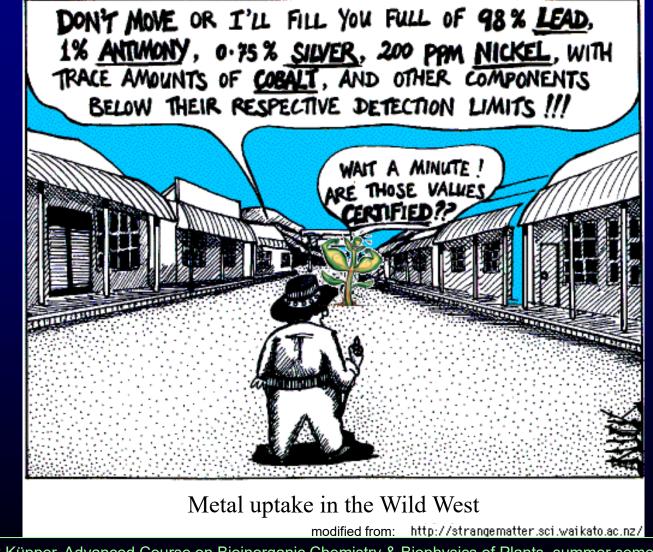
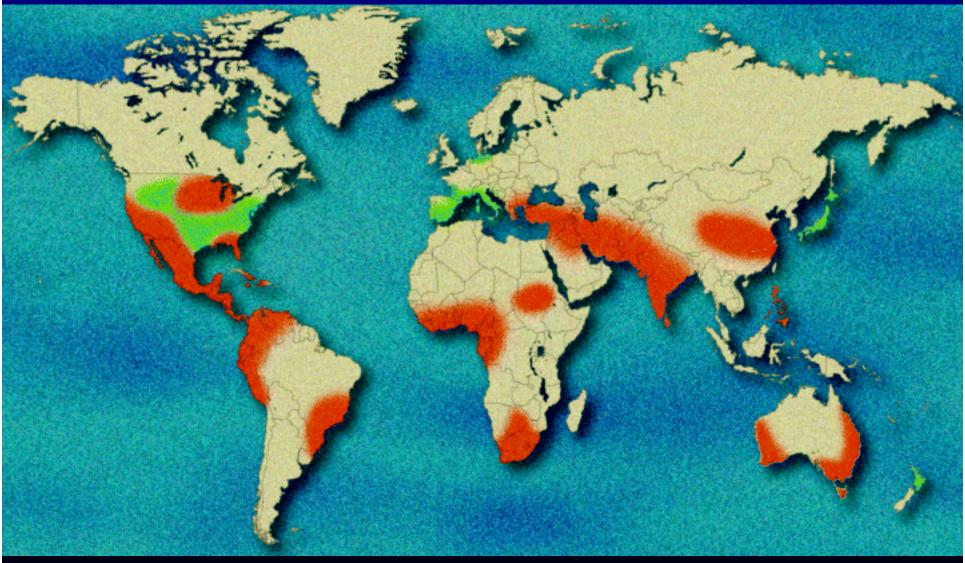
Trace metals as micronutrients



Hendrik Küpper, Advanced Course on Bioinorganic Chemistry & Biophysics of Plants, summer semester 2021

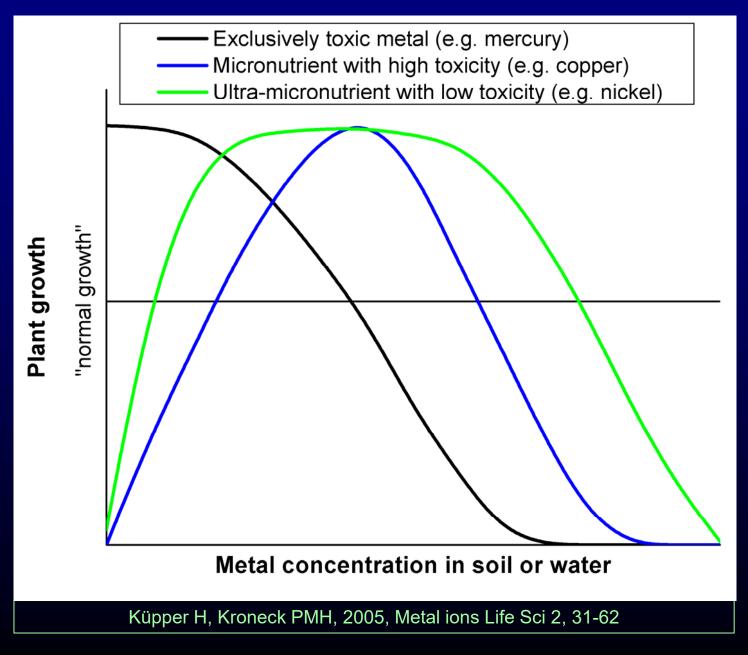
Heavy metal deficiency as a global problem of agriculture

green = moderate zinc deficiency; red = severe zinc deficiency



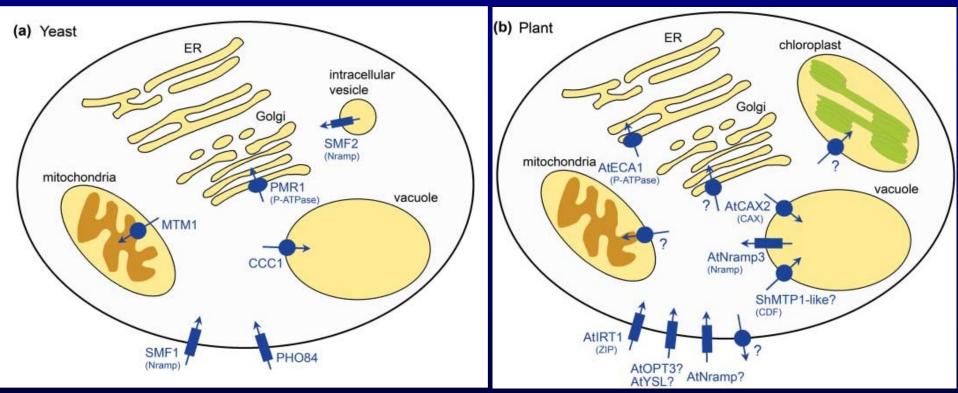
From: Alloway BJ. 2001. Zinc the vital micronutrient for healthy, high-value crops. Brussels, Belgium: International Zinc Association.

Dose-response principle for heavy metals



Mechanisms of metal uptake in Eucaryotes: Main families of metal transport proteins

