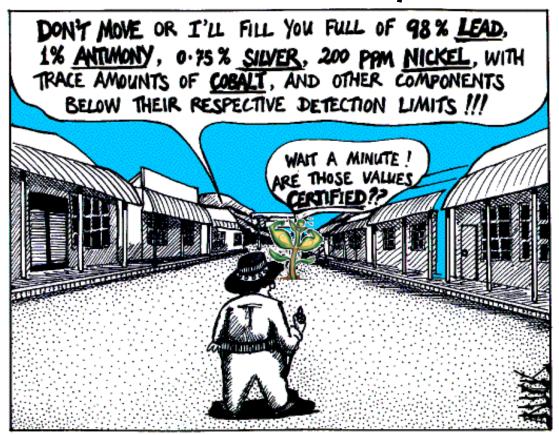
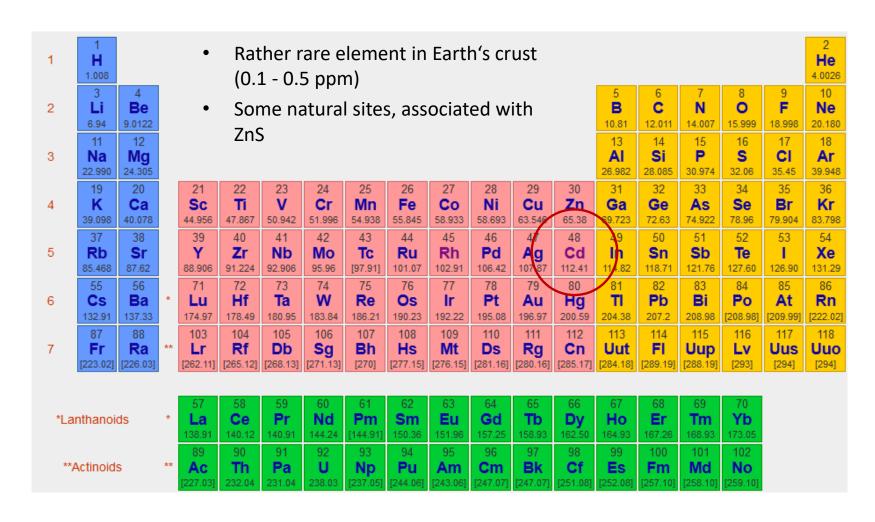
Heavy Metals and Plants - a complicated relationship Cadmium toxicity



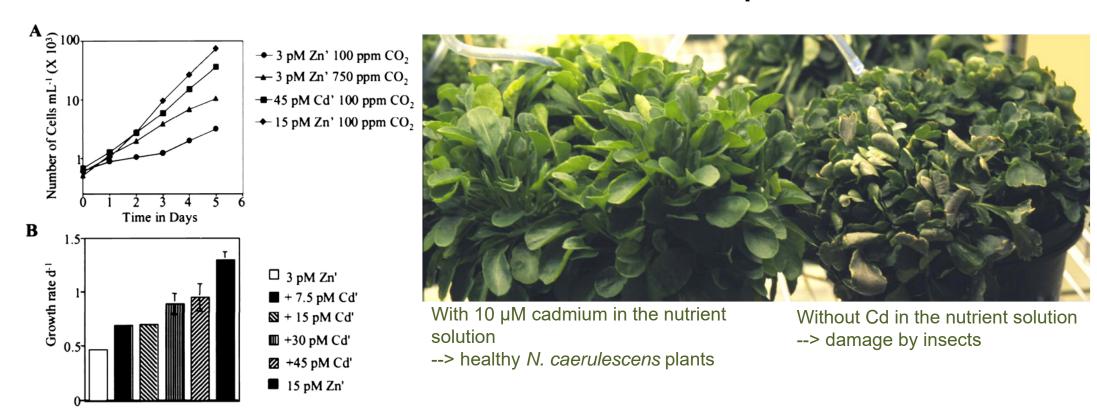
Heavy metal-hyperaccumulation in the Wild West

modified from: http://strangematter.sci.waikato.ac.nz/

Cadmium in nature



Cadmium as a micronutrient: the exception...

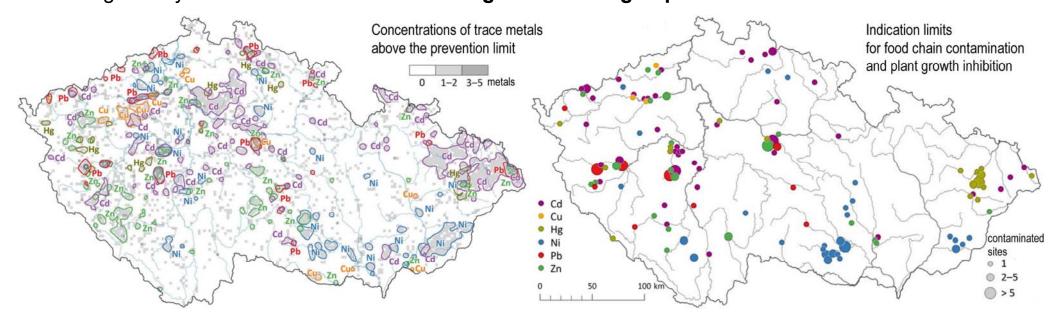


Cadmium as Plant-micronutrient in *Thalassiosira weissflogii*. A, B: growth of the algae. (Lane and Morel, 2000, PNAS97)

