# The Role of Beneficial Elements in Triggering Adaptive Responses to Environmental Stressors and Improving Plant Performance



#### Fernando Carlos Gómez-Merino and Libia Iris Trejo-Téllez

Abstract Aluminum (Al), cerium (Ce), cobalt (Co), iodine (I), lanthanum (La), sodium (Na), selenium (Se), silicon (Si), titanium (Ti), and vanadium (V) are emerging as novel biostimulants that may enhance crop productivity and nutritional quality while improving responses to environmental stimuli and stressors in some plant species. These beneficial elements are not essential for most plants, but when supplied at low dosages, they help improve their growth, development, and yield quality by stimulating different molecular, biochemical, and physiological mechanisms triggering adaptive responses to challenging environments. When plants are exposed to environmental cues such as drought, heavy metal toxicity, low temperatures, saline soils, pest insects, or pathogens, beneficial elements may induce tolerance, resistance, or defense responses that allow plants to achieve acclimation to such stressors. Enhancement of nutrient uptake, synthesis of antioxidants and osmoprotectants, stimulation of secondary metabolism and signaling cascades, and reduction of senescence are among the responses boosted by beneficial elements when applied at low dosages. Nevertheless, beneficial elements may trigger hormesis in plants, a biphasic dose response with at low-dose stimulation or beneficial effect and a high-dose inhibitory or toxic effect. Thus, when properly applied, beneficial elements may have great potential to cope with some of the most daunting challenges facing humanity, such as climate change and food production under restrictive conditions for the growing human population. In this chapter, we mainly focus on the positive effects of beneficial elements on plant performance in restrictive environments and discuss some of the challenges of using these elements as biostimulants

**Keywords** Climate change · Environmental stressors · Plant nutrition · Nonessential elements · Biostimulants · Hormesis · Innovation · Beneficial elements

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# 1 Introduction

Plants need essential elements to ensure successful growth and development during both vegetative and reproductive stages. Essential elements are classified as macronutrients and micronutrients, depending on the amounts contained in plant tissues. Macronutrients are represented by elements which are generally found in plants at concentrations greater than 0.1% of dry matter weight (DMW, >1000 mg kg<sup>-1</sup>), consisting of nitrogen (N), phosphorus (P), potassium (K), sulfur (S), calcium (Ca), and magnesium (Mg). Micronutrients are represented by chlorine (Cl), boron (B), copper (Cu), iron (Fe), manganese (Mn), molybdenum (Mo), nickel (Ni), and zinc (Zn); these nutrients are typically found at concentrations lower than 0.01% DMW, (<100 mg kg<sup>-1</sup>DMW) (Pilon-Smits et al. 2009; Alcántar-González et al. 2016). These 14 nutrients, along with the elements carbon (C), hydrogen (H), and oxygen (O), are broadly accepted as essential for all plant species (Kirkby 2012; Alcántar-González et al. 2016).

Beneficial elements are not essential for most plants, but they can promote growth and be essential for some plant species under specific conditions (Pilon-Smits et al. 2009). When supplied at low dosages, they have a favorable impact on some vital processes and can also stimulate the mechanisms of resistance to biotic and abiotic stresses or promote the uptake of other nutrients (Trejo-Téllez et al. 2016). Additionally, beneficial elements can compensate for or remedy the toxic effects of other elements, and they can also, in some cases, provide certain functions of essential nutrients, such as the maintenance of osmotic pressure (Trejo-Téllez and Gómez-Merino 2012), or induce adaptive plant responses to adverse environmental phenomena (Pilon-Smits et al. 2009).

This chapter describes the effects of the beneficial elements identified so far, that is, aluminum (Al), cerium (Ce), cobalt (Co), iodine (I), lanthanum (La), sodium (Na), selenium (Se), silicon (Si), titanium (Ti), and vanadium (V), on the physiology of plants, with special emphasis on the induction of adaptive responses to challenging environments.

## **2** The Ten Beneficial Elements

The ten beneficial elements recognized until now have been shown to improve plant growth, production, and yield quality, as well as ameliorate the responses of plant to different environmental stress factors. A summary of the main functions of these elements in plants is displayed in Fig. 1.



**Fig. 1** Overview of the mechanisms responsible for the stimulating effects of the ten beneficial elements Al, Ce, Co, I, La, Na, Se, Si, Ti, and V on plant growth and stress responses. Main groups of plants on which beneficial effects of these elements have been documented are displayed at the heading of each box text

## 2.1 Aluminum (Al)

Aluminum is the most abundant metal in the Earth's crust (comprising about 7% of its mass), and its solubility increases with decreasing soil pH (Dong et al. 2002). While in acidic soils (pH less than 5) Al may inhibit root growth and display toxic effects to plants, it can be a beneficial element for some plant taxa under certain conditions (Moreno-Alvarado et al. 2017). One of the best-known examples of the beneficial effect of Al is observed in hydrangea (*Hydrangea macrophylla* Thunb. Ser.), since supplying them with different concentrations of Al turns them from pink (50 mg kg<sup>-1</sup> DBW) to blue (4000 mg kg<sup>-1</sup> DBW), which is attributed to the formation of a colloidal complex or to the combination of Al with a pigment called delphinidin (Trejo-Téllez et al. 2016), an anthocyanin responsible for the pigments in the plant's epidermal or subepidermal cells.

In rhododendron (*Melastoma malabathricum* L.), supplying Al in the complete nutrient solution enhances root development and plant growth (Watanabe et al. 2005). In rose (*Rosa* spp.) cv. "Cherry Brandy," supplying  $Al_2(SO_4)_3$  significantly increased vase life and improved postharvest quality, because it may maintain the flower's fresh matter weight (FMW) and may increase the chlorophyll content in leaves (Jowkar et al. 2012). According to Seyf et al. (2012a), the application of

0, 150, and 300 mg  $L^{-1}$  Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub> in "Boeing" roses increased vase life from 9 to 12 and 12.3 days, respectively, and increased flower diameter compared to control plants. In lisianthus (Eustoma grandiflorum Raf. Shinn.), the application of 150 mg  $L^{-1}$ Al sulfate to flowers prolonged vase life from 8 to 15 days, and FMW continued to increase up to 8 days after the start of the experiment (Li-Jen et al. 2001). In tuberose (*Polianthes tuberosa* L.) cv."Single," the application of 50 and 100 mg  $L^{-1}$ aluminum sulfate extends the vase life to 11.5 and 12 days, respectively (Mohammadi et al. 2012a). Furthermore, Al increased protein content and reduced FMW losses. The foliar application of 0.5, 1.0, and 1.5 g  $L^{-1}$  potassium aluminum sulfate [KAl(SO<sub>4</sub>)<sub>2</sub>] in sampaguita (Jasminum sambac L.) twice a day increased vessel life and FMW (Acero et al. 2016). The application of aluminum sulfate and 8% sucrose increases the quality and durability of rose cy. "Maroussia" stems in postharvest (Basaki et al. 2013). Likewise, De la Cruz-Guzmán et al. (2007) found that treatment with 0.6 g  $L^{-1}$  Al<sub>2</sub>(SO<sub>4</sub>)<sub>3</sub>, in rose cv. "Royalty," reduces FMW loss during the vase period. Therefore, Al has great potential as a senescence retardant in cut flowers and also enhances their quality.

In medicinal plants such as chamomile (*Matricaria chamomilla*.), 60  $\mu$ M Al increases soluble phenol and flavonoid contents in shoots and free amino acids in roots (Kováčik et al. 2010), which are antioxidant compounds that help plants overcome some stress factors. In silver birch (*Betula pendula* Roth.), the application of 2 and 5 mg L<sup>-1</sup> Al enhances leaf growth (Kidd and Proctor 2000). In soybean (*Glycine max* L. Merr.) plants exposed to 1.0  $\mu$ M Cd and 150  $\mu$ M Al at pH 4.0, the malondialdehyde (MDA, a lipid peroxidation marker) content and superoxide dismutase (SOD) and peroxidase (POD) enzyme activities increased, which shows that Cd and Al are synergistic and stimulate antioxidant mechanisms (Shamsi et al. 2008). In maize (*Zea mays* L.), the application of 48  $\mu$ M Al increased leaf growth rates, as a consequence of an increase in protein synthesis and a reduction in ubiquitin-mediated proteasomal degradation of growth-repressing proteins, such as DELLA in plants, and consequently promoted growth (Conti et al. 2014; Wang et al. 2015).

# 2.2 Cerium (Ce)

Cerium levels in soil range from 2 to 150 ppm and average 50 ppm (Trejo-Téllez et al. 2016). As a beneficial element, Morales et al. (2013) reported that adding 125 mg kg<sup>-1</sup>CeO<sub>2</sub> nanoparticles (nCeO<sub>2</sub>) to cilantro (*Coriandrum sativum* L.) plants produces longer roots and increases catalase (CAT) enzyme activity in shoots and that of POD in roots.

Tomato (*Solanum lycopersicum* L.) seeds treated with  $CeO_2$  nanoparticles (<10 mg L<sup>-1</sup>) develop seedlings with more extensive root hairs than the control. However, substantially higher Ce concentrations were detected in the fruits exposed

to 10 mg  $n\text{CeO}_2\text{L}^{-1}$ , compared with controls (Wang et al. 2012), shedding light on the long-term impact of  $n\text{CeO}_2$ on plant health and its implications for our food safety and security. Importantly, second-generation seedlings grown from seeds collected from treated parent plants with  $n\text{CeO}_2$  (treated second-generation seedlings) were generally smaller and weaker, as indicated by their smaller biomass, lower water transpiration, and slightly higher reactive oxygen species content (Wang et al. 2013a).

In spinach (*Spinacia oleracea* L.) plants grown in Mg-deficient medium and treated with CeCl<sub>3</sub>, Ce stimulated the activity of nitrate reductase, nitrite reductase, glutamate dehydrogenase, glutamate synthase, urease, and glutamic-pyruvic transaminase, which are key for N metabolism, suggesting that Ce may partially replace Mg functions to transform inorganic N to organic N, but the mechanisms underlying such responses need further study (Yin et al. 2009).

