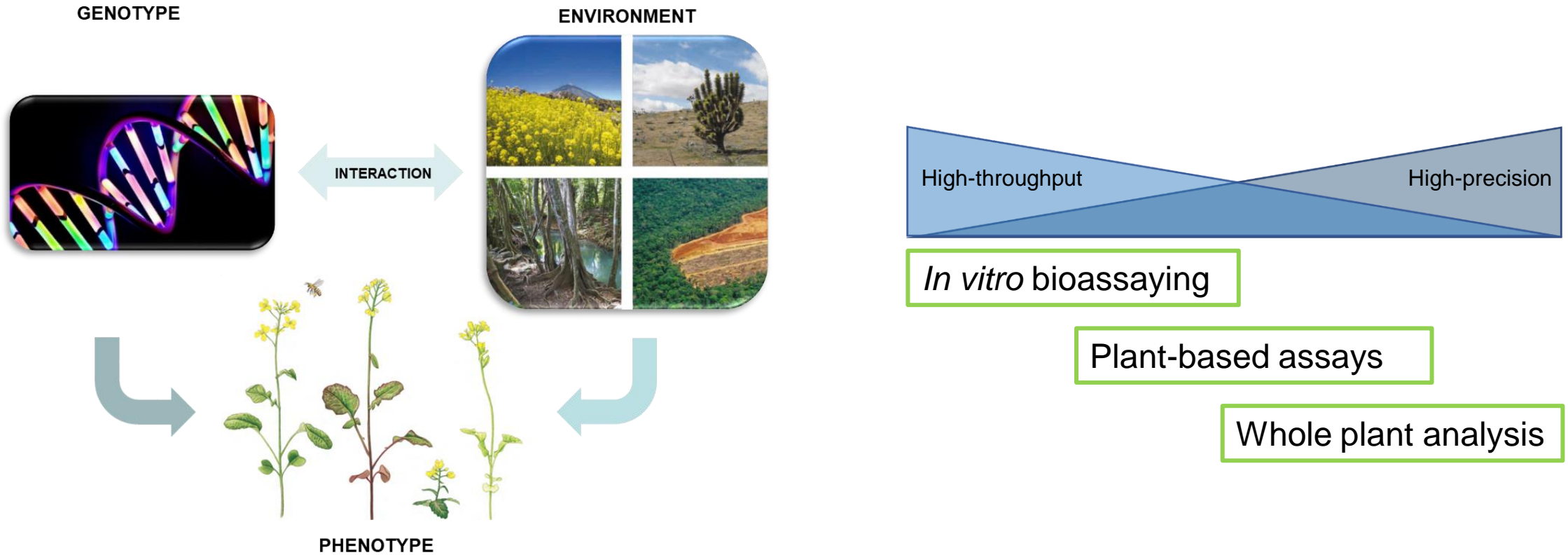
**Phenotyping**

**Plant genetics & engineering**

**Recombinant proteins**

**Phytochemistry**

## Monitoring of environmental interactions of plants



Key words: Phenotype / Non-invasive / Bioassaying / Automation / High-throughput/precision / controlled conditions



# What can we do

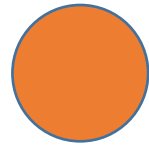


# olophen

phenotyping  
technologies  
& methods



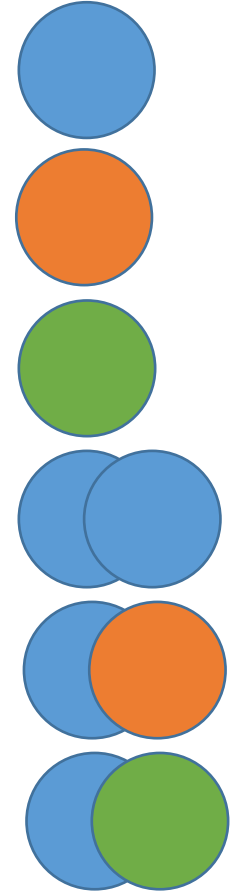
- High-throughput



- High-precision



- Affordable

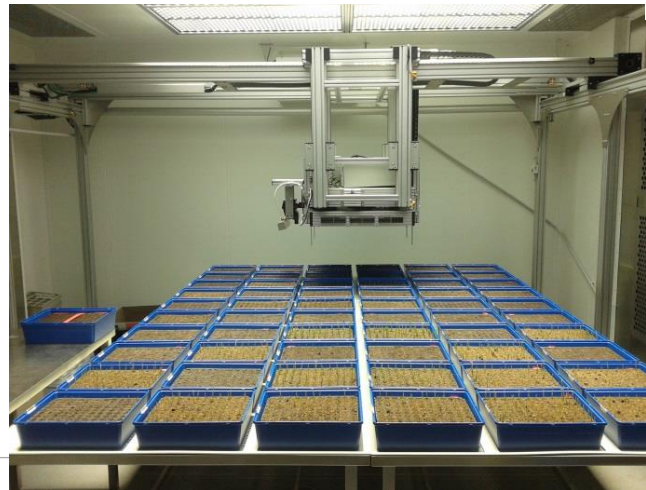
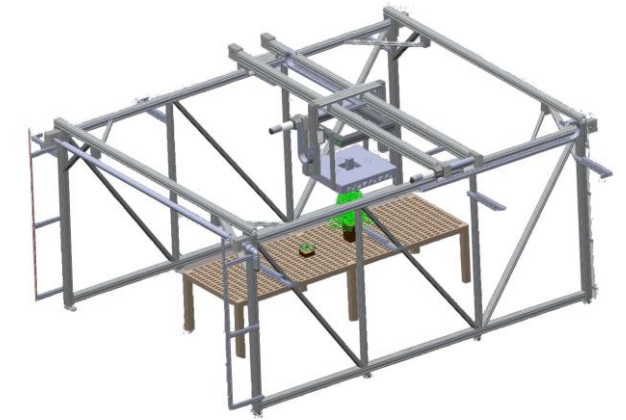


