The SbMT-2 Gene from a Halophyte Confers Abiotic Stress Tolerance and Modulates ROS Scavenging in Transgenic Tobacco

Amit Kumar Chaturvedi, Manish Kumar Patel, Avinash Mishra*, Vivekanand Tiwari, Bhavanath Jha*

Abstract

Heavy metals are common pollutants of the coastal saline area and Salicornia brachiata an extreme halophyte is frequently exposed to various abiotic stresses including heavy metals. The SbMT-2 gene was cloned and transformed to tobacco for the functional validation. Transgenic tobacco lines (L2, L4, L6 and L13) showed significantly enhanced salt (NaCl), osmotic (PEG) and metals (Zn++, Cu++ and Cd++) tolerance compared to WT plants. Transgenic lines did not show any morphological variation and had enhanced growth parameters viz. shoot length, root length, fresh weight and dry weight. High seed germination percentage, chlorophyll content, relative water content, electrolytic leakage and membrane stability index confirmed that transgenic lines performed better under salt (NaCl), osmotic (PEG) and metals (Zn++, Cu++ and Cd++) stress conditions compared to WT plants. Proline, H2O2 and lipid peroxidation (MDA) analyses suggested that the role of SbMT-2 in cellular homeostasis and H2O2 detoxification. Furthermore in vivo localization of H2O2 and O2−; and elevated expression of key antioxidant enzyme encoding genes, SOD, POD and APX evident the possible role of SbMT-2 in ROS scavenging/detoxification mechanism. Transgenic lines showed accumulation of Cu++ in root and Cd++ in root while Zn++ in stem under stress condition. Under control (unstressed) condition, Zn++ was accumulated more in root but accumulation of Zn++ in stem under stress condition suggested that SbMT-2 may involve in the selective translocation of Zn++ from root to stem. This observation was further supported by the up-regulation of zinc transporter encoding genes NtZIP1 and NtHMA-A under metal ion stress condition. The study suggested that SbMT-2 modulates ROS scavenging and is a potential candidate to be used for phytoremediation and imparting stress tolerance.

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* Email: avinash@csmcri.org (AM); bjha@csmcri.org (BJ)

Introduction

Abiotic stresses such as salinity, drought, temperature and heavy metals have significant effect on the agricultural production over the years [1]. Most often, plants may encounter these abiotic stresses simultaneously, resulting in the substantial loss in agricultural productivity. Over the past two centuries, increased industrial and anthropogenic activities viz. mining, irrigation with sewage effluents/waste waters, use of phosphate fertilizers are the major sources of metal contamination to soil [2-3]. Salinity, drought and heavy metals have similar consequences ensuing oxidative stress, disturbances in ionic homeostasis and generation of reactive oxygen species (ROS) [1,4].

Metallothioneins (MTs) are the group of polypeptides which can bind with heavy metals through their thiols via chelation and involved in the homeostasis of essential metals (Cu and Zn) and cellular detoxification of nonessential metals (Cd and Hg) [5,6]. Characteristics of metallothioneins are the cysteine (Cys) residue which is the basis of its classification. Based on the arrangement of Cys residues MTs are divided into two classes [7]. Plant MTs belong to Class II and further classified into four types (type 1 to 4) based on the position and allocation of cysteine residues [7]. Spatial expression of all four MT type has been reported which is localized to root, stem, leaves and developing seeds [7,8] and show differential tolerance to different metals. Although there are several reports regarding the possible role of MTs in plants but its physiological role is still not fully conclusive because of difficulties in its isolation and stability [7,9]. Over-expression of MTs in various model systems like Arabidopsis, tobacco, yeast and E. coli established its functional role in homeostasis and tolerance to Cu, Zn and Cd [10,11,12,13], high salinity, drought, low temperature, heavy metal ions, abscisic acid (ABA) and ethylene [12,14]. Besides above, MTs are reported in the inhibition of root elongation [15], fruit ripening, seed development [7] and provide disease resistance against pathogen attack [16].

Heavy metals affect the plant at cellular, biochemical and molecular level causing the oxidative stress. These toxic metals generate free radicals and reactive oxygen species (ROS) which damages the cell membrane, nucleic acids and photosynthetic pigments [17]. The equilibrium between ROS generation and quenching is prerequisite for the cell survival which is maintained by the intricate anti-oxidative system comprising of enzymatic [superoxide dismutase (SOD), catalase (CAT), peroxidase (POD)],...
and non-enzymatic (ascorbate, glutathione and phenolic compounds) systems [4,12,17,18]. Under stress conditions the activity of this antioxidative system gets increased due to the increase of free radical formation [17].