In maize and mung bean (*Vigna radiata* L. Wilczek), Ce shows favorable effects on the absorption of other nutrients when applied at concentrations below 0.2  $\mu$ M (Diatloff et al. 2008). In *Arabidopsis thaliana* L. Heynh., the concentration of Ca<sup>2+</sup>in protoplasts increased by applying 0.1 mmol Ce<sup>3+</sup>, showing that this beneficial element can regulate metabolism, growth, and development through changes in the concentration of Ca<sup>2+</sup> (Liu et al. 2011). Furthermore, Ce stimulates growth of lettuce (*Lactuca sativa* L.) cv. "Regina" seedlings (Barbieri et al. 2013), while in cowpea (*Vigna unguiculata* L. Walp.), the application of 0.713–17.841  $\mu$ M cerium nitrate [Ce(NO<sub>3</sub>)<sub>3</sub>] increases chlorophyll content, relative yield, and nitrate reductase activity (Shyam and Aery 2012).

In wheat (*Triticum aestivum* L.), the application of 125, 250, and 500 mg nCeO<sub>2</sub> per kg soil improved plant growth, shoot biomass, and grain yield by 9.0%, 12.7%, and 36.6%, respectively, in comparison to the control. As well, Ce nanoparticles increased linolenic acid by up to 6.17% but decreased linoleic acid by up to 1.63%, compared to the other treatments. The findings suggest the potential of nanocerium to modify crop physiology and food quality with unknown consequences for living organisms (Rico et al. 2014). On the other hand, Wu et al. (2014) reported that the alleviation of Cd toxicity by cerium in rice (*Oryza sativa* L.) seedlings is related to improved photosynthesis, elevated antioxidant enzymes, and decreased oxidative stress.

In barley (*Hordeum vulgare* L.), the application of 500 mg kg<sup>-1</sup> nCeO<sub>2</sub> promoted plant development resulting in a 331% increase in shoot biomass compared with the control, though these plants did not form grains. Moreover, 250 mg kg<sup>-1</sup> nCeO<sub>2</sub> enhanced grain Ce accumulation by as much as 294%, with a remarkable increases in P, K, Ca, Mg, S, Fe, Zn, Cu, and Al (Rico et al. 2015).

In turfgrass (*Poa pratensis* L.) seedlings, pretreatment with Ce(NO<sub>3</sub>)<sub>3</sub> decreased the MDA content and electrolyte leakage and increased the FMW and DMW under Cu stress. Furthermore, Ce alleviated oxidative damage by regulating the metabolism of ascorbate and glutathione under Cu stress, and Ce had an important role in the acquisition of Cu tolerance in this species (Liu et al. 2016).

# 2.3 Cobalt (Co)

The Co concentration in plants normally ranges between 0.1 and 10 ppm considering DMW, although hyperaccumulator plants of the families Lamiaceae, Scrophulariaceae, Asteraceae, and Fabaceae can accumulate more than 1000 ppm of this element in leaves (Pilon-Smits et al. 2009). In higher plants, Co adheres strongly to the roots and is absorbed from the soil solution through passive transport. Since Co shows chemical similarity with nickel (Ni), it is possible that the two elements enter the cell through the same types of membrane transporter proteins (Chen et al. 2009).

Cobalt concentrations may increase under Fe deficiency, since Co can compete with Fe for the active sites of the transporter IRT1 (Baxter et al. 2008). In fact, Gad (2012) proved that a sufficient supply of Co significantly decreases the Fe content in groundnut (*Arachis hypogaea* L.) seeds, and therefore both elements are antagonists and compete for the same transporters.

At low concentrations, Co can have beneficial effects, especially in legumes. In pea (*Pisum sativum* L.), applying 8 ppm Co to the soil increased growth, nodule number and weight, plant nutrient levels, and yield and seed quality, which can be attributed to the importance of Co for *Rhizobium* populations living in the roots of these plants (Gad 2006). Co is a component of cobalamin (vitamin B12), which is required to activate enzymes related to N fixation in symbiotic microorganisms (Palit et al. 1994).

Likewise, the application of 8 ppm Co significantly increased nitrogenase activity and nodule number and weight, especially when N was applied at 75% and 100% in groundnut (Gad 2012). In the plant, Co improved growth and yield indicators and the contents of N, P, K, Mn, and Zn. Cobalt contents in seeds ranged between 2.3 and 3.5 ppm, which is within the range reported for other plants. The application of Co can contribute to a more efficient use of N, with a savings of up to 25% of the N applied (Gad 2012).

In lily (*Lilium* spp.) cv. "Star Fighter," floral stems treated with preservative solutions including 0.1 and 0.2 mM Co increased floral longevity in 61.1% and 44%, respectively, while in the cv. "Star Gazer," the application of 0.1 mM Co enhanced this variable by 19.7% (Mandujano-Piña et al. 2012). In cut marguerite (*Argyranthemum* sp.) flowers, applying 1 and 2 mM Co increased vase life by 5 days compared to the control containing only distilled water (Kazemi 2012). Cobalt and nickel (2.5 mM Co + 2 mM Ni + 2 mM salicylic acid with 2.5% sucrose) increased the vase life of lily cv. "Prato" due to improved membrane stability and reduced oxidative stress damage during flower senescence. In addition, these elements reduce the loss of anthocyanins (Kazemi and Ameri 2012). In carnation (*Dianthus caryophyllus* L.), Co retards senescence since it reduces ethylene production, an effect very similar to that shown with the application of Ni in this species (Jamali and Rahemi 2011).

In tuberose, application of 300 mg  $L^{-1}$  cobalt chloride (CoCl<sub>2</sub>) stimulated vase life (10.66 days) and water uptake (1.53 mL g<sup>-1</sup> FMW) and reduced FMW losses

(19.99 g). Moreover, the application of 400 mg  $L^{-1}$  increased the content of carotenoids (0.40 g) and proteins (31.10%) in petals (Mohammadi et al. 2012b). In tomato, the application of 50 mg kg<sup>-1</sup> Co to the soil increased the contents of N, P, K, Cu, Fe, Mn, and Zn in leaf tissue (Jayakumar et al. 2013).

In cut roses, cobalt chloride inhibited vascular blockage in the stem and maintained a high water flow rate through stems, leading to significant water uptake by flowers. The best effects were observed with 200 mg  $L^{-1}$  CoCl<sub>2</sub> in the vessel solution (Aslmoshtaghi et al. 2014).

In gladiola (*Gladiolus grandiflorus* Hort.) cv. Borrega Roja, the application of Co (0, 0.3 and 0.6 mM) significantly increased water absorption in flower stems, while the lowest percentage of weight loss was recorded in fresh rods treated with 0.3 mM Co. Furthermore, the low concentration of Co significantly increased N content in stems and leaf concentration of chlorophyll. The total dry weights of leaves and stems were higher with the treatment of 0.3 mM Co (Trejo-Téllez et al. 2014).

### 2.4 Iodine (I)

In terms of plant nutrition, iodine has been little studied in crop species (Smolen and Sady 2011b), though it can be transported from the underground (roots) to the aboveground (shoots) parts of the plants (Ashworth 2009). As a beneficial element, iodine can promote growth, induce tolerance mechanisms to cope with stress, and trigger antioxidant capacity of the plant (Medrano-Macias et al. 2016). Iodine has the ability to advance the flowering process in fruit tree species, as a result of increased photosynthetic activity, producing a greater accumulation of sugars (Landini et al. 2012).

Blasco et al. (2011) determined that applications of 40  $\mu$ M of iodate significantly improve nitrogen-use efficiency and nitrogen metabolism, which increase lettuce productivity and quality. The application of 0.05% (w/v) iodine salts in the nutrient solution caused the potato (*Solanum tuberosum* L.) tubers to absorb 272 mg 100 g<sup>-1</sup> FMW and tomato fruits 527 mg 100 g<sup>-1</sup> FMW of IO<sub>3</sub><sup>-</sup>, respectively (Caffagni et al. 2011).

After the foliar application of 25 g L<sup>-1</sup> I or soil application of 90 g kg<sup>-1</sup> I in crops grown in the open field, the maximum iodine content ranged between 9.5 and 14.3  $\mu$ g 100 g<sup>-1</sup> for plum and nectarine fruits, to 89.4 and 144.0  $\mu$ g 100 g<sup>-1</sup> for potato tubers and tomato fruits, respectively (Caffagni et al. 2012). In hydroponic culture, fresh fruits managed to accumulate up to 2423  $\mu$ g 100 g<sup>-1</sup> of iodine. In all cases, iodine was mainly accumulated in the leaves.

In spinach, the application of 1–2 mg dm<sup>-3</sup> iodine and 1 g dm<sup>-3</sup> sucrose significantly increased the content of this element, the N and the oxalate content in leaves (Smolen and Sady 2011a). Iodine synergistically improves the uptake of Mg, Na, and Ce, as well as of Fe, and reduces chromium (Cr) uptake. After the

application of 2 mg dm<sup>-3</sup> I in the soil, a greater accumulation of Na, Fe, Zn, and Al and a reduced concentration of P, S, Cu, and barium (Ba) were observed (Smolen and Sady 2011b).

The beneficial effect of foliar-applied iodine was also observed in "Golden Delicious" apple (*Malus domestica* Borkh.) trees grafted on M.9 by applying an organic-mineral liquid fertilizer (Biojodis) at a concentration of 5 L ha<sup>-1</sup> diluted in water (600 L ha<sup>-1</sup>), which improved fruit yield, diameter, and uniformity of fruits (Szwonek 2009).

In soybean seeds, the application of  $20 \ \mu M \ IO_3^-$  enhanced the expression of more proteins in comparison to the control. Furthermore, when iodine (20, 40, and 80  $\mu M \ IO_3^-$ ) was applied to the seeds, the activity of antioxidant enzymes such as SOD, ascorbate peroxidase (APX), and glutathione reductase (GR) was boosted, counteracting the toxic effects of 100 mM Cd (Gupta et al. 2015).

No negative effects of iodine fertilization (5 kg ha<sup>-1</sup> I, either as KI or as KIO<sub>3</sub>) were noted with respect to carrot (*Daucus carota* L.) yield, while higher accumulation and uptake by leaves and storage roots of iodine were obtained after the application of KI than KIO<sub>3</sub>, which improved the biofortification of carrot storage roots (Smolen et al. 2016).

