



Comparative Transcriptome Profiling Under Cadmium Stress Reveals the Uptake and Tolerance Mechanism in *Brassica juncea*

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Abstract

Cadmium (Cd) is a biologically non-essential and phytotoxic heavy metal pollutant. In this study, we estimated the Cd accumulation potential of Indian mustard and identified factors responsible for Cd uptake, tolerance, and detoxification. Eight transcriptomic libraries were sequenced and ≈ 230 million good quality reads were generated. The alignment rate against *B. juncea* reference genome V1.5 varied in the range of 85.03–90.06%. Comparative expression analysis using DESeq2 revealed 11,294 genes to be significantly differentially expressed under Cd treatment. The agriGO singular enrichment analysis revealed genes related to response to chemical, oxidative stress, transport, and secondary metabolic process were upregulated, whereas multicellular organismal development, developmental process, and photosynthesis were downregulated by Cd treatment. Furthermore, 616 membrane transport proteins were found to be significantly differentially expressed. Cd-related transporters such as metal transporter (Nramp1), metal tolerance protein (MTPC2, MTP11), cadmium-transporting ATPase, and plant cadmium resistance protein (PCR2, PCR6) were upregulated whereas cadmium/zinc-transporting ATPase (HMA2, HMA3, HMA4), high-affinity calcium antiporter (CAX1), and iron transport protein (IRT1) were downregulated by Cd treatment. A total of 332 different gene-networks affected by Cd stress were identified using KAAS analysis. Various plant hormones signaling cascades were modulated suggesting their role in Cd stress tolerance. The regulation overview using MapMan analysis also revealed gene expression related to plant hormones, calcium regulation, and MAP kinases were altered under Cd stress.

Keywords Cadmium · *Brassica juncea* · Phytoremediation · Transcriptome · Pathway analysis

Introduction

Cadmium (Cd) is a toxic heavy metal (HM) and a widespread environmental pollutant. The concentration of Cd in soil is increasing over the years, for example, it is reported to increase by 0.2% per year in Scandinavia (Jarup 2003). Although the sources of Cd are both natural and anthropogenic, the contribution of the latter far exceeds the former. The anthropogenic factors for Cd pollution are atmospheric deposition, metal smelting, fuel combustion, application of

phosphate fertilizers, incineration and dumping of Cd-polluted waste, and sewage and sludge disposal. Cd can immobilize in soil by binding to organic matter. The agricultural sites polluted with Cd are of major concern as being highly soluble Cd is readily absorbed by plants and enters the food chain (Sarwar et al. 2010). The dietary consumption of such Cd-containing crops can cause chronic toxicity in human. Besides being carcinogenic, Cd intoxication can damage the kidney, liver, bones, and lungs (Rahimzadeh et al. 2017). Cd is a non-essential element for plant growth and can cause stunting, leaf rolls, chlorosis, oxidative stress, genotoxicity and inhibition of the photosynthesis, respiration and nitrogen metabolism in plants (Andresen and Küpper 2013).

Remediation of environmental pollutants is a financial and technical challenge. The physicochemical methods like excavation, adsorption, biosorption, ion-exchange, precipitation, and so on, are neither cost-effective nor suitable for widespread pollutants like cadmium. Plants that accumulate HMs in their aboveground tissue have the potential to

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be used for HM remediation. The technology of removing pollutants from the environment with the help of plants is termed as phytoremediation (Cunningham and Berti 1993). The ideal plant for phytoremediation purposes should be able to (1) tolerate the pollutant, (2) accumulate the pollutant in an easily harvestable part, (3) adapted to grow locally, (4) and accumulate high biomass. The idea of phytoremediation is not new, hitherto the true potential of phytoremediation remains to be achieved. For its successful implementation, selection of suitable plant species plays an important role. Members of the family Brassicaceae are known to accumulate various metal (loid)s in the aboveground tissue. Previous studies have suggested Indian mustard (*Brassica juncea*) as a potential species for phytoremediation. Indian mustard is reported to accumulate various metal(loid)s, for example, Cd, As, Pb and Cu in its aboveground part (Goswami and Das 2015).

Cd toxicity can generate oxidative stress in plants although it is not a redox-active element. The accumulated Cd ions disturb the redox balance resulting in the generation of reactive oxygen species (ROS) which further leads to the destruction of cellular and metabolic function. The level of enzymatic and non-enzymatic antioxidants is reported to increase in response to Cd exposure to alleviate the oxidative damage (Ahmad et al. 2011; Choudhury and Kumar 2004). Stress perception is important in providing tolerance towards a particular stress. The Cd stress is reported to activate various kinases of the MAPK signaling cascade which further phosphorylate numerous substrates including transcription factors (DalCorso et al. 2010; Liu et al. 2010). The MAPK signaling cascade in plants is involved in the perception of environmental and developmental stimuli (Jonak et al. 2002). Calcium ions (Ca^{2+}) are also shown to be involved in signal cascades of environmental stimuli. Studies have shown the protective role of Ca against Cd toxicity (Wang and Song 2009). Ca modulates the activity of target proteins and protein kinases via direct binding or through calmodulin (CALM) (Roberts and Harmon 1992). Another consequence of Cd toxicity is the modulation of plant hormone signaling. The level of abscisic acid (ABA) is shown to be increased under Cd stress in oilseed rape (*Brassica napus* L.) (Yan et al. 2016). Transcriptomic studies have identified transcription factors belonging to several families, for example, WRKY, ethylene-responsive factor (ERF) and myeloblastosis protein (MYB) to control gene expression under Cd stress. Previous studies were limited to biochemical analysis, determination of Cd bio-accumulation or characterization of a limited number of genes for Cd tolerance and accumulation (Mobin and Khan 2007; Qadir et al. 2004; Zhu et al. 1999). The underlying mechanism and the gene networks involved in Cd tolerance and accumulation in Indian mustard remain to be identified. With the advances in high-throughput technology, it has become possible to study the whole stress

transcriptome for non-modal crops. In the present study, we investigated Indian mustard cultivar RLM514 for understanding the molecular mechanism of Cd tolerance and identification of modulated gene networks using RNA-seq. The cultivar RLM514 has been tested to accumulate arsenic (As) in a separate study conducted in our laboratory. Thus, the cultivar RLM514 constitutes a fascinating plant material to study uptake, translocation, and detoxification of metal(loid)s and can contribute significantly to phytoremediation of sites contaminated with multiple metal(loid)s.