Though MTs are known to be involved in abiotic/metal stresses and have been reported from several plant species, there are a few reports on halophytes under heavy metal stress [19]. *Salicornia brachiata* is an extreme halophyte and frequently exposed to heavy metals in coastal areas. The plant has nutritional value, unique oligosaccharide profile and requires NaCl for *in vitro* regeneration [20–22]. The plant is considered as a model for the study of tolerance mechanism and several stress responsive genes have been isolated and characterized [13,23–28]. In our previous study, *SbMT*-2 gene was isolated and physiological role was determined in *E. coli* which showed its role in the homeostasis and detoxification of Zn, Cu and Cd ions [13]. The gene (*SbMT*-2) was considered as a potential candidate to be utilized for the genetic engineering of plants for phytoremediation of heavy metals and stress tolerance. Therefore in the present study *SbMT*-2 gene was transformed into tobacco plants. Fourteen independent transgenic lines (T0) were raised and preliminarily screened by PCR using gene specific primers (data not shown). These lines were grown in containment facility and seeds (T1) were collected. T1 Seeds were germinated on hygromycin containing media and selected T1 transgenic lines were confirmed by PCR using gene specific primers (Figure 1b). Expected size of amplicon was found in all lines except L11. Based on *SbMT*-2 (Figure 1c) gene expression level, four independent T1 transgenic lines; L2, L4, L6 and L13 were selected for the further morpho-physio-biochemical analyses. All transgenic lines showed high gus and *SbMT*-2 gene expression (Figure 1c and 1d).

Southern blot confirmed single and double copy gene integration to L2, L4, L6 and L13 lines, respectively (Figure 1e).

Growth parameters in T1 transgenic lines

Transgenic lines L2, L4, L6 and L13 showed high percentage of seed germination in all stress treatments (metals- Zn, Cu and Cd; NaCl and PEG) compared to wild type (WT) plants (Figure S1). Transgenic lines grown under different stress treatments were comparatively healthier than WT plants (Figure 2) and showed enhanced growth parameters under stress conditions (Figure 3). Shoot and root lengths were found significantly higher in transgenic lines compared to WT plants in all stress treatments. Similarly, fresh and dry weights of transgenic lines were significantly higher than WT plants. Among all stress treatments studied, transgenic lines showed better growth in osmotic stress followed by metal and NaCl stress.

Results

Over-expression of the *SbMT*-2 gene

A plant expression vector (Figure 1a), harboring *SbMT*-2 gene driven by CaMV35S promoter was constructed and transformed to tobacco plants. Fourteen independent transgenic lines (T0) were raised and preliminarily screened by PCR using gene specific primers (data not shown). These lines were grown in containment facility and seeds (T1) were collected. T1 Seeds were germinated on hygromycin containing media and selected T1 transgenic lines were confirmed by PCR using gene specific primers (Figure 1b). Expected size of amplicon was found in all lines except L11. Based on *SbMT*-2 (Figure 1c) gene expression level, four independent T1 transgenic lines; L2, L4, L6 and L13 were selected for the further morpho-physio-biochemical analyses. All transgenic lines showed high gus and *SbMT*-2 gene expression (Figure 1c and 1d). Southern blot confirmed single and double copy gene integration to L2, L4, L6 and L13 lines, respectively (Figure 1e).

Figure 1. Analyses of transgenic tobacco plants. (a) Schematic map of plant expression gene construct pCAMBIA1301- *SbMT*-2. (b) Confirmation of transgenic lines by PCR amplification of *uidA*, *hptII* and *SbMT*-2 genes (Lane M is molecular weight markers, Lane PC is positive control of PCR using plasmid pCAMBIA1301- *SbMT*-2 as template, Lane WT is wild type control plant (non-transgenic) and Lane L is transgenic lines). (c) Over-expression of the *SbMT*-2 gene in transgenic lines compared to wild-type control plants (Lane WT is wild type control plant and Lane L is transgenic lines) analyzed by semi-quantitative Rt-PCR, where *actin* was used as internal gene control. (d) Histochemical GUS staining of WT and transgenic plants. (e) Southern blot analysis (Lane PC is positive control plasmid pCAMBIA1301- *SbMT*-2, Lane WT is wild type control plant (non-transgenic) and Lane L is transgenic lines). doi:10.1371/journal.pone.0111379.g001

The *SbMT*-2 Gene Modulates ROS Scavenging
Leaf senescence assay and chlorophyll content

Stress tolerance of T1 transgenic lines was studied by leaf disc senescence assay and chlorophyll estimation. In leaf discs assay, stress-induced necrosis resulted in the decrease of chlorophyll content was lower in the SbMT-2 over-expressing lines compared to WT plants (Figure 4). The damage caused by stress treatments was visualized by the degree of bleaching of leaf tissues and it was evident that the transgenic plants (L2, L4, L6 and L13) had a better ability to tolerate osmotic stress followed by metal and NaCl stress (Figure 4). The chlorophyll content in the WT plants reduced significantly with stress treatments while the transgenic lines (L2, L4, L6 and L13) retained higher chlorophyll contents than WT plants (Figure 5a).

