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Pim de Voogt *Editor*

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Foreword

International concern in scientific, industrial, and governmental communities over traces of xenobiotics in foods and in both abiotic and biotic environments has justified the present triumvirate of specialized publications in this field: comprehensive reviews, rapidly published research papers and progress reports, and archival documentations. These three international publications are integrated and scheduled to provide the coherency essential for nonduplicative and current progress in a field as dynamic and complex as environmental contamination and toxicology. This series is reserved exclusively for the diversified literature on “toxic” chemicals in our food, our feeds, our homes, recreational and working surroundings, our domestic animals, our wildlife, and ourselves. Tremendous efforts worldwide have been mobilized to evaluate the nature, presence, magnitude, fate, and toxicology of the chemicals loosed upon the Earth. Among the sequelae of this broad new emphasis is an undeniable need for an articulated set of authoritative publications, where one can find the latest important world literature produced by these emerging areas of science together with documentation of pertinent ancillary legislation.

Research directors and legislative or administrative advisers do not have the time to scan the escalating number of technical publications that may contain articles important to current responsibility. Rather, these individuals need the background provided by detailed reviews and the assurance that the latest information is made available to them, all with minimal literature searching. Similarly, the scientist assigned or attracted to a new problem is required to glean all literature pertinent to the task, to publish new developments or important new experimental details quickly, to inform others of findings that might alter their own efforts, and eventually to publish all his/her supporting data and conclusions for archival purposes.

In the fields of environmental contamination and toxicology, the sum of these concerns and responsibilities is decisively addressed by the uniform, encompassing, and timely publication format of the Springer triumvirate:

Reviews of Environmental Contamination and Toxicology [Vol. 1 through 97 (1962–1986) as Residue Reviews] for detailed review articles concerned with any aspects of chemical contaminants, including pesticides, in the total environment with toxicological considerations and consequences.

Bulletin of Environmental Contamination and Toxicology (Vol. 1 in 1966) for rapid publication of short reports of significant advances and discoveries in the fields of air, soil, water, and food contamination and pollution as well as methodology and other disciplines concerned with the introduction, presence, and effects of toxicants in the total environment.

Archives of Environmental Contamination and Toxicology (Vol. 1 in 1973) for important complete articles emphasizing and describing original experimental or theoretical research work pertaining to the scientific aspects of chemical contaminants in the environment.

The individual editors of these three publications comprise the joint Coordinating Board of Editors with referral within the board of manuscripts submitted to one publication but deemed by major emphasis or length more suitable for one of the others.

Coordinating Board of Editors

Preface

The role of *Reviews* is to publish detailed scientific review articles on all aspects of environmental contamination and associated (eco)toxicological consequences. Such articles facilitate the often complex task of accessing and interpreting cogent scientific data within the confines of one or more closely related research fields.

In the 50+ years since *Reviews of Environmental Contamination and Toxicology* (formerly *Residue Reviews*) was first published, the number, scope, and complexity of environmental pollution incidents have grown unabated. During this entire period, the emphasis has been on publishing articles that address the presence and toxicity of environmental contaminants. New research is published each year on a myriad of environmental pollution issues facing people worldwide. This fact, and the routine discovery and reporting of emerging contaminants and new environmental contamination cases, creates an increasingly important function for *Reviews*. The staggering volume of scientific literature demands remedy by which data can be synthesized and made available to readers in an abridged form. *Reviews* addresses this need and provides detailed reviews worldwide to key scientists and science or policy administrators, whether employed by government, universities, nongovernmental organizations, or the private sector.

There is a panoply of environmental issues and concerns on which many scientists have focused their research in past years. The scope of this list is quite broad, encompassing environmental events globally that affect marine and terrestrial ecosystems; biotic and abiotic environments; impacts on plants, humans, and wildlife; and pollutants, both chemical and radioactive; as well as the ravages of environmental disease in virtually all environmental media (soil, water, air). New or enhanced safety and environmental concerns have emerged in the last decade to be added to incidents covered by the media, studied by scientists, and addressed by governmental and private institutions. Among these are events so striking that they are creating a paradigm shift. Two in particular are at the center of ever increasing media as well as scientific attention: bioterrorism and global warming. Unfortunately, these very worrisome issues are now superimposed on the already extensive list of ongoing environmental challenges.

The ultimate role of publishing scientific environmental research is to enhance understanding of the environment in ways that allow the public to be better informed or, in other words, to enable the public to have access to sufficient information. Because the public gets most of its information on science and technology from internet, TV news, and reports, the role for scientists as interpreters and brokers of scientific information to the public will grow rather than diminish. Environmentalism is an important global political force, resulting in the emergence of multinational consortia to control pollution and the evolution of the environmental ethic. Will the new politics of the twenty-first century involve a consortium of technologists and environmentalists, or a progressive confrontation? These matters are of genuine concern to governmental agencies and legislative bodies around the world.

For those who make the decisions about how our planet is managed, there is an ongoing need for continual surveillance and intelligent controls to avoid endangering the environment, public health, and wildlife. Ensuring safety-in-use of the many chemicals involved in our highly industrialized culture is a dynamic challenge, because the old, established materials are continually being displaced by newly developed molecules more acceptable to federal and state regulatory agencies, public health officials, and environmentalists. New legislation that will deal in an appropriate manner with this challenge is currently in the making or has been implemented recently, such as the REACH legislation in Europe. These regulations demand scientifically sound and documented dossiers on new chemicals.

Reviews publishes synoptic articles designed to treat the presence, fate, and, if possible, the safety of xenobiotics in any segment of the environment. These reviews can be either general or specific, but properly lie in the domains of analytical chemistry and its methodology, biochemistry, human and animal medicine, legislation, pharmacology, physiology, (eco)toxicology, and regulation. Certain affairs in food technology concerned specifically with pesticide and other food-additive problems may also be appropriate.

Because manuscripts are published in the order in which they are received in final form, it may seem that some important aspects have been neglected at times. However, these apparent omissions are recognized, and pertinent manuscripts are likely in preparation or planned. The field is so very large and the interests in it are so varied that the editor and the editorial board earnestly solicit authors and suggestions of underrepresented topics to make this international book series yet more useful and worthwhile.

Justification for the preparation of any review for this book series is that it deals with some aspect of the many real problems arising from the presence of anthropogenic chemicals in our surroundings. Thus, manuscripts may encompass case studies from any country. Additionally, chemical contamination in any manner of air, water, soil, or plant or animal life is within these objectives and their scope.

Manuscripts are often contributed by invitation. However, nominations for new topics or topics in areas that are rapidly advancing are welcome. Preliminary communication with the Editor-in-Chief is recommended before volunteered review manuscripts are submitted. *Reviews* is registered in Web of Science™.

Inclusion in the Science Citation Index serves to encourage scientists in academia to contribute to the series. The impact factor in recent years has increased from 2.5 in 2009 to almost 4 in 2013. The Editor-in-Chief and the Editorial Board strive for a further increase of the journal impact factor by actively inviting authors to submit manuscripts.

Amsterdam, The Netherlands
January 2015

Pim de Voogt

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Gene Expression Profiling in Fish Toxicology: A Review

Girish Kumar and Nancy D. Denslow

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1 Introduction

Pollution of the aquatic environment, due to the growing levels of contaminants associated with human activity such as industry, mining, agriculture, and household waste production, has become a serious problem worldwide (Conti et al. 2012; Copat et al. 2012a, b). Aquatic organisms, such as fish, accumulate pollutants directly from contaminated water and indirectly via the food chain (Sasaki et al. 1997). Bio-concentration of potentially harmful substances—e.g. heavy metals, pesticides, polychlorinated biphenyls (PCBs), polycyclic aromatic hydrocarbons (PAHs) etc., in fish present a major threat to human health. Since fish occupy the top of the aquatic food chain, they are widely used as a bio-indicator to evaluate the health of aquatic ecosystem (Camargo and Martinez 2007).

Use of fish in toxicity studies can be traced back to 1800s when Penny and Adams (1863) used the chemicals (used in dye-works) to study their effects on goldfish (*Carassius auratus*). Although basic research in toxicology began in the 1800s, test methodology was established, especially on fish, only in the 1940s (Hunn 1989). Subsequently, considerable effort was made and a standardized test method was published in 1960 (American Public Health Association et al. 1960). Since the 1960s, rapid development in toxicity testing changed toxicology from a descriptive science to one in which mechanisms of action of toxicants are recognized. In subsequent decades, with increased emphasis on the use of molecular biology techniques, toxicological studies further expanded in many areas including chemical carcinogenesis and xenobiotic metabolism, among many others. In recent years, analysis of gene expression has become the method of choice for many toxicological studies. This is due to the fact that toxicant exposure often alter gene expression profile as cells regulate certain genes to protect cellular structures and repair damage (Causton et al. 2001). Analysis of these gene expression data can provide information regarding contaminant exposure and potential effects at higher biological levels (Garcia-Reyero et al. 2008). Gene expression profile, therefore, provides a sensitive measurable endpoint for toxicity and thus can serve as early warning of a specific biological endpoint.

Until 1992, Northern blot and subtractive hybridization were the only methods used in gene expression analysis. Although these methods are still quite useful and reliable, they are tedious, time consuming and difficult to perform. Additionally, they require large amounts of mRNA that can be limited in many situations. In 1992, polymerase chain reaction (PCR) based alternative methods—real time quantitative PCR (RT-qPCR) (Porcher et al. 1992) and differential display reverse transcription PCR (DDRT-PCR) (Liang and Pardee 1992) were developed. These methods allow the identification of differentially expressed genes in a less labor-intensive fashion than the methods mentioned above. RT-qPCR now represent the method of choice for analyzing gene expression of a small number of genes in thousands of samples. In the last decade, use of microarrays and next-generation sequencing (NGS) technologies paved the way for a paradigm shift from the analysis of a selected set of genes to a whole-genome screening of potentially all

expressed genes. However, so-called close-end transcriptomic technique, microarray require prior knowledge of gene sequences and are therefore unsuitable for non-model species where limited or no sequence information is available. To overcome this limitation, pyrosequencing technology was utilized to create a transcriptome-wide microarray from a non-model species (Garcia-Reyero et al. 2008). Nevertheless, in recent years, NGS technology has become a method of choice for transcriptome profiling due to its decreasing costs and high-throughput.

In past, many reviews were published to discuss and describe the various aspects of fish toxicology using a particular model species (Ankley and Villeneuve 2006), technique (Ju et al. 2007; Williams et al. 2014) or toxicant (Larkin et al. 2003; Garcia-Reyero and Denslow 2006). However, no attempt has been made to present a comprehensive review on application of gene expression in fish toxicology. Therefore, the objective of this paper is to review the application of gene expression profiling techniques in fish toxicology and to discuss the biological insights already gained from these techniques in fish species.

2 Use of Northern Blotting in Fish Toxicology

Northern blotting was one of the first molecular biology techniques used in fish toxicology to determine the relative abundance of gene transcripts in response to toxicants (Chen 1983). In this technique, electrophoretically separated bands of RNA are hybridized with DNA fragments labeled with radioactive nucleotides complementary to the gene(s) of interest to determine the differentially expressed gene (Alwine et al. 1977).

In the beginning, Northern blot analysis was used to study the transcriptional induction of vitellogenin (VTG) gene in response to endocrine disrupting chemical (EDC), 17 β -estradiol (E₂) (Chen 1983). Since then, use of Northern blots have been expanded to a large number of EDCs to study the gene expression profile of many biomarker genes (Flouriot et al. 1997; Arukwe et al. 2001, 2002; Bowman et al. 2002; Yadetie and Male 2002; Larkin et al. 2003).

Vitellogenin, the egg yolk precursor protein, is synthesized in all oviparous vertebrates. It is normally produced by females in response to estradiol during oogenesis. However, it also gets synthesized in livers of males upon exposure to estrogen or an estrogen mimic chemical. This characteristic of VTG has been explored using several fish species for detecting the presence of xenoestrogens from contaminated aquatic environments (Sumpter and Jobling 1995; Folmar et al. 1996; Denslow et al. 1997a, b). In order to find out the presence of estrogenic compounds in effluents of pulp and paper mills, juvenile whitefish (*Coregonus lavaretus* L. s.l.) were exposed to diluted effluents from three operating pulp, paper, and paperboard mills in Southern Lake Saimaa, Finland (Mellanen et al. 1999). Northern blot analysis showed increased VTG mRNA levels suggesting that the effluent of pulp and paper mills were a source of estrogenic contaminants. In

addition to the VTG mRNA, the presence of estrogen or estrogen mimic chemicals in aquatic environments also cause increase in expression levels of choriogenin mRNA (Murata et al. 1997; Arukwe et al. 2001) and estrogen receptor (ER) mRNA (Flouriot et al. 1997; Bowman et al. 2002). Therefore, these genes can be used as a potential biomarker to identify the presence of estrogenic chemicals in aquatic environment.

Alkylphenols such as 4-nonylphenol (NP) are one of the wide variety of environmental chemicals that have estrogenic effects. The estrogenic effects of NP have been assessed through gene expression levels of VTG and zona radiata (ZR) proteins (Jobling et al. 1996; Arukwe et al. 1997; Oppen-Berntsen et al. 1999; Jung et al. 2006). However, to enhance the understanding of the mechanisms of actions and to develop new potential biomarkers for xenoestrogens, Yadatie and Male (2002) studied the effects of NP (50 and 125 mg/kg body weight) on mRNA levels of leutinizing hormone beta (LH β) in male and female juvenile of Atlantic salmon (*Salmo salar*). Northern blot analysis revealed significant induction of mRNA levels of LH β in female juvenile salmonids (up to sixfold), suggesting that NP has the potential to perturb the regulation of LH β gene expression by mimicking E₂ and thus can be used as a biomarker for pituitary effects of xenoestrogens. Nevertheless, one should always take into consideration that too much circulating E₂ can signal the pituitary to shut down by feedback inhibition (Kriegsfeld et al. 2006).

Northern blot analysis has also been used to study the down-regulation of biomarker genes in response to exposure to chemicals. For instance, treatment of mature female tilapia (*Oreochromis niloticus*) with high levels of androgen (17- α -methyltestosterone) (20 mg/kg, in 50 % ethanol/saline (5 mg/ml)) resulted in a pronounced decline in VTG mRNA levels and serum E₂ level (Lazier et al. 1996). These results suggest the presence of an inhibitory site of action at the hypothalamic-pituitary axis, probably through an androgen receptor or through an estrogen receptor after local aromatization of 17 α -methyltestosterone (Lazier et al. 1996). In another study, authors observed reduction in transthyretin (TTR) expression in males of gilthead seabream (*Sparus aurata*) after estrogen treatment (Funkenstein et al. 2000). Furthermore, Northern blot analysis of hepatic retinol-binding protein (RBP) revealed reduced level of mRNA expression in adult male gilthead seabream (Funkenstein 2001) and rainbow trout (Sammar et al. 2001) after E₂ treatment. Since the E₂ levels in fish during the period between vitellogenic growth and final oocyte maturation decreases, the down-regulation of RBP and TTR genes in response to estrogen treatment in the above studies indicate the role of TTR and RBP in transport of maternal thyroid hormones and retinol (Funkenstein 2000, 2001).

Cytochrome P450s play a pivotal role in metabolism of toxic compounds in vertebrates and hence the expression of P450s influenced by exposure to xenoestrogens and other environmental contaminants (Williams et al. 1998). The effect of E₂ and testosterone on expression of P450s (CYP2M1, CYP2K1, CYP3A27) was studied in rainbow trout (Buhler et al. 2000). Gene expression analysis suggested that exposure of fish to endocrine disruptors can adversely affect a number of physiological processes through mechanisms involving altered levels of expression

of specific P450 isozymes mRNA (Buhler et al. 2000). Furthermore, a higher levels of cytochrome P450 1A1 (CYP1A) mRNA was observed when fish were exposed to β -naphthoflavone (Leaver and George 2000) and polybrominated diphenyl ethers (Boon et al. 2002). The reason for increased levels of CYP1A transcripts in above mentioned studies is due to the fact that polycyclic aromatic hydrocarbons (PAHs) transcriptionally activate CYP1A by binding to Aryl hydrocarbon Receptor (AhR). In another study, Ferraris et al. (2005) attempted to investigate the molecular mechanisms of the dicarboximide fungicide iprodione (Ip) in rainbow trout. Findings of the study revealed that Ip-mediated CYP1A3 gene induction involves the activation of AhR complex via phosphorylation-dephosphorylation reactions.

Northern blot analysis is not only used to measure expression level of mRNA but also to validate gene induction profiles of DDRT-PCR and microarrays. One of the main advantages of this method is that it can detect the size of the mRNA and alternatively spliced transcripts (if present). However, this technique can be used to measure the expression levels of only a small number of genes. In addition, it is labor intensive and have limited resolution. Therefore in recent years, it has been largely replaced by newer methods, such as DDRT-PCR, SSH, RT-qPCR, microarrays and NGS.

3 Use of DDRT-PCR in Fish Toxicology

The technique of DD RT-PCR was developed by Liang and Pardee in 1992. Since then this method has gained popularity among researchers and is now used to identify and isolate differentially expressed genes in several experimental systems. In this technique reverse transcription of mRNA is done with anchored oligo-dT primers, followed by PCR amplification using anchor primer and a limited number of short arbitrary primers. Amplified products are then visualized either by radio-activity or fluorescence.

In fish toxicology, DDRT-PCR has been applied to examine the alterations in gene expression following chemical exposure in natural and experimental environments. For example, gene expression patterns in sheepshead minnows (*Cyprinodon variegatus*) were determined thorough *in vivo* exposure to natural and anthropogenic estrogens (Denslow et al. 2001a, b). Based on the results authors concluded that E₂, EE₂ (17 α -ethinyl estradiol), and DES (diethylstilbestrol) act through the same pathway when administered through the water at low concentrations. In largemouth bass changes in gene expression induced by exposure to natural estrogen (Bowman et al. 2002) and paper and pulp mill effluents (Denslow et al. 2004) were measured. Differential expression of genes (VTG, ER, ERp72 ZP2, fibrinogen γ) in response to E₂ at different time series (0, 6, 12, 24, 48 h) confirmed the idea that gene regulation by E₂ is complicated and time dependent (Bowman et al. 2002). Measurements of changes in gene expression induced by

exposure to paper mill effluents resulted in an up-regulation of CYP1A and down-regulation of genes normally expressed during the reproductive season in females. These antiestrogenic changes suggest that effluents of paper and pulp mills could decrease the reproductive success in affected populations (Denslow et al. 2004).

Carginalo et al. (2002) investigated the ability of cadmium (Cd) to affect gene transcription in Antarctic icefish (*Chionodraco hamatus*) using DDRT-PCR. Results indicated that Cd has both inhibitory effect as well as positive modulator of gene expression. The inhibitory effect of Cd may be due to an improper folding of zinc finger domains and/or oxidation of the cysteine residues while positive effect may be attributed to the fact that binding of Cd to zinc containing proteins release this metal which act as a positive modulator of gene expression. DDRT-PCR has also been used to study the contaminant effects of PAHs such as anthracene (Peterson and Bain 2004) and pyrene (Roling et al. 2004) in mummichogs (*Fundulus heteroclitus*). Genes demonstrating differential expression after PAHs (pyrene and anthracene) exposure suggested that multiple biomarkers are required to get insight into the mechanism of toxicity and information on potential effects of toxicants. In another study, altered gene expression in killifish (*F. heteroclitus*) was studied by collecting the samples from a highly contaminated site (Elizabeth River, VA, USA), and non-contaminated reference site (King's Creek, VA, USA). Seventy-four differentially expressed mRNAs were identified including sex and population specific suggesting that sex of the fish and population differentiation play a role in response to contaminant exposure (Meyer et al. 2005).

In tilapia, lindane exposure caused differential expression of 50 different genes, four of these showed homology with immunoglobulin heavy chain, coagulation factor V, casein kinase 2 alpha, and receptor protein- tyrosine-like phosphatase (Colli-Dula et al. 2009). These findings suggest that lindane triggers the expression of genes that are involved in immunological and stress responses in fish. Woźny et al. (2012) used DD PCR to study the molecular background of zearalenone (ZEA) toxicity in rainbow trout. A total of 59 differentially expressed cDNA were identified, five of these genes (β -contractin, bccip, α -enolase, protein C, and ferritin heavy chain subunit) were successfully confirmed by RT-qPCR. Differential expression of above mentioned genes indicates that ZEA interferes with blood coagulation and different cellular components (i.e. cytoskeleton).

The DD RT-PCR is an effective and powerful approach for detecting novel genes involved in toxicant effect pathways and identifying new molecular biomarkers. However, it has not been widely applied in fish toxicology research due to its several inherent limitations. One of the major limitations is the large percentage of false-positive results generated by the PCR step. In addition, the identification of differentially expressed genes remains tedious and confirmation of the gene expression is required by RT-qPCR. For these reasons, in the mid-2000s, DDRT-PCR technique was superseded by RT-qPCR, DNA microarrays and RNA-Seq.

4 Use of SSH in Fish Toxicology

Suppression subtractive hybridization (SSH) is a method used to distinguish between two mRNA samples that are either overexpressed or exclusively expressed in one sample compared to another. This technique is based on selective amplification of target cDNA and simultaneously suppressing the amplification of non-target cDNA (Diatchenko et al. 1996). SSH combines normalization and subtraction in a single procedure. This dramatically increases the probability of obtaining low-abundance differentially expressed cDNA. However, one of the major limitations of SSH is that it compares gene transcripts from only two samples at one time and therefore it is time consuming and labor intensive. Another limitation is that the process is leaky and thus increases the probability of obtaining non-specific genes. Therefore, everything needs to be validated by RT-qPCR. Nevertheless, SSH has been employed by many research groups to investigate various aspects of aquatic toxicology. For example, Brown et al. (2004) studied the response of male plaice (*Pleuronectes platessa*) to EE₂ exposure. The results showed a selective down-regulation of some egg proteins (VTG, ZRP) mRNA responsible to maintain the quality of eggs. Thus, exposure of fish to EE₂ may cause reproductive failure by diminishing the quality of eggs. In the subsequent year, Volz et al. (2005) assessed the qualitative gene expression changes in whole brain, liver, and testis of adult male Japanese medaka in response to 2, 3, 7, 8-Tetrachlorodibenzo-*p*-dioxin (TCDD). CYP1A mRNA was found significantly higher in TCDD-exposed brain and liver. However, testis specific CYP1A mRNA was not significantly induced, suggesting that TCDD has tissue specific effect on fish. In another study, Li et al. (2008) examined the effects of exposure to polychlorinated biphenyls (PCBs) on false kelpfish (*Sebastes marmoratus*). Functional analysis of differentially expressed genes were found to be associated with long-term potentiation and neurotransmitter release, neuroendocrine, mitosis and cell proliferation, energy-related metabolism, general metabolism, signal protein, hemopoiesis system and immune system. However, only 45 of the 108 gene sequences could be identified with homologous database sequences. Therefore, further investigation is required to understand the effects of PCB-induced toxicity in fish. Furthermore, SSH was applied to isolate and enrich the differentially expressed genes from juvenile samples of European seabass (*Dicentrarchus labrax*) collected at the reference site and at a contaminated site. Altered expression in genes involved in energy metabolism, immune system activity and antioxidant response was identified (Nogueira et al. 2010). Based on the results, the authors suggested that these changes due to environmental contamination could affect the population in future.

In order to study the toxic effects of Cd, Dangre et al. (2010) identified over 700 Cd responsive cDNAs in sheepshead minnow. These genes were found to be broadly involved in protective and repair functions as well as cellular and metabolic processes. In addition, the authors found that Cd exposure down-regulated both embryonic α and β globin in sheepshead minnow. These results support the

perspective that Cd exerts its toxicity through the modulation of multiple physiological processes.

To elucidate the genes in response to tetrodotoxin (TTX), Matsumoto et al. (2011) examined the hepatic gene expression profile of marine puffer fish (*Takifugu rubripes*) by intramuscular administration of 0.50 mg TTX/kg. A high level of up-regulation was observed in hepcidin precursor gene. In addition, induction of complement C3, serotransferrin, apolipoprotein A-1, high temperature adaptation protein Wap65-2, complement C7, fibrinogen beta chain, and heat-shock protein 4 were also obtained. Increased expression of these genes confirmed that the exposure of TTX increases the gene expression of the acute-phase response proteins. SSH was also used to identify alterations in gene transcription of silver carp (*Hypophthalmichthys molitrix*) and bighead carp (*Aristichthys nobilis*) after exposure to Microcystin-LR (MCLR). In silver carp, sequencing analysis and homology searches revealed 75 unique differentially expressed genes in response to MCLR (Qu et al. 2011). Of the 75 unique genes, 38 were identified with functions of immune response, transport and cell metabolism. Some genes (*Fs59*, *Fs70*, *Rs15*, and *Rs2*) showed the time-dependent transcriptional in silver carp livers. In the case of bighead carp, Cai et al. (2012) tested the gene expression profile of cytidine deaminase (*cda*) gene. Altered expression profile of the *cda* gene was inversely proportional to time and directly proportional to dose of MCLR in most of the tested tissue. These findings indicate that the effects of MCLR are highly complex and in future more cellular pathways affected by MCLR could be identified.

5 Use of RT-qPCR in Fish Toxicology

Reverse transcription-quantitative PCR (RT-qPCR) is an accurate and sensitive tool to analyze and validate differential gene expression. This technique came into light in early 1990s (Higuchi et al. 1993) and became popular among researchers for analyzing gene expression of a moderate number of genes in anywhere from a small number to thousands of samples. Despite the limitation in the number of genes that can be analyzed per sample, RT-qPCR has been widely used to study the toxic effects of a large number of chemical pollutants including heavy metals, natural and synthetic steroids, pesticides, herbicides, fungicides, PCBs, perfluorononanoic acid (PFNA), PAHs, dioxins, microcystin, brominated and organophosphorus flame retardants, pharmaceuticals and personal care products, and manufactured nanoparticles.

5.1 Endocrine Disrupting Chemicals (EDCs)

To determine the effects of EDCs on gene expression, Celius et al. (2000) quantified the mRNA levels in E₂ treated (0.01, 0.1, 1.0 or 10 mg/kg body weight) rainbow trout and reported induction of ZR and VTG mRNA. However, induction

of ZR-gene occurred at all concentrations of E₂ while VTG was only significantly induced at the highest dose (10 mg/kg). The more sensitive nature of ZR suggests that ZR-gene is more suitable indicator than VTG to estrogenic substances. Hogan et al. (2008) investigated the sensitivity of three hormone responsive genes (VTG, spiggin, AR β) in threespine stickleback (*Gasterosteus aculeatus*) by exposing the fish to 1, 10, and 100 ng/L of methyltestosterone or E₂ for 7 days. Tissue-specific induction in expression of VTG (liver) and spiggin (kidney) mRNA in male and female fish respectively suggest that expression pattern of these genes can be used to identify compounds causing reproductive dysfunction in fishes.

Furthermore, effects of EDCs (bisphenol A, nonylphenol, octylphenol) and trace metals (Cd, Cu, and Zn) were investigated in different tissues (brain, eye, gonad, intestine, liver, muscle, and skin), development stages (embryonic and larval), and gender types (hermaphrodites and secondary males) of mangrove killifish (*Kryptolebias marmoratus*) (Rhee et al. 2009). The RT-qPCR analysis showed suppressive effects of EDCs on expression of metallothionein (MT) gene. Based on the findings, authors suggested that the expression level of MT may be affected if metals are present in water along with EDCs. In another study, exposure of sex steroids (estradiol, 2-methyl-testosterone) in Chinese rare minnow (*Gobiocypris rarus*) showed up-regulation of *foxl2* gene after treatment with estradiol and down-regulation after treatment with 2-methyl-testosterone (Jiang et al. 2011). These results suggest that *foxl2* mRNA expression is regulated by sex hormones, and therefore *foxl2* can be used as a molecular indicator in monitoring for environmental endocrine disruptors.

In an attempt to study the effects of E₂ (10, 100, and 1000 μ g/L) on stress related genes, Woo et al. (2012) investigated the transcriptional changes in intestinal, liver and muscle tissues of Japanese medaka. RT-qPCR analysis showed a tissue specific alteration in mRNA levels of catalase (CAT), glutathione peroxidase and ubiquitin genes. The differential expression of a range of stress-related genes in this study suggests that transcriptional changes in these genes could be used as a rapid assay to study the effects of E₂ exposure (Woo et al. 2012).

Attempts were also made to study the effects of EDCs on embryonic development and sexual differentiation. For instance, Lei et al. (2013) used RT-qPCR to evaluate reproductive toxicities in Japanese medaka by exposing the fertilized eggs to 1, 10, 100, and 1000 ng/L of β -estradiol 17-valerate (EV) for 15 days. Results showed an up-regulation of VTG-I in liver of males while down-regulation of ER α and VTG-I in liver of female individuals. These findings suggest that EV is a reproductive toxicant and cause reproductive failure in fish species.

In another study, zebrafish (*Danio rerio*) embryos and larvae were used to assess how EDCs (EE₂, genistein, fadrozole) interfere with embryogenesis (Santos et al. 2014). RT-qPCR analysis revealed alterations in the expression levels of the hormone receptors genes (*esr* and *ar*) and apoptosis-related genes (*p53* and *c-jun*) at different time points (between 2 and 144 h post-fertilization), suggesting a bidirectional communication between hormones action and apoptosis pathways on early development. These results also suggest that exposure to EDCs poses a threat to

aquatic organisms during embryogenesis by impairing physiological processes essential to a normal development and, consequently, survival.

In a more recent study, Schiller et al. (2014) proposed a novel method of determining the effects of substances with estrogenic and anti-androgenic activity by comparing the embryos of two species (medaka and zebrafish). Patterns of gene expression between the two species were compared by exposing the embryos (medaka: 7-day post fertilization; zebrafish: 48 and 96 h post fertilization) to estrogenic (EE₂) and anti-androgenic (flutamide) reference compounds and five test compounds with endocrine activities (bisphenol A, genistein, prochloraz, linuron, and propanil). Estrogenic responses were comparable between the two species while responses to anti-androgenic chemicals differed between medaka and zebrafish. While responses to anti-androgenic chemicals were more sensitive in medaka embryos, a broader range of morphological effects were observed in zebrafish embryos. Therefore, together medaka and zebrafish embryos would be beneficial for testing effects of EDCs in early life states.

5.2 Metals, Metallic Nanoparticles and Organometals

To assess the risk of metal ions in fishes inhabiting metal contaminated area, MT gene has been used as a biomarker. For example, metal ions (Cu²⁺, Cd²⁺, Hg²⁺, Ni²⁺, Pb²⁺, Zn²⁺) exposure in *Tilapia* showed increased hepatic and gills MT mRNA levels. However, renal MT mRNA levels did not show significant induction in response to Cd²⁺ and Zn²⁺ metal ions, suggesting that gene expression of MT is tissue dependent (Cheung et al. 2004). In another study, Chan et al. (2006) investigated the induction of MT gene in zebrafish following the administration of different metal ions (Cu²⁺, Cd²⁺, Hg²⁺, and Zn²⁺). RT-qPCR analysis showed highest level of MT mRNA induction (40–50 fold) in embryo-larvae in response to Hg²⁺ followed by Cd²⁺. However, in the *in vitro* study of liver cell-line (ZFL), Cd²⁺ was the most potent inducer of MT mRNA (up to 250 folds), followed by Cu²⁺ and Zn²⁺ (50–100 folds). These results showed that ZFL is more sensitive to metal exposure and thus it is a better endpoint to study metal contamination. Furthermore, MT gene expression was studied in metal exposed (Cu, Zn, Cd) mangrove killifish. Gene expression analysis showed that Cd is strongest inducer of MT gene (Rhee et al. 2009). The contrasting results obtained in above mentioned studies suggest that induction of MT gene in response to metals is influenced by tissue, sex, and species of fish.

In order to understand the effect and mode of action (MoA) of uranium (U), juvenile Atlantic salmon were exposed 0.25, 0.5 and 1.0 mg/L waterborne depleted uranyl acetate (Song et al. 2012). The hepatic transcriptional response showed induction of genes involved several major toxicological pathways including the anti-oxidant defense, DNA damage and repair, apoptosis and protein degradation. Although no significant adverse effects were observed after relatively short

exposure (48 h), results suggested that long term exposure to U may cause adverse effects such as DNA damage in aquatic organisms.

The transcriptional response to mercury chloride (HgCl_2) exposure in zebrafish larvae revealed that a concentration between 20 and 30 $\mu\text{g/L}$ in aquatic environment may cause oxidative stress by modulating activity and expression of P2X7 receptor (Cruz et al. 2013). Cu is known to induce apoptotic cell death in fish species. To find out the apoptotic pathways triggered by Cu, zebrafish were exposed to 12.5 and 100 $\mu\text{g/L}$ of Cu during 6, 12, 24 and 48 h. Based on the mRNA expression analysis, authors proposed that Cu initiate apoptosis via intrinsic pathway (caspase-9), through p53 activation; then by the extrinsic pathway (caspase-8) and finally by the caspase-independent pathway (AIF) (Luzio et al. 2013). In another study, Jing et al. (2013) evaluated the potential and applicability of heat shock protein (HSP70) gene to monitor heavy metal stress (Cu, Cd) in white cloud mountain minnow (*Tanichthys albonubes*). The results indicated dose-dependent expression of HSP70 mRNA. However, gene expressions did not show consistent changes between transcription and translation levels. Therefore, both transcriptional and translational variation mechanisms should be taken into account while analyzing metal stress response pathways. In a more recent study, Arini et al. (2015) demonstrated that genetic response to Cd and Zn contamination is tissue- and metal-dependent. Based on the results of the study, authors suggested that in case of binary contamination (Zn and Cd), Zn plays a protective role and prevents against oxidative stress while Cd induces oxidative stress and generate detoxification mechanisms.

To investigate the toxic effects of manufactured nanomaterials (MNs) in aquatic environment, Pham et al. (2012) studied gene expression pattern of eight stress related genes (*MT*, *GST*, *p53*, *HSP70*, *TF*, *Orla C3-I*, *Chg-L*, and *vtg-I*) after exposing Medaka to silver nitrate or a silver nanoparticle (Ag-NP). Significant induction of MT and GST (Glutathione S-transferase) mRNA suggest that the Ag-NPs cause oxidative and inflammatory stress in fish. Similarly, increase in expression of CAT, GST, and SOD (Superoxide dismutase) mRNA were observed when juvenile tilapia were exposed to titanium-dioxide NPs (TiO_2 -NPs) (Varela-Valencia et al. 2014). However, exposure to rutile phase TiO_2 -NPs generated a faster and stronger response in expression of CAT and GST gene than anatase. This suggests that rutile causes the production of higher levels of ROS than anatase.

5.3 Pesticides and Herbicides

The intensive use of pesticides and herbicides in modern agricultural system has resulted in serious environmental problems. Researchers have demonstrated that these agricultural chemicals adversely affect the immune system, early developmental stages and sexual differentiation in fish species (Li et al. 2013a; Zhang et al. 2013; Glisic et al. 2016; Wang et al. 2015a). Gene expression analysis using RT-qPCR has shown that exposure of fish to chlorpyrifos (CPF) and atrazine (ATR) causes down-regulation of acetylcholinesterase (AChE)

mRNA, an enzyme that helps in regulation of cholinergic nervous transmission (Xing et al. 2010). Thus, the alteration in expression of AChE mRNA could be used to reveal toxic mechanisms related to cholinergic signaling. In another study, CPF exposure to common carp resulted in increased IgM and complement C3 expression, indicating that CPF causes immunotoxicity to common carp (Li et al. 2013a). Transcriptional profiles of zebrafish embryos (24 h post-fertilization) after an acute exposure (48 and 72 h) to ATR resulted in significant changes in *cyp1a*, *cyp3a65* genes (Glisic et al. 2016). However, ATR failed to perturb transcription of GST isoforms and antioxidant enzymes (AOE) genes. This suggests that GST and AOE genes are not reliable markers of acute ATR exposure.

Contamination of water by organochlorine pesticides and brominated and organophosphorus flame retardants also causes toxicity in fish species. For example, induction of VTG mRNA levels in response to these chemicals suggests that they have estrogenic activity and can impair fish reproduction (Chow et al. 2011; Wang et al. 2015a). Monocrotophos (MCP), an organophosphorus pesticide has also been found to have significant estrogenic properties. MCP exposure in zebrafish stimulated *foxl2* and suppressed *dmrt1* transcription factors (Zhang et al. 2013). Differential expression of these genes indirectly elevated the gonadal (*cyp19a1a*) and brain (*cyp19a1b*) aromatase gene expression. Gonadal aromatase promotes the phenotypic feminization while brain aromatase, which is thought to be involved in reproductive physiology via the brain-pituitary-gonadal axis, causes indirect impact on sexual differentiation. MCP exposure also induces E₂ levels and promotes steroidogenesis by inducing transcription of CYP11A1, CYP17, and CYP19A genes (Wang et al. 2015b). Flame retardants (brominated and organophosphorus) have been identified to cause cardiac toxicity in fish (Wu et al. 2013; Du et al. 2015). For example, hexabromocyclododecane (HBCD) exposure of zebrafish embryos resulted in elevated expression of the *Tbx5* and *Nkx2.5* genes, which are mainly responsible for alteration of heart rate (Wu et al. 2013). Bisphenol-A (BPA), a derivative of bisphenol was found to be toxic in early development of fish. The up-regulation of *sp4* in response to BPA suggests its role in altering brain development (Lam et al. 2011).

5.4 Miscellaneous

Dioxins such as 2, 3, 7, 8-tetrachlorodibenzo-*p*-dioxins (TCDDs) are of great concern as environmental contaminants. TCDD cause a wide range of adverse effects in vertebrates. In general, fishes are highly sensitive to TCDD exposure, especially during early developmental stages (Peterson et al. 1993). RT-qPCR analysis in seabream (*Pagrus major*) embryos revealed that TCDD-induced early stage toxicity is caused by increased expression of *rsAHR2* and *CYP1A* genes (Yamauchi et al. 2006). Additionally, TCDD exposure causes modest changes in

expression of microRNAs (*miR-451*, *23a*, *23b*, *24* and *27e*) that are critical for hematopoiesis and cardiovascular development (Jenny et al. 2012). Furthermore, TCDD causes cardiac toxicity in fish embryos by suppressing transcription of a battery of cell cycle related genes (Chen 2015). RT-qPCR analysis has also been used in determining the toxic effects of PCBs in fishes by measuring the CYP1A mRNA transcript abundance (Dixon et al. 2002; Zanette et al. 2009).

Gene expression profiling in response to pharmaceuticals and their metabolites and compounds used in personal care products have shown that these contaminants cause adverse effects on aquatic organisms through the cellular toxicity, *p53* related genotoxicity and estrogenic effects (Hong et al. 2007; Yamauchi et al. 2008). In aquatic environment fish are also exposed to toxins such as microcystin (MC) and cylindrospermopsin (CYN) released by cyanobacterial blooms. Gene expression analysis have shown that MC disrupts thyroid hormone homeostasis in zebrafish (Yan et al. 2012). In another study, Li et al. (2013b) suggested that metabolism and detoxification of MCs in zebrafish is done by increased expression of CYP3A65 and its receptor PXR (pregnane X receptor) gene. Authors also found that expression of CYP3A65 and PXR gene is regulated by miRNA (*dre-miR-27b*). CYN exposure has been shown to alter the activity and gene expression of some of the glutathione related enzymes in tilapias (Gutiérrez-Praena et al. 2013). The elevated mRNA levels of GST in liver and kidney of tilapia indicates that CYN cause hepatocyte and renal injury to fish.

Fan et al. (2010) characterized ten nervous system genes (*syn2a*, *mbp*, *gap43*, *elavl3*, *nkx2.2a*, *ngn1*, *α 1-tubulin*, *nestin*, *shha*, *gfap*) to study the effects of ethanol in developmental neurotoxicity. The expression profiles of these genes were altered after exposure to sublethal doses of ethanol, a known developmental neurotoxicant, to zebrafish embryos/larvae. Presence of some of the affected genes (*α 1-tubulin*, *ngn1*, *mbp*, and *gfap*) exclusively in the nervous system suggest that gene expression profile to these genes can be used as an endpoint for efficient screening for developmental neurotoxicity using zebrafish. Yuan et al. (2014) evaluated the toxic effects of benzo[a]pyrene (BaP) on developmental stage of Chinese rare minnows. Simultaneous up-regulation of the *cyp* (*cyp1a*, *1b1*, and *1c1*) and ATP-binding cassette (ABC) transporter (*abcc1*, *abcc2*, and *abcg2*) gene transcripts suggests that these genes work in a coordinated manner to detoxify the effects of BaP in early developmental stages of fish.

