Effect of Metal Stress due to Strontium and The Mechanisms of Tolerating it by *Amaranthus caudatus* Linn

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**Abstract**

Heavy metals are major environmental pollutants when present in high concentration in soil and have toxic effects on growth and development of plants. Experiment were carried out to find out the effect of different levels of strontium element commonly used in the fireworks – cottage industries of this area and the predominant pollutant on growth, biochemical and enzymatic characteristics of *Amaranthus caudatus* Linn, widely cultivated in this area. This study is aimed at assessing the stress tolerant ability of *Amaranthus caudatus* Linn. The seedlings of *Amaranthus caudatus* L. were treated with various concentration of strontium (2 mM, 4 mM, 6 mM, 8 mM, 10 mM) for ten days and its effect on the morphometric, biochemical and enzymatic characters were analyzed. After ten days of treatment, the growth parameters such as leaf area, fresh weight, dry weight, shoot and root length were found decreased than in the control. Biochemical characteristics such as the content of chlorophyll, carotenoid, soluble sugar and protein also decreased with the increase in the concentration of strontium. In contrary, the content of free amino acid, proline, leaf nitrate and the activities of anti-oxidative enzymes such as catalase and peroxidase were found increased with the increase in the concentration of strontium while the activity of nitrate reductase was found decreased. The result suggest that *Amaranthus caudatus* Linn. has been affected adversely by metal stress due to strontium but at the same time the plant adopts mechanisms such as accumulation of anthocyanin and enhanced activities of antioxidant enzymes to overcome the ill effects of the metal ions.

**Keywords**: Strontium; *Amaranthus caudatus* L.; Free amino acid; Proline; Catalase; Peroxidase

**Introduction**

Heavy metals are defined as group of elements that have specific weights higher than about 5 g/cm³. Iron, Mn, Mo, Ni, Zn and Cu are essential micronutrients required for normal growth and metabolic processes in plants [1]. Cd, Pb, Cr and Hg are nonessential and highly toxic to plants [2]. Heavy metals inhibit physiological processes such as respiration, photosynthesis, cell elongation and affect plant water relationship as well as mineral nutrition [3]. The most common heavy metal contaminants are Cd, Cr, Cu, As, Hg, Pb and Ni. Other naturally occurring metallic elements with high molecular weight are also considered as important pollutants. These elements occur either naturally in soil or get deposited through the use of agricultural chemicals, urban waste and polluted water [4]. As metals cannot be broken down, when concentrations within the plant exceed optimal levels, they adversely affect the plant both directly and indirectly. Some of the direct toxic effects caused by high metal concentration include inhibition of cytoplasmic enzymes and damage to cell structures due to oxidative stress [5]. Heavy metal toxicity is known to injure cell membranes, reduces transpiration, causes the breakdown of the protein synthesis, damages the photosynthesis apparatus, inhibits the photosynthetic rate, and affects the activity of several enzymes [6]. Heavy metals are known to interfere with chlorophyll synthesis either through direct inhibition of an enzymatic step or by inducing deficiency of an essential nutrient [7]. Reactive oxygen species (ROS) are continuously produced at low level during normal metabolic processes [8]. But in biological systems, increasing the synthesis of ROS is one of the initial responses to different stress factors [9]. ROS induce damage to the biomolecules through peroxidation of membrane lipids, alteration of protein functions, DNA mutation, damage to chlorophyll and disruption of metabolic pathways (electron transport chain and ATP production) [10]. Therefore the tolerance of plants to stress conditions depends on their ability to make balance between the production of toxic oxygen derivatives and capacity of antioxidative defense systems. Therefore, plants have complex ROS scavenging mechanisms at the molecular and cellular levels. These mechanisms with inhibition or slowing the oxidation of biomolecules and oxidative chain reactions [11] decrease the cellular oxidative damage and increase resistance to heavy metals. Pb induces toxicity in plants in terms of their growth, development, and biochemical attributes. Primary effects of Pb toxicity in plants include stunted root growth, probably due to inhibition of cell division in root tips. Secondarily, it induces oxidative stress via reactive oxygen species generation and results in cellular damage [12]. Farrag et al. [13] noticed the stress of heavy metals on some physiological parameters of *Amaranthus hybridus, Chenopodium ambrosioides, Mentha longifolia* and *Typha domingensis*. The expression of proteins in stressed plant was significantly decreased at all the levels compared to unstressed plant samples. Heavy metals increase the activities of catalase, glutathione peroxidase and glutathione reductase. Recently Muradoglu et al. [14] reported that the excessive Cd reduced chlorophyll contents, increased antioxidant enzyme activities and change plant nutrition concentrations in both roots and leaves. The higher concentration of Cd has negative effect on chlorophyll content and nearly decreased 30% of plant growth in strawberry. The present experiment was undertaken to investigate the changes in the level of growth, biochemical aspects, and enzymatic characteristics in *Amaranthus caudatus* Linn. treated with strontium and to assess its stress tolerating efficacy. This crop may further be