A recent review on the use of iodine to biofortify and promote growth and stress tolerance in crops published by Medrano-Macias et al. (2016) stated that this element has strong interactions with Fe, Mn, Cu, and V, either directly in plant metabolism or indirectly through the microbiome of the plant. In general, good results are obtained regarding biofortification when applied to the soil as KIO<sub>3</sub> in concentrations of 7.5 kg ha<sup>-1</sup>, 10 mg kg<sup>-1</sup> soil in pots, or  $10^{-6}$ – $10^{-5}$  M in the nutrient solution. Leaf spray with KI at 0.5 kg ha<sup>-1</sup>gave good results. With higher concentrations, the response is variable: negative, neutral, or positive, depending on the plant species (Medrano-Macias et al. 2016).

#### 2.5 Lanthanum (La)

Lanthanum is considered a beneficial element as it enhances the uptake of essential nutrients such as K, Ca, and Mg (Wahid et al. 2000). In rice, the application of  $0.1 \text{ mM La}(\text{NO}_3)_3$  increased germination rate and biomass accumulation in plantlets (Liu et al. 2012a). Liu et al. (2013) found that applying 0.05 mmol La promotes root growth and that this element tends to accumulate in the cell wall of the root. In addition, La treatments affect the accumulation of K, Mg, Ca, Na, Fe, Mn, Zn, Cu, and Mo in the root and thus plant growth.

The application of 5 up to 50  $\mu$ M La in the nutrient solution stimulates the growth of maize, mung bean (*Vigna radiata* L. Wilczek), and black gram (*V. mungo* L. Wilczek) plants and the germination percentage, root length, shoot length, as well as FMW and DMW in all three crops (Chaturvedi et al. 2014). In tobacco (*Nicotiana tabacum* L.), the application of 5–20 mg L<sup>-1</sup> LaCl<sub>3</sub> in the Hoagland solution gradually increased dry matter accumulation and chlorophyll content,

which decreased when the concentration of LaCl<sub>3</sub> exceeded 50 mg L<sup>-1</sup> (Chen et al. 2001). Best results were observed with 20 mg L<sup>-1</sup>, since the synthesis and activity of choline, Mg<sup>2+</sup>-ATPase, and phosphorylation were significantly activated. Diatloff et al. (2008) reported that concentrations of La below 0.2  $\mu$ M produced positive effects on maize and mung bean, although at higher concentrations it decreases absorption of Ca, Na, Zn, and Mn.

In cucumber (*Cucumis sativus* L.),  $La^{3+}$  reduces the levels of Na, Mg, Cl, K, and Ca while increasing Mn and Fe levels. Also, the effects of  $La^{3+}$  on ion absorption show similarity to those of  $Ca^{2+}$ , indicating that La affects physiological mechanisms in the plant by regulating the levels of Ca; optimal growth occurred at a concentration of 0.02 mM  $La^{3+}$  (Zeng et al. 2000).

In cucumber plants, La has been found to be involved in ion transport and also modifies the absorption and distribution of Se, Co, V, and technetium (Tc), affecting growth and absorption of other elements that influence the physiology of cells and biochemical functions in this plant (Huang et al. 2003). In addition, Shi et al. (2005) reported that low concentrations of La (0.002 and 0.02 mM LaCl<sub>3</sub>) promote growth of cucumber plantlets and increase chlorophyll and carotenoid contents, while La was found to be involved in activating antioxidant enzymes such as POD, CAT, and SOD, as well as in reducing MDA contents.

According to Liu et al. (2012b), the Ca level decreases slightly with 0.2 mM  $La^{3+}$ ; with 1.0 mM $La^{3+}$ , oscillations of  $Ca^{2+}$  were observed; and at 2.0 mM $La^{3+}$ , there was an increase in  $Ca^{2+}$ , indicating that  $La^{3+}$  participates in signal transduction networks mediated by calmodulin (CaM) and that it can enter the root through the cell membrane and intracellular  $Ca^{2+}$  channels.

In tulip (*Tulipa gesneriana* L.) cv. "Ile de France," the diameter and length of the floral stem were higher when plants were ferti-irrigated with 10  $\mu$ M La (Ramírez-Martínez et al. 2009). In the same ornamental species, Ramírez-Martínez et al. (2012) observed that La stimulates the accumulation of Ca, K, and La itself at concentrations of 10 and 20  $\mu$ M.

Yan et al. (2007) found that adding 20 mg  $L^{-1} La^{+3}$  decreased cell membrane permeability and the contents of MDA, H<sub>2</sub>O<sub>2</sub>, and proline in soybean plants exposed to UV-B radiation (280–320 nm; 0.15–0.45 W cm<sup>-2</sup>). Moreover, the activity of the enzymes CAT and POD was higher in La-treated plants, demonstrating the activation of antioxidant mechanisms.

In a study aimed at evaluating the bioaccumulation of La and its effects on growth and mitotic index of soybean, plants were exposed to increasing concentrations of La (0, 5, 10, 20, 40, 80, and 160  $\mu$ M) in the nutrient solution for 28 days. Roots accumulated 60-fold more La than shoots. La deposition occurred mainly in cell walls and in crystals dispersed in the root cortex and in the mesophyll. The application of La resulted in increased contents of essential nutrients such as Ca, P, K, and Mn, whereas Cu and Fe levels decreased. Furthermore, low La concentrations (i.e., 5–10  $\mu$ M La) stimulated the photosynthetic rate and total chlorophyll content and led to a higher incidence of binucleate cells, resulting in a slight increase in root and shoot biomass. At higher La levels (i.e., 20–160  $\mu$ M La), soybean growth was reduced, as a result of ultrastructural modifications in the cell wall, thylakoids, and chloroplasts and the appearance of c-metaphases (de Oliveira et al. 2015).

In rangpur lime (*Citrus limonia* Osbeck), 50 mg LaCl<sub>3</sub> increased mass and height of plants with a consequent increase of dry matter, suggesting the use of La as fertilizer in citrus, especially when used at low concentrations (Turra et al. 2015).

Recently, García-Jiménez et al. (2017) reported that the application of 10  $\mu$ M to four sweet pepper (*Capsicum annuum* L.) varieties significantly increased seedling height, shoot diameter, number of flower buds, number of leaves, and leaf area, though it did not affect dry biomass accumulation. Furthermore, La stimulated the biosynthesis of chlorophyll a and b and total chlorophylls, total soluble sugars, and soluble protein concentration.

## 2.6 Selenium (Se)

In seleniferous soils, most plant species contain from 1 to 10 ppm Se, while hyperaccumulators (such as those belonging to the genera *Stanleya* and *Astragalus*) can accumulate from 1000 to 15,000 ppm Se (0.1-1.5% Se) (Pilon-Smits et al. 2009).

The application of Se at low concentrations can increase tolerance to oxidative stress induced by UV radiation, retard senescence, and promote growth. In addition, Se can regulate water content under drought conditions (Germ et al. 2007). In ryegrass (*Lolium perenne* L.), the application of 0.1 and 1 mg kg<sup>-1</sup> Se activated antioxidant mechanisms and increased glutathione peroxidase (GPX) activity while reducing senescence processes and promoting plant growth (Hartikainen et al. 2000).

In canola (*Brassica napus* L.), Se can increase seed yield and enhance its nutritional value (Hajiboland and Keivanfar 2012). In carrot, the combination of KI and Na<sub>2</sub>SeO<sub>4</sub> stimulated uptake, accumulation, and storage of both I and Se, and the consumption of 100 g FMW carrot produced by plants fertilized with KI + Na<sub>2</sub>SeO<sub>3</sub> and KIO<sub>3</sub> + Na<sub>2</sub>SeO<sub>3</sub> can provide 100% of the I and Se levels recommended for human nutrition (Smolen et al. 2016).

Because Se increases the absorption of heavy metals like lead (Pb) in common coleus (*Coleus blumei* Benth.), this beneficial element can be useful in stimulating phytoremediation mechanisms in environments contaminated by heavy metals (Yuan et al. 2013).

The foliar application of 10 mg  $L^{-1}$ Se as Na<sub>2</sub>SeO<sub>4</sub> to soybean cv. "Olna" plants increased respiration potential, especially in young plants (Mechora and Germ 2010). In melon (*Cucumis meloL*.) seedlings under salt stress, supplying 2–8 mM Se improved growth and triggered antioxidant mechanisms by inhibiting lipid peroxidation and increasing the enzymatic activity of SOD and POD (KeLing et al. 2013). Likewise, exogenous Se alleviates salt stress in maize via the improvement of photosynthetic capacity, the activities of antioxidant enzymes, and the regulation of  $Na^+$  homeostasis (Jian et al. 2017).

In maize, applying 5  $\mu$ M dm<sup>-3</sup> Se stimulated growth and root elongation (Hawrylak-Nowak 2008). Furthermore, applications of up to 15  $\mu$ M Se (as selenite or selenate) in lettuce improved plant growth, while no major changes in oxidative state, pigment concentration, and sulfur accumulation were observed (Hawrylak-Nowak 2013). Moreover, the application of Na<sub>2</sub>SeO<sub>4</sub> (0, 5 and 10  $\mu$ M) in cucumber plants treated with Cd (0, 25 and 50  $\mu$ M) reduced Cd uptake and lipid peroxidation as well, while plasma membrane was more stable, suggesting a beneficial effect of Se in plants exposed to Cd (Hawrylak-Nowak et al. 2014).

In the grass *Stylosanthes humilis* Kunth. grown in acidic soils with high concentrations of toxic aluminum  $(Al^{3+})$ , applying up to 1  $\mu$ M Se activated antioxidant mechanisms, and Se itself contributed to remove reactive oxygen species (Ribeiro et al. 2011).

Selenium has also been associated with fruit maturation. In peach cv. "Suncrest" grafted onto GF 677 (*Prunus persica* L. Batsch. × *Prunus amygdalus* Stokes) and pear (*Pyrus communis* L.) cv. "Conference" grafted onto BA 29 (*Cydonia oblonga* Mill.) receiving 0.1 and 1 mg L<sup>-1</sup> Na<sub>2</sub>SeO<sub>4</sub> in leaves or 1 mg L<sup>-1</sup> Na<sub>2</sub>SeO<sub>4</sub> in fruits, Se kept firmness and delayed fruit maturation, which prolonged fruit storage (Pezzarossa et al. 2012). Similar effects have been reported in tomato, since the application of 1 mg L<sup>-1</sup> Na<sub>2</sub>SeO<sub>4</sub> reduced lipid peroxidation, increased the activity of antioxidant enzymes such as SOD and GPX, and improved fruit quality during storage (Zhu et al. 2016).