To study the oxidative stress response of nitrite, gene expression profile of antioxidant genes were evaluated in Wuchang bream (*Megalobrama amblycephala*). The acute nitrite exposure in juvenile fish resulted in the formation of excess of reactive oxygen species (ROS) and reduced expression of antioxidant genes (*MaCAT*, *MaGPx1*, *MaCu/Zn-SOD*). The reverse correlation between ROS and antioxidant genes suggest that imbalance between ROS and antioxidant defenses is the mechanism underlying nitrite toxicity in Wuchang bream (Sun et al. 2014). In another study, toxic effect of perfluorononanoic acid (PFNA) was examined in zebrafish larvae. Gene expression analysis showed an up-regulation of genes involved in oxidative stress and apoptotic pathways (Yang et al. 2014). Furthermore, to evaluate the adverse effects and intrinsic toxicological properties

of tributyltin (TBT), Tian et al. (2015) studied the gene expression profile of aromatase gene in male guppies (*Poecilia reticulata*). RT-qPCR analysis revealed that TBT exposure suppresses the expression of two isoforms of aromatase. Suppression of aromatase causes the disturbance in reproductive behavior of fish by endocrine-disrupting action.

6 Use of Microarrays in Fish Toxicology

Microarrays (also commonly known as DNA arrays, DNA chip or biochip) are a high throughput technology that are used to measure expression levels of thousands or even tens of thousands of genes simultaneously. This allows researchers to study the effects of toxicants on the expression of virtually all known genes within a single microarray (Ramsay 1998). However, one of the major limitations for the use of microarrays in fish toxicology is the lack of sequence information for non-model species. Nevertheless, ESTs and custom arrays based on genes expressed in response to specific contaminants have been developed and used with microarray to provide the genome wide expression data. Although use of microarray technology in molecular biology research started in mid-1990s (Ramsay 1998), the first use of microarray in fish was done by Gracey and co-workers in 2001 (Gracey et al. 2001). Since then microarrays have been used in large number of fish species to evaluate the gene expression responses to toxicants and to identify genes and molecular pathways involved in fish responses to toxicants.

6.1 Metals, Metallic Nanoparticles and Organometals

In European flounder (*Platichthys flesus*) microarrays were applied to investigate the effect of acute exposure to Cd (Sheader et al. 2006). The mRNA expression analysis detected up-regulation of 27 transcripts (including Cu/Zn SOD, thioredoxin, a peroxiredoxin, and a GST) and down-regulation of CYP1A. Alteration in these genes indicates that Cd exposure cause oxidative stress in flounder. In another study, fibroblast cell line (SAF1) from gilthead bream was exposed to sub-toxic levels of Cu, Zn, and Cd (Minghetti et al. 2011). Microarrays analysis showed an increased mRNA levels of MT and GST genes in response to Cu, Zn, and Cd. However, only ATP7A mRNA levels were increased in Cu treated SAF1. Findings of this study suggest that expression of copper transporting ATPase, ATP7A, may be regulated at the transcriptional level directly by Cu.

The toxic effect and mode of action (MoA) of uranium (U) was studied in the brain of the zebrafish by exposing the fish to 15 and 100 µg/L of waterborne U for 3 and 10 days (Lerebours et al. 2010). A total of 56 transcripts expressed differentially and were categorized into eight functional classes with the most significantly affected being the olfactory system. Results of this study suggest that genes in the

olfactory region may be used as a sensitive transcriptional biomarkers of U water-borne exposure. Song et al. (2014) expanded their previous study (Song et al. 2012) of U toxicity in Atlantic salmon using microarrays in combination with RT-qPCR. Hepatic transcriptional responses showed that U exposure affects various toxicity pathways involving mitochondrial function, oxidative stress, nuclear receptor signaling, and organ damage. In another study, genome-wide gene expression responses to metal poisoning and mechanistic insights into metal toxicity were evaluated in whole adult male zebrafish following exposure to three metals (nickel, cobalt, chromium) (Baer et al. 2014). Based on the expression data authors identified changes in gene expression of group of genes involved in adaptive responses, cell cycle regulation, apoptosis and metabolic depression.

Research has shown that aggregated nanoparticles and release of dissolved metal ions produce different toxicity effects in aquatic organisms. For example, global gene expression analysis in gills of zebrafish following exposure to nanocopper, nanosilver, and nanotitania produced a distinct gene expression profile for each of nanometal or soluble metal, suggesting that each exposure is producing biological response by a different mechanism (Griffitt et al. 2009). Jovanovic et al. (2011) examined the acute toxicity of engineered nanoparticles (titanium dioxide/TiO₂ and hydroxylated fullerenes/C₆₀ (OH)₂₄) in zebrafish embryos. The nanoparticle exposure caused changes in genes related to circadian rhythm, cell kinase activity, intracellular trafficking, and immune response. However, expression patterns were similar for down-regulated genes but different for upregulated genes. Finding of this study suggest that different nanomaterials may have similar suppression but different up-regulation patterns of transcriptome changes. Furthermore, sublethal effects of nano-Ag and dissolved Ag were examined on rainbow trout (Gagné et al. 2012). Gene expression analysis revealed that genes involved in inflammation are sensitive to nano-Ag exposure while oxidative stress and protein stability related genes are sensitive to dissolved Ag. This shows that the toxicity of Ag differ depending on the presence of Ag nanoparticles and aggregates.

Methylmercury (MeHg), a known neurotoxicant, is widely distributed contaminant polluting many aquatic environments. To study its toxic effects, fathead minnows (*Pimephales promelas*) were fed with food containing MeHg (Klaper et al. 2006). Gene expression analysis revealed a marked up-regulation in VTG mRNA in males and a significant decline in VTG gene expression in female fish. This differential expression indicates that MeHg cause endocrine disruption in fish. In an expansion to this study, Klaper et al. (2008) used 15,000 gene microarray to investigate the impact of acute and chronic MeHg exposures in male gonad and liver tissues of fathead minnows. Gene expression analysis showed significant differences in differentially expressed genes in each of the tissues in response to acute and chronic treatment. In general, genes associated with stress, adaptation, and the immune system were altered. In a more comprehensive study, Yadetie et al. (2013) prepared Atlantic cod 135 k oligonucleotide arrays to study the mechanisms of toxicity and range of molecular targets of MeHg. In addition to the known effects of MeHg such as oxidative stress, gene transcripts coding for

enzymes involved in metabolism of amino acids, fatty acids and glucose were up-regulated. These results suggest that exposure to MeHg may cause disruption of nutrient metabolism in fish species.

6.2 Endocrine Disrupting Chemicals

Usefulness of microarrays has been also demonstrated in studying the effects of EDCs in fish species. For example, Larkin et al. (2003) used estrogen-responsive gene array of sheepshead minnow (SHM) to study the effects of E₂, EE₂, DES, *p*-nonylphenol (PNP), methoxychlor (MXC), and endosulfan (ES). Gene expression analysis showed that E₂, EE₂, DES, and MXC have similar genetic signatures. However, the number of genes examined in this study were limited (30 genes). Knoebel et al. (2006) expanded the existing gene array for SHM and used it to examine temporal changes in gene expression for male SHM exposed to 100 ng E₂/L for five time points between 0 and 48 h. Authors identified five patterns of temporal induction in genes involved in oocyte development. However, this difference in temporal expression was lost at higher concentration (500 ng E₂/L) probably due to feedback effects in brain. Findings of this study suggest that exposure to high levels of environmental contaminants may affect the normal ordered expression of genes required for reproduction.

In another study, Moens et al. (2007a) developed a cDNA microarray of 398 different gene fragments for the evaluation of endocrine disruption in common carp (*Cyprinus carpio*). Gene expression profiles of fish treated with E₂ resulted in a total of 35 differentially expressed genes, whereas cortisol exposure affected only three genes. These results indicate the discriminating power of the developed array and its usefulness to describe the toxicological mode of action of EDCs. In order to identify novel genomic responses to EE₂ exposure, Martyniuk et al. (2007) utilized microarrays in conjunction with RT-qPCR. Gene expression analysis in liver and telencephalon of male zebrafish identified tissue specific gene responses. For example, genes involved in protein metabolism, carbohydrate metabolism were down-regulated in the liver but were induced in the telencephalon.

Furthermore, a cDNA microarray consisting of 9692 clones was developed for threespined stickleback and used to assess gene expression responses to the dibenzanthracene (DbA) and E₂ as single and binary treatments (Geoghegan et al. 2008). Induction of CYP1A and ER α mRNA was observed in both male and female individuals in response to DbA and E₂ respectively. However, exposure to E₂ and binary mixture showed a different expression pattern in male and female sticklebacks for VTG, zona radiata and chorion protein mRNA. VTG, zona radiata and chorion protein mRNA was induced in male fish exposed to E₂ and a binary mixture while females showed induction only with the binary mixture. The disparity in expression pattern of genes in male and female in response to E₂ is most likely because of high endogenous levels of E₂ in female fish (Geoghegan et al. 2008).

Wei et al. (2008) used a custom cDNA microarray containing 1773 unique genes to study the effects of perfluorooctanoic acid (PFOA) in male and female rare minnows. Gene expression analysis revealed the inhibition of thyroid hormone biosynthesis genes and induction of estrogen-responsive genes strongly suggesting that PFOA is an EDC.

Wang et al. (2011) investigated the effects of trilostane on the transcriptional regulatory dynamics of zebrafish. Differential expression analysis revealed that trilostane-induced genes are broadly involved in cell proliferation, differentiation, migration, and apoptosis. Additionally, authors concluded that trilostane-induced effects on fish endocrine functions are not confined to the HPG-axis alone. Further, to strengthen the toxicogenomic study, Pham et al. (2011) constructed cDNA microarrays from Japanese medaka to screen biomarkers for E₂, NP, and 2-chlorophenol (2CP). To study the toxicological potency of DES in fish, Adedeji et al. (2012) exposed sexually mature fathead minnows of both sexes to 1, 10, or 100 ng of DES/L of water in a flow-through system. Evaluation of hepatic transcript profile revealed induced mRNA level of VTG and ER α in both sexes at DES concentrations ≥ 10 ng/L suggesting concentration dependent change in gene expression.

6.3 Miscellaneous

Exposure of fish to TCDD not only causes alteration in gene expression but multiple injuries in the heart, liver, kidney, intestine, and ovary (Volz et al. 2006; Liu et al. 2014). Microarray analysis of 175-gene array from medaka revealed that alteration in hepatic gene expression and histological changes are strongly dependent on dose and time (Volz et al. 2006). Liu et al. (2014) showed that altered genes in response to TCDD were involved in pathways associated with cardiac necrosis/cell death, cardiac fibrosis, renal necrosis/cell death, and liver necrosis/cell death. In another study, Hofsteen et al. (2013) revealed that TCDD exposure inhibit the regeneration of heart muscle and also alters the pattern of gene expression induced by wounding at the ventricle in zebrafish. Based on the results authors speculated that TCDD exposure inhibits a process that is important for progenitor cells needed for both early heart development and regeneration.

Utility of microarray in longjaw mudsucker (*Gillichthys mirabilis*) revealed that exposure of fish to hypoxic environment first suppresses the energy-requiring processes like protein synthesis and locomotion in skeletal muscle and then induces genes in liver needed for anaerobic ATP production and for gluconeogenesis (Gracey et al. 2001). Martinovic et al. (2009) showed that hypoxia caused alteration in reproduction of zebrafish by reorganization of lipid transport and mechanisms involved in hydroxyl steroid dehydrogenase alterations and down-regulation of contractile elements.

In order to assess the potential of microarray technique for environmental monitoring, Williams et al. (2003) sampled European flounder from the polluted

Tyne estuary. The inductions of CYP1A, UDP-glucuronosyltransferase (UDPGT) and aldehyde dehydrogenase mRNA in the liver of flounder suggested that Tyne estuary is contaminated with polycyclic aromatic hydrocarbons (PAHs). Moens et al. (2007b) conducted a chronic toxicity test by exposing common carp to effluent for 21 days under flow-through conditions. Microarray analysis revealed that effluent treatment affected molecular pathways associated with the energy balance of the fish, including changes in carbohydrate and lipid metabolism, as well as digestive enzyme activity. Furthermore, Lie et al. (2009) investigated the effects of contaminants on gene expression in two natural populations of Atlantic cod from western Norway. The up-regulation of CYP1A, heme oxygenase, ferritin, and metallothionein gene suggests the heavy metal pollution in sampling sites.

Other studies using microarray technology in fish include the examination of the toxic effects on development and survival of zebrafish embryos in response to a wider range of ecotoxins (2, 4-dichlorophenol, 3, 4-dichloroaniline, pentachlorophenol, and cadmium chloride) (Sawle et al. 2010), dithiocarbamates (DTCs) (van Boxtel et al. 2010), flusilazole (Hermesen et al. 2012), dimethylformamide (DMF), dimethylsulfoxide (DMSO) (Turner et al. 2012) and PAHs (Goodale et al. 2013; Jayasundara et al. 2015). Tilton et al. (2005) used rainbow trout as a research model for study of a fungicide metabolite, aflatoxin B1 (AFB1) induced hepatocarcinogenesis. The altered gene expressions were found to be similar to those observed in humans and other mammalian models. Thus, rainbow trout can be used as model to study the carcinogenesis in humans and other mammals. Martyniuk et al. (2010) examined the toxic effects of sub-chronic dietary exposures to 2.95 mg dieldrin/kg feed in the neuroendocrine brain of largemouth bass. Microarrays, proteomics, and pathway analysis revealed that sub-chronic exposure to dieldrin alters the abundance of mRNAs and proteins in the hypothalamus that are associated with cell metabolism, cell stability and integrity, stress, and DNA repair. The application of microarrays has also been used for the characterization of gene expression patterns associated with exposure to ethynylestradiol, 2,2,4,4-tetrabromodiphenyl ether, diquat, chromium VI, BaP (Hook et al. 2006), 2, 4-dinitrotoluene (2, 4-DNT) (Wintz et al. 2006), perfluorooctane sulfonate (PFOS) (Hagenaars et al. 2008), domoic acid (Lefebvre et al. 2009), polyfluorinated and perfluorinated compounds (PFCs) (Wei et al. 2009), cyclotrimethylenetrinitramine (RDX) (Gust et al. 2011) and brominated flame retardant mixture (Williams et al. 2013).

7 Use of NGS in Fish Toxicology

The advent of next-generation sequencing (NGS) technologies has driven a massive acceleration in research and development. The decreasing per base cost of raw sequence and increasing throughput has made it affordable even for non-model species. Although microarrays are still the primary technology for transcriptome analysis, in recent years RNA-seq using NGS has been widely used to study the

effects of chemicals and pollutants on fish species. This is largely because RNA-seq does not require prior sequence knowledge and is able to detect even low expressing genes. This increased sensitivity allows for the identification of novel biomarkers. In addition, RNA-seq experiments require less RNA input compared to microarrays (Wang et al. 2009).

In one of the first toxicological studies in fish using NGS, Pierron et al. (2011) examined the effects of Cd and Cu exposure on yellow perch (*Perca flavescens*). Transcriptome analysis in treated fish revealed a decrease in mRNA levels of genes associated with protein biosynthesis, immune system, and lipid and energy metabolism. Based on the results the authors suggested that down-regulation of these genes could be due to an impairment of bile acid metabolism by Cd and silencing of genes by epigenetic modification of histones and DNA. In a subsequent year, Huang et al. (2012) studied the transcriptional response of marine medaka (*Oryzias melastigma*) embryos after PFOS exposure. Differential expression of genes related to neurobehavioral defects, mitochondrial dysfunction, and the metabolism of proteins and fats provided the useful data for identifying affected pathways and possible mechanisms of PFOS in marine fish. In another study, the toxicological effects of TCDD, a well-known teratogen, on development of zebrafish was examined using RNA-seq supplemented with microarrays and RT-qPCR (Jenny et al. 2012). Expression profiles of miRNA suggested that TCDD exposure causes abnormal phenotypes associated with hematopoiesis and cardiovascular development. In a similar study, RNA-seq was used to identify miRNAs as potential biomarker for monitoring stress responses towards acidic aluminum-rich water in Atlantic salmon (Kure et al. 2013).

Garcia de la serrana et al. (2012) successfully applied the RNA-seq technology to study the effects of deepwater horizon (DH) oil release on gulf killifish (*Fundulus grandis*). Altered expression of a large set of genes (1070 up and 1251 down-regulated) indicates that exposure to DH oil mediates a broad and complex genomic response including activation of AHR and related pathways associated with xenobiotic hydrocarbons. RNA-seq has also been used to investigate sublethal effects of oil sands process-affected water (OSPW) in fathead minnows (Wiseman et al. 2013). Based on abundances of transcripts in livers of fathead minnows exposed to untreated OSPW, authors proposed that OSPW cause toxicity through a mechanism involving reactive oxygen species (ROS) and apoptosis.

In order to find out the transcriptional responses to Hg, liver transcriptome of gadoid fish (*Brosme brosme*) was sequenced (Olsvik et al. 2013). Findings of this study suggested that Hg exposure affect the lipid metabolism and beta-oxidation in liver. Webster et al. (2013) used the RNA-seq to investigate the molecular mechanisms of metal (water-borne mixture of metals, including copper and zinc) tolerance in brown trout (*Salmo trutta*). Based on functional analysis, authors suggested that metal- and ion-homeostasis pathways could be the most important mechanisms contributing to the metal tolerance in brown trout. Du et al. (2014) extended the RNA-seq technology to study the gene expression in response to loading stress in Japanese grenadier anchovy (*Coilia nasus*). Based on pathway enrichment analysis of loading stress-responsive unigenes, authors concluded that loading stress induces

liver injury through the mitochondrial apoptosis pathway. In a most recent study, Bahamonde et al. (2015) used RNA-seq in conjunction with microarray to study the molecular signaling cascades associated with intersex in rainbow darter (*Etheostoma caeruleum*) exposed to municipal wastewater. Differential expression analysis of genes involved in sex differentiation (*sox9*, *foxl2*, and *dmrt1*) and reproduction (*esr1*, *esrb*, *ar*, *vtg*, *cyp19a1*, and *cyp11a*) showed an intermediate expression level in intersex males when compared to phenotypic males and females. Thus, RNA-seq can be used to discriminate pollutant-exposed males without intersex from those males with intersex.

8 Toxicogenomics Databases and Bioinformatics

A rapid progress in toxicogenomics approaches in the last decade has generated a wealth of gene expression data. There was a growing desire in scientific community to make these data sets publicly available once research findings have been published. Efforts were made in this direction and several public databases (ArrayExpress, Gene Expression Omnibus (GEO), Comparative Toxicogenomics Database (CTD), Center for Information Biology gene Expression database (CIBEX), Sequence Read Archive (SRA)) for functional genomics were established. The ArrayExpress (<http://www.ebi.ac.uk/arrayexpress>) (Brazma et al. 2003) and GEO (<http://www.ncbi.nlm.nih.gov/geo/>) (Edgar et al. 2002) are repositories for microarray and high-throughput sequencing based genomics experiments. The CTD (<http://ctdbase.org/>) (Mattingly et al. 2003) was developed to formalize, harmonize and centralize the information on numerous genes and proteins responding to environmental toxic agents across diverse species.

To facilitate and promote the sharing of high quality and well-annotated microarray data within the scientific community, members of Microarray Gene Expression Data (MGED) Society (<http://www.mged.org>) laid down and published standardizations for microarray experiments and data description called MIAME (Minimum Information About a Microarray Experiment). In Japan, along with the MGED activity, Ikeo et al. (2003) developed a gene expression database called CIBEX, with a data retrieval system in compliance with MIAME. In 2009, database for next-generation sequencing data, SRA, was established as a part of the International Nucleotide Sequence Database Collaboration (INSDC) (Shumway et al. 2010). Similar to MIAME guidelines for microarray data, guidelines for high-throughput sequencing data, Minimum Information about Sequencing Experiment (MINSEQE; <http://www.fged.org/projects/minseqe/>) were proposed.

Some other toxicogenomics resources such as External RNA Controls Consortium (ERCC) was developed to verify technical performance and to interpret quality of gene expression data. To further strengthen the toxicogenomics resources, MicroArray Quality Control project (MAQC) was initiated which is working to improve application of the microarray and next-generation sequencing technologies by providing large reference datasets and developing guidelines for

toxicogenomics data analysis. In another project called Percellome Toxicogenomics Project (PTR), a systematic method to normalize the mRNA expression values from DNA microarrays and RT-qPCR in a “per one cell” basis was developed. This method helps to monitor the time- and dose-dependent alteration of gene expression in response to various chemicals.

In toxicogenomics, bioinformatics approaches are used to identify chemical classes, toxic effects of a new compounds and pathways affected by a chemical stressors. The clustering method is used to compare gene expression profile of two samples or relative intensity of a single sample in response to a particular chemical. This approach groups the genes within or between samples to find the chemicals with similar toxic mechanisms (Waring et al. 2001). The connectivity mapping approach is used to find the toxic effects of a new chemical by comparing the gene expression profile to the compounds with similar gene expression profile whose toxicity is already known. Bioinformatics tools are also used to discern mechanisms of action of toxic responses. In mechanistic analysis, gene annotation databases such as Gene Ontology (GO, <http://geneontology.org/page/go-database>), pathway databases such as Kyoto Encyclopedia of Genes and Genomes (KEGG, <http://www.genome.jp/kegg/>) and WikiPathways (<http://www.wikipathways.org>) are commonly used. However, in recent years CTD has evolved as premier toxicology resource for mechanistic analysis of toxicogenomics data (Davis et al. 2015). Although the quantity of toxicogenomics data are increasing at an exponential rate, there is a lack of capacity in computational and bioinformatics tools. Therefore, new and efficient data-mining programs and comparative genomic tools are required to explore and interpret data in a time frame compatible with the rate at which they are generated.

9 Summary and Conclusion

Gene expression analysis is one of the most widely used techniques in detecting and elucidating molecular responses to toxicants. Over the last 10 years by the introduction and widespread adoption of toxicogenomic techniques in fish toxicology has made a paradigm shift from studying expression patterns of single or few molecular endpoints to genome wide expression response. Nevertheless, toxic effects of chemicals and pollutants have been investigated on only a few fish species (Tables 1, 2, 3, 4, 5, and 6). This is largely because of limited genomic information available for non-model fish species. Even for fish species in which the genome has been sequenced, poor annotation of the genome often poses difficulty in interpretation of the biological significance of differentially expressed gene sets. In recent years, however, great efforts have been made on sequencing the genomes of economically important aquaculture species. In this effort 12 fish species have been fully sequenced and assembled (<http://www.ensembl.org>). In addition, 22 fish species have been annotated by the NCBI Eukaryotic Genome Annotation Pipeline (<http://www.ncbi.nlm.nih.gov>). Furthermore, recent advances in NGS technologies

Table 1 Examples of studies using Northern blotting in fish toxicology

Toxicants/compounds	Species	Tissue	Target gene	Major findings	Reference
17 α -methyltestosterone (17 α MT)	<i>Oreochromis niloticus</i>	Liver	Vitellogenin (VTG)	Inhibition of VTG gene expression	Lazier et al. (1996)
Effluents from pulp, paper, and paperboard mills	<i>Coregonus lavaretus</i>	Liver	VTG	Increased vitellogenin gene expression	Mellanen et al. (1999)
17 β -estradiol (E ₂)	<i>Sparus aurata</i>	Liver	VTG, transthyretin (TTR)	Induction of VTG mRNA level and reduction of TTR mRNA levels by E ₂ treatment	Funkenstein et al. (2000)
β -naphthoflavone (BNF)	<i>Pleuronectes platessa</i>	Gill, heart, liver	Cytochrome P450 1A (CYP1A)	Up-regulation of CYP1A after BNF treatment, with greatest response in gill followed by heart and liver	Leaver and George (2000)
E ₂ , nonylphenol (NP)	<i>Oncorhynchus mykiss</i>	Liver	Zona radiate (ZR), VTG	Up-regulation of ZR and VTG mRNA	Arukwe et al. (2002)
E ₂ , ethinylestradiol (EE ₂)	<i>Cyprinodon variegatus</i>	Liver	VTG	Induction of VTG mRNA in liver	Bowman et al. (2000)
E ₂	<i>Micropterus salmoides</i>	Liver	VTG, estrogen receptor (ER)	Hepatic ER and VTG mRNA are coordinately regulated following acute E ₂ exposure (2 mg/kg)	Bowman et al. (2002)
Polybrominated diphenyl ethers (PBDEs)	<i>Salmo salar</i>	Blood, liver, fillet, brain	CYP1A, VTG, ZR	Up-regulation of CYP1A, VTG and ZR gene	Boon et al. (2002)
2,3,7,8-Tetrachlorodibenzo- <i>p</i> -dioxin (TCDD)	<i>Danio rerio</i>	Embryo	CYP1A	Hypoxia decreased TCDD induction of CYP1A mRNA, and decreased the potency of TCDD in causing edema	Prasch et al. (2004)
4-Nonylphenol (4-NP)	<i>Sebastes schlegelii</i>	Liver	VTG	4-NP disrupt the reproductive system of immature rockfish by suppressing the expression of vitellogenesis	Jung et al. (2006)

Table 2 Examples of studies using DDRT-PCR in fish toxicology

Toxicants/ compounds	Species	Tissue	Target gene	Major findings	Reference
E ₂ , EE ₂ , diethyl- stilbestrol (DES)	<i>Cyprinodon variegatus</i>	Liver	48 genes	E ₂ , EE ₂ and DES act through the same pathway when administered through the water at low concentrations	Denslow et al. (2001a, b)
Cadmium (Cd)	<i>Chionodraco hamatus</i>	Liver	Seven genes	Up-regulation of gene coding for heat-shock protein HSP70 and down- regulation of transferrin gene	Carginale et al. (2002)
Effluents from pulp and paper mills	<i>Micropterus salmoides</i>	Liver	CYP1A	Decrease in repro- ductive success in females	Denslow et al. (2004)
Anthracene	<i>Fundulus heteroclitus</i>	Liver	26 genes	CYP2N2 and two expressed sequence tags (15C1 and 18C2) act as good indica- tor of anthracene exposure	Peterson and Bain (2004)
Lindane	<i>Oreochromis niloticus</i>	Liver	50 genes	Lindane triggers the expression of genes involved in immunological and stress responses	Colli-Dula et al. (2009)
Zearalenone (ZEA)	<i>Oncorhynchus mykiss</i>	Liver, ovary	59 genes	ZEA interferes with blood coagu- lation and different cellular components	Woźny et al. (2012)

offers opportunity of using high-quality genomic and transcriptomic data in non-model fish species.

It is evident that most of the researchers have used single compounds or simple mixtures to explore the relationship of gene expression and contaminant exposure in laboratory conditions. However, in natural environments, pollutants most often are present as complex mixtures rather than discrete chemicals. Therefore, future research should aim to understand the synergetic effects of multiple environmental stressors under field conditions. Furthermore, since chemical stressors invoke both toxic and adaptive processes, genetic variation of individual fish should also be taken into account for interpretation of alteration in gene expression profile

Table 3 Examples of studies using suppression subtractive hybridization (SSH) in fish toxicology

Toxicants/compounds	Species	Tissue	Target gene	Major findings	Reference
EE ₂	<i>Pleuronectes platessa</i>	Liver	75 genes	Selective down-regulation of egg proteins, potentially diminishing the quality of eggs	Brown et al. (2004)
2,3,7,8 Tetrachlorodibenzo- <i>p</i> -dioxin (TCDD)	<i>Oryzias latipes</i>	Brain, liver, and testis	42 genes	Up-regulation of CYP1A mRNA in TCDD-exposed brain and liver	Volz et al. (2005)
Polychlorinated biphenyls (PCBs)	<i>Sebastes marmoratus</i>	Brain	108 genes	PCBs cause differential expression of genes with function of neurotransmitter release, neuroendocrine, mitosis and cell proliferation, energy-related metabolism, general metabolism, signal protein, hemopoiesis system, immune system, and structure	Li et al. (2008)
Domestic effluents	<i>Dicentrarchus labrax</i>	Liver	CYP1A	Altered expression in genes involved in energy metabolism, immune system activity and antioxidant response	Nogueira et al. (2010)
Cadmium (Cd)	<i>Cyprinodon variegatus</i>	Larvae	700 genes	Cd responsive genes are broadly involved in protective and repair functions as well as cellular and metabolic processes	Dangre et al. (2010)
Microcystin-LR (MCLR)	<i>Hypophthalmichthys molitrix</i>	Liver	75 genes	Differential expression of genes with functions of immune response, transport and cell metabolism	Qu et al. (2011)
Tetrodotoxin (TTX)	<i>Takifugu rubripes</i>	Liver	1048 genes	High level of up-regulation in hepcidin precursor gene	Matsumoto et al. (2011)

Table 4 Examples of studies using reverse transcription quantitative polymerase chain reaction (RT-qPCR) in fish toxicology

Toxicants/ compounds	Species	Tissue	Target gene	Major findings	Reference
E ₂ , α -zearalenol	<i>Oncorhynchus mykiss</i>	Liver	Zona Radiata (Zr), VTG	Induction of Zr and VTG	Celius et al. (2000)
Cu ²⁺ , Cd ²⁺ , Hg ²⁺ , Ni ²⁺ , Pb ²⁺ , Zn ²⁺	<i>Tilapia aurea</i> , <i>T. nilotica</i>	Liver, kidney, gill	MT	Metal contamination induce hepatic and renal MT mRNA levels	Cheung et al. (2004)
Bisphenol A, nonylphenol, octylphenol	<i>Kryptolebias marmoratus</i>	Brain, eye, gonad, intestine, liver, muscle, skin	MT	Down-regulation of metallothionein gene in both male and female individuals	Rhee et al. (2009)
Atrazine, chlorpyrifos	<i>Cyprinus carpio</i>	Brain, muscle	Acetylcholinesterase (AChE)	Down-regulation of AChE	Xing et al. (2010)
Estradiol, 2-methyl-testosterone	<i>Gobiocypris rarus</i>	Eye, brain, gill and gonads	<i>foxl2</i>	Up-regulation of <i>foxl2</i> with estradiol and down regulation with 2-methyl-testosterone	Jiang et al. (2011)
β -estradiol, 17 α -valerate (EV)	<i>Oryzias latipes</i>	Liver	Estrogen receptor α (ER- α) and vitellogenin-1 (VTG-1)	EV is a reproductive toxicant and cause reproductive failure in fish species	Lei et al. (2013)
Hg	<i>Danio rerio</i>	Larvae	P2X7R	Mercury cause toxicity by modulating P2X7R in zebrafish larvae	Cruz et al. (2013)
Cylindrospermopsin (CYN)	<i>Oreochromis niloticus</i>	Liver, kidney	GST	CYN cause hepatocyte and renal injury to fish	Guiérrez-Praena et al. (2013)
Benzo[a]pyrene (BaP)	<i>Gobiocypris rarus</i>	Embryo	<i>cyp1a</i> , <i>lcl</i> , <i>gxm</i> , <i>ugt1a</i> , <i>abcb1</i> , <i>abcc1</i> , <i>abcc2</i> , <i>abcg2</i>	<i>cyp1a</i> , <i>lcl</i> and <i>abcc1</i> , <i>abcc2</i> , <i>abcg2</i> genes work in a coordinated manner to detoxify the effects of BaP in early developmental stage of fish	Yuan et al. (2014)
Nitrite	<i>Megalobrama amblycephala</i>	Liver	MaCAT, MaGPx1, MaCu/Zn-SOD	Imbalance between ROS and antioxidant defenses cause nitrite toxicity in <i>M. amblycephala</i>	Sun et al. (2014)
Tributyltin (TBT)	<i>Pocilia reticulata</i>	Brain	Aromatase	Suppression of aromatase cause the disturbance in reproductive behavior of fish by endocrine-disrupting action	Tian et al. (2015)

Table 5 Examples of studies using microarrays in fish toxicology

Toxicants/compounds	Species	Tissue	Target gene	Major findings	Reference
Hypoxia	<i>Gillichthys mirabilis</i>	Liver, brain, skeletal, and cardiac muscle	126 genes	Hypoxic environment first suppress the energy-requiring processes and then induce the genes needed for anaerobic ATP production and for gluconeogenesis	Gracey et al. (2001)
E ₂ , EE ₂ , DES, PNP, MXC, and ES	<i>Cyprinodon variegatus</i>	Liver	30 genes	E ₂ , EE ₂ , DES, and MXC have similar genetic signatures	Larkin et al. (2003)
Cadmium	<i>Platichthys flesus</i>	Liver	CYP1A, Cu/Zn SOD, thioredoxin, a peroxiredoxin and a GST	Cadmium exposure cause oxidative stress in flounder	Sheader et al. (2006)
Methylmercury (MeHg)	<i>Pimephales promelas</i>	Liver	76 genes in female and 42 in males	MeHg cause endocrine disruption in fish	Klaper et al. (2006)
Perfluorooctanoic acid (PFOA)	<i>Gobiocypris rarus</i>	Liver	124 genes in male and 171 in female	Inhibition of thyroid hormone biosynthesis genes and induction of estrogen-responsive genes	Wei et al. (2008)
Perfluorooctane sulfonate (PFOS)	<i>Cyprinus carpio</i>	Liver	1632 genes	PFOS affects the genes involved in energy metabolism, reproduction and stress response	Hagenaars et al. (2008)
PCBs, PAHs, Cu, and Cd	<i>Gadus morhua</i>	Liver, gill, and kidney	CYP1A, heme oxygenase, ferritin, and metallothionein	Microarrays can be used in diagnostic tool for environmental monitoring	Lie et al. (2009)

Diieldrin	<i>Largemouth bass</i>	Hypothalamus	274 genes	Sub-chronic exposure to diieldrin alters the genes associated with cell metabolism, cell stability and integrity, stress, and DNA repair	Martyniuk et al. (2010)
Cu, Zn, and Cd	<i>Sparus aurata</i>	Fibroblast cell line (SAF1)	MT, GST, ATP7A	Increase in mRNA levels of MT and GST genes	Minghetti et al. (2011)
Titanium dioxide/TiO ₂ and hydroxylated fullerenes/C ₆₀ (OH) ₂₄	<i>Danio rerio</i>	Embryo	2693 genes	Nanoparticle exposure alters expression pattern in genes related to circadian rhythm, cell kinase activity, intracellular trafficking and immune response	Jovanovic et al. (2011)
E ₂ , nonylphenol (NP) and 2-chlorophenol (2CP)	<i>Oryzias latipes</i>	Liver	1509 ESTs	Providing information of 1509 high-quality ESTs including 260 new EST sequences to O. latipes dbEST and GenBank of the NCBI	Pham et al. (2011)
Silver nanoparticles	<i>Oncorhynchus mykiss</i>	Liver	207 genes	Toxicity of Ag differ depending on the presence of Ag nanoparticles and aggregates	Gagné et al. (2012)
Uranium	<i>Salmo salar</i>	Liver	927 genes	Uranium exposure affects various toxicity pathways involved in mitochondrial functions, oxidative stress, nuclear receptor signaling, and organ damage	Song et al. (2014)

Table 6 Examples of studies using next-generation sequencing technology in fish toxicology

Toxicants/compounds	Species	Tissue	Platform	Major findings	Reference
Cadmium (Cd) and Copper (Cu)	<i>Perca flavescens</i>	Liver	Roche 454 GS-FLX	Chronic exposure of Cd and Cu decrease in the transcription levels of genes involved in protein biosynthesis, immune system, and lipid and energy metabolism	Pierron et al. (2011)
Perfluorooctane sulfonate (PFOS)	<i>Oryzias melastigma</i>	Embryo	Illumina HiSeq 2000	PFOS exposure cause neurobehavioral defects, mitochondrial dysfunction and the metabolism of proteins and fats	Huang et al. (2012)
2,3,7,8-Tetrachlorodibenzo- <i>p</i> -dioxin (TCDD)	<i>Danio rerio</i>	Embryo	SOLID	TCDD exposure causes modest changes in expression of miRNAs, including some (<i>miR-451</i> , <i>23a</i> , <i>23b</i> , <i>24</i> , and <i>27e</i>) that are critical for hematopoiesis and cardiovascular development	Jenny et al. (2012)
Deepwater horizon oil	<i>Fundulus grandis</i>	Liver	Illumina GAIIX	DH oil exposure cause broad and complex genomic response including the expected AHR-mediated response, CYP genes, immune response, and choriogenin family genes	Garcia de la serrana et al. (2012)
Water-borne mixture of metals, including copper and zinc	<i>Salmo trutta</i>	Gills, liver and kidney	Illumina GAIIX	Metal- and ion-homeostasis pathways are likely to be the most important mechanisms contributing to the metal tolerance in brown trout	Webster et al. (2013)
Mercury (Hg)	<i>Brosme brosme</i>	Liver	Illumina HiSeq 2000	Hg have toxic effects on lipid metabolism and oxidative stress	Olsvik et al. (2013)
Acidic aluminum-rich water	<i>Salmo salar</i>	Muscle	Illumina GAIIX	Identification of specific miRNAs as potential early markers for stress	Kure et al. (2013)
Oil sands process-affected water (OSPW)	<i>Pimephales promelas</i>	Liver	Illumina HiSeq 2000	Exposure of OSPW alters gene expression of genes involved in oxidative metabolism, oxidative stress, apoptosis, and immune function	Wiseman et al. (2013)
Loading stress	<i>Coilia nasus</i>	Liver	Illumina HiSeq 2000	Loading stress induce liver injury through the mitochondrial apoptosis pathway	Du et al. (2014)
Municipal wastewater	<i>Etheostoma caeruleum</i>	Ovary, testis	Roche 454 GS-FLX	Intermediate expression level of genes involved in sex differentiation (<i>sox9</i> , <i>foxl2</i> and <i>dmrt1</i>) and reproduction (<i>esr1</i> , <i>esrb</i> , <i>ar</i> , <i>yfg</i> , <i>cyp19a1</i> , and <i>cyp11a</i>) in intersex males compared to phenotypic males and females	Bahamonde et al. (2015)

following chemical exposure. Although gene expression analyses provide the information about immediate toxic effects of pollutants and new hypotheses about their functions, it does not, as such, give functional information. Analysis of protein concentrations in a cell can give insights regarding gene function, interacting genes, and gene pathways. Therefore, multidisciplinary approaches should be implemented in monitoring the acute and chronic adverse effects of pollutant to understand the complexity of anthropogenic impacts on fish species.

Conflict of Interest The authors declare that they have no conflict of interest.

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Review of the Ecotoxicological Properties of the Methylenedianiline Substances

T. Schupp, H. Allmendinger, B.T.A. Bossuyt, B. Hidding, B. Tury, and R.J. West

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1 Introduction

The methylenedianiline substances (MDAS) are a family of high production volume chemicals with annual world production volume estimated to exceed four million metric tons per year (Carvajal-Diaz 2015). More than 98 % of this production is consumed as an intermediate for the production of the methylenediphenyl diisocyanate (MDI) substances, which are important monomers for the versatile thermoset polymer group of polyurethanes. Minor amounts of the MDAS are also consumed in manufacture of high performance polyimide fibers, and in manufacture of other specialty chemicals and resins.

The industrial-scale production of the MDAS occurs via a condensation reaction of aniline and formaldehyde, as shown in Fig. 1.

The bulk product of this reaction is commonly referred to as polymeric methylenedianiline (pMDA), and the relative proportions of the illustrated components can be controlled by adjusting the ratio of aniline and formaldehyde reactants. It should be noted here that the name “polymeric MDA” does not necessarily indicate that this reaction mixture meets the current OECD definition of “polymer”. In many instances, due to > 50 % of composition coming from 4,4'-methylenedianiline, the “polymeric MDA” reaction product would not meet this OECD definition (<http://www.oecd.org/env/ehs/oecddefinitionofpolymer.htm>). The pMDA reaction products can be further isolated or purified by fractional distillation, making possible any number and combination of the substances listed in Table 1 (Alport et al. 2003). The 2:1 condensation products are sometimes called “2-ring-MDA” but are hereafter referred to as methylenedianiline (MDA). When isolated from the bulk pMDA reaction mixture the 2-ring MDA will typically occur

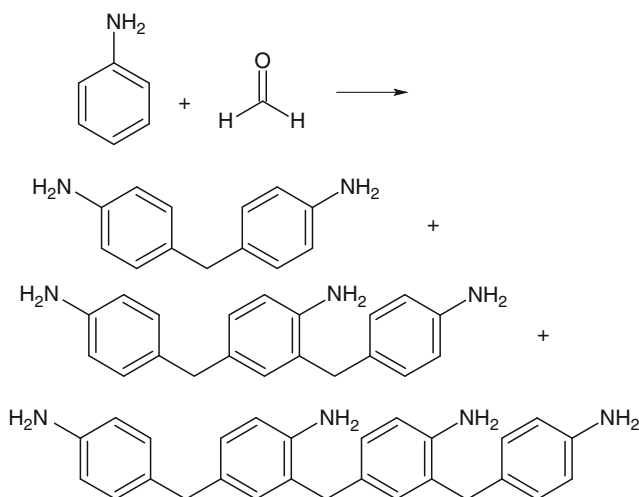
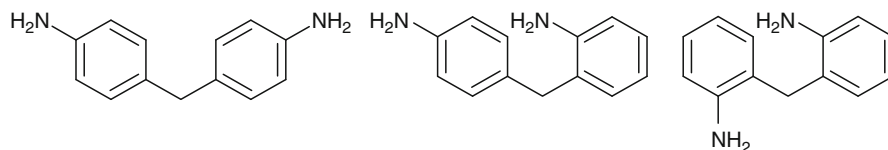


Fig. 1 Generalized commercial synthesis route of the methylenedianiline substances (MDAS)

Table 1 Identity of the commercially-relevant methylenedianiline substances (MDAS)

Substance	Chemical name	Chemical Abstracts Registry Number	Synonyms	Typical % in MDA	Typical % in pMDA
Polymeric Methylene-dianiline (pMDA)	Formaldehyde, polymer with benzenamine	25214-70-4	pMDA	N/A	100
Methylenedianilines (MDA)	Benzenamine, 4,4'-methylenebis-	101-77-9	4,4'-MDA	90–95	50
	Benzenamine, 2, 4'-methylenebis-	1208-52-2	2,4'-MDA	2–5	
	Benzenamine, 2,2'-methylenebis-	6582-52-1	2,2'-MDA	<1	
Oligomeric methylenedianilines	3-ring	N/A	oMDA	N/A	25
	4-ring			N/A	12
	5-ring			N/A	6
	Higher oligomers			N/A	<6

**Fig. 2** Molecular structures of the 4,4'-; 2,4'-; and 2,2'-isomers of methylenedianiline

as a mixture of three positional isomers (Fig. 2), where the 4,4'-MDA is the predominant isomer representing more than 90–95 % of the isomer mixture, with the 2,4'-MDA and 2,2'-MDA making up 2–5 % and less than 1 % of the mixture, respectively. The higher oligomer components of pMDA are sometimes named “3-ring-MDA”, “4-ring-MDA” and so on, and are hereafter referred to as oligomeric MDA or oMDA.