Materials and Methods

Plant Material and Growth Condition

For the present study, we used genotype RLM514 of Indian mustard. Seeds were surface sterilized using 3% H_2O_2 and subsequently rinsed thoroughly with distilled H_2O . Seeds were then germinated on trays containing autoclaved vermiculite and allowed to grow for 14 days in controlled conditions (14 h photoperiod; at 28 ± 2 °C).

Cadmium Treatment and Bioaccumulation Estimation

The seedlings grown as discussed above were uprooted carefully and washed thoroughly with dH_2O . The seedlings were then transferred to hydroponic solutions and divided into two groups, that is, control and Cd-treated. The control group was grown without any Cd whereas the Cd-treated group was supplied with different concentrations of Cd (250 μM , 500 μM , 750 μM and 1000 μM) as CdCl_2 . After 72 h, seedlings were taken out of the hydroponic solutions and washed thoroughly with dH_2O . The root and shoot tissues were then oven dried separately for 48 h at 60 °C. A total of 100 mg of the dried samples were digested using a microwave in 8 mL mix of concentrated HNO_3 and H_2O_2 (3:1) and subsequently diluted with milliQ, filtered through Whatman filter paper and the amount of Cd bioaccumulated was determined using an atomic absorption spectrometer (Shimadzu Model AA-7000).

RNA Sequencing

After 72 h of treatment, the roots and shoots of control and maximum Cd-accumulating groups (750 μM) were harvested and ground using liquid nitrogen. Total RNA was isolated from the ground tissues using a NucleoSpin RNA Plant kit (Macherey–Nagel), as per the manufacturer's instructions. The quantity of RNA was estimated on a NanoDrop 2000 spectrophotometer (Thermo Fisher Scientific) and the quality was confirmed using agarose gel electrophoresis. The

RNA integrity was determined using Bio-analyzer (Agilent Technologies). The mRNA was isolated from the total RNA using oligo-dT beads using a TruSeq RNA sample preparation kit (Illumina). Subsequently, cDNA was synthesized and sequencing libraries were constructed using random fragmentation followed by adapter ligation and PCR amplification. A total of eight sequencing libraries were generated consisting of two biological replicates from the root and shoot tissues of control and Cd-treated groups. Prepared libraries were quantified using qPCR according to the Illumina qPCR Quantification protocol guide and sequenced on the HiSeq platform (Illumina Inc.) using 100 bp paired-end sequencing generating 20–25 million reads per library.

Transcriptome Assembly

The FastQC (version 0.11.4) was used to check the quality of raw reads (Andrews 2010). Afterward, the Trimmomatic tool (version 0.32) was used for trimming adapters and low quality reads (Phred Score ≤ 20) (Bolger et al. 2014). Paired-end reads having less than a 100 bp sequence length were removed from further analysis. Trimmed paired-end reads were aligned against the *B. juncea* reference genome V1.5 (Yang et al. 2016) using HISAT (Galaxy Version 2.0.5.1). The aligned reads were assembled to form potential transcripts using StringTie (Galaxy Version 1.3.3) (Pertea et al. 2015). CD-HIT-EST was used to make the master transcriptome (transcriptome consisting of all the 8 samples) non-redundant at 90% sequence identity cut-off (Huang et al. 2010).

Comparative Expression Analysis

The abundance of potential transcripts was determined from alignment files using HTSeq (Anders et al. 2015). For comparative expression analysis, the count tables generated using HTSeq were analyzed using DESeq2 (Love et al. 2014). Comparisons were made between control and Cd-treated tissues. The transcripts with a false discovery rate (FDR) below 0.05 and \log_2 fold change is greater than 2 or less than -2 were considered significantly differentially expressed.

Functional Annotations

A BLASTx homology search was performed against the TAIR10 database (<https://www.arabidopsis.org>) to determine the functional description of differentially expressed transcripts and against the transporter classification database (TCDB) for identification of the transporters (<http://www.tcdb.org/>) (e -value $< 10^{-3}$). Gene ontology (GO) analysis was performed using agriGO singular enrichment analysis (SEA) with FDR < 0.05 (Du et al. 2010) and visualized using

REVIGO (Supek et al. 2011). Various regulatory and metabolic pathways were determined and visualized using KAAS (KEGG Automated Annotation Server) (Moriya et al. 2007) and MapMan (Version 3.6.0) (Thimm et al. 2004).

Real-Time Quantitative PCR (RT-qPCR)

The root and shoot tissues from control and Cd-treated groups were harvested in liquid nitrogen after 72 h of Cd treatment. The harvested tissues were then utilized for total RNA isolation using the NucleoSpin RNA Plant kit (Macherey–Nagel), as per the manufacturer's instructions. The quality and quantity of RNA were estimated using agarose gel electrophoresis and NanoDrop 2000 spectrophotometer (Thermo Fisher Scientific) respectively. RNase-free water was used to dilute and adjust RNA concentration (100 ng/ μ L) in each sample. Later, cDNA synthesis was performed using a high-capacity cDNA reverse transcription kit (Applied Biosystems) as per the manufacturer's instructions. Primer-pairs (Online Resource 1) were designed from selected transcripts using Primer3Plus (Untergasser et al. 2007). PowerUp SYBR Green Master Mix (Applied Biosystems) was used to perform RT-qPCR. The comparative gene expression was calculated using the $2^{-\Delta\Delta CT}$ method (Rao et al. 2013). A housekeeping gene (actin) was used as a reference gene for calculations.

Results

Cadmium Bioaccumulation

After 72 h, the active accumulation of Cd in shoots increased linearly with increase in treatment concentration until it reached a maximum value (1057.92 mg/Kg) at 750 μ M of Cd (Fig. 1a) whereas in roots it was 36703.063 mg/ Kg at 500 μ M of Cd (Fig. 1b). The results of the present study are in line with the study in which Cd accumulation increased with an increasing concentration of treatment in *Sedum alfredii* and reached a maximum of 5659.334 mg/Kg at 500 μ M of Cd (Ni and Wei 2003). In another hydroponics study, Cd accumulation in roots was detected to be more than 5000 mg/Kg and 15,000 mg/Kg in poplar and willow clones, respectively (Zacchini et al. 2009).