#### 2.7 Silicon (Si)

Silicon constitutes between 0.1% and 10% of the DMW of higher plants and its accumulation can vary significantly among species. Importantly, Si-deficient plants are brittle and susceptible to fungal infections (Ma and Yamaji 2006).

Silicon can counteract the toxic effects of elements such as Al and Mn, confer resistance against pests and diseases, and even allow the formation of nanostructures using organic compounds, enzymes, or organisms as catalysts (Raya and Aguirre 2009). Silicon is absorbed in a pH range of 2–9, being taken up by the roots in the solution as monosilicic acid (Si(OH)<sub>4</sub>) to be accumulated in the epidermal cells of leaves (Borda et al. 2007).

The beneficial effects of Si are associated with its high deposition in plant tissues, improving their strength and rigidity (Ma and Yamaji 2006). It is also possible that Si plays an active role in resistance to plant diseases by stimulating defense mechanisms. Also, Si can play an important role in resistance to abiotic stress factors such as heavy metal toxicity, salinity, and drought and can reduce the generation of reactive oxygen species, due to the increased activity of antioxidant enzymes (Balakhnina and Borkowska 2013).

In forage oat (*Avena sativa* L.), applying 100 mg kg<sup>-1</sup> (116 g per pot) of monosilicic acid in the pre-sowing period increased height and dry matter production as a result of better nutritional absorption promoted by Si (Borda et al. 2007). Moreover, Si stimulated cell elongation and turgor and improved the conversion of assimilates. In bitter gourd (*Momordica charantia* L.) plants under saline stress (50 mM NaCl), the application of increasing concentrations of Si (1–5 mM) stimulated the germination rate and index, seedling vitality, and antioxidant enzyme activities of SOD, POD, and CAT (Wang et al. 2010).

By adding 100, 150, and 200 mg  $L^{-1}$  potassium silicate (K<sub>2</sub>SiO<sub>3</sub>), carnation flower cv. "Harlem" improved vase life as a result of a significant reduction in ethylene production (Jamali and Rahemi 2011). The application of 2.5 mM Si together with 3 mM acetylsalicylic acid reduces wilting in carnation flowers, retards chlorophyll and carbohydrate degradation, and reduces the activity of oxidase enzymes (Kazemi et al. 2012).

The application of Si increases the amount of  $O_2$  in leaves, stems, and roots, which causes the rhizosphere to oxidize. Thus, the elements Fe and Mn are oxidized, which prevents excessive uptake of these elements by the plant (Furcal-Beriguete and Herrera-Barrantes 2013). In common borage(*Borago officinalis* L.), Si plays a detoxifying role when the plant is under aluminum stress because it stimulates the synthesis of phenolic compounds and proline (Shahnaz et al. 2011).

In maize, application of Si by seed priming improved growth of stressed plants (exposed to alkaline stress induced by 0, 25, 50, and 75 mM Na<sub>2</sub>CO<sub>3</sub>) while enhancing the leaf relative water content and levels of photosynthetic pigments, soluble sugars, soluble proteins, total free amino acids, and K<sup>+</sup>, as well as activities of SOD, CAT, and POD enzymes. Moreover, Si supplement resulted in a decrease in the contents of proline, MDA, and Na<sup>+</sup>, which together with an enhanced K<sup>+</sup> level led to a favorable adjustment of K<sup>+</sup>/Na<sup>+</sup> ratio, in stressed plants relative to plants treated with alkaline stress alone. These findings confirm that Si plays a pivotal role in alleviating the negative effects of alkaline stress on maize (Abdel Latef and Tran 2016). Similarly, Marxen et al. (2016) showed that application of 0.4 and 17.3 t ha<sup>-1</sup> Si (as silica gel) to rice plants cv. "Khang Dan 18" increased Si contents in plant tissues, as well as biomass production and grain yield.

### 2.8 Sodium (Na)

Sodium is one of the most studied ions in plant biology due to its toxic effects, although at low concentrations its beneficial effect has also been proved. In fact, in some plants with C4 photosynthetic metabolism, Na is considered an essential element (Kronzucker et al. 2013), and its benefits are more evident in conditions

of potassium deficiency (Schulze et al. 2012). Sodium also increases the biosynthesis of amino acids, especially proline (Jouyban 2012).

Salt stress can promote growth in some crops such as wheat, while in rice the low yield caused by salinity is mainly associated with the reduction in tillers and an increase in sterile spikelets in some cultivars (Läuchli and Grattan 2007).

According to Lee and van Iersel (2008), salinity induced by NaCl has the potential to act as a growth regulator. In fact, the application of 60–120 mM NaCl increases the height of faba bean (*Vicia faba* L.) plants (Abdul Qados 2011). Importantly, salinity may lead to toughening of tomato fruit skin. Accordingly, Silva et al. (2015) reported a linear correlation between thickness of the subepidermis and salinity of the irrigation water (up to 12.61 dS m<sup>-1</sup>). Interestingly, the tougher tomato skin obtained under conditions of salinity is attributed to increased number of hypodermal cell layers rather than to changes in cell wall composition.

In a proteomic study using two genotypes of Indian mustard (*Brassica juncea* L. Czern.) displaying contrasting sensitivity to salt stress, Yousuf et al. (2016) reported differential expression of 21 salt stress-responsive proteins associated with various functional processes, including osmoregulation, photosynthesis, carbo-hydrate metabolism, ion homeostasis, protein synthesis and stabilization, energy metabolism, and antioxidant defense system. Salt-tolerant genotype (CS-52) showed a relatively higher expression of proteins involved in turgor regulation, stabilization of photosystems and proteins, and salt compartmentalization, as compared to salt-sensitive genotype (Pusa Varuna). These results suggest that modulating the expression of salt-responsive proteins can pave the way for developing salt tolerance in the Indian mustard plants.

According to Lee and van Iersel (2008), the quality of Chrysanthemum x *morifolium* Ramat. cut flower is improved by applying 1 g  $L^{-1}$ NaCl in irrigation water. In cut roses cv. "Avalanche," the application of 250 mg  $L^{-1}$  sodium benzoate improved vase life by reducing ethylene production (Imani et al. 2012). Similarly, applying 20  $\mu$ M of sodium nitroprusside (SNP) in rose and 60  $\mu$ M in sunflower (Helianthus annuus L.) increased vase life (Nazirimoghaddam et al. 2014). In cut rose cv. "Utopia" treated with 50  $\mu$ M SNP, a higher soluble protein content, an increased solution uptake rate by the flower stems, and an improved FMW ratio were observed, while vase life increased from 11 to 13.3 days compared to the control (Seyf et al. 2012b). Moreover, in giant chincherinchee (Ornithogalum saundersiae Bak.) plants grown in pots receiving either 100 or 200 mM NaCl weekly, Na increased chlorophylls and carotenoids contents, as well as N, K, Na, and Cl concentrations in leaves (Salachna et al. 2016). In purpletop vervain (Verbena bonariensis L.), the application of 200 mM NaCl enhanced Mn contents in leaves, whereas neither P, K, Mg, Cu, Zn, and Fe contents nor the initiation of flowering was affected (Salachna and Piechocki 2016).

# 2.9 Titanium (Ti)

Titanium began to gain importance in plant biology studies in the 1930s and is now considered a beneficial element (Carvajal and Alcaraz 1998). This element is not toxic to animals or to humans, and at low concentrations it is beneficial for plants since it triggers physiological mechanisms leading to better growth and development under certain environmental conditions (Jaberzadeh et al. 2013). So far, limited information is available regarding critical levels of Ti in plant toxicity.

Apart from the application of Ti as conventional reagent, the use of Ti nanoparticles (nTi) is currently gaining more importance. A recent review of Ti as a beneficial element described that seeds treated with *n*Ti suspensions exhibited increased germination rates, enhanced root lengths, or improved seedling growth in different plant species. Furthermore, the application of *n*Ti increased plant tolerance to abiotic and biotic stresses, including cold, drought, Cd toxicity, and bacterial spot disease caused by *Xanthomonas perforans* (Lyu et al. 2017).

In oat plants (*Avena sativa* L.)cv. "Zlat'ák," the effect of Ti is considerably weaker if it is applied on leaves than if being added to the nutrient solution (Kuzel et al. 2003). Importantly, the action of Ti on plant physiology can be explained by a hormetic effect. The application of 960 g ha<sup>-1</sup> Ti to tomato increases the content of N, P, Ca y Mg, and that of K with 80 g ha<sup>-1</sup> Ti,which demonstrated the importance of Ti in plant nutrition and the quality of this vegetable (Kleiber and Markiewicz 2013).

Under drought stress, application of 0.02% of titanium dioxide nanoparticles in wheat increases gluten and starch content (Jaberzadeh et al. 2013). Furthermore, the application of 1200 and 1500 mg L<sup>-1</sup>Ti in canola seedlings produced greater root development and bud growth (Mahmoodzadeh et al. 2013). In addition, TiO<sub>2</sub> has the potential to be used as an alternative for the control of bacterial blight caused by *Xanthomonas* in zonal geranium (*Pelargonium x hortorum* L. H. Bailey) and leaf spot in poinsettia (*Euphorbia pulcherrima* Willd. ex Klotzsch), since treatments using this compound at concentrations of 25 and 75 mM showed a reduction in lesions of 85% and 93%, respectively (Norman and Chen 2011).

Two applications of  $\text{TiO}_2$  at 125 cm<sup>3</sup> ha<sup>-1</sup> in cowpea in the 1st and 2nd year of production improved development and yield and reduced the severity of foliar and pod diseases compared to a single application at lower concentration. Application of TiO<sub>2</sub> increased cowpea yield from 8.74% to 36.11% and from 10.33% to 51.31%, respectively, in the 2 years of evaluation (Owolade and Ogunleti 2008).