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useful in soil reclamation through the process of phytoremediation. In the present day context our study is very crucial because the soil in our area remains highly polluted due to strontium indiscriminately used in fireworks industries which effects most commonly cultivated *Amaranthus caudatus Linn.* - the chief crop of vegetable source of the locals. Thus there is an urgent necessity to understand the ill effects of strontium on the crop mentioned and the necessary steps to taken to overcome the problem.

**Materials and Methods**

**Plant material**

*Amaranthus caudatus Linn.* was selected for present study for its economic importance since it is cultivated in large scale in Virudhunagar district, Tamil Nadu, India.

**Experimental design**

Healthy viable and uniform sized seeds of *Amaranthus caudatus Linn.* were selected and were surface sterilized for 20 min with 1% (v/v) sodium hypochlorite, and then washed several times with distilled water. Both control and experimental plants were allowed to grow in the soil. After ten days of growth, they were treated with various concentration of strontium (2 mM, 4 mM, 6 mM, 8 mM,10 mM v/v) for seven days and the growth parameters such as root length, shoot length, leaf area, fresh weight and dry were analyzed. The various biochemical and enzyme activities were estimated following the methods proposed by those mentioned within the bracket. Chlorophyll and carotenoids [15], Anthocyanin [16], Total soluble sugar [17], Protein content [18], amino acid content [17], Proline [19], leaf nitrate [20] *in vivo* nitrate reductase activity [21], peroxidase and catalase activity [22].

**Statistics Analysis**

The Growth parameters were determined with ten independent replicates. Biochemical characters and enzymatic assay were carried out for five times. The data reported as mean ± SE and within parentheses represent the per cent activity. Statistical analysis (One way ANOVA – Tukey test) was applied using the statistical package, SPSS 16.0.

**Results and Discussion**

The results obtained have been summarized and discussed as follows.

**Effect of strontium on plant growth**

*Amaranthus caudatus Linn.* seedling exposed to different concentration of strontium exhibited inhibition of morphometric characters such as root length, shoot length, leaf area, fresh weight and dry weight compared to the control. The reduction of seedling growth was 41%, 69%, 35%, 30% and 23% respectively at 10mM concentration compared to the control (Figure 1 and Table 1). Reduced root length is due to reductions in both new cell formation and cell elongation in the extension region of the root [23]. The reduced root length of plants on metal treatments could be due to the reduction in mitotic cell division in meristematic zone [24]. This is in agreement with the findings of Soleimani et al. [25] who stated that the amount of accumulated Pb in tall fescue and Bermude grass (*Cynodon dactylon*) were higher in roots compared to the shoots. Likewise, similar response to lead treatment was previously noticed in water hyacinths. The plant growth was significantly inhibited (50%) at 1000 mg/L Pb concentration. Similar observations were also recorded under *Mentha arvensis* [26]. The same trend was observed earlier by other workers [27-30]. Accumulation of Pb was high in root than in shoot tissues [31]. Similar changes in the content by various metal treatments were recorded by Muradoglu et al. [14] with cadmium.

**Effect of strontium on photosynthetic pigments**

Chlorophyll is the important photosynthetic pigment which plays a vital role in the photosynthetic process. A significant decrease in Chlorophyll *a, b* and total chlorophyll contents was observed in the range of 64%, 74%, 43%, respectively at the maximum concentration (Figure 2 and Table 2). Similarly, carotenoids content also got significantly decreased to the range of 62% (Figure 2 and Table 2). But in case of anthocyanin it was significantly increased. This reduction in the growth and photosynthetic pigments could be due to the disturbance in photosystem I and induced activity of chlorophyllase enzyme [32].