In the evaluation of physical-chemical and toxicological properties of chemical substances, it is desirable to conduct testing on commercially-relevant substances which occur at a high-purity or as a single-component. For this reason, and because it is the most commercially prominent of the methylenedianiline substances, the 4,4'-MDA substance has been intensely investigated for its physical-chemical and toxicological properties. While some properties have been studied for representative pMDA mixtures, very few studies have been conducted on the less prominent isomers of MDA and apparently none on the isolated oMDA homologues. Thus, properties of these latter substances are often inferred (i.e., by read-across) from those of 4,4'-MDA. Though mammalian toxicological hazards are not a subject of this review, it should be mentioned that 4,4'-MDA is classified by several

authorities as a genotoxic carcinogen. In 2001, the European Union issued the EU risk assessment report on 4,4'-MDA which provided a summary of all physical-chemical and hazard property data known by then (European Union 2001). Since that time, and in preparation for the registration under commission regulation 1907/2006 in the EU, some more data on 4,4'-MDA were generated. Robust summaries of these past and more recent studies are now available to the public via the European Chemicals Agency (ECHA) web site (<http://echa.europa.eu/web/guest/information-on-chemicals/registered-substances>). In addition, the government of Canada has recently completed a draft screening level risk assessment for a grouping of the MDA, pMDA, and MDI substances (Government of Canada 2014). These and other regulatory assessments can provide good overviews of the available physical-chemical, health, and environmental properties of the MDAS family, as well as their assessed potential exposures and risks. This current review provides a deeper focus and summary of the known ecotoxicological properties of the MDAS family as available from published and private company studies available through year 2014. A similar in-depth review and summary of environmental fate properties for the MDAS family, including the aspect of bioaccumulation, will be reviewed in a subsequent review article of this journal.

The majority of studies cited in this review and in the aforementioned regulatory assessments of the MDAS have been commissioned by the International Isocyanates Institute, Inc. (III; <http://diisocyanates.org/>) and its private company members, and have not been published previously. For such cited references which are not from publically available sources, the ECHA web site (ECHA 2014a, b) provides robust study summaries for these studies.

2 Physical-Chemical Properties

The knowledge of the physical-chemical properties of the MDAS and their environmental behavior is essential to the complete understanding of the ecotoxicological effects of this substance family. The physical-chemical properties of 4,4'-MDA and pMDA are summarized in Table 2. These data for 4,4'-MDA are summarized in the EU risk assessment report (European Union 2001; McNabb *et al.* 1999) and in the more recent robust study summaries provided by ECHA (2014a). The substance tested consisted of 97.39 % 4,4'-MDA, 1.98 % 2,4'-MDA, 50 ppm 2,2'-MDA, and less than 10 ppm aniline.

Physical-chemical properties for pMDA are not that uniform and depend on the composition, as pMDA is an oligomeric mixture which varies slightly in composition based on the ratio of its reactants (Table 1). Robust study summaries of these pMDA properties are also provided by ECHA (2014b).

As weak bases, the water solubilities of the MDAS depend on the pH. Though the individual molecules possess two or more primary amino-groups, their pKa

Table 2 Summary of relevant physical-chemical properties of the 4,4'-methylenedianiline (MDA) and polymeric methylenedianiline (pMDA) substances

Property	4,4'-Methylenedianiline (MDA)	References	Polymeric Methylenedianiline (pMDA) ^a	References
Physical state at 20 °C	White solid	ECHA (2014a)	Viscous liquid	ECHA (2014b)
Melting point (°C)	90–92	ECHA (2014a)	Glass transition at 0.8 and –2.7	ECHA (2014b)
Boiling point (°C)	398 ± 5 @ 101.3 kPa	ECHA (2014a)	410.6 @ 101.3 kPa	ECHA (2014b)
Bulk density (g/cm ³ , 20 °C)	1.150	ECHA (2014a)	1.150	ECHA (2014b)
Vapor pressure (Pascal)	2.5 × 10 ^{−4} (25 °C)	ECHA (2014a)	<1 × 10 ^{−4} (20 °C) 1.6 × 10 ^{−4} (50 °C)	ECHA (2014b)
Water solubility (g/L at 25 °C)	pH 5.3 = 2.2 pH 7 = 1.01 pH 9 = 0.84	ECHA (2014a)	pH 7 = 0.36–1.22	ECHA (2014b)
Octanol-water partition coefficient (Log Pow)	1.55	ECHA (2014a)	1.2–2.7	ECHA (2014b)
Dissociation constant (pKa at 20 °C)	4.96	ECHA (2014a)	Not determined	N/A

^apMDA sample containing 58 % 2-ring-, 23 % 3- ring-, 10 % 4-ring- and 3 % 5-ring-isomers; 6 % higher oligomers

values are essentially indistinguishable by the spectrophotometric method of OECD Guideline 112 (OECD 1981). The apparent equivalence of the pKa values for these amino groups is owed to the fact that delocalization of the positive charge of the corresponding ammonium ions across the molecule is restricted by the methylene group which bridges the aromatic rings.

3 Environmental Behavior

The environmental fate behaviors of the MDAS are complex, and while summarized in a separate review, is it relevant to mention here the aspects which influence stability and bio-availability associated with ecotoxicological study. Briefly, like other primary aromatic amines, the MDAS react with natural organic matter in surface waters, sediment, and soil, forming covalent nitrogen-carbon bonds. The 1,4-addition of primary aromatic amines to α,β -unsaturated carbonyl compounds such as quinones is known to serve as a sink for sequestration of these amines in the environment (Parris 1980; Weber et al. 1996; Colón et al. 2002). A generalization of this reaction is illustrated in Fig. 3. Ferulic acid, a derivative of cinnamic acid which for example represents about 1 % wt of cereal plant dry mass, is another

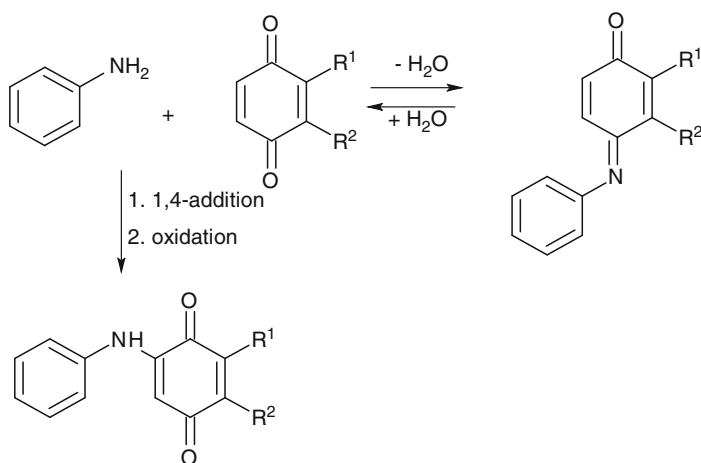


Fig. 3 Representative reaction of primary aromatic amines with quinones in humic matter, where R^1 and R^2 indicate substituent groups of the macromolecular organic matter

reaction partner for covalent binding of primary aromatic amines in the environment, as demonstrated by Tatsumi et al. (1994).

In soil degradation tests, 4,4'-MDA showed an organic carbon-normalized adsorption coefficient (K_{OC}) of 3800 and 5680 in anaerobic and aerobic soils, respectively, after an equilibration time of 8 h (Cowen et al. 1998). The 8 h value achieved approximately 90 % of the value measured after 7 days equilibration time. After 7 days, part of the MDA could be desorbed when equilibrated with fresh 0.1 M CaCl_2 for 24 h. The high affinity of MDA to soil is most likely attributable to reversible and irreversible binding of the aromatic amino group to organic matter (Tatsumi et al. 1994; Parris 1980; Weber et al. 1996; Li et al. 2000; Colón et al. 2002).

From these examples of primary aromatic amine reactivity in environmental media, it is obvious that the availability of free MDAS has to be checked in ecotoxicological experiments to be able to draw correct and robust conclusions associated with their exposure with organisms. Vice versa, for environmental risk assessment, the potential biological (non)-availability of MDAS in eco-systems needs to be considered when data from controlled, somewhat artificial laboratory experiments are taken as benchmarks.

For the performance of ecotoxicological tests, MDAS usually need to be dissolved in solvents. Although reactions may be possible, for example, Schiff base formation, MDA was shown to be sufficiently stable in acetone, ethyl acetate, acetonitrile and diluted phosphoric acid (Cowen et al. 1996).

4 Ecotoxicological Properties

The ecotoxicological properties of MDA will be summarized in the following chapters, with each chapter allocated to the aquatic, benthic, and terrestrial environmental compartments and their associated standardized tests/species.

4.1 Toxicity to Aquatic Organisms

In the following, the aquatic taxa bacteria, algae, daphnia and fish will be addressed. Data are summarized in Table 3.

4.1.1 Tests with Bacteria

Fujiwara (1981) investigated the effect of MDA on *Escherichia coli* at 25 °C. In physiological saline, 100 mg/L MDA caused a measurable reduction in cell count over 5 days, whereas 50 mg/L was the no-observed-effect concentration (NOEC). In nutrient broth, 100 mg/L MDA did not cause a reduction in cell count compared to the control.

Caspers et al. (1986) investigated the inhibition of activated sludge by 4,4'-MDA. The composition of the test substance was 99.7 % 4,4'-MDA, 0.25 % 2,4'-MDA and 0.05 % 2,2'-MDA. The test was performed according to guideline OECD No. 209 (1984). After a 3 h contact time, oxygen consumption was measured for 0.5 h. Test concentrations were 1, 10 and 100 mg/L (nominal). At each concentration tested, the inhibition was 15 %. The MDA concentration was below the limit of solubility, and the lack of a dose–response is somewhat odd, but the validity of the test system was checked with the positive control 3,5-Dichlorophenol. The authors concluded the 3 h-EC₅₀ > 100 mg/L. It should however be noted that EC₅₀ values for bacteria are dependent on the biomass loading of the tested inoculum. Evidence for inhibition of bacterial respiration has been observed in biodegradation screening tests employing as little as 10 mg/L 4,4'-MDA and 30 mg/L (dry solids) activated sludge (unpublished data of The Dow Chemical Company).

Bringmann and Meinck (1964) reported that MDA inhibits the digestion of glucose by *Pseudomonas putida* at a concentration of about 15 mg/L. The digestion of glucose resulted in a decrease of the pH value from pH = 7.5 to pH = 6.0. The lowest test concentration, which resulted in a higher pH value at the end of the test compared to the control, was called inhibition concentration. Details on the composition of the test substance are not provided.

Kaiser and Palabrica (1991) investigated the effect of 4,4'-MDA (99 % pure) on the luminescent bacterium *Photobacterium phosphoreum* (i.e., *Vibrio fischeri*). The

Table 3 Summary of 4,4'-methylenedianiline (MDA) toxicity to aquatic organisms

Species	Endpoint and duration	Result (mg/L)	References
Bacteria <i>Escherichia coli</i>	5 days, cell count	LOEC = 100 NOEC = 50	Fujiwara (1981)
Bacteria activated sludge	3 h, respiration inhibition,	EC ₁₅ = 1 EC ₅₀ > 100	Caspers et al. (1986)
Bacteria <i>Pseudomonas putida</i>	16 h, growth	EC ₂₀ = 15 ^a	Bringmann and Meinck (1964)
Marine bacteria <i>Vibrio fischeri</i>	0.5 h, luminescence	EC ₅₀ = 6.6	Kaiser and Palabrica (1991)
Bacteria <i>Ochrobactrum anthropi</i>	35 days, respiration inhibition	EC ₅₀ = 53	Kim et al. (2002)
Fungus <i>Aspergillus</i> sp.	35 days, respiration inhibition	EC ₅₀ = 53	Kim et al. (2002)
Green algae <i>P. subcapitata</i>	72 h	E _b C ₅₀ = 5.34 E _r C ₅₀ = 14.4 NOEC _b = 0.93 NOEC _r = 9.3	Mitsubishi Chemical Safety Institute (2008a)
Green algae <i>S. subspicatus</i>	96 h; growth	EC ₂₀ = 31 ^a	Bringmann and Meinck (1964)
Green algae <i>S. subspicatus</i>	72 h; E _r C ₅₀	21	Rufli and Mueller (1985a)
Green algae <i>S. subspicatus</i>	72 h;	E _b C ₅₀ = 9.8 E _r C ₅₀ = 11.0 E _b C ₁₀ = 2.4 E _r C ₁₀ = 0.3	European Union (2001)
Green algae <i>C. pyrenoidosa</i>	N.L. ^b	1–10	Rhône-Poulenc Chimie (1977)
Cyanobacterium <i>S. cedrorum</i>	N.L. ^b	1–10	Rhône-Poulenc Chimie (1977)
Marine Diatom <i>N. fustulum</i>	N.L. ^b	10–100	Rhône-Poulenc Chimie (1977)
Crustacean <i>Daphnia magna</i>	EC ₅₀	5.7 (24 h)	Rufli and Mueller (1985c)
Marine Crustacean <i>Moina macrocopa</i>	EC ₅₀	2.3 (24 h)	Fujiwara (1982)
Marine Crustacean <i>Moina macrocopa</i>	14-days	0.15	Fujiwara (1982)
Crustacean <i>Daphnia magna</i>	EC ₅₀	0.25 (48 h)	Bringmann and Meinck (1964)
Crustacean <i>Daphnia magna</i>	EC ₅₀	8.08 (24 h); 2.47 (48 h)	Mitsubishi Chemical Safety Institute (2008b)
Crustacean <i>Daphnia magna</i>	EC ₅₀	0.19–0.6 (24 h); 0.019–0.06 (48 h)	Mitsubishi Chemical Safety Institute (2008c)
Crustacean <i>Daphnia magna</i>	21-day	NOEC = 0.00525 LOEC = 0.0182	Mitsubishi Chemical Safety Institute (2008c)

(continued)

Table 3 (continued)

Species	Endpoint and duration	Result (mg/L)	References
Crustacean <i>Daphnia magna</i>	EC ₅₀	1.5 (24 h); 0.35 (48 h)	Salinas (2011)
Crustacean <i>Daphnia magna</i>	EC ₅₀	2.19 (24 h); 0.46 (48 h)	Salinas (2012) ^c
Mollusca <i>Limnea stagnalis</i>	20-h LC ₅₀	210	Rhône-Poulenc Chimie (1977)
Zebrafish <i>Danio rerio</i>	96-h LC ₅₀	42	Rufli and Mueller (1985b)
Rainbow trout <i>Oncorhynchus mykiss</i>	96-h LC ₅₀	39	Rufli and Mueller (1985c)
Zebrafish <i>Danio rerio</i>	96-h LC ₅₀	65	Caspers et al. (1986)
Golden orfe <i>Leuciscus idus</i>	96-h LC ₅₀	53	Munk and Kirsch (1988)
Medaka <i>Oryzias latipes</i>	96-h LC ₅₀	20.6	Mitsubishi Chemical Safety Institute (2008d)

^aAssumed to be an EC₂₀ (see text)^bNoxious limit (see text)^c56.9 % 4,4'-MDA, 7.1 % 2,4'-MDA, 26 % 3-ring MDA, 10 % 4-ring MDA and higher oligomers

EC₅₀, determined as the concentration resulting in 50 % reduction in light emission compared to control after 30 min contact time, was 6.6 mg/L.

In the scope of biodegradation tests with 4,4'-MDA, Kim et al. (2002) demonstrated that at a concentration of 30 mg/L MDA, CO₂ evolution reached about 70–80 % of the theoretical yield over 35 days with non-adapted activated sludge as well as with enrichment cultures of *Ochrobactrum anthropi* or *Aspergillus* sp. These enrichment cultures were generated by the incubation of soil extracts taken from areas where dehydrated activated sludge of the waste water treatment plant was buried with MDA as the sole carbon source. However, at concentrations of 50, 100 or 300 mg/L MDA, CO₂ evolution reached levels of about 50, 20 and 5 % after 35 days, respectively. Data were presented in diagrams and, therefore, any further interpretation has to be regarded with care. If it is assumed that 30 mg/L MDA had no effect on the microorganisms, dosages of 50, 100 and 300 mg/L reduced the CO₂ evolution down to levels of 71, 21 and 7 % for *O. anthropi* and to 57, 14 and 7 % for *Aspergillus* sp. A rough logit analysis of the graphical data given in the paper results in EC₅₀ values of about 53 mg/L for *O. anthropi* and *Aspergillus* sp., however the concentration of biomass associated with these tests was not quantitatively determined. From the dose–response curves it is not clear whether or not 50 mg/L represent already an inhibition of the microorganisms or whether there is simply a substrate overload; 100 mg/L, however, is clearly inhibiting for *Aspergillus* sp., and for *O. anthropi* the formation of CO₂ comes to a halt after

25 days. Further, at 30 mg/L the CO₂ production rate is virtually the same between the non-adapted sludge and the enrichment cultures; this result indicates toxicity of MDA to MDA-digesting microorganisms.

4.1.2 Tests with Algae

The toxicity of 4,4'-MDA to a freshwater single-celled algae (*Pseudokirchneriella subcapitata*; previously *Selenastrum capricornutum*) was investigated by the Mitsubishi Chemical Safety Institute in 2002, and an English translation was made available as International Isocyanate Institute report 11545 (Mitsubishi Chemical Safety Institute 2008a). The test was performed according to OECD Guideline No. 201 (2006). The purity of the test substance was 99.6 %, and test concentrations were checked by HPLC-UV. The test substance concentration was constant (± 20 %) over the exposure time. In the European Union, the median growth rate inhibition concentration (E_rC_{50}) is the preferred endpoint; in this article, the endpoint of median biomass inhibition concentration (E_bC_{50}) is provided additionally. Against biomass, after 72 h the E_bC_{50} was 5.34 mg/L (95 % C.I. 3.57–7.98 mg/L) and the $NOEC_b$ was 0.93 mg/L. In terms growth rate, the E_rC_{50} was 14.4 mg/L (24–72 h) and the $NOEC_r$ was 9.3 mg/L.

In a study with the colonial freshwater alga, *Scenedesmus subspicatus*, the 72 h E_rC_{50} was 21 mg/L (95 % C.I. 11–29 mg/L) for 4,4'-MDA (Rufli and Mueller 1985a). The test was performed according to OECD Guideline 201 (1981). The test substance was named TK 10504 (commercial grade) and claimed to be 4,4'-MDA. Data on purity are not provided. However, the European Union cites this report (2001) and mentions a purity of 95.5–98 %. Results were based on measured concentrations and the data were provided; the analytical method is not mentioned.

Caspers and Mueller (1992) investigated the toxicity of pMDA on *Scenedesmus subspicatus*. After 72 h exposures, the E_bC_{10} and E_bC_{50} values were 2.4 and 9.8 mg/L, and the E_rC_{10} and E_rC_{50} values were 0.3 and 11.0 mg/L, respectively.

Bringmann and Meinck (1964) reported a limit concentration of 31 mg/L for the inhibition of growth for *Scenedesmus* for 4 days exposure. It is assumed that this level is equal to an EC_{20} .

A pMDA sample containing 60 % wt 4,4'-MDA was tested against various algae/cyanobacteria (Rhône-Poulenc Chimie 1977). In those tests, a “noxious limit” was established, which was a concentration where the growth of the organisms was inhibited against a control. Tests were stated to be “run for a time sufficient to detect multiplication of organisms in the control”. For the green alga *Chlorella pyrenoidosa* and the blue-green alga *Synechocystis cedrorum* (i.e., *Cyanobacterium cedrorum*) the noxious limit was in the range from 1.0 to 10.0 mg/L. For the marine diatom *Nitzschia frustulum* the noxious limit fell between 10.0 and 100.0 mg/L. Test concentrations were verified to be controlled within initial nominal values using a photometric method.

As the Mitsubishi studies are the only ones conducted and documented under Good Laboratory Practice (GLP) standards for quality assurance (Mitsubishi Chemical Safety Institute 2008a) these data could be regarded as the most reliable for environmental risk assessment. The data generated by Rhône-Poulenc Chimie (1977) add value in so far as there seems not to be a significant greater sensitivity of cyanobacteria and diatoms compared to green algae.

4.1.3 Tests with Crustacea

With the freshwater crustacean *Daphnia magna*, data from several acute and chronic test exposures are available. Rufli and Mueller investigated the acute toxicity of 4,4'-MDA against *Daphnia magna* (Rufli and Mueller 1985b) according to OECD Guideline 202 (1981). The exposure time was 24 h, whereas the current guideline requires a 48 h exposure. Due to a resulting odd dose–response behavior (Fig. 4), an EC_{50} could not be established. The authors reported 24 h EC_0 and EC_{100} values of < 3.2 and > 100 mg/L, respectively. However, taking the three lowest concentrations only, a more meaningful dose–response relationship becomes evident, and the 10 mg/L exposure would appear to cause about 75 % immobility. Exposure concentrations of this study are based on measured concentrations, but the analytical method was not mentioned in that report. By probit analysis, an EC_{50} of 5.7 mg/L can be calculated; however, using this reduced data set a 95 % C.I. cannot be derived.

Fujiwara (1982) reported a 24 h EC_{50} of 2.3 mg/L for *Moina macrocopa*, a common aquatic invertebrate in the Asia-pacific region which tolerates high salinity (95 %-C.I.: 1.8–3.0 mg/L). The test substance was claimed to be extra-pure with no

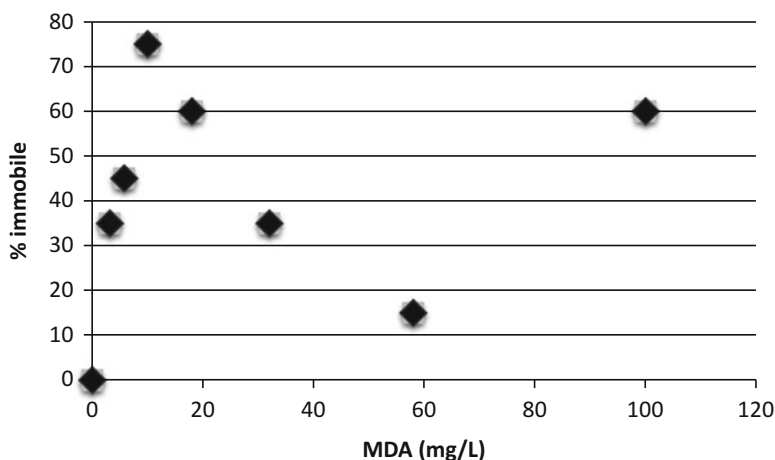


Fig. 4 Immobility responses associated with acute (24 h) exposures of 4,4'-MDA to *Daphnia magna* (Rufli and Mueller 1985b)

further details provided. The composition and pH of the test water were described. For each exposure concentration, 4 replicates of 5 animals each were prepared. The reference toxicant potassium dichromate ($K_2Cr_2O_7$) served as positive control. There is no information concerning an analytical method. Therefore, results reported are most likely based on nominal concentrations.

In an OECD Guideline 202 study conducted in compliance with GLP, the acute EC_{50} for *Daphnia magna* was 2.47 mg/L after an exposure time of 48 h (Mitsubishi Chemical Safety Institute 2008b). After 24 h, the test solutions were renewed (semi-static). The 24 h EC_{50} was 8.08 mg/L (95 % C. I.: 5.23–12.8 mg/L). After an exposure time of 48 h, the highest concentration at which no effects were observed was 0.2 mg/L, and the EC_{100} was 200 mg/L. The 95 % C.I. for the 48 h EC_{50} was 1.27–4.4 mg/L. Concentrations of the test substance were verified analytically with HPLC-UV. The purity of the test-substance was 99.6 %. Another more recent OECD Guideline 202 (OECD 2004) study on the acute toxicity of 4,4'-MDA to *Daphnia magna* was issued in 2011 (Salinas 2011). The 48 h EC_{50} was determined as 0.35 mg/L (95 % C. I. 0.18–0.71 mg/L), whereas the 48 h EC_0 and 48 h EC_{100} were 0.22 and 2.2 mg/L, respectively. The purity of the test substance was 98 %. According to HPLC-MS analyses, impurities consisted of an isomer to 4,4'-MDA, as well as some 3-ring MDA and N-methylated by-products. The concentrations of the test solutions maintained over the exposures were verified with HPLC-UV. This test was performed under static conditions.

The acute toxicity of pMDA to *Daphnia magna* was checked in an OECD Guideline 202 test (Salinas 2012). The test substance was characterized by IR, UV, HPLC-MS, GC and 1H -NMR. The tested product consisted of 55.9 % 4,4'-MDA, 7.1 % 2,4'-MDA, 0.3 % 2,2'-MDA, 26.1 % 3-ring MDA and 9.9 % 4-ring MDA and 0.7 % impurities, from which N-methyl-methylene-4,4'-dianiline and N-formyl-methylene-4,4'-dianiline could be identified. To avoid potential for physical toxicity associated with undissolved particulate matter in test solutions, pMDA was dissolved in peroxide-free THF ($H_2O_2 \leq 1$ ppm) and the THF was evaporated in a rotating round bottom flask. M4 medium was added to the dried flask and slowly shaken (50 rpm) for 24 h in the dark. Finally, the resulting test solutions were filtered through a 0.2 μm membrane. Analyses were performed to determine total dissolved organic carbon (DOC) and 4,4'-MDA concentrations by HPLC-UV. The derived 48 h EC_{50} of 0.46 mg/L (nominal) or 0.26 mg/L (as measured 4,4'-MDA), indicate that the acute toxicity of pMDA to *D. magna* is in the same order of magnitude as that for pure 4,4'-MDA.

In addition to the aforementioned acute tests with aquatic crustacea, two chronic exposure tests have been conducted to determine potential effects on survival and reproduction. Fujiwara (1982) performed a 14 days chronic test with *Moina macrocopa*. The test period covered three generations of the organism. Animals were not older than 24 h at the beginning of the test. Per test concentration, four replicates with ten individuals each were prepared. The test solutions were renewed every 48 h, and the organisms were fed with unicellular green algae daily. The 14 days NOEC for reproduction was 0.15 mg/L. There is no information concerning

an analytical method; therefore, results reported are most likely based on nominal exposure concentrations.

In 2002, Mitsubishi Laboratories performed a chronic study (21 day) with *Daphnia magna* according to GLP and OECD Guideline 211 (1998); the data were made available in English in 2008 by the International Isocyanate Institute (Mitsubishi Chemical Safety Institute 2008c). Test solutions were renewed daily. The purity of the test substance was 99.6 %, and exposure concentrations were verified using HPLC-UV analyses. In the scope of that test it was observed that the 48 h EC₅₀ was between 0.019 and 0.06 mg/L; the authors reported a 21 day NOEC of 0.00525 mg/L and a lowest observed effect concentration (LOEC) of 0.0182 mg/L based on the reproduction endpoint.

Concerning acute toxicity and LC/EC50 values for crustacea, *D. magna* is the most sensitive of all species tested with the MDAS, owing to the 48 h EC₅₀ of 0.019–0.06 mg/L (Mitsubishi Chemical Safety Institute 2008c) and the results of the Salinas (2011) study, yielding a 48 h-EC₅₀ of 0.35 mg/L. This latter value is not significantly different to that of pMDA (0.46 mg/L), tested in the same laboratory. The highest 48 EC₅₀ value reported for daphnia is 2.47 mg/L (Mitsubishi Chemical Safety Institute 2008b). Therefore, the MDA acute toxicity to crustacea spreads over two orders of magnitude if results 48h data from acute and chronic tests are considered; focused on data from acute tests only, the difference between 48 h EC₅₀ values is about 10. Shortcomings in the tests performed were not identified, and impurities are not a likely explanation as this bulk industrial chemical commodity is synthesized via a well known, standard route. The 24 h EC₅₀ values for K₂Cr₂O₇ were in the range required by OECD Guideline 202. In the Mitsubishi Chemical Safety Institute acute and chronic tests (2008b, c), the water was renewed daily. The only remaining difference identified is the fact that in the chronic test, the daphnids were fed daily with *Chlorella vulgaris*. It remains to be found out whether this difference to the acute test is at least partially responsible for the differences in results.

4.1.4 Test with Vertebrates (Fish)

Rufli and Mueller reported the acute toxicity of MDA against zebrafish (*Danio rerio*) (1985c). The static test was performed according to OECD Guideline 203 (1984). This study resulted in a 96 h LC₅₀ of 42 mg/L (95 % C.I.: 35–51 mg/L). Results were based on measured concentrations; however neither the analytical method for concentrations verification nor the accurate purity of the test substance was given. Rufli and Mueller (1985d) also reported the acute toxicity of 4,4'-MDA against rainbow trout (*Oncorhynchus mykiss*; formerly *Salmo gairdneri*), following OECD Guideline 203 (1984) under a static exposure procedure. The 96 h LC₅₀ was 39 mg/L (95 % C.I.: 20–134 mg/L). Results are based on measured concentrations, but again neither the analytical method for concentrations verification nor the accurate purity of the test substance was given.

A further report on the toxicity of MDA against zebra fish is reported by Caspers et al. (1986), following OECD Guideline 203 (1984). The authors report a 96 h LC₅₀ of 65 mg/L, and 96 h LC₀ and LC₁₀₀ values of 40 and 70 mg/L, respectively. The test substance consisted of 99.7 % 4,4'-MDA, 0.25 % 2,4'-MDA and 0.05 % 2,2'-MDA. Maintenance of the exposure concentrations over 96 h was verified using dissolved organic carbon (DOC) analyses.

Munk and Kirsch (1988) reported the acute toxicity of MDA against golden orfe (*Leuciscus idus*). The test procedure followed the German standard DIN 38412 (DIN 1980). Chloro acetamide was used as a positive control. The dissolved oxygen content of the water was throughout > 60 % of the air saturation value, pH was maintained at 8, and the water temperature was held at 20 ± 1 °C. Water hardness and salinity were kept in the required range. Purity of the test substance was claimed to be at least 96 %. The 96 h LC₅₀ was determined by graphical interpolation to be 53 mg/L. The 96 h LC₀ was 21.5 mg/L and the 96 h LC₁₀₀ was 100 mg/L, where derivation of these endpoints is based on nominal concentrations. At higher concentrations, the test substance was noted to have precipitated from solution.

In an English-translated summary report of original studies of the Chemicals Inspection and Testing Institute-Japan, a 48 h LC₅₀ of 32 mg/L was reported for the orange-red killifish, *Oryzias latipes* (CITI 1992). The testing regime was semi-static, with renewal of the exposure solutions every 8–16 h, and followed the Japanese standard JIS K 0102-1986-71. The Mitsubishi Chemical Safety Institute similarly investigated the acute toxicity of MDA with *Oryzias latipes* in 2002 according to OECD Guideline 203 (1992) and in compliance with GLP. The translated report was issued by the International Isocyanate Institute (Mitsubishi Chemical Safety Institute 2008d). Applying a semi-static method with daily renewal of the test solutions, a 96 h LC₅₀ of 20.6 mg/L was reported (95 % C.I.: 16.7–25.3 mg/L). These results were based on measured concentrations (HPLC-UV) and the test substance had a purity of 99.6 %.

4.1.5 Tests with Mollusca

The acute toxicity of a pMDA sample containing 60 % 4,4'-MDA was examined using 20 h static aquatic exposures to the Great Pond Snail (*Limnea stagnalis*). The tests were performed in reconstituted river water in the dark at 20 °C, with pH of 8.0 and dissolved oxygen maintained at ≥ 80 % air saturation. A cohort of 6–8 days post-hatch snails was investigated for egg laying performance, and a 1–4 days post-hatch cohort was checked for survival. The LC₅₀ for the post exposure 4 days survival of young snails was 210 mg/L, and for egg laying performance (fecundity) the LC₅₀ was 220 mg/L (Rhône-Poulenc Chimie 1977). Aside from this study, no other studies of MDAS with freshwater or marine mollusca are known. Based on this single study, this animal phylum would appear to be less sensitive to the MDAS substances than are the crustacea.

4.2 Toxicity to Benthic Organisms

While 4,4'-MDA was under evaluation in the European Union, effects of chemicals to benthic sediment organisms gained interest. This was particularly true for substances highly hydrophobic and/or reactive substances such as the MDAS which have high affinity for sediment and associated organic matter. In a research program of the German Umweltbundesamt (UBA; Environmental Protection Agency), the sediment blackworm *Lumbriculus variegatus* was revealed to be particularly sensitive against primary aromatic amines (Riedhammer and Schwarz-Schulz 2001). As a consequence, 4,4'-MDA was tested against the sediment organisms *Lumbriculus variegatus*, *Chironomus riparius* and *Hyalella azteca*, representing three different taxa of sediment organisms (oligochaete, insects and amphipods) and playing an important role in the aquatic food chain and in organic matter turnover. Data concerning benthic organism are summarized in Table 4.

4.2.1 Tests with an Oligochaete

The benthic oligochaete, *L. variegatus* (blackworm) inhabits a wide range of sediment types and is typically present in shallow water sediments of slowly flowing rivers which are rich in organic matter. It is feeding on mainly subsurface organic matter of dead and decaying organisms, and because primary aromatic amines are known to be covalently bound with such organic matter, the potential effects on this organism are of relevance. Egeler and Ginzburg (2001) investigated the toxicity of 4,4'-MDA to *L. variegatus*. The study was conducted in compliance with GLP, but since the now standardized OECD Guideline 225 (2007) for sediment-water *Lumbriculus* toxicity tests was still under development, and

Table 4 Summary of toxicity of 4,4'-methylenedianiline (MDA) to sediment organisms

Species	Duration	Endpoint and value (mg/L)	References
Oligochaete <i>Lumbriculus variegatus</i>	28 days	NOEC = 25.2 LOEC = 50.3	Egeler and Ginzburg (2001)
Oligochaete <i>Lumbriculus variegatus</i>	28 days	NOEC < 3.75 ^a , NOEC = 30 ^b LOEC = 60 ^b	Egeler (2002)
Insect larvae <i>Chironomus riparius</i>	28 days	NOEC = 500 LOEC = 1000	Egeler and Gilberg (2005a)
Amphipod <i>H. azteca</i>	28 days	NOEC = 41.4 ^c NOEC = 90.9 ^d LOEC = 90.9 ^c LOEC = 200 ^d	Egeler and Gilberg (2005b)

^aBased on statistics (see text)

^bBiological interpretation (see text)

^cSurvival

^dBiomass

because *Lumbriculus* turned out to be most sensitive sediment organism against MDA with a non-monotonic dose–response, a more detailed explanation on test design and results will be provided in the following section.

In a first test with *L. variegatus*, the artificial sediment consisted of 75 % quartz sand, 20 % clay and 5 % sphagnum peat. The peat content was chosen to represent a moderate content of organic matter as indicated at that time. Further, between 0.05 and 1 % pure CaCO_3 , 0.4 % TetraMin[®] fish food (Tetra Werke, Melle, Germany) and 46 % water were added per sediment dry weight. While the test was running, further TetraMin[®] was added on the surface of the sediment at the beginning of weeks 2, 3 and 4. Reconstituted water was used according to OECD guideline 203 (1992) Annex 2. During the test, the temperature was maintained at $20 \pm 2^\circ\text{C}$, and a 16/8 h light (100–1000 Lux)/dark cycle was applied. Air was slowly passed through the aqueous layer at one bubble per second. As MDA can bind covalently to sediment, spiked sediment was allowed to equilibrate for 2 days before the test organisms were added. This equilibration time was chosen to ensure that added organisms were exposed against both organic matter-bound and freely dissolve MDA. Per dose group, three replicate exposures were prepared with ten synchronized worms added to each of the 1 L test vessels, filled with equilibrated sediment and reconstituted water. Nominal concentrations of 4,4'-MDA (purity 97.9 %) were 0, 1.6, 3.1, 6.3, 12.6, 25.2, 50.3 and 100.7 mg/kg sediment dry weight. The spiking solutions, overlying water, and sediment pore-water were analyzed by HPLC-UV. During the test, the dissolved oxygen in the overlying water was at least 74 % of the air saturation value, and the pH was kept between 6.0 and 7.0. At the end of the 28 days exposure period, the number of worms and total worm dry weight were analyzed by ANOVA and Dunnett's *t*-test. In the control group, mortality did not exceed 7 %. For reproduction (total number of worms and number of regenerated worms), biomass, and number of worms with new posterior and anterior end, a clear dose–response pattern could not be observed (Figs. 5–7). Due to the reactivity of MDA with organic matter, and possibly also due to its biodegradation in the sediment-water test system, its concentration in the aqueous phase declined over

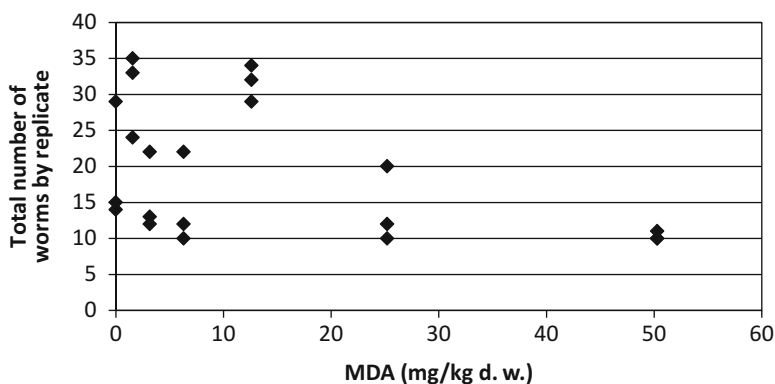


Fig. 5 Representative plot of number of *L. variegatus* worms recovered from replicate 4,4'-methylendianiline (MDA) sediment-water exposure vessels after 28 days (adapted from Egeler and Ginzburg 2001)

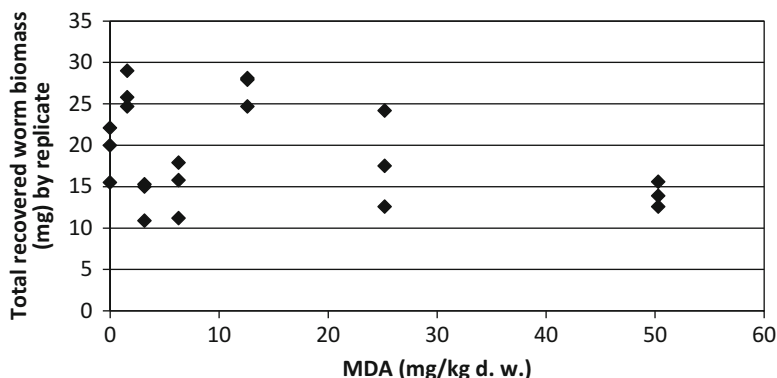


Fig. 6 Representative plot of total worm biomass recovered from replicate 4,4'-methylenedianiline (MDA) sediment-water exposure vessels after 28 days (adapted from Egeler and Ginzburg 2001)

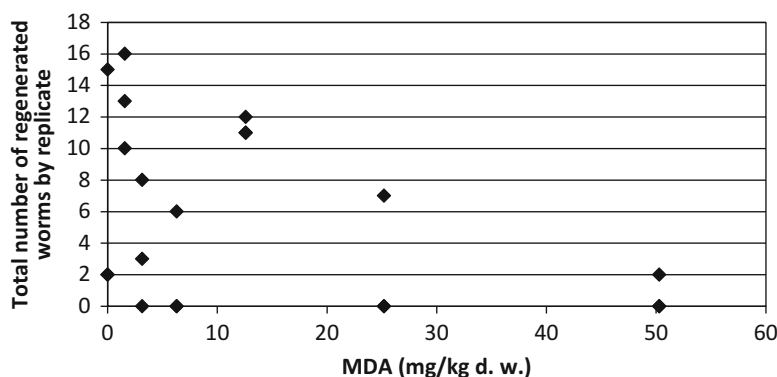


Fig. 7 Representative plot of total of total number of *L. variegatus* worms with new posterior/anterior ends recovered from replicate 4,4'-methylenedianiline (MDA) sediment-water exposure vessels after 28 days (adapted from Egeler and Ginzburg 2001)

time. The mean recovery of MDA was about 4 % of that initially applied after 28 days. Based on nominal concentrations, the NOEC and LOEC values for biomass were 50.3 and 100.7 mg/kg d. w., respectively, whereas for reproduction the values were 25.2 and 50.3 mg/kg d. w. Based on measured concentrations after 28 days, the NOEC and LOEC were 2.1 and 4.1 mg/kg d. w. against biomass and 1.0 and 2.1 mg/kg d. w. for reproduction. For lower nominal concentrations, extractable MDA was not detectable at the end of the test period (<0.1 mg/kg d. w.). In freshly-spiked sediment, recovery of MDA was 82–102 % of the applied concentration. Therefore, the extraction and analytical methods as such were effective. The disappearance of the MDA in these test systems was most likely attributable to its irreversible binding to organic matter of the formulated sediment, and perhaps also biodegradation in both sediment and water layers. In the previously described test with *L. variegatus*, it was speculated the iterative feeding with

TetraMin[®] could predispose the test organisms towards selective feeding on this uncontaminated food versus the MDA-associated sediment organic matter. In such a case, the true toxicity of MDA-associated sediment organic matter may be misrepresented. A second study was performed to check whether sediment amended with an organic matter food source— here nettle powder (*Urtica* sp.)— instead of semi-continuous feeding would deliver different results (Egeler 2002). Nominal concentrations of 4,4'-MDA (purity 97.9 %) were 0, 3.75, 7.5, 15, 30 and 60 mg/kg sediment dry weight. Due to high variability among replicates observed in the previous test, six replicates were used for the control and three replicates for each exposure concentration. The test solutions used to spike the formulated sediment with MDA were verified by HPLC-UV, but further analyses of pore-water and overlying water was deemed not necessary due to the reactivity of MDA to sediment organic matter.