The response of potato, barley, and wheat plants to titanium application is almost negligible under N deficiency. However, when there is sufficient N, responses are more evident (Tlustos et al. 2005). In mung bean, foliar application of 10 mg L<sup>-1</sup> TiO<sub>2</sub> increased shoot (17.0%) and root (49.6%) lengths, chlorophylls, as well as total protein contents in leaves. Furthermore, Ti application increased microbial populations in the rhizosphere and the activity of enzymes involved in the P nutrient cycle, such as acid phosphatase, alkaline phosphatase, phytase, and dehydrogenase (Raliya et al. 2015). Foliar application of Ti-ascorbate at concentrations of 25, 50,

75, and 100 mg L<sup>-1</sup> improved height of geranium cv. "Elite Cherry," petunia (*Petunia x hybrida* hort. ex E.Vilm.) cv. "Celebrity White," pansy (*Viola x wittrockiana* Gams.) cv. "Delta Premium Marina," and snapdragon (*Antirrhinum majus* L.) cv. "Montego Purple" (Whitted-Haag et al. 2014).

Just recently, Andersen et al. (2016) evaluated the application of different concentrations of Ti dioxide nanoparticles (nTiO<sub>2</sub>) on the germination of ten crop species. They reported that the application of 500 µgmL<sup>-1</sup> nTiO<sub>2</sub> increased the percentage of germination in cabbage, and root length in onion, while applying 1000 µg mL<sup>-1</sup> enhanced root length both in cucumber and onion. In barley the application of 500 or 1000 mg kg<sup>-1</sup> nTiO<sub>2</sub> to the soil stimulated plant growth and reduced the toxic effects induced by nCeO<sub>2</sub> (Marchiol et al. 2016).

It is important to note that Ti and nTi may have neutral or negative effects on plant physiology, which may be attributed to several factors including differences in plant species, physiological status of plants at the time being evaluated, seed quality, nanoparticle sizes and their uniformity, and experimental objectives and methods. Furthermore, attention does need to be given to the fate and consequence of applied nTi within the environment and food chain (Lyu et al. 2017).

#### **2.10** Vanadium (V)

The biological importance of V can be divided into three levels, depending on the daily intakes and tissue contents: nutritional (intakes of  $\mu$ g a day), pharmacological (mg a day), and toxicological (mg kg<sup>-1</sup> food DMW) (Antal et al. 2009). Because of its toxic, mutagenic, genotoxic, and even carcinogenic potential, V has been the subject of numerous public health studies worldwide (Rodríguez-Mercado and Altamirano-Lozano 2006), while its beneficial effects on plants have been sparsely addressed. In the field of plant physiology, one of those first reports on this element showed that soybean plants grown in oxisols develop normally even at concentrations of 75 mg kg<sup>-1</sup> V in the soil (Wang and Liu 1999).

Vanadium has been considered as either beneficial or as a secondary metabolism elicitor in plants, but the mechanisms involved are not yet fully understood (Saco et al. 2013). This element is a metal widely distributed both in nature and in biological systems and is also one of the trace elements present in fossil fuels (Rodríguez-Mercado and Altamirano-Lozano 2006).

Antal et al. (2009) studied 56 medicinal plant species to determine V contents, finding the highest V contents in flowering aerial parts, with an average of 763  $\mu$ g kg<sup>-1</sup>DMW, followed by the leaves (682  $\mu$ g kg<sup>-1</sup> DMW), roots (600  $\mu$ g kg<sup>-1</sup> DMW), flowers (352  $\mu$ g kg<sup>-1</sup> DMW), and fruits (112  $\mu$ g kg<sup>-1</sup> DMW). Of the plants analyzed, lemon thyme (*Thymus pulegioides* L.) displays a particular capacity to accumulate this element, while other species like *Geum urbanum* L., *Urtica dioica* L., *Hypericum perforatum* L., and *Valeriana officinalis*. also tend to accumulate this element (Antal et al. 2009).

In soybean, the application of two different sources of V increased chlorophyll contents, as well as fresh and dry biomass (Sozudogru et al. 2001). The application of 240  $\mu$ M V in common bean (*Phaseolus vulgaris* L.) cv. "Contender" caused thicker roots, where it accumulated more than in leaves (Saco et al. 2013). In mustard (*Brassica campestris* L. ssp. *chinensis* cv. "Parachinensis") and tomato (*Solanum lycopersicum* L.), Vachirapatama et al. (2011) showed that the application of 20 mgL <sup>-1</sup>NH<sub>4</sub>VO<sub>3</sub> improves growth in both species; V accumulates mostly in root in comparison to leaf, stem, or fruit. In wheat, the application of 40  $\mu$ M V effectively improved the antioxidant defense system to alleviate the oxidative damage induced by Cu (Wang et al. 2013b). In pennyroyal (*Mentha pulegium*L.), the application of 10, 20, and 40 mg L<sup>-1</sup>NH<sub>4</sub>VO<sub>3</sub> increased root DMW, while V concentration was higher in roots than in shoots. Furthermore, V did not affect K, Ca, Mn, and Zn concentrations in roots, while a reduction of Ca, K, Mg, and Mn concentration was observed in leaves at high V application rates (Akoumianaki-Ioannidou et al. 2015).

Root uptake and translocation efficiency of V do not significantly vary with the species, whereas its translocation to plant aerial parts depends on individual plant response. Future studies are necessary to determine the effect of V-status on different taxa, while the accumulation and transfer of vanadium within the food chain remain a daunting task (Qian et al. 2014).

A summary of the levels at which beneficial elements have been applied in various model and cultivated species is presented in Table 1.

# 3 Effects of Beneficial Elements on Plant Nutrition as a Mean to Overcome Abiotic Stress

In several modern plant nutrition approaches, beneficial elements are considered part of nutrient management in an increasing number of crop plants. Some of these elements, including Ce, Co, I, Na, Se, Si, and Ti, have been proved to increase crop yield. Some others such as Al and La have been less studied, and their influence on plant yield and nutrient status are little known. Cerium raises P, K, Ca, Mg, S, Fe, Zn, Cu, and Al concentrations in barley leaves and grains (Rico et al. 2015), which boosts plant growth, yield, and grain quality. Cobalt enhances N, P, K, Cu, Fe, Mn, and Zn concentrations in plant tissues not only in tomato (Javakumar et al. 2013) but also in groundnut (Gad 2012) and pea (Gad 2006), which results in increased yields and harvest quality. Iodine has been found to improve N, Mg, Na, Ce, and Fe uptake in fruits (Szwonek 2009), lettuce (Blasco et al. 2011), spinach (Smolen and Sady 2011a, b), and tomato (Caffagni et al. 2012) while enhancing productivity and crop quality. Lanthane increases Mn and Fe contents in cucumber (Zeng et al. 2000), displays a synergic effect with Ca and K uptake in tulip (Ramírez-Martínez et al. 2012), and increases Ca, P, K, and Mn concentrations in soybean (de Oliveira et al. 2015), which renders better plant growth and productivity. Selenium enhances P and Ca uptake, though it also reduces K absorption in maize (Hawrylak-Nowak 2008).

Beneficial element	Species studied	Level evaluated	Beneficial level	Application system	Reference
A	Silver birch (Betula pendula Roth.)	0, 2, 5, 10, 15, 25, and 35 mg Al(NO3)3L-1	2 and 5 mg $L^{-1}$	Nutrient solution	Kidd and Proctor (2000)
	Lisianthus (Eustoma grandiftorum Raf. Shinn.) cv. "Hei Hou"	0, 50, 100, and 150 mg $L^{-1}$ Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub>	$150 \text{ mg L}^{-1}$	Floral preservative solution	Li-Jen et al. (2001)
	Rhododendron (Melastoma malabathricum L.)	0 and 0.5 mM AlCl <sub>3</sub>	0.5 mM	Nutrient solution	Watanabe et al. (2005)
	Rose (Rosa spp.) cv. "Royalty"	0 and 600 mg L <sup>-1</sup> Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub> + 4.5% sucrose	$600 \text{ mg L}^{-1}$	Preservative solution	De la Cruz- Guzmán et al. (2007)
	Soybean (Glycine max L. Merr.)	0 and 1.0 $\mu$ M CdCl <sub>2</sub> + 150 $\mu$ M AlCl <sub>3</sub>	150 μM	Culture solution	Shamsi et al. (2008)
	Chamomile ( <i>Matricaria</i> <i>chamomilla</i> L.)	0, 60, and 120 μM AlCl <sub>3</sub>	60 µМ	Nutrient solution	Kováčik et al. (2010)
	Rose (Rosa spp.) cv. "Cherry Brandy"	$0, 100, 200, and 300 \text{ mg } \text{L}^{-1}$ Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub>	$300 \text{ mg L}^{-1}$	Preservative solution	Jowkar et al. (2012)
	Rose (Rosa hybrida) cv. "Boeing"	$0, 150, and 300 \text{ mg } \mathrm{L}^{-1}$ Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub>	150 and 300 mg $L^{-1}$	Floral preservative solution	Seyf et al. (2012a)
	Tuberose (Polianthes tuberosa L.) cv. "Single"	$0, 50, 100, \text{ and } 150 \text{ mg } \text{L}^{-1}$ Al <sub>2</sub> (SO <sub>4</sub> ) <sub>3</sub>	$100 \text{ mg L}^{-1}$	Floral preservative solution	Mohammadi et al. (2012a)
	Rosa hybrid (Rosa spp.) cv. "Maroussia"	0 and 600 mg $L^{-1}Al_2(SO_4)_3$	$600 \text{ mg } \text{L}^{-1}$ + 8% sucrose	Floral preservative solution	Basaki et al. (2013)
	Maize (Zea mays L.)	0 and 48 µM AlCl <sub>3</sub>	48 µM	Nutrient solution	Wang et al. (2015)
	Sampaguita ( <i>Jasminum</i> sambac L.)	0, 500, 1000, and 1500 mgL <sup>-1</sup> KAl(SO <sub>4</sub> ) <sub>2</sub>	0.5, 1.0,  and  1.5  g L $^{-1}$ sprayed twice a dav	Foliar application with spraying solution	Acero et al. (2016)

(continued)