example: manganese transport in yeast and plants

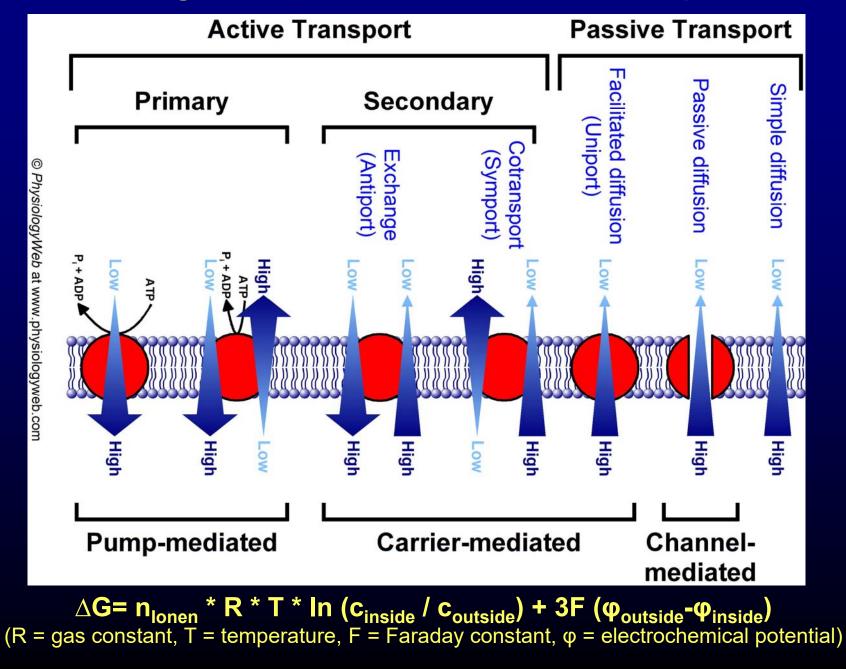


From: Pittman JK, 2005, NewPhytol167, 733-742

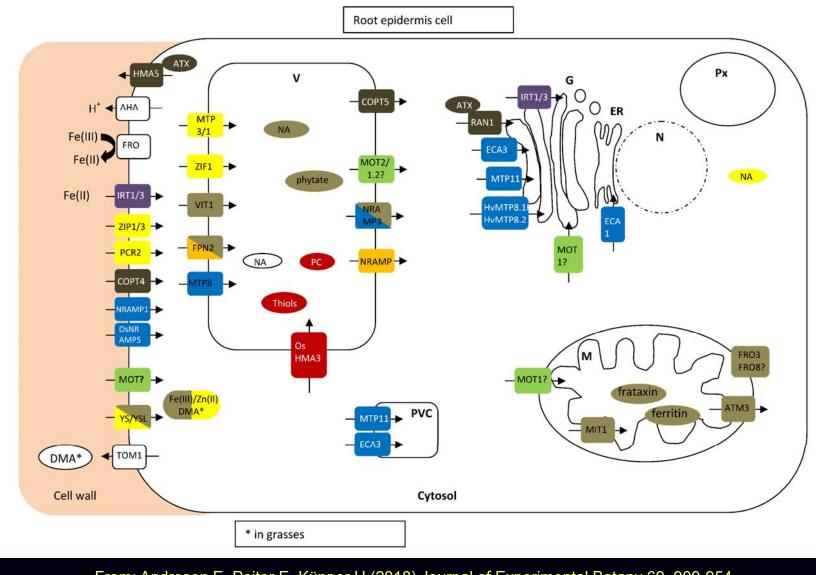
4 main families of transmembrane metal transport proteins

- P-type ATPases
- cation diffusion facilitators (CDF-transporters)
- ZRT-/IRT-like proteins (ZIP-transporters)
- > Natural resistance associated Macrophage proteins (Nramp-transporters)

Energetics and variants of metal transport

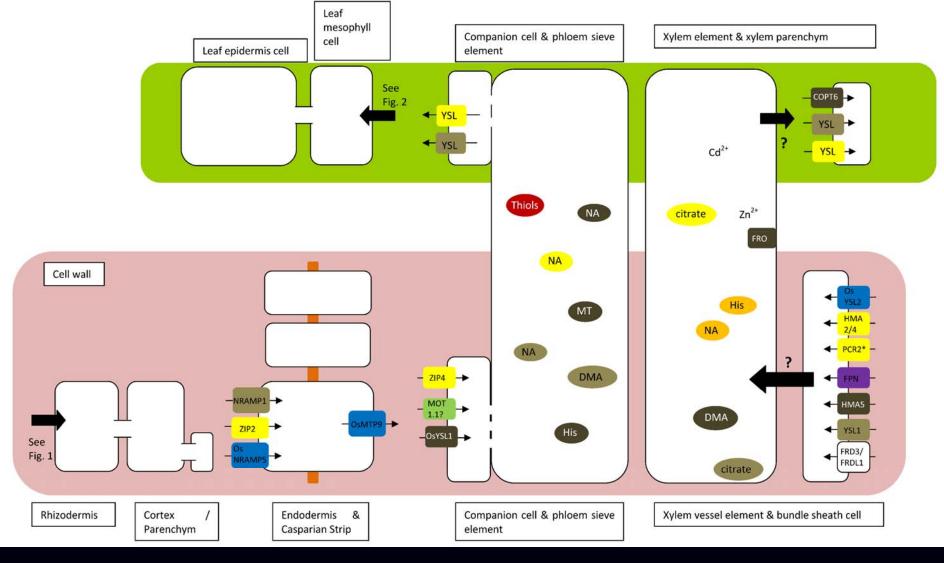


Mechanisms of metal uptake in plants: Different transport steps require different transporters 1) Root uptake and intracellular distribution



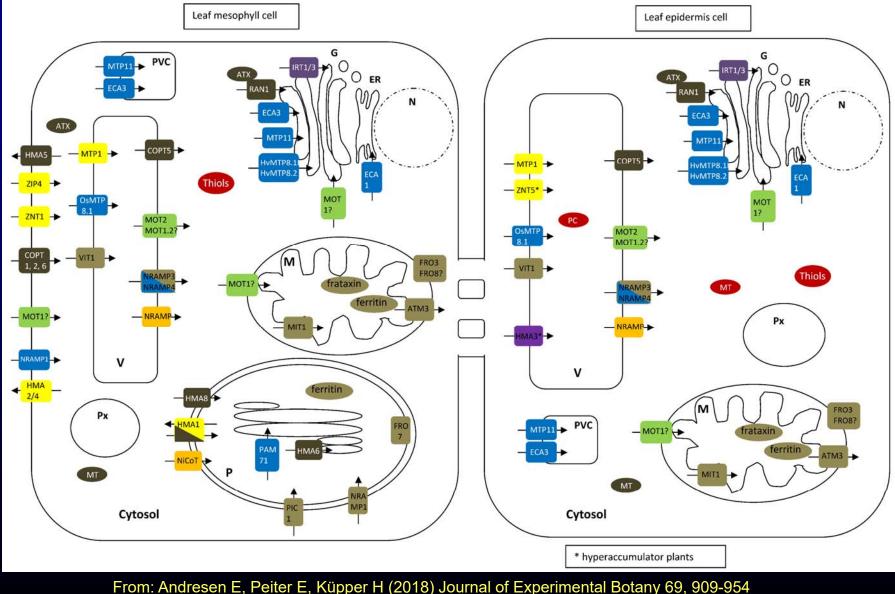
From: Andresen E, Peiter E, Küpper H (2018) Journal of Experimental Botany 69, 909-954

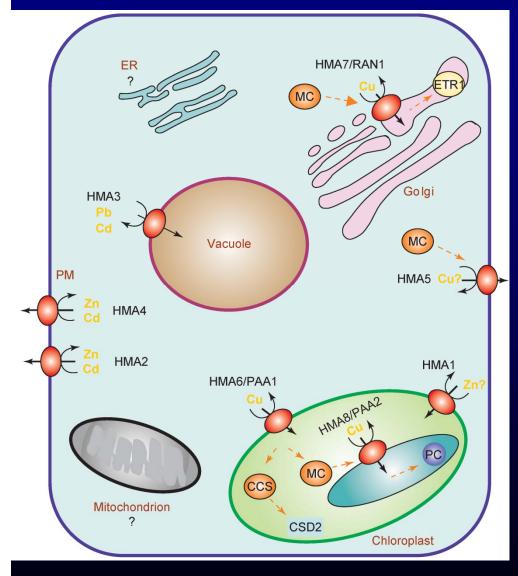
Mechanisms of metal uptake in plants: Different transport steps require different transporters 2) Translocation. Root-to-shoot: Xylem, shoot-to-root: phloem



From: Andresen E, Peiter E, Küpper H (2018) Journal of Experimental Botany 69, 909-954

Mechanisms of metal uptake in plants: Different transport steps require different transporters 2) Distribution in shoot cells



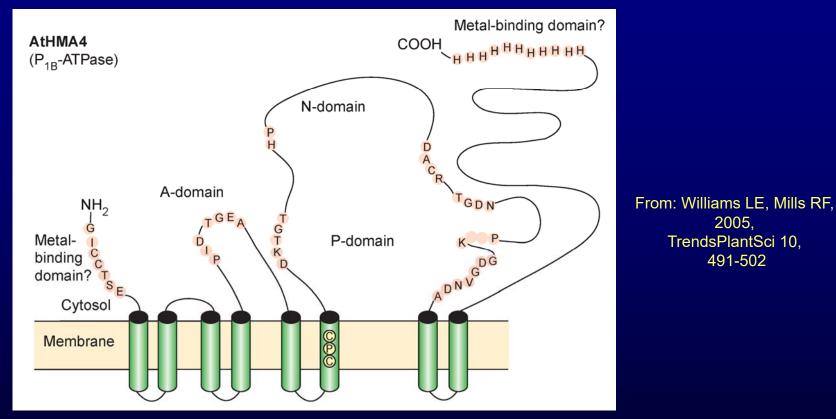


From: Williams LE, Mills RF, 2005, TrendsPlantSci10, 491-502

Functions (concluded mostly from differential expression studies)
> translocation into the root xylem, that means out of root cells (→ e.g. HMA4)
> xylem unloading in shoots
> intracellular metal sequestration e.g. in

the vacuole

 transport into the chloroplast, inside the chloroplast into the thylakoids
 transport into the Golgi apparatus