A dose–response curve could not be fitted to the data due to high variations between dose groups (Figs. 8 and 9). For the total recovered biomass endpoint, the lowest test concentration of 3.75 mg/kg d. w. was the LOEC after 28 days. For reproduction, the 28 days NOEC was 3.75 mg/kg and the 28 days LOEC was 7.5 mg/kg d. w. sediment. For regenerated worms with new anterior and posterior ends, the effect of MDA was more prominent. In the control vessels, the number of complete new worms ranged from 2 to 8. In all exposure vessels, there was at maximum one complete new worm.

The results from these two tests with *L. variegatus* would indicate that the feeding regimen employed can influence the derived effective concentrations for biomass, reproduction, and regeneration of this sediment-feeding worm. The test which employed feeding via the overlying water appeared to result in higher effective concentrations for the MDA substance. However, it cannot be determined how difference in feeding modes among these two studies could influence other factors such as microbial activity (biodegradability MDA) and ammonia production

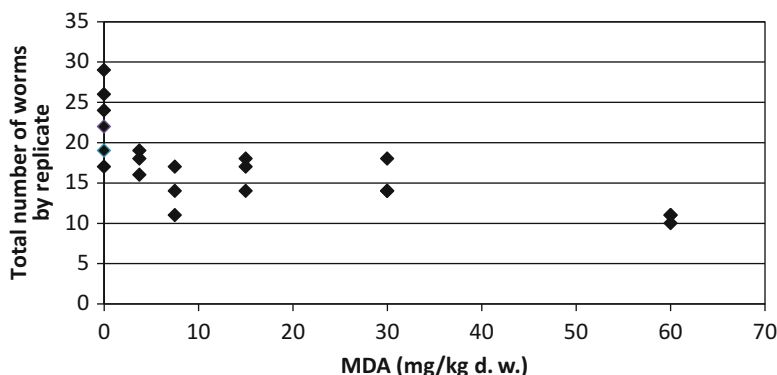


Fig. 8 Representative plot of total of total number of *L. variegatus* worms recovered from replicate 4,4'-methylendianiline (MDA) sediment-water exposure vessels after 28 days (adapted from Egeler 2002)

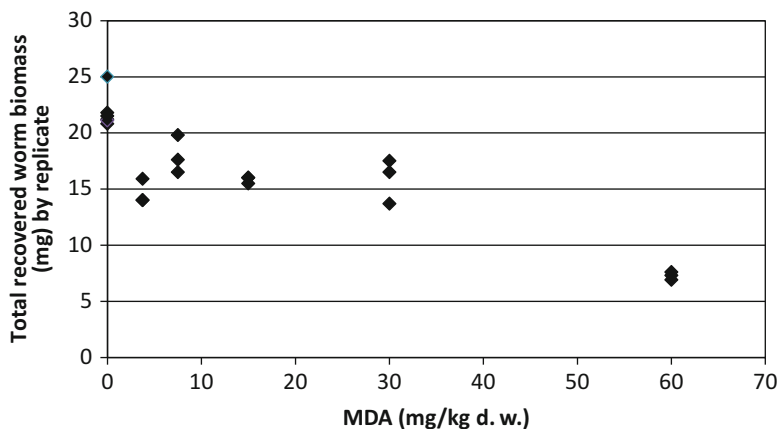


Fig. 9 Representative plot of total worm biomass recovered from replicate 4,4'-methylenedianiline (MDA) sediment-water exposure vessels after 28 days (adapted from Egeler 2002)

as confounding toxicant in these sediment-water test systems. Nevertheless, this benthic oligochaete was shown to be most sensitive to MDA in both tests, compared to other species evaluated.

4.2.2 Tests with an Insect Larvae

Larvae of the harlequin fly (*Chironomus riparius*) are living in and on sediments, and are feeding on organic matter deposited in the sediment during this life cycle stage. As such, they too are a relevant test organism for assessing potential hazard of substances such as the MDAS which bind with sediment organic matter. In an OECD Guideline 218 study, Egeler and Gilberg (2005a) tested the chronic toxicity of 4,4'-MDA (purity 97.9 %) to *C. riparius*. Prior to test substance addition, the sediment was spiked with 0.4–0.5 % stinging nettle powder as the sole food source. As West et al. (2004) discovered, the spiking of sediment with nettle powder can produce significant amounts of ammonia. As a result, in this study a tighter control of pH values was maintained to maximize speciation of any ammonia produced as the less toxic ammonium ion. In addition, aeration of the overlying water layer was doubled from that employed in the *L. variegatus* tests from one to two bubbles of air per second, so that any un-ionized ammonia (gas) would be more readily expelled. Larvae were added 2 days after spiking the sediment with MDA. This procedure ensured that part of the test substance is already bound to sediment, while another part is still available in the aqueous phase. Nominal test concentrations of 4,4'-MDA (purity 97.9 %) were 62.5, 125, 250, 500 and 1000 mg/kg d. w. sediment. Due to the experience with previous tests, MDA was analyzed in stock solutions only, but not in sediment and overlying water. The pH was kept in the range of 7.3–8.3 by occasional addition of diluted HCl (pH up to 9 would be acceptable

according to the guideline). The ammonium content in the overlying water ranged from 0.2 to 16.1 mg/L in the exposures employed in this test. For the emergence ratio endpoint, the 28 days NOEC was 500 mg/kg dry weight, and 1000 mg/kg d. w. was the 28 days LOEC. For the development rate endpoint, 1000 mg/kg d. w. was determined to be the 28 days NOEC. Thus, the larvae of the *C. riparius* insect appear to be much less sensitive to MDA than is the oligochaete *L. variegatus*.

4.2.3 Test with an Amphipod

A third organism has been tested with chronic exposures to MDA, to enable an analysis of species sensitivity across three taxa of benthic organisms and their associated feeding modes. The amphipod crustacean *Hyalella azteca* shows some subsurface deposit feeding, but has a stronger affinity to feeding on organic matter deposits at the surface of sediments than the previously describe benthic organisms. The *H. azteca* organism is very sensitive to ammonia (Whiteman et al. 1996); the sensitivity depends on the pH value of the test medium. At a nominal concentration of 50 mg/L NH_4^+ (148 mg/L NH_4Cl) in the overlying water, all exposed amphipods survived a 4 days exposure period in M4 medium at pH values not higher than 8.5. When the pH value was raised to 9, all exposed organisms died within 24 h (Egeler 2004).

Egeler and Gilberg (2005b) tested the chronic toxicity of 4,4'-MDA to *H. azteca*. The testing regime was performed in compliance with GLP and followed OECD Guideline 218 (OECD 2004). The chironomus test conditions were adapted to testing with *H. azteca*, as an OECD guideline for testing with this organism does not exist. In this adapted test, the amount of stinging nettle powder incorporated into the sediment was reduced from the guideline recommended 0.5 % d.w. to 0.25 % d.w. This was thought to reduce potential for ammonia formation from this N-containing organic matter, and 0.25 % cellulose (which contains no nitrogen) was added to replace amount of eliminated nettle. These sediment-incorporated food sources were equilibrated in the test system before addition of the test substance. Nominal concentrations of 4,4'-MDA (purity 97.9 %) were 8.5, 18.8, 41.3, 90.9 and 200.0 mg/kg sediment d. w.; spiking solutions were analyzed by HPLC-UV, but no analysis was performed for the sediment and the overlying water due to the known reactivity of MDA to sediment organic matter. After a 2 day equilibration period organisms were added, and pH was controlled daily between 7.4 and 8.3 by occasional addition of dilute HCl. After 28 days, the NOEC and LOEC for survival were 41.3 and 90.9 mg/kg d. w., respectively. With regard to the amphipod length and total biomass, the 28 days NOEC and 28 days LOEC were 90.9 and 200 mg/kg d. w., respectively.

4.2.4 Additional Tests on Feeding Mode and Ammonia Formation

Over the course of the aforementioned testing of MDA with benthic organisms, the potential confounding of results from feeding of the organisms, the feed types used, and potential for associated ammonia formation from decay of the food was questioned. Additional studies were conducted to improve the understanding of how these test parameters may have influenced results and interpretations of the aforementioned tests. West et al. (2004) reported that spiking of sediment with nettle powder (*Urtica* sp.) at the beginning of formulated sediment tests results in a stronger and earlier increase in ammonia concentration than feeding via the overlying water with semi-continuous addition of TetraMin[®]. The discovery of these feeding-related influences on ammonia production prompted further investigation using a factorial design study to investigate the influence of different feeding parameters on the water quality in sediment tests with pre-spiked sediments (Egeler and Gilberg 2005c). The food spiked into the sediment was either *Urtica*: cellulose = 1:1 (wt:wt) or cereal leaves (wheat; *Triticum aestivum*), test-substance (4,4'-MDA, 97.9 %) was added or not (10 mg/kg d. w.), aqueous medium was either M4 (Elendt medium) or a pH-reduced medium (by reduction of NaHCO₃), and organisms added were either the blackworm *L. variegatus* or the amphipod *H. azteca*. The tests were run for 28 days. For *L. variegatus*, cereal leaves was shown to be the best food source concerning total number of worms and biomass. The *urtica* powder/cellulose combination food source resulted in a significant reduction of biomass, especially when the M4 medium was modified. For *H. azteca*, survival and length were not influenced by any of the feeding factors evaluated, but the biomass was significantly reduced when the modified M4 medium was used. The ammonia concentration in the pore water and overlying water was significantly higher when cereal leaves were used as food source. The presence of MDA had no measureable influence on the ammonia content. This can be regarded as an indication that nitrification is not inhibited by up to 10 mg/kg d. w. MDA in the formulated sediment.

In the previously-described second test with the blackworm *L. variegatus* (Egeler 2002), pH values were taken at day 1 and 28 and found both to be below pH 8. However, pH values were not recorded in between and ammonia was not monitored. In sediment spiked with nettle powder, the pH exceeded a value of 8 after 48 h, and then declined to levels below 8 after 11 day, influenced by addition of diluted HCl. Ammonia, however, achieved the highest concentration at day 4 of the experiment (West et al. 2004). The authors further demonstrated that ammonia formation is less a problem in case of semi-continuous feeding with TetraMin[®]. As a result, the lower NOEC and LOEC in the second study with *L. variegatus* are probably attributable to interference by ammonia. Although the number of worms with new posterior and anterior end seems to be most sensitive parameter in the second study, results from the first study demonstrated that this endpoint suffers from a high variability even in the control. As a result, this endpoint is not suitable

for the interpretation of the toxicity of MDA to *L. variegatus* under the test design employed.

Sediment tests performed with MDA generally showed a high variability in outcomes. This was at least partly attributable to the confounder ammonia production. In addition, MDA can be regarded as a “difficult substance” for sediment testing. For example, in an OECD Guideline 308 study simulating the fate of radiolabeled MDA in a surface water and sediment system (Schaefer and Ponizovsky 2013), there was no MDA detectable in the aquatic layer 7 days after test initiation. After 100 day, 5.7 % of the MDA was transformed to CO₂, about 6 % MDA metabolites were detectable in the water, 12 % MDA products were extractable from the sediment and 90 % remained unextractable (retrievability about 114 %). Especially in the first few days after addition to the sediment, the MDA suffers rapid transformations against the time frame of the chronic sediment tests. The parent substance and the metabolites are assumed to show different toxicity against the sediment organisms. The design as used was chosen as such because it is not known whether MDA itself or one of the transformation products is most critical. The variability in test outcomes observed seems to be attributed to subtle variations in conditions of these dynamic test systems.

4.3 Toxicity to Terrestrial Organisms

The reactivity of the MDAS with natural organic matter indicates the potential for their association with surface soils should direct exposures occur as a result of spillages, or indirect exposures such as a hypothesized hydrolysis and deposition of atmospheric emissions of MDI substances (Government of Canada 2014). Data with soil organisms are summarized in Table 5.

4.3.1 Tests with Soil Bacteria

Soil bacterial serve a critical role in the cycling of nitrogen in the environment. The decay of nitrogen-containing organic matter can result in the release of ammonia, and this ammonia can then be oxidized by specific nitrifying bacteria to nitrite and nitrate. Because several aromatic amine substances were shown to behave as inhibitors of these nitrifying bacteria in soil (Zhang et al. 2010), the potential for MDAS to have similar effect was evaluated using a modification to the OECD Guideline 216 (2000). The standard version of this OECD test recommends that plant material is added to soil and to check for the formation of nitrate. The principle of this test is that as heterotrophic soil bacteria decompose this plant material to release ammonia, the amount of nitrate formed from this ammonia is dependent upon the activity of the specific denitrifying bacterial population in the soil. A reduction in the rate or total mass of nitrate production in soil treated with the test substance, compared to that in an untreated control soil, is interpreted as

Table 5 Summary of acute and chronic toxicity studies for 4,4'-methylenedianiline (MDA) with soil organisms

Organism	Endpoint	Value (mg/kg d. w.)	References
Oats <i>Avena sativa</i>	17-days growth	EC ₅₀ = 353	Van der Hoeven et al. (1992a)
Lattuce <i>Lactuca sativa</i>	17-days growth	EC ₅₀ = 128	
Radish <i>Raphanus sativus</i>	6-days emergence	EC ₂₀ = 96 ^a EC ₅₀ = 331 ^a	Kim et al. (2002)
Earthworm <i>Eisenia fetida</i>	14 days mortality	EC ₅₀ = 444	Van der Hoeven et al. (1992b)
Earthworm <i>Eisenia fetida</i>	56 days reproduction/ mortality	EC ₅₀ = 333	Moser and Hamberge (2012)
		EC ₁₀ = 11	
		LOEC = 18	
Springtail <i>Folsomia candida</i>	Reproduction 28 days	NOEC = 562	Moser and Schott (2011)
		LOEC = 1000	
Bacteria Nitrification in soil	NH ₄ ⁺ depletion; NO ₃ ⁻ formation	NOEC = 1000	Schwarz (2013)
Nitrification in activated sludge	NH ₄ ⁺ depletion	IC ₅₀ = 61 mg/L	

^aIn water

potential inhibition of these nitrifying soil bacteria. However, recognizing that the MDAS are reactive aromatic compounds, and nitrite, formed transiently during nitrification of ammonia, may react directly with the MDAS by forming azo-compounds or nitroso-aromatics. If such reactivity were to occur under conditions of the standard OECD 216 test, the resulting reduction in nitrate formation may be mis-interpreted as inhibition of nitrifying bacteria. In a screening study to examine potential for this reactivity, (unpublished data of BASF Polyurethanes, Gericke 2010) the stability of 4,4'-MDA was observed in an aquatic solutions at pH = 6 in the presence and absence of equimolar amounts of sodium nitrite; the decay of nitrite was not investigated. The solutions were incubated at about 22 °C in the dark. The results of this screening test, shown in Fig. 10, indicated an instability (reactivity) of 4,4'-MDA in the presence of nitrite under these slightly acidic conditions. In the presence of nitrite, the initially colorless 4,4'-MDA solution became yellow after 7 days.

With this knowledge, an OECD Guideline 216 study was performed in a modified way (Schwarz 2013), where instead of plant material the soil was spiked with about 75 mg ammonium sulfate per 5 g dry soil. Over the test period, both the decline of ammonium and the formation of nitrate were monitored. The soil was spiked with 0, 10, 20, 50, 125, 250, 500 or 1000 mg 4,4'-MDA per kg soil (d. w.). The EC₁₀ value was shown to be above 1000 mg/kg after 14 and 28 days in terms of ammonia depletion, as well as in terms of nitrate formation. Only after 7 days, the EC₁₀ was between 500 and 1000 mg/kg, indicating a potential minor and transient

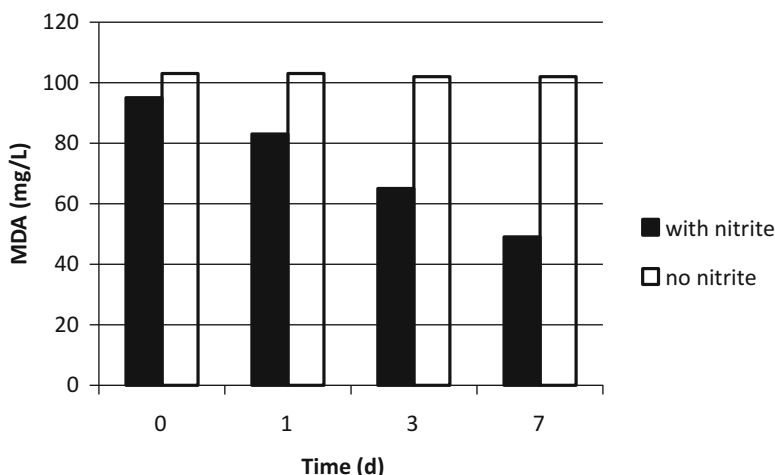


Fig. 10 Time-dependent concentrations of 4,4'-MDA in aqueous solution in the presence (*black bars*) and absence (*white bars*) of nitrite at pH = 6

effect on the soil bacteria. The amount of MDA measured in aqueous soil extracts taken for nitrate analyses was shown to rapidly decline and was about 14–60 % of the initial lowest and highest MDA concentrations tested, respectively, at the beginning of this test. At the end of the test (28 days), recoveries of the MDA were < 10 % of their initial nominal values. This result is in line with observations in the previously discussed sediment tests with MDA, where an irreversible absorption of MDA to soil organic matter along with biodegradation are the most likely explanation disappearance from the test systems. So, a limited inhibition of nitrification was observable only at the beginning at the highest soil load (1000 mg/kg d. w.), when some freely available MDA was likely still available. The quantitative formation of nitrate further indicates that transiently-formed nitrite was not significantly reacted with MDA, as was shown to be the case in pH 6 aqueous solution (Gericke 2010).

The ISO 9509 test (2006) can be used to check the nitrification inhibition in activated sludge. Initial cursory tests were inconclusive as 17 mg/L seemed to be the IC₅₀ concerning ammonium decay, but it was difficult to discern even a 10 % inhibition of nitrate formation (Schupp 2014). Therefore, a definitive test was performed with activated sludge of the municipal STP of the city Steinfurt, Germany. The sludge (3 g solids per liter, dry wt.) was spiked with ammonium sulfate and 0, 9.4, 18.7, 47 and 93.5 mg/L 4,4'-MDA (98 % pure). The N-Allyl-thiourea substance, a known inhibitor of nitrification, served as control. The nutrient solution contained 2.71 g/L NH₂SO₄ and 5.47 g/L NaHCO₃ and was diluted 1:10 for the test runs. Tests were run in duplicate. After 4 h incubation at 20 °C, samples were taken, filtered through a 0.46 µm membrane, and ammonium ions were analyzed photometrically after reaction with K₂HgI₄. A blank experiment proved that MDA does not interfere with this photometrical method. In this test, MDA shows moderate

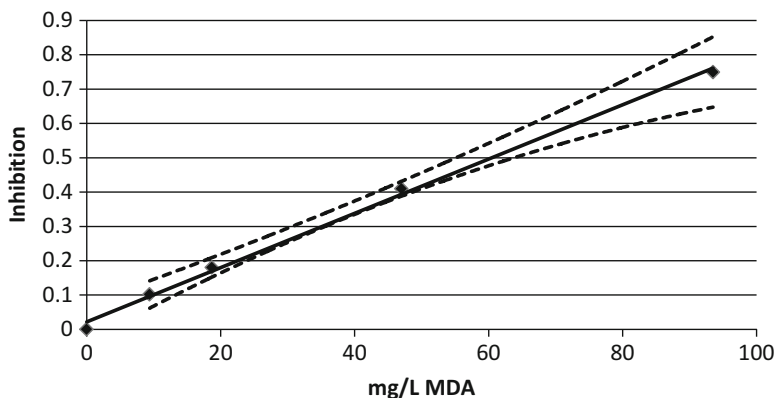


Fig. 11 Inhibition of ammonium nitrification by MDA in an ISO 9509 test

nitrification inhibition with $IC_{50} = 61$ mg/L (Fig. 11). HPLC-UV analysis of the 47 mg/L solution after the test showed that MDA was neither absorbed on sludge nor did it suffer other routes of decay.

4.3.2 Toxicity to Terrestrial Plants

The potential effect of MDA on the emergence and growth of monocot (oat, *Avena sativa*) and dicot (lettuce, *Lactuca sativa*) vascular plants was investigated according to OECD Guideline 208 (Van der Hoeven et al. 1992a). The purity of the test substance was 99.5 %. Oat was sown at 1 cm depth, and lettuce on the surface of a semi-natural soil. The soil was previously spiked with sand that was coated with MDA. For the latter, MDA was dissolved in acetone, the sand soaked with this MDA-solution, and the acetone was then evaporated under a stream of nitrogen. Tests were run for 17 days and five replicates at ten seeds per concentration (0, 3.2, 10, 32, 100, 320 and 1000 mg/kg d. w. soil, nominal concentrations). Short-term endpoints were associated with concentrations at which no effects were observed, and the lowest concentration at which effects were observed by pair-wise binomial test (2×2 contingency table).

For the emergence endpoint, concentrations were defined as 320 and 100 mg/kg d. w. at which no effects were observed for 17 days exposures to oat and lettuce, respectively. For growth, the corresponding concentrations showing no effects were 100 and 10 mg/kg d. w. for oat and lettuce, and the corresponding EC_{50} values were 353 and 128 mg/kg d. w., where these endpoint values are based on nominal concentrations. The dicotyledonous lettuce, therefore, is somewhat more sensitive to 4,4'-MDA than the monocotyledonous oat.

In another seed germination test, Kim et al. (2002) investigated the influence of 4,4'-MDA dissolved in water and poured onto filter paper in petri-dishes on the germination of radish seeds. The precise species, not mentioned in the paper,

probably was *Raphanus sativus*. After 6 days at 20 °C, germination was assessed visually. The reliability of this test may be questioned, as the authors claim to have used MDA solutions up to 1 % MDA (10 g/L) which is clearly beyond the water solubility of MDA (Table 2). Data were presented in graphs only, and germination is estimated to be 86, 48, 29, 14, 5 and 0 % at 0.01, 0.02, 0.06, 0.1, 0.5 and 1.0 % MDA in water, respectively. If the two highest concentrations are omitted due to potential insolubility of MDA, the 6 days EC₅₀ is interpreted to be 331 mg/L, and a 6 days EC₂₀ is 96 mg/L is estimated by logit analysis.

4.3.3 Toxicity to the Earthworm *Eisenia fetida*

Van der Hoeven et al. (1992b) investigated the acute toxicity of 4,4'-MDA against *E. fetida* after 14 days exposure according to OECD Guideline 207 (1984), using nominal test concentrations of 0, 18, 32, 56, 100, 320 and 560 mg/kg d. w. applied to five replicates per concentration. The 4,4'-MDA (99.5 % purity) was dissolved in acetone and added to artificial soil. Subsequently, the acetone was evaporated under a stream of nitrogen. The 14 days LC₅₀ was determined to be 444 mg/kg d. w.

Moser and Hamberge (2012) investigated the effect of 4,4'-MDA on the reproduction of *E. fetida* according to OECD Guideline 222 (2004) and in compliance with GLP. Stock solutions of MDA (purity 98 %) were prepared in ethanol, and the concentrations were verified by HPLC-UV. Defined volumes of the MDA solutions were added to quartz sand, and the ethanol was evaporated. The coated quartz sand was used incorporated into the artificial soil, containing 5 % sphagnum peat. Nominal concentrations were 18, 32, 56, 100, 180 and 320 mg/kg d. w., with four replicates used at each concentration and ten worms per replicate. Worms were fed with cow manure which was free of contamination and veterinary pharmaceuticals. At day 56, test vessels were analyzed. For the mortality endpoint, the 56 days NOEC was 180 mg/kg d. w., the 56 days EC₁₀ was 92.06 mg/kg d. w., and the 56 days EC₅₀ was 333 mg/kg d. w. soil. For adult biomass endpoint, the 56 days NOEC and 56 days LOEC values were 32 and 56 mg/kg d. w., respectively. For reproduction, the 56 days LOEC was 18 mg/kg d. w. As no NOEC (reproduction) could be derived the 56 days EC₁₀ of 11.2 mg/kg (95 % C.I.: 0.6–21.3 mg/kg d. w.) was statistically determined instead.

4.3.4 Toxicity to the Collembolan Species *Folsomia candida*

The effect after 4 weeks of exposure to 4,4'-MDA on the collembolan species *F. candida* was investigated by Moser and Schott (2011) according to OECD Guideline 232. The 4,4'-MDA test substance (98 % pure), dissolved in ethanol, was spiked to artificial soil containing 5 % sphagnum peat. Spiking solutions were analyzed by HPLC-UV. Nominal concentrations were 56.2, 100, 178, 316, 562 and 1000 mg/kg d. w. soil. For adult mortality, there was no clear dose–response; the highest mortality of 20 % was observed at 316 mg/kg; for reproduction, the 28 days

NOEC was 562 mg/kg and the 28 days LOEC was 1000 mg/kg d. w. The reference substance boric acid generated the expected results.

4.3.5 Acute Toxicity to Birds

Hurlbut et al. (1983) report an acute LD₅₀ of 148 mg/kg for redwinged blackbirds (*Agelaius phoeniceus*), fed with grains containing 4,4'-MDA. Data on purity of the substance and stability of MDA in/on the food are not provided, and many experimental details are lacking. The reliability of the data generated, therefore, is questionable.

4.4 Potential for Endocrine-Modulating Effects and Brief Overview of the Toxicity to Mammalia

A review of data on endocrine activity of MDA was issued by Jaeger and Collins (2011) and Jaeger et al. (2012). A summary of the reviewed findings is presented as follows:

In yeast receptor binding studies, 4,4'-MDA and pMDA showed neither estrogenic nor androgenic activity and no anti-estrogenic activity in the dose range of 10^{-7} to 0.1 mmol/L; however, 4,4'-MDA and pMDA showed apparent anti-androgenic activity at concentrations of 10^{-3} and 10^{-2} mmol/L, respectively (Kolle and Landsiedel 2010a, b, c, d). Since MDA has potential to react directly with the endogenous hormone (testosterone) and with proteins which are the construct of the androgen receptor, the apparent anti-androgenic activity associated with high MDA concentrations could be an artifact of this reactivity and not necessarily indicative of receptor-mediated antagonism.

In a subacute oral study, castrated female rats received 150 and 200 mg/kg MDA for up to 14 days (Tullner 1960). The body weight decreased whereas uterine, thyroid and adrenal weights increased. The adrenals showed extensive lipid accumulation, and thyroid follicles contained little or no colloid.

Intravenous administration of 50 and 100 mg/kg MDA caused a transient reduction in 17-hydroxycorticoid output in the dog (Tullner 1960).

In ovariectomized rabbits, subcutaneous administration of MDA caused a progestational response. However, in ovariectomized and adrenalectomized rabbits, MDA did not cause a response, whereas subcutaneous progesterone did. Cortisol did suppress the progestational response to MDA, but not that due to progesterone (Tullner 1960).

Rats received 0, 80, 400 or 800 ppm MDA in drinking water for 3 months (Ciba-Geigy 1982). Besides hematotoxic and hepatotoxic effects, thyroid follicular epithelial cells were hypertrophic, and the glandular structures showed a diffuse hypertrophy and colloid depletion. In another subchronic drinking water study

with rats, similar effects were reported (DHHS NTP 1983). In both studies, gonads and accessory organs were not affected.

In mice, MDA caused mainly hepatotoxicity; goiters with papillary hyperplasia and vacuolization of the colloid were observed in one high dose male and female animal (DHHS NTP 1983).

In chronic drinking water studies, MDA caused thyroid follicular cell adenomas and carcinomas in rats and mice (DHHS NTP 1983). This effect might be attributable to irreversible inhibition of thyroid peroxidase by the reactive MDA substance at high dose levels. In a later study, MDA was shown to be an effective inhibitor of hog thyroid peroxidase (TPO) in vitro with an IC₅₀ of 0.06 μ M (Freiberger 1994). However, direct genotoxicity of MDA in thyrocytes was demonstrated by a positive comet assay (Martelli et al. 2002).

5 Summary

Concerning chronic toxicity, *D. magna* is the most sensitive species tested against MDA aquatic exposures, with a 21 day-NOEC of 0.00525 mg/L. Exposure of daphnids takes place via the aquatic phase. Other species of the same phylum (*Arthropoda*) appear to be less sensitive albeit with exposures via soil or sediment, with a 28 days-NOEC of 562 mg/kg d. w. soil (*F. candida*) and 41.3 mg/kg d. w. sediment (*Hyalella azteca*), for reproductive and survival endpoints, respectively. Also for acute toxicity, *D. magna* is more sensitive than the other species, with an 48 h-EC₅₀ that spreads over two orders of magnitude, ranging from 0.019 to 2.7 mg/L; the reason for this large difference is not clear at the moment. Data laying at both ends of the range are from the same lab, testing the same charge of MDA, and the only difference visible so far is the feeding mode. That is, daily feeding with algae resulted in an 48 h EC₅₀ of 0.019–0.06 mg/L, whereas the 48 h EC₅₀ from the acute test with no feeding is 2.47 mg/L. Future tests might investigate whether intake of MDA via the food chain results in a significantly higher body burden in the daphnids than intake via water, only. Fish show a more uniform reaction to MDA, with 96 h-LC₅₀ values ranging from about 20 to 60 mg/L; chronic data for fish are not available. Acute toxicity data for algae and cyanobacteria are in the range of 1–10 mg/L; based on growth-rate, the 72 h-NOEC or EC₁₀ of MDA to algae is 0.3–9.3 mg/L.

For sediment organisms, the blackworm *L. variegatus* shows the highest sensitivity to MDA with NOEC values between ≤ 3.75 mg/kg and 30 mg/kg d. w., followed by the amphipod *H. azteca*. The higher sensitivity of *L. variegatus* in the second study compared to the first study is obviously attributable to the different feeding regimes (semi-continuous feeding against pre-spiked sediment). One argument might be that semi-continuous feeding allows the organisms to avoid the contaminated food. However, a change from semi-continuous feeding to sediment pre-spiked with nettle powder (*Urtica* sp.) results in an earlier and much stronger increase in ammonia concentration in the system. This became apparent after both

studies on *the blackworm* were finalized. The ammonia 96 h-EC₅₀ for the *blackworm* is 0.69 mg/L at pH = 8.2, and the 96 h-EC₁₀ at pH = 8.2 is 0.33 mg/L (Hickey and Vickers 1994). As a result, the lower NOEC and LOEC in the second study with *L. variegatus* are probably attributable to interference by ammonia.

MDA binds irreversibly to soil and sediment which may explain the general, but not uniform lower sensitivity of soil and sediment organisms against aquatic organisms. The intrinsic toxicity of MDAS, beyond that attributed to a baseline narcosis mode of action, is imparted by the reactivity of the primary aromatic amine groups. Thus, reaction of these amine groups with organic matter (macromolecules) would be expected to coincide with reduced reactivity, toxicity, and bio-availability. However, species with intense soil or sediment contact (*L. variegatus* and *E. fetida*) show in general lower NOEC values than those organisms with less direct contact (3.75 and 11 mg/kg d. w., respectively). On the one hand it may be hypothesized that this intense contact to soil-bound MDA is one reason for the higher sensitivity; on the other hand, metabolic capacity against MDA of the organisms tested is unknown at this point in time and might as well explain differences in species sensitivity. For plants there are only acute data available; for the terrestrial species tested, *L. sativa* is more sensitive to MDA than *E. fetida*.

Limited aquatic data available so far do not indicate that the toxicity of pMDA is different from that of MDA. In addition, the limited set of data generated with the marine *M. macrocopa* (crustacean, acute and chronic test), *N. fustulum* (diatom) and *V. fischeri* (bacteria) do not indicate that sea water organisms are more sensitive to MDA than fresh water organisms.

In mammals, MDA is unlikely to interact directly with the endocrine-mediated homeostatic, growth, and developmental pathways; interaction with the adrenergic system cannot be ruled out, and apparent effects of MDA on the thyroid hormone system have been demonstrated. MDA inhibits the thyroid peroxidase enzyme which might contribute to the thyroid gland tumors observed in chronic studies with rats and mice. Some anti-androgenic activity in vitro did not prevail in the in vivo studies with rats and mice. Main target organs in mammals are the liver and the thyroid gland. MDA shows genotoxicity in different test systems and is a carcinogen by GHS category 1b.

6 Conclusion

The toxicity of 4,4'-Methylenedianiline, its isomers and homologues (MDA substances; MDAS) against organisms in the environment was iteratively tested over at least four decades. This review underlines the importance of accurate substance identification and analyses that accompany ecotoxicity tests for substances which can react with or in environmental compartments. Especially for the sediment and soil compartment that absorb MDA reversibly and irreversibly, the influence of these processes on the biological availability and toxicity of MDA and its products

requires careful consideration. Sediment tests with MDA were performed at a time where the increase in knowledge concerning reactive substance was on a steep slope, and the guideline for *L. variegatus* was under development. This, together with the dynamics of the MDA molecule in the sediment, may explain variations in findings of the sediment tests. There is a remarkably high variation in toxicity values in acute daphnia studies. Whereas most values are in the range of 0.46–5.7 mg/L which represent the typical variation in ecotoxicology there is one study showing significant lower EC50 values in the range of 0.019–0.06 mg/L, thus indicating that the factors governing ecotoxicity of MDAS to daphnids is currently not fully understood.

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Conflict of interest: T. Schupp worked for BASF, a MDA-producer, until 2012; H. Allmendinger is a consultant for Currenta GmbH & Co. OHG; B.T.A. Bossuyt is working for Huntsman, a MDA-producer; B. Hidding is working for BASF, a MDA-producer; B. Tury is a consultant for International Isocyanates, Inc.; R.J. West is working for Dow Chemical Company, a MDA-producer.

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Cadmium Bioavailability, Uptake, Toxicity and Detoxification in Soil-Plant System

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1 Introduction

Cadmium (Cd) accumulation in plants and the subsequent trophic transfer along the food chain poses a serious threat to both ecosystem and human health (Zhang et al. 2014; Goix et al. 2014). Human exposure to Cd in excessive concentrations has the potential to cause cancer, bone lesions, lung insufficiency, cancer, teratogenic effects, renal disturbances, anemia, hypertension and weight loss (Gad 2014). Because of its potential to cause such adverse effects, Cd is classed as potential human carcinogen (group 2B) by the US Environmental Protection Agency (EPA), a human carcinogen (group 1) by the International Agency for Research on Cancer of the World Health Organization (WHO), and ranked No. 7 among the top 20 priority hazardous substances by the Agency for Toxic Substances and Disease Registry (ATSDR 2012). The main route of Cd exposure to humans is through the diet (Chaney 2012). This is well exemplified by the *Itai-Itai* (ouch ouch) disease outbreak in Japan, which caused softening of the bones and kidney failure in those affected from the consumption of Cd contaminated rice (Li et al. 2011). Because of the potential health risks, Cd contamination in rice has been of mounting concern in China (Fang et al. 2014). The use of Cd has been restricted in many European countries owing to its high potential public health impacts.

Although Cd is extremely rare in the Earth's crust, various anthropogenic activities (e.g., Zn mining and smelting, application of sewage sludge/compost as well as fertilizer, pesticide and insecticide applications) (Rao and Kashifuddin 2012) have resulted in elevated Cd-soil concentrations. Cadmium is easily taken up by plants because of its high mobility within the soil-plant system (Nedjimi and Daoud 2009; Gill et al. 2012). Cadmium phyto-uptake is affected by a number of factors related to soil properties (soil particle size, pH, temperature, cation exchange capacity) and plant physiology (root surface area, rate of root exudation and transpiration) (Rosén et al. 2012). The majority of plant species accumulate Cd in root tissues, with only very small portions being transferred to the shoots (Podazza et al. 2012). However, high Cd levels in aerial plant parts are observed in Cd hyperaccumulators (Douchiche et al. 2012; Huguet et al. 2012).

Excessive Cd accumulation in plant tissue has the potential to result in numerous morphological, physiological, and biochemical toxic effects. Cadmium is not known to participate in any vital process in living organisms. Because of its capacity to elicit adverse effects, several literature reviews have been devoted to different aspects of Cd phytotoxicity (Page et al. 1987; Holmgren et al. 1993; Kabata-Pendias 1993; Toppi et al. 2012; Gallego et al. 2012). Toxicity of Cd to plants is generally associated with disruption of water and nutrient uptake and transport (Kabata-Pendias 1993; Zhang et al. 2010; Douchiche et al. 2012), alteration of nitrogen metabolism (Dguimi et al. 2009), disruption of ATPase activity (Lin et al. 2012), reduced photosynthesis, plant growth and respiration, dysfunction of the plant photosynthetic machinery in chloroplasts and stomatal closure (Hasan et al. 2011). Other symptoms of Cd toxicity may include reduced plant length and browning of roots, leaf rolling, chlorosis, and necrosis (Najeeb et al. 2011).

At the cellular level, excessive Cd concentrations can cause chromosomal aberrations and alteration of cell cycles and division (Inglot et al. 2012; Filipič 2012) as well as enhanced production of reactive oxygen species (ROS) (Lin et al. 2012; Gill et al. 2013). Overproduction of ROS may cause cell death due to oxidative processes such as DNA and RNA damage, enzyme inhibition, protein oxidation and membrane lipid peroxidation (Tian et al. 2011; Shahid et al. 2013a, 2014a). To minimize Cd-induced ROS formation, plants possess several defense strategies, including sequestration of the metal within vacuoles and chelation by organic molecules (Hossain et al. 2012). As a secondary defense mechanism, plants can also activate antioxidants to fight against the overproduction of Cd-induced ROS (Chen et al. 2012; El-Boshy et al. 2014). Several genes have been suggested to play a role in Cd detoxification and/or transport (Fässler et al. 2011).

The basic concepts of Cd bioavailability, uptake, toxicity and detoxification in soil-plant system have been well documented in studies carried out and published before 2000. These earlier findings/facts have been summarized in several books and review articles (e.g., Page et al. 1987; Holmgren et al. 1993; Kabata-Pendias 1993; Das et al. 1997). During the last decade, several articles have additionally been produced which focus on the biogeochemical behavior of Cd in the soil-plant system, highlighting the mechanisms involved at the cellular level. The present review summarizes the latest data (published after 2000) on Cd bioavailability in soil, plant uptake and translocation, toxicity, and detoxification – aspects which are highly relevant to soil remediation and risk assessment. The review is divided into eight sections: (1) introduction; (2) global use of Cd; (3) behavior of Cd in soil; (4) soil-plant transfer of Cd; (5) Cd phytotoxicity; (6) Cd detoxification mechanisms; (7) hermetic effects, and; (8) conclusions and perspectives.