The protective function of anthocyanin against the stress conations is fairly clears [33]. Similarly, observations have been made in many plants species [34-36]. Other results that support what has been shown here, are those by Kapoor et al. [37] with his study on *Brassica juncea* L. under Cd stress, Rastgoo et al. [38] with their study on *Aeluropus littoralis* under copper, nickel and zinc. It has been proposed that Cu at toxic concentration interferes with enzymes associated with chlorophyll biosynthesis and protein composition of photosynthetic membranes [39]. Many studies have shown the decrease of photosynthetic pigments under high concentration of metals [40-47].

**Effect of strontium on biochemical characters**

The total soluble sugar and protein contents got significantly decreased with increasing concentration of strontium. At 10 mM concentration the levels of reduction was 63% and 71% respectively while leaf nitrate, free amino acids, proline (Figure 3 and Table 3) contents was significantly increased with the increasing metal concentration. Under the heavy metal treatment, there was a considerable reduction in the growth and photosynthetic pigments, which could be due to the disturbance in photosystem I and induced activity of chlorophyllase enzyme. This disturbance paralleled with the reduction in sugar content that could be attributed to reduction in chlorophyll contents of the leaf and also a decline in protein. This change might have already affected the photosynthetic activity in the plant and hence the reduction in carbohydrate contents [32]. Similar observation was reported by Somashekararia et al. [48] in *Phaseolus vulgaris* and Neehu et al. (2000) in *Vicia faba* treated with Cd. A reduction in leaf protein indicated the reduction in RUBACase, which caused a reduction in photosynthetic activity, which in turn, affects the total soluble sugar
Impact of various concentration of strontium on the pigments and morphometric characteristics of Vigna mungo were not utilized in the protein synthesis. The notable decrease in the nitrate sources when 10 mM concentration of strontium was noticed. There is a dramatic rise in the availability of free amino acid is significantly high. It may be due to destruction of protein or due to the biosynthesis of amino acid from the nitrate sources which were not utilized in the protein synthesis. Accumulation of proline has been frequently used as biochemical marker for water stress in plants [51]. The free proline has been shown to protect plants against free radicle induced damage by quenching of singlet oxygen [52]. The results of the present study was confirmed by Ravikumar and Thamizhiniyzn [54] who observed proline changes in black gram seedlings by under in Pb stress. Moreover, Sharma [55] in his study on Brachythecium populare proved the impact of heavy metals on some physiological parameters like total chlorophyll, sugar, protein content and confirmed the inhibitory effect of heavy metal on biochemical contents.

Effect of strontium exposure on antioxidant enzyme activities

The notable decrease in in vivo nitrate reductase activity of 63% at 10 mM concentration of strontium was noticed. There is a dramatic rise in

### Table 1: Impact of various concentration of strontium on the morphometric characteristics of *Amaranthus caudatus* Linn.

<table>
<thead>
<tr>
<th>Growth Parameters</th>
<th>Control</th>
<th>2 mM</th>
<th>4 mM</th>
<th>6 mM</th>
<th>8 mM</th>
<th>10 mM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Root length (cm)</td>
<td>11.33 ± 0.176 (100)</td>
<td>10.43 ± 0.088a* (92)</td>
<td>9.56 ± 0.084a* (78)</td>
<td>8.46 ± 0.066a* (69)</td>
<td>7.26 ± 0.054a* (59)</td>
<td>5.06 ± 0.145a* (41)</td>
</tr>
<tr>
<td>Shoot length (cm)</td>
<td>15.48 ± 0.240 (100)</td>
<td>14.70 ± 0.057a* (95)</td>
<td>13.66 ± 0.066a* (98)</td>
<td>12.76 ± 0.088a* (93)</td>
<td>11.76 ± 0.088a* (76)</td>
<td>10.60 ± 0.057a* (69)</td>
</tr>
<tr>
<td>Leaf area (cm²)</td>
<td>6.88 ± 0.026 (100)</td>
<td>5.89 ± 0.038a* (86)</td>
<td>4.93 ± 0.026a* (72)</td>
<td>3.84 ± 0.023a* (56)</td>
<td>3.23 ± 0.003a* (47)</td>
<td>2.38 ± 0.293a* (35)</td>
</tr>
<tr>
<td>Fresh Weight (gm)</td>
<td>1.99 ± 0.052 (100)</td>
<td>1.77 ± 0.098a* (89)</td>
<td>1.33 ± 0.294a* (70)</td>
<td>0.87 ± 0.005a* (44)</td>
<td>0.81 ± 0.046a* (41)</td>
<td>0.59 ± 0.008a* (30)</td>
</tr>
<tr>
<td>Dry Weight (gm)</td>
<td>0.38 ± 0.014 (100)</td>
<td>0.24 ± 0.008a* (64)</td>
<td>0.20 ± 0.008a* (54)</td>
<td>0.18 ± 0.003a* (48)</td>
<td>0.15 ± 0.005a* (39)</td>
<td>0.086 ± 0.003a* (23)</td>
</tr>
</tbody>
</table>