Küpper H, Kroneck PMH (2004) MIBS 44 (Sigel et al., eds), chapter 5

Cadmium pollution in the environment

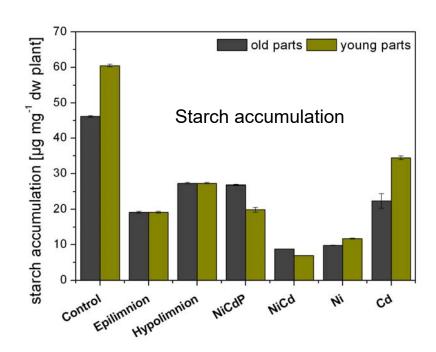
- → Itai-itai disease (japanese ouch-ouch sickness), 1 of the 4 big pollution diseases in Japan
- → Mass cadmium poisoning in Japan: severe pains in joints and spine, softening of the bones, kidney failure
- → Cadmium release into rivers by mining, the mining companies were successfully sued for the damage
- → In Europe: Cd from contamination of fertilisers in Agriculture
- → Mechanistic knowledge of metal(loid) toxicity is required for realistic environmental risk assessment
- → Revealing toxicity mechanisms is essential for targeted breeding of plants that are less vulnerable to stress

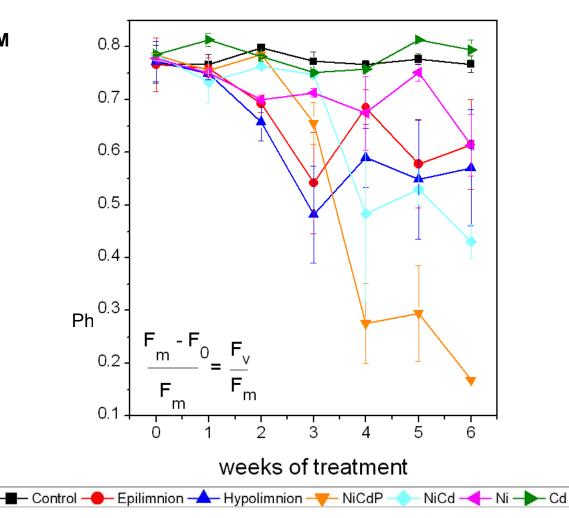


Maps from Bednarova et al. (2016) Spatial distribution and risk assessment of metals in agricultural soils. Geoderma 284, 113-121

Metal toxicity induced inhibition of photosynthesis at synergistically acting nanomolar Cd+Ni concentrations

- -Ceratophyllum demersum plants treated with natural or simulated lake water containing only 3 nM Cd²⁺ and 300 nM Ni²⁺ already show inhibition
- inhibition by Ni+Cd combination treatment much stronger than by the single metals → synergistic effect!





Andresen E, Opitz J, Thomas G, Stärk H-J, Dienemann H, Jenemann K, Chang C, Küpper H (2013) Aquatic Toxicology 142-143, 387-402

Growth inhibition

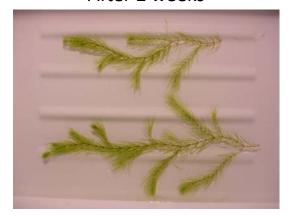
Before treatment start



After 1 week



After 2 weeks



After 3 weeks



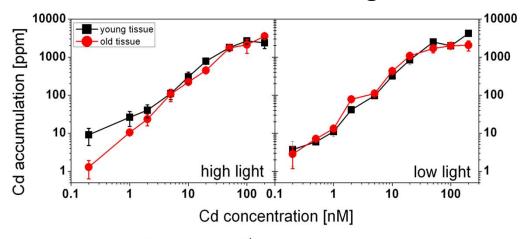
After 4 weeks

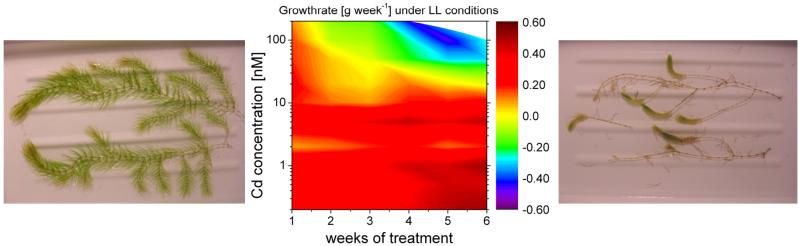


C. demersum treated with 200 nM Cd for 4 weeks

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.

Accumulation of Cd and growth





→ Already low nanomolar Cd is toxic for sensitive plants

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.

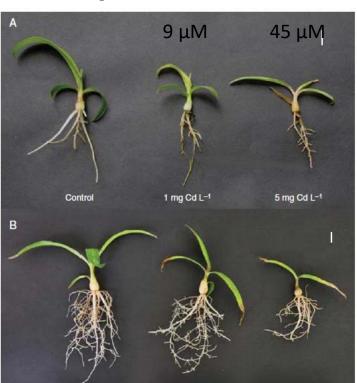
Cd toxicity in plants

- 1) Roots
- 2) Photosynthesis
- 3) Reactive oxygen species
- 4) Genotoxicity
- 5) Stress prevention

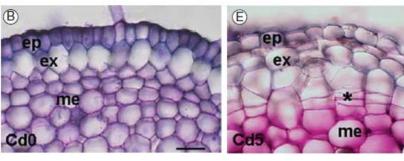
Cadmium toxicity in plants – 1: Roots

First organ which gets affected

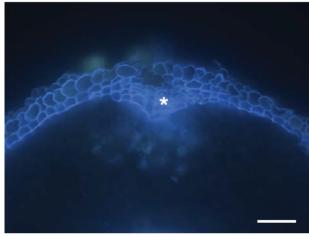
Reduced growth after Cd treatment



More layers of hypodermal periderm



More layers & suberized cell walls (*) like after injury of root surface

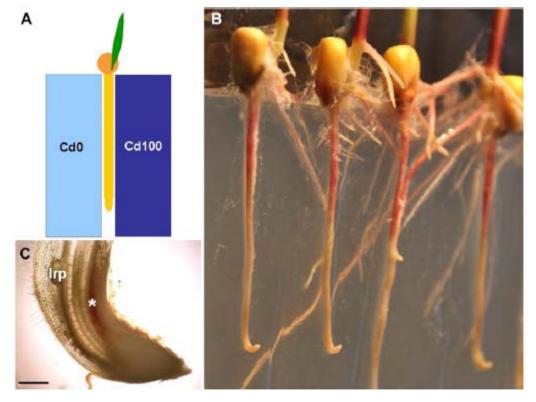


Lux et al., Annals of Botany 107:285-292, 2011

- Maize seedlings with proper roots placed between 2 agar blocks
- one of which contained Cd (50 or 100 μ M), grown in phytochamber under nature-like conditions