Palacký University  
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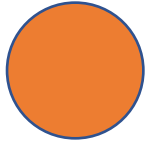
EDUCATION  
PARTNER



- Controlled conditions
- plant growth sensor (RGB top view high-resolution camera with homogenous LED lightning)
- sensors of physiological responses:
  - FluorCam unit – Chl fluorescence kinetic analysis
  - hyperspectral unit (VIS 380-900 nm)
- capacity: 7.5 square metres (528 culture multiwell plates, 64 trays, 1280 standardized Arabidopsis pots)

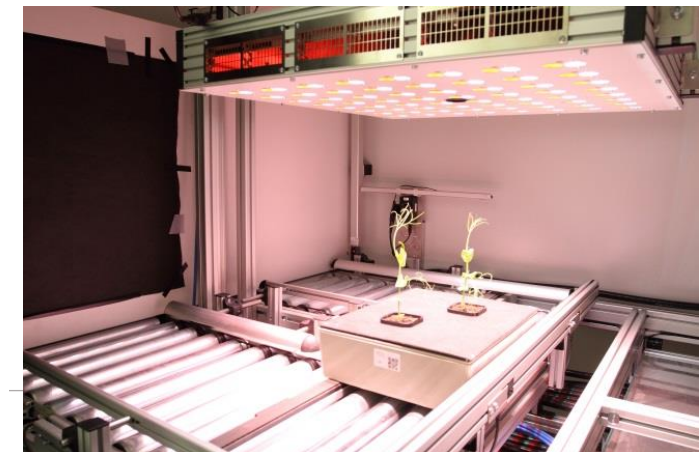
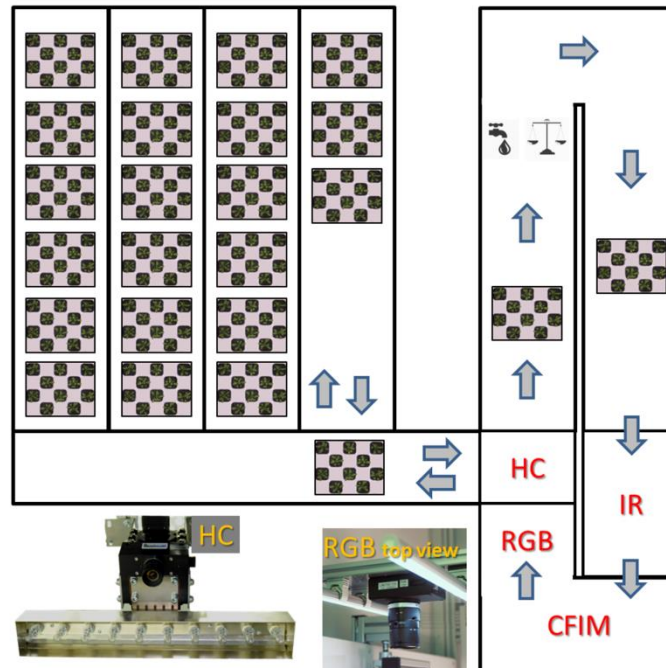
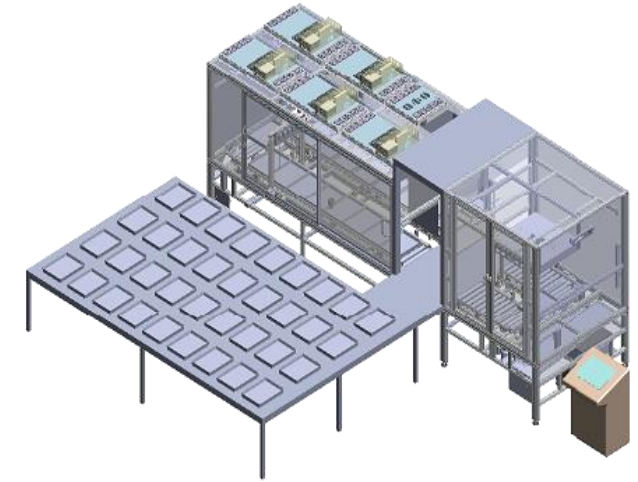
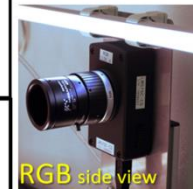
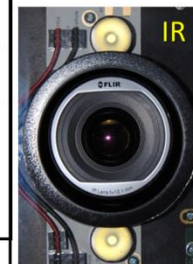




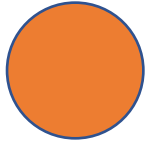


- Controlled conditions
- three RGB cameras, FluorCam, thermoimaging, acclimation cabinet, automatized pot weighing and watering
- capacity: 640 plants for top-view experiments, 64-32 plants for three-views experiments

Humplík JF, Lazar D, Husičková A, Spíchal L (2015) Automated phenotyping of plant shoots using imaging methods for analysis of plant stress responses – a review. *Plant Methods*, 11:29.