Relative water content, Electrolyte leakage and membrane stability index

The relative water content (RWC) was found almost similar (insignificant difference) in transgenic and WT plants under un-stressed condition (control), however RWC was significantly higher in transgenic lines compared to WT plants under all stress treatments (Figure 5b). Membrane permeability measured by electrolyte leakage (EL) was found significantly stable in transgenic lines, which showed reduced electrolyte leakage compared to WT plants under stress condition (Figure 5c). Similarly, membrane stability index (MSI) of transgenic lines was significantly higher compared to WT under stress treatments (Figure 5d). RWC, EL and MSI evident that transgenic lines are thriving well in stress conditions compared to WT plants.

Localization of O$_2^-$ & H$_2$O$_2$ and H$_2$O$_2$, proline & MDA content analysis

Leaves of transgenic lines and WT plants, subjected to different stress treatments showed in vivo localization of O$_2^-$ and H$_2$O$_2$ (Figure 6). It was observed that leaves of WT plants showed more accumulation of O$_2^-$ and H$_2$O$_2$ content compared to transgenic lines under various stress conditions. Transgenic lines showed significantly lower accumulation of H$_2$O$_2$, proline and MDA contents under stress condition compared to WT plants (Figure 7). Lower content of H$_2$O$_2$ exhibited the better ROS system in transgenic lines compared to WT plants. Lower accumulation of proline and MDA revealed that transgenic lines have higher osmoprotectants and lower lipid peroxidation, respectively compared to WT plants under stress condition.

Ion content analysis

Metallothioneins play a vital role in ion homeostasis and heavy metal binding. Metal ion sequestration was analyzed by ICP, which revealed high accumulation of metal ions in transgenic lines compared to WT plants (Figure 8). It was observed that metal ion contents were approximately same (insignificant difference) in WT and transgenic lines at control (unstressed) condition (Figure 8a). Under stress condition, metal ion accumulation was significantly higher in transgenic lines compared to WT plants (Figure 8b). Among different tissues, Zn accumulation was higher in shoot followed by root and leaves. However, high Cu and Cd contents were detected in roots followed by stem and leaves. Furthermore high affinity of SbMT-2 was observed with Zn ions. Compared to control condition, Na$^+$ contents increased in transgenic as well as WT plants under NaCl stress treatment, however K$^+$ contents were decreased (Figure 8a). Similarly, transgenic lines showed higher K$^+$/Na$^+$ ratio compared to WT plants under NaCl stress condition (Figure 8b).

Transcript analysis of genes encoding metal transporters and antioxidative enzymes

Ion content analysis suggested that SbMT-2 may involve in the selective translocation of Zn$^{2+}$ from root to stem. To further support this observation, expression analysis of zinc and heavy metal transporters encoding genes (NtZIP1 and NtHMA-A) were studied under the metal stress (Zn, Cu and Cd) treatments. Expression of genes NtZIP1 and NtHMA-A involved in heavy metal (Zn) translocation were up-regulated in transgenic lines compared to WT plants (Figure 9). Among different metal stress, the NtZIP1 gene showed maximum up-regulation under Zn stress (Figure 9a). Transgenic lines, I2, I4, I6 and L13 showed about 10-, 13-, 8- and 6-fold expression compared to their respective control plants. However, NtZIP1 expression was also up-regulated in WT plant. Similarly, the NtHMA-A gene was also up-regulated under metal stress compared to WT and control condition (Figure 9b). In contrast to other metal stress, inconsistent expression was observed among transgenic lines and WT under Cu stress treatment.