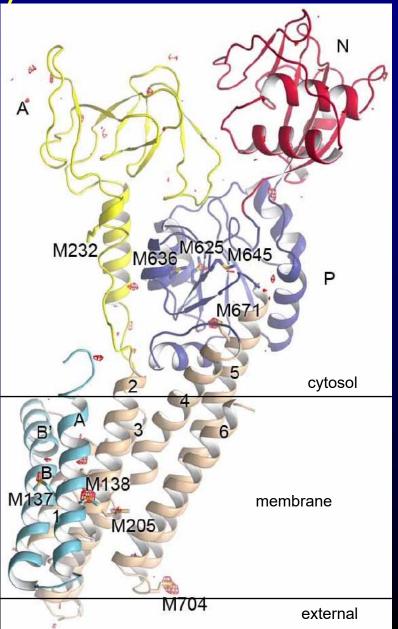


Sequence characteristics

- > CPx-motif in 6th transmembrane helix (\rightarrow Name!)
- Variable number of transmembrane helices
- > MANY histidines and cysteines in sequence (\rightarrow e.g. 58 Cys in TcHMA4)
- Metal binding domain at N-terminus (in cytosol)
- Histidine repeat at C-terminus (in cytosol)

Structural characteristics, from an X-ray structure of a bacterial protein

- Iarge cytosolic domain
- electronegative funnel connects membrane surface to high-affinity Zn-binding site
- the size and structure of the channel suggests that it interacts with aqueous Zn²⁺, not a larger complex
- high affinity Zn-binding site contains two cysteins

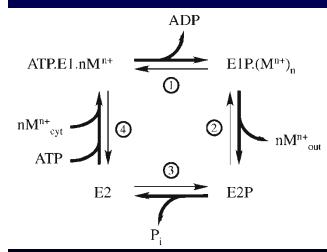


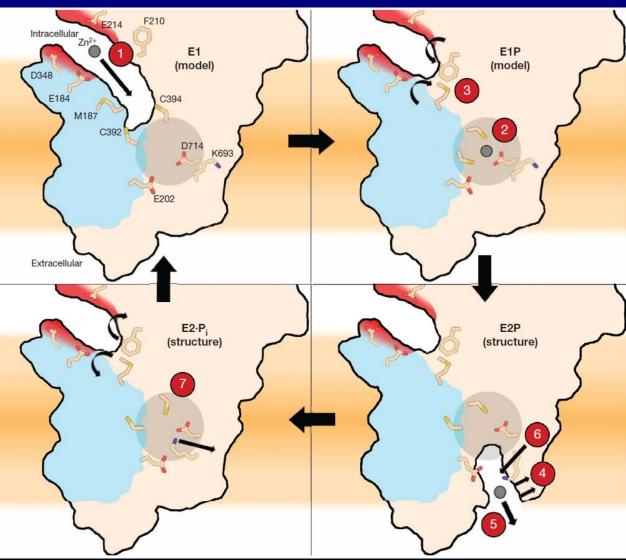
From: Wang K et al, 2014, Structure and mechanism of Zn-transporting P-type ATPases. Nature 514, 518-

Mechanism

Zn is guided into binding pocket by negatively charged residues

- binding pocket closes after ATP binding
- pore opens on other side of protein, release of Zn²⁺
- pore closes after ATP hydrolysis





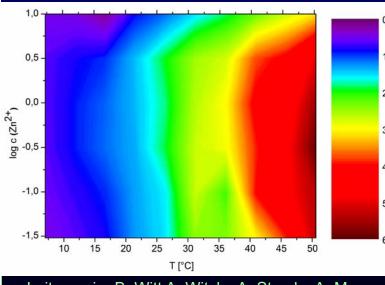
From: Argüello JM et al., 2007, Biometals, DOI 10.1007/s10534-006-9055-6

From: Wang K et al, 2014, Structure and mechanism of Zntransporting P-type ATPases. Nature 514, 518-

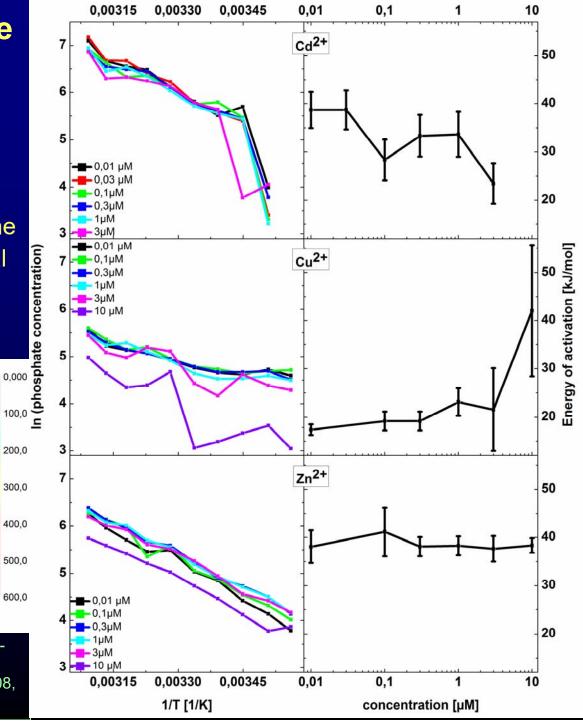
Activation and substrate inhibition of TcHMA4

- Activation energies for TcHMA4 (CPx = P_{1B} ATPase) are similar to other metal ATPases.

- Activation energy changes with the concentration and type of the metal to be pumped.

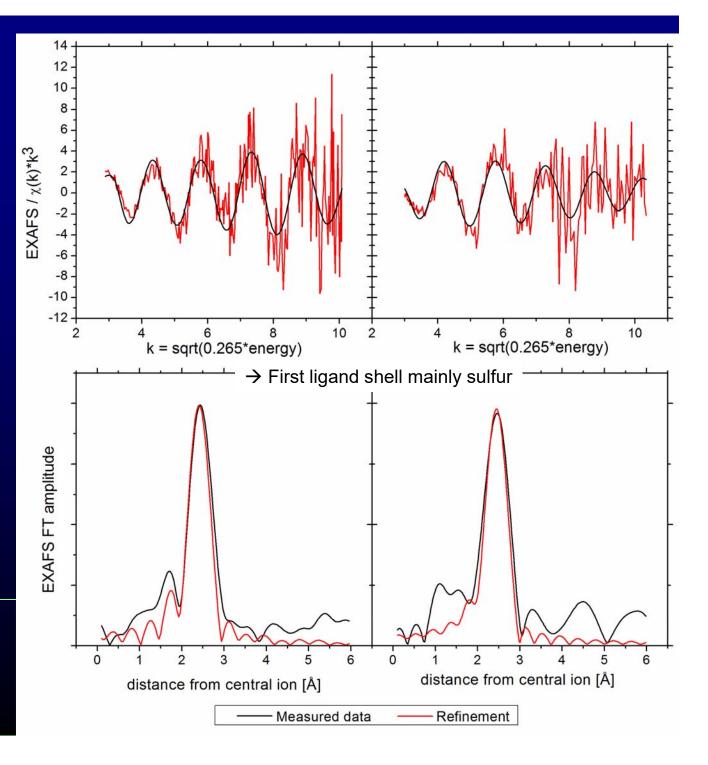


Leitenmaier B, Witt A, Witzke A, Stemke A, Meyer-Klaucke W, Kroneck PMH, Küpper H (2011) Biochimica et Biophysica Acta (Biomembranes) 1808, 2591-2599