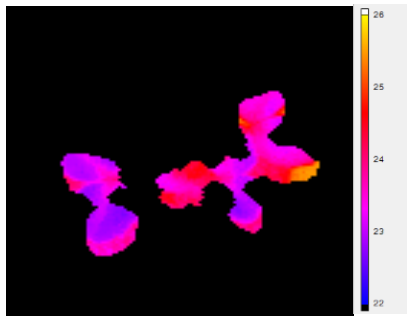




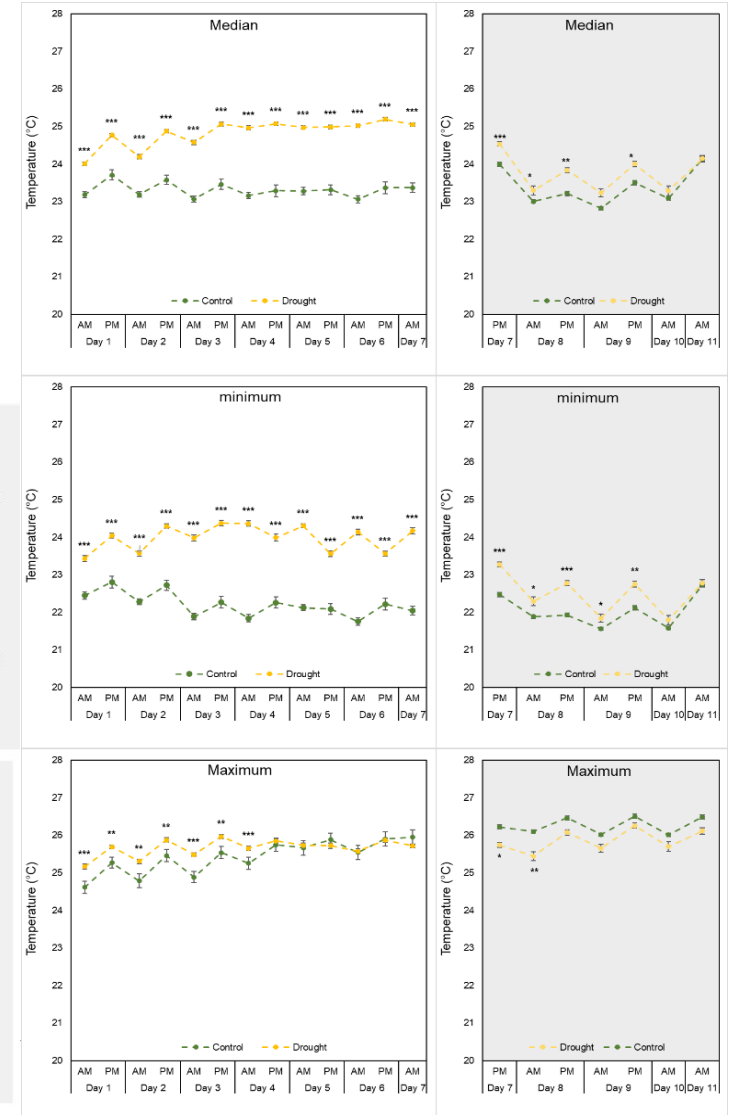
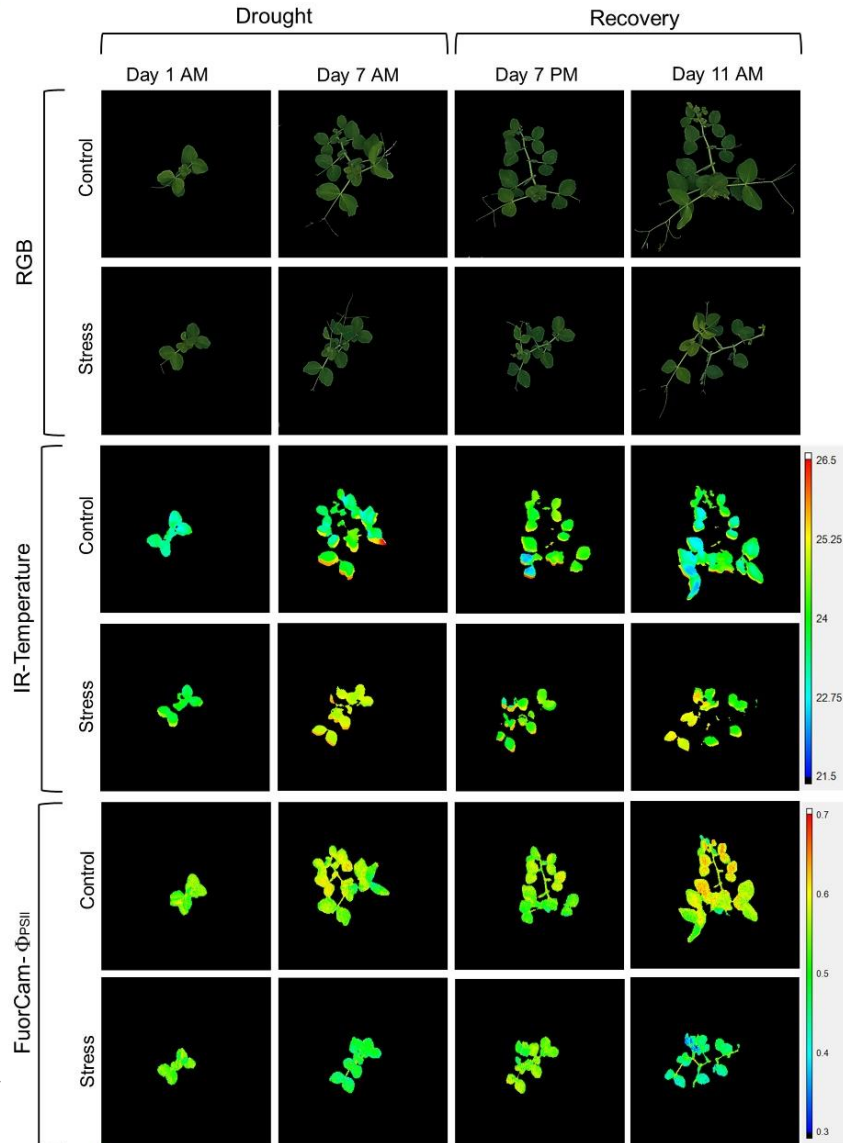
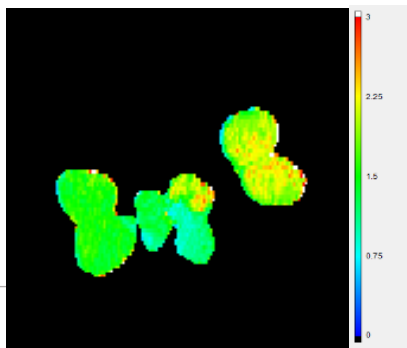
RGB