Table 1   (co	intinued)				
Beneficial element	Species studied	Level evaluated	Beneficial level	Application system	Reference
Ce	Maize (Zea mays L.) cv. "Hycorn 82" and mung bean (Vigna radiata L. Wilczek) cv. "Berken"	0, 0.2, 1.0, and 5.0 μM Ce (NO <sub>3</sub> ) <sub>3</sub>	< 0.2 µM	Continuously flowing solu- tion culture units	Diatloff et al. (2008)
	Spinach (Spinacia oleracea L.)	0 and 15 µM CeCl <sub>3</sub>	15 μM CeCl <sub>3</sub>	Culture solution	Yin et al. (2009)
	Arabidopsis thaliana (L.) Heynh.	0, 0.1, 0.5 and 1 mM Ce(NO <sub>3</sub> ) <sub>3</sub>	0.1 mM	Protoplasts solution	Liu et al. (2011)
	Cowpea (Vigna unguiculata L. Walp.)	0, 0.713, 3.568, 17.841, 89.206, and 446.030 μM Ce (NO <sub>3</sub> ) <sub>3</sub>	0.713, 3.568, 17.841 μΜ	Pots with soil (silty sand)	Shyam and Aery (2012)
	Lettuce (Lactuca sativa L.) cv. "Regina"	0, 5, 10, 15, 20, and 25 mg $L^{-1}$ Ce(NH <sub>4</sub> ) <sub>2</sub> (NO <sub>3</sub> ) <sub>6</sub>	$15 \text{ mg L}^{-1}$	Seed immersed in aqueous solutions	Barbieri et al. (2013)
	Cilantro ( <i>Coriandrum sativum</i> L.)	0, 62.5, 125, 250, and 500 mg kg <sup>-1</sup> nCeO <sub>2</sub>	$125 \text{ mg kg}^{-1}$	Organic potting soil	Morales et al. (2013)
	Tomato (Solanum lycopersicum L.)	0 and 10 mg $L^{-1}nCeO_2$	< 10 mg L <sup>-1</sup>	Germination solution and hydroponic solution	Wang et al. (2013a)
	Wheat (Triticum aestivum L.)	0, 125, 250, and 500 mg kg <sup>-1</sup> nCeO <sub>2</sub>	$500 \text{ mg kg}^{-1}$	Potting soil	Rico et al. (2014)
	Rice (Oryza sativa L.)	0 and 100 μM CdCl <sub>2</sub> + 10 μM CeCl <sub>3</sub>	10 µM	Nutrient solution and foliar application	Wu et al. (2014)
	Barley (Hordeum vulgare L.)	0, 125, 250, and 500 mg kg <sup>-1</sup> $nCeO_2$	$250 \text{ mg kg}^{-1}$	Potting soil	Rico et al. (2015)

Pea (Pisum sativum L.)	0 and 8 mg kg <sup><math>-1</math></sup> CoSO <sub>4</sub>	$8 \text{ mg kg}^{-1}$	Potting soil in greenhouse and field experiment	Gad (2006)
Carnation (Dianthus caryophyllus L.) cv. "Harlem"	$\begin{bmatrix} 0, 50, 75, \text{ and } 100 \text{ mg } \text{L}^{-1} \\ \text{CoCl}_2 \end{bmatrix}$	$100~{ m mg~L}^{-1}$	Treatment solution	Jamali and Rahemi (2011)
Groundnut (Arachis hypogaea L.)	0 and 8 mg $kg^{-1}CoSO_4$	$8 \text{ mg kg}^{-1}$	Field experiments	Gad (2012)
Lily (Lilium spp.) cv. "Star Fighter" and cv. "Star Gazer"	0, 0.1, 0.2, 0.4, and 0.8 mM CoCl <sub>2</sub>	0.1 and 0.2 mM	Preservative solution	Mandujano-Piña et al. (2012)
Tuberose (Polianthes tuberosa L.)	0, 200, 300, and 400 mg $L^{-1}$ CoCl <sub>2</sub>	300 and $400$ mg L <sup>-1</sup>	Preservative solution	Mohammadi et al. (2012b)
Marguerite (Argyranthemum sp.)	0, 1, and 2 mM $CoCl_2$	2 mM	Preservative solution	Kazemi (2012)
Lily (Lilium spp.) cv. "Prato"	0, 1, and 2 mM $CoCl_2$	2.5 mM Ni + 2 mM Co + 2 mM salicylic acid with 2.5% sucrose	Preservative solution	Kazemi and Ameri (2012)
Tomato (Solanum lycopersicum L.)	0, 50, 100, 150, 200, and 250 mg kg <sup>-1</sup> CoCl <sub>2</sub>	$50 \text{ mg kg}^{-1}$	Pot contained soil	Jayakumar et al. (2013)
Rose (Rosa spp.) cv. "Red one"	0, 100, 200, and 300 mg $L^{-1}$ CoCl <sub>2</sub>	$200 \text{ mg L}^{-1}$	Preservative solution	Aslmoshtaghi et al. (2014)
Gladiola ( <i>Gladiolus</i> grandiflorus Hort.) cv. Borrega Roja	0, 0.3, and 0.6 mM CoCl <sub>2</sub> 6H <sub>2</sub> O	0.3 mM	Preservative solution	Trejo-Téllez et al. (2014)
				(continued)

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The Role of Beneficial Elements in Triggering Adaptive Responses to...

Table 1 (co	ontinued)				
Beneficial					
element	Species studied	Level evaluated	Beneficial level	Application system	Reference
I	Apple (Malus domestica	0, 1.5–2.40 and 3.0–4.8 mg L	$3.0-4.8 \text{ mg L}^{-1}$	Sprayed with foliar	Szwonek (2009)
	Borkh.) cv. "Golden Delicious/	<sup>-1</sup> organic-mineral		treatments	
	M.9"	liquidfertilizer			
	Lettuce (Lactuca sativa L.)	0, 20, 40, and 80 µM KI and	≤40 μM	Pots with vermiculite as	Blasco et al. (2011)
	cv. "Longifolia"	KIO <sub>3</sub>		asubstrate	
	Potato tubers (Solanum	0, 0.05, and 0.1% KI (w/v) or	0.05% KI, 0.1 and	Pot contained soil irrigated	Caffagni et al.
	tuberosum L.) and tomato	0, 0.05, 0.1, 0.2, and 0.5%	0.2% KIO <sub>3</sub>	with treatments	(2011)
	(S. lycopersicum L.)	KIO <sub>3</sub> (w/v)			
	Spinach (Spinacia oleracea L.)	0, 1, and 2 mg dm <sup><math>-3</math></sup> KI	$2 \text{ mg I} + 1 \text{ g dm}^{-3}$	Containers filled with silt	Smolen and Sady
	cv. "Olbrzym Zimowy"		sucrose	loam in plastic tunnel	(2011a)
	Spinach (Spinacia oleracea L.)	0, 1, and 2 mg dm <sup><math>-3</math></sup> KI	$2 \text{ mg dm}^{-3}$	Containers filled with silt	Smolen and Sady
	cv. "Olbrzym Zimowy"			loam in plastic tunnel	(2011b)
	Tomato (Solanum	Foliar application:2500 mg L	Hydroponic culture:	Foliar spray and soil fertilizer	Caffagni et al.
	lycopersicum L.)	<sup>-1</sup> I; and soil application: 90 g	1, 2, and 5 mM KI	in field experiments and	(2012)
		kg <sup>-1</sup> I crystalline fertilizer		hydroponic culture in	
				greenhouse	
	Soybean (Glycine max	0, 20, 40, and 80 μM KIO <sub>3</sub>	20 μM	Pots carrying soil and cow	Gupta et al. (2015)
	L. Merr.)			dung manure (ratio 3:1)	
	Carrot (Daucus carota L.)	0 and 2.5 kg ha <sup><math>-1</math></sup> KI and KIO <sub>3</sub> ,	$2.5 \text{ kg ha}^{-1} \text{ KI} +$	Field study	Smolen et al.
	cv. "Kazan F <sub>1</sub> "	applied twice	Na <sub>2</sub> SeO <sub>3</sub>		(2016)

Cucumber (Cucumis sativus	0, 0.02, and 2 mM LaCl3	0.02 mM	Quartz sand irrigated with	Zeng et al. (2000)
L.)			treatments	
Tobacco (Nicotiana tabacum L.)	$\begin{bmatrix} 0, 5, 10, 20, 50, and 100 \text{ mg L} \\ ^{-1}\text{LaCl3} \end{bmatrix}$	$20 \text{ mg L}^{-1}$	Hydroponic with nutrient solution	Chen et al. (2001)
Cucumber (C. sativus L.)	0, 0.002, 0.02, 0.2, and 2 mM	0.2 and 2 mM	Nutrient solution	Huang et al. (2003)
Cucumber (C. sativus L.)	0, 0.002, 0.02, 0.2, and 2 mM LaCl3	0.002 and 0.02 mM	Spraying twice daily	Shi et al. (2005)
Soybean ( <i>Glycine max</i> L. Merr.) cv. "Kennong"	0, 10, 20, 30, 40, and 50 mg L <sup>-1</sup> LaCl3	$20 \text{ mg L}^{-1}$	Sprayed solution on leaves	Yan et al. (2007)
Maize (Zea mays L.)	0, 0.2, 1.0, and 5.0 µM La	< 0.2 µM	Continuously flowing solu-	Diatloff et al.
cv. "Hycorn 82" and mung bean (Vigna radiata 1. Wilrzek) cv. "Berken"	(NO <sub>3</sub> ) <sub>3</sub>		tion culture units	(2008)
Tulip (Tulipa gesneriana) cv. "Ile de France"	$\frac{0, 5, 10, 20, 30, and 40 \mu M La}{(NO_3)_3}$	10 µM	Pots irrigated with treatments	Ramírez-Martínez et al. (2009)
Tulip (Tulipa gesneriana)	0, 5, 10, 20, 30, and 40 μM LaCl3 and La(NO <sub>3</sub> ) <sub>3</sub>	10 and 20 µM	Pots irrigated with treatments	Ramírez-Martínez et al. (2012)
Rice (Oryza sativa L.) cv. "Shengdao 16"	0, 0.05, 0.1, 0.5, 1.0, and 1.5 mM La(NO <sub>3</sub> ) <sub>3</sub>	0.1 mM	Application in basal medium	Liu et al. (2012a)
Rice (Oryza sativa L.) cv. "Shengdao 16"	0, 0.05, 0.1, 0.5, 1, and 1.5 mM La(NO <sub>3)3</sub>	0.05 mM	Application in basal medium	Liu et al. (2013)
Maize (Zea mays L.), mung bean (Vigna radiata	0, 5, 10, 15, 20, 25, 30, 35, 40, 45, and 50 μM La <sub>2</sub> O <sub>3</sub>	50 µM	Pots with nutrient solution	Chaturvedi et al. (2014)
				(continued)