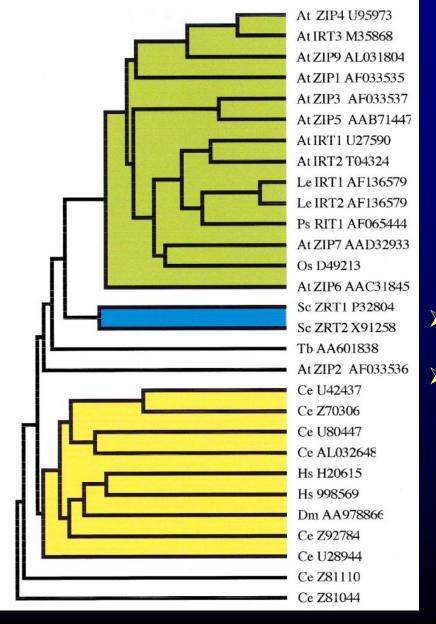
EXAFS-analysis of TcHMA4

- at low Cd concentrations, the first ligand shell in this ATPase consists mainly of S (thiol groups from some of the 58 cysteines in the sequence)



Barbara Leitenmaier, Annelie Witt, Annabell Witzke, Anastasia Stemke, Wolfram Meyer-Klaucke , Peter M.H. Kroneck, Hendrik Küpper (2011) Biochimica et Biophysica Acta (Biomembranes) – 1808, 2591-2599

Mechanisms of metal uptake in plants (II) ZIP-transporters

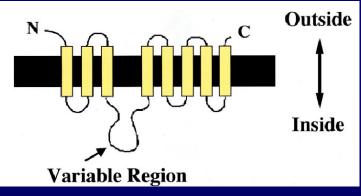


Likely Functions (deduced from expression studies)

uptake of metals into cells over the cytoplasmatic membrane
 abundant in all eucaryotes, incl. humans, plants and fungi

From:Guerinot ML, 2000, BBA 1465, 190-8

Mechanisms of metal uptake+compartmentation in plants (II) ZIP-transporters



From:Guerinot ML, 2000, BBA 1465, 190-8

Cd influx Cd inf

Functions suggested by expression studies)

- uptake of metals into cells over the cytoplasmatic membrane
- abundant in all eucaryotes, incl. humans, plants and fungi

Structure predicted by sequence

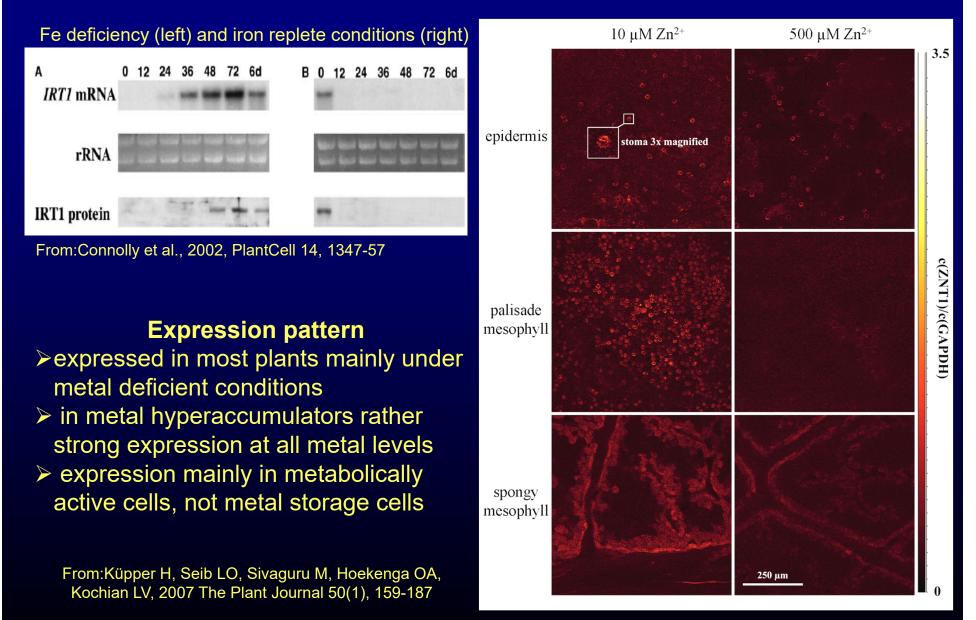
- usually 8 transmembrane helices, one long variable region, predicted to be in the cytoplasm
 309-476 amino acids
- still no complete 3D structure available

Characteristics revealed by yeast expression studies

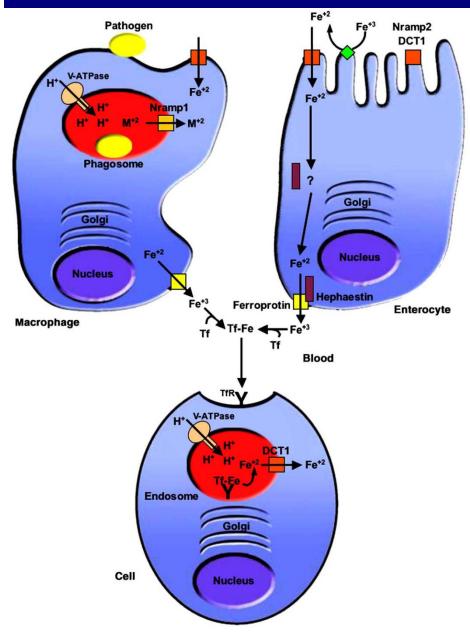
- High affinity and saturable kinetics for selected metal (e.g. Zn in ZNT1)
- Lower affinity uptake for related metals (e.g. Cd in ZNT1)

From:Pence NS et al., 2000, PNAS 97, 4956-60

Mechanisms of metal uptake+compartmentation in plants (II) Transcriptional regulation of ZIP transporters



Mechanisms of metal uptake in plants (III) Natural resistance associated macrophage proteins (Nramps)

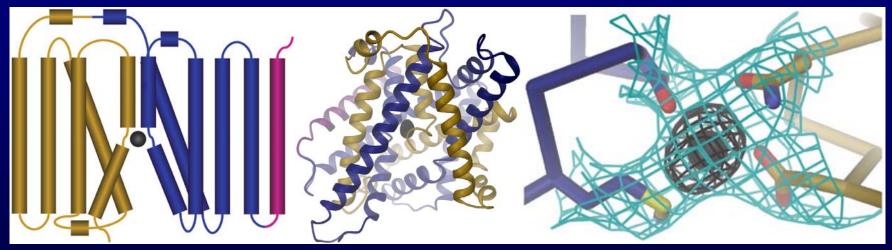


Likely Functions (deduced from expression studies)

- ➢Discovered to have a role in the immune response of animals (→ name!)
- abundant in all eucaryotes, incl. humans, plants and fungi
- predicted to play a role in uptake into the cell as well as trafficking inside the cell

From: Nevo Y, Nelson N, 2006, BBA 1763, 609-620

Mechanisms of metal uptake in plants (III) Natural resistance associated macrophage proteins (Nramps)

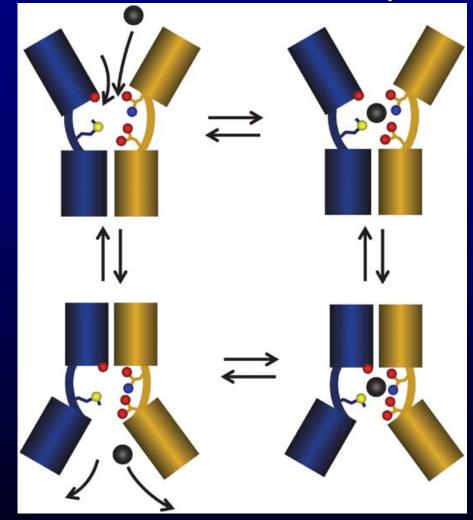


From: Ehrnstorfer IA, Geertsma ER, Pardon E, Steyaert J, Dutzler R. Nat Struct Mol Biol. 21, 990-6