Common antioxidant enzymes involved in ROS scavenging mechanism are superoxide dismutase (SODs), peroxidase (POD) and ascorbate peroxidase (APX). Transcript analysis reveals that expression of the NtSOD gene increased concomitantly with stress treatments (except NaCl) in transgenic lines compared to WT plants (Figure 10a). About 3.5-fold expression was observed in L4.
line under Zn and osmotic stress, while about 2-fold expression was detected under Cu and Cd stress compared to control condition (untreated plants). Remaining L2, L6 and L13 lines showed higher relative expression of NtSOD gene under different stress conditions compared to WT plants. Surprisingly down-regulation of gene was found under NaCl stress in transgenic as well as WT plants compared to control (untreated) plants. Relative fold expression of NtPOD gene was found higher in transgenic lines compared to WT plants under metal stress treatments (Figure 10b). Transgenic line L4 showed maximum gene expression about 10-fold under Zn, Cu and Cd stress. However, NtPOD gene expression was down-regulated in WT and transgenic lines under NaCl and PEG stress treatments. Elevated NtAPX gene expression was observed in transgenic lines (except L2 under NaCl and PEG) compared to WT plants under NaCl, osmotic and metal stress conditions (Figure 10c).

**Discussion**

Our previous study revealed that heterologous expression of SbMT–2 gene not only augments zinc and copper tolerance but also increases metal ion sequestration in *E. coli* cells [13]. Further, in this study, in planta functional validation of SbMT–2 gene has been elucidated. The SbMT–2 gene was over-expressed in tobacco and its functional role was studied in different abiotic stresses viz. salt (200 mM NaCl), osmotic (10% PEG), Zn (5 mM), Cu (0.2 mM) and Cd (0.2 mM) stress for 3 weeks. Morphological variation was not observed in transgenic lines over-expressing SbMT-2 gene compared to WT plants (Figure 2). Transgenic lines showed higher seed germination percentage, increased root length, shoot length, fresh weight (FW), dry weight (DW) and chlorophyll content under different stress conditions compared to WT plants (Figure 3 and 5a) which reveal that SbMT-2 leads to overcome the

![Figure 3](https://example.com/image3.png)

**Figure 3. Analysis of SbMT-2 transgenic tobacco lines under different stress condition.** Comparison of (a) shoot length (cm), (b) root length (cm), (c) fresh weight (mg) and (d) dry weight (mg) of wild type (WT) and T1 transgenic lines (L2, L4, L6 and L13) grown under control, salt (200 mM NaCl), osmotic (10% PEG), Zn (5 mM), Cu (0.2 mM) and Cd (0.2 mM) stress for 3 weeks. Graph represents the mean ± SE (of three replicates; n = 3) followed by similar letters are significantly different according to Tukey HSD at *P* < 0.05.

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![Figure 4](https://example.com/image4.png)

**Figure 4. Leaf disc assay of transgenic tobacco lines (T1) for different stress tolerance.** Leaf discs of WT, L2, L4, L6 and L13 transgenic lines respectively were floated in different stress solution for 8 days.

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deleterious effect of salt, drought and metal stresses. Previously, ectopic expression of \textit{BjMT2} gene conferred the increase in percentage seed germination, fresh weight (FW), dry weight (DW) and chlorophyll content in \textit{Arabidopsis} under Cu and Cd stress [15]. Similar results were also observed under salt, drought, Zn and Cu stress in transgenic tobacco plants over-expressing clustered \textit{OsMT1e-P} metallothionein gene [29].

Salt, drought and heavy metal stress are multigenic in nature, causing osmotic stress and thus create physiological drought conditions for plants. Transgenic lines displayed increase in relative water content under stress (Figure 5b) which indicates that \textit{SbMT-2} gene may help in water retention to counteract the osmotic shock. Abiotic stress causes the perturbation in metabolic balance of the cell, resulting in the enhanced production of ROS, which in-turn damages cell membranes, nucleic acids and chloroplast pigments [15,30]. Electrolyte leakage (EL) and membrane stability index (MSI) is the indicator of cell membrane stability. In the present study electrolyte leakage was lower in transgenic lines while MSI was higher under stress condition (Figure 5c-d) which suggests the role of \textit{SbMT-2} gene in cell membrane protection. It was further supported by lipid peroxidation analysis, H$_2$O$_2$ and proline quantification, in which high MDA, H$_2$O$_2$ and proline contents were found in WT plants compared to transgenic lines under stress condition (Figure 7). These result evident the possible role of \textit{SbMT-2} in ROS scavenging/detoxification and maintaining the cellular homeostasis during stress condition. Furthermore, \textit{in vivo} localization of H$_2$O$_2$ and O$_2^-$ under different abiotic stress (Figure 6), confirmed the role of \textit{SbMT-2} gene in ROS scavenging.