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Table 1 (cc	ntinued)				
Beneficial		-			, ,
element	Species studied	Level evaluated	Beneficial level	Application system	Keterence
	L. Wilczek)and black gram (V. mungo)				
	Soybean (Glycine max	0, 5, 10, 20, 40, 80, and 60 µM	$\leq 10 \ \mu M$	Pots containing nutrient	de Oliveira et al.
	L. Merr.) cv. "BRSMG760SRR"	La(NO <sub>3</sub> ) <sub>3</sub>		solution	(2015)
	Rangpur lime (Citrus limonia	0, 50, 100, 200, and 400 mg	50 mg	Polypropylene tubes with	Turra et al. (2015)
	Osbeck)	LaCl3in 100 mL of water		substrates	
	Horseradish (Armoracia	0, 20, 100, and 300 mg $L^{-1}$	$20 \text{ mg L}^{-1}$	Nutrient solution	Zhang et al. (2016)
	rusticana G. Gaertn., B. Mey.	LaCl <sub>3</sub>			
	& Scherb.)				
	Bell pepper (Capsicum	0 and 10 µM LaCl <sub>3</sub>	10 µM	Nutrient solution	García-Jiménez
	annuum L.)				et al. (2017)
Na	Chrysanthemum (Chrysanthe-	0, 1000, 3000, 6000, and	$1000 \text{ mg L}^{-1}$	Pots with soilless substrate	Lee and van lersel
	mum x morifolium Ramat.)	$9000 \text{ mg } \mathrm{L}^{-1} \mathrm{NaCl}$			(2008)
	cv. "Yellow blush"				
	Faba bean (Vicia faba L.)	0, 60, 120, and 240 mM NaCl	60 mM	Pots with vermiculite	Abdul Qados (2011)
	Rose (Rosa hybrida)	0, 150, 200, and 250 mg $L^{-1}$	$250 \text{ mg L}^{-1}$	Preservative solution	Imani et al. (2012)
	cv. "Avalanche"	sodium benzoate			
	Rose (Rosa hybrida)	0, 50, and 100 $\mu$ M sodium	50 µM	Pretreatment for 24 h in pre-	Seyf et al. (2012b)
	cv. "Utopia"	nitroprusside		servative solution	
	Rose (Rosa hybrida)	0, 20, 40, and 60 μM sodium	20 µM	Preservative solution	Nazirimoghaddam
		nitroprusside			et al. (2014)
	Sunflower (Helianthus annuus	0, 20, 40, and 60 μM sodium	60 µM	Preservative solution	Nazirimoghaddam
	L.)	nitroprusside			et al. (2014)

	Lisianthus (Eustoma grandiflorum Raf. Shinn.)	0, 20, 40, and 60 μM sodium nitroprusside	40 µM	Preservative solution	Nazirimoghaddam et al. (2014)
	Cherry tomato (Solanum lycopersicum L.)	0, 0.87, and 28.6 mM Na <sup>+</sup> , saline water	28.6 mM Na <sup>+</sup>	Sandy soil	Silva et al. (2015)
	Purpletop vervain (Verbena bonariensis L.)	0 and 200 mM NaCl	200 mM	Pots with deacidified peat	Salachna and Piechocki (2016)
	Giant chincherinchee ( <i>Ornithogalum saundersiae</i> baker.)	0, 100, and 200 mM NaCl	100 or 200 mM	Pots with a mixture of peat and fertilizer	Salachna et al. (2016)
	Indian mustard (Brassica juncea L. Czern.)	0 and 150 mM NaCl	150 mM	Nutrient solution	Yousuf et al. (2016)
Se	Ryegrass (Lolium perenne L.)	0, 0.1, 1, 10, and 30 mg kg <sup>-1</sup> H <sub>2</sub> SeO <sub>4</sub>	$1 \text{ mg kg}^{-1}$	Pot with coarse-textured soil	Hartikainen et al. (2000)
	Maize (Zea mays L.) cv. "Złota Karłowa"	0, 5, 25, 50, and100 μMdm <sup>-3</sup> Na <sub>2</sub> SeO <sub>3</sub>	5 μM dm <sup>-3</sup>	Nutrient solution	Hawrylak-Nowak (2008)
	Soybean ( <i>Glycine max</i> L. Merr.) cv. "Olna"	0 and 10 mg $L^{-1}Na_2SeO_4$	$10 \text{ mg L}^{-1}$	Foliar sprayed with aqueous solution	Mechora and Germ (2010)
	Townsville stylo (Stylosanthes humilis Kunth Hester)	1 $\mu$ M and 0.1 mM Na <sub>2</sub> SeO <sub>4</sub>	1 µM	Seedling in Petri dishes with test solution	(2011)
	Peach grafted onto GF 677 (Prunus persica L.Batsch x Prunus amygdalus Stokes) and pear (Pyrus communis L.)	0, 0.1, and 1 mg $L^{-1}$ Na <sub>2</sub> SeO <sub>4</sub> via foliar or 1 mg $L^{-1}$ Na <sub>2</sub> SeO <sub>4</sub> via fruit	$1 \text{ mg L}^{-1}$	Via foliar and fruit application	Pezzarossa et al. (2012)
	Canola (Brassica napusL.) cv. "RGS"	0, 10, and 20 $\mu g$ plant <sup>-1</sup> Na <sub>2</sub> SeO <sub>4</sub>	10 and 20 µg	Foliar application	Hajiboland and Keivanfar (2012)
	Lettuce (Lactuca sativa L.) cv. "Justyna"	0, 2, 4, 6, 15, 20, 30, 40, and 60 μM Na <sub>2</sub> SeO <sub>4</sub> and 0, 2, 4, 6, 10, 15, 20, 25, and 30 μM Na <sub>2</sub> SeO <sub>3</sub>	<15 µM Na <sub>2</sub> SeO <sub>4</sub> or Na <sub>2</sub> SeO <sub>3</sub>	Nutrient solution	Hawrylak-Nowak (2013)
					(continued)

<b>Table 1</b> (co	ontinued)				
Beneficial					
element	Species studied	Level evaluated	Beneficial level	Application system	Reference
	Melon (Cucumis melo L.)	0, 2, 4, 8, and 16 $\mu M \ Na_2 SeO_3$	2, 4, and 8 μM	Nutrient solution	KeLing et al. (2013)
	Coleus (Coleus blumei Benth.)	0, 0.1, 0.5, 1, 2.5, and 5 mM Na <sub>2</sub> SeO <sub>3</sub>	1 mM	Hydroponic system in nutri- ent solution	Yuan et al. (2013)
	Cucumber (Cucumis sativus L.) cv. "Polan F1"	0, 5, and 10 $\mu$ M Na <sub>2</sub> SeO <sub>4</sub>	10 µM	Nutrient solution	Hawrylak-Nowak et al. (2014)
	Tomato (Solanum lycopersicum L.) cv. "Provence"	0, 0.1, 1, and 10 mg L <sup>-1</sup> Na <sub>2</sub> SeO <sub>4</sub>	$1\mathrm{mgL^{-1}}$	Foliar spray	Zhu et al. (2016)
	Carrot (Daucus carota L.) cv. "Kazan F1"	0 and 0.5 kg $ha^{-1}Na_2SeO_4$ or $Na_2SeO_3$	$0.5 \mathrm{kg}\mathrm{ha}^{-1}\mathrm{Na}_2\mathrm{SeO}_3$	Soil fertilization	Smolen et al. (2016)
Si	Forage oat (Avena sativa L.)	$0,50,100,150,and200mgkg^{-1}H_4O_4Si$	$100 \text{ mg kg}^{-1}$	Soil fertilization in pots	Borda et al. (2007)
	Bitter gourd (Momordica charantia)	$0, 1, 2, 3, and 5 mM K_2SiO_3$	3 mM	Aerated solution	Wang et al. (2010)
	Carnation (Dianthus caryophyllus L.) cv. "Harlem"	$0, 100, 150, and 300 \text{ mg } \text{L}^{-1}$ K <sub>2</sub> SiO <sub>3</sub>	$300 \text{ mg L}^{-1}$	Floral preservative solution	Jamali and Rahemi (2011)
	Starflower (Borago officinalis L.)	0, 0.5, 1, 1.5, and 2 mM Na <sub>2</sub> (SiO <sub>2</sub> ) <sub>3</sub>	0.5 and 1.5 mM	Pots with vermiculite	Shahnaz et al. (2011)
	Carnation (Dianthus caryophyllus L.)	0, 1.5, 2.5, and 3.5 mM Si	2.5 mM	Floral preservative solution	Kazemi et al. (2012)
	Rice (Oryza sativa L.) cv. "Dongjin"	0, 0.5, 1, and 2 mM $Na_2SiO_3$	0.5 mM	Nutrient solution	Kim et al. (2014)
	Snapdragon (Antirrhinum majus L.) cv. "Montego Purple"	0, 50, 100, 150, and 200 mg L <sup>-1</sup> NaSiO <sub>3</sub>	$150 \mathrm{mg}\mathrm{L}^{-1}$	Foliar application	Whitted-Haag et al. (2014)