→ Roots bending towards the Cd-containing agar → due to growth stop on the Cd-side & continued growth on control-side

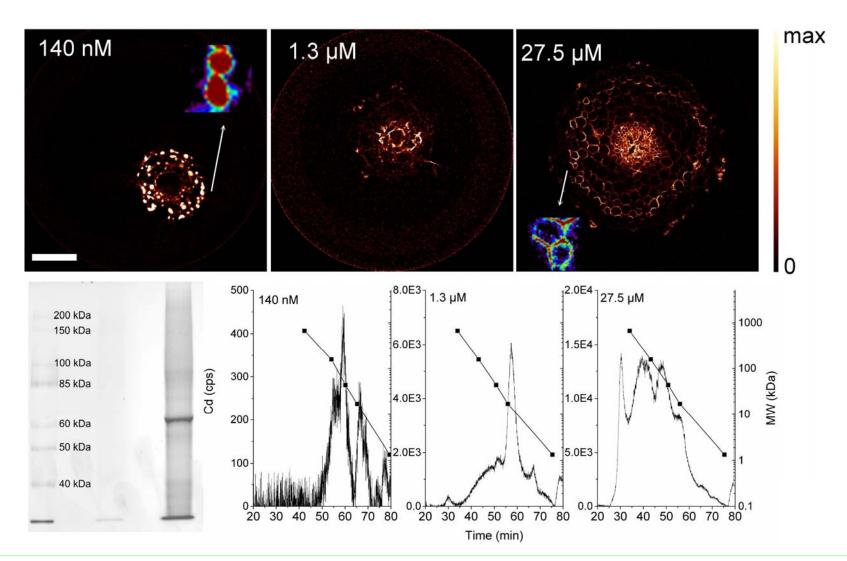
Lignification on Cd-exposed side (*) and initiation of lateral root primordium (Irp)



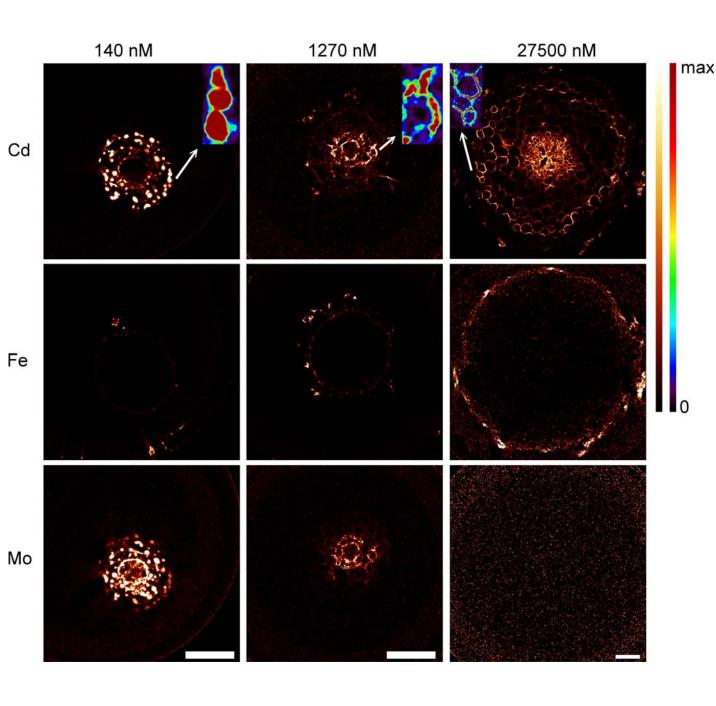
Lux et al., Journal of Experimental Botany 62(1): 21-37, 2011

Mechanisms of sublethal vs. lethal Cd toxicity in soybean roots

- → change of intracellular localisation: symplastic vs. apoplastic
- → change of tissue localisation: central cylinder vs. unspecific
- → change of target proteins: few vs. unspecific
- → Drastically different mechanism of toxicity at rarely studied sublethal vs. frequently studied lethal concentrations!



Andresen E, Flores-Sanchez IJ, Brückner D, Bokhari SNH, Falkenberg G, Küpper H* (2023) Journal of Hazardous Materials 442, 130062.



Where does the Cd go? Sub-cellular distribution (measured at Cd Kα with 0.6μm resolution at beamline PETRA P6)

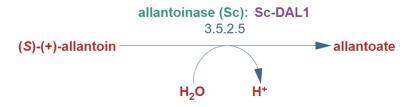
3- phase response to Cd toxicity

- A) Initially, at sublethal concentrations with low toxicity, Cd is sequestered into vacuoles of the root parenchyma
- B) At sublethal concentrations with high toxicity, stronger accumulation starts in cell walls in the vascular bundle, starting to block micronutrient (Fe, Mo) uptake
- C) At lethal concentration, Cd accumulates unspecifically in cell walls throughout the root diameter, and blocks transport of micronutrients (Fe & Mo)