Although role of metallothionein genes in ROS scavenging and detoxification have studied [12,14,16,29,31] but the molecular mechanism of ROS detoxification/scavenging is still unknown. ROS, being a signaling molecule play an important role in development and regulation of different metabolic processes whereas ROS toxicity resulted into the oxidative damage to the cell membrane and its components [30]. Plant cell maintains a stringent regulation over its production and scavenging. In \textit{Arabidopsis} about 150 genes are involved in the homeostasis of ROS which comprised of ROS scavenging enzymes and ROS producing proteins [30]. Superoxide dismutase (SOD), peroxidase (POD) and ascorbate peroxidase (APX) are important ROS scavenging enzymes activated under different stress conditions to maintain the ROS homeostasis. Expression of genes encoding these ROS scavenging enzymes was studied in transgenic lines and compared with WT plants (Figure 10) to confirm the role of the \textit{SbMT-2} gene in ROS scavenging and detoxification. Higher relative fold expression of \textit{NtSOD} and \textit{NtPOD} genes were found in transgenic lines under stress conditions, however similar result was also observed with the \textit{NtAPX} gene except under salt stress. Moreover, high expression was detected in metal stress compared to salt and osmotic stress, which may be because of availability of cofactors of the enzymes.

Metallothioneins are involved in essential-metal homeostasis and impart protection against heavy metal toxicity by sequestra-
Plants maintain a high K⁺/Na⁺ ratio to combat with the deleterious effect of salinity. In present study transgenic lines over-expressing SbMT-2 gene showed a high K⁺/Na⁺ ratio under stress compared to WT plants and control condition (Figure S2). It provides further evidence that SbMT-2 may have role in ionic homeostasis and detoxification of H₂O₂ and thus impart salt tolerance. Transgenic lines showed accumulation of Cu²⁺ and Cd²⁺ in root compared to stem and leaves under stress condition. However, the SbMT-2 gene over-expression leads to the accumulation of Zn²⁺ in stem compared to control condition where it was accumulated more in roots (Figure 8). Zinc ion accumulation and tolerance varies among plant species and depends on MT type. The result exhibits SbMT-2 may involve in the selective translocation of Zn²⁺ from root to stem. Furthermore, the result also suggests that SbMT-2 is involved in Cu and Cd binding and accumulation rather than translocation, as observed with Zn ions. Previously, it was reported that SbMT2 protein exhibited high binding affinity and sequestration for Zn²⁺ compared to Cu and Cd ions [13].

In order to support this observation, expression of type 1B heavy metal–transporting P-type ATPases (P-Type ATPase) transporter and zinc specific transporter encoding gene NtHMA-A and NtZIP1 were analyzed under metal stress treatments (Figure 9). Up-regulation of genes in transgenic lines under metal stress especially, zinc compared to control and WT plants provides a supporting evidence that expression of ShMT-2 gene may influence metal transporters and thus translocation of Zn was observed in the study. The qPCR analysis revealed higher up-regulation of these genes in transgenic lines compared to WT plants (about 6 to 10 fold up-regulation of NtZIP1 gene in transgenic lines compared to about 2.5 fold of WT plants) under metal stress condition and it might be due to introgression of SbMT-2 gene. Furthermore, higher down-regulation of these genes (NtZIP1 and NtHMA-A) in transgenic plants compared to WT was observed under de-stress treatments, performed by re-culturing the plants in un-stressed conditions (Table S1). Results further suggest that the ShMT2 gene may influence the expression of these genes.

It was observed that expression of MT type 2 gene PsMT(A1) of Pisum sativum enhanced metal tolerance in white poplar and accumulation of zinc and copper (in leaves and roots) respectively [11]. Expression of seed specific MT gene MT4a in Arabidopsis increased Cu⁺⁺ accumulation but did not show any response under Zn stress [10]. The OsMT1a, a type 1 metallothionein gene, involved in the Zn⁺⁺ accumulation and thus provides tolerance to the transgenic rice [12]. Elsholtzia haichowensis metallothionein 1 (EhMT1) over-expression in tobacco plants enhances copper tolerance and accumulation in root cytoplasm [32] however, expression of BjMT2 gene in A. thaliana showed copper and cadmium tolerance [15]. Sequestration, translocation and thereby accumulation are the important mechanism used by plants for the phytorextraction [33] and therefore ShMT-2 gene may be utilized for the phytoremediation.

**Conclusion**

In conclusion, the present study provides an useful insight that ShMT-2 may involve in maintaining the cellular homeostasis by modulating ROS scavenging/detoxification during stress conditions and thus impart tolerance to salt and osmotic stress. It was observed that ShMT-2 provides protection against heavy metal toxicity by metal ions accumulation and may be involved in the
selective translocation of Zn$^{++}$ from roots to stem (Figure 11). It is speculated that SbMT-2 gene is a potential candidate for introgression to crop plants for imparting stress tolerance and phytoremediation.