IR



FluorCam



What if  
you want?

$$1 G \times 1 E \times 10 R = 10 \text{ plants}$$

$$1 G \times 2 E \times 10 R = 50 \text{ plants}$$

$$5 G \times 2 E \times 10 R = 100 \text{ plants}$$

$$1 C \times 5 G \times 2 E \times 10 R = 100 \text{ plants}$$

$$5 C \times 5 G \times 2 E \times 10 R = 500 \text{ plants}$$

$$3 c \times 5 C \times 5 G \times 2 E \times 10 R = 1500 \text{ plants}$$

$$3 c \times 5 C \times 5 G \times 2 E \times 3 e \times 10 R = 4500 \text{ plants}$$

$$3 c \times 5 C \times 5 G \times 2 E \times 3 e \times 50 R = 22500 \text{ plants}$$



**G** genotype

**E** environmental condition

**C** compound

**c** compound concentration

**e** environmental condition level

**R** repetition

**T** timepoint

What if you want do it in 2 weeks?



- Analyses of effect on shoot area of *Arabidopsis*
  - Stimulation/Inhibition of shoot growth
  - normal conditions / Interaction with stress conditions
  - Salt, temperature, nutrition, drought, chemicals, pathogen response

METHODS ARTICLE

Front. Plant Sci., 04 October 2017 | <https://doi.org/10.3389/fpls.2017.01702>

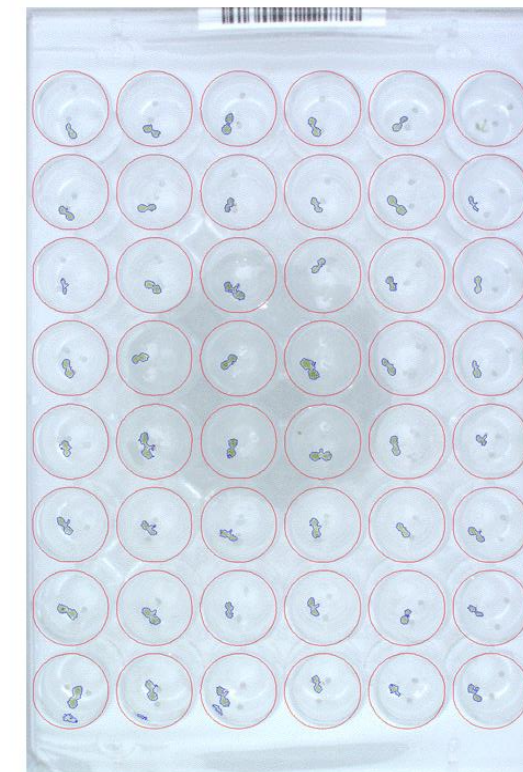
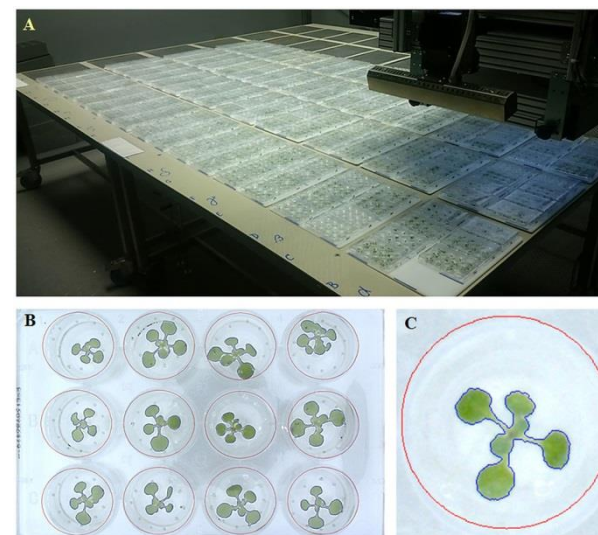


## An Automated Method for High-Throughput Screening of *Arabidopsis* Rosette Growth in Multi-Well Plates and Its Validation in Stress Conditions