Transcriptome Assembly and Functional Annotation

We obtained ≈ 230 million good quality reads (20–30 million paired-end reads per library) from 8 eightlibraries generated from the root and shoot tissue. For reference-guided transcriptome assembly, the reads were aligned against the *B. juncea* reference genome (V1.5). The alignment rate varied in the range of 85.03–90.06% (Online Resource 2).

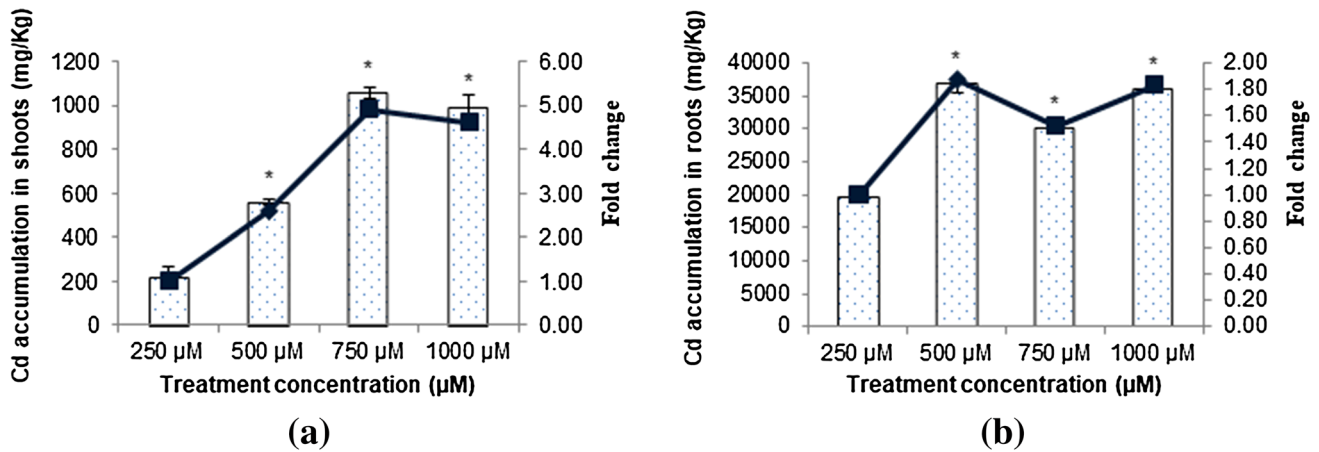


Fig. 1 Cadmium accumulation by RLM514 in **a** shoot and **b** root tissues at different concentrations of treatment. Values are mean \pm standard deviation of three replicates

The estimation of the global quality of the whole RNA-seq dataset and reproducibility of biological replicates is critical for gene-expression analysis. There are no clear standards but it is expected that in a principal component analysis (PCA) biological replicates will cluster together. The PCA performed using the alignment files clustered the biological replicates together also the different tissues, that is, root and shoot, were clustered on different quadrants (Fig. 2). The results of PCA analysis authenticate the reliability of biological replicates used in the present study (Conesa et al. 2016).

For comparative expression, comparisons were made among Cd-treated and control tissues. We found 11,294 genes to be significantly differentially expressed (\log_2 fold change is > 2 or < -2 at $FDR < 0.05$) (Fig. 3) out of which 911 (8.06%) were mutually expressed in root and shoot, whereas 7370 (65.26%) were expressed only in root and 2101 (18.60%) were exclusive to shoots. We found 2,021 (67.09%) genes to be upregulated, whereas 992 (32.90%) were downregulated in the shoot. Similarly, 4682 (56.53%) genes were upregulated and 3599 (43.46%) were downregulated in root (Fig. 4). A total of 18,184 transcripts mapping to differentially expressed genes (DEGs) were identified by a search against the *Arabidopsis thaliana* TAIR10 protein database (<https://www.arabidopsis.org>) for functional annotation of genes.

Transporters

A total of 4670 transcripts from DEGs mapped to 616 different membrane transport proteins (Online Resource 3) Further the transport proteins were assigned to seven classes, that is, channels/pores (148), electrochemical potential-driven transporters (200), primary active transporters (131), group translocators (12), transmembrane electron carriers (13), accessory factors involved in transport (40) and

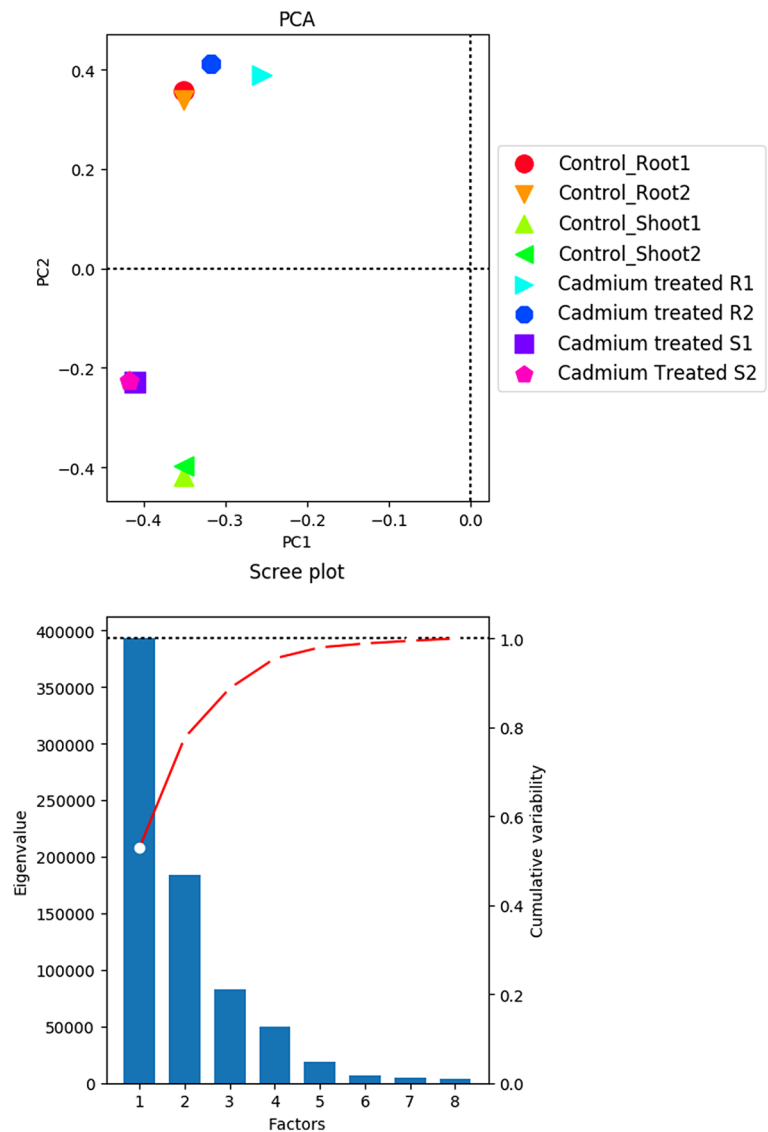
incompletely characterized transport systems (65). We found Cd-related transporters such as metal transporter (Nramp1), zinc transporter (ZIP8), metal tolerance protein (MTPC2, MTP11), zinc-induced facilitator (ZIF1), cadmium-transporting ATPase, plant cadmium resistance protein (PCR2, PCR6), metal-nicotianamine transporter (YSL2) to be upregulated and cadmium/zinc-transporting ATPase (HMA2, HMA3, HMA4), high affinity calcium antiporter (CAX1), iron transport protein (IRT1), zinc transporter 1 precursor (ZRT/IRT-like protein 1) were downregulated in response to Cd treatment. Furthermore, 130 transcripts mapped to different ABC transporters among which MDR (15 transcripts) were upregulated while other ABC transporters such as ABCA12, ABCB2, ABCC5 and ABC transporter G family members were downregulated in response to Cd.