Structure of a bacterial protein with high similarity to eukaryotic proteins

- > 11 transmembrane helices, 10 of them with conserved sequence
- Iong loops on both sides of the membrane
- > main metal binding site in the middle of transmembrane helices 1 and 6
- metal (in this case Mn²⁺) coordination in main binding site by one methionine-S and oxygens from alanine, aspartate and asparagine

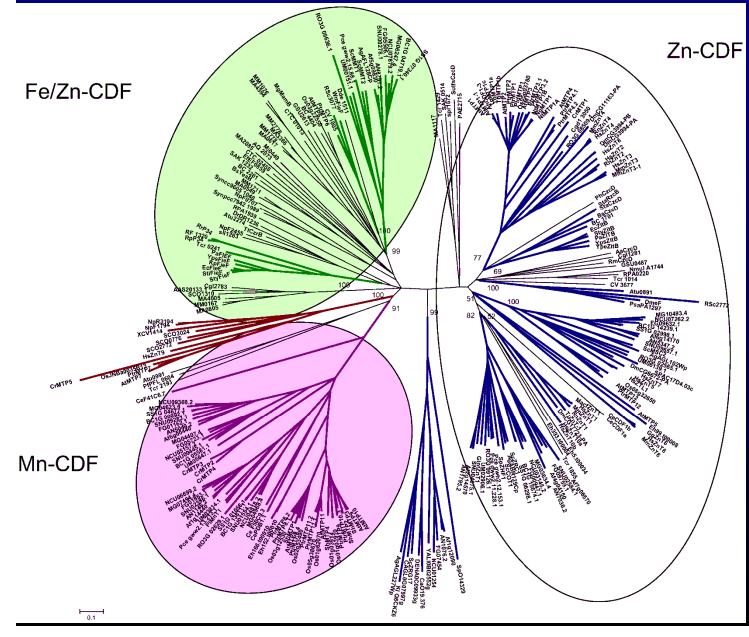
Mechanisms of metal uptake in plants (III) Natural resistance associated macrophage proteins (NRAMPs)



Mechanism deduced from crystal structure and enzyme kinetic studies

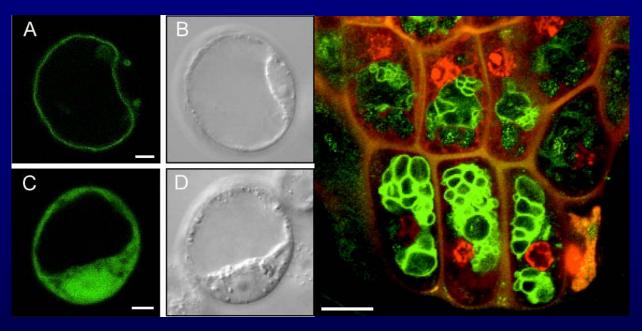
- proton symport with the electrochemical gradient drives metal translocation against the gradient
- binding of metal and proton induces a conformational change of the two halves of the helices 1 and 6 around a hinge in the metal binding site
- the conformational change closes the pore on the outer side and opens a pore on the inner side
- the opening of the intracellular pore releases metal and proton.

From: Ehrnstorfer IA, Geertsma ER, Pardon E, Steyaert J, Dutzler R. 2014. Nat Struct Mol Biol. 21, 990-6



 abundant in all eucaryotes, incl. humans, plants and fungi
 different subfamilies

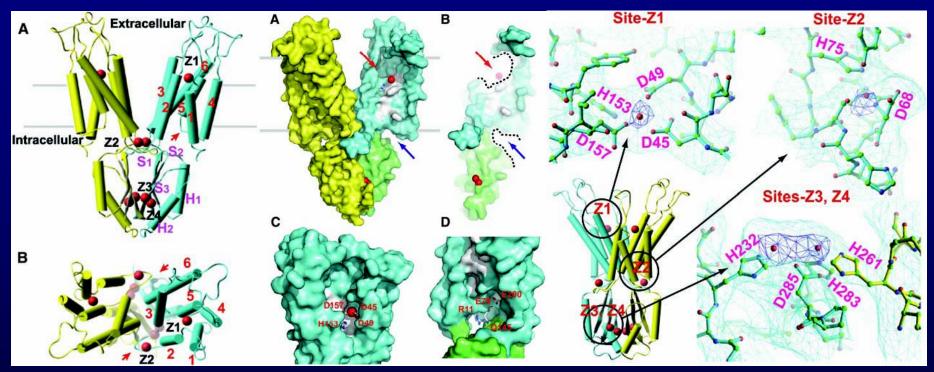
From: Montanini B et al., 2007, BMCGenomics 8, 107



From: Kobae et al., 2004, PlantCellPhysiol 45, 1749-58 From: Blaudez D et al., 2003, PlantCell 15, 2911-28

Functions concluded from expression studies (localisation and overexpression/knockout phenotypes)

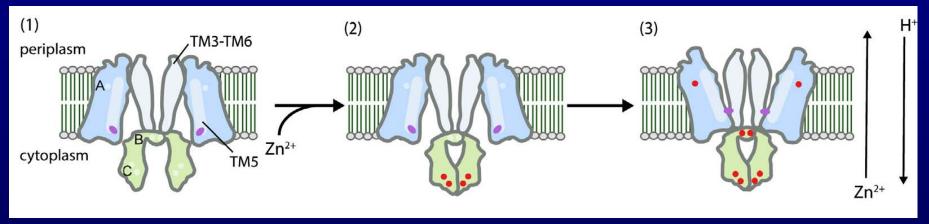
- Metal detoxification
- Sequester the metals in intracellular compartments (mainly vacuole)



From: Lu M, Fu D. 2007. Structure of the zinc transporter YiiP. Science 317, 1746-8

Structure of a bacterial Zn-transporting CDF (YiiP) similar to others

- MANY histidines in sequence used for metal binding
- 6 transmembrance helices per protein, active form is dimer held together by four Zn²⁺ in cytoplasmic domain
- > 2 further metal binding domains in the protein at both sides of the membrane

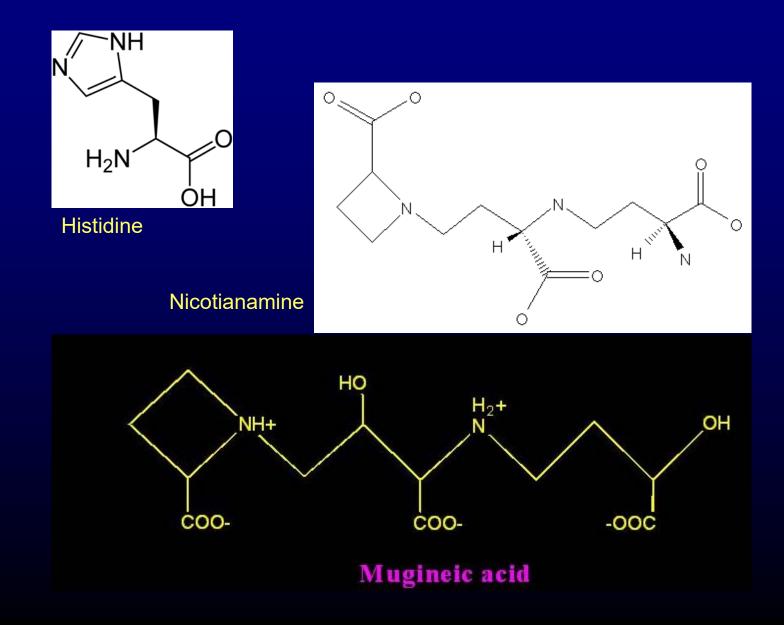


From: Kolaj-Robin O, Russell D, Hayes KA, Pembroke JT, Soulimane T. 2015. FEBS Lett. 589, 1283-95