Nuria De Diego<sup>1</sup>, Tomáš Fůrst<sup>1</sup>, Jan F. Humplík<sup>1,2</sup>, Lydia Ugena<sup>1</sup>, Kateřina Podlešáková<sup>1</sup> and Lukáš Spíchal<sup>1\*</sup>

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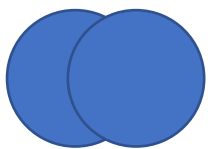


„This approach will allow simultaneous testing of a large number of potentially bioactive compounds in a wide range of concentrations and/or genotypes, under various growth conditions.“ (De Diego et al., 2017)





# In vitro bioassays – Shoot growth response



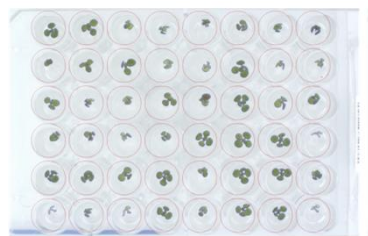
48-well plates      528 plates      25344 plants      7 days

1 day / 1 plate

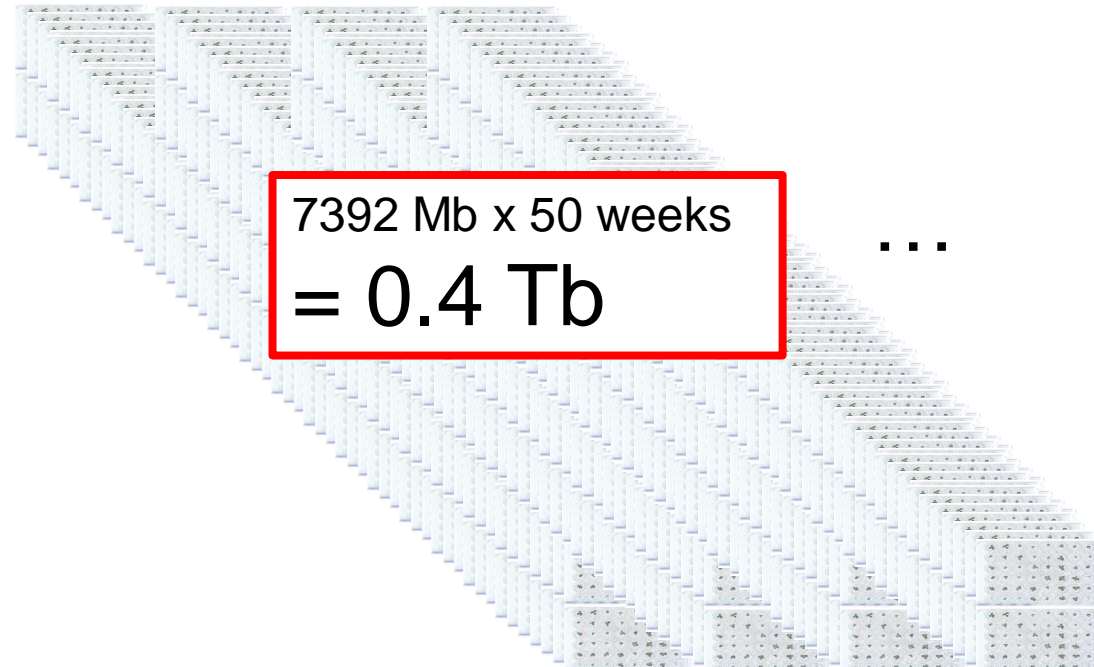
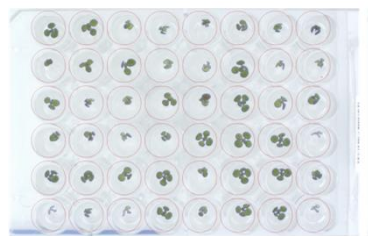
1 week / 1 plate

1 week / 528 plates

a.m.



p.m.



7392 Mb x 50 weeks  
= 0.4 Tb

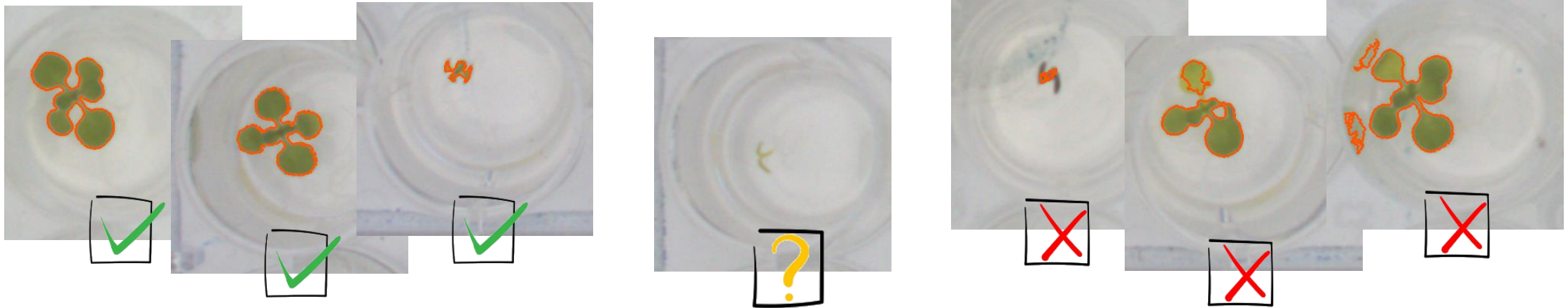
$$2 \times 10 \text{ Mb} = 20 \text{ Mb}$$