Gene Ontology (GO) and Gene-Network Analysis

Gene ontology (GO) analysis of DEGs revealed 174 and 81 significant GO terms ($FDR < 0.05$) were upregulated and downregulated, respectively in Cd-treated shoots. In addition, 210 and 55 significant GO terms were upregulated and downregulated in Cd-treated roots (Online Resource 4). Overall, the genes associated with response to a stimulus, jasmonic acid (JA), localization, and secondary metabolism were upregulated, whereas multi-organism process, developmental process, reproduction, and starch metabolism were downregulated in Cd-treated tissues (Fig. 5).

We found 332 different pathways in KAAS analysis to be affected by Cd stress (Online Resource 5) which include glutathione (GSH) metabolism (87 transcripts), phenylpropanoid biosynthesis (112 transcripts), MAPK signaling pathway (125 transcripts), calcium signaling pathway (14 transcripts), and plant hormone signal transduction (184 transcripts). MapMan analysis of regulation overview also

Fig. 2 Principal component analysis (PCA) and scree plot for top eight principal components



revealed the expression of various transcripts mapping to plant hormones, calcium regulation, MAP kinases, and protein degradation were altered under Cd stress (Fig. 6).

A total of 97 transcripts mapping to the MAPK signaling pathway were upregulated and 28 were downregulated (Online Resource 6). The genes MAP kinase substrate 1 (MKS1), WRKY transcription factor 33 and 22 (WRKY33, 22), mitogen-activated protein kinase kinase 4/5 (MKK4/5), aminocyclopropane-1-carboxylate synthase (ACS), serine/threonine-protein kinase OXI1 (OXI1), mitogen-activated protein kinase kinase 9 (MKK9), calmodulin (CALM), respiratory burst oxidase (RBOH) mapping to MAPK signaling in plants were upregulated. Three genes related to calcium signaling viz., phosphatidylinositol phospholipase C, delta (PLCD), mitochondrial calcium uniporter (MCU) and voltage-dependent anion channel protein 2 (VDAC2) were upregulated in Cd-treated tissues (Online Resource 7). A

total of 136 transcripts mapping to plant hormone signal transduction were upregulated and 48 transcripts were found to be downregulated by Cd treatment.

Exposure to Cd in plants is well known to induce oxidative stress although it is a non-redox metal and unable to carry out Fenton-type reactions (Laspina et al. 2005). The level of ROS is maintained carefully inside a plant cell to avoid the oxidative stress. Cd can inhibit as well as stimulate several antioxidative enzymes. To avoid the oxidative damage due to ROS, plants employ various enzymatic and non-enzymatic antioxidants. A total of 50 transcripts mapped to peroxidases out of which 28 were upregulated. The gene expression of non-enzymatic antioxidants such as coumarins and flavonoids was also modulated by Cd exposure. We found 23 differentially expressed transcripts mapping to flavonoid biosynthesis out of which ten were upregulated in Cd-treated tissue. Among the transcripts

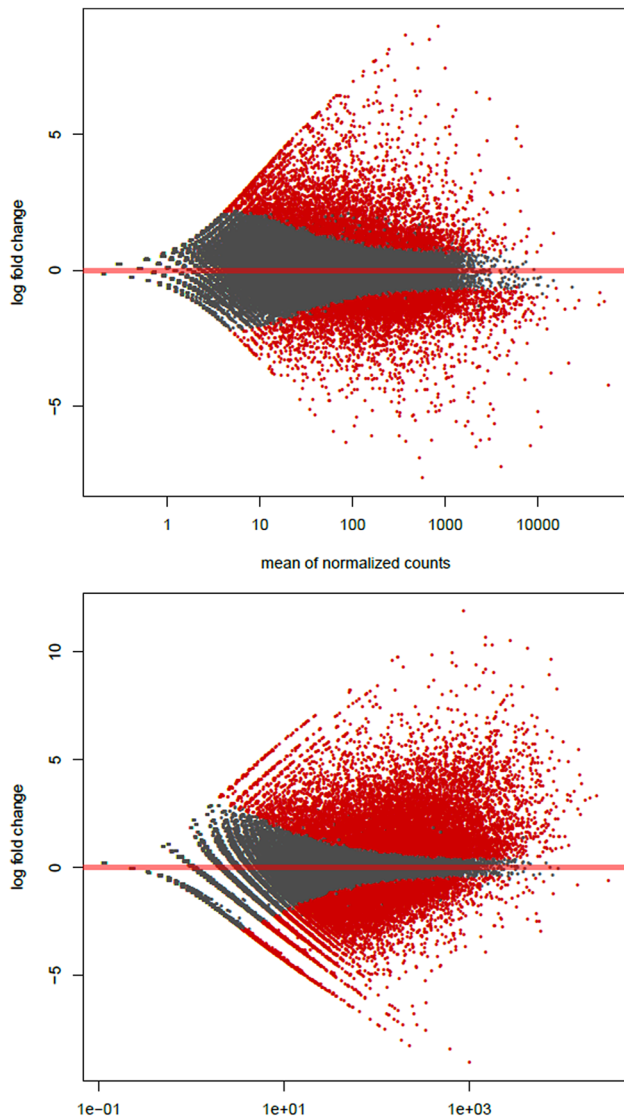


Fig. 3 MA plots of DEGs in **a** root and **b** shoot tissues. Red dots in the plots represents significantly differentially expressed genes, that is, log₂ fold change is > 2 or < -2 at FDR < 0.05

