Antioxidant defense in rat tissues after supplementation with organic form of manganese

Aleksandra Jankovic, Milica Vucetic, Ana Stancic, Vesna Otasevic, Bato Korac1,2, Biljana Buzadzic1

1 University of Belgrade, Institute for Biological Research "Sinisa Stankovic", Department of Physiology, 11060 Belgrade, Serbia.
2 University of Belgrade, Faculty of Biology, 11000 Belgrade, Serbia.

Corresponding author: Biljana Buzadzic, PhD
University of Belgrade
Institute for Biological Research "Sinisa Stankovic" Department of Physiology
Bulevar despot St芬a 142
11060 Belgrade, Serbia
Tel: (381-11)-2078-307
Fax: (381-11)-2761-433
E-mail: buzadzic@ibi.bg.ac.rs

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Abstract: This study investigated the effect of Mn chelate supplementation on the activity of antioxidant enzymes: copper-zinc superoxide dismutase (CuZnSOD and MnSOD), catalase (CAT), glutathione peroxidase (GSH-Px), glutathione reductase (GR), glutathione S-transferase (GST) and the amount of glutathione (GSH) in the blood, liver, brain, heart and small intestine of the rat. Increased Mn intake of 34% above daily intake (0.051 ± 0.045 mg Mn / kg b.wt.) induced changes in the antioxidant defense in the tissues of the rat.

Decreased activity of peroxides-converting enzymes CAT and GSH-Px was found in all examined tissues (except GSH-Px activity in the liver). The amount of GSH was increased in plasma and liver, while it was decreased in heart and brain of supplemented animals. Treatment with Mn increases MnSOD activity in the liver, and CuZnSOD activity in the heart. The observed changes in antioxidant defense are closely connected with the newly established redox profile in the examined tissues after Mn supplementation.

Key words: Manganese; metal chelate; antioxidant enzymes; glutathione.

INTRODUCTION

Manganese is one of several first-row transition elements that assist in various metabolic and structural roles in biological system. It is used to give structural support to proteins and it is a cofactor in chemical transformation that include hydrolytic and redox reactions. Some of the enzymes that contain manganese are arginase, pyruvate carboxylase, glutamine synthetase and the enzymes that catalyze redox-based chemical transformations: manganese superoxide dismutase (MnSOD), manganese peroxidase, and manganese dioxygenase [1].

No cases of human manganese deficiency have been recorded [2]. In animals, manganese deficiency can cause impaired growth, skeletal abnormalities, and altered metabolism of carbohydrates and lipids [3]. Food items are the most important sources of essential trace elements. Mineral requirements are highly dependent on the physiological and health status of the animals and humans. In many situations additional mineral supplementation is recommended. The effort has been made to provide minerals in correct quantities and in the most available forms to the animals and humans. As a consequence there has been increasing interest in the role of organic sources of trace elements, particularly Cu, Mn, Se, Zn, and Fe, which function at the active sites of metal ion-dependent enzymes, such as SOD, catalase (CAT) and glutathione peroxidase (GSH-Px). With respect to human and animal nutrition, the potential benefit to health of antioxidants supply, either through dietary means or as nutritional supplements, came in the focus of the research in the last years.

Many factors affect mineral requirements, including age, amount and chemical forms of elements and their interrelationships with other nutrients. Because many factors are involved in bioavailability of dietary inorganic trace elements, the alternative approaches and sources were examined. The ability of metals to both ‘heal’ and ‘harm’, has led to the development of a wide range of mechanisms that regulate metal uptake and distribution in tissues. It is known that gastrointestinal absorption for organometallic species is more efficient than inorganic forms [8].

In order to increase the content and bioavailability of trace minerals some authors tried to increase the total level of minerals in plant foods, and at the same time to increase the concentration of compounds which promote their uptake (ascorbic acid), and/or to decrease the concentration of compounds which inhibit their absorption (phytic acid or phenolic compounds) [9]. The inclusion of organic complexes or chelated mineral products into supplements has been suggested