with Si Abdel Latef and Tran (2016)	on and pots Marxen et al. (2016)	ion Owolade and Ogunleti (2008)	ion Norman and Chen (2011)	plants Norman and Chen (2011)	I fertigation Kleiber and Markiewicz (2013)	insions with Mahmoodzadeh et al. (2013)	ion Jaberzadeh et al. (2013)	ion Whitted-Haag et al. (2014)	ion Whitted-Haag et al. (2014)	ion Raliya et al. (2015)	(continued)
Seed priming solution	Soil fertilizatic with soil	Foliar applicat	Foliar applicat	Spraying onto	Rockwool and system	Seeds in suspe treatments	Foliar applicat	Foliar applicat	Foliar applicat	Foliar applicat	
1.5 mM	$_{-1}^{0.4}$ and 17.3 Mg ha	$125 \text{ cm}^3 \text{ ha}^{-1}$	75 mM	25 and 75 mM	960 g ha <sup>-1</sup>	$2000 \text{ mg L}^{-1}$	0.02%	50 and 75 mg $L^{-1}$	$75 \mathrm{mg}\mathrm{L}^{-1}$	$10 \text{ mg L}^{-1}$ 10 mg L	
0 and 1.5 mM Na <sub>2</sub> SiO <sub>3</sub>	$\begin{bmatrix} 0, 0.4, \text{ and } 17.3 \text{ Mg ha}^{-1} \text{silica} \\ \text{gel} \end{bmatrix}$	0, 62, and 125 cm <sup>3</sup> ha <sup>-1</sup> TiO <sub>2</sub>	0, 25, and 75 mM TiO <sub>2</sub>	0, 25, and 75 mM TiO <sub>2</sub>	0, 80, 240, 480, and 960 g $ha^{-1}$ Tytanit® fertilizer	10, 100, 1000, 1200, 1500, 1500, 1700, and $2000 \text{ mg L}^{-1} n \text{TiO}_2$	0, 0.01, 0.02, and 0.03% <i>n</i> TiO <sub>2</sub> and bulk Ti	0, 25, 50, 75, and 100 mg L <sup>-1</sup> Ti-ascorbate (Tytanit®)	0, 25, 50, 75, and 100 mg L <sup>-1</sup> Ti-ascorbate (Tytanit®)	0 and 10 mg L <sup><math>-1</math></sup> ordinary TiO <sub>2</sub> or $n$ TiO <sub>2</sub>	
Maize (Zea mays L.)	Rice (Oryza sativa L.) cv. "Khang Dan 18"	Cowpea (Vigna unguiculata L. Walp.) cv. "Ife Brown"	Geranium ( <i>Pelargonium x hortorum</i> L. H. Bailey) cv. "Patriot Bright Violet"	Poinsettia ( <i>Euphorbia</i> <i>pulcherrima</i> Willd. ex Klotzsch) cv. "Snowcap"	Tomato (Solanum lycopersicum L.) cv. "ISI 68249"	Canola (Brassica napusL.) cv. "RGS003"	Wheat (Triticum aestivum L.) cv. "Pishtaz"	Geranium( <i>Pelargonium x hortorum</i> L. H. Bailey)cv. "Elite Cherry"	Snapdragon (Antirrhinum majus L.) cv. "Montego Purple"	Mung bean (Vigna radiataL. Wilczek)	
		H									

Table 1 (cc	ntinued)				
Beneficial element	Species studied	Level evaluated	Beneficial level	Application system	Reference
	Timothy (Phleum pratenseL.)	0, 0.2, 0.4, and 0.8 dm <sup>3</sup> ha <sup>-1</sup> Tytanit®	0.4 and $0.8$ dm <sup>3</sup> ha <sup>-1</sup>	Foliar fertilization in field	Radkowski et al. (2015)
	Cabbage (Brassicaoleracea L.) cv. "Dutch premium"	0, 250, 500, and 1000 $\mu g \text{ mL}^{-1}$ $n \text{TiO}_2$	500 µg mL <sup>-1</sup>	Seeds in treatment suspension	Andersen et al. (2016)
	Cucumber (Cucumis sativus L.) cv. "Straight eight"	0, 250, 500, and 1000 $\mu g \text{ mL}^{-1}$ $n \text{TiO}_2$	500 and 1000 $\mu g \; mL_{-1}$	Seeds in treatment suspension	Andersen et al. (2016)
	Oat (Avena sativa L.)	0, 250, 500, and 1000 $\mu g \text{ mL}^{-1}$ $n \text{TiO}_2$	500 and 1000 $\mu g \; mL_{-1}$	Seeds in treatment suspension	Andersen et al. (2016)
	Barley (Hordeum vulgare L.)	0, 500, and 1000 mg kg <sup>-1</sup> nTiO <sub>2</sub>	$1000 \text{ mg kg}^{-1}$	Mixture of soil and nTiO <sub>2</sub>	Marchiol et al. (2016)
>	Soybean ( <i>Glycine max</i> L. Merr.)	0, 5, 10, 15, 30, 50, or 75 mg kg <sup>-1</sup> NH <sub>4</sub> VO <sub>3</sub>	15 mg kg <sup>-1</sup> in fluvo- aquic soil and 50 mg kg <sup>-1</sup> in red earth soil	Mixture of soil and V	Wang and Liu (1999)
	Soybean (Glycine max L. Merr.) cv. "Corsoy"	$0, 0.5, 1.0, and 2.0 \text{ mg kg}^{-1}$ Na <sub>3</sub> VO <sub>4</sub>	1 mg kg <sup>-1</sup> with farmyard manure	Mixture of soil with V	Sozudogru et al. (2001)
	Chinese green mustard (Bras- sica campestris L. subsp. Chinensis) cv. "Parachinensis"	$_{-1}^{}$ NH_4 VO_3 $^{-1}$ NH_4 VO_3 $^{-1}$	$1-20 \text{ mg L}^{-1}$	Nutrient solution	Vachirapatama et al. (2011)
	Tomato (Solanum lycopersicum L.)	0, 1, 10, 20, 40, and 80 mg $L^{-1}$ NH <sub>4</sub> VO <sub>3</sub>	1 and 10 mg $L^{-1}$	Hydroponics solution	Vachirapatama et al. (2011)
	Common bean ( <i>Phaseolus</i> vulgaris L.) cv. "Contender"	0, 160, 240, 320, and 400 μM VOSO <sub>4</sub>	240 µM	Vermiculite and watered with treatments	Saco et al. (2013)
	Wheat (Triticum aestivum L.) cv. "Liaochum 9"	0 and 40 $\mu$ M Na <sub>3</sub> VO <sub>4</sub>	40 µM	Nutrient solution	Wang et al. (2013b)
	Pennyroyal (Mentha pulegium L.)	$0, 5, 10, 20, and 40 \text{ mg } \text{L}^{-1}$ NH <sub>4</sub> VO <sub>3</sub>	$10-40 \text{ mg L}^{-1}$	Pots with peat and perlite	Akoumianaki- Ioannidou et al. (2015)

Application of Se increases its concentrations in fruit trees (Pezzarossa et al. 2012), lettuce (Hawrylak-Nowak 2013), and carrot (Smolen et al. 2016), which in turn improves plant growth and development. Under saline stress, Si reduces Na uptake and increases that of K, in order to maintain a better K/Na ratio in maize (Abdel Latef and Tran 2016), while in rice, Si increases P, N, and Mg uptake, though reduces that of K, which dramatically improves plant growth, yield, and grain quality (Marxen et al. 2016). Sodium increases N and K contents in Ornithogalum saundersiae Baker. (Salachna et al. 2016), which improves vessel life and flower quality. Titanium may improve crop performance through stimulating the activity of certain enzymes, enhancing chlorophyll content and photosynthesis, promoting nutrient uptake, strengthening stress tolerance, and improving crop yield and quality. These benefits lie in its interaction with other nutrient elements, especially Fe. Fe and Ti have synergistic and antagonistic relationships, depending on the Fe status of the plant. When plants experience Fe deficiency, Ti may enhance Fe uptake and utilization and subsequently improving plant growth. When Ti concentration is high in plants, Ti competes with Fe for ligands or proteins. The competition could be severe, resulting in Ti phytotoxicity (Lyu et al. 2017). Moreover, Ti has been shown to increase N, P, K, Ca, and Mg contents in tomato (Kleiber and Markiewicz 2013). Vanadium does not affect Ca, K, Mn, and Zn concentrations in roots, though it does reduce Ca, K, Mg, and Mn concentrations in leaves of Mentha pulegium L. (Akoumianaki-Ioannidou et al. 2015). Importantly, its role in productivity, nutrient uptake, and mobility within the plant deserves further investigation. All these effects of beneficial elements ameliorate the responses of plants to environmental stressors. A summary of the main effects of the beneficial elements on plant nutrition and their functions on plant performance is presented in Fig. 2.

#### 4 Conclusions and Perspectives

Aluminum, cerium, cobalt, iodine, lanthanum, sodium, selenium, silicon, titanium, and vanadium have been shown to have beneficial effects in some species of model and cultivated plants. The positive effects of these elements on plants include improved yield and postharvest quality, absorption of other nutrients, and activation of mechanisms of defense against pests and diseases and resistance or tolerance to abiotic stress factors such as heavy metals, drought, and salinity. In some cases, these elements can replace some biological functions of other essential nutrients in plant metabolism.

In every case, the beneficial effects of these elements are always observed when they are used at low concentrations. Since they trigger hormetic effects, at high doses, these elements can disrupt the homeostasis of the plant and cause deleterious effects.

The great challenges facing humanity such as population growth, climate change, pollution, and the depletion and degradation of natural resources make it necessary



Fig. 2 Beneficial effects of Al, Ce, Co, I, La, Na, Se, Si, Ti, and V on plant nutrition and their impacts on plant production

to find new methods of sustainable crop production. In this context, beneficial elements offer an alternative little explored until now.

To date, well-supported experimental evidence demonstrates the positive effects of beneficial elements in different plant species. Nevertheless, more in-depth research is still needed in order to know their action based on the plant genotypes used, the production systems, the chemical forms in which they should be applied, and, above all, the optimal doses and phenological stages in which their beneficial effects are more evident and their application cost-effective. Importantly, nanotechnology has the potential to positively impact the agrifood sector, minimizing adverse problems of agricultural practices on environment and human health and improving food security and productivity while promoting social and economic equity. However, acquisition of knowledge and developments of methods for risk and life-cycle assessment of nanomaterials, nanopesticides, and nanofertilizers, as well as assessment of the impacts on nontarget organisms (i.e., other plants, soil microbiota, and bees), and the regulations about the use of nanomaterials require further attention (Amenta et al. 2015; Fraceto et al. 2016).

In addition to the ten beneficial elements described herein, recent reports indicate that other elements such as silver (Ag), chromium (Cr), fluorine (F), and tungsten (W) may also have potential benefits in crops, but research on them is still in its infancy.

Because the responses that trigger beneficial elements differ among families, genera, and species of plants, it is crucial to explore the underlying genetic and molecular foundations that explain the positive effect of these elements, which constitute an area of great interest for future studies. Food security, sustainability, and efficient use of current inputs eminently justify such an approach.

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