Mechanism concluded from structure and kinetic studies

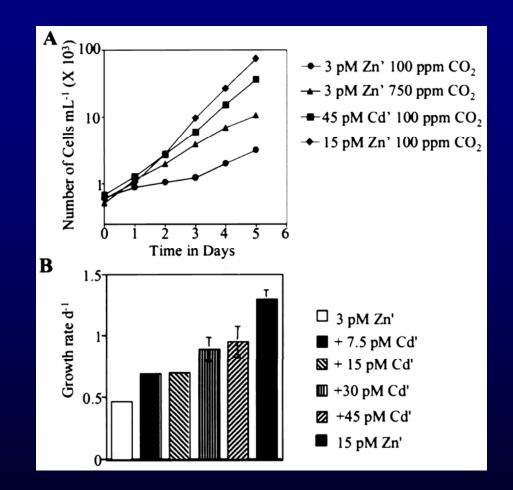
- Proton-metal antiport
- The exact movements are still discussed as the only available complete crystal structure is rather low resolution (3.8A)
- > Metal binding causes a conformational change of the cytoplasmic domain
- > The conformational change leads to release of the metals on the outside of the cell

Mechanisms of metal uptake in plants (V): Long-distance transport ligands



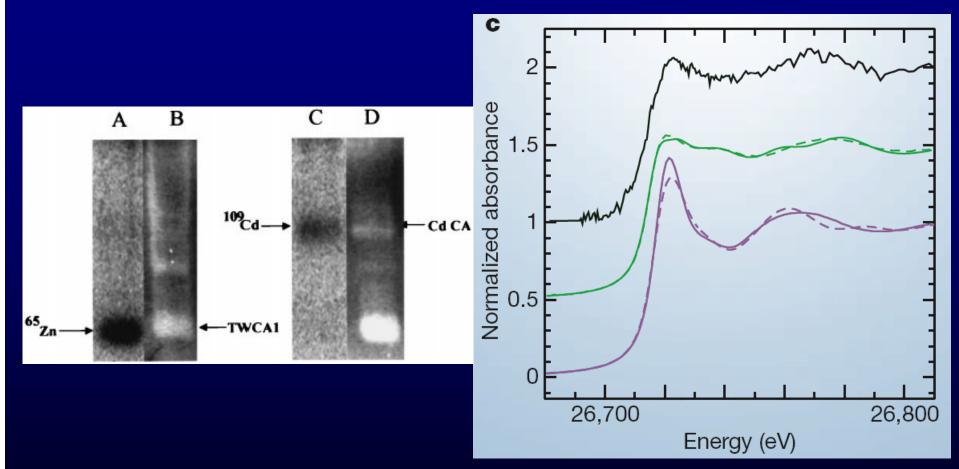
Now let's have a look at a few plant micronutrients

Cadmium as a micronutrient



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

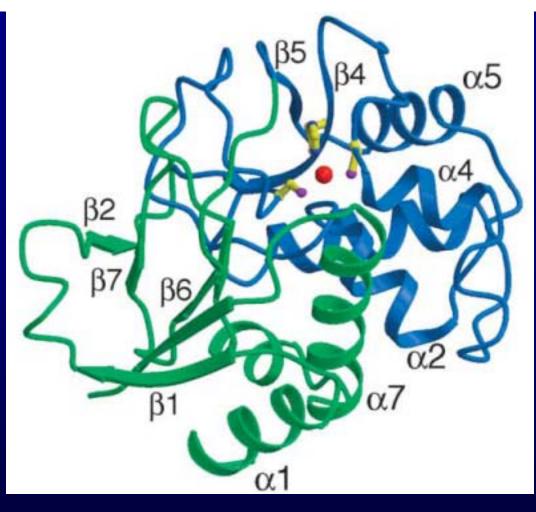
Carboanhydrase from *Thalassioria weissflogii*: An enzyme with cadmium in its active centre



Size of the cadmium-carboanhydrase in comparison to the normal Zn-carboanhydrase (Lane and Morel, 2000, PNAS Vol. 97) EXAFS-spectrum of the isolated Cdcarboanhydrase (Lane et al., 2005, Nature Vol. 435)

Properties and structure of the Cd-carboanhydrase

(Xu et al., 2008, Nature 452, pp 56-61)



• Cd-CA can bind both Cd and Zn. Activity with Zn somewhat higher, but activity with Cd much higher than for regular Zn-carboanhydrases.

•Cd-CA has 7 α -helices and 9 β -sheets, Cd at the lower end of a funnel-like binding pocket

•Cd²⁺ is bound via three conserved amino acid residues: 2x cystein and 1x histidin, plus 1x Water (\rightarrow tetrahedral coordination). Further fixed water molecules nearby

Cadmium deficiency in the Cd/Zn-hyperaccumulator Thlaspi caerulescens



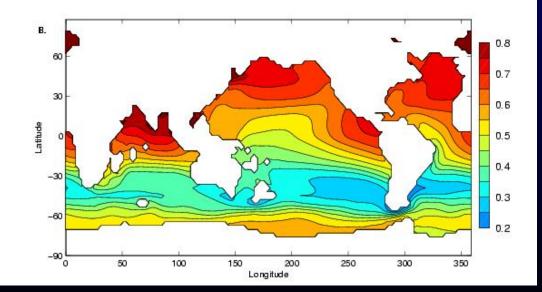
With 10 µM cadmium in the nutrient solution --> healthy plants

Without Cd in the nutrient solution --> damage by insects

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

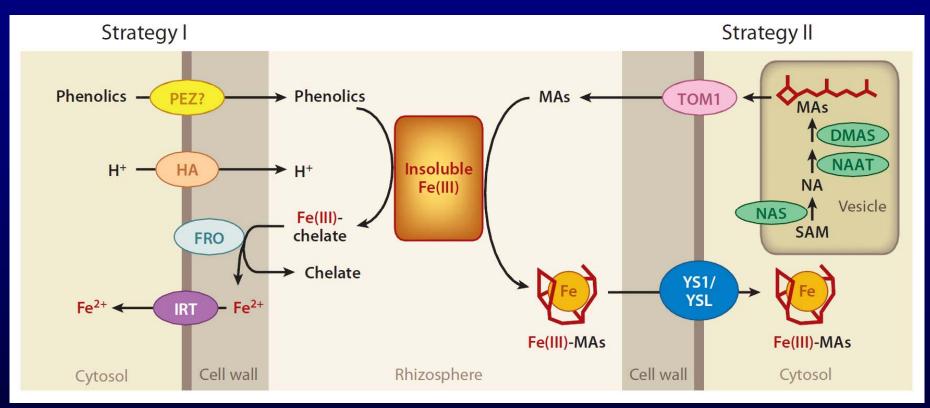
Iron

A. 60 0.8 30 0.6 Latitude 0.4 -30 0.2 -60 0 -90 L 0 100 150 350 50 200 300 250 Longitude



Iron concentrations at the surface (top picture) and in 1000m depth (bottom picture)

Mechanisms of iron uptake in plants Strategies of iron efficiency

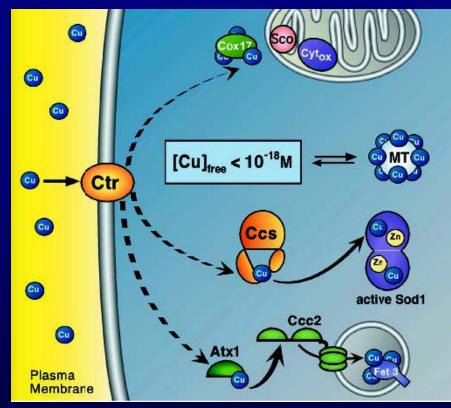


From: Kobayashi T, Nishizawa NK. 2012. Ann Rev Plant Biol 263, 131-152

Strategies of making insoluble Fe(III) bioavailable

- Strategy I (most plants): use mostly of soil acidification and iron reductase at root surface
- Strategy II (grasses): use of secreted iron ligand mugineic acid

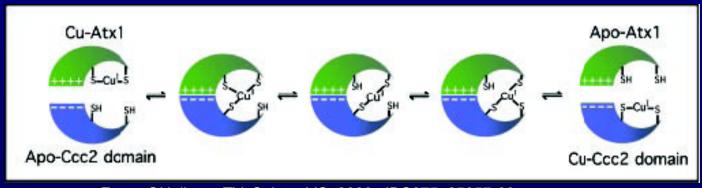
Copper delivery inside cellular compartments