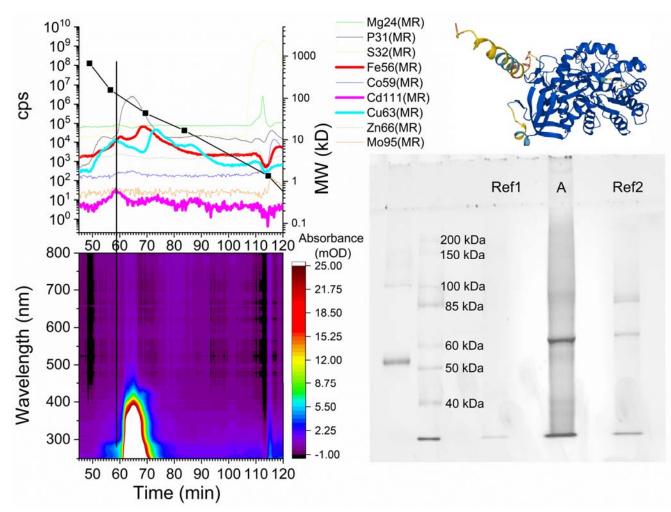
Andresen E, Flores-Sanchez IJ, Brückner D, Bokhari SNH, Falkenberg G, Küpper H (2023 Sublethal and lethal Cd toxicity in soybean roots specifically affects the metabolome, Cd binding to proteins and cellular distribution of Cd. Journal of Hazardous Materials 442, 130062 https://doi.org/10.1016/j.jhazmat.2022.130062

To which exact proteins does the Cd bind at sublethal concentrations? → protein purification, identification and verification

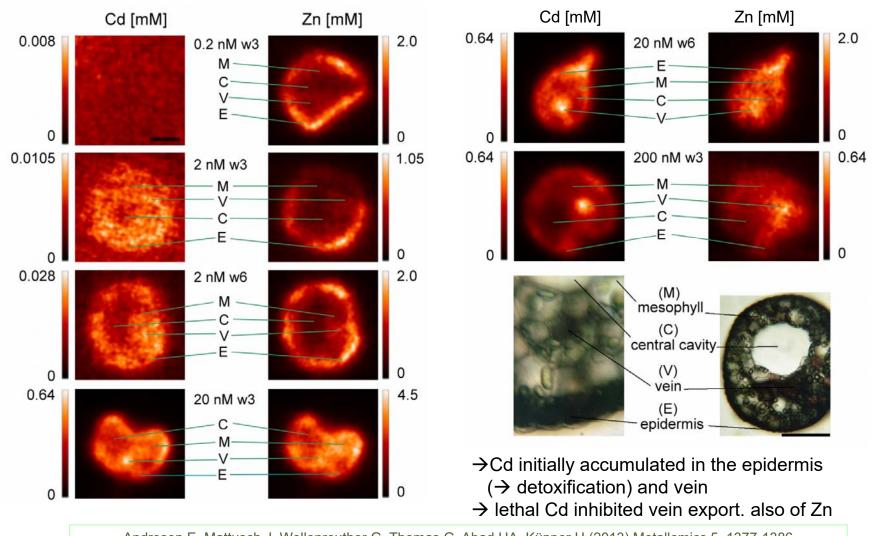
- → highest affinity targets among soluble proteins in roots are allantoinase; protein disulfide isomerase-like proteins (PDI), aquaporin-like and peroxidase-like proteins,
- → verified for allantoinase by metabolomics, there it most likely replaces the Zn²+ in the active centre



→ others to be investigated in future research



Distribution of Cd and its effect on Zn distribution in the non-accumulator shoot model *Ceratophyllum demersum*



Andresen E, Mattusch J, Wellenreuther G, Thomas G, Abad UA, Küpper H (2013) Metallomics 5, 1377-1386

Changing distribution of Cd and micronutrients under Cd toxicity

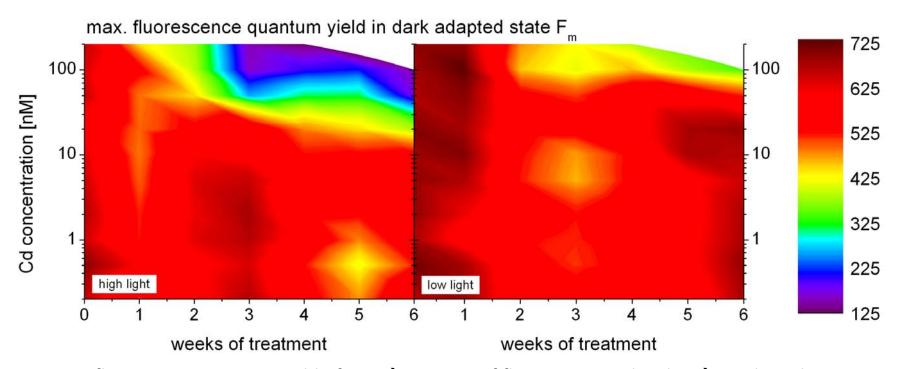
Increasing Cd:

- Roots: more aplastic Cd binding at high Cd
- Roots and leaves: more Cd in vein
- Roots and leaves: inhibition of micronutrient (Fe, Mo, Zn) transmembrane transport by Cd
- Leaves: Increased sequestration into non-photosynthetic tissues
- Leaves: re-distribution of Zn → inhibited export out of vein

Cadmium toxicity in plants – 2: Photosynthesis

- Indirect measurement:
 - Growth, O₂ production / CO₂ consumptio: balance with O₂-consuming and CO₂ producing (photo-)respiration
 - Diminishing the Chl/pigment/protein content
- Direct: Photosynthetic paramters via Chlorophyll fluorescence measurement