**Methods**

**Construction of plant transformation vector and tobacco transformation**

The SbMT-2 cDNA was amplified using forward (MT-2F: 5’TCTGAGATGTCTCTTGTGTTGGAATC-3’) and reverse MT-2R: 5’-GGTACCTATTGGCAAGTGCAAGGGTTG-3’) primers containing XhoI and KpnI sites respectively. The amplicon was digested with XhoI/KpnI and cloned in pRT101 vector [34]. Thereafter, the gene cassette “CaMV35S-SbMT-2” was excised with PstI and cloned into the pCAMBIA1301 vector. The resulting vector was mobilised into Agrobacterium tumefaciens (EHA105) and further transformed to tobacco (Nicotiana tabacum cv. xanthii) plants according to the standard protocol [35].

**Confirmation of transgenic lines and expression of transgene**

Total genomic DNA was isolated from leaves of transgenic and wild type (WT; control/untransformed) tobacco plants (T1) using Qiagen DNeasy plant mini kit and quantified by Nanodrop Spectrophotometer (ND1000, Wilmington, USA). The integration of transgene in different lines was confirmed by PCR analyses using the SbMT-2 gene, reporter gene gus and selection marker gene hptII specific primers (Table 1). Presence and expression of the SbMT-2 gene in selected lines (L2, L4, L6 and L13) were confirmed by Southern blot and semi-quantitative RT-PCR, respectively.

Genomic DNA (20 μg) from WT and transgenic lines L2, L4, L6 and L13 was digested with HindIII, separated on agarose (0.7%) by electrophoresis and transferred to a Hybond N+ membrane (Amersham Pharmacia, UK) using alkaline transfer buffer (0.4 N NaOH with 1 M NaCl). DNA blot was hybridized with PCR-generated probe for the uidA gene labeled with DIG-11-dUTP following pre-hybridization and hybridization carried out at 42°C overnight in DIG EasyHyb buffer solution [36–39]. The hybridized membrane was detected by using CDP-Star chemiluminescent as substrate, following manufacturer user guide (Roche, Germany) and signals were visualized on X-ray film after 30 min.

Total RNA was extracted from transgenic tobacco (T1) and wild-type plants using Qiagen RNeasy plant mini kit and the cDNA was made by reverse transcription with Superscript II RT (Invitrogen, USA). The actin gene was used as an internal control and both genes (SbMT-2 and actin) were amplified using gene specific primers (Table 1). Histochemical GUS assay was performed with leaves as described by Jefferson [40].

Analyses of transgenic plants under different abiotic stress

Transgenic and wild type tobacco lines were maintained under controlled containment facility. Further morphological and physio-biochemical analyses were performed with T1 transgenic lines under different abiotic stress treatments.

**Growth parameters**

Seeds of transgenic lines (L2, L4, L6 and L13) and WT plants were germinated on MS medium supplemented with 200 mM NaCl, 10% PEG, 5 mM ZnSO4, 0.2 mM CuSO4 or 0.2 mM CdCl2 under culture room conditions and the percentage of seed germination was calculated after 20 days.

Different growth parameters were measured under same stress conditions using T1 seedlings. Seeds of transgenic lines were germinated on the MS media supplemented with 20 mg/l hygromycin, while seeds of WT plant were germinated on MS media only. WT and hygromycin positive T1 seedlings were transferred to MS medium supplemented with 200 mM NaCl, 10% PEG, 5 mM ZnSO4, 0.2 mM CuSO4 or 0.2 mM CdCl2 after eight days and grown for further 21 days. Growth
parameters; shoot length, root length, fresh weight and dry weight were recorded and compared with wild type plants.

**Leaf senescence assay and chlorophyll estimation**

Leaf discs from 30 days old WT and transgenic plants (L2, L4, L6 and L13) were used for the stress tolerance assay. Healthy leaves of similar age were detached and leaf discs (of 5 mm in diameter) were punched out. About 8 discs of each plant were floated in 5 ml ½ Hoagland media (control) supplemented with 200 mM NaCl, 10% PEG, 1 mM ZnSO₄, 0.5 mM CuSO₄ or 0.5 mM CdCl₂ for 7 days. The effect of these treatments on leaf discs were assessed by observing phenotypic changes.