2 Global Use of Cadmium

Cadmium is a naturally occurring element in the Earth's crust and waters. Cadmium ore is not mined for production because more than enough is produced as a byproduct from mining, smelting, and refining of zinc sulphide ore (Rao et al. 2010). For this reason, global Cd production depends more on Zn refining than marketplace demand. The proportion of Cd and Zn varies from mine to mine, ranging from 0.07 % to 0.83 % with an average value of 0.23 % (OECD 1994). Recycling of steel and iron scrap is the secondary source for Cd production (about 10–15 % of consumption) (OECD 1994).

Despite its numerous industrial, agricultural and domestic uses, Cd was not widely utilized until World War-I (Rao and Kashifuddin 2012). Owing to its useful properties, Cd is now used in several industrial activities. Today, the major global consumption of Cd (almost 80 %) can be attributed to its use in rechargeable nickel–Cd batteries (USGS 2015). However, nickel–Cd batteries are gradually being phased out and replaced by nickel metal hydride batteries. Cadmium was often used in the steel industry for protecting steel from corrosion by Cd electroplating. Cadmium also has a number of other uses, including the preparation of alloys (due to its good fatigue resistance and a low coefficient of friction) and as a stabilizer for plastics (Scoullou et al. 2001; Nzengue et al. 2015). Cadmium telluride thin-film photovoltaics are used as a substitute to the traditional silicon-based solar cells (USGS 2015), and are preferred for large-scale, ground-mounted utility systems and commercial rooftop applications (USGS 2015).

Between 2001 and 2003, Cd production decreased from 18,700 to 15,000 thousand metric tons, but increased to 18,800 thousand metric tons in 2004. After 2004, production of recoverable Cd increased gradually, reaching 22,200 thousand metric tons in 2014 (USGS 2015; Fig. 1), with China, Korea and Japan being the major producers (USGS 2015). Given that Cd-production in mining increased overall by 15 % between 2001 and 2014 alone (Fig. 1), it is very important to understand the biogeochemical behavior of Cd, and the impact that increased Cd concentrations will have on the environment.

3 Behavior of Cd in Soil

3.1 Cd Level and Sources in Soil

Cadmium was first discovered in Germany simultaneously by Stromeyer and Hermann in 1817 as a by-product of the Zn refining process. Approximately 700 times less abundant than Zn, Cd ranks as the 65th most abundant element in the Earth's crust (Emsley 2011), with a lithosphere concentration of approximately 0.1–0.2 mg/kg (Heinrichs et al. 1980). While Cd can be found together with the Zn ore spalerite, greenockite (CdS) is the only known Cd-based ore (Rao et al. 2010).

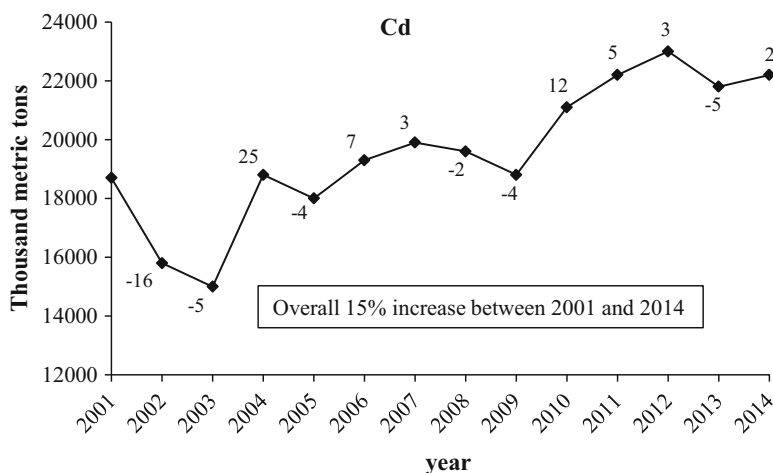


Fig. 1 Annual world mine production of Cd in thousand metric tons (source, [USGS 2015](#)). The *plus and minus values/labels* in the graph line indicate the % increase and decrease of Cd production. Despite restriction on the use of Cd in many European countries owing to its high potential public health impacts, the production of Cd has increased by 17% between 2001 and 2013

The natural concentration of Cd in soil is between 0.07 to 1.1 mg/kg in many parts of the world (Thornton 1992). While natural concentrations vary with soil type, higher Cd concentrations can be expected for soils overlying sedimentary rocks (which contain the highest Cd concentrations ranging between 0.3 to ≥ 15 mg/kg) (Traina 1999), and soils impacted by Cd-phosphate fertilizers (Liu et al. 2012). Except for soil and coal, concentrations of Cd for different constituents of the Earth's crust (Table 1) generally do not exceed 0.6 mg/kg. Cadmium concentrations higher than 10 mg Cd/kg soil can be toxic to plants (Holmgren et al. 1993).

According to an extensive survey carried out in the United States with 3305 soil samples collected from agricultural areas in 36 states, Cd soil levels varied from 0.005 to 2.4 mg/kg, with mean values of 0.27 mg/kg (Holmgren et al. 1993). According to FOREGS, total Cd concentrations in European agricultural soil range between 0.06 to 0.6 mg/kg (Salminen et al. 2005). Similarly, concentrations ranged from < 0.1 to 2.9 mg Cd/kg for surface soils collected from 237 farms in Ohio (Logan and Miller 1983), and 0.06 to 0.74 mg/kg for 16 soil series in Minnesota (Pierce et al. 1982). Burau et al. (1973) reported that Cd concentrations ranged from 0.05 to 10.1 mg/kg for 177 agricultural soils developed from Monterey shale deposits in Salinas Valley, California. The variation in Cd concentrations among different soils reflects in part the concentration differences in the parent material from which these soils are formed. To minimize the risk of elevated Cd concentrations in edible produce and Cd-induced phytotoxicity, it is recommended that soils containing more than 1 mg Cd/kg dry soil not be used for agricultural or horticultural purposes (Louwagie et al. 2009), and that remedial action be taken for

Table 1 Cadmium ranges/level in different constituents of Earth's crust

Constituents of Earth's crust	Cd level (mg/kg)	References
Lithosphere	0.1–0.2	Heinrichs et al. (1980)
Ultramafic rocks	0.03–0.05	Mesjasz-Przybyłowics et al. (2004)
Mafic rocks	0.18	Ivanov (1996)
Intermediate rocks	0.13	Ivanov (1996)
Clay, clay shales	0.3	Ivanov (1996)
Sandstones	0.01–0.41	Page et al. (1987)
Limestones	0.001–0.5	Page et al. (1987)
Granite	0.001–0.6	Page et al. (1987)
Basalt	0.006–0.6	Page et al. (1987)
Coal	1.0	Tauber (1988)
Subsoil	0.09	FOREGS
Topsoil	0.15	FOREGS
Soil	0.07–1.1	Thornton (1992)

soils containing 5–20 mg/kg Cd (Eikmann et al. 1993). Unfortunately, several previous studies have shown high Cd contamination in agricultural soils where food has been grown (Zhao et al. 2015).

Volcanoes and weathering of rocks are the major natural sources of Cd mobilisation from the Earth's crust (Choppala et al. 2014; Zhao et al. 2015). Collectively, natural sources are estimated to release about 150–2600 tonnes of Cd to air (Nordic Council of Ministers 2003). In 1983, about 140–1500 tonnes of Cd was emitted to the atmosphere from volcanoes (Nriagu 1989).

Relative to natural processes such as volcanic activity and the weathering of rocks, anthropogenic activities such as mining, combustion of fossil fuels, metal production and other industrial and agricultural practices have been responsible for the increased Cd levels in the environment (Choppala et al. 2014; Agnieszka et al. 2014; Zhao et al. 2015). Industrial emission, animal manures, sewage sludge and phosphate fertilizers are reported to have increased Cd concentrations in agricultural soils (Rao and Kashifuddin 2012). Munshower (1977), for example, reported high Cd levels (29 mg/kg) in Montana up to two km northeast of a Pb-Zn smelter, which decreased with distance and reached near background levels after 24 km. Recently, Shahid et al. (2013b) reported high levels of Cd (70.2 mg/kg) within a 40 km² area surrounding the Metaleurop smelter near Evin-Malmaison (North of France).

Although atmospheric deposition has decreased during the last two to three decades in developed countries, it remains a major source of Cd release (e.g., 2500–15,000 tonnes of Cd annually) to agricultural soils (Shahid et al. 2013b; UNEP 2010). The use of wastewater for crop irrigation without any prior treatment in peri-urban areas is another major source of Cd for agricultural soil. Singh et al. (2010) reported 0.02 mg/l of Cd in wastewater of Varanasi, India. Despite low levels of Cd in wastewater, sorption of Cd by the soil may result in Cd

accumulation over time. Soil amendments such as municipal sewage sludge (MSS) may also contain high levels of Cd (e.g., 2–3500 mg Cd/kg sludge) (Logan and Chaney 1984; Page et al. 1987). It is estimated that the application of MSS @ 5 tons/ha, for MSS containing 10–15 mg/kg Cd, can lead to an increase in the surface soil Cd concentration of 10–15 %. Landfills and various deposits of discarded products further release about 7500–29,500 tonnes of Cd per year to soil (UNEP 2010), as do agrochemicals used for enhanced crop production (e.g., 8.9 g ha⁻¹ year⁻¹ from phosphorus fertilizers; Bramley 1990). Among the fertilizers, those that are phosphate-based and manufactured from seabed sediments with high Cd content have the potential to contain the highest Cd concentrations (Louwagie et al. 2009). In various countries of Europe (Denmark, Ireland, Austria, Finland, Greece and the United Kingdom), Cd is estimated to have increased by 7–43 % as a result of the past 100 years of fertilizer application (UNEP 2010).

3.2 Bioavailability of Cd in Soil

In order to estimate the ecological risk associated with Cd contamination in soil, it is necessary to understand the fraction of total Cd available for uptake by biota. Total Cd concentrations in soil do not necessarily reflect Cd phytoavailability (Duplay et al. 2014) because different chemical forms (chemical speciation) and the varying distribution of Cd (adsorbed or free) affect its phytoavailability. Cadmium is relatively available for plant uptake because it is predominantly found in soil bound to the exchangeable solid phases (Degryse et al. 2012) and thus readily released into the soil solution. In the soil solution, Cd is mainly present as the Cd²⁺ ion, or as inorganic or organic complexes. In the solid phase, Cd may be reversibly bound to soil particulates, such as organic matter or Fe and Mn oxides. Plant uptake of Cd occurs mainly through contact with porewater, which is the net result of Cd partitioning between solid and liquid phases of soil. However, recent studies have also reported metal uptake by plants via foliar transfer (Uzu et al. 2010; Schreck et al. 2012a, b). According to Xiong et al. (2014a), foliar uptake of Cd and Pb due to particulate matter deposition is of particular importance near recycling factories.

The chemical forms and partitioning of Cd in soil is governed by a number of reactions that involve precipitation/dissolution, adsorption/desorption, and Cd–ligand complex formation (Bolan et al. 2013). These dynamic processes are strongly affected by variations in redox conditions (Zhang et al. 2012), cation exchange capacity (Jiang et al. 2012), soil pH (Saeki and Kunito 2012), soil texture (Qi et al. 2012), biological and microbial conditions (Joubert et al. 2007), metal burdens (Gao et al. 2009), organic and inorganic ligands (Niazi et al. 2011a; Shahid et al. 2014b; Austruy et al. 2014), competing cations (Helios-Rybacka and Wójcik 2012) and temperature (Silber et al. 2012) (Fig. 2).

The chemical speciation of Cd, and its partitioning in soil, is a key factor in controlling both the fate and toxicity of Cd in the soil-plant system (Pizzol et al. 2012; Landrot et al. 2012). It is well-known that total soil Cd concentrations

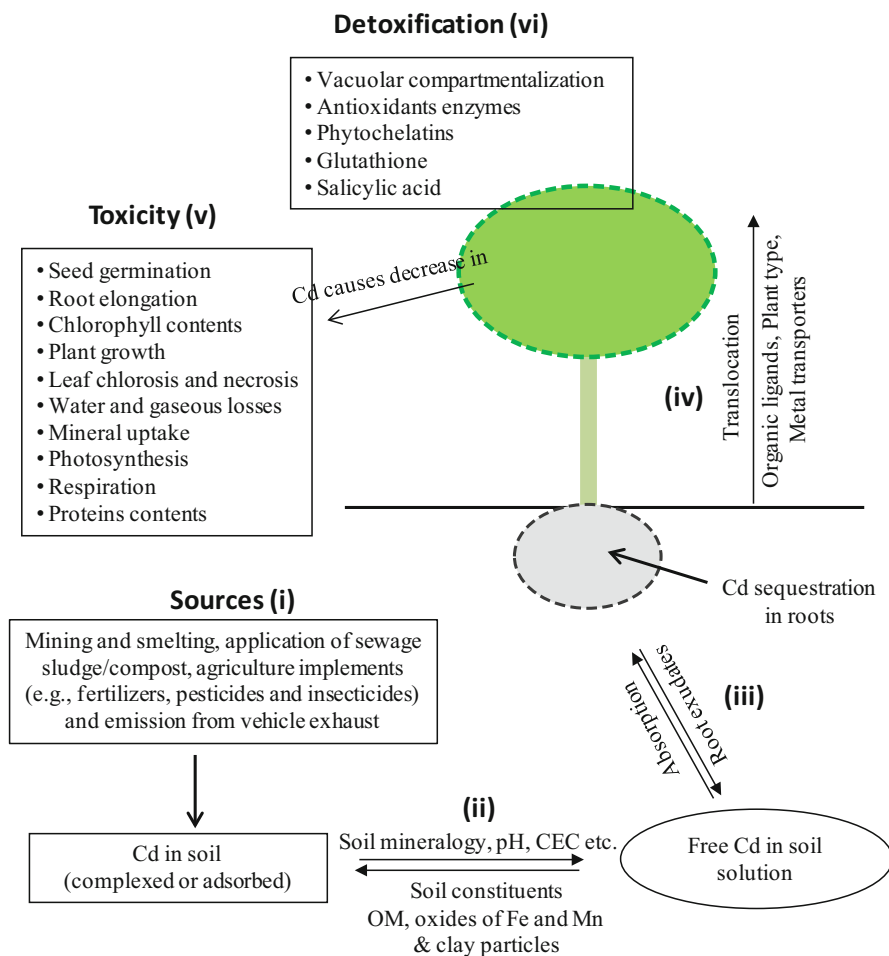


Fig. 2 Biogeochemical behavior of Cd in soil-plant system. (i) Accumulation of Cd in soil from different natural and anthropogenic sources, (ii) effect of soil physic-chemical properties on the complexation, mobility and bioavailability of Cd in soil, (iii) Cd uptake by plants via root transfer and major sequestration (>90 %) in root tissues, (iv) translocation to aerial parts depending on the nature of plant and applied Cd level and (v) toxic effects of Cd to plants and (VI) detoxification mechanisms involving the role of antioxidants, phytochelatins, glutathione and salicylic acid

tend to be a poor proxy for predicting Cd mobility, solubility, extractability, and bioavailability (Udovic and McBride 2012). This is because the total sum of all chemical forms of Cd are rarely 100 % bioavailable (Udovic and McBride 2012), and the biogeochemical behavior of Cd depends largely on the free Cd concentration (Dabrin et al. 2012). For these reasons, knowledge of Cd speciation is considered necessary for informing soil remediation and risk assessment studies (Niazi et al. 2011b; Shahid et al. 2011, 2012a; Bade et al. 2012).

Traditionally, analytical techniques used to assess metal speciation in soil and water have included anodic stripping voltammetry, ion selective electrodes, competitive ligand equilibration/adsorptive stripping cathodic voltammetry, and sorption onto C_{18} columns (Cobelo-García et al. 2005). However, these techniques are very complex and time consuming. When actual measurements are not feasible, computer speciation models (capable of performing complex calculations) offer a cost-effective alternative (Shahid et al. 2012b, 2014b), although site-specific validation is required to confirm the accuracy of the model's output. The most widely used speciation models include: the Windermere Humic Aqueous Model VI (WHAM VI), CHESS, CHEAQS, PHREEQC and VISUAL MINTEQ (Tipping et al. 1998; Parkhurst and Appelo 1999; Gustafsson, 2008; Antunes and Kreager 2014; Shahid et al. 2014d). These models offer the advantage of desk-top computing and are appropriate for both small and large scale applications. However, the accuracy of the estimated speciation may vary among models and chosen input values (e.g., Cd concentration range, DOC concentration and quality, consideration of precipitation reactions).

3.3 Effect of Soil Chemical Properties on Cd Bioavailability

3.3.1 Effect of Soil pH on Cd Bioavailability

Soil pH is one of the most important parameters governing Cd speciation, partitioning, and bioavailability (Ardestani and van Gestel 2013), as evidenced by the negative correlation established between soil pH and Cd mobility/phytoavailability (Bilodeau-Gauthier et al. 2011). Cadmium is reported to exist in different chemical forms under different soil pH conditions. In most soils, 99 % of the total Cd is associated with soil colloids. Cadmium may occur as cationic species: $CdHS^+$, $CdOH^+$, $CdHCO_3^+$, $CdCl^+$, and as anionic species: $Cd(HS)_4^{2-}$, $Cd(OH)_3^-$, $Cd(OH)_4^{2-}$, $CdCl_3^-$ (Kabata-Pendias and Sadurski 2004). Of the Cd in soil solution, it has been estimated that between 55 and 99 % exists in free ionic form (Kabata-Pendias 1993). At low pH, Cd in soil solution is predominately present as Cd^{2+} , $CdSO_4$ or $CdCl^+$, whereas in alkaline solutions the less bioavailable $CdHCO_3^+$, $CdCO_3$ or $CdSO_4$ species predominate (Sauvé et al. 2000). Cadmium solubility greatly depends on the acidity of the soil solution. For example in mineral soils having a pH range of 4.0–4.5, a decrease in pH of merely 0.2 units results in a 3–5 times increase in the labile Cd pool. This is in contrast to an increase in pH within the basic pH range, which results in a greater extent of Cd adsorption on to soil particles with a subsequent reduction in plant uptake of Cd, Zn, and Pb (Kuo et al. 1984). Douay et al. (2009) and Fritsch et al. (2010) reported that the spatial distribution of extractable and total Cd levels in 262 soil samples over an area of 40 km² in Northern France greatly varied in accordance with soil pH and other soil properties. Because of the relationship between pH and Cd bioavailability, one strategy used for Cd phytoremediation has been to lower soil pH to enhance

plant Cd uptake (Shahid et al. 2011). In contrast, for both plant and groundwater protection, increasing soil pH is likely to be an effective measure to reduce phytoavailability, since Cd precipitation and adsorption on soil particles should minimize soil solution Cd concentrations (Sauvé et al. 2000; Qi et al. 2012). Although impacts of soil pH on the formation of Cd precipitates vary with soil type, high redox potential and low soil pH have generally been shown to restrict the formation of Cd-precipitates (Li et al. 2011). In acidic soils, Cd solubility is mainly controlled by soil organic matter and oxides and hydroxides of Fe and Mn (Jiao et al. 2012). Above pH 7.5, sorbed and precipitated Cd (e.g., as CdCO_3 , and possibly $\text{Cd}_3(\text{PO}_4)_2$) is not readily mobilized (Kabata-Pendias 2011). In calcareous soils where Cd-carbonate complexes predominate (Renella et al. 2004), a large proportion (e.g., 35 %) of Cd may additionally be present as Cd^{2+} and CdHCO_3^+ (Hirsch and Banin, 1990).

Under conditions of low soil pH, Cd bioavailability and mobility is higher due to Cd transformation from immobile forms such as Fe and Mn oxides and carbonates to more bioavailable and exchangeable forms (Xian and Shokohifard 1989; Li et al. 2014). Soil pH may decrease or increase Cd bioavailability in soil indirectly via transformation of Fe and Mn minerals. Iron oxides have a significant capacity to sorb and immobilize Cd (Liu et al. 2014a, b). Thus, secondary Fe minerals formed by Fe(III) reducing bacteria and the Fe bacteriogenic oxides formed by Fe(II)-oxidizing microorganisms can reduce Cd bioavailability (Cortés-Martínez et al. 2004; Muehe et al. 2013), whereas the reduction of Cd bearing Fe(III) minerals by Fe(III)-reducing bacteria can increase Cd bioavailability (Muehe et al. 2013). Different mechanisms have been proposed for the negative correlation between soil pH and Cd dissolution/desorption. Soil pH governs the degree of net charge associated with the solid-phase: the higher the pH the more negatively charged the solid-phase surface and vice versa. Under acidic soil conditions, H^+ ions compete with Cd for binding sites, which results in Cd desorption from soil particles into solution. In contrast, under alkaline conditions, the solid-phase exchange sites are readily available for metal cation binding. Additionally, at high pH levels, Cd is adsorbed more strongly by the solid phase due to hydrolysis of Cd to hydroxy species (Barrow 1986) and higher specific adsorption of Cd (He et al. 2001), thus, making metal desorption more difficult at high pH (Tudoreanu and Phillips 2004).

3.3.2 Effect of Soil Organic Matter on Cd Bioavailability

Soil organic matter, originating from the decomposition of microbial, plant and animal material, plays an important role in governing Cd mobility/bioavailability (Quenea et al. 2009) as it readily forms complexes with Cd. The effect of humic substances (HSs) on metal bioavailability varies with its form in soil (i.e., either suspended or dissolved), as well as with the concentration, source, and physico-chemical quality (e.g., molar mass, humification stage, types of functional groups) (Cabaniss et al. 2000; Calace and Petronio 2004). Humic substances have been

reported to bind Cd^{2+} to a greater extent than the major inorganic ligands, especially at high pH values (Reuter and Perdue 1977). Compared to the humic acid component of HSs, fulvic acids are more reactive due to their smaller size, higher oxygen content, and higher cation exchange capacity (Zeng et al. 2002; Shahid et al. 2012d). Soils with a higher organic matter content are more effectively able to reduce Cd uptake by plants as a result of Cd-sorption (onto carboxylic and phenolic hydroxyl groups (Dumat et al. 2006; Shahid et al. 2012d)) and subsequent removal of Cd from the soil solution.

Since dissolved HSs form soluble complexes with Cd, soil amendments with OM may also enhance Cd mobility and uptake by plants (Almås and Singh 2001). This may in part be why some authors have reported a decrease in metal bioavailability in the presence of HS (Guerrero et al. 2000), while others have shown an increase (Sánchez-Marín et al. 2010). In addition to these direct impacts, OM can also influence Cd bioavailability by altering the soils' physico-chemical properties such as CEC, pH, particle size distribution, porosity and cracking pattern, microbial and enzyme activities, and soil solution composition (Turrión et al. 2012).

3.4 Influence of Soil Microbial Activity on Cd Behavior in Soil

Soil microbial activity is known to increase Cd availability via excretion of organic acids and subsequent solubilization of Cd-bearing minerals (Wang et al. 2009; Liu et al. 2014a, b; Ahmad et al. 2015). Soil amendment with Cd-solubilizing microorganisms such as plant growth-promoting bacteria (PGPR) is a very effective technique for enhancing Cd bioavailability (Belimov et al. 2005). Xiong et al. (2008) for example, observed significant increases in the uptake of Cd, Zn, and Pb by *Sedum alfredii* Hance when soils were inoculated with rhizosphere microorganisms. Similarly, the capacity of *Thlaspi caerulescens* to extract Cd, Pb and Zn has been found to increase following microbial inoculation of the soil (Epelde et al. 2010). This finding supports the work of Duponnois et al. (2006), who showed that inoculation of *Sorghum bicolor* with Cd-tolerant bacteria significantly increased Cd uptake, and also the work of Sheng and Xia (2006), who observed that soil inoculation with isolates containing bacteria caused a 74 % increase in the Cd content of *Brassica napus*. Arbuscular mycorrhizal fungi appear to play a similar role, as suggested by the results of Long et al. (2010), who found that inoculation of *Perilla frutescens*, *Litsea cubeba*, *Dysphania ambrosioides*, *Phytolacca Americana* and *Rehmannia glutinosa* resulted in enhanced uptake of Cd, Zn, Cu and Pb.

In contrast, some soil-inoculated microorganisms, such as *Azospirillum lipoferum*, *Arthrobacter mysorens*, *Agrobacterium radiobacter* (Belimov et al. 2004), and arbuscular mycorrhizae (Karagiannidis and Nikolaou 2000) have been shown to decrease Cd phytoavailability. Recently, Dary et al. (2010) reported

a decrease in the uptake of Cu, Cd and Pb by *Lupinus luteus* when inoculated with *Brady rhizobium*. The inoculation of Cd-polluted soil with *Ralstonia eutropha* has also been shown to reduce Cd uptake by *Nicotiana bentaminana* as a result of Cd immobilization (Valls et al. 2000). Tiwari et al. (2008) found that 11 bacterial strains of *Typha latifolia* reduced Cd bioavailability in fly ash. Similarly, *Pseudomonas aeruginosa* was also reported to decrease Cd uptake by >50 % in *Cucurbita maxima* in Cd-contaminated soil (Sinha and Mukherjee 2008). The decrease in bioavailability of Cd by soil microorganisms might be due to Cd accumulation by the microbes themselves, or Cd immobilization in the soil. The latter hypothesis is supported by the work of Xu et al. (2012), who found that bacteria transformed 8–25 % of Cd to a less bioavailable organic-bound fraction.

4 Soil-Plant Transfer of Cd

4.1 Molecular Understanding of Cd Uptake by Plants

Although Cd is a non-essential element, its phytoaccumulation factor index (ratio of Cd level in plant to soil) is high, and may surpass that of many essential nutrients (Gonçalves et al. 2009). Cadmium does not possess any specific cell entry pathway. Cadmium enters the plants from soils mainly via root uptake in the form of accidental transport by specific and non-specific transporters of essential elements (Llugany et al. 2012), such as Fe^{2+} , Ca^{2+} , Zn^{2+} , Cu^{2+} and Mg^{2+} (Roth et al. 2006; Mendoza-Cózatl et al. 2011). To date, no Cd-specific transporters have been unambiguously identified.

Transcriptional and post-transcriptional gene regulations responsible for encoding specific and non-specific transporters are complex. ZIP (ZRT, IRT-like protein) family transporter for Fe^{2+} , Zn^{2+} and Mn^{2+} have been reported to mediate Cd uptake in plant roots (Mendoza-Cózatl et al. 2011). It has been demonstrated that Cd enters root cells via ZIP transporters or through cation channels (Verbruggen et al. 2009; Gallego et al. 2012). OsIRTs is also suggested to contribute to Cd uptake by plants (Nakanishi et al. 2006; Lee and An 2009). The role of OsIRTs in Cd uptake varies with plant type. For example, OsIRTs has been found to have no effect on Cd uptake by Prayon ecotype (*T. caerulescens*), which is less capable of accumulating Cd (Lombi et al. 2002). Cadmium can also enter plant root cells by Yellow-Stripe 1-Like proteins as Cd-chelates (Curie et al. 2009) and NRAMP (natural resistance-associated macrophage protein) family transporters. Similarly, TcZNT1 has been shown to mediate low-affinity Cd uptake in *T. caerulescens* (Pence et al. 2000), and a low affinity cation transporter (LCT1) cDNA of *Triticum aestivum* has been reported to enhance Cd uptake in yeast cells (Clemens et al. 1998). It has been proposed that LCT1 functions as a Ca transport system in plants and might contribute to Cd transport (Clemens et al. 1998). Despite these advances in our understanding, the mechanisms/processes involved in Cd uptake by plants are not yet well understood.

4.2 Competition Between Cd and Cations for Uptake

Although further research is still needed, we know that metal-metal interactions influence Cd uptake. The presence of La^{3+} , Ca^{2+} , Cu^{2+} , Fe^{2+} , Zn^{2+} or Mn^{2+} in the rhizosphere solution, for example, have been shown to inhibit Cd uptake by plant roots (Lux et al. 2011). These cations likely affect Cd uptake as a result of competition for ion transport specific channels (Wojas et al. 2009). This hypothesis is supported by the work of Li et al. (2012) who showed that pre-treatment with the Ca^{2+} channel blockers LaCl_3 and verapamil significantly reduced Cd^{2+} influx into roots. Although selenium would not be expected to compete with Cd for uptake in the same manner as cationic metals, Hu et al. (2014) similarly reported reduced uptake (44 %) of Cd by *Oryza sativa* in the presence of selenium.

Ion-specific interactions of trace metals with Cd, and the subsequent impact on plant-Cd uptake, vary with plant genotype and the type and concentration of metals present in the soil solution (or rhizosphere) (Lugon-Moulin et al. 2004). Cadmium and Zn, for example, have been found to behave alike in soil- water- plant system owing to their chemical similarities (Mengel et al. 2001). The similarities between these two metals result in a protective effect of Zn against Cd uptake, whereby plants generally absorb more Cd when the soil solution concentration of Zn is low (Mengel et al. 2001). Owing to this antagonistic relationship, Cd concentrations in plant roots have been shown to decrease by up to 50 % in the presence of Zn (Lugon-Moulin et al. 2004). Thus, in the context of food chain transfer, agricultural soils or waters containing moderate concentrations of Zn are likely to afford some protection against crop contamination by Cd.

Cadmium in the rhizosphere or soil solution can interfere with the uptake of essential nutrients by plants (Castillo-Michel et al. 2009). This interaction is of public concern (Gonçalves et al. 2009), as Cd can cause severe nutrient deficiencies in plants (Dong et al. 2006). This in turn can result in physiological disorders and a diminution of growth, with specific impacts differing among plant species, cultivar, and plant tissues (Liu et al. 2003). Using energy-dispersive X-ray spectroscopy Vaculík et al. (2012) revealed that Cd disturbed the distribution of essential elements in plant tissues. They reported that Ca, Mg, Si, Na and Zn accumulated mainly in the peripheral bark, S and K in the phloem, and Cd in both vascular tissues. Jiang et al. (2004) reported that Cd reduced the uptake of K, P, Fe, Zn and Ca in the roots, and K, P, Cu and Ca in the shoots of *Brassica juncea*. In a study with *Brassica campestris*, Zhu et al. (2004) reported that Cd exposure resulted in an increase in the K concentration in the roots and decrease in the shoots, with the opposite trend being observed for P, Mg, B, Fe and Mn. For Zn, Ca and S, concentrations increased in both roots and shoots. Nedjimi and Daoud (2009) reported that the addition of Cd to the culture medium decreased both root and shoot concentrations of Ca and K in *Atriplex halimus*. Sun and Shen (2007) also reported a decrease in Fe, Mn, P, Mg and S concentrations under Cd stress in *Brassica oleracea*. Cherif et al. (2011) further reported a decrease in Zn uptake by

Solanum lycopersicum in the presence of Cd. Based on the available data, no definitive conclusion can be drawn regarding the impact of Cd on essential nutrients.

4.3 Cadmium Sequestration in Plant Roots

Subcellular accumulation of Cd in plants varies greatly not only among plant species but also among cultivars and genotypes of the same species (Xin et al. 2013), and the plants' Cd detoxification and tolerance mechanisms. For most plants, Cd tends to accumulate in the roots, with only a small portion being translocated to aerial parts (Ahmad et al. 2011). Generally, Cd contents in plants decrease in order of roots > stems > leaves > fruits > seeds (Benavides et al. 2005). Some species can sequester high concentrations of Cd in their roots with reduced translocation to shoots, or remobilize Cd from leaves to roots through the phloem (Mendoza-Cózatl et al. 2011). These are referred to as low Cd-accumulating plants. Recently, Zhou et al. (2015) reported 60–90 % accumulation of Cd in roots of 32 hybrid rice (*O. sativa*) cultivars—a percentage similar to that observed by Gonneau et al. (2014) (i.e., 50–80 %) for Cd accumulation in shoots of 22 populations of *Noccaea caerulea*. The concentration of Cd in plant roots can be tenfold higher than in the shoot (Wang et al. 2009), suggesting that roots serve as an effective barrier to Cd translocation. The efficacy of the root cell walls in trapping Cd is attributed to the number of available functional groups (e.g., hydroxyl, carboxyl) that are able to form complexes with trace metals (Nishizono et al. 1987). For example in *Arabidopsis thaliana*, Cd was mainly bound to O/N groups in the cell wall and only a minor fraction was bound to S-containing groups (Isaure et al. 2006).

Cadmium accumulation in plant roots varies with plant type and rhizosphere Cd concentration. For example, more Cd accumulates in the cell-wall fraction of *Zea mays* than *Vicia faba* seedlings (Lozano-Rodriguez et al. 1997). Wu et al. (2005) showed that 51 % of the Cd existed as a soluble form and 36 % was present in cell walls in barley roots. Similarly, Cd is mainly concentrated in the apical cells of plant roots at low soil Cd concentration or in the proximal subapical region (Seregin et al. 2004). Cadmium mainly accumulates in nuclei and vacuoles, while a lesser amount of Cd may also be found in the plastids and cytoplasm of plant roots (Liu et al. 2007). Toppi et al. (2012) reported that Cd was concentrated in cell walls and intercellular spaces, as well as in vacuoles of cortical root cells of *Daucus carota*. Electro-dense aggregates seen using transmission electron microscopy have been found between the plasmalemma and cell wall in the outermost tissues of root, whilst very few are present in the vascular cylinder (Lux et al. 2011; Toppi et al. 2012).

4.4 Cadmium Translocation to Shoots

Cadmium movement from roots to aerial tissues occurs via transpiration-driven xylem loading, with symplastic and/or apoplastic transport (Hasan et al. 2009; Kranner and Colville 2011) of free Cd^{2+} , or Cd complexed with various chelates (Maestri et al. 2010; Lux et al. 2011). In the studies of Zhao et al. (2006), Cd shoot concentration decreased in response to a decrease in transpiration. The important role of transpiration in governing Cd translocation has been reiterated by Kranner and Colville 2011, who found that the seed Cd content differed for two rice cultivars having different transpiration rates. The process of Cd translocation to the shoot tissues is also known to occur very quickly, as evidenced by the considerable amount of Cd observed in xylem sap and shoot tissue of *O. sativa* only 1 h after Cd treatment with positron emitting radio isotopes (Fujimaki et al. 2010). Thus, Cd is translocated with water flowing through the vascular system to the leafy tissues, where water then evaporates and Cd is left to accumulate (Krzesłowska et al. 2010).

The proportion of absorbed Cd in the plant aerial parts varies according to plant characteristics (species, age, cultivar) (Foucault et al. 2013; Koren et al. 2013; Zhou et al. 2015; Van der Vliet et al. 2007). For instance, Cd translocation from roots to shoots was lower in *Solanum torvum* compared to *Solanum melongena* (Yamaguchi et al. 2011; Lux et al. 2011). Similarly, *Lupinus albus* has been found to accumulate less Cd in its shoots than *Lupinus angustifolius* and *L. luteus* (Egle et al. 2003). Pectin-rich layers of the collenchyma cell walls of the leaf veins have been reported to be the main storage sites of Cd in *Salix viminalis* (Vollenweider et al. 2006). In *A. halleri*, Cd accumulated mainly in leaf veins and trichomes (Isaure et al. 2006). Cadmium was mainly stored in the stem pitch and cortex of hyperaccumulating ecotype of *S. alfredii*, while in the non-accumulating ecotype, Cd was mainly localised in vascular bundles (Tian et al. 2011). Wójcik et al. (2005) reported that *N. caerulescens* accumulated Cd in vacuoles of spongy and palisade mesophyll cells.

The exact role of different chelators in Cd translocation has not yet been elucidated. Organic ligands such as citrate and malate are known to form weak complexes with Cd (attributed to their low binding affinities), especially at cytosolic pH values (Rascio and Navari-Izzo 2011). These and other chelating compounds are involved not only in metal hyperaccumulation and tolerance (Haydon and Cobbett, 2007), but also in overcoming metal diffusion limitations which may otherwise restrict Cd uptake (Antunes and Hale, 2006). Amino acids, such as histidine and nicotinamine, similarly play key roles in Cd translocation owing to their ability to form stable complexes with Cd and other divalent cations (Rascio and Navari-Izzo 2011).

Xylem loading plays an important role in Cd accumulation within aerial plant parts. Low Cd loading into the xylem sap in the roots has been associated with subsequent low Cd transport to aerial parts (Hasan et al. 2009; Verbruggen et al. 2009; Santos-Echeandía et al. 2010). A number of important membrane

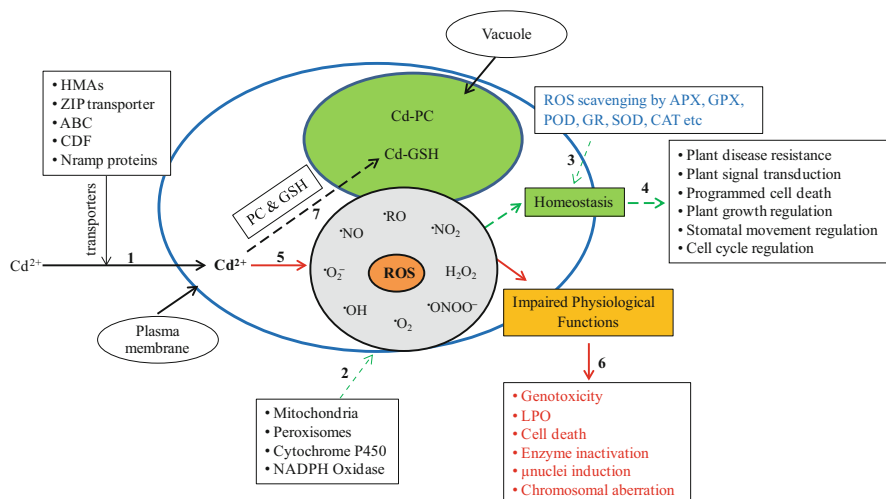


Fig. 3 Schematic representation showing possible Cd entrance, accumulation toxicity and detoxification in plant cell. Cadmium entrance across plasma membrane is mediated by membrane transporter gene families (1). Under normal conditions, ROS are produced in different cell organelles (2) and an optimal ROS level (homeostasis) is maintained by antioxidant enzymes (3). These ROS are involved in different essential roles in living organisms (4). Presence of Cd may result in increased production of ROS (5), which may lead to dramatic physiological challenges to the plant termed as “oxidative stress” (6). Alternatively, Cd may be sequestered into vacuole by PC and GSH (7). *Green arrows* indicate toxic effects of Cd whereas *green arrows* indicate Cd detoxification. HMAs, heavy metal ATPases; ZIP, ZRT and IRT-like protein; ABC, ATP binding cassette; CDF, cation diffusion facilitator; NRAMP, natural resistance associated macrophage protein; PCs, phytochelatin; GSH, glutathione; ROS, reactive oxygen species; APX, ascorbate peroxidase; GPX, guaiacol peroxidase; POD, Peroxidase; GR, glutathione reductase; SOD, superoxide dismutase; CAT, catalase; LPO, lipid peroxidation

transporter gene families, identified and characterized in recent years, are thought to be involved in Cd loading into the xylem sap and Cd transport. These gene families include the ATP binding cassette (ABC) superfamily, HMA (heavy metal ATPase), ZIP (ZRT, IRT-like protein), NRAMP (natural resistance-associated macrophage protein), YSL (yellow-stripe-like transporter), NAS (nicotinamine synthase), SAMS (S-adenosyl-methionine synthetase), FER (ferritin Fe (III) binding), CDF (cation diffusion facilitator), NRT (nitrate transporter) and IREG (iron regulated transporter) (Memon and Schröder 2009; Maestri et al. 2010) (Fig. 3).

Similarly, several studies have reported the involvement of P1B-ATPase and also ZIP family transporters in Cd transfer across the plasmalemma into the shoots (Hanikenne et al. 2008; Wong and Cobbett 2009), using energy provided by ATP hydrolysis to transport Cd against its electrochemical gradient. The IRT1 gene, which is transcriptionally responsive to Fe deficiency, can also transport Cd to plant shoot tissues (Cohen et al. 1998), and genes from the NRAMP family are thought to play a role in Cd accumulation in Arabidopsis seedlings (Thomine et al. 2000). Certain members of the Cation Diffusion Facilitator (CDF) family function in Cd

efflux or vacuolar uptake (Williams et al. 2000), while OsHMA3 has been reported to regulate xylem Cd transport by regulating vacuolar sequestration of Cd in root cells (Miyadate et al. 2011). Over-expression of HMA3 has been found to improve Cd tolerance and accumulation in *Wassilewskija* ecotype (Morel et al. 2009) and the gene is further thought to be responsible for differences in Cd accumulation in grains of two rice varieties (Ueno et al. 2010). Owing to its role of xylem Cd loading and transportation, HMA3 has been reported to play a key role of Cd hyperaccumulation in *Arabidopsis* and *N. caerulescens*. However, it is not yet well known how these metal transporters are regulated at the transcriptional level. Given the involvement of HMA3 in Cd hyperaccumulation, genetic research focusing on HMA3 may lead to new approaches to enhancing Cd accumulation and tolerance of non-hyperaccumulating plants.

