Cd Subcellular Localization in Root Tips of *Hordeum vulgare*

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Abstract

The effects of Cd on the subcellular localization in root tip cells of *Hordeum vulgare* were investigated by Energy dispersive x-ray analysis (EDXA) in order to further understand Cd toxic mechanisms in plants. EDXA showed that Cd ions were localized in meristem, elongation, and mature zone in the root tips. In transverse section of the mature zone, Cd was accumulated in epidermal, cortical, and vessel cells, and the level of Cd is in the order: epidermal cells < vessel cells < cortical cells. In cortical cells Cd ions were observed in cytoplasm and walls.

Keywords: *Hordeum vulgare*, cadmium (Cd), subcellular localization, root

Introduction

Naturally, cadmium (Cd) is present in trace concentrations and is a widespread heavy metal released into the environment by natural sources, agriculture, and manifold industrial uses [1]. Cd accumulation in the environment now has a major environmental and human health problem [2]. Cd is a non-essential element for plant growth and can be dangerous for plants. Cd toxicity in plants is well documented, including inducement of low mitotic index and pycnosis, disturbance of mitosis [3-5], and inhibition of plant root and shoot growth [6-7].

Cd can be absorbed and accumulated in plant tissues where roots are the primary site of accumulation [6, 8]. Plant roots are one of the organs most sensitive to environmental stress. After Cd is absorbed by roots, they can be deposited in different tissues. Knowledge about its deposition and distribution in tissue compartments can be gained through energy-dispersive x-ray analyses (EDXA). However, only a few investigations have been carried out using EDXA to localize intracellular Cd.

Higher plants such as *Allium cepa, Vicia faba, Arabidopsis Thailand*, and *H. vulgare* provide a useful genetic system for screening and monitoring environmental pollutants [9-11]. *H. vulgare* is well known as an excellent model plant and a useful biomarker for detecting heavy metal pollution in laboratories [11]. The aim of our study was to further understand Cd subcellular localization in root tip cells of *H. vulgare* under Cd stress by means of electron microscopy with EDXA. The data obtained here are useful for better understanding the mechanisms of Cd-induced cell toxicity.

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**Methods**

**Culture Condition and Cadmium Treatment**

Healthy equally-sized seeds of *H. vulgare* were collected and soaked in distilled water for 24 h before starting the experiments. After the seeds were germinated in moistened gauze in the dark at 23ºC for 12 h, they were treated with 50 μM Cd for 48 h. Cadmium chloride (CdCl₂) was used in the present investigation.

**Fluorescence Labelling of Cd**

*H. vulgare* intact roots treated with or without 50 μM Cd for 48 h were stained using the Cd-specific probe Leadmium™ Green AM solution (Molecular Probes, Life Technologies, California, USA) according to the manufacturer’s instructions to visualize the Cd absorption and distribution. Roots were put into 20 mM Na₂-EDTA for 15 min at room temperature, and then they were washed with the ddH₂O three times (3×10 min). A stock solution of Cd-specific probe was made by adding 50 μL of dimethyl sulfoxide into a vial of the dye. This stock solution was then diluted 1:10 with 0.85 % NaCl. The roots were immersed in the diluted stock solution at 37ºC for 2 h in the dark, and then were washed with 0.85 % NaCl three times. Then the roots were stored in the dark at 4ºC until fluorescence from the labeled Cd was visualized under a confocal laser scanning microscope (Nikon ECLIPSE 90i) using an exciter at 488 nm and a barrier at 590/50 nm.

**Sample Preparation for Scanning Electron Microscope**

Elemental distribution and composition of experimental plants was determined from samples of freeze-dried root materials. Ten root samples of 1 cm