$$2 \times 7 \times 10 \text{ Mb} = 140 \text{ Mb}$$

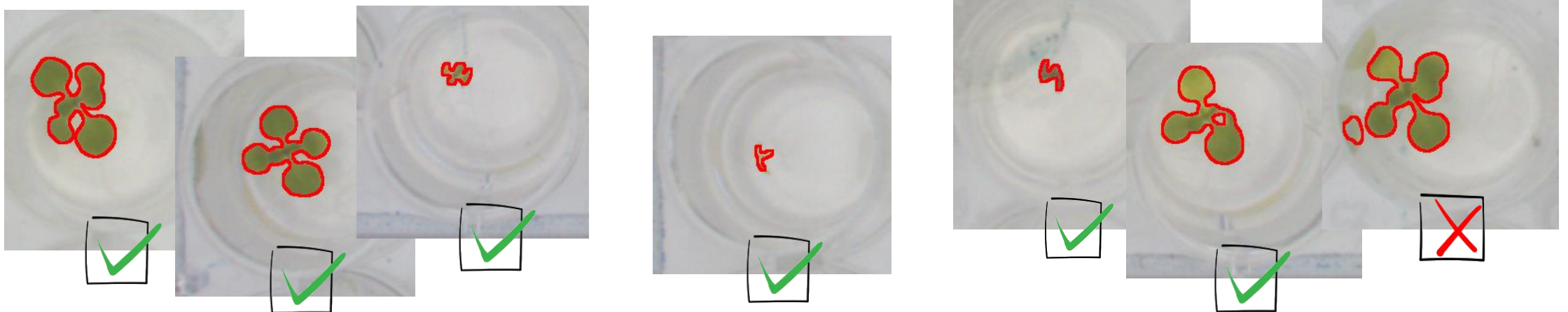
$$2 \times 7 \times 528 \times 10 \text{ Mb} = 7392 \text{ Mb}$$



- Color based analysis



- Neural network



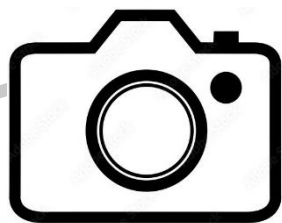
Because the model is only as good as you train it





# What we want to do



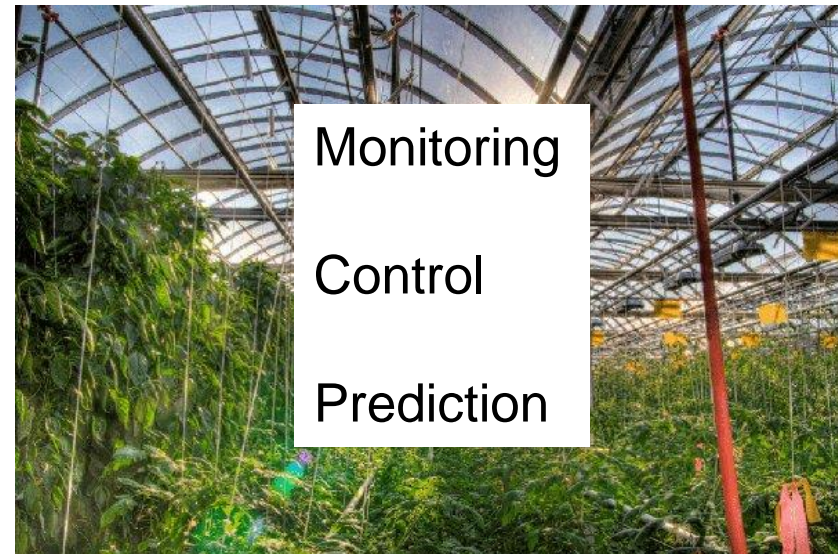
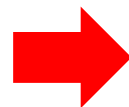