Hyperaccumulators are plants that accumulate large metal concentrations in above-ground tissues (Arshad et al. 2008; Shahid et al. 2012a), have a phytoaccumulation factor index (ratio of Cd level in plant to soil) >1 and a translocation factor index (ratio of Cd level in plant shoot to roots) >1 , have a high metal tolerance, and are fast growing with good biomass production. Hyperaccumulators can accumulate high levels of Cd in aerial tissues partly as a result of enhanced expression of metal transporter genes (Verbruggen et al. 2009; Santos-Echeandía et al. 2010). During the last three decades, the identification of metal hyperaccumulation traits for phytoremediation has been the focus of intensive research (Niazi et al. 2011c, 2012; Llugany et al. 2012; Sabir et al. 2015). To date, more than 500 hyperaccumulating species (belonging to 45 families) have been reported worldwide, yet the number that tolerate and accumulate Cd specifically remains largely unknown (Zhou and Song 2004). So far, identified Cd hyperaccumulators include *T. caerulescens* (Baker et al. 1994), *Arabidopsis halleri* (Dahmani-Muller et al. 2001), *Solanum nigrum* (Xiao et al. 2010), *S. alfredii* (Sun et al. 2007), *Arabis paniculata* (Qiu et al. 2008), *S. nigrum* (Sun et al. 2008), *Lonicera japonica* (Liu et al. 2009b), *Bidens pilosa* (Sun et al. 2009), *Arthrocnemum macrostachyum* (Redondo-Gómez et al. 2010) and *Viola baoshanensis* (Wu et al. 2010). These plants can solubilize soil Cd through root exudates (Redondo-Gómez et al. 2010), thereby enhancing Cd uptake and translocation.

Increased accumulation of Cd especially in aerial parts of hyperaccumulator plants is highly useful for the remediation of Cd-polluted sites. Hyperaccumulators generally have well-established detoxification mechanisms and can therefore accumulate high levels of Cd without showing any physio-morphological toxicity symptoms. Some hyperaccumulators are native to areas with high soil metal concentration, and thus grow well in soils with high Cd levels. On the other hand, plants that do not efficiently cope with Cd and other pollutants are termed as 'sensitive'. These plants do not possess well-established detoxification mechanisms and are often used in risk assessment studies. In fact, metal exposure to these sensitive plants (such as *V. faba*), even at low levels, can induce several toxic effects resulting in reduced plant growth (Shahid et al. 2014d).

4.5 Cadmium Accumulation in Edible Plant Parts

Cadmium translocation to plant parts that are animal feed and/or food products (fruit) may pose serious health concerns. Therefore, there is increasing concern regarding food safety due to environmental pollution by Cd, especially since the *Itai-Itai* disease was reported in the late 1950s (Hagin and Kono 1955; Tang et al. 2016). For populations not subjected to a Cd-contaminated environment, consumption of Cd contaminated food (vegetables, cereals, pulses and legumes) is the main source of Cd exposure in animals and humans (Järup and Akesson 2009; Mombo et al. 2016). In the United States (US), the consumption of contaminated food is the largest source of Cd intake in children and non-smoking adults (Tavarez et al. 2015). It is estimated that in the US, the average Cd consumption via contaminated food (primarily cereal grain and vegetables) is between 8 and 30 μg of Cd per day per person (Järup and Akesson 2009; Schwartz and Reis 2000; Clemens et al. 2013). Human exposure to excess Cd via contaminated food causes severe harm to organs such as kidney, liver and lungs, which elevates the risk of developing cancer (Järup and Akesson 2009; Shahid et al. 2014c; Baldantoni et al. 2016). For these reasons, Cd uptake and accumulation in edible plants and its possible effects on human health have received considerable attention (Garate et al. 1993; Akoumianakis et al. 2008; Mombo et al. 2016).

Since Cd is toxic to humans at lower concentrations than plants, plants that appear healthy are not necessarily safe for human consumption. To decrease the potential risks associated with excessive Cd intake via consumption of contaminated food, maximum allowable values of Cd in edible plant parts have been established by the Codex Alimentarius Commission of Food and Agriculture Organization of World Health Organization (FAO/WHO, CODEX 2006) (Table 2). Threshold values of Cd in edible plant parts vary with food type (cereal, legumes or

Table 2 Threshold values of Cd in edible plant parts established by the Codex Alimentarius Commission of FAO/WHO (CODEX 2006)

Food	Threshold values (mg/kg)	Remarks
Cereals, Pulses and Legumes	0.1 ^a	Excluding Bran and Germ, Wheat Grain, Rice, Soybeans and Peanuts
Wheat Grain and Rice	0.2 ^b	Including Bran and Germ
Soybeans and Peanuts	0.2 ^b	
Vegetable, including potatoes (edible part)	0.5 ^b	Excluding Leafy Vegetables, Fresh Herbs, Stem and Root Vegetables, Fungi, Tomatoes and Peeled Potatoes
Peeled Potatoes, Stem and Root Vegetables	0.1 ^b	Excluding Celeriac
Leafy Vegetables, Fresh Herbs, Fungi and Celeriac	0.2 ^b	

^aIndicates guideline level

^bIndicates maximum level

vegetables) as well as the edible/consumed plant part (grain, root, stem or leaf) (Table 2). The tolerable limit of Cd intake by humans is $< 70 \mu\text{g day}^{-1}$ (FAO/WHO 1978), and the provisional tolerable weekly intake is $7 \mu\text{g/kg}$ body weight. Several recent studies have reported higher accumulation of Cd in edible plant parts than the maximum allowable values set by the Codex Alimentarius Commission—thus indicating a potential human health threat (Table 2). Among plants consumed by humans, mushrooms, rice, lettuce and other leafy vegetables are reported to be the main source of Cd exposure (Table 3). Recently, Schlecht and Ina (2015) found that 54 % of mushrooms collected from different locations near high traffic areas in Berlin-Germany had Cd levels that exceeded the threshold values. However, it must be kept in mind that Cd uptake, translocation and compartmentation (in edible or other plant parts) in crops vary greatly among cultivars within the same species (Simmons et al. 2005; Marmiroli et al. 2014). For example Yu et al. (2006) reported high variation of Cd accumulation in rice grains ($0.30\text{--}2.19 \text{ mg kg}^{-1}$) among 43 cultivars when grown on Cd contaminated soil. Shi et al. (2015) also showed variation in Cd concentration ($0.14\text{--}0.22 \text{ mg kg}^{-1}$) in rice grains among 12 cultivars. Therefore, in addition to crop type, cultivar and species type must be considered in risk assessment studies and in setting food consumption guidelines.

4.6 Foliar Uptake of Cd

Nowadays, metal recycling and production processes release a large proportion of fine and ultra-fine metallic particulate matter (PM) containing heavy metals into the atmosphere due to the use of thinner and more effective filters (Schreck et al. 2014). Since these fine particles containing Cd and other heavy metals are readily transported over long distances, biomonitoring studies are currently being used to assess metal pollution via atmospheric deposition. Health risks associated with atmospheric Cd pollution arise mainly from inhalation of particles and consumption of polluted vegetables (Morman and Plumlee 2013). Although root-metal uptake is considered the main uptake pathway for plants, foliar uptake has also been identified as a potentially significant route of Cd exposure (Shahid et al. 2013b; Schreck et al. 2014; Xiong et al. 2014a, b; Uzu et al. 2010). Recently, Schreck et al. (2013) suggested that root and shoot metal transfer pathways are independent, with an additive effect in terms of toxicity. The idea of foliar metal uptake grew in the 1950s subsequent to the comment by Michigan State University that “a leaf is a very efficient organ of absorption”. However, unlike Cd uptake by roots, which has been widely studied (Vaculík et al. 2012), comparatively few studies have investigated the uptake of atmospherically deposited metals by leaves (Uzu et al. 2010).

While foliar uptake of Cd is generally negligible, under certain situations (especially near mines), foliar uptake should be considered (Schreck et al. 2012a). As accumulation is roughly a function of the deposited amount of pollutants, the exposure time and compound effects of climatic factors are of prime importance. In experimental work using labelled Cd in the soil to distinguish between soil and

Table 3 Cadmium concentrations in edible plant parts of different vegetables and crops, reported in studies published between 2011 and 2015

Crop	Cd exposure level (mg/kg)	Cd in edible plant part (mg/kg)	Medium	References
<i>Lettuce</i>	0.8	0.73	Soil	Mombo et al. (2016)
<i>Celery</i>	0.8	0.4	Soil	Mombo et al. (2016)
<i>Celeriac</i>	0.8	0.7	Soil	Mombo et al. (2016)
<i>Chard</i>	0.6	0.4	Soil	Mombo et al. (2016)
<i>Lactuca sativa</i>	42	9	Soil	Baldantoni et al. (2016)
<i>Cichorium endivia</i>	42	4.5	Soil	Baldantoni et al. (2016)
<i>Lettuce</i>	1.87	3.62	Soil	Tang et al. (2016)
<i>Cacao beans</i>	2.6	3	Soil	Chavez et al. (2015)
<i>Solanum lycopersicum</i>	100 µM	13	Soil	Hédiji et al. (2015)
<i>Mushrooms</i>	5.23	2.88	Soil	Liu et al. (2015)
<i>Alchemilla xanthochlora</i>	16.3	8	Soil	Quezada-Hinojosa et al. (2015)
<i>Cynosurus cristatus</i>	16.3	9	Soil	Quezada-Hinojosa et al. (2015)
<i>Hypericum maculatum</i>	16.3	3	Soil	Quezada-Hinojosa et al. (2015)
<i>Wild edible mushrooms</i>	NM	10	Soil	Schlecht and Saumel (2015)
<i>Wheat</i>	13.7	0.22	Soil	Shi et al. (2015)
<i>Brassica napus</i>	0.3	6	Soil	Wu et al. (2015)
<i>Brown rice</i>	0.1 mmol/L	3.4	Hydroponics	Xue et al. (2014)
<i>Lettuce</i>	NM	1.8	Foliar	Xiong et al. (2014)
<i>Radish</i>	NM	1.6	Foliar	Xiong et al. (2014)
<i>Parsley</i>	NM	0.9	Foliar	Xiong et al. (2014)
<i>Brassica campestris L.</i>	1.59–5.18	1.08	Soil	Mahmood and Malik (2014)
<i>Brassica rapa L.</i>	1.59–5.18	0.36	Soil	Mahmood and Malik (2014)
<i>Spinacia oleracea L.</i>	1.59–5.18	1.9	Soil	Mahmood and Malik (2014)
<i>Green vegetables</i>	0.52–1.61	1	Soil	Chen et al. (2014a, b)
<i>Garlic chives</i>	0.52–1.61	0.5	Soil	Chen et al. (2014a, b)
<i>Spinach</i>	0.52–1.61	1.1	Soil	Chen et al. (2014a, b)
<i>Chinese cabbage</i>	0.52–1.61	0.55	Soil	Chen et al. (2014a, b)
<i>Garden pea</i>	0.52–1.61	0.55	Soil	Chen et al. (2014a, b)
<i>Rice grains</i>	6 mg/cm ³	7	Soil	Pereira et al. (2011)
<i>Lettuce</i>	6 mg/cm ³	130	Soil	Pereira et al. (2011)

NM not mentioned

airborne Cd in several crops, the the impact of airborne Cd was not negligible, even in rural areas (Hovmand et al. 1983). Harrison and Chirgawi (1989) used growth cabinets with filtered air and demonstrated that the atmospheric contribution to the contamination of vegetables was as large as 23 % for Cd.

While more research is needed to further elucidate the mechanisms involved in metal foliar transfer (Zangi and Filella 2012), the two major mechanisms involved in foliar Cd uptake are thought to include (1) deposition and internalization through the lipophilic cuticle and (2) penetration through stomata openings (Schreck et al. 2012a, Barber et al. 2004). According to Chamel et al. (1991), penetration through the cuticle would consist of four steps: sorption in the cuticle, diffusion through the cuticle, desorption in the apoplast and absorption by the subjacent cells. After this penetration phase, active transport within cells (symplastic pathway), transport in the vascular system (phloem or xylem) and transport between the phloem and xylem intervene. Metal uptake via root or shoot transfer may have different effects on metal localisation in different plant tissues, metal speciation (Sarret et al. 2013), and phytotoxicity. Moreover, plant response (detoxification and tolerance mechanisms) may also vary with the type of Cd exposure (root or shoot). Additional studies are required to compare the two pathways of Cd accumulation (i.e., root or shoot), and subsequent speciation, localization, toxicity and detoxification in plant tissues—especially near industrial and mining areas.

The processes of foliar metal uptake and cuticular penetration strictly depend on a number of factors, such as the physicochemical properties of the compound and cuticle, the surface area and morphology of the plant leaf, surface texture (roughness and pubescence), plant habitus (evergreen or deciduous), the physical and chemical forms of the deposited element, duration of exposure, gas exchange and environmental conditions (Uzu et al. 2010; McLaughlin et al. 2011; Calderón-Preciado et al. 2013). The adsorption of Cd on plant foliage depends on its concentration in air as particulate matter (PM). Climatic conditions also influence the efficacy of foliar Cd transfer through direct effects on the physico-chemical properties of the leaf surface, and by affecting biological processes in the plant, both over the short- and long-term. The immediate conditions of light, temperature and humidity at the time of foliar Cd application affect plant metabolic status and hence may directly influence absorption processes across the leaf surface and from within the internal leaf spaces. Schreck et al. (2012a) found metal accumulation in lettuce shoots to be linked to precipitation (correlation coefficient: 0.83), with high metal accumulation occurring during periods of greatest rainfall. This was possibly due to enlargement of the cuticle, pores, and stomata openings (Schreck et al. 2012a) which would have facilitated leaf penetration by the metal. However, climatic processes such as rainfall and wind can also remove Cd from shoot surfaces (Rodrigo et al. 1999; Staelens et al. 2008). In the case of vegetables, washing of samples prior to analysis resulted in a 25 % decrease in foliar metal concentration (Schreck et al. 2012a).

5 Toxic Effects of Cd on Plants

The processes related to Cd toxicity in plants are complex (Bandyopadhyay and Mukherjee 2011). Cadmium accumulation in plants, even at low levels, adversely affects plant growth and morphology (Hasan et al. 2009; Saidi et al. 2014; Shahid et al. 2015b). Above the toxic threshold, Cd can negatively affect morphological, physiological, and biochemical functions (Benavides et al. 2005; Deng et al. 2014; Sergeant et al. 2014). Cadmium may cause plant death when its level increases above 5–10 mg Cd g⁻¹ leaf dry weight (Gill et al. 2013). Data summarized in Table 4 (published after 2010) shows toxic effects of Cd in different plant species under various applied levels of Cd. The key symptoms of Cd-induced toxicity in plants are chlorosis, inhibition of photosynthesis, stunted growth, inactivation of enzymes in CO₂ fixation, root elongation, altered chloroplast ultrastructure, impaired seedling development, cell division and transpiration, leaf epinasty, lipid peroxidation, and disturbance of sulfur and nitrogen metabolism (Das et al. 1998; Benavides et al. 2005; Gill et al. 2013; Ran et al. 2014).

Cadmium has also been reported to reduce the uptake and transport of essential nutrients (phosphorus, calcium, magnesium and potassium) (Das et al. 1998), and to interfere with nitrate uptake and translocation to aerial parts (Hernandez et al. 1996) via the reduction of nitrate reductase activity (Mathys 1975). Cadmium toxicity can influence plasma membrane permeability by inducing lipid peroxidation (Fodor et al. 1995). At the cellular level, Cd causes increased generation of reactive oxygen species (ROS), interrupts cell redox status, deteriorates biological macromolecules (lipids, proteins and nucleic acids), dismantles membranes, and cleaves DNA-strands (Shahid et al. 2014f). The Cd-induced changes in biomembranes alter their permeability to protons, nutrient and water, and also affect the activity of membrane-bound enzymes (Belkadhi et al. 2015).

5.1 Effects of Cd on Seed Germination and Plant Growth

Although the reason is not yet well understood, one of the major consequences of the cumulative impacts of Cd exposure is a decrease in seed germination. This has been reported in Cd-exposed *Lycopersicon esculentum*, *Cucumis sativus*, *Lactuca sativa*, *D. carota*, *O. sativa*, *Z. mays*, *Hordeum vulgare*, *Pinus halepensis*, *Elsholtzia argyi*, *Spartiana alterniflora* and *T. aestivum* (Wang et al. 2012; Irfan et al. 2014; Ehsan et al. 2014).

Since Cd has a tendency to preferentially accumulate in the roots (Gill et al. 2012) it is not surprising that root growth and anatomy (Lux et al. 2010) are two characteristics adversely affected by Cd exposure. At the biochemical level, Cd has been shown to induce root cell wall lignification (Vaculík et al. 2012), premature lignification and xylogenesis (Ďurčáková et al. 2007), root browning, and reduced root length (Cherif et al. 2011; Wang et al. 2012) (Table 4). However,

Table 4 Toxic effects of Cd on plant growth. Data is collected from manuscripts published between 2010 and 2015

Cd exposure level	Duration (days)	Cd toxicity	Plant species	Growth medium	References
500 μM	7	Decreased plant biomass and photo-synthetic pigments	<i>Gossypium hirsutum</i>	Hydroponics	Daud et al. (2015)
100, 500 μM	8, 13, 25	Decreased plant growth, chlorophyll, nitrates and iron content	<i>Nicotiana tabacum</i>	Hydroponics	Iannone et al. (2015)
200 mg/kg	30	Reduced Rubisco activity, chlorophyll contents, gas exchange parameters, leaf nitrogen content and plant growth	<i>Triticum aestivum</i>	Soil	Khan et al. (2015)
1000 μM	45	caused electrolyte leakage, decreased plant biomass, relative water content, free proline, total soluble sugars and starch concentrations	<i>Triticum aestivum</i>	Hydroponics	Rady and Hemida (2015)
50 μM	1 h	Decreased quantum yields of PSII and protein contents	<i>Populus nigra</i>	Hydroponics	Lomaglio et al. (2015)
50 μM	14	Plant height, root length leaf area, and number of leaves	<i>Brassica napus</i>	Hydroponics	Ehsan et al. (2014)
1 mM	7, 14	Root and shoot length, leaf area per plant, and plant dry mass, number of pods per pot, pod yield per pot, and pod protein	<i>Phaseolus vulgaris</i>	Hydroponics	Howladar et al. (2014)
20 μM	4	Root and leaf fresh weight, chlorophyll and carotenoid content	<i>Helianthus annuus</i>	Hydroponics	Saidi et al. (2014)
120, 240, 480 μM	30, 60	Plant length, fresh mass, dry root and shoot mass, and leaf area	<i>Brassica juncea</i>	Soil	Irfan et al. (2014)
50 mM	30	Rubisco activity and net photosynthesis	<i>Brassica juncea</i>	Hydroponics	Asgher et al. (2014)

(continued)

Table 4 (continued)

Cd exposure level	Duration (days)	Cd toxicity	Plant species	Growth medium	References
5, 10, 20, 40 μM	7	Root mass, leaf mass, total mass, and leaf width.	<i>Ceratopteris pteridoides</i>	Hydroponics	Deng et al. (2014)
20 μM	56	Plant growth. Necrosis of youngest leaves was also observed.	<i>Populus tremula</i>	Hydroponics	Sergeant et al. (2014)
100, 500 mM	15	Fresh and dry biomass, root diameter, root surface area, number of root tips, and root volume.	<i>Brassica napus</i>	Hydroponics	Ali et al. (2013)
500, 1000 μM	56	Shoot length, leaf area per plant, and fresh and dry plant biomass.	<i>Triticum aestivum</i>	Hydroponics	Agami and Mohamed (2013)
1, 5 μM	35	Plant height, root length, number of leaves per plant, and leaf area.	<i>Gossypium hirsutum</i>	Hydroponics	Farooq et al. (2013)
30 $\mu\text{mol/L}$	7	Root and shoot length, and root and shoot biomass.	<i>Hibiscus cannabinus</i>	Hydroponics	Feng-tao et al. (2013)
15, 29, 43, 58 μM	190	Shoot and root length, fresh and dry mass, leaf area, chlorophyll content, and photosynthesis-related traits.	<i>Solanum lycopersicum</i>	Hydroponics	Hayat et al. (2012)
50 or 200 mM	7	Shoot length, root and shoot fresh weight, and chlorophyll a and b content. Chlorosis was also observed.	<i>Brassica juncea</i>	Hydroponics	Mohamed et al. (2012)
50–100 μM	15	Chlorophyll and carotenoid levels.	<i>Arabidopsis thaliana</i>	Hydroponics	Martínez-Peñalver et al. (2012)
36 μM	1, 2, 3, 4, 7, 14	Increased lateral root formation, cell-to-cell separation events leading to schizogenous oil duct formation.	<i>Daucus carota</i>	Hydroponics	Toppi et al. (2012)

(continued)

Table 4 (continued)

Cd exposure level	Duration (days)	Cd toxicity	Plant species	Growth medium	References
120, 241, 483 μM	30	Plant growth, net photosynthetic rate, stomatal conductance, intercellular CO_2 , chlorophyll content.	<i>Lepidium sativum</i>	Hydroponics	Gill et al. (2012)
10 μM	27	chlorophyll content, xylem-vessel number, and diameter of mature leaves. Plant death, water and gaseous losses in mature leaves were also observed.	<i>Myriophyllum alterniflorum</i>	Hydroponics	Delmail et al. (2011)
10 $\mu\text{mol/L}$	7	Phytosynthetic pigmentation.	<i>Solanum lycopersicum</i>	Hydroponics	Cherif et al. (2011)
483, 966 μM	10	Root and shoot fresh weight.	<i>Brassica juncea</i>	Hydroponics	Ahmad et al. (2011)
15, 29, 43, 58 μM	60	Chlorophyll content, transpiration rate, stomatal conductance, and internal CO_2 concentration.	<i>Solanum lycopersicum</i>	Hydroponics	Hasan et al. (2011)
10, 25, 50 μM	14	Root and shoot fresh and dry weight, biomass, growth rate, and chlorophyll content.	<i>Tagetes patula</i>	Hydroponics	Liu et al. (2011)
0–97 μM	7	Photosynthetic pigmentation, chlorophyll a and b, total chlorophyll ratios.	<i>Groenlandia densa</i>	Hydroponics	Yılmaz and Parlak (2011)
200–500 μM	14	Fresh and dry biomass.	<i>Juncus effusus</i>	Hydroponics	Najeeb et al. (2011)
0.5 mM	1, 3, 7	Plant growth	<i>Solanum lycopersicum</i>	Hydroponics	Monteiro et al. (2011)
400 μM	14	Root elongation and plasma membrane integrity.	<i>Sedum alfredii</i>	Hydroponics	Tian et al. (2011)
0–30 μM	45	Root and shoot growth.	<i>Solanum nigrum</i>	Hydroponics	Fidalgo et al. (2011)
5 μM	1, 5, 10, 15, 25	Plant height, root length, and biomass. Leaf necrosis and root browning were also observed.	<i>Hordeum vulgare</i>	Hydroponics	Chen et al. (2010)

(continued)

Table 4 (continued)

Cd exposure level	Duration (days)	Cd toxicity	Plant species	Growth medium	References
50, 100, 200 μ M	28	Plant growth, plant height, number of tillers, water content, and chlorophyll content.	<i>Achnatherum inebrians</i>	Hydroponics	Zhang et al. (2010)

most studies carried out regarding risk assessment and Cd toxicity to plants after 2010 were carried out using a hydroponic (as opposed to soil) medium (Table 4).

At lower exposure levels (e.g., 0.5 μ M), Cd has been reported to enhance the production of root hairs proximal to the root apex in various plants (Kopitke et al. 2010). However, the hindrance of root hair production has also been reported at high exposure levels (e.g., 0.5 mM) (Gratão et al. 2009). Increased root diameter can also result from Cd exposure (Maksimović et al. 2007; Lux et al. 2011), possibly attributed to an increase in parenchyma cell size, which enhances the resistance to radial flows of water and solutes (Maksimović et al. 2007).

In addition to effects on roots, Cd exposure is also known to negatively affect plant growth and biomass (Rady 2011) (Table 4). Based on the results of light microscopy work with *D. carota*, Corguinha et al. (2012) proposed that Cd affects plant growth in two separate phases, with the first phase impacting the functional and cytological events (1–4 days of Cd exposure), and the second phase impacting cell to cell separation, cell hypertrophy, and thiol-peptide content (4–14 days of Cd exposure). Cadmium is also known to inhibit photosynthesis by a variety of means including: reducing the transcription of genes which are related photosynthesis such as *rbcL*, *psaB* and *psbA* (Qian et al. 2010); inactivating Rubisco and phosphoenol pyruvate carboxylase (Bashir et al. 2013); inducing lipid peroxidation (Iannone et al. 2010), interfering with protochlorophyllide production (Prasad 2004), reducing the activity of enzymes involved in CO₂ fixation and inhibiting chlorophyll biosynthesis (Stobart et al. 1985); enhancing proteolysis (Pena et al. 2007); interfering with leaf transpiration through changes in stomatal conductance and stomatal density (Souza et al. 2011; Barylka et al. 2001) and; disturbing nutrient metabolism and plant antioxidant machinery (Gill and Tuteja 2010). For *B. napus*, Cd application has also been found to reduce the concentration of leaf pigments, including lutein, neoxanthin, β -carotene and violaxanthin (Barylka et al. 2001).

Generally, photosystem II (PS II) is more affected by Cd toxicity than photosystem I (PS I) (Baker, 1991). However, the impact of Cd toxicity on PS systems I and II is not still clear. In a time course experiment, Herbert et al. (2006) revealed that Cd downregulated many genes involved in glucosinolate biosynthesis and photosynthesis within hours of Cd exposure, while genes involved in phenylpropanoid metabolism, cell wall metabolism and sulfur metabolism were rapidly induced. Ultimately, high levels of Cd exposure, as well as long term exposure of elevated Cd, may cause plant death (Chen et al. 2010; Mohamed et al. 2012).

5.2 Cadmium Genotoxicity

Cadmium has the capacity to affect plant genetic material (Liu et al. 2012; Andresen and Küpper 2013; Xie et al. 2014) in a variety of ways, including the reduction of nuclear DNA content (Monteiro et al. 2012), and DNA damage that includes modified bases, single- and double-strand breaks, DNA-protein cross-links, 8-hydroxyguanine, DNA demethylation and base oxidation (Liu et al. 2005), and sister chromatid exchange (Unyayar et al. 2006). Cadmium-induced DNA damage may occur indirectly by activating and deactivating levels of radical scavengers, intracellular enzymatic processes, and the defense mechanism of target cell. It is believed that Cd damage to DNA is associated with the inhibition of 8-oxoguanine-DNA glycosylase activity (Kovalchuk et al. 2005). Cadmium may be transported across cellular/nuclear membranes and disrupt the DNA repair and replication mechanisms by binding directly or indirectly to DNA and/or protein (Liu et al. 2012). Cadmium exposure may also cause the breakdown of 1300 bp, a ladder-like degradation of DNA (Andosch et al. 2012). Some authors, however, believe that Cd alone could not induce DNA damage, but that it instead acts in combination with other genotoxic substances (Bandyopadhyay and Mukherjee 2011).

Cadmium can cause DNA damage/oxidation indirectly via ROS production. The hydroxyl radical (HO^\bullet) is highly electrophilic and can oxidize DNA by reacting with pyrimidine aromatic rings. The HO^\bullet radical preferentially attacks position 5 of cytosine and thymine and can also attack the hydrogen atom of the thymine methyl group, resulting in the formation of an allylic radical (which can in turn form peroxy radicals by reacting with oxygen) (Nzengue et al. 2015). The peroxy radicals are dismutated to hydroperoxides, which can induce the protonation reaction (Tremblay et al. 1999), mainly in the presence of metals. The purine bases of DNA are more perceptive to Cd-induced ROS, particularly those with lower oxidation potential. Generally, ROS induced formation of 8-OHdG starts by transferring an electron to guanine, resulting in the formation of a radical cation. This radical cation loses a proton to generate a neutral radical, which acts as an oxazolone precursor. Alternatively, the radical formed from guanine may produce a neutral radical intermediate by hydration, which is reduced or oxidized respectively to generate 2,6-diamino-4-hydroxy-5-formamidopyrimidine and 8-OHdG (Nzengue et al. 2015).

In the absence of DNA repair, the formation of 8-OHdG may result in the GC to TA transversion, which is involved in the development of cancer (Feng et al. 2003). Formation of 8-OHdG is often used as biomarker of DNA damage (Nzengue et al. 2015). Cd-induced ROS can also cause DNA damage indirectly by reacting with proteins and generating 8-OHdG (Luxford et al. 2002). For example, peroxy radicals of protein can cause oxidative lesions like 8-OHdG, whereas alkoxy radicals of protein can generate DNA adducts (Luxford et al. 2002). DNA damage can also occur as a result of Cd-induced lipid peroxidation, since by-products of lipid peroxidations such as 4-hydroxy-2-nonenal and malondialdehyde are capable

of reacting with DNA bases (Stone et al. 1990). Malonedialdehyde can react with nucleophilic atoms of cytosine, adenine and guanine by its two electrophilic carbonyl bonds (Blair 2001), which lead to the formation of a number of adducts, such as pyrimidopurinones.

Cadmium-induced DNA damage has been reported for different plant species studied using various genotoxicity techniques. Comet assays were used by Bandyopadhyay and Mukherjee 2011 for a study involving *Nicotiana tabacum* and *Allium cepa*. Liu et al. (2005 and 2009), in contrast, used random amplified polymorphic DNA (RAPD) profiles and found that Cd exposure caused loss of normal bands, appearance of new bands, and variation in band intensity. Zhang et al. (2012) used microarray analysis and observed 28 up-regulated genes and 19 down-regulated genes in Cd-exposed *O. sativa* roots. Monteiro et al. (2012) used full peak coefficient of variation (FPCV) and reported increased FPCV values by Cd in *Thlaspi arvense* and *T. caerulescens*.

Another reported symptom of Cd toxicity is the disruption of cell mitosis via micronucleus formation, chromosomal fragmentation and aberrations, loss of acentric fragments during meiosis and restrictions to mismatch repair causing mutations (Jin et al. 2003). Micronuclei emanate as a result of mitotic anomalies or chromosomal breaks that need a passage through mitosis to be recognizable (Al-Sabti and Metcalfe 1995). Cadmium-induced micronuclei are most often associated with processes involving ROS-induced redox shuttling (Circu and Aw 2010). In a 5-day experiment, Cd exposure increased micronucleus frequency in *V. faba* root tips (Foltête et al. 2012).

Cadmium may also interfere with RNA metabolism, trigger programmed cell death, cause chromatin fragmentation (Marcano et al. 2002; Behboodi and Samadi 2004), and impact genetic functioning by affecting protein structure. Protein carbonylation (or oxidative protein modification) is commonly observed after Cd exposure (Kranner and Colville 2011). In *C. sativus* seedlings, Cd exposure resulted in a two-fold increase in protein carbonylation (Gonçalves et al. 2007), which is known to inactivate proteins and cross-linking due to the conversion of side-chains of Lys, Arg, Pro and Thr to aldehyde or keto groups (Rinalducci et al. 2008).

5.3 Cadmium-Induced Oxidative Stress and Lipid Peroxidation

Plant organelles such as chloroplasts, peroxisomes and mitochondria are the main sites of ROS generation (~90 %) in plant cells (Lushchak 2011; Del Río 2011). Reactive oxygen species are generally short lived, unstable and chemically highly reactive molecules due to the presence of unpaired valence shell electrons (Andresen and Küpper 2013; Shahid et al. 2014f). ROS include: singlet oxygen ($^1\text{O}_2$), hydroxyl (HO^\bullet), superoxide anion ($\text{O}_2^{\bullet -}$), hydrogen peroxide (H_2O_2), peroxy (RO_2^\bullet), alkoxyl (RO^\bullet) radicals, and organic hydroperoxide (ROOH).

Cadmium-induced ROS production is common, and one of the earliest biochemical changes exhibited by plant subjected to Cd stress (Kung et al. 2014; Cuypers et al. 2011; Andresen and Küpper 2013; Farooq et al. 2013; Pourrut et al. 2008; Shahid et al. 2014f, 2015a). Recently, Shahid et al. (2012d) reported in a time course experiment that metal-induced ROS production occurred immediately after metal concentrations reached significant levels in roots (after 1 h of exposure). Moreover, metal-induced ROS production may occur in phases (e.g., two bursts of overproduction of ROS, occurring at 1 and 12 h). Data published after 2000 regarding Cd-induced ROS production (mainly in the form of H_2O_2 and $\text{O}_2^{\cdot -}$) for different plant species and various exposure durations and levels of Cd are summarized in Table 5.

ROS production is known to be an unavoidable consequence of aerobic metabolism (Shahid et al. 2014d, e). Although Cd is not a redox active metal and unable to produce ROS directly via Fenton and Haber-Weiss reactions, indirect ROS production has been reported in Cd-exposed plants (Gallego et al. 1996; Hasan et al. 2009; Iannone et al. 2010). Cadmium causes three waves of ROS, i.e., hydrogen peroxide (H_2O_2) by NADPH oxidase, superoxide anions ($\text{O}_2^{\cdot -}$) in mitochondria and fatty acid hydroperoxide (Garnier et al. 2006). Direct interaction between Cd and enzyme binding sites resulting from the affinity of Cd for thioyl-, histidyl- and carboxyl-groups is proposed to be responsible for Cd-induced ROS production. These interactions cause enzyme inactivation which in turn shatters ROS balance and results in oxidative stress (Cuypers et al. 2011). Cadmium also causes increased ROS production via the binding and consumption of GSH and its derivatives, which are reported to restrict ROS production by binding reactive transition elements (Hodoshima et al. 2007). In addition to these, plasma-membrane-bound NADPH oxidase has been shown to be involved in Cd-induced oxidative stress for *Pisum sativum* (Rodríguez-Serrano et al. 2006), *N. tabacum* (Olmos et al. 2003; Garnier et al. 2006) and *S. nigrum* (Deng et al. 2010).

Plant NADPH oxidases are homologs of the mammalian NADPH oxidase catalytic subunit pg91phox and are called Rboh—respiratory burst oxidase homologs (Jakubowska et al. 2015). Activation of NADPH oxidase involves the translocation and association of cytoplasmic proteins with membrane bound components. The proteins involved in the activation of NADPH oxidase include two small GTP-binding proteins (Rap1A, p21rac), two plasma membrane-associated proteins (p22phox, gp91phox) and three cytosolic proteins (p47phox, p67phox, p40phox) (Pourrut et al. 2008). NADPH oxidase generates unstable superoxide radical anion ($\text{O}_2^{\cdot -}$) by catalyzing oxygen reduction using NADPH as the electron donor. Studies with diphenylene iodonium and imidazole as an inhibitor of NADPH oxidase and NaN_3 as an inhibitor of peroxidase revealed that NADPH oxidase and peroxidase are the main sources of Cd-induced ROS in plants (Chou et al. 2012). Similar to animals, the mitochondrial electron transfer chain is the main place of Cd-induced ROS production (Heyno et al. 2008).

The essential roles of ROS in living organisms include: programmed cell death via signal transduction, initiation of defense metabolism under stress, growth regulation, regulation of stomatal movement, fruit ripening and senescence, and

Table 5 Cadmium-induced ROS production in different plant species. Data is collected from manuscripts published after 2000

ROS	Plant species	Cd exposure level	Culture	Duration (days)	References
H ₂ O ₂ , O ₂ ⁻	<i>Gossypium hirsutum</i>	500 µM CdCl ₂	Hydroponics	7	Daud et al. (2015)
H ₂ O ₂	<i>Helianthus annuus</i>	5 µM CdCl ₂	Hydroponics	14	Hawrylak-Nowak et al. (2015)
H ₂ O ₂	<i>Nicotiana tabacum</i>	100, 500 µM CdCl ₂	Hydroponics	8, 13, 25	Iannone et al. (2015)
H ₂ O ₂	<i>Triticum aestivum</i>	200 mg/kg CdCl ₂	Soil	30	Iqbal et al. (2015)
H ₂ O ₂	<i>Populus nigra</i>	50 µM CdSO ₄	Hydroponics	1 h	Lomaglio et al. (2015)
H ₂ O ₂	<i>Triticum aestivum</i>	1000 µM CdCl ₂	Hydroponics	45	Rady and Hemida (2015)
H ₂ O ₂ , O ₂ ⁻	<i>Brassica campestris</i>	50 µM CdCl ₂	Hydroponics	24 h	Wu et al. (2014)
H ₂ O ₂ , O ₂ ⁻	<i>Arabidopsis thaliana</i>	80 mg/L CdCl ₂	Hydroponics	1, 3, 6, 12, 24 h	Gu et al. (2014)
H ₂ O ₂	<i>Brassica juncea</i>	50 µM CdCl ₂	Hydroponics	15	Asgher et al. (2014)
H ₂ O ₂	<i>Ganoderma lucidum</i>	1000 µM CdCl ₂	Hydroponics	5	Kung et al. (2014)
H ₂ O ₂ , O ₂ ⁻	<i>Cynodon dactylon</i>	750 mg/kg CdCl ₂	Soil	21	Shi et al. (2014)
H ₂ O ₂	<i>Arabidopsis thaliana</i>	5, 10 µM CdSO ₄	Hydroponics	1	Keunen et al. (2013)
H ₂ O ₂	<i>Phaseolus vulgaris</i>	20 µM CdCl ₂	Hydroponics	3	Saidi et al. (2013)
H ₂ O ₂	<i>Gossypium hirsutum</i>	1, 5 µM CdCl ₂	Hydroponics	35	Farooq et al. (2013)
H ₂ O ₂	<i>Arabidopsis thaliana</i>	50, 100 µM CdCl ₂	Hydroponics	3, 9, 24 h	Martínez-Peñalver et al. et al. (2012)
H ₂ O ₂	<i>Chlorella vulgaris</i>	100 µM Cd (NO ₃) ₂	Hydroponics	24 h	Piotrowska-Niczyporuk et al. (2012)
H ₂ O ₂ , O ₂ ⁻	<i>Gracilaria dura</i>	0.4 mM CdCl ₂	Hydroponics	4	Kumar et al. (2012)
H ₂ O ₂	<i>Brassica juncea</i>	890, 1779 µM CdSO ₄	Hydroponics	45	Ahmad et al. (2011)
H ₂ O ₂	<i>Medicago sativa</i>	750 mg/kg CdCl ₂	Soil	60	Antolín et al. (2010)
H ₂ O ₂ , O ₂ ⁻	<i>Solanum nigrum</i>	200 µM CdCl ₂	Hydroponics	3	Deng et al. (2010)
H ₂ O ₂	<i>Brassica juncea</i>	200 µM CdCl ₂	Hydroponics	2, 6, 12, 24 h	Guan et al. (2009)
NO [•] , H ₂ O ₂ , O ₂ ⁻	<i>Pisum sativum</i>	50 µM CdCl ₂	Hydroponics	14	Rodríguez-Serrano et al. (2009)

(continued)

Table 5 (continued)

ROS	Plant species	Cd exposure level	Culture	Duration (days)	References
OH [•] , H ₂ O ₂ , O ₂ ^{•-}	<i>Ceratophyllum demersum</i>	10 µM CdCl ₂	Hydroponics	7	Aravind et al. (2009)
H ₂ O ₂	<i>Triticum aestivum</i>	50, 250 µM CdCl ₂	Hydroponics	24 h	Singh et al. (2008a)
H ₂ O ₂ , O ₂ ^{•-}	<i>Arabis paniculata</i>	22, 44, 89, 178 µM CdCl ₂	Hydroponics	24 h	Qiu et al. (2008)
NO [•]	<i>Triticum aestivum</i>	1000 µM CdCl ₂	Hydroponics	5	Groppa et al. (2008)
H ₂ O ₂ , O ₂ ^{•-}	<i>Solanum tuberosum</i>	100 µM CdCl ₂	Hydroponics	1 h	Heyno et al. (2008)
H ₂ O ₂	<i>Vigna mungo</i>	25, 50, 100 mg/kg CdCl ₂	Soil	30	Singh et al. (2008b)
O ₂ ^{•-}	<i>Mytilus galloprovincialis</i>	27–52 ng/g	Hydroponics	7	Koutsogiannaki et al. (2006)
H ₂ O ₂	<i>Nicotiana tabacum</i>	5 mM CdCl ₂	Hydroponics	7	Olmos et al. (2003)
O ₂ ^{•-}	<i>Lupinus luteus</i>	100 µM CdCl ₂	Hydroponics	6	Kopyra and Gwóźdź (2003)
H ₂ O ₂	<i>Pisum sativum</i>	50 µM CdCl ₂	Hydroponics	14	Romero-Puertas et al. (2002)
O ₂ ^{•-}	<i>Oryza sativa</i>	100, 500 µM Cd(NO ₃) ₂	Hydroponics	5	Shah et al. (2001)

O₂^{•-}; superoxide anion, HO[•]; hydroxyl, H₂O₂; hydrogen peroxide, NO[•]; nitric oxide

alleviation of seed dormancy (Hasan et al. 2009; Shahid et al. 2014f and references therein). Recently, it was reported that ROS (H₂O₂) act as signaling molecules to emanate defense mechanisms against toxic metals including Cd. Hydrogen peroxide, for example, can activate protein kinases, which in turn regulate gene transcription by repressing or activating the transcription factors (Pandey and Somssich 2009). Several studies have reported Cd-induced activation of protein kinase transcripts via ROS accumulation in plants (Lushchak 2011; Zhang et al. 2012).