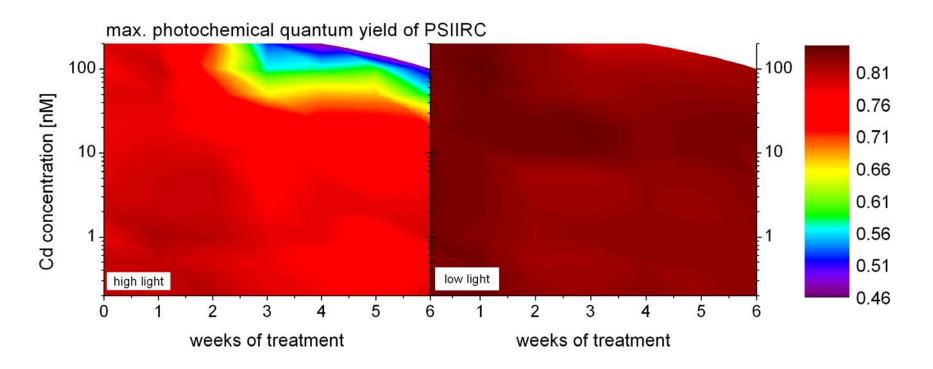
Cd affects Photosynthetic light reactions in C. demersum



- Max. fluorescent quantum yield of PSII → amount of fluorescent molecules → LL plants have bigger antenna systems
- Reduced F_m towards higher concentrations and longer treatment duration → decreased Chl content
- Reduction in week 1 due to acclimation to Cd + HL

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.

Cd affects Photosynthetic light reactions in C. demersum



Reduced variable fluorescence $F_v/F_m = (F_m-F_0) / F_m \rightarrow$ decreased activity of PS II towards higher concentrations and longer treatment duration

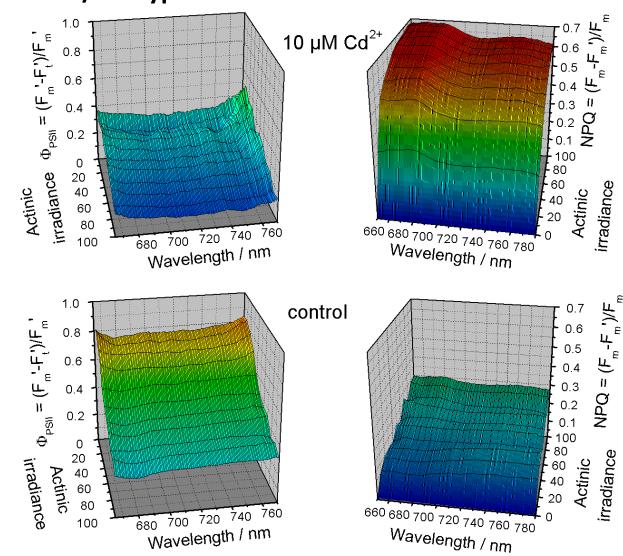
HL much more affected than LL

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.

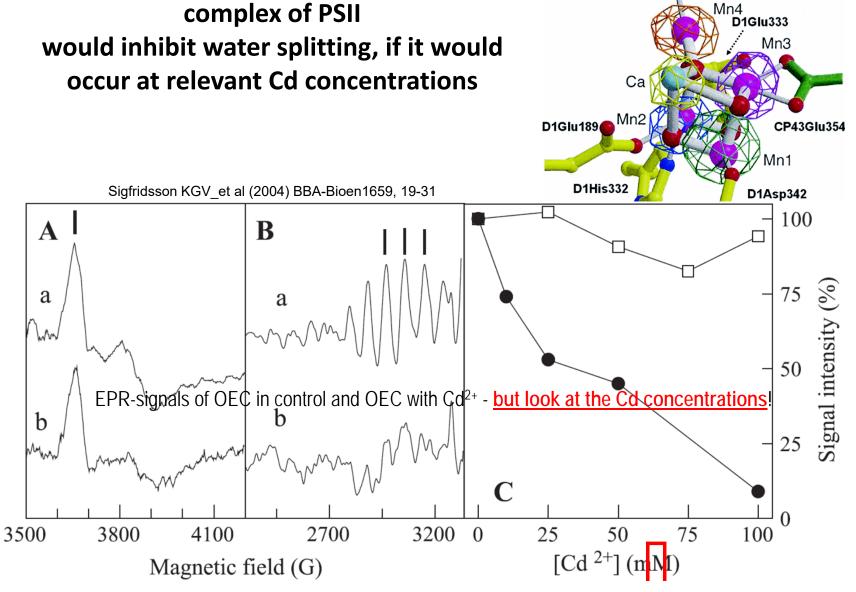
Cd-stress in the Zn-/Cd-hyperaccumulator *T. caerulescens*

Spectral changes of PSII activity parameters:

- inhibition of electron flow visible over the whole excitation spectrum
- → RC-related
- increase of NPQ with peak at 680nm → Antenna-related

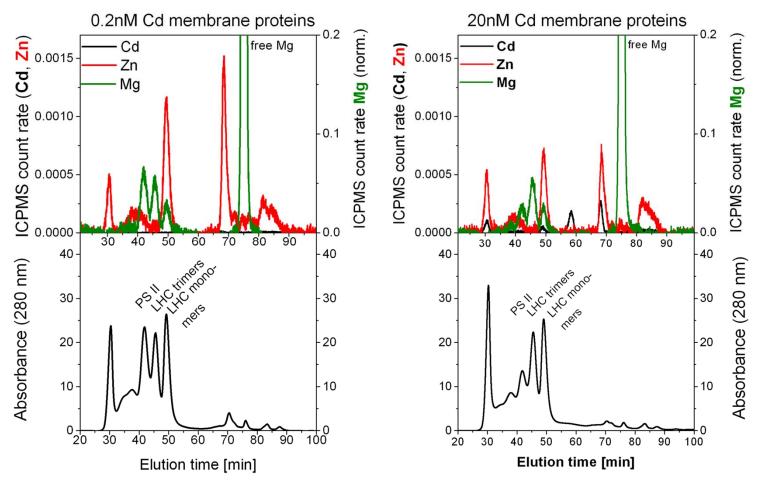


Substitution of Ca by Cd in water splitting complex of PSII would inhibit water splitting, if it would