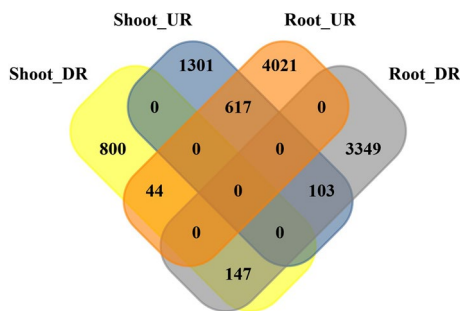


Fig. 4 DEGs distribution in Cd-treated root and shoot tissues. Root_UR and Root_DR represent no. of DEGs upregulated and downregulated in Cd-treated roots in comparison to control tissues. Shoot_UR and Shoot_DR represent no. of DEGs upregulated and downregulated in Cd-treated shoots in comparison to control tissues

mapping to GSH metabolism, the gene expression of a total of ten important components including glutathione S-transferase (GST), glutathione synthase (GSS) and gamma-glutamyltranspeptidase (GGT) found to be modulated by cadmium (Online Resource 8). A total of 66 different transcripts encoding GST were found to be significantly upregulated.

Among the plant hormone signal transduction, four genes (auxin-responsive protein IAA (AUX/IAA), auxin response factor (ARF), auxin-responsive GH3 gene family (GH3), and SAUR family protein (SAUR)) related to the auxin signaling cascade were modulated in Cd-treated tissues (Online Resource 9). In cytokinin (CKK) signaling, three genes (CRE1 (cytokinin response 1), type-A *Arabidopsis* response regulators (A-ARR), type-B *Arabidopsis* response regulators (B-ARR)) and two genes related to gibberellin signaling (gibberellin receptor (GID1) and phytochrome-interacting factor 4 (PIF4)) were differentially expressed in Cd-treated tissues. We also detected downregulation of adenylate dimethylallyltransferase (cytokinin synthase) (IPT) and upregulation of PP2C. Similarly, we found 5,6,3 and 3 differential genes related to ethylene, brassinosteroid (BR), jasmonic acid (JA) and salicylic acid signaling. The genes mapping to ethylene receptor (ETR), ethylene-insensitive protein 3 (EIN3), ethylene-responsive transcription factor 1 (ERF1), ethylene-responsive transcription factor 1 (ERF1) were upregulated and ethylene-insensitive protein 2 (EIN2) were downregulated by Cd treatment. EIN2 and EIN3 play a central role in the ethylene signaling. The plant steroid hormones, brassinosteroids (BRs), play important roles in alleviating biotic and abiotic stress (Rajewska et al. 2016). We found two brassinosteroid-insensitive genes, BRI1 and BIN2, known components of BR signaling, to be modulated under Cd treatment. BRI1, essential for BR perception, was downregulated, whereas BIN2, involved in negative regulation of BR signaling, was upregulated in Cd-treated tissues (Nam and Li 2002). The gene expression of transcription factors brassinosteroid resistant 1 and 2 (BZR1 and 2) was upregulated, although BIN2 is its negative regulator. The gene expression of xyloglucan:xyloglucosyl transferase (TCH4), cyclin D3 (CYCD3), the response genes of the BR signaling pathway was also upregulated in Cd-treated tissue. TCH4 encodes for cell wall protein involved in cell wall modification in response to abiotic stress, whereas CYCD3 promotes supernumerary cell division (Sun et al. 2017). The gene expression of all the three components jasmonic acid-amino synthetase (JAR1), jasmonate ZIM domain-containing protein (JAZ) and MYC2 were upregulated by Cd treatment. JAR1 conjugates JA with isoleucine (Ile) which further promotes proteasomal degradation of JAZ and hence releases transcription factors such as MYC2 involved in the induction of early JA-responsive genes. The regulatory protein (NPR1) is a critical component of SA signaling and