Optical sensors

wearable sensors

Environmental sensors

AI



Monitoring

Control

Prediction





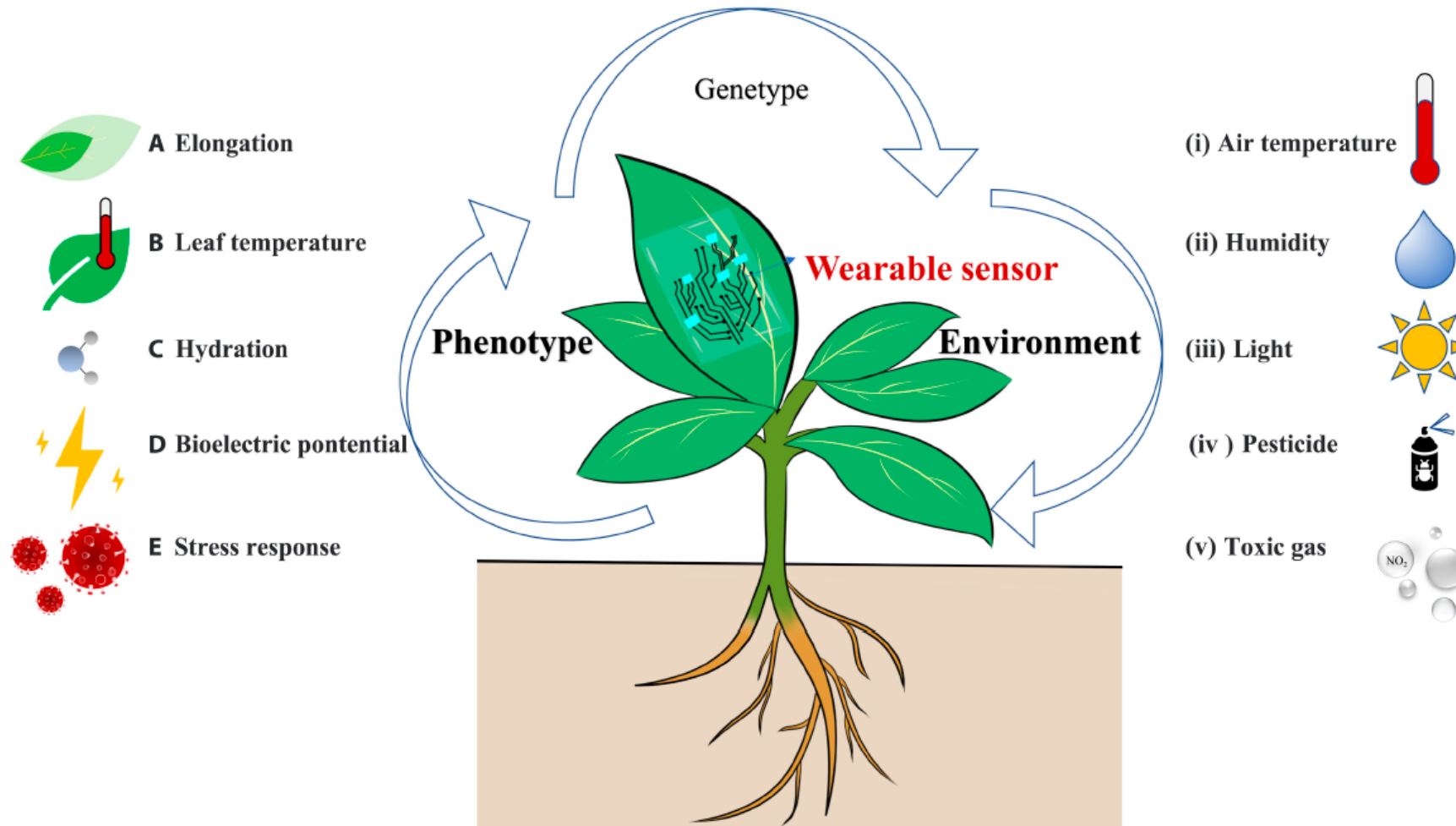
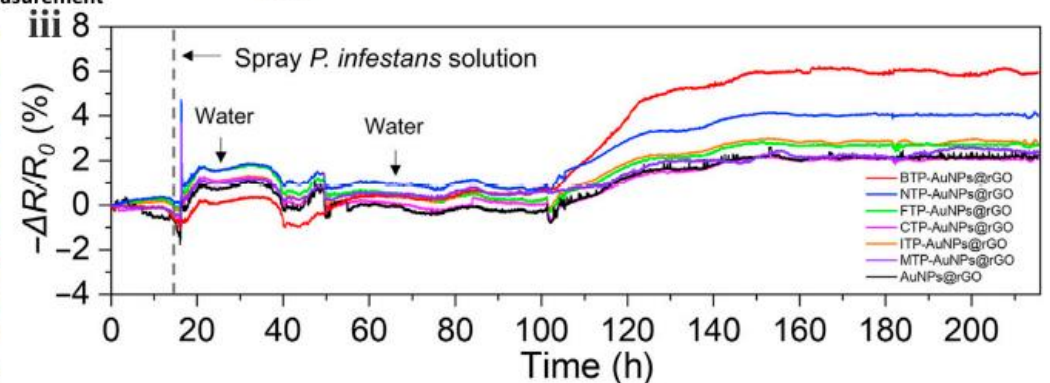
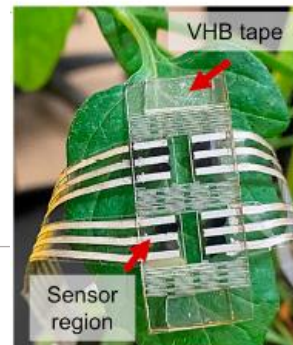
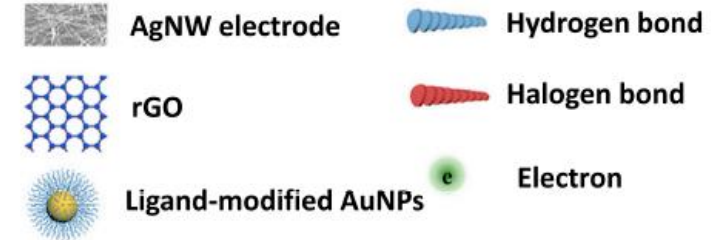
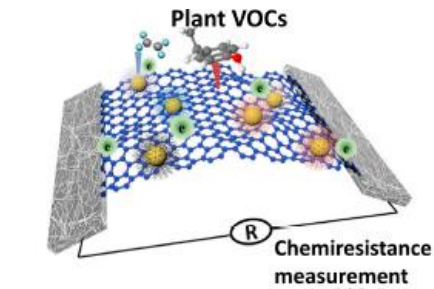
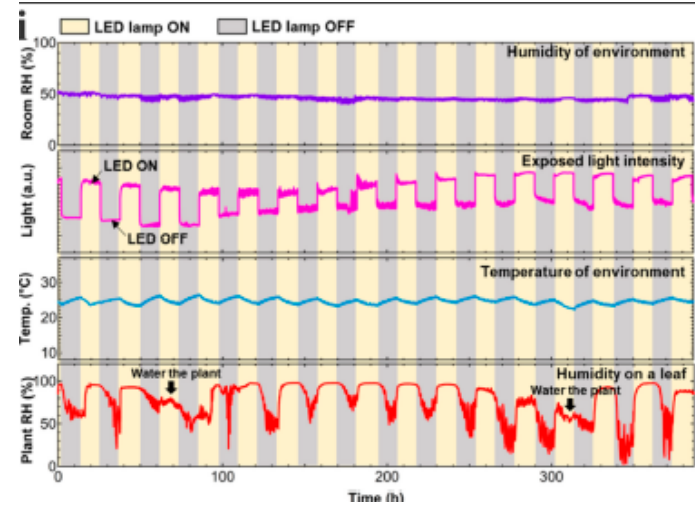
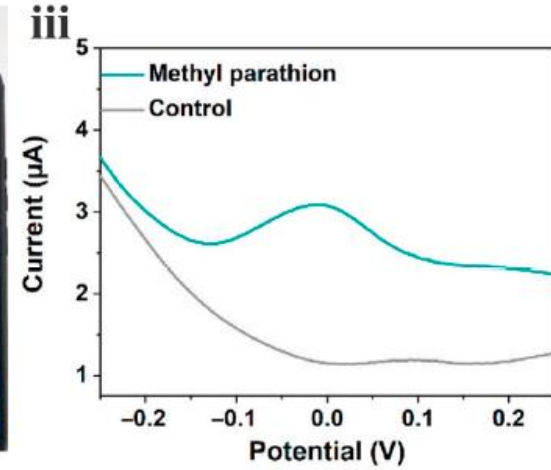
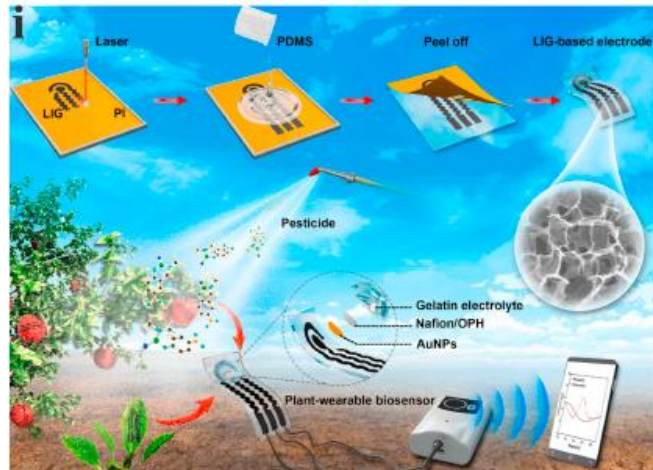
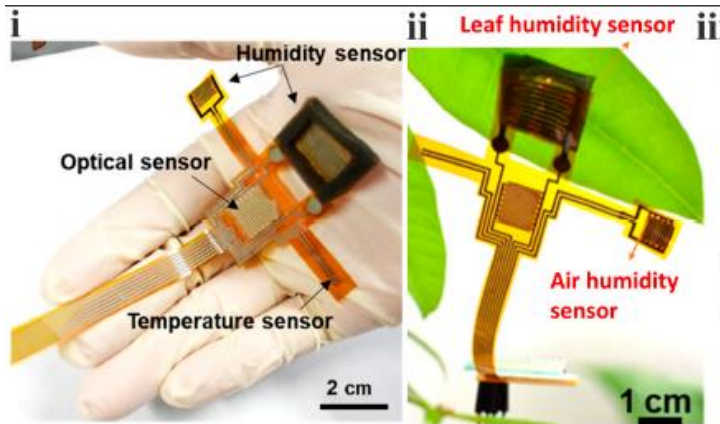