Leaf discs (control and treated) were further subjected to chlorophyll isolation. Leaf discs were thoroughly homogenized in chilled N, N-dimethylformamide (DMF) at 4°C and thereafter centrifuged at 10,000 g for 10 min. Supernatant was aspirated out and O.D. was recorded at 664 and 647 nm. Total chlorophyll content was calculated per gram fresh weight of tissue according to Porra et al. [41].

**Relative water content, Electrolyte leakage and Membrane stability index**

WT and T₁ seedlings (L2, L4, L6 and L13) were transferred to ½ Hoagland hydroponics culture and maintained for 20 days. Healthy young plants of same age and size were collected for each treatment (24 h for 200 mM NaCl and 0.5 mM CuSO₄; 12 h for 10% PEG, 1 mM ZnSO₄ and 0.5 mM CdCl₂).

About 100 mg (FW) leaves of control and treated plants were submerged into the deionised water and after 12 h turgid weight (TW) were recorded. Samples were further kept at 80°C for 48 h.
to record the dry weight (DW). The relative water content was calculated as: 
\[ RWC \, (\%) = \left( \frac{FW - DW}{TW - DW} \right) \times 100. \]

Collected leaves were washed thoroughly with deionized water to remove surface-adhered electrolytes. Samples were kept in closed vials containing 10 ml deionised water and incubated at 25°C on a rotary shaker for 24 h. Subsequently, the electrical conductivity (EC) of the solution (\( L_t \)) was determined using conductivity meter (SevenEasy, Mettler Toledo AG 8603, Switzerland). Samples were autoclaved at 120°C for 20 min, cooled up to 25°C and electrical conductivity (\( L_0 \)) was determined. The electrolyte leakage was determined: 
\[ \text{Electrolyte leakage} \, (\%) = \left( \frac{L_t}{L_0} \right) \times 100. \]

Healthy young leaves of stress-treated plants were taken and membrane stability index (MSI) was determined [42]. Leaves (200 mg) were kept in close vials containing 10 ml deionized water. A set of vials was incubated at 40°C for 30 min while second set of vials was incubated at 100°C for 10 min. Electrical conductivity was recorded for both sets (\( L_1 \) for 40°C while \( L_2 \) for 100°C) and MSI was calculated using the formula as: 
\[ MSI = \left[ 1 - \left( \frac{L_1}{L_2} \right) \right] \times 100. \]

**In vivo localization of O$_2^-$ and H$_2$O$_2$**

Histochemical staining was performed for the *in vivo* detection of O$_2^-$ and H$_2$O$_2$ using nitro-blue tetrazolium (NBT) and 3, 3-diaminobenzidine (DAB), respectively [43]. The presence of O$_2^-$ in transgenic and WT leaves exposed to different stresses (as described above) was detected by immersing the leaf samples in NBT solution (1 mg ml$^{-1}$ in 10 mM phosphate buffer; pH 7.8) at room temperature for 2 h and then illuminated for 12 h in light until blue spots appeared. For the localization of H$_2$O$_2$, treated leaves were incubated in DAB solution (1 mg ml$^{-1}$ in 10 mM phosphate buffer; pH 3.8) at room temperature for 6 h in dark, thereafter exposed to the light until brown spots were appeared. Leaf samples were treated with absolute ethanol (for bleaching chlorophyll contents) before the documentation.
Quantification of H$_2$O$_2$, proline and lipid peroxidation

The H$_2$O$_2$ and free proline content in leaf samples (WT and transgenic lines under different stress conditions) were measured [44,45]. Proline and H$_2$O$_2$ levels were calculated by the standard curve, prepared against known concentration of proline or H$_2$O$_2$ measured at 520 and 560 nm absorbance respectively. Lipid peroxidation was estimated by determining the concentration of malondialdehyde (MDA) produced by thiobarbituric acid (TBA) reaction [46]. Leaf samples were extracted in 2 ml 0.1% trichloroacetic acid (TCA) and 0.5 ml extract was reacted with 2.0 ml of TBA reagent followed by boiling at 95°C for 30 min. Samples were cooled at ice, centrifuged at 10000 g for 5 min and absorbance of the supernatants was measured at 440 nm, 532 nm and 600 nm.

Ion content analysis

For ion content analysis, tissues (root, shoot and leaves) from 4-week-old plants grown in hydroponic medium and treated with 200 mM NaCl, 10% PEG 6000, 1 mM ZnSO$_4$, 0.5 mM CuSO$_4$ or 0.5 mM CdCl$_2$ were washed with deionised water, dried in hot air oven for 48 h at 70°C and digested with 4 ml perchloric acid-nitric acid solution (3:1). The solution was heated to dry and further diluted to 25 ml with deionised water. Ion contents were measured by inductively coupled plasma optical emission spectrometer (Optima2000DV, PerkinElmer, Germany).