From: OHalloran TV, Culotta VC, 2000, JBC275, 25057-60

Confusing large number of names for homologous proteins in different organisms
 REALITY: just 3 really different (non-homologous) Cu-chaperones are well known, some more proteins are postulated to be Cu-chaperones

Copper delivery to the Golgi and thylakoid: ATX1 = HAH1 = ATOX1 = CopZ ≈ CCH (a) occurrence

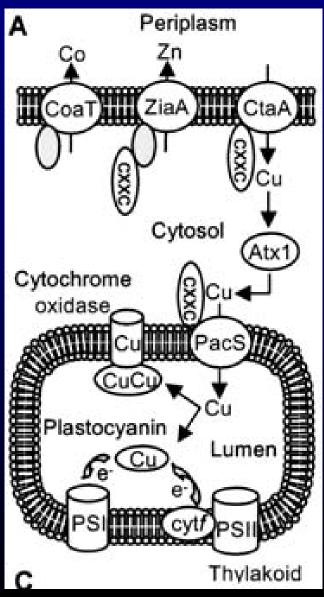


From: OHalloran TV, Culotta VC, 2000, JBC275, 25057-60

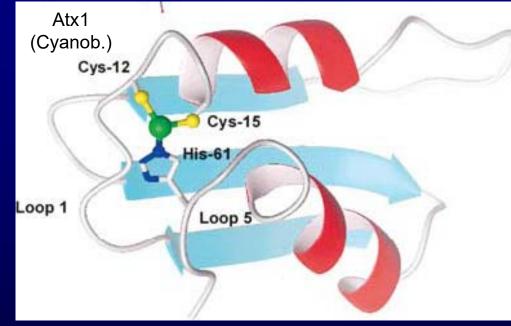
- ATX1 found in yeast originally as a gene involved in protection against oxidative damage;
- human homologue: HAH1 = ATOX1
- bacterial homologue: CopZ
- cyanobacterial and homologues: Atx1
- similar to plant CCH

Copper delivery to the Golgi and thylakoid: ATX1 = HAH1 = ATOX1 = CopZ (b) function in bacteria, cyanobacteria+plants

- CopZ in non-photosynthetic bacteria donates Cu to CopY transcription factor? Or Cu delivery to Cuefflux transport ATPase CopB or copper influx ATPase CopA?
- Atx1 found to specifically shuttle copper to an intracellular CPx-type copper ATPase the thylakoid in cyanobacteria+plants
- CtaA+PacS ATPases deliver Cu for plastocyanine across thylakoid membrane



Copper delivery to the Golgi and thylakoid: ATX1 = HAH1 = ATOX1 = CopZ = Atx1 (c) Cu-binding in bacterial+cyanobacterial+plant version

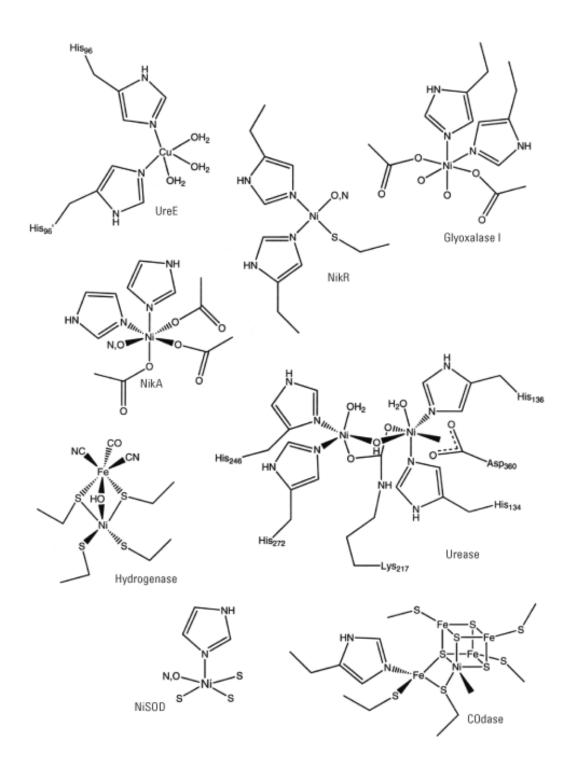


From: Borrelly GPM, et al., 2004, BiochemJ378, 293-7

>Atx1 binds a single Cu(I) ion like ATX1

Atx1: like in the yeast+animal proteins, Cu-binding via two Cys in the sequence MT/HCXXC, <u>BUT</u> additional histidine61 from loop 5

- ➤ the additional histidine shifts Atx1 binding affinity towards CtaA by reducing affinity for PacS → trafficking of Cu(I) from one CtaA to the PacS
- > other features like in yeast+animal proteins



Examples of nickel complexes in proteins

Characteristics

- Nickel is usually bound by nitrogen (mainly histidine), sulphur (cysteine) and oxygen ligands
- usually 5 (4-6) ligands

From: Carrington PE et al, 2002, EnvHealthPers110, 705-

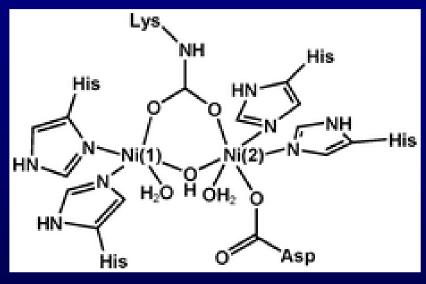
Best known (and most important?) Ni-enzyme: Urease a) Function and occurrence

$$H_{2}N \xrightarrow{0} H_{2} + H_{2}O \xrightarrow{\text{urease}} NH_{3} + H_{2}N \xrightarrow{0} OH$$

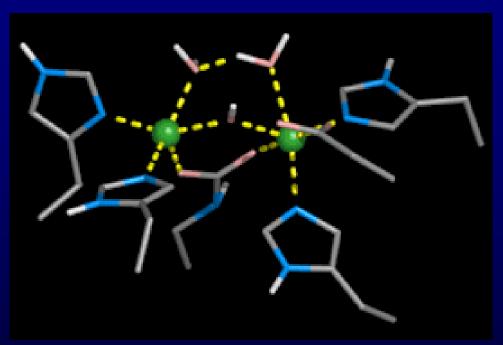
$$H_{2}N \xrightarrow{0} OH + H_{2}O \xrightarrow{0} NH_{3} + H_{2}CO_{3}$$

- Catalyses the decomposition of urea into carbon dioxide and ammonia
- In most organisms (plants, fungi, bacteria)
- Very important for metabolism: urea toxicity is one of the main mechanisms of damage caused by nickel deficiency in plants
- Very specific for urea as a substrate
- > Rather fast: turnover rate k_{cat} around 3,500 s⁻¹

Urease b) Active site



From: Lee WZ et al., 2008, DaltonTrans, 2538-41



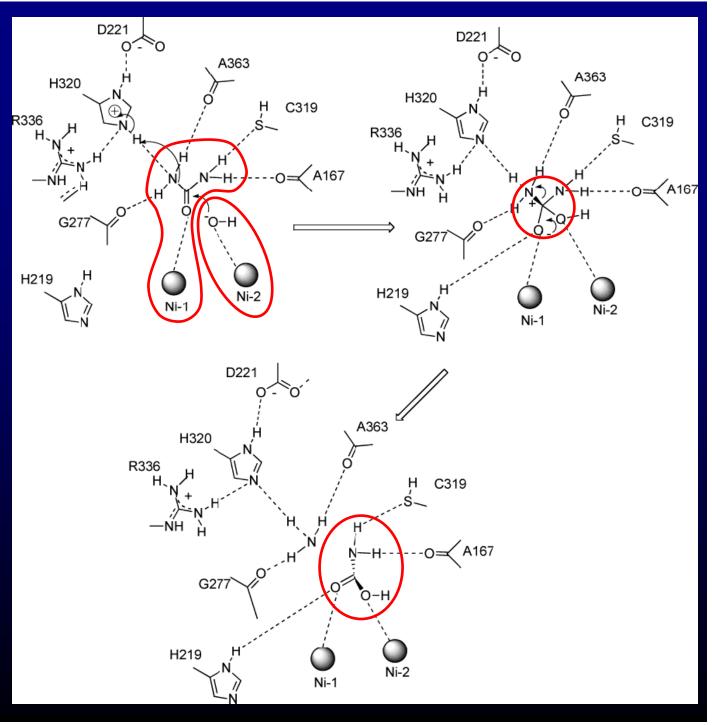
- ➤ two Ni²⁺ ions
- > one Ni²⁺ bound by three fixed ligands (2 His-N and 1 Lys-O) and one water
- the other Ni²⁺ bound by four fixed ligands (2 His-N, 1 Asp-O and 1 Lys-O) and 1 water
- > the two nickels are bridged by a water molecule