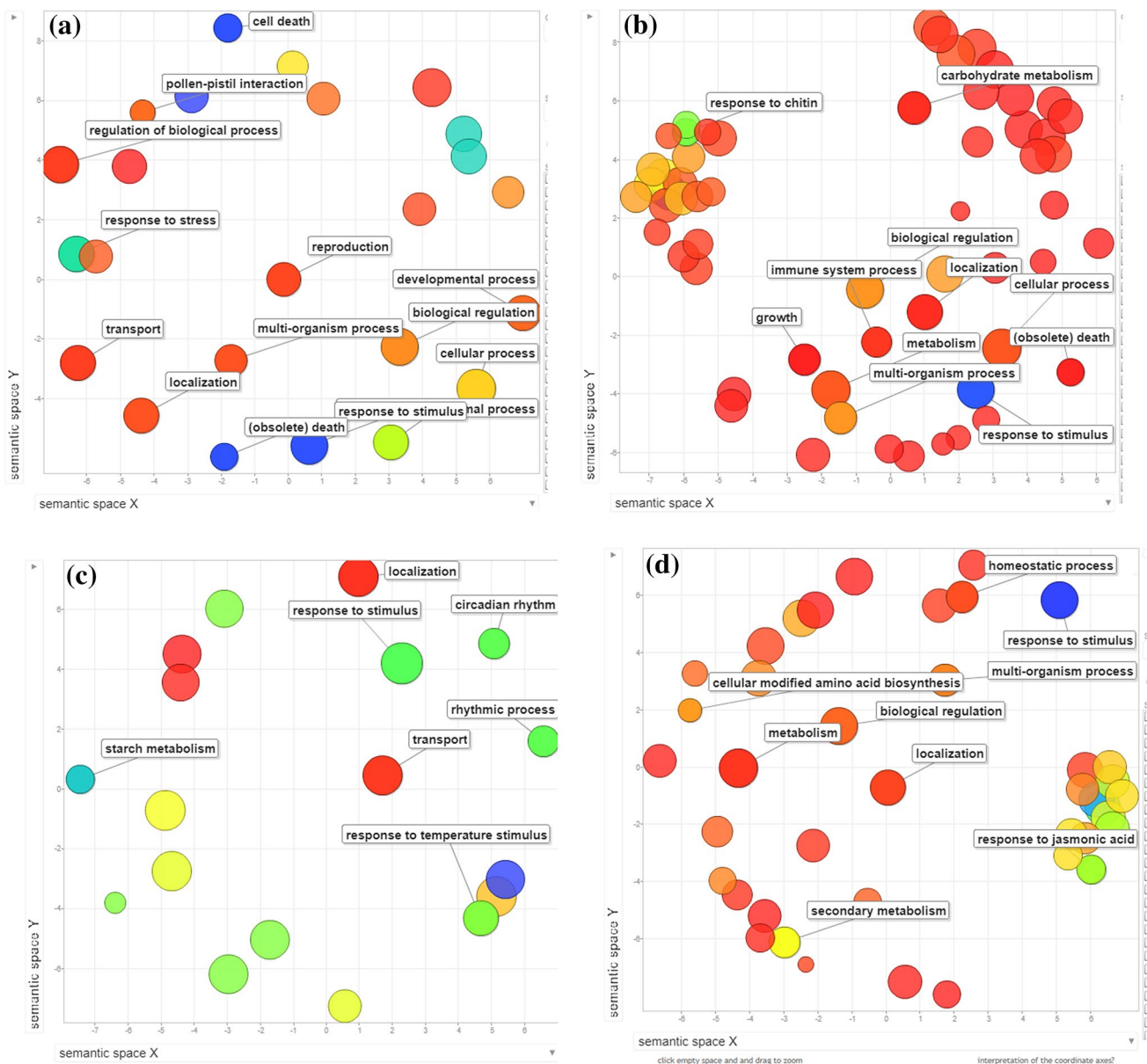


Fig. 5 Statistically significant gene ontology (biological process) terms associated with DEGs in **a** downregulated in roots, **b** upregulated in roots, **c** downregulated in shoots, **d** upregulated in shoots as

pathogenesis-related protein 1 (PR1) is an SA response gene. NPR1 and PR1 were upregulated by Cd treatment.

Gene Expression Analysis Using Quantitative Real-Time PCR (RT-qPCR)

We performed RT-qPCR on 13 selected genes mapping to MAPK signaling, calcium signaling, plant hormone signal transduction, and GSH metabolism to validate the comparative expression analysis (Online resource 1). The RNA-seq data from comparative expression analysis and RT-qPCR

obtained from REVIGO at allowed similarity 0.5. Colours represent the p values (Blue and green bubbles represent more significant p values than the orange and red bubbles). (Color figure online)

data showed a positive correlation ($r=0.648$) which further confirms the results of the present study (Fig. 7).

Discussion

Cadmium Uptake and Sequestration is Facilitated Through Various Transporters

Metal transporters play important roles in plant nutrition and metal homeostasis (Nelson 1999). Furthermore, these

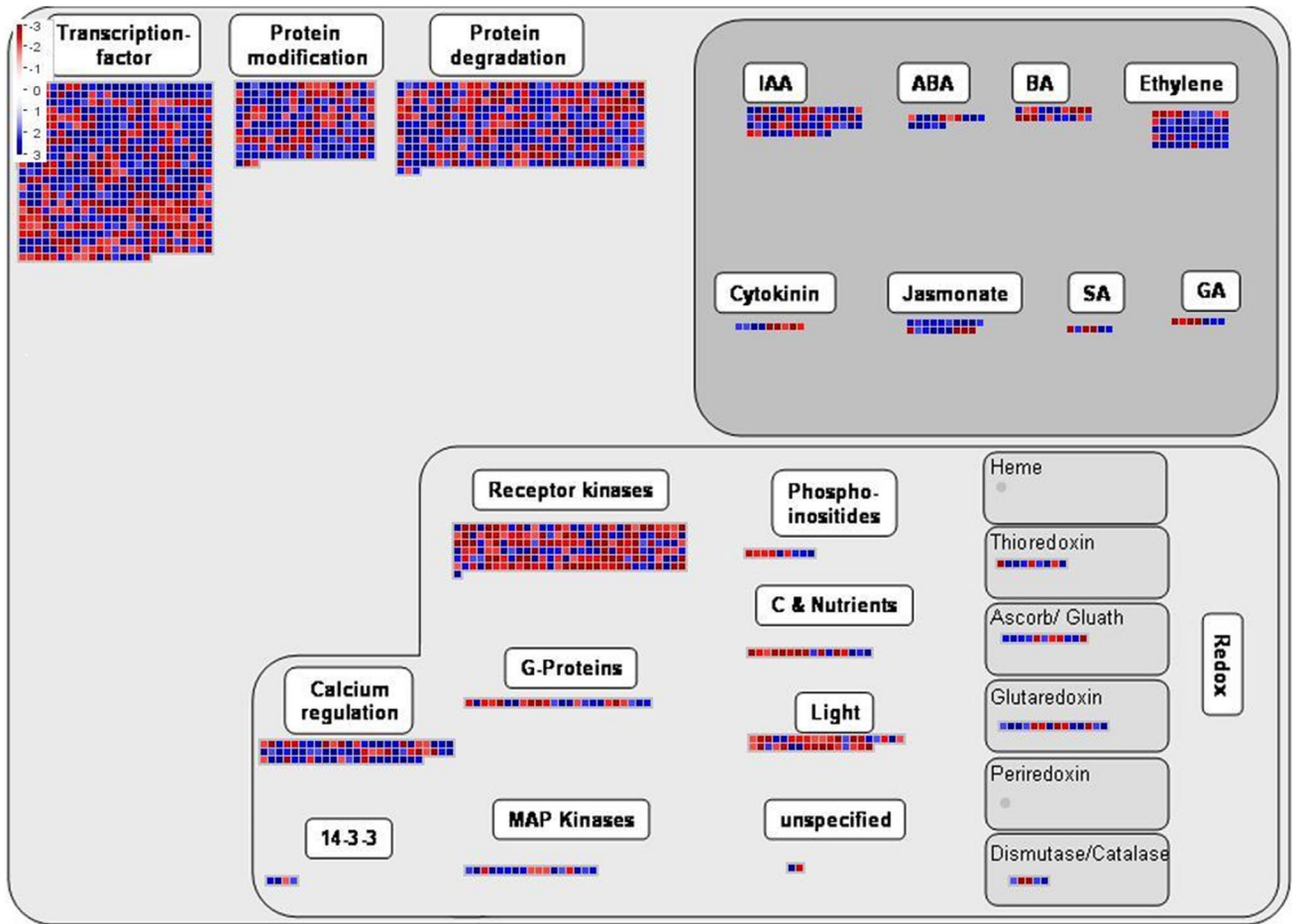
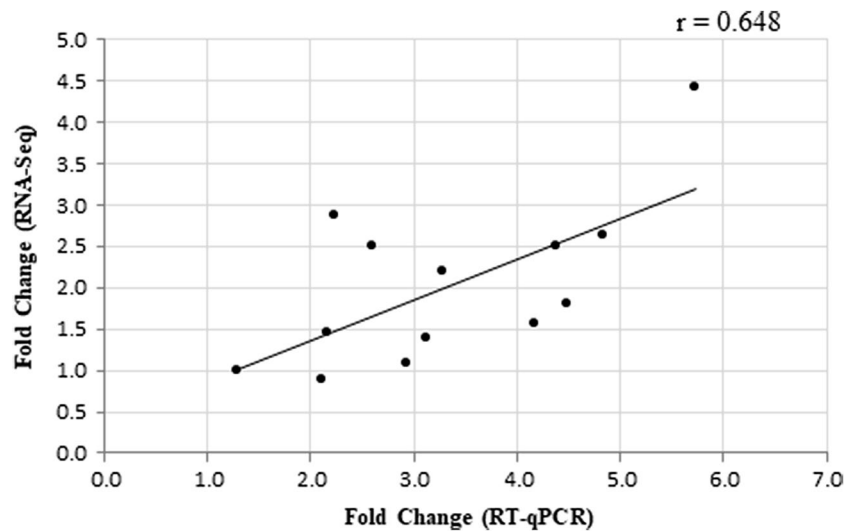