An imbalance between ROS production and elimination takes place after Cd exposure, leading to dramatic physiological challenges in the plant termed as “oxidative stress”. Interaction of ROS with biomolecules and the resulting toxicity vary with the nature of target molecules and ROS. Although ROS can interact with and modify all cellular components, some are more susceptible. ROS-induced degradation/oxidation/inactivation is more pronounced for enzymes containing active thiol groups such as glyceraldehyde-3-phosphate dehydrogenase or [Fe–S]-clusters such as fumarate hydratase and aconitase, 6-phosphogluconate dehydratase (Lushchak 2011; Shahid et al. 2014f). ROS-induced degradation of most cellular

constituents is irreparable. The few biomolecules that can be repaired include DNA and cysteine and methionine residues in proteins (Lushchak, 2007). A range of physiological dysfunctions are thus observed due to ROS interaction with lipids, proteins and DNA.

Recent research suggests that Cd-induced ROS causes changes in cellular redox homeostasis in favor of the pro-oxidants, mainly by activating the peroxidation of membrane lipids (Körpe and Aras 2011; Andresen and Küpper 2013). Lipid peroxidation is the process whereby ROS generated by Cd or other heavy metals oxidize lipids and their esters in cell membranes (Hasan et al. 2009; Shahid et al. 2014f). Cadmium-induced lipid peroxidation has been reported in many studies involving different plants (Liptáková et al. 2013); it occurs commonly in polyunsaturated fatty acids and involves a process of initiation, progression and termination (Shahid et al. 2014f). Lipid peroxidation results in a reduction of plant development and the generation of highly cytotoxic compounds (Márquez-García et al. 2011; Pourrut et al. 2011, 2013). This process leads to oxidative damages that compromise the general redox homeostasis (Cuypers et al. 2011) and alteration to the lipidic environment which impairs its fluidity and intrinsic membrane protein activities (Quartacci et al. 2006; Shahid et al. 2014f). ROS-induces changes to biomembranes resulting from Cd toxicity affects membrane permeability to nutrients, water and protons.

Cadmium toxicity also impacts lipid biochemistry by inhibiting biosynthetic pathways which may lead to a reduction in unsaturated fatty acids (Mohamed et al. 2012). Under Cd stress, plant species including *B. juncea*, *Capsicum annum*, *Karelian birch* and *T. aestivum* balance ROS production by regulating the composition of membrane lipids (e.g. neutrallipids, phospholipids and galactolipids) (Belkadhi et al. 2015). Moreover, Cd may substitute Ca from essential sites on the endoplasmic reticulum membranes, tonoplast, apoplast and mitochondrial (Breckle and Kahle 1991). Cadmium diffusion into cytosol activates phospholipases, which results in membrane discharge of linolenic (C18:3) and linoleic (C18:2) acids serving as substrates for ROS or lipoxygenase (Chmielowska-Bąk et al. 2014), which also has been reported in many species such as *O. sativa* and *H. vulgare* (Belkadhi et al. 2015). ROS production also has a major impact on the activation of membrane-bound enzymes (Ca^{2+} -ATPase and H^{+} -ATPase).

6 Cadmium Detoxification in Plants

Plants naturally possess a well-organized antioxidant defense system for managing overproduction of ROS and evading oxidative stress (Anjum et al. 2012; Shahid et al. 2014f). Inherent to this defence system are mechanisms to increase: Cd elimination, Cd redistribution to insensitive tissues including sequestration in vacuoles, cellular Cd binding to cell walls, and complexation of Cd by organic ligands. These mechanisms are aided by metabolic compounds such as

phytochelatins (PCs), glutathione (GSH), carotenoids and tocopherols, and enzymatic antioxidant systems including superoxide dismutases (SOD and EC 1.15.1.1), catalase (CAT and EC 1.11.1.6), peroxidase (POD, EC 1.11.1.7), ascorbate peroxidase (APX, EC 1.11.1.11), guaiacol peroxidase (GPX, EC 1.11.1.7), glutathione reductase (GR, EC 1.6.4.2), monodehydroascorbate reductase (MDHAR, EC 1.6.5.4) and dehydroascorbate reductase (DHAR, EC 1.8.5.1). These mechanisms often act simultaneously to scavenge free ROS and can have immediate (PCs) or long-term (gene encoding) effects. However, the plasticity of induction of a specific defense mechanism and its efficiency depends on plant species and maturity, metal type and exposure duration (Andresen and Küpper 2013; Shahid et al. 2014f). Together, the increased levels of the defence-related metabolic compounds and antioxidant enzymes lead to increased stress tolerance against heavy metal-induced ROS (Pourrut et al. 2011).

6.1 Vacuolar Compartmentalization

Vacuolar compartmentalization is important for Cd homeostasis and detoxification as it reduces the movement of free Cd in the cytosol (Mendoza-Cózatl et al. 2011; Choppala et al. 2014). Cadmium sequestration in vacuoles may occur via PC-dependant or -independent pathways (Wojas et al. 2009; Khoudi et al. 2012), with direct Cd transport facilitated by Cd/H⁺ antiport activity (Salt and Wagner 1993) driven by trans-membrane H⁺ gradients. Increased Cd sequestration in vacuoles has been reported due to the expression of CAX1 variant in petunia (Wu et al. 2011). There are several molecules involved in vacuolar Cd sequestration, but analyses from various plant species have shown that PCs and glutathione (GSH) are the main metal–ligand molecules involved.

6.1.1 Phytochelatins

Phytochelatins (PCs) are a family of peptides with the structure (c-Glu-Cys)_n-Gly, where n represents the repetition of g-Glu–Cys, which tends to vary from 2 to 11 (Akhter et al. 2012). As with other metals, PCs play an important role in Cd detoxification (Márquez-García et al. 2011; Choppala et al. 2014). Phytochelatin synthesis under metal stress is perhaps one of the most well studied mechanisms mediating metal detoxification in plants and some yeast (Mendoza-Cózatl et al. 2011). Phytochelatins are reported to neutralize harmful effects by chelating metals with their SH groups. Of all the heavy metals, Cd²⁺ has been observed to invoke the strongest activation of the phytochelatin synthase (PCS) and thus PC production (Akhter et al. 2012) due to the high affinity of the thiol groups for Cd on the cysteine residues (Chekmeneva et al. 2011). Chen et al. (2007) studied Cd–PC complexes in *Brassica chinensis* and observed binding stoichiometries of 1:1 to 3:1 depending on the availability of thiol groups and Cd²⁺ in the cytosol.

The process of PC formation subsequent to Cd exposure is complex (Lima et al. 2012) and includes Cd absorption, GSH synthesis, and PCS activation followed by synthesis of the PC oligomeric chains (Ogawa et al. 2011). During PC synthesis, a g-Glu-Cys unit from GSH is first added to the GSH molecule itself and then to the elongating PC chain (Rea et al. 2004; Ramos et al. 2007). Although the PCS enzyme is thought to be self-regulated (due to the synthesis of PC following Cd exposure (Loeffler et al. 1989)), its mechanism of self-regulation remains unknown. While protein phosphorylation has been shown to prompt PCS synthesis during GSH production (Wang et al. 2009), there are no known reports of protein phosphatases involved in PC regulation. Huang et al. (2012) reported that Cd tolerance and accumulation failed when GSH and PC synthesis was blocked by L-buthionine sulfoximine, and also when PC synthesis was blocked in the PCS-deficient *cad1-3* mutant.

Once formed, Cd-PC complexes are transported through the tonoplast and compartmentalized in metabolically inactive sites (Najmanova et al. 2012) such as vacuoles (Fig. 3) (Lux et al. 2011), where they are promptly sequestered (Verbruggen et al. 2009) or released to apoenzymes. Transport of the Cd-PC complex to vacuoles is thought to be facilitated by ABC transporters (ATP-binding cassette transporters) (Prévéral et al. 2009), which for *O. sativa* seedlings, are encoded by *OsPDR5/ABCG43* (Oda et al. 2011). A half-size ABC transporter (Hmt1; Heavy metal tolerance 1) has been identified in *Schizosaccharomyces pombe* as a vacuolar Cd-PC transporter involved in Cd tolerance (Salt and Rauser 1995). It is unclear whether Hmt1 function is GSH-dependent or PC-dependent.

Two independent groups of ABC transporters have been identified as being involved in the mediation of vacuolar PC transport in *Arabidopsis* (*MRP1/ABCC1* and *MRP2/ABCC2*) and *S. pombe* (*Abc2*) (Mendoza-Cózatl et al. 2010). These transporters (*ABCC1* and *ABCC2*) have been found to mediate the sequestration of metal-glutathione complexes into vacuoles (Rea 2007). Sequestration of Cd by the PC complex depends on the complex stability; with Cd-PC complexes having acid-labile sulphur possessing an improved stability and a higher Cd sequestering efficiency. Complex stability has also been found to increase with chain length up until PC₃, beyond which the number of thiol groups (i.e., ≥ 4) saturate the tetrahedral coordination of Cd²⁺ (Chekmeneva et al. 2011).

Moreover, PCs have also been found to be involved in the long-distance transport of Cd from root to shoot (Mendoza-Cózatl et al. 2011). Using Extended X-ray Absorption Fine Structure (EXAFS), Vogel-Mikus (2010) showed that 60 % of Cd is coordinated with PCs in *Thlaspi praecox*. This is in addition to the earlier work of Belleghem et al. (2007) who identified sulfur-Cd complexes in the cytoplasm of phloem-loading cells of *A. thaliana* (consistent with the idea that thiols (PCs) mediate long-distance Cd transport). Long-distance Cd transport via xylem sap is mediated by ZIP transporters, xylem loading Heavy Metal ATPases (*HMA2* and *HMA4*), and PC transporters *ABCC1* and *ABCC2* (Mendoza-Cózatl et al. 2011). These PC-Cd complexes loaded into the phloem are more likely to be sequestered in root vacuoles (Mendoza-Cózatl et al. 2011). While the mechanisms involved in long-distance Cd translocation from root to shoot are not yet completely

understood, it is possible that PC-mediated movement of Cd out of the plant shoots could result in more systemic disruption of essential processes (if the plant is unable to compartmentalize Cd and render it less biologically reactive).

Although Cd-induced production of PC in plants is well reported (Vurro et al. 2011; Ogawa et al. 2011; Son et al. 2012), some authors argue that PC is not involved at all in Cd detoxification (Inouhe et al. 2000). Ebbs et al. (2002) showed that PC synthesis had no role in Cd tolerance in *T. caerulescens*. Zhang et al. (2010) also showed no significant role of PC in Cd tolerance in *S. alfredii*. In fact, Cd is reported to induce PC synthesis only at high concentrations (Lugon-Moulin et al. 2004)—consistent with the theory of Salt and Wagner (1993) who argue that it would only be at high levels of Cd exposure that PCs might play a role. Thus the exact role of PCs in Cd detoxification for plants remains unclear. As the expression and role of PCs in Cd tolerance quite likely vary with plant type (e.g., metal sensitive or tolerant), continued research is needed to further our understanding of the role of these compounds—particularly for plant species which can be used for phytoremediation.

6.1.2 Glutathione

Glutathione (GSH) is a low molecular weight tri-peptide which is the major source of non-protein thiol in plant cells (Márquez-García et al. 2011; Pourrut et al. 2013). Owing to its unique nucleophilic and redox properties, GSH plays a key role in physiological functions such as transport of GSH-conjugated amino acids and intracellular redox state regulation (Deng et al. 2010). As an important antioxidant, GSH takes part in bio-reductive reactions as an ROS scavenger to relieve cells from oxidative stress (Park et al. 2012; Bashir et al. 2013), becoming oxidized to GSSG in the process (Shahid et al. 2014f). The antioxidant properties of GSH depend on the oxidation of the SH group, giving rise to GSSG (Márquez-García et al. 2011). Glutathione synthesis takes place in two steps which are ATP-dependent. These two steps are catalyzed by glutamylcysteine synthetase (ECS) and glutathione synthetase (GS) (May et al. 1998; Fässler et al. 2011). GSHI and GSHII genes are respectively involved in encoding these two enzymes (Zhu et al. 1999). Therefore, enhanced generation of GSH due to the overexpression of g-glutamylcysteine synthetase (g-ECS) leads to increased tolerance of Cd. Overexpression of GS has also been shown to result in higher GSH levels and Cd tolerance (Zhu et al. 1999; Lugon-Moulin et al. 2004). Nowadays these genes are used in genetic engineering for increased phytoaccumulation and Cd tolerance. Under Cd stress, GSH production may decrease or increase. Glutathione contents increase during the initial metal exposure phase but then return to control levels or lower (Chen et al. 2012). The decrease in GSH content might be due to its high rate of consumption as a reductant in antioxidant systems as metal load increases with concentration and exposure duration (Mishra et al. 2006; Chen et al. 2012).

Transgenic plants are capable of producing high GSH levels which can (but do not always) enhance tolerance to Cd and other metals (Xiang et al. 2001; Choppala

et al. 2014). Indeed, GSH alone is not capable of providing enough protection against trace metal toxicity, especially under conditions of high metal loading (Noctor et al. 1998). Different physiological, biochemical and genetic studies suggest that GSH acts as a substrate for PC synthesis (Cobbett, 2000; Márquez-García et al. 2011). Glutathione is also involved in Cd-GSH transport to vacuoles via ABC transporters (Fig. 3) (Lugon-Moulin et al. 2004). Cadmium produces an imbalance of the available GSH pool because of its high affinity for thiol groups. In this way, Cd disturbs the AsA–GSH cycle which has been suggested as the source of H_2O_2 removal in many plants (Semane et al. 2007).

6.2 Antioxidant Enzymes

Cadmium is well known to enhance the activities of antioxidant enzymes—enzymes which tend to be electron donors that react with ROS to form innocuous end products (e.g., water) (Shahid et al. 2014f; Abbas et al. 2015). Several studies have reported Cd-induced activation of antioxidant enzymes (Table 6), possibly via modulation of gene expression (Anjum et al. 2012) or Cd-induced restriction of enzyme inhibitors (Ali et al. 2013; Deng et al. 2014; Chen et al. 2014a, b). Cadmium may also enhance antioxidant enzymes activity indirectly by increasing the concentration of their substrates (Anjum et al. 2012). Antioxidative enzymes activated in response to Cd stress for different plant species are highlighted in Table 6.

Because of their ability to detoxify ROS, antioxidative enzymes provide an important line of defense against Cd toxicity (Iqbal et al. 2010). The following describes some of the principal known antioxidant enzymes:

Superoxide Dismutase (SOD) is a key antioxidant enzyme which defends plants against ROS by removing $O_2^{\bullet -}$ by catalyzing its dismutation (i.e., one $O_2^{\bullet -}$ is reduced to H_2O_2 and other oxidized to O_2) (Noctor and Foyer 1998; Hasan et al. 2011). H_2O_2 , which itself may serve as a signaling molecule in the activation of antioxidative enzymes (Chou et al. 2012), is subsequently detoxified by enzymes such as GPX, APX, and CAT. Cadmium-induced activation of SOD is reported in different plant species (Saidi et al. 2014; Deng et al. 2014; Ehsan et al. 2014; Irfan et al. 2014; Howladar 2014).

Catalase (CAT) can convert H_2O_2 to O_2 and H_2O (Chen et al. 2012). CAT has very high catalytic rates but generally low substrate affinities, since the reaction requires the simultaneous access of two H_2O_2 molecules (Willekens et al. 1997). Several studies have shown Cd-induced increases in CAT activity in plants (Saidi et al. 2013; Chen et al. 2014a, b; Shi et al. 2014; Agami and Mohamed 2013).

Guaiacol Peroxidase (GPX) requires a reductant such as guaiacol to reduce H_2O_2 to H_2O (Gill et al. 2012). Activation of GPX under Cd stress is reported by several authors (Monteiro et al. 2011; Wang et al. 2012; Farooq et al. 2013; Ehsan et al. 2014).

Table 6 Cadmium-induced activation of different enzymes in different plant species. Data is collected from manuscripts published between 2010 and 2015

Enzymes	Plant species	Cd exposure level	Culture	Duration (days)	References
SOD, CAT, POD, GR	<i>Gossypium hirsutum</i>	500 μM CdCl_2	Hydroponics	7	Daud et al. (2015)
SOD, CAT, APX, GPX	<i>Nicotiana tabacum</i>	100, 500 μM CdCl_2	Hydroponics	8, 13, 25	Iannone et al. (2015)
SOD, CAT, POD, PPO	<i>Zea mays</i> and <i>Helianthus annuus</i>	10, 25 mg kg^{-1}	Soil	60	Aghababaei and Raiesi (2015)
PROX, GPX, GR	<i>Triticum aestivum</i>	200 mg/kg CdCl_2	Soil	30	Iqbal et al. (2015)
GR, APX, NADPH oxidase, SOD	<i>Medicago sativa</i>	6, 30 μM CdCl_2	Hydroponics	7	Flores-Caceres et al. (2015)
SOD, CAT, APX, GR	<i>Trifolium repens</i>	100 μM CdCl_2	Hydroponics	7	Liu et al. (2015)
CAT, POX, APX	<i>Triticum aestivum</i>	1000 μM CdCl_2	Hydroponics	45	Rady and Hemida (2015)
SOD, POD, CAT	<i>Ricinus communis</i>	2, 5 mg L^{-1} CdCl_2	Hydroponics		Zhang et al. (2015)
GPX, DHAR, LOX, AAO	<i>Hordeum vulgare</i>	15 μM CdCl_2	Hydroponics	3, 6 h	Tamás et al. (2015)
GST, CAT, GPX, APX, GR	<i>Triticum aestivum</i>	50 μM Cd (NO_3) ₂	Hydroponics	7	Kovács et al. (2014)
SOD, CAT, POD, APX, GR	<i>Helianthus annuus</i>	20 μM CdCl_2	Hydroponics	4	Saidi et al. (2014)
CAT, SOD, POD	<i>Ceratopteris pteridoides</i>	0, 5, 10, 20, 40 μM	Hydroponics	7	Deng et al. (2014)
SOD, GPX, CAT, APX	<i>Brassic napus</i>	10, 20 μM CdCl_2	Hydroponics	56	Ehsan et al. (2014)
SOD, CAT, POX	<i>Brassica juncea</i>	25, 50, 100 mg/kg CdCl_2	Soil	30, 60	Irfan et al. (2014)
SOD, POD, GR, CAT	<i>Phaseolus vulgaris</i>	90 μM , 1 M CdCl_2	Soil	35	Howladar (2014)
SOD, POD, CAT	<i>Phanerochaete chrysosporium</i>	1–500 μM Cd (NO_3) ₂	Hydroponics	1	Chen et al. (2014a, b)
CAT, APX, GPX	<i>Brassica juncea</i>	5, 35 mg/kg CdCl_2	Soil	60	Armas et al. (2014)
SOD, CAT, POD, GR	<i>Cynodon dactylon</i>	750 mg/kg CdCl_2	Soil	21	Shi et al. (2014)

(continued)

Table 6 (continued)

Enzymes	Plant species	Cd exposure level	Culture	Duration (days)	References
SOD, CAT, POX	<i>Triticum aestivum</i>	500, 1000 μM CdCl_2	Hydroponics	56	Agami and Mohamed (2013)
CAT, SOD, POD	<i>Arabidopsis thaliana</i>	150 μM CdCl_2	Hydroponics	7	Tao et al. (2013)
CAT, APX	<i>Phaseolus vulgaris</i>	20 μM CdCl_2	Hydroponics	3	Saidi et al. (2013)
SOD, GPX, CAT, APX	<i>Gossypium hirsutum</i>	1, 5 μM CdCl_2	Hydroponics	35	Farooq et al. (2013)
APX, SOD, POD	<i>Brassica napus</i>	100 and 500 μM CdCl_2	Hydroponics	15	Ali et al. (2013)
SOD, GR, APX, CAT	<i>Arabidopsis thaliana</i>	50 μM CdCl_2	Hydroponics	3, 5	Bashir et al. (2013)
SOD, CAT, POD, GR	<i>Hibiscus cannabinus</i>	20, 50, 80, 120 μM CdCl_2	Hydroponics	21	Feng-tao et al. (2013)
SOD, CAT, GR, GST, APX	<i>Lemna gibba</i>	0.05, 0.5, 5, 10, 20 mgL^{-1} Cd (NO_3) ₂	Hydroponics	7	Parlak and Yilmaz (2013)
SOD, APX, GR	<i>Gracilaria dura</i>	0.4 mM CdCl_2	Hydroponics	4	Kumar et al. (2012)
APX, MDHAR, DHAR, GR, GST	<i>Helianthus annuus</i>	10 mg/kg Cd	Soil	3	Nehnevajova et al. (2012)
SOD, GR, APX	<i>Lepidium sativum</i>	25, 50, 100 mg kg^{-1}	Soil	30	Gill et al. (2012)
POD, APX, CAT, SOD	<i>Triticum aestivum</i>	0.10 mmol/l CdCl_2	Hydroponics	10	Guo et al. (2012)
SOD, POX, APX, CA	<i>Solanum lycopersicum</i>	3, 6, 9, 12 mg/kg Cd	Soil	20	Hayat et al. (2012)
CAT, POX	<i>Brassica juncea</i>	50, 200 μM Cd (NO_3) ₂	Hydroponics	7	Mohamed et al. (2012)
SOD, GR, APX	<i>Erica andevalensis</i>	0.53, 5.3, 53, 530 μM CdCl_2	Hydroponics	5	Márquez-García et al. (2012)
SOD, GPOX, APX, CAT	<i>Vicia faba</i>	6 mmol/l CdCl_2	Hydroponics	15	Wang et al. (2012)
SOD, GR, APX, CAT, GST	<i>Groenlandia densa</i>	0.05, 0.5, 5, 10, 20 mgL^{-1} Cd (NO_3) ₂	Hydroponics	7	Yilmaz and Parlak (2011)
SOD, POD, CAT, APX, GR	<i>Juncus effusus</i>	10, 50 and 100 μM Cd (NO_3) ₂	Hydroponics	14	Najeeb et al. (2011)

(continued)

Table 6 (continued)

Enzymes	Plant species	Cd exposure level	Culture	Duration (days)	References
SOD, SOD, CAT, APX, GR, SPX	<i>Arabidopsis thaliana</i>	5 μM Cd	Hydroponics	3	Vanhoudt et al. (2011)
SOD, GPX, CAT, APX, GR,	<i>Solanum lycopersicum</i>	0.5 mM CdCl ₂	Hydroponics	7, 20, 36	Monteiro et al. (2011)
SOD, CAT, GPOX	<i>Sedum alfredii</i>	400 μM Cd	Hydroponics	1	Tian et al. (2011)
APX, GR, SOD, APX	<i>Tagetes patula</i>	10, 20, 50 μM CdCl ₂	Hydroponics	2	Liu et al. (2011)
SOD, CAT, APX, GR	<i>Solanum lycopersicum</i>	10 μM CdCl ₂	Hydroponics	7	Cherif et al. (2011)
APX, MDHAR, DHAR, SOD, GST	<i>Typha latifolia</i>	10, 50, 100, 250 μM CdCl ₂	Hydroponics	3	Lyubenova and Schröder (2011)
SOD, APX, CAT, GR	<i>Brassica juncea</i>	890, 1779 μM CdSO ₄	Hydroponics	45	Ahmad et al. (2011)
SOD, CAT, APX	<i>Solanum nigrum</i>	7.5, 15, 30 μM CdSO ₄	Hydroponics	45	Fidalgo et al. (2011)
SOD, POX, APX, CA	<i>Solanum lycopersicum</i>	3, 6, 9, 12 mg kg ⁻¹	Soil	60	Hasan et al. (2011)
POD, CAT	<i>Amaranthus hybridus</i>	30, 60, 90, 120, 150, 180 mg kg ⁻¹ CdCl ₂	Soil	90	Zhang et al. (2010)
SOD, GR, APX	<i>Atriplex halimus</i>	100 μM CdCl ₂	Hydroponics	112	Lefèvre et al. (2010)

SOD superoxide dismutase, APX ascorbate peroxidase, GPX guaiacol peroxidase, CAT catalase, PPO polyphenol oxidase, GR glutathione reductase, AsA ascorbic acid, GSH glutathione, GST Glutathione S-transferase, ACOX acyl co-A oxidase, POD peroxidase, DHAR dehydroascorbate reductase, MDHAR monodehydroascorbate reductase, LOX lipoxigenase, AAO ascorbic acid oxidase, SDH succinate dehydrogenase, PROX proline oxidase

Ascorbate Peroxidase (APX) is a potential participant in the ascorbate-glutathione cycle (Gill et al. 2012), and is located mainly in the cytoplasm, chloroplasts and other cellular organelles where APX along with glutathione reductase (GR) control cellular redox status. Ascorbic peroxidase requires two molecules of ascorbate to reduce H₂O₂ to O₂ and H₂O (Noctor and Foyer 1998), which once oxidized, is reduced by GSH (generated from oxidized glutathione (GSSG) by GR). Cadmium can activate APX (Nehnevajova et al. 2012; Parlak and Yilmaz 2013; Ali et al. 2013; Ehsan et al. 2014; Saidi et al. 2014).

Glutathione Reductase (GR) helps in maintaining a high GSH/GSSG ratio (Foyer et al. 1997), which is crucial for protection against oxidative damage. The

activation of GR by Cd has been reported by Bashir et al. (2013), Saidi et al. (2014) and Howladar (2014).

Ascorbic Acid (AsA) is the most abundant antioxidative enzyme in plants and participates in H_2O_2 detoxification via the AsA-GSH cycle which involves (among other things) AsA, GSH, APX, and GR (Noctor and Foyer 1998; Chao et al. 2010). AsA is not only involved in H_2O_2 detoxification, but also reacts with O_2^- , OH^- and lipid hydroperoxides (Saidi et al. 2013). The exogenous application of AsA reduces Cd toxicity as well as abiotic stresses in plants (Jin et al. 2013). Ascorbate oxidase (AAO), present in plant apoplasts, causes the oxidation of AsA to MDHA using oxygen (Chao et al. 2010).

Peroxidases (PODs) are antioxidant enzymes that catalyze the H_2O_2 -dependent oxidation of a number of substrates, especially phenolic compounds (Saidi et al. 2013). Cadmium-induced oxidative stress increases POD activity in plants (Gallego et al. 1996; Lee et al. 2010; Ali et al. 2013; Deng et al. 2014; Howladar 2014).

Dehydroascorbate reductase (DHAR) is a very important antioxidant in plants, which reduces dehydroascorbate to maintain an appropriate level of ascorbate (ASC), using GSH as the reducing substrate (Nehnevajova et al. 2012). However, no gene or cDNA has been cloned for plant DHAR (Urano et al. 2000). The ASC produced by DHAR is used by APX as a substrate for H_2O_2 detoxification. DHAR is reported to contribute to Cd detoxification via its role in the ROS scavenging cycle (Chen et al. 2010; Nehnevajova et al. 2012).

6.3 Salicylic Acid (SA)

Salicylic acid (SA) is a hormone-like signaling molecule which influences several physiological and developmental processes in plants. Salicylic acid is involved in the regulation of seed germination, plant growth, flowering, fruit yield, and glycolysis (Guo et al. 2009) and is also known to increase and elicit specific responses as a result of Cd and other trace-metal-induced stress (Ahmad et al. 2011; Tao et al. 2013). Salicylic acid has been shown to reduce Cd-induced oxidative stress in plants (Guo et al. 2007; Tamás et al. 2015) and thereby increase Cd tolerance (Metwally et al. 2003). The protective role of SA primarily includes the induction of gene expression, regulation of antioxidants and ROS, membrane stabilization, and absorption/distribution of essential nutrients (Belkadi et al. 2015). Plant SA content has been shown to increase in a wide range of species after exposure to Cd and other heavy metals (Tao et al. 2013; Kovács et al. 2014). While the means by which this endogenous signalling molecule is synthesized and regulated in response to abiotic stress (especially in crop plants) requires further study, Dempsey et al. (2011) proposed that SA is synthesized via two distinct pathways: (1) compartmentalized enzymatic pathways that use the metabolite chorismate, and (2) a phenyl-propanoid pathway that uses L-phenylalanine, and synthesizes SA via cinnamate.

Exogenous applications of SA have been shown to mitigate decreases antioxidant enzyme activity and reduce Cd-induced production of TBARS, H_2O_2 and O_2^- and lipid peroxidation in *O. sativa* (Panda and Patra 2007). Belkadhi et al. (2013) showed that exogenous SA protected membranes against a loss of integrity and lipid peroxidation by increasing fatty acid unsaturation in *Linum usitatissimum* roots. Later, Belkadhi et al. 2015 further reported that exogenous SA also protected phospholipids of *L. usitatissimum* by decreasing the percentages of monogalactosyldiacylglycerol, phosphatidylcholine, phosphatidylethanolamine and phosphatidylglycerol. The net result of exogenous SA application is the alleviation of Cd-induced root growth inhibition and overall attenuation of Cd toxicity—the latter of which has been reported for a variety of plants (e.g., *V. faba* (Popova et al. 2009), *O. sativa* (Guo et al. 2007), *B. juncea* (Ahmad et al. 2011), *G. max* (Drazic and Mihailovic 2005), *L. usitatissimum* (Belkadi et al. 2010), *H. vulgare* (Tamás et al. 2015), *Cannabis sativa* (Shi et al. 2009), *Z. mays* (Kranterev et al. 2008), *T. aestivum* (Agami and Mohamed 2013) and *Pisum sativum* (Popova et al. 2009)). Salicylic acid is reported to reduce Cd toxicity to plants not only at the level of antioxidant defense but also by altering other mechanisms of Cd detoxification which are not yet well known (Gallego et al. 2012).

Contrary to the above, some authors have postulated that SA itself may cause oxidative stress. Pal et al. (2002) reported that despite decreasing Cd uptake, exogenous SA application enhanced Cd-induced phytotoxicity. Recently, Liu et al. (2011) showed that pretreatment of *Ricinus communis* with SA alone reduced plant biomass, photosynthetic rate, stomatal conductance and intercellular CO_2 concentration. They also reported that SA pretreatment under Cd stress caused a significant decrease in plant biomass, photosynthetic rate, stomatal conductance and intercellular CO_2 concentration. Endogenous application of SA may act as a signaling molecule to initiate Cd-induced oxidative stress in *Arabidopsis thaliana* (Zawoznik et al. 2007). Collectively, these studies suggest that the presence of SA (endogenous or exogenous applications) in plants may have contrasting effects in alleviating Cd toxicity, likely due in part to plant type, applied level and form of Cd, and concentration of SA applied.

7 Hormetic Effect of Cadmium Toxicity

Cadmium can have adverse effects on the morphological, physiological, and biochemical development and function in plants. The adverse effects of Cd on plants over the complete dose-response range tend to be linearly related to Cd exposure (Gill et al. 2012; Hayat et al. 2012; Irfan et al. 2014). However, some studies have reported a hormetic effect whereby Cd is toxic at higher applied levels but acts as a growth stimulator at low levels. This biphasic dose-response phenomenon is often described by an inverted U-shaped curve (Jia et al. 2013). Hormetic U-shaped growth effects have frequently been reported for plants exposed to low

doses of toxic metal ions such as Cd, Cr, Al, and Pb (Poschenrieder et al. 2013; Calabrese and Baldwin 1999; Calabrese and Baldwin 2003a, b), and have also been reported for malondialdehyde and electrolyte leakage under low Cd exposure. The hormetic effect is not necessarily seen in studies focusing on more highly toxic levels of Cd.

Based on the current knowledge of Cd toxicity and detoxification in plants, hormetic effects of Cd may in part be attributed to adaptive compensatory processes and/or defence mechanisms (Jia et al. 2013; Tamás et al. 2015). This implies the existence of metal-sensing systems that regulate gene expression via cellular signaling transduction cascades. Growth stimulation in plants due to ROS-induced programmed cell death is an important mechanism in developmental processes (Gill and Tuteja 2010). Low level Cd exposure to *T. aestivum* has been shown to prompt a hormetic effect by stimulating plant growth (Lin et al. 2007a, b) which is accompanied by a decrease in ROS production and increase in glutathione reductase activity and GSSG/GSH ratio. An increase in SA concentration (which can similarly stimulate growth) has also been reported in plants when exposed to low levels of Cd (Keunen et al. 2013).

Although perhaps less common, defense or growth-stimulating effects may also be observed in response to higher Cd doses—occurring, for example, in plants pre-acclimated to Cd exposure. The hormetic effect of Cd may also vary with plant species, growth stage, exposure duration, and the time taken by the plant to activate the full suite of metal tolerance mechanisms (Küpper et al. 2007). However, the majority of studies did not investigate the physiological and molecular mechanisms responsible for Cd-induced plant growth stimulation. This is especially the case in studies of Cd toxicity or hyperaccumulation, which instead focused on (1) soil-plant transfer of Cd, (2) accumulation of Cd in different plant tissues, and (3) Cd-induced toxicity to basic growth parameters. For this reason, the response mechanisms of plants subjected to low levels of Cd stress requires further evaluation.

8 Conclusions and Perspectives

This review highlighted the biogeochemical behavior of Cd in soil-plant systems. Cadmium adsorption/desorption in soil, mobility towards plant roots and phyto-uptake varies greatly according to the physico-chemical environment. Factors which influence Cd uptake include (but are not limited to) Cd exposure concentration and chemical speciation, soil properties (which influence Cd lability and bioavailability), exposure duration, and plant characteristics (e.g., species type, life stage, and overall fitness). Cadmium uptake by plants, which constitutes the first step of its entrance into the terrestrial food chain, occurs via specific and non-specific transporters of essential nutrients. Inside the plant, Cd mainly accumulates in root cells and a small proportion is translocated to the shoots. Limited translocation of Cd to shoot tissues is due to the blockage of Cd by the Casparian

band within the endodermis; that which does make it through is translocated to shoot tissues mainly via different membrane transporter gene families.

Excessive Cd accumulation in plant tissue induces toxicity, which may be manifested as visual impairments such as seed development and germination, root elongation, leaf chlorosis and necrosis, and stunted growth. Excessive Cd accumulation also impairs mineral nutrition and water balance, lamellar organization and chlorophyll production in the chloroplast, cell division, and membrane permeability and structure (by reacting with functional groups of different enzymes). Moreover, Cd inhibits the activity of many enzymes by replacing essential ions and reacting with the active sites of ATP or ADP. Cadmium-induced overproduction of ROS causes DNA damage and lipid peroxidation.

Plants have well-organized enzymatic and non-enzymatic antioxidant defense systems (i.e., defence-related genes and metabolites) which are elicited in response to Cd exposure and collectively used to mitigate toxicity. Plant compartmentalization and detoxification responses include Cd retention in the cell wall (as the first barrier against Cd stress), Cd chelation by PC and subsequent compartmentalization of the PC-Cd complex, and the induction of stress signaling molecules such as salicylic acid. The plasticity of induction of a specific defense mechanism by plants and their efficiency depends on plant species (sensitive or tolerant), exposed Cd level and exposure duration. Because Cd has various negative impacts on environmental sustainability, we believe it deserves to be evaluated in supplementary studies in order to strengthen our understanding about its behavior in soil-plant systems.

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Radionuclides: Accumulation and Transport in Plants

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Highlights

1. Bioavailability of radionuclides in environment.
2. Green plants for restoring balance/their application for radionuclide accumulation.

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3. Rhizosphere, rhizobacteria and metal transporters used in radionuclide remediation.
4. Radionuclide exposure and plant responses.
5. Radio-phytoremediation—an approach to decontaminate radionuclides using plants.

1 Introduction

With the discovery of X-rays in 1895 and radioactivity in the subsequent year, awareness of radiation and radioactivity became obvious and a new branch of science was started and subsequently developed into its deliberate use. However, the term radionuclide is much newer. Referring to an atomic species having precise constitution of its nucleus containing a certain number of nucleons (neutrons (N) and protons (Z) and its nuclear energy state), in 1947, Dr. T.P. Kohman of University of Chicago proposed the name nuclide. Thus, a nuclide consists of strongly bound protons and neutrons often referred to as nucleons (Kohman 1947). Nuclides may be stable and exist for an indefinite period of time (common material) or unstable. Usually, elements having atomic number higher than 83 (heavier than bismuth ^{83}Bi), have only unstable nuclei with excess energy available. This surplus energy within a nucleus is either transformed internally or passed on to a newly formed radiation particle within the nucleus. This kind of nuclides with potential to undergo radioactive decays is called radionuclide. The decay goes together with the release of ionizing radiations like gamma ray(s) and/or subatomic, high speed alpha or beta particles. The nuclides are ultimately converted to stable nuclides. Ionizing radiation is capable of generating ions by displacing electrons in the living matter (like in DNA) and thereby disturbing its function (<http://water.epa.gov/drink/contaminants/basicinformation/radionuclides.cfm>, accessed 24.08.2015). However, exposure to radioactivity is a common and natural phenomenon. Apart from about 20 % anthropogenic sources (especially diagnostic imaging like X-rays, CT scans etc.) (Brenner and Hall 2007), most of the radiation exposures are usually natural (like cosmic radiation, radon gas from rocks and soil, or ^{40}K through foods). Water from natural sources may contain radionuclides due to dissolution of such materials from the earth crust or sporadic release from laboratories or nuclear power plants. Though, in nature, unstable nuclides with long half-lives, so called primordial radionuclides, are also present. Especially ^{235}U has been widely used for different applications in defence and civilian sectors. It is best used in nuclear power plants (due to low cost consumption), where, theoretically 1 kg of U (approx. 1500 tonnes of coal equivalent) can produce about 20 terajoules (2×10^{13} J) of energy.

However, extensive use and indiscriminate or improper disposal of wastes, insufficient decontamination at mining sites, release of production tails or decommissioning sites of such material are a serious environmental concern and can lead to contamination by radioactive substances. And, radiation is produced by

spontaneous decay of radioactive materials, the amount of which and power for penetration within the body may vary with each type. However, all kinds of ionizing radiations produce health effects that may vary in different tissues and even in different individuals. Exposure to radiation and radionuclides (radiation emitters) are well-known to affect the entire body while inhalation or ingestion may affect diverse tissues inside the body (<http://www.medindia.net/patients/patientinfo/radiation-hazards.htm>). Ionizing radiation is capable of ionizing many atoms or molecules and destroying molecular bonds. For example, due to a nuclear explosion or accident at a nuclear power plant, radioactive caesium or iodine can be released into the environment, which may directly or indirectly get accumulated within the body, for example through the food chain. For quantification of the radionuclide uptake efficiency the soil-to-plant transfer factor (TF) is widely used to analyse radiological human dose via the ingestion pathway (Chakraborty et al. 2013). The TF is basically the concentration of element in plant (Bq kg^{-1}) divided by concentration of the same element in soil (Bq kg^{-1}). The TF may vary depending upon the element, soil pH and its texture, solid/liquid distribution coefficient, exchangeable K^+ , organic matter and availability of the element at the root zone (Chakraborty et al. 2013). Starting from a given contamination level the total effective dose equivalent (TEDE) to individuals living in the neighbouring areas of waste-disposal/decommissioning sites can be calculated using models of biosphere and living habits. The Nuclear Regulatory Commission (NRC) has developed some sophisticated methodologies for the assessment of probable dose to man from evaluations of groundwater and soil contamination, irrigation processes, subsequent to food-chain pathways, of crops and forage and lifestyle of potentially exposed individuals (USNRC 2003; Napier et al. 2005, 2007). Above said chains, we want to focus following on soil to plant transfer of such radionuclides at contaminated sites (Fig. 1). Uptake of the radionuclides by plants depends upon several factors including mode of interaction with the materials and physiological characteristics of the species and factors like concentrations, bioavailability, and mobility of radionuclides in surface and subsurface geologic systems (Napier et al. 2005; Gupta and Walther 2014; Walther and Gupta 2015).