Urease c) Mechanism

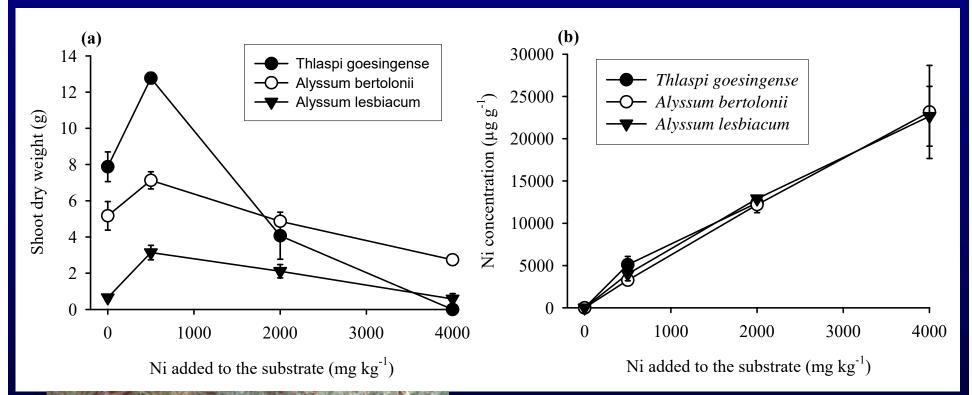
Steps

- Nickel1 binds urea at the oxygen
- Nickel 2 binds water
- tetrahedral intermediate
- after cleavage of the C-N bond, carbamic acid is bound to the nickels

From: Karplus PA, Pearson MA, Hausinger RP, 1997, Acc Chem Res 30, 330-37, modified by Estiu G, Merz KM, 2004, JACS126, 11832-42



Nickel deficiency in Ni-Hyperaccumulators

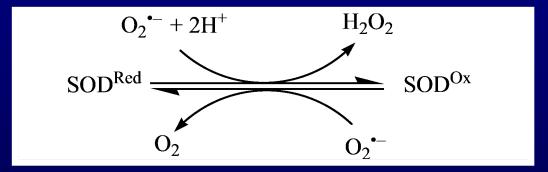




Alyssum lesbiacum

Küpper H, Lombi E, Zhao FJ, Wieshammer G, McGrath SP (2001) J Exp Bot 52 (365), 2291-2300

Nickel superoxide dismutase (a) Function and occurrence



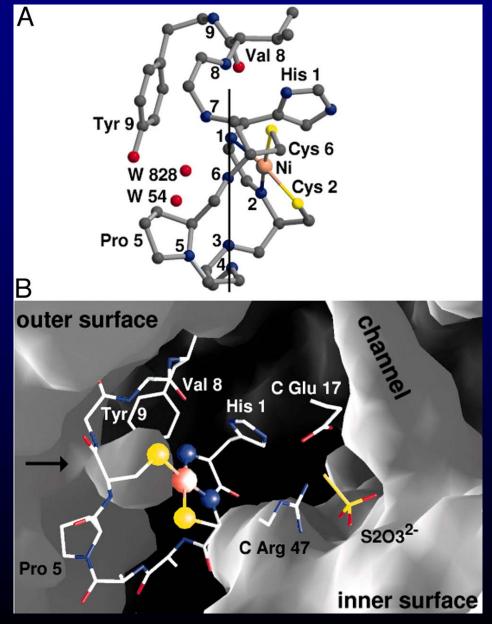
Characteristics

- Catalyses the detoxification of superoxide (O_2^-) by disproportionation into dioxygen (O_2) and hydrogen peroxide (H_2O_2)
- Ni-SOD is found in cyanobacteria and in Streptomyces (eubacteria), other SOD's are usually Cu/Zn, Fe or Mn enzymes

2. Nickel superoxide dismutase (c) active site

Characteristics

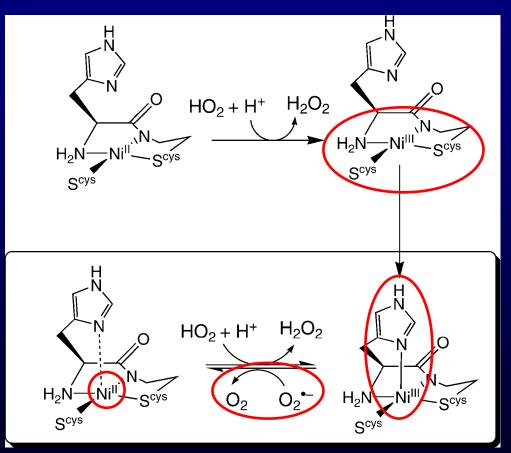
- Ni 5-6-coordinated, axial Nligand(s) artefactually (x-ray damage!) lost when Ni³⁺ is reduced to Ni²⁺ during x-ray data collection
- Ni coordination by the amino group of His-1, the amide group of Cys-2, and the thiolate group of Cys-2 and of Cys-6
- sulphur (thiolate) ligation makes otherwise biologically redox-indert nickel redox-active (2+/3+)
- ➤ Accessibility of active site limited by Pro-5 and Tyr-9 → specificity for small molecules!



2. Nickel superoxide dismutase (d) Mechanism

Steps and Characteristics

- Oxidation of the four-coordinate Ni²⁺ center to Ni³⁺ (unusual!)
- Rapid imidazole coordination. Once the imidazole is coordinated to the oxidized Ni³⁺ center it will remain ligated throughout catalysis
- ➢ Re-reduction of Ni³⁺ to Ni²⁺
- ➤ the axial H(1) imidazole enhances the activity of NiSOD e.g. by reducing structural reorganisation during catalysis (→ enhances speed!)



From: Neupane KP et al., 2007, JACS129, 14605-18

Zinc efficiency

Zn-inefficient

From: Hacisalihoglu G, Kochian, LV. How do some plants tolerate low levels of soil zinc? Mechanisms of zinc efficiency in crop plants. New Phytologist 159 (2), 341-350.

Zn-efficient

1II) Subcellular Zn Compartmentation, II) Zn Uptake and Translocation

IV) Biochemical Zn Utilization

I) Zn 🔳 **Bioavailability**

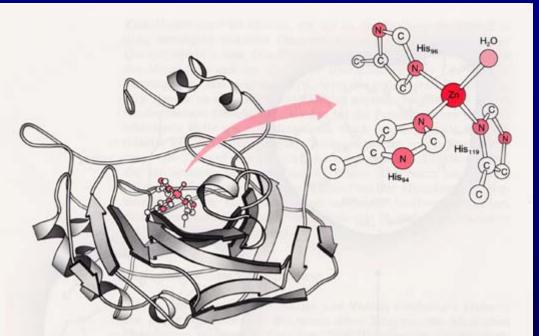
Emerging field: Role of trace metals in plant defence response to biotic stress

- Local Zn mobilization in response to pathogen Phomopsis in soybean roots revealed by µXRF imaging of living roots.
- Still unknown (ongoing work): regulatory mechanism, genes involved,...

Zn0.4 Ni Fe Zn Cu Mn Zn1.5 Zn Ni Fe Cu Mn Zn4 Zn Fe Cu Mn NI 300 µ

Morina F, Mijovilovich A, Koloniuk I, Pecnik A, Novak O, Gruz J, Küpper H (2021) Journal of Experimental Botany DOI: doi.org/10.1093/jxb/erab052

Selected important plant enzymes with zinc in their active centre



Carboanhydrase→ details in the lecture about photosynthesis related metal proteins

Zinc finger-motive

Tyrosin phosphatase

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