Transcriptional regulation of metal transporters and antioxidative enzymes

Total RNA was isolated from the stressed and control plants by using RNeasy plant mini kit (Qiagen) following the manufacturer’s instructions and quantified with Nanodrop spectrophotometer (NanoDrop, USA). The cDNA was prepared using 2 µg of total RNA with a SuperScript RT III first-strand cDNA synthesis kit (Invitrogen, USA). The expression pattern of genes; NtZIP1, NtHMA-A, encoding metal transporters [47] and NtSOD, NtPOX and NtAPX, encoding antioxidant enzymes [48] which involved in heavy metal transportation and ROS scavenging, respectively were analyzed under different stress by using gene-specific primer pairs (Table 1), while the gene actin was used as an internal control. The quantitative real time PCR (qRT-PCR) was performed in a Bio-Rad IQ5 detection system (Bio-Rad, USA) with QuantiFast Kit (Qiagen, USA). Fold expression was calculated by the method described by Livak and Schmittgen [49] and specificity of qRT-PCR was monitored by melt curve analysis.

Statistical analysis

Each experiment was carried out in three replicates. All data was expressed as mean ± SD and subjected to analysis of variance (ANOVA) to determine the significance of difference between the means of WT and transgenic plants of each treatment group. A Tukey HSD multiple comparison of mean test was used, significant differences were considered at $P<0.05$ and indicated by similar letters.
Table 1. Primer sets used for the confirmation and transcript analysis of transgenic lines.

| Genes | Primer sequences (5’ to 3’) | PCR conditions |
|-------|-----------------------------|----------------|
| uida  | F: GATCGCGAAAACGTGGGAAT    | 95°C, 5min; 34 Cycles: 95°C, 1min; 60°C, 1min; 72°C, 2 min; Extension: 72°C, 7 min |
|       | R: TGAAGCCTCCGACAACTTAC     |                |
| hptl  | F: TTCTTTGCTGCCGACGAGTG     | 95°C, 5min; 34 Cycles: 95°C, 1min; 55°C, 1min; 72°C, 2 min; Extension: 72°C, 7 min |
|       | R: ACAAGCTCTCCGACCGATG      |                |
| SbMT-2| F: AGGTTCGAAAGTGTCC          | 95°C, 5min; 35 Cycles: 95°C, 30sec; 60°C, 45sec; 72°C, 3 min; Melt curve: 60–95°C |
|       | R: TCCCAGTGTCACGTGCTAC      | with 0.5 C/cycle increment |
| Actin | F: CGTTTGATCCTGCTGGTGT      |                |
|       | R: CAGCAAGCCGAGGACATAG      |                |
| NZIP1 | F: TGGTTGCTACCTGCTGAGAT     |                |
|       | R: CGAAGAGCTCAAGACTGAGA     |                |
| NHMA-A| F: ACAATGGCTGGCAAACCAA      |                |
|       | R: TTTCCTGAGGGACAGTCTC      |                |
| NISOD | F: AGCTACATGAGCCATTCTCC     |                |
|       | R: CCGTGAAGGCACGACCTTC      |                |
| NIPOD | F: CTTGGAACACGACCGTTCTCT    |                |
|       | R: TGCGTATCGCCGTTCTCTCTCT   |                |
| NAPX  | F: CAAATGAAGAGAAACTCAGAGGA  |                |
|       | R: CAGCCTTGAGGGCTCATGGTACCG |                |

Supporting Information

Figure S1  Percentage of seed germination under different stress condition. Seeds of WT and transgenic lines were germinated on MS media supplemented with 200 mM NaCl, 10% PEG, 5 mM Zn, 0.2 mM Cu or 0.2 mM Cd. Graph represents the mean ± SD (of three replicates; n = 3) followed by similar letters are significantly different according to Tukey HSD at P<0.05. (TIF)

Figure S2 Sodium and potassium ion content in transgenic tobacco lines under different stress condition. Comparison of (a) Na⁺ and K⁺ content and (b) K⁺/Na⁺ ratio in WT and transgenic lines in root, stem and leaf under stressed (200 mM NaCl) and un-stressed conditions. Graph represents the mean ± SD (of three replicates; n = 3) followed by similar letters are significantly different according to Tukey HSD at P<0.05. (TIF)

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Author Contributions

Conceived and designed the experiments: AM BJ. Performed the experiments: AKC MKP VT. Analyzed the data: AM AKC. Wrote the paper: AKC AM.
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