D1Asp170

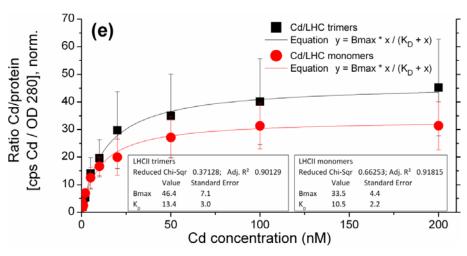
Cd stress in *C. demersum*: Incorporation of Cd into proteins in LL analysed by metalloproteomics via HPLC-ICP-MS



→ Cd binding to main light harvesting antenna LHCII in LL

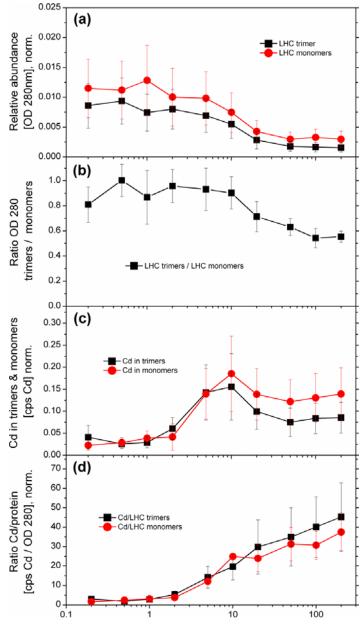
Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.

Example of metal toxicity in the nanomolar range in "normal" plants: Incorporation of Cd into LHCII in LL

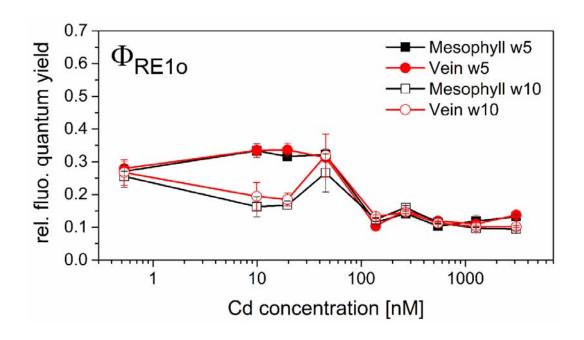


- → Cd binding to LHCII causes disintegration of trimers
- → Cd bind to LHCII with dissociation constants in the low nanomolar range → diminished photosynthesis despite funtional reaction centres!

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.



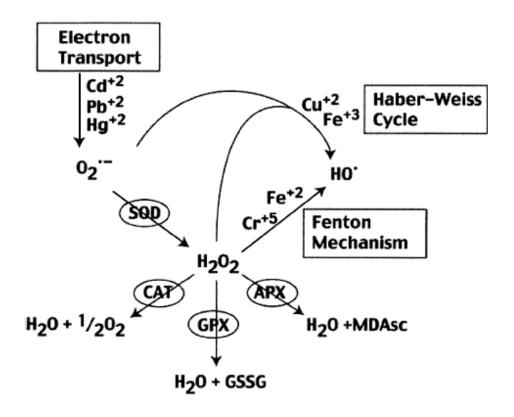
Chronic Cd toxicity in the nanomolar range in soybean plants: additional inhibition of electron transfer to PSI



→ Cd inhibits electron flow to PSI already in the environmentally relevant nanomolar range

ROS and Cadmium

Cadmium redox inert → No Fenton reaction!



Fenton:

Fe(II) +
$$H_2O_2 \rightarrow \rightarrow$$

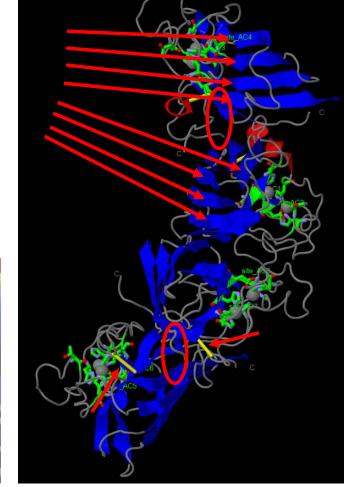
Fe(III) + $R^* + OH^- + H_2O$

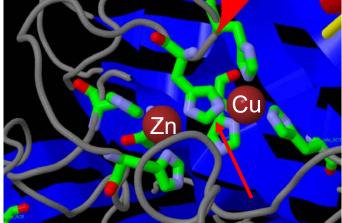
"Biggest source of ROS in animal cells" ...

→ NEVER shown in vivo!

Possible target for Cd toxicity leading to ROS: Superoxide dismutase (SOD), in plants a Cu/Zn enzyme

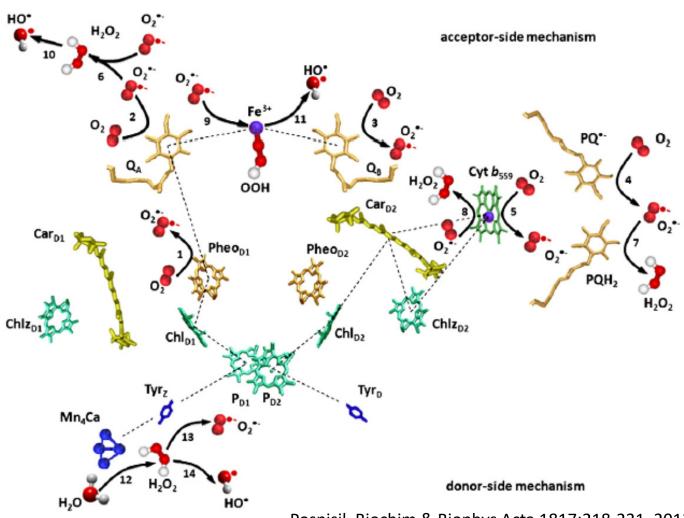
Substitution of Zn by Cd in SOD may contribute to oxidative stress during Cd toxicity