Fig. 6 MapMan regulation overview map showing the differential expression of transcripts under Cd treatment. Colour of small squares represents differential expression on the logarithmic scale, blue represents downregulated transcripts and red represents upregulated transcripts

Fig. 7 A positive correlation between RNA-seq and RT-qPCR data of selected genes



transporters mediate the uptake of metal ions from the surroundings and detoxification via vacuolar compartmentalization. Toxic metals like Cd can enter the plant cell via the transporters of essential elements like Ca, Zn and iron (Fe). The metal transporter encoding the gene Nramps in *Arabidopsis* can transport both Fe and Cd (Thomine et al. 2000). The ZIP gene family can transport a variety of metal ions including Cd, Zn, Mn and Fe (Guerinot 2000). The comparative expression analysis suggests that Nramp1 and ZIP8 could be the major Cd influx transporters present in the plasma membrane of Indian mustard. Further, the Cd²⁺ ions from the cytoplasm are then sequestered into vacuoles with the help of vacuolar transporters (Fu et al. 2017). Among the Zn transporters present in the vacuolar membrane, ZIF1 and MTPC2 could be involved in Cd sequestration in the present study. Besides the upregulation of transcripts encoding Cd-influx transporters, the upregulation of efflux transporters such as PCR2, PCR6, YSL2, and probable cadmium-transporting ATPase could also play an important role in improving Cd tolerance by pumping the toxic Cd ions out of the cell. The dual localization of AtPCR2 in the epidermis and vascular tissue attributes to its contrasting roles, that is, as Zn-efflux transporter and Zn-translocator (Song et al. 2010). Similarly, PCR2 can also aid in the translocation of Cd from root to shoot. YSL2 is known to transport nicotinamine (NA)-metal complexes laterally in the vasculature (DiDonato et al. 2004). The upregulation of transcripts encoding YSL2 is consistent with its role in imparting Cd tolerance. It is plausible that as a detoxification strategy, the gene expression of a few transporters such as HMA, IRT, ZRT/IRT-like protein1 and CAX1 was downregulated in the present study. The HMAs are type 1_B P-type ATPases and their role in maintaining metal homeostasis is well established (Miyadate et al. 2011; Williams and Mills 2005). Sequence analysis of these P-type ATPases revealed that HMA2, HMA3, and HMA4 in *Arabidopsis* are closely related and essential for Zn homeostasis (Cobbett et al. 2003). Several studies reported AtHMA2, AtHMA4 and OsHMA3 to be involved in xylem loading and translocating Cd from root to shoot (Hussain et al. 2004; Miyadate et al. 2011). IRT1 and ZRT/IRT-like protein1 are important metal homeostasis proteins known to transport metals including Cd, Fe and Zn (Vert et al. 2002). CAXs are antiporters shown to transport Cd into the vacuole and export H⁺ (Hirschi et al. 2000; Koren'kov et al. 2007).

Cadmium-Induced Oxidative Stress Management

The HM Cd is capable of producing oxidative stress in plants. Plant species that are tolerant to Cd tend to manage the oxidative stress via upregulating gene expression of various enzymatic and non-enzymatic antioxidants. The results of the present study are in line with the results of

several studies exhibiting the upregulation of antioxidants in response to Cd exposure. Among the enzymatic antioxidants CAT, DHAR, APX, and GR play important roles in alleviating oxidative stress and transduction of stress stimuli. Several studies have shown the importance of non-enzymatic antioxidants like flavonoids and coumarins in oxidative stress management (Agati et al. 2012; Gacche and Jadhav 2012; Kostova et al. 2011; Pietta 2000). The upregulation of these non-enzymatic antioxidants in the present study indicates their role in Cd-induced oxidative stress management. GSH is central in controlling the differential expression of several anti-oxidants including glutathione reductase, superoxide dismutase, phenylalanine ammonia lyase, and chalcone synthase. Several studies have reported a positive relation between Cd treatment, GSH content and GST activity (Nocito et al. 2011; Yang et al. 2018b; Zhand et al. 2013; Zhang and Ying 2008). Inside a plant cell, Cd-ions can bind to GSH and phytochelatin with high affinity leading to the formation of Cd-GSH/PC complexes which are then transported to vacuoles or apoplasts for detoxification. Glutathione S-transferase (GST) contributes to detoxification by catalyzing complexation of GSH with Cd and also helps in reducing the Cd-induced oxidative stress.