2 Natural Occurrence and Exposures

All types of soil and rocks contain uranium, the concentration of which is ranging from 0.003 mg kg^{-1} in meteorites to 120 mg kg^{-1} in phosphate rock. As a normal constituent of earth's crust, rock phosphate (U resources in phosphate rocks are estimated at $9 \times 10^6 \text{ MT}$) deposits usually contain several million tons of uranium (U), radium (Ra), and thorium (Th) which leads to potential exposure of the environment to radiation (Gupta et al. 2014). Commonly studied U decay series involves the following steps:

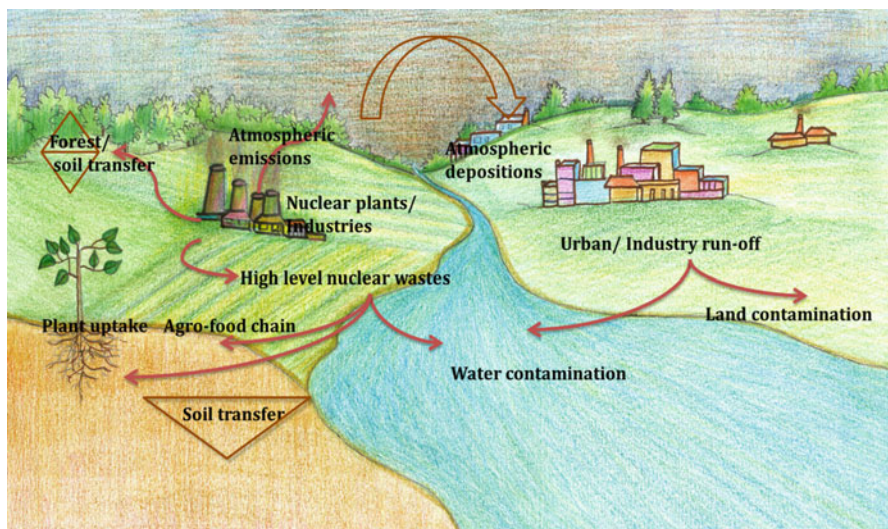
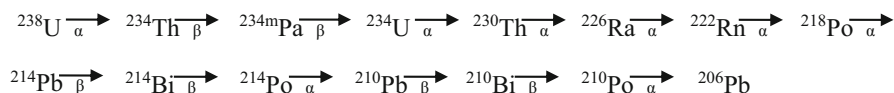


Fig. 1 Potential radionuclide emissions and contamination pathways in the environment. Incorporation of radioactive materials may occur either as NORM (Naturally Occurring Radioactive Materials) or TENORM (Technologically Enhanced Naturally Occurring Radioactive Materials). Radioactive materials once enter into the environment gets absorbed into the soil or adsorbed to the different soil organic matter or mixed up with the water sources. Atmospheric emissions from different nuclear power plants, industries lead to deposition and pollution of the environment. Plants usually take up such elements leading to incorporation in the food chain. However, transfer factor of the radionuclides depend upon several factors including soil quality, texture and type, organic/inorganic materials, availability of elements etc



High levels of dose rate are found above soils predominated by granite rocks or mineral sand in comparison to the other soil types. Inhabitants of communities settled on this type of soil (like monazite based sand beaches of Espirito Santo and Rio de Janeiro in Brazil) receive doses many times the average global radiation level (Eisenbud and Gesell 1997). Additional sources of radiation are up-concentrated when earth crust products (oil, coal, coal ash, minerals, and phosphate based fertilizers etc.) are extracted refined or used, which is called naturally-occurring radioactive materials (NORM). Anthropogenic activities like sewage sludge treatment, mining of radioactive materials, concentrate or expose radioactive materials that occur naturally in ores, soils, water, or other natural materials are the sources of production of NORM (IAEA 2014) or technologically enhanced NORM. Radionuclides of concern found in NORM mainly include isotopes of U, Th, Ra, Rn, Pb and Po. Reports suggest that, based on samples collected from 15 different countries, the average concentrations of ${}^{40}\text{K}$, ${}^{238}\text{U}$ and ${}^{232}\text{Th}$, in coal are 50, 20, and 20 Bq kg^{-1} (1.35, 0.54, and 0.54 nCikg^{-1}), respectively

(however, the concentration may vary considerably in different coal mines). Similarly, crude oil and natural gases are another source of radiation, where, approximately (average estimation), 40 pCi L^{-1} (1.5 Bq L^{-1}) of Ra may be found at well heads of natural gas. Further, LPG (liquefied petroleum gas) processing and blending tends to enhance concentrations of radon and its products ^{210}Pb , ^{210}Po (Eisenbud and Gesell 1997). Uranium contamination in soil and water has been increased in a number of regions throughout the world due to technological progressions related to establishment of nuclear power plants and its fuel cycle (e.g., mining and milling of U ore and its waste), combustion of fossil fuels (e.g., coal), increased production of phosphate fertilizers through phosphate rock mining and other uses for warheads and present as in the form of TENORM, which leads to potential ecological risks to the biota through radiation as well as chemical toxicity (Sheppard 2001).

Phosphogypsum (mostly calcium sulfate) is a by-product of a wet-acid based process during the generation of phosphoric acid in the phosphate based fertilizer industry. Large quantities of deposits of phosphogypsum are a potential source of enhanced natural radiation and for heavy metals. Approximately 70 % of calcium phosphate along with different impurities like radionuclides is present in high grade ores of phosphate rock (PR) for the production of phosphoric acid and those impurities ultimately end up in the phosphogypsum. USEPA (<http://www.epa.gov/radiation/neshaps/subparttr/about.html>) reports suggest the concentration of ^{238}U and ^{226}Ra in phosphogypsum of central Florida were about 10 times for U and 60 times for ^{226}Ra higher than the average background levels in soil. The concentration may vary according to the geographical area and quality of the PR ore. Mineral sands (sands with a specific gravity of more than 2.9), produced through the erosion of rocks (naturally or anthropogenically through blasting etc.), are also a major source of ^{232}Th and ^{238}U (UNSCEAR 1993: <http://www.unscear.org/unscear/en/publications/1993.html>).

3 Use of Green Plants for Restoring Balance and Its Applications

Phytoremediation (Ancient Greek: phyto- ‘plant,’ and Latin remedium- ‘restoring balance’), according to a recent definition by Landmeyer (2011), is the “application of plant-controlled interactions with groundwater and organic and inorganic molecules at contaminated sites to achieve site-specific remedial goals”. Cleaning up of the environment through plants is rendered by diverse environmental pollution problems through direct uptake of toxic chemical(s), followed by subsequent transformation, transport, and their accumulation in less toxic forms (Schnoor et al. 1995). Furthermore, remediation processes are being improved by plants, roots release of exudates and enzymes that induce microbial diversity at

rhizosphere and biochemical activity in the bulk soil and mineralization (Macek et al. 2000; Chatterjee et al. 2013b).

Phytoremediation techniques are becoming a popular alternative to the conventional energy and instrument intensive, chemical-based expensive restoration techniques of vast polluted areas of land and water (Padmavathiamma and Li 2007; Lone et al. 2008). Use of green living systems for environmental remediation was first reported in 1948 for accumulation of nickel by the plant *Alyssum bertolonii*; however, the concept received momentum later. Since the last two decades phytoremediation work has got much attention throughout the globe (Gupta 2013).

After the reactor accident at Chernobyl, Ukraine in 1986 that caused the release of large quantities of radioactive particles into the environment and resulted in some 30 deaths mostly due to deterministic radiation effects and more due to long-term (stochastic) effects like thyroid cancers and leukaemia in the case of clean-up workers (UNSER report on 'The Chernobyl accident' from- <http://www.unscear.org/unscear/en/chernobyl.html>), the scientific community of the world started thinking to use plants for radionuclide decontamination of soil and water. In 1998, Consolidated Growers and Processors (CGP), PHYTOTECH, and the Ukraine's Institute of Bast Crops, employed the phytoremediation process using industrial hemp (*Cannabis*) as most efficient plant useful for eliminating toxins such as metals, solvents, pesticides, explosives etc. from the contaminated topsoil. Continuing to this application of plant based green technology, US Department of Defense and USEPA jointly developed plant-based clean-up approaches to large-scale clean-up projects, where, specific plants were used to remove toxins and certain metals (Rai and Pal 1999).

Use of plants for removal of inorganic components or toxic elements like Hg, Cd, As, Cr, Cs, Pb and Sr involves extraction and translocation of toxic cations or oxyanions to above ground tissues by converting the element to a less toxic chemical species (Meagher 2000; Chatterjee et al. 2012). However, for organic compounds like polychlorinated biphenyl (PCBs), polycyclic aromatic hydrocarbons (PAH), dioxin, and trichloroethylene, the objective of phytoremediation processes is to totally mineralize them into relatively nontoxic constituents, such as carbon dioxides, nitrate and ammonia (Cunningham et al. 1996). Plants deal with such components through the strategies of stabilization, exclusion, detoxification and/or their storage in specific cells or cell organelles (vacuoles, cell walls). These strategies are variable with regard to the component characteristics and the processes termed as: phytostabilization, phytoextraction, phytovolatilization, rhizofiltration, phytodegradation, and phytostimulation (Marques et al. 2009; Chatterjee et al. 2013b). Furthermore, plants also develop defence mechanisms through the production of different anti-stress proteins like metallothioneins and phytochelatins etc.

Hyperaccumulator plants, that actively uptake exceedingly large amounts of one or more heavy metals (at concentrations 100–1000-fold higher than those found in non-hyperaccumulating species) from the soil, without exhibiting any symptoms of phytotoxicity, are especially suited candidates for phytoremediation (Reeves 2006). Plants belonging to the families of Brassicaceae, Papilionaceae, Caryophyllaceae,

Poaceae and Asteraceae are most important and offer best potential for heavy metal phytoremediation. Amongst these, species belonging to the Asteraceae family show bio-removal potential of heavy metals and radionuclides, such as Sr, Cs and U. Plants like *Lactuca sativa* L., *Silybum marianum* Gaertn., *Centaurea cyanus* L., *Carthamus tinctorius* L. from Asteraceae, can uptake ^{134}Cs very efficiently; high biomass producing varieties of *Helianthus annuus* and *H. tuberosus* are also suitable for the phytoremediation of polluted sites (Tang and Willey 2003). Phytoremediation of radionuclides has many advantages over the traditional treatments. Firstly, in phytoremediation the soil is treated in situ, which does not cause further disruption to the soil dynamics. Secondly, plants are established, they remain for consecutive harvests to continually remove the contaminants. Last but not least, phytoremediation reduces the time of workers who are supposed to expose to radionuclides. Finally, phytoremediation can be used as a long term treatment that can provide an affordable way to restore radionuclide contaminated areas (Gupta and Walther 2014). Table 1 summarizes the successful used radionuclide phytoremediator plant species (aquatic/terrestrial).

4 Rhizosphere, Rhizobacteria and Metal Transporters Used in Remediation Purposes

Plant roots secrete exudates in the adjacent soil matrix at the rhizosphere (region of soil at the root-soil interface, which is subjected to root secretions and related soil microorganisms). This mechanism helps in metal-chelation to control the entry of metal within plant through the root cell. Plants root exudates normally increase the abundance of soil microflora (bacterial and fungal communities) by 1-4 orders of magnitude compared to the surrounding bulk soil that helps to degrade diverse pollutants (Anderson et al. 1994). Root exudates include: diffusates (e.g. amino or organic acids, water, inorganic ions, and sugars etc.), excretions (e.g. bicarbonates, protons, and carbon dioxide etc.), secretions (e.g. mucilage, siderophores, and allelopathic compounds etc.) which help to modulate soil microflora community composition to have a better range of metabolic capabilities with unique gene pool (Hall 2002; LeDuc and Terry 2005). The availability of U and ^{137}Cs to plants can be enhanced using citric acid and ammonium nitrate, respectively, in soil. However, soil augmentation may be done in a properly managed fashion, as for the inherent risks associated with the application which may in turn contaminate soil or ground-water (Prasad 2011). Plant cell walls operate as cation exchanger, sharing or excluding diverse metals, to retain the homeostasis in the cell (Rauser 1999).

On the other hand, plant growth-promoting rhizobacteria (PGPR) play a crucial role in plants growth by countering physiological stress in contaminated soils. PGPR generally performs rhizospheric colonization by stimulating the production of plant growth regulators like indole acetic acid, gibberellic acid, ethylene and cytokinins contributing to plant health and better response. Further, rhizobacteria

Table 1 Selected plant list which are successful used as a Cs, U and Ra phytoremediator species. (After Eapen et al. 2007; Chakraborty et al. 2013; Favas and Pratas 2013; Fukuda et al. 2014; Hu et al. 2014; Favas et al. 2014)

Plant species	Family	Radionuclide	Habitat
<i>Amaranthus retroflexus</i>	Amaranthaceae	^{137}Cs	Terrestrial
<i>Apium nodiflorum</i>	Apiaceae	U	Aquatic
<i>Atriplex canescens</i>	Amaranthaceae	^{226}Ra	T
<i>Batrachospermum virgato-decaisneanum</i>	Batrachospermaceae	^{137}Cs	A
<i>Bertholletia excelsa</i>	Lecythidaceae	^{226}Ra	T
<i>Brassica juncea</i>	Brassicaceae	^{137}Cs	T
<i>Brassica oleracea</i>	Brassicaceae	^{137}Cs	T
<i>Callitriche stagnalis</i>	Plantaginaceae	U	A
<i>Carex buxbaumii</i>	Cyperaceae	^{226}Ra	T
<i>Carthamus tinctorius</i>	Asteraceae	^{134}Cs	T
<i>Centaurea acyanus</i>	Asteraceae	^{134}Cs	T
<i>Chloroidium saccharophilum</i>	Chlorophyta	^{137}Cs	T
<i>Dryopteris scottii</i>	Dryopteridaceae	^{226}Ra	T
<i>Eleocharis dulcis</i>	Cyperaceae	U	A
<i>Fontinalis antipyretica</i>	Fontinalaceae	U	A
<i>Helianthus annuus</i>	Asteraceae	U	T
<i>Helianthus tuberosus</i>	Asteraceae	U	T
<i>Hydrilla verticillata</i>	Hydrocharitaceae	U	A
<i>Iris pseudacorus</i>	Iridaceae	^{226}Ra	T
<i>Juncus inflexus</i>	Juncaceae	^{226}Ra	T
<i>Lactuca sativa</i>	Asteraceae	^{134}Cs	T
<i>Lemna aoukikusa</i>	Tracheophyta	^{137}Cs , U	A
<i>Lemna gibba</i>	Araceae	U	A
<i>Lemna minor</i>	Araceae	U	A
<i>Nymphaea violacea</i>	Nymphaeaceae	^{137}Cs	A
<i>Phalaris arundinacea</i>	Poaceae	^{137}Cs	T
<i>Phaseolus acutifolius</i>	Fabaceae	U	T
<i>Phragmites australis</i>	Poaceae	^{228}Th , ^{226}Ra	T
<i>Pteridium aquilinum</i>	Pteridaceae	^{226}Ra	T
<i>Pteris multifida</i>	Pteridaceae	^{226}Ra	T
<i>Silybum marianum</i>	Asteraceae	^{134}Cs	T
<i>Typha latifolia</i>	Typhaceae	U, ^{226}Ra	A
<i>Zostera japonica</i>	Zosteraceae	U	A
<i>Zostera marina</i>	Zosteraceae	U	A

may also secrete antibiotics, phosphate solubilising substances, hydrocyanic acid, siderophores, 1-aminocyclopropane-1-carboxylic acid (ACC) to increase bioavailability and root absorption of different metals (Meyer 2000; Davies et al. 2001). In a nickel-contaminated soil, Burd et al. (1998) reported that after the addition of *Kluyvera ascorbata* SUD165/26 an associated rhizobacteria, germination and

growth of Indian mustard (*Brassica juncea*) seeds increase by 50–100 % with respect to the control plants.

Root cell plasma membranes harbour a number of metal transporter-proteins which play important roles in heavy metal homeostasis. Among different families of transporters, NRAMP (natural resistance-associated macrophage protein), ZIP (Zinc importer) families (ZRT, IRT-like Protein; [ZRT-Zinc regulated transporter, IRT-iron regulated transporter]), CDF (cation diffusion facilitator) family, heavy metal ATPases (HMAs) family like P1B-ATPases, copper transporter (COPT) family proteins, ATP-binding cassette (ABC) transporters, ABC transporters of the mitochondria (ATM), Ca^{2+} cation antiporter (CAX), multidrug resistance-associated proteins (MRP), yellow-stripe-like (YSL) pleiotropic drug resistance (PDR) transporters are well studied transporters (Dubey 2011; Huang et al. 2012; Gupta et al. 2013). Histidine-rich domain of ZIP family transporters is supposed to get activated in response to divalent metals and their uptake. As for example, AtZIP4 (*Arabidopsis thaliana* ZIP4) proteins help in Zn transport and Cd uptake from soil to root cells and Cd transport from root to shoot (Krämer et al. 2007). While, transport of Fe, and other metals like Mn^{2+} , Zn^{2+} , and Cd^{2+} in root cells was being carried out by IRT1 in *A. thaliana* (Nishida et al. 2008). HMAs family transporters (P1B-type ATPases) are basically internal transporters that load Cd and Zn metals from the surrounding tissues into the xylem and perform as an efflux pump (Krämer et al. 2007). AtHMA3 transporter of tonoplast membrane in *A. thaliana* sequesters a wide variety of heavy metals and its over-expression raises the tolerance to heavy metals like Cd, Co, Pb, and Zn (Manara 2012; Gupta et al. 2013). Strategies for plants to isolate metals from dynamic cytosol and metabolically active cellular compartments include vacuolar sequestration using proton pumps like vacuolar proton-ATPase (V-ATPase) and vacuolar proton-pyrophosphatase (V-PPase) (Dalcorso et al. 2010).

5 Radio-Phytoremediation: An Approach to Decontaminate Radionuclides Using Plants

Radionuclides and/or toxic elements contaminated soils, sediments, surface water and groundwater's remediation by using green plants is widely reported (Nishita et al. 1958; IAEA 1989; Soudek et al. 2004, 2006). However, radionuclide decontamination has its inherent problem of radioactivity itself. Plants require a long period of time to contact with a contaminant to evolve the ability to hyperaccumulate radioactive material (Fig. 2). The concentrations, mobility, and bioavailability of radionuclides depend upon several factors. These include the quality, quantity and the rate of release of radionuclides present at the source; hydrological factors, like dispersion, advection, and dilution; geochemical processes, such as complexation at aqueous phase, pH, adsorption/desorption, solid/liquid distribution coefficient, reduction/oxidation (redox), ion exchange,

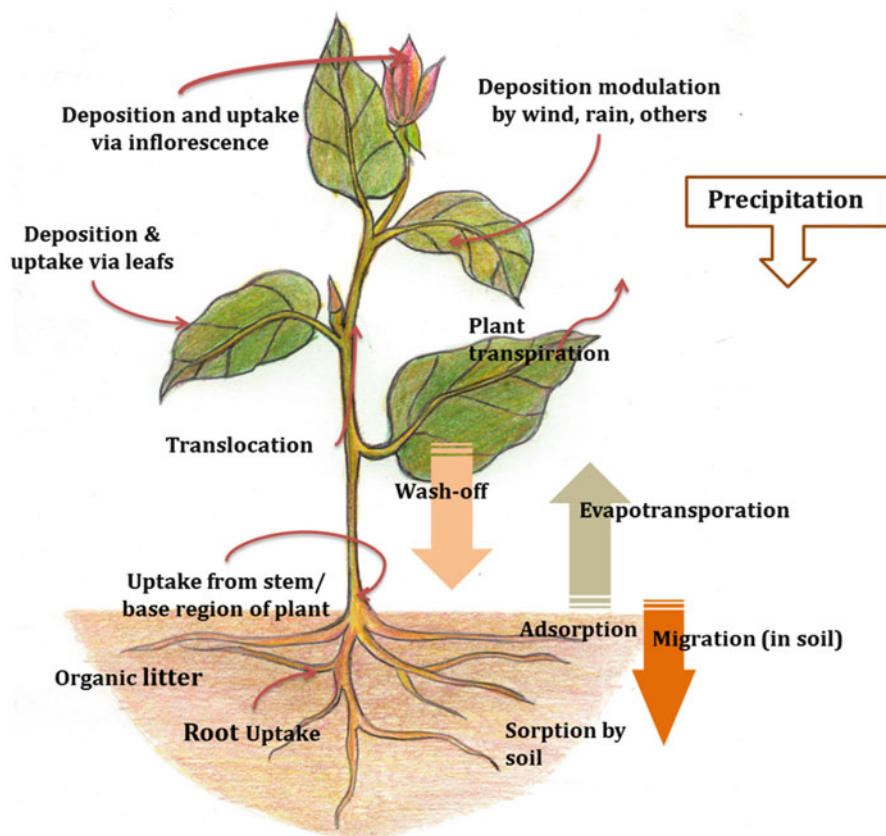


Fig. 2 Radionuclide contamination pathways in plants. The root uptake of radionuclide depends upon various factors like the condition of metal with clay/soil particle/minerals and its availability to the plants. The translocation and deposition of the same may take place in the leaf or inflorescence. Again, metals present in the atmosphere also get deposited on different plant body surfaces (like leaf surfaces). The concentration of deposition varies and modulated through the factors like wind, rain etc. Wash-off metals further deposited into soil and sorbed by soil particles, and available for plants uptake

precipitation/dissolution, diffusion, colloid-facilitated transport, exchangeable potassium ion distribution, anion exclusion and organic matter contents (Albrecht et al. 2002; Napier et al. 2012; Chakraborty et al. 2013; Hegazy et al. 2013). Absorption and distribution of contamination in plants may take place either through direct (exposures at aerial organs like leaf, stem, and tendrils) or indirect (through root systems in soil related contamination) routes, which varies considerably in different plant species especially in case of long-lived radionuclides (Din et al. 2010). Further, soil properties/texture (like drying and subsequent cracking of soils) due to biological activity, colloid-facilitated transport in soil may augment the mobility and/or affectivity of radionuclides (Napier et al. 2012).

In the environment, a number of radionuclides (e.g. Tc, U and Pu) may be present in more than one oxidation state, and adsorption and precipitation behaviour of individual states may vary considerably. Thus, transfer pathways and efficiencies of radionuclides vary widely according to (1) chemical nature and reactivity of the isotope that may affect the solubility and transport of the isotope through soil pore (includes pore structures) water within the root (rhizosphere) of the plant, (2) exposure route (e.g. root versus foliar exposure) of the contaminant, (3) plant itself (species, physical stature, age, root structure, and root-shoot ratio etc.), (4) requirements and availability of nutrient(s) of the plant (chemical resemblance of the isotope to a nutrient) and (5) level of toxins and pathogens in the soil.

It is depending upon the water cycle and the compositions of the flora which are intrinsically coupled, thus interactions may occur within these water bodies or in terrestrial ecosystems. However, the distribution and productivity of terrestrial vegetation is mostly dependent upon the equilibrium of water. Plant communities and root zone (rhizosphere) of the plant along with the soil microorganisms play important roles in evapo-transpiration generation of runoffs (Gerten et al. 2004), metal uptake and remediation. Ehlken and Kirchner (2002) suggested that the concentration of a radionuclide in a plant or plant part is linearly related to its concentration in soil within the rhizosphere (same kind of relation is also true for transfer of heavy metals and pesticides in plants) (Gast et al. 1988; Trapp et al. 1990). However, interestingly, soil-to-plant transfer factors for a number of long-lived radionuclides (as for example, radiocaesium uptake from agro-soil) may vary considerably up to three orders of magnitude (Frissel 1992; Nisbet and Woodman 2000; Ehlken and Kirchner 2002). Again significant variation is also present in case of longevity of plant roots in different plant species, their development and function is being responded by genetically determined range to their environment (Clausnitzer and Hopmans 1994).

6 Bioavailability of Radionuclides

In soil, most of the elements are usually present, from where plants acquire essential elements as micronutrients. Bioavailability of trace substances in soil is a complex phenomenon, where competitions of varied substances along with metal-metal interactions take place. Transfer dynamics from soil to plant is usually calculated for trace substances whose behaviour in the soil-plant-rhizosphere system largely depends on the concentrations of macro-nutrients present. It is reported that an activity concentration in a soil solution of 1 BqL^{-1} of ^{90}Sr or ^{137}Cs corresponds to ca. $2 \times 10^{-15} \text{ MolesL}^{-1}$ though the average concentrations of chemical homologues K, Ca, and Mg, respectively, in soil solution are in the order of 1 mMoles L^{-1} (Robson and Pitman 1983; Ehlken and Kirchner 2002). Uptake of radionuclides like radioactive caesium and strontium and heavy metals by plants is thus affected by several factors naturally present in the soil (Lorenz et al. 1994). Cation exchange reactions in the soil milieu determine the actual concentration of

an ion in soil-solution which is competitive in nature. Accordingly, co-precipitation also depends on the concentrations of competing substances in solution (as for example, sorption of radiostrontium is dominated by reversible exchange with major cations like Ca^{2+}). While strontium is exchanged in preference to Ca in minerals, the preference of the same metal reverses in the presence of organic matter (Valcke and Cremers 1994). The sorption of radiocaesium in soils is determined by ion exchange to a few sites, as for example, weathered mica which are accessible only to poorly hydrated cations and shows high selectivity for Cs^+ over K^+ and NH_4^+ (Comans and Hockley 1992; Ehlken and Kirchner 2002). In a recent report by Voronina et al. (2015), they reported a comparative study of selectivity and reversibility of radiocaesium and radiostrontium sorption by natural aluminosilicates (glauconite and clinoptilolite) as well as modified ferrocyanide sorbents, it was shown from their experiment that modification of natural aluminosilicates by ferrocyanides allows to increase distribution coefficients of caesium by 100–1000 times, to improve sorption capacity and to make sorption of caesium more selective and almost irreversible. They recommended the use of modified aluminosilicates for remediation of radioactively contaminated lands.

Some metals (like Zn, Cu) are essential for plant growth and sustenance, however, higher levels of essential or non-essential metals cause toxicity and inhibition of growth in most of the plants (Hall 2002; Lasat 2002). Absorption in plants is a vital issue, where the root zone (rhizosphere) region along with soil microorganisms interacts with the elements for bio-availability and uptake of the same (Wenzel et al. 2003). Elements are typically co-transported across the plasma membrane in the form of cations. Toxic heavy metals which do not have any known biological function (viz. As, Cd, and Pb) are also transported through the common transportation system (Manara 2012). Generally- in plants, root secrete exudates in adjacent soil matrix which may help in chelation of unwanted metals and to prevent transportation of these metals inside the cell (Marschner 1995). As for example, root exudates like histidine (His) prevent Ni uptake from the soil (Salt et al. 2000), whereas, pectic sites and different extracellular carbohydrate molecules present on the cell wall play an important role in immobilization of toxic heavy metal ions (Manara 2012). In metal contaminated soil, plants act upon as either “accumulators” or “excluders” for particular metal(s), where the excluders restrict metal uptake into their biomass and accumulators amass (high concentration) metals or contaminants into their aerial tissues like leaf tissues after breakdown of the same (Tangahu et al. 2011). However, chemical forms of metals may vary as bound component to soil particle and/or precipitated form, which mostly make a major portion of the metal(s) that is insoluble or unavailable to plants (Chatterjee et al. 2013a). Several factors involve in controlling uptake, translocation, and storage of different metals including toxic elements into the plant body. In soil microorganisms, plant induce pH changes and redox reactions, plants also produce chelating agents and exudates and a group of transporters embedded in plant cell plasma membrane, it is the major regulator for several functions. The metal transporter proteins like proton pumps –ATPases (that generate electrochemical gradients and consume energy), co-and anti-transporters proteins (that use the

electrochemical gradients generated by $-ATPases$), and ion channel proteins (that facilitate the transport of ions into the cell) involved in metal uptake and translocation (Tangahu et al. 2011).

7 Radionuclide Exposure and Plant Responses

Plants have been observed to take up many cations present in their root region irrespective of biological requirement. When plants are exposed to ionizing radiation, molecular and cellular effects are induced directly through damage of macromolecules or indirectly through water radiolytic reactions producing reactive oxygen species (ROS) (Gupta and Walther 2014). By energy transfer from the radiation field to plant tissue, ionizing radiation can directly induce DNA strand breaks, lipid oxidation, or enzyme denaturation. ROS are also produced under natural metabolism and their functions in plants are as signaling molecules, regulating normal growth and development and they are useful in stress responses. Plants also possess an antioxidative defense system comprising enzymes (e.g., superoxide dismutase (SOD) and catalase (CAT)) and metabolites (e.g., ascorbate and glutathione) to regulate the amount of ROS in cells (Gupta and Sandalio 2012). Plant metal tolerance mechanisms requires the coordination of complex physiological and biochemical processes. Plants employ various strategies to cope with toxic effects of radionuclides and heavy metals or metalloids, in general, and radionuclides stress either by avoidance (restricting the metal uptake), or by tolerance (survive in the presence of high internal metal concentration). Plants also restrict metal stress by the mechanisms like reducing the concentration of metal entering into the cell by extracellular precipitation, biosorption to cell walls, reduced uptake, and/or by increased efflux (Gupta and Walther 2014). However, tolerating metal stress involves elaborate physiological responses like intracellular chelation through synthesis of amino acids, organic acids, glutathione (GSH), and/or by metal-binding ligands such as metallothioneins (MTs) and phytochelatins (PCs), vacuolar compartmentalization, and up-regulation of antioxidant defense and glyoxalase systems to counter the deleterious effects caused by ROS (Gupta and Sandalio 2012).

8 Phytoremediation of Selected Radionuclides

8.1 *Caesium*

Caesium (Cs) has the chemical similarity to potassium (K) and thus absorption and translocation by plants is common. Contamination of Cs (^{134}Cs half-life: $T_{1/2} = 2.06$ years and ^{137}Cs - $T_{1/2} = 30.04$ years) leads to long persistence

in soil and environment (Koarashi et al. 2012; Kamei-Ishikawaa et al. 2013). This alkaline metal is present in solution as free hydrated cation Cs^+ . However, plant uptake of Cs was first reported by Collander in early 1940s (Collander 1941) and was thought to be coupled as a nutrient analogue with the same K^+ uptake transporter, especially during low K concentrations (Menzel 1954; Zhu and Smolders 2000). It is apparent that K^+ is the most significant cation amongst all other alkaline metals and compounds that competes with Cs^+ uptake (Zhu and Shaw 2000). Plant families such as Brassicaceae, Amaranthaceae and Chenopodiaceae are the major Cs accumulators. For plant uptake of Cs, K^+ concentration in soil is the most important factor. Thus, both selection of plant and potassium concentration in the soil should be taken into consideration for phytoremediation of Cs from soil (Buysse et al. 1996).

Cs uptake in plants is not yet been fully elucidated on a microscopic level. However, it has been observed that, low K concentration in soil helps to uptake Cs through K uptake system. Thus, higher K concentration suppresses the Cs uptake (Zhu and Smolders 2000). Studies on discrimination factor on Cs/K revealed that plant roots usually absorb Cs less ably than K like nutrient analogue (Smolders et al. 1996). Two K^+ transport pathways (K^+ channels and K^+ transporters) are involved with Cs^+ transportation, where multiple genes may be involved. As for example, in *A. thaliana*, high affinity K^+ transporter family (e.g. AtHAK5) and voltage-insensitive cyclic nucleotide gated channels, AtCNGCs, are involved in the Cs^+ uptake (Kanter et al. 2010; Kobayashi et al. 2010; Yamashita et al. 2014).

Specific absorption mechanism in soil by plant may depend upon several factors like availability and exchangeability of Cs along with the particular salt content and its interionic effects, soil type, water submergence time etc. (Tensho et al. 1961, Mimura et al. 2001). Salts like ammonium molybdophosphate (AMP) and rubidium chloride augment in uptake of Cs from soil. The application of AMP for partitioning and recovery of radioactive ^{137}Cs from HNO_3 and NaNO_3 rich liquid waste is well documented (Mimura et al. 2001).

However, a number of conditions and physiological parameters of plants are involved that lead to differential uptake in different species. Among these factors, plants demand for potassium and its rooting pattern for its growth strategies are most important, along with the factors like presence of mycorrhizal fungi, rate of root growth, efficiency of ion transporters that are present on plasma membranes of root cells etc. (Broadley and Willey 1997; Zhu and Smolders 2000). Halophytes of family Chenopodiaceae are long been known as Cs accumulators, however, their capability to discriminate K and Na from Cs for uptake from salty soil is still discussed controversially (Flowers et al. 1986; Zhu and Smolders 2000).

To understand the genetic variations in plants for uptake of Cs is very important for selection of suitable crops grown in soils and their accumulation pattern within the edible parts. This kind of data may be helpful to minimize its transfer to food chain. Further, non-edible plants may also be explored to decontaminate the soil contaminated with this radionuclide (Zhu and Smolders 2000). Reports showed that red root pigweed (*Amaranthus retroflexus*) is a good accumulator of ^{137}Cs with higher uptake rate and plant growth, as also shown during the phytoremediation practices for Cs-contaminated soil in the vicinity of Chernobyl, Ukraine

(Lasat et al. 1997, 1998; Dushenkov et al. 1999). Kang et al. (2012) showed that napier grass (*Pennisetum purpureum* Schum.) under hydroponic condition is also a good accumulator of Cs and may be applied for phytoremediation. To reduce the total time required for phytoremediation, various chemical and biological amendments may be useful in speeding up phytoextraction process. Reports suggest that the application of NH_4NO_3 , $(\text{NH}_4)_2\text{SO}_4$ could increase phytoextraction, while application of potassium based components to the soil should be minimized (Lasat et al. 1997, 1998; Dushenkov et al. 1999; Zhu and Smolders 2000). Further, genetic alteration and special plant breeding by selection of suitable plant taxa may be helpful to achieve the goal.

In a recent report by Yamashita et al. (2014) on the deposition of radionuclides (radiocesium- ^{134}Cs , ^{137}Cs) in different plant species after the accident of the Fukushima Daiichi Nuclear Power Station, Japan in March 2011, it is shown that the transfer factors of radiocesium from soil to plant ($[\text{Cs}]_{\text{plant}}/[\text{Cs}]_{\text{soil}}$) vary widely in different plants. However, plants like *Athyrium myosotis*, *Dryopteris tokioensis* and *Cyperus brevifolius* exhibited relatively high TF values (more than 0.4).

8.2 Uranium

U has six naturally known isotopes ^{233}U to ^{238}U with half-lives varying between 69 years and 4.47 billion years. Among three primordial natural isotopes, ^{238}U is the most abundant isotope (~99.275 % of the uranium found in nature) followed by ^{235}U (~0.720 %), and ^{234}U (~0.005 %). Radioactivity and metal toxicity of U makes it a pollutant of concern to the environment. Naturally, U is released into water or soil through the weathering of rocks in the oxidized zone of the terrestrial near-surface environment. In an almost constant isotopic ratio $^{238}\text{U}(137.88):^{235}\text{U}(1)$, release of U to water takes place through natural geochemical process, while ^{234}U is produced by the radioactive decay of ^{238}U . A range of physicochemical forms like free metal ion (U^{4+} or UO_2^{2+}), inorganic compounds (e.g., uranyl carbonate or uranyl phosphate), mineralogical matrices and soil humic substances (e.g., uranyl fulvate or humate) in dissolved, colloidal, and/or particulate forms regulate the speciation and bioavailability of U (Markich 2002). Biologically dissimilatory metal reducing bacteria (DMRB) and sulfate reducing bacteria (SRB) play an important role in U(VI) reduction to U(IV) in anaerobic environments (Liu et al. 2002; Payne et al. 2002). Though U has no known essential biological functions, it is taken up by a variety of mineral and other surfaces by the presence of dissolved calcium by inducing the formation of ternary uranyl-calcium carbonate complexes. However, redox state of U distribution is a consequence of the redox potential in solution, predominantly the U(VI) to U(IV) ratio, usually regulate its solubility and potential for migration and formation of soluble UO_2 (Fredrickson et al. 2000; Stewart 2008). Organic acids like citric acid were found to be most effective in increasing U accumulation in plants. Huang et al. (1998) showed that U accumulation in shoot of *Brassica juncea* and *B. chinensis* increased from less than 5 mg kg^{-1} to more than

5000 mg kg⁻¹ in citric acid-treated soils, where soil concentration of total U was 750 mg kg⁻¹.

Chemical and civil engineering based U remediation have several apprehensions for its high economic and energy implications, chemicals that are used for the process and their fates, specificity on the characteristics of contaminant, etc. On the other hand, phytoextraction of uranium contaminated soil and water have recently gained importance as it provides a low-cost alternative to currently employed remediation procedures. Phytoremediation processes of uranium generate a minimum amount of secondary waste(s) that may otherwise produce large amounts of heavy metal laden leachate. Results of the study by Huang and his colleagues for phytoextraction of uranium through amendment of soil using citric acid, however, showed the process to be highly effective in triggering U hyperaccumulation in selected plant species (Huang et al. 1998). The application of citric acid transiently reduced the soil pH that enhances desorption of soil U leading to more root absorption. Favas and Pratas (2013) reported that plants like *Callitriche stagnalis*, *Apium nodiflorum* and *Fontinalis antipyretica* have the capacity to accumulate significant amounts of U. These authors also suggested *C. stagnalis*, *A. nodiflorum* are keystone species which can be used for different phytoremediation applications for removal of U from the contaminated sites for their efficient rooting capability, high biomass and bio-productivity and the positive correlation with the uranium concentration in waters (Favas and Pratas 2013; Pratas et al. 2012). Duckweed (*Lemna gibba* and *L. minor*) was also reported to have good ability to accumulate U in the standing water sources (Mkandawire and Dudel 2005; Favas and Pratas 2013). Further, greenhouse experiments with *B. juncea* and *H. annuus* showed the phytoremediation of uranium-contaminated soil is enhanced in *B. juncea* when the soil is augmented with citric acid buffer (pH 4.8) that caused an enhancement of the desorption of U in the soil (Dushenkov 2003; Huhle et al. 2008).

8.3 Radium

Radium (²²⁶Ra), discovered by Marie Curie and Pierre Curie in 1898, is the product of ²³⁸U decay series but is 2.7 million times more radioactive than the same molar amount of natural U due to its shorter half-life. In the last nine steps of the fourteen steps ²³⁸U decay series, Ra decay (Ra emanation) occurs producing exradio (²²²Ra gas), Ra A (²¹⁸Po), Ra B (²¹⁴Pb), Ra C (²¹⁴Bi) etc. until stable ²⁰⁶Pb is reached. Ra is a very important radionuclide due to its (inclusive isotopes ²²⁶Ra and ²²⁸Ra) presence in all three natural decay series, relatively long half-lives. It has a high mobility in different environmental conditions like uranium mining and processing industries, phosphate and gold mining areas. Ra is a well-known stressors of human body and accumulates in bone (IAEA 2002, 2014). In general, naturally four Ra isotopes are present in the environment: uranium decay series- ²²⁶Ra, thorium decay series ²²⁸Ra and ²²⁴Ra and actinium decay series ²²³Ra. Among these, ²²⁶

Ra and ^{228}Ra have half-lives of 1600 and 5.75 years, respectively (IAEA 2002, 2014). Alkaline earth metal Ra is a readily reactive element that reacts with soil and its organic matter. However, absorption of Ra by plants from root zone decreases with high soil sulphate content, increasing pH and increased exchangeable Ca (IAEA 2014). ^{226}Ra is mostly studied among its other isotopes. In plants, grown in natural environment, Ra concentrations vary from 0.0044 to 52 Bq kg⁻¹ (DW), with higher concentrations reported in Brazil nut (Simon and Ibrahim 1990).

Wetland plants like *Typha latifolia*, *Phragmites australis*, *Juncus inflexus*, *Carex buxbaumii* and, *Iris pseudacorus* showed interesting result in the accumulation of Ra from a constructed wetland contaminated with U/Ra heavy metals (Soudek et al. 2006, 2007). Macrophytic algae, members of Characeae family have also been shown as hyper-accumulators of radium, the mechanism of which needs to be ascertained properly (Kalin et al. 2002). However, Ra may be loaded into the calcium-rich lattice and/or form of BaSO₄ substitute crystals in algal tissue (Kalin et al. 2002; Kunze et al. 2007).

9 Summary and Conclusion

The two most crucial factors for successful implementation of specific plants for restoration of radionuclide contaminated sites are (1) ability to uptake the radioactive material to relatively high levels (2) without affecting the growth or high biomass production. Potential plants should usually accumulate radionuclide in the above ground parts at a level that exceed the soil concentration. There are several factors that limit the application of plant based techniques for remediation, like the rapid incorporation of the metal within/in clay/soil particle/minerals which may limits its availability to the plants. However, phytoremediation might be the most valuable easy technique for restoration of radionuclide contaminated soil if selected cultivars are used. Further, selection of plants, genetic manipulation of different transporters, elucidation of anti-stress factors in plant physiology are the main future area of comprehensive research that will prove useful to counter our own poised nuclear age.

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