Spinach SOD, From: Kitagawa Y et al., 1991, J Biochem 109, 477-85, images generated with Jena 3D viewer

Photosynthesis-related ROS



Pospisil, Biochim & Biophys Acta 1817:218-231, 2012

ROS and Cadmium

ROS production



Removal of ROS



- Cd interferes with photosynthesis / respiration → electrons transferred to O₂
- Cd replaces Zn in SOD (e.g.) → less functional SOD

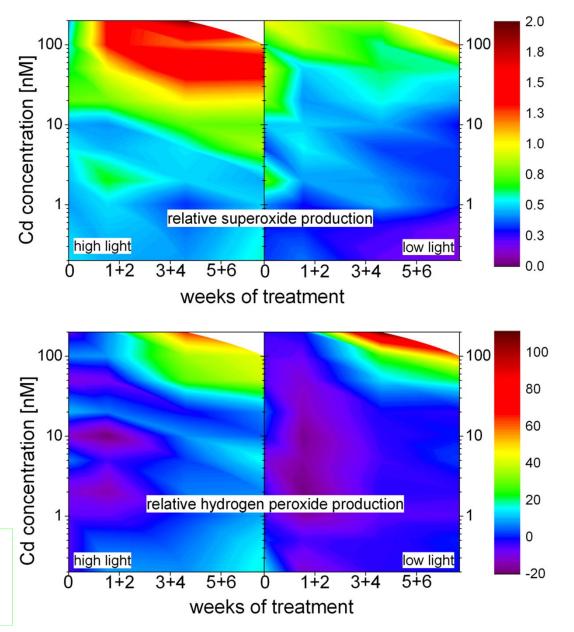
• In response antioxidant enzymes



Production of reactive oxygen species (ROS) during Cd-stress in HL vs. LL

- → Cd-induced formation of ROS stronger in HL
- → superoxide formation starts at lower Cd concentrations than peroxide formation

Andresen E, Kappel S, Stärk HJ, Riegger U, Borovec J, Mattusch J, Heinz A, Schmelzer CEH, Matoušková Š, Dickinson B, Küpper H (2016) New Phytologist 210, 1244-1258.



Influence of Cd on antioxidant enzymes

- Lower Cd concentrations and shorter treatment duration tend to increase the antioxidant system
- Longer exposure and higher Cd concentrations lead to decreased activity or content of the antioxidants

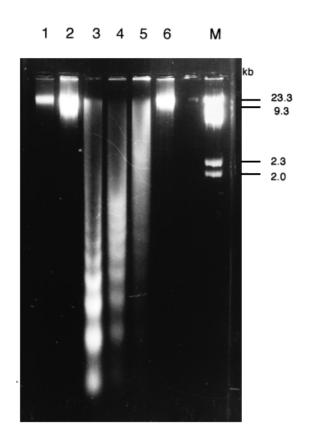
Cadmium toxicity in plants – 4. Genotoxicity

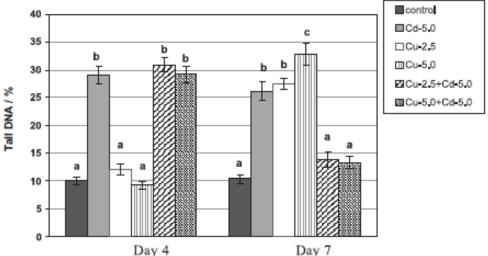
- Induction of DNA damage by
 - direct interaction with the nucleotides
 - modifications like base and sugar lesions, DNA strand breaks, destruction of DNA-protein crosslinks etc.
 - inhibiting DNA repairing enzymes
 - Induction of ROS, ROS lead to lipid peroxidation, which causes membrane damage and production of mutagenic aldehydes

Methods to detect Genotoxicity

- DNA Analyses
 - Gelelectrophoresis and Comet Assay
 - Random amplification of polymorphism DNA (RAPD)
- DNA / Chromosome Analyses
 - Micronuclei formation
 - Sister chromatid exchange
 - Chromosomal aberrations
- Upregulation of DNA-related / repairing enzymes

DNA disruption – Gelelectrophoresis and Comet Assay



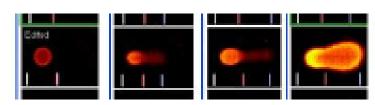


Lemna minor treated with μM concentrations of Cd and Cu. Tail DNA (Comet - length) increase due to metal treatment.

Cvjetko et al., Arh Hig Rada Toksikol 61:287-296, 2010

1: DNA from control plant, 2:-6: DNA from plants treated with Cd 10, 50, 75, 100, 1000 μM

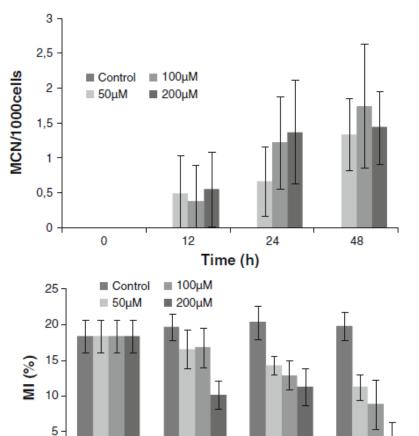
Fojtova & Kovarik, Plant, Cell & Envir. 23:531-537, 2000



From the CometAssay Manual, Trevigen^R

Micronuclei & Mitotic index

48

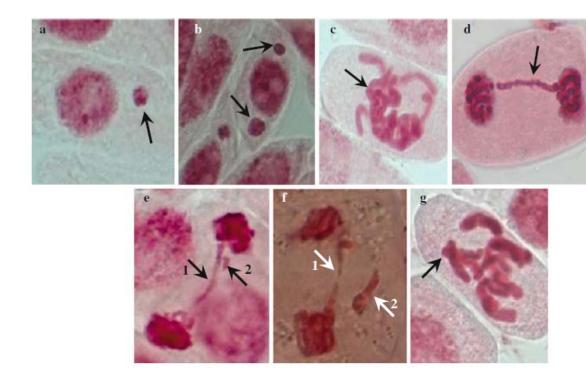


12

Time (h)