Fig.1. Wearable sensors for monitoring plant phenotypes and environment.

Phenotypes	Sensor	Plant	Ref.
Elongation	A Ti/Au-based strain sensor	Barley and lucky bamboo	[50]
	A chitosan-based sensor	Cucumber	[54]
	A carbon nanotube/graphite-based strain sensor	<i>Cucurbita pepo</i> , <i>Solanum melongena</i> L.	[39]
Temperature	A liquid-alloy-based sensor	Sprout	[55]
	A tag sensor	/	[56]
	A “dust” network of wireless sensors	Melon	[40]
Hydration	An RFID-based system	Pumpkin	[58]
	A PI-based sensor	Tobacco	[64]
	A GO-based humidity sensor	<i>Epipremnum aureum</i>	[41]
Bioelectric potential	A graphene-based sensor	Maize	[69]
	A Cu-based flexible electronic sensor	Watermelon	[70]
	BDD electrodes	<i>Opuntia</i>	[73]
	BDD /Nafion and BDD/ Vylon electrodes	<i>Aloe</i> and <i>Opuntia</i>	[42]
	Thermogel-based morphable ionic electrodes	Sunflower and tobacco	[48]
Stress response	Self-adhering electrodes	<i>Dionaea muscipula</i> , <i>Arabidopsis thaliana</i> , and <i>Codariocalyx motorius</i>	[74]
	A graphene-based sensor array	Tomato leaf	[80]
	Conductive polymer electrodes	<i>Hosta</i> and pothos seedling	[82]
	Conductive polymer electrodes	Grape leaf	[44]







- Nuria De Diego
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*Jan Zdražil*