Cadmium Stress Perception and Signaling

Exposure to toxic substances leads to changes in the gene expression pattern to mitigate the stress stimuli. A number of signaling pathways lead to differential gene regulation. The mitogen-activated protein kinase (MAPK) pathway and calcium (Ca) signaling pathways are the important pathways involved in transduction of external stress stimuli to the nucleus for a pertinent response (Huang et al. 2012; Price et al. 1994). The MAPK signaling cascade modulates homeostasis of ROS, cell death, defense response to pathogens, stress adaptation and tolerance in plants. Cd-induced ROS is detected by OXI1 (oxidative signal-inducible 1) which further stimulates various serine/threonine kinases of MAPK signaling (Rentel et al. 2004). These proteins can stimulate the rapid signal transduction of a downstream mitogen-activated protein kinase (MAPK) cascade. In the present study, the MAPK signaling cascade activated a wide range of kinases and transcription factors such as WRKY 22/29 and 33, MYC2 and ERF1 which further can induce a wide array of stress and growth responses. Previous studies have also reported the activation of the MAPK signaling by cadmium (Lin and Aarts 2012; Yang et al. 2018a; Yeh et al. 2007). The concentration of Ca²⁺ and Ca-binding proteins (calmodulin; CALM) are known to be altered under abiotic stress (Yang and Poovaiah 2003). MCU increases cellular concentrations of Ca²⁺ and CALM binds to Ca²⁺ which brings about conformational changes in it. CALM and Ca are hypothesized to play roles in Cd signaling (Suzuki et al.

2001). The gene expression of CALM was upregulated in Cd-treated tissues which can regulate the plant's response to Cd via a variety of mechanisms including regulation of expression, ion transport, stress tolerance, and metabolism.

Plant Hormones Signaling in Response to Cadmium

Plant hormones are small signaling molecules known to regulate varied physiological processes including responses to biotic and abiotic stresses (Davies 2010). Several studies have exhibited their role in tolerance towards Cd stress in plants (Guo et al. 2007, Hsu and Kao 2003, 2005). Modulation of gene expression of phytohormone signaling components indicates their involvement in managing Cd stress in the present study. Results indicate that the plant hormones (IAA, ABA, JA, ethylene, BR, CK, and GA signaling) were active in response to Cd stress in Indian mustard. In rice under Cd stress, the phytohormones including CKs, IAA, ABA, JA, and GA were detected, suggesting their role in Cd stress response (Cai et al. 2015). IAA signaling was activated and the gene expression of an auxin-responsive gene belonging to the family GH3 and SAUR were upregulated which could be producing proteins involved in imparting Cd tolerance (Minglin et al. 2005). IAA was shown to alleviate Cd toxicity in eggplant by reducing Cd uptake and enhancing antioxidant activity/production (Singh and Prasad 2015).

The expression of transcripts mapping to components of ABA signaling was regulated under Cd stress. PP2C is a key repressor of ABA signaling and regulates its downstream responses. Out of the transcripts mapping to PP2C, 17 were upregulated and 2 downregulated which could be a possible mechanism to maintain homeostasis and regulating the ABA amount in response to Cd stress. ABA was shown to impart Cd tolerance via reducing root to shoot translocation of cadmium (Hsu and Kao 2003). JAs are known to play roles in Cd stress response (Singh and Shah 2014). JA was shown to alleviate Cd-induced oxidative stress via activation of antioxidant machinery and accumulation of phytochelatins (Maksymiec et al. 2007; Yan et al. 2013). MYC2 is a master regulator of JA signaling and involved in crosstalk between plant hormones such as ABA, SA, GA, and IAA. The upregulation of EIN3 and its target ERF1 in Cd-treated tissues exhibited that ethylene signaling is active under Cd stress. Activation of transcription factors like ERF1 then regulate the expression of downstream ethylene-responsive genes (Abozeid et al. 2017; Kendrick and Chang 2008). Ethylene is suggested to enhance the Cd stress response via controlling GSH content and the oxidative stress profile in *A. thaliana* (Schellingen et al. 2015). BRs are polyhydroxy steroidal plant hormones known to improve various abiotic stresses including Cd tolerance via increasing osmolytes such as proline content and increasing activities of antioxidative enzymes in *B.juncea* and *Phaseolus vulgaris* (Hayat

et al. 2007; Rady 2011). Cd treatment leads to downregulation of an enzyme involved in CKK synthesis (cytokinin synthase) and upregulation of an enzyme that catalyzes the degradation of cytokinins (cytokinin dehydrogenase; CKX). The decrease in cytokinin (CKK) contents aid in reducing transpiration driven Cd translocation and hence providing tolerance towards Cd toxicity (Veselov et al. 2003).

Conclusion

Indian mustard exhibited the potential for Cd remediation as evident from bioaccumulated of a large amount of Cd in shoots. Huge reprogramming at the transcriptomic level was exhibited by Cd treatment. Various transporters involved in Cd uptake and efflux were identified. GSH is involved in detoxification of Cd ions via sequestration into vacuoles. MAPK signaling and Ca signaling were found to play important roles in Cd stress perception. Furthermore, plant hormone signaling was found to be affected by Cd which contributes to Cd tolerance through varied ways such as enhancing osmolyte content, antioxidant enzyme activity or reduction in root to shoot Cd translocation. The present study has provided a thorough understanding of the molecular mechanisms involved in Cd uptake, tolerance, and detoxification in Indian mustard.

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Author Contributions PB designed and conceived of the study. ST analyzed and interpreted data, drafted the manuscript. SC helped in the acquisition of data and data analysis. PB and SC coordinated further in improving the manuscript. All the authors have read and approved the final manuscript.

Data Availability The raw sequence data of the libraries are available from the NCBI SRA under the study with Accession Nos.: SRP152398 and SRP126203.

Compliance with Ethical Standards

Conflict of interest Authors declare that there is no conflict of interest.

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