- Vicia root meristem cells; established assay
- Micronuclei formation due to malfunctioning cell division
- Dose and time dependent
 - $-\,$ Cd treatment increases MCN at high μM Cd
- Mitotic index: ratio of cells in metaphase stage to all cells
 - $-\,$ Cd treatment reduces MI at high μM Cd

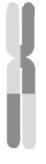
Chromosomal aberrations



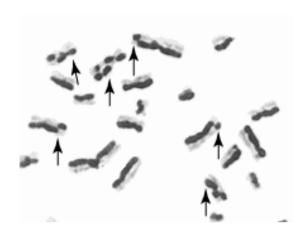
Vicia root meristem cells a, b, e, f = 50 μ M Cd c, d, g= 200 μ M Cd

a & b = micronuclei c = sticky chromosome d = chromosome bridge e = "" + break f = "" +isolated chromosome g = laggered chromosome in metaphase

Sister chromatid exchange



- -Exchange of identical parts of both sister chromatids in the same chromosome after / during DNA replication
- -As DNA sequence identical, exchange does not lead to genetic information change (≠ crossing over)
- -Happens in normal cells, but enhanced after treatment with toxic / radioactive substances



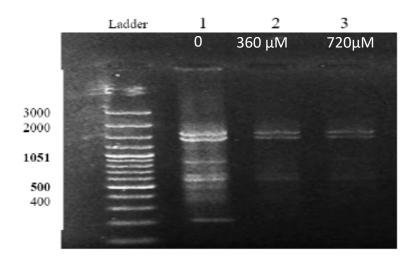
Test substance	Concentration	SCEs/metaphase (mean ± SE)	MI (mean ± SE)
Negative Control (Hoagland's nutrient Solution)	0	6.26 ± 0.29	11.80 ± 0.11
Cadmium nitrate (μM)	50	6.63 ± 0.3	9.70 ± 0.26***
	100	$7.43 \pm 0.04**$	$7.16 \pm 0.32***$
	200	$7.90 \pm 0.07***$	1.16 ± 0.03***
Positive control (Cyclophosphamide, µg/mL)	5	12.11 ± 0.06 ***	2.36 ± 0.31***

^{** (}P < 0.01) and *** (P < 0.001) compared with negative control.

From: http://www.siteklabs.com/GenTox/MammalianCellCytogenetics.html

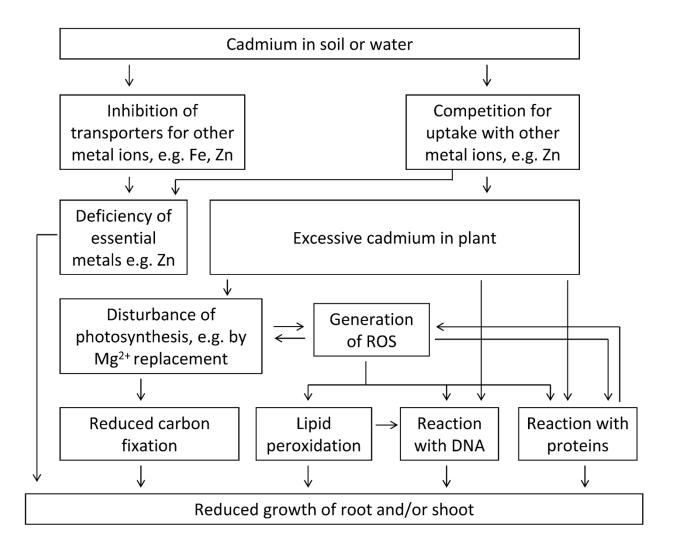
Random amplification of polymorphism DNA analyses (RAPD)

- Cd interacts with DNA / induces mutation
- New / disappearing primer binding sites
- → New / disappearing bands on gel
- But mostly seen at very high μM Cd!



	Treatments					
		40 (mgL ⁻¹) Cd concentration		80 (mgL ⁻¹) Cd concentration		
Primers names	Total bands in control	Appearance of new bands	Disappearance of control bands	Appearance of new bands	Disappearance of control bands	
OPA-2	1679, 1500, 1205, 874, 657, 603, 513, 221		1205, 874, 513, 221	900	1205, 874, 513, 221	

Shahrtash et al., J of Cell & Molecular Research 2(1):42-48, 2010



Andresen E, Küpper H (2013) Cadmium Toxicity in Plants. In: Cadmium: From Toxicity to Essentiality, "Metal Ions in Life Sciences Vol. 11;

Summary Cd toxicity

- Threshold concentration for first toxic effects (Ceratophyllum, soybean): 20nM
 - Most fluorescence parameters, growth, pigments,
 - First site of inhibition: photosynthetic apparatus
 - ROS generation due to photosynthesis malfunction
 - Direct inhibition of PSIIRC only pronounced in HL
- In LL binding of Cd to LHCII already below 20nM
 - Toxicity more pronounced under HL conditions
 - Chl in LHCII acts as buffer under LL
- Slightly higher but still nanomolar concentrations:
 - selective Cd binding to few enzymes like allantoinase also in roots → disturbance of root metabolism
- Migh micromolar concentrations
 - many unselective effects, also genotoxicity

All slides of my lectures can be downloaded from my workgroup homepage

Biology Centre CAS → Institute of Plant Molecular Biology → Departments

→ Department of Plant Biophysics and Biochemistry,

or directly

http://webserver.umbr.cas.cz/~kupper/AG_Kuepper_Homepage.html