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**Fig. 1.** EDXA spectra taken from the site of the SEM micrographs, showing Cd localization in different zones of root tip cells in *H. vulgare* exposed to 50 μM Cd for 48 h: A) Intact root. Image was taken at X4 magnification (Scale bar = 100 μm); B) Mature zone (Scale bar = 100 μm); C) Elongation zone (Scale bar = 100 μm); D) Meristem zone (Scale bar = 20 μm). Mt = meristem zone, E = elongation zone, M = mature zone. Scale bar = 100 μm. “*” site of the analysis; x-axis - energy [keV].
length were cut from the root tips of H. vulgar, exposed to 50 μM Cd for 48 h, and rapidly frozen in liquid nitrogen and lyophilized. Cross-sections of the roots (about 1 mm) were coated by gold using a sputter/coater (EMITECH K550X, Quorum Group, England). The energy-dispersive x-ray microanalytical studies were carried out using scanning electron microscopy (FEI Nova NANOSEM 230, Oregon, USA) provided with energy dispersive x-ray spectrum analysis (EDXA) (Genesis Apollo 10, EDAX, USA). The spectra were collected at 20 KV in an x-ray and x-ray detector equipped with a super ultra thin window. The collection time of spectra was 30-40s. The Cd content in the roots was shown by wt% (weight percent), which means concentration percent on the basis of weight (or mass) of a particular element relative to total element weight (or mass).

**Results**

The results from EDXA showed subcellular localization of Cd on the surface of root tip cells in H. vulgar exposed to 50 μM Cd for 48 h. The level of Cd in the different regions (meristem, elongation, and mature zone) was in the order: elongation zone (1.54 Wt%) > meristem zone (1.36 Wt%) > mature zone (0.62 Wt%) (Fig. 1A-D). In the transverse section of the meristem the protoderm, ground meristem, and procambium can be distinguished in very close proximity to the apical meristem (Fig. 2A). The Cd content was in the order: procambium (1.61 Wt%) > protoderm (1.33 Wt%) > ground meristem (1.29 Wt%) (Fig. 2B-D). In transverse section of the mature zone, epidermis, cortex and vascular cylinder can be easily distinguished from each other (Fig. 3A). The level of Cd in epidermal cells (0.54 Wt%) was low in comparison with cortical and vessel cells (Fig. 3B, E). In cortical cells more Cd ions were observed in the cytoplasm (2.92 Wt%) than those in the cell walls (1.86 Wt%) (Fig. 3C-D). In the vascular cylinder we observed that the Cd concentration in vessel walls (2.7 Wt%) was high when compared with the adjacent parenchyma cells (1.67 Wt%) (Fig. 3E-F). From a transverse section point of view, the Cd level in mature zone was high in comparison with the meristem zone.

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**Fig. 2.** EDXA spectra taken from the sites of the SEM micrographs, Cd localization in meristem zone in root tip cells of H. vulgar exposed to 50 μM Cd for 48 h: A) Transverse section of meristem zone (Scale bar = 100 μm); B) Protoderm (Scale bar = 30 μm); C) Ground meristem (Scale bar = 30 μm); D) Procambium (Scale bar = 20 μm). **"site of the analysis; x-axis - energy [keV]. G = ground meristem, Pc = procambium, P = protoderm.**
Discussion

The roots are the sole organ penetrating the soil, directly contacting with toxic metals. It is expected that they are the first to undergo toxicity and develop the responding symptoms. Therefore, evaluating the action mechanisms of Cd toxic to plant root tip cells and their consequences on Cd localization and genetic material, as performed in the present investigation, is very important.

EDXA as an analytical technique is very useful for analyzing localization of elements in biological specimens at the subcellular level. Cd ions were observed in elon-

Fig. 3. EDXA spectra taken from the sites of the SEM micrographs, Cd localization in mature zone in root tip cells of *H. vulgare* exposed to 50 μM Cd for 48 h: A) Transverse section of mature zone (Scale bars = 100 μm); B) Epidermal cells (Scale bar = 50 μm); C–D) Cortical cells; C) Cytoplasm; D) Cell wall (Scale bars = 50 μm); E) Vessel cells (Scale bar = 20 μm); F) Parenchyma cells (Scale bar = 20 μm). **“site of the analysis; x-axis - energy [keV]. C = cortex, E = epidermis, V = vascular cylinder.**
gation, meristem, and mature zone in root tips of *H. vulgare* exposed to Cd in the present investigation. From a transverse section point of view, the Cd level in the mature zone was high in comparison with the meristem zone due to its perfect conducting system. Root hairs of barley play an important role in the acquisition of Cd from soil. These results are consistent with those of Zheng et al. [12] and Balestri et al. [13], who observed that root hairs contribute significantly to Cd uptake in barley. Root hairs are found only in the maturation region of the root. They are outgrowths and have a large surface area, which makes absorbing both water and minerals more efficient using osmosis. The abundant channels and/or transporters in the plasma membrane of root hair cells can facilitate the uptake of Cd. In addition, hairy roots can contribute to the hyperaccumulation of Cd in root cell walls of *Noccaea* (Thlaspi) caerulescens [14]. Thus, the higher root hair-mediated Cd influx may be a key contributor to the hyperaccumulation of Cd [15].