Values in parameters indicate percent activity; values are represents means of five observation. Values in parentheses are activity with respect to control. Mean (±) SE a - refers to values compared with control in various concentrations of metals, a* - refers to significant (P ≤ 0.05 – Tukey test). a - refers to non – significant

### Table 2: Impact of various concentration of strontium on the pigments characteristics of *Amaranthus caudatus* Linn.

<table>
<thead>
<tr>
<th>Growth Parameters</th>
<th>Control</th>
<th>2 mM</th>
<th>4 mM</th>
<th>6 mM</th>
<th>8 mM</th>
<th>10 mM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chlorophyll a</td>
<td>15.28 ± 0.199 (100)</td>
<td>13.38 ± 0.122a* (88)</td>
<td>11.97 ± 0.063a* (78)</td>
<td>9.69 ± 0.053a* (63)</td>
<td>7.81 ± 0.143a* (51)</td>
<td>5.55 ± 0.048a* (36)</td>
</tr>
<tr>
<td>Chlorophyll b</td>
<td>10.23 ± 0.104 (100)</td>
<td>8.81 ± 0.010a* (87)</td>
<td>7.35 ± 0.121a* (72)</td>
<td>6.30 ± 0.122a* (61)</td>
<td>4.42 ± 0.072a* (43)</td>
<td>2.67 ± 0.257a* (28)</td>
</tr>
<tr>
<td>Total Chlorophyll</td>
<td>25.51 ± 0.271 (100)</td>
<td>23.03 ± 0.132a* (91)</td>
<td>20.34 ± 0.074a* (79)</td>
<td>19.3 ± 0.254a* (76)</td>
<td>17.76 ± 0.056a* (64)</td>
<td>14.56 ± 0.070a* (57)</td>
</tr>
<tr>
<td>Carotenoids</td>
<td>7.85 ± 1.722 (100)</td>
<td>6.04 ± 0.276a* (77)</td>
<td>5.39 ± 0.289a* (68)</td>
<td>5.27 ± 0.037a* (54)</td>
<td>4.82 ± 0.003a* (46)</td>
<td>3.75 ± 0.003a* (38)</td>
</tr>
<tr>
<td>Anthocyanin</td>
<td>4.22 ± 0.058 (100)</td>
<td>2.62 ± 0.045a* (118)</td>
<td>3.71 ± 0.213a* (167)</td>
<td>4.63 ± 0.029a* (208)</td>
<td>4.86 ± 0.016a* (219)</td>
<td>5.68 ± 0.023a* (256)</td>
</tr>
</tbody>
</table>

Values in parameters indicate percent activity; values are represents means of five observation. Values in parentheses are activity with respect to control. Mean (±) SE a - refers to values compared with control in various concentrations of metals, a* - refers to significant (P ≤ 0.05 – Tukey test). a - refers to non – significant