The results here revealed that the level of Cd in epidermal cells of the mature zone in root tips of *H. vulgare* exposed to Cd was low when compared with cortical and vessel cells. We suggest that Cd ions in the epidermal cells after Cd stress may be absorbed quickly by root hairs, then more Cd ions are accumulated in cortical cells. This result is in conflict with the findings of Seregin et al. [16]. More studies, however, are required in this direction.

The limited translocation of Cd from cortex into the vessel cells is believed to result from the barrier function of the root endodermis: its Casparian strips bar Cd entrance into the central cylinder [17]. The results from Seregin and Ivanov [15] showed that the insignificant metal translocation into the stele are probably related to the Casparian strip barrier in the endodermal cells, suggesting that predominant metal accumulated in all tissues of the root apex as the Casparian strips, and had not as yet developed in the dividing and elongating cells. At nonlethal concentrations Cd does not practically reach the stele because the Casparian strips were already formed in the root hair zone [16]. However, it is not known whether the Casparian strips are the only limiting factor restricting Cd transport across the endodermis into the central cylinder tissues. The results of this investigation indicated that Cd is bound to cell walls in root tips of barley, which is consistent with the findings of other workers, suggesting that cell walls are the main site of Cd accumulation [6, 18-20]. We suppose that once excessive Cd ions enter the cytoplasm, a defence mechanism becomes activated, protecting the cells against Cd toxicity. The root is the first plant tissue in direct contact with metal ions in the growth medium and plays a major role in metal tolerance [21]. Cd accumulation due to metal exclusion in the root should theoretically be the best strategy for defense against Cd toxicity. As the first barrier to prevent metals from entering a cell, the cell wall is a pivotal site for Cd storage in plants [22-23]. Recent findings have demonstrated that in order to enhance its heavy-metal accumulation capacity the cell wall is even actively modified, which protects the protoplast. So metal-exposed plants can increase the amount of polysaccharides that bind heavy metal ions [24]. Santos et al. [25] found that Cd increased cell wall thickness in *Gracilaria domingensis* treated with 100, 200, and 300 μM of Cd for 16 d, suggesting that such an increase in cell wall thickness can be interpreted as a defense mechanism against exposure to Cd [26]. It is probable that the activity of Dictyosomes more intense in plants exposed to Cd and that this results in the large production of vehicles, which then format the matrix content of the cell wall [26]. Cytochemical evidence also confirms that cysteine-rich proteins, commonly referred to as phytochelatins, are localized in electron-dense granules in root cell walls of *A. cepa* [27], which seems to contribute substantially to Cd detoxification.

In cortical cells more Cd ions were observed in the cytoplasm than those in the cell walls, suggesting that cytoplasm is one of the main storage sites of Cd in the root tip cells of *H. vulgare*, which supports the findings of Liu and Kottke [27], Liu et al. [20], and Ali et al. [28]. In the cytoplasm, vesicles arising from the endoplasmic reticulum and dictyosomes also contain Cd deposits. Usually, several vesicles are gradually fused together, producing a bigger cytoplasmic vacuole. The increased vacuolation in Cd-exposed cells can prevent the circulation of free Cd ions in the cytoplasm and forces them into a limited area [27, 29], which can reduce Cd toxicity. Cd remaining in the cytoplasm could be detoxified in the cytoplasm by high-affinity ligands, which is considered to be an important mechanism of heavy-metal detoxification and tolerance [30].

**Conclusions**

Based on the information provided in this article, it is concluded that 1) Meristem, elongation, and the mature zone are the main sites of Cd uptake and 2) Cytoplasm and walls are the main Cd storage sites.

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**References**