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catalase and peroxidase activities after exposure to metal stress (Figure 4 and Table 4). Catalase activity was found significantly increased in all the experiment plants than the control plants. The increase was about 35% at 6 mM. Peroxidase activity was significant increased in all treatments. The increase was about 33% at 6 mM. Plant cells are equipped with a protective system including antioxidant enzymes like catalase and peroxidase which can flush free radicals [56]. Reduction in in vivo NR activity with increased concentration of cadmium on Vigna radiata has been reported earlier [57]. Catalase is an antioxidant and scavenging enzyme and it is activity was found increased with the increasing concentration of strontium. Catalase is special type of peroxidase enzyme which catalase the degradation of $\text{H}_2\text{O}_2$, which

<table>
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<th>6 mM</th>
<th>8 mM</th>
<th>10 mM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total Soluble sugar (mg/gLFW)</td>
<td>7.77 ± 0.030 (100)</td>
<td>6.90 ± 0.032a* (89)</td>
<td>5.92 ± 0.017a* (76)</td>
<td>4.92 ± 0.020 a* (63)</td>
<td>3.83 ± 0.085 a* (49)</td>
<td>2.92 ± 0.017 a* (37)</td>
</tr>
<tr>
<td>Total Soluble Protein (mg/gLFW)</td>
<td>4.85 ± 0.049 (100)</td>
<td>4.35 ± 0.033a* (90)</td>
<td>3.41 ± 0.088a* (70)</td>
<td>2.75 ± 0.047a* (57)</td>
<td>2.02 ± 0.030 a* (42)</td>
<td>1.41 ± 0.012 a* (29)</td>
</tr>
<tr>
<td>Amino acid (µ mole/g LFW)</td>
<td>2.37 ± 0.204 a* (100)</td>
<td>2.68 ± 0.191 a* (112)</td>
<td>3.07 ± 0.057a* (123)</td>
<td>3.32 ± 0.128a* (135)</td>
<td>4.31 ± 0.147a* (148)</td>
<td>4.75 ± 0.017 (165)</td>
</tr>
<tr>
<td>Proline (µ mole/g LFW)</td>
<td>3.27 ± 0.025 (100)</td>
<td>3.84 ± 0.008a* (117)</td>
<td>4.505 ± 0.035a* (121)</td>
<td>5.43 ± 0.001a* (128)</td>
<td>5.83 ± 0.003 a* (142)</td>
<td>7.44 ± 0.005 a* (152)</td>
</tr>
<tr>
<td>Leaf Nitrate (µg/gLFW)</td>
<td>5.34 ± 2.100 (100)</td>
<td>5.84 ± 0.026a* (107)</td>
<td>6.44 ± 0.057a* (115)</td>
<td>6.75 ± 4.150a* (124)</td>
<td>7.77 ± 0.033 a* (143)</td>
<td>8.44 ± 0.103 a* (155)</td>
</tr>
</tbody>
</table>

Values in parameters indicate percent activity; values are represents means of five observation. Values in parentheses are activity with respect to control. Mean (±) SE a - refers to values compared with control in various concentrations of metals, a* - refers to significant (P ≤ 0.05 – Tukey test). a# - refers to non–significant

Table 3: Impact of various concentration of strontium on the biochemical characteristics of Amaranthus caudatus (L.)

Figure 3: Effect of the various concentration of strontium on the biochemical characters of Amaranthus caudatus Linn.

Figure 4: Effect of the various concentration of strontium on the enzymatic characters of Amaranthus caudatus Linn.
is a natural metabolite to toxic plants [58]. Sharma [55] in his study on Brachythecium populare, proved that the under the heavy metal stress, catalase and peroxidase enzymes activities were significantly high. In addition, certain changes was observed earlier under cadmium on strawberry [14]. Results of the study by Farrag et al. [13] on heavy metal analysis with an increase in catalase, peroxidase enzyme activity and increasing the activity of antioxidant enzymes and accumulation of antioxidant compounds, protein but a significant enhancement of anthocyanin, proline, free amino acids, leaf nitrate. Likewise the activity of nitrate reductase got significantly reduced while activity of catalase and peroxidase were increased [59]. Our result suggest that strontium in high concentration makes the soil toxic to the plants and results in growth inhibition, structural damage, decline in physiological and biochemical activities of plants. As the same time plants adopt certain damage controlling mechanisms such as increasing the activity of antioxidant enzymes and accumulation of anthocyanin as a protective measure helping them to overcome the ill effect of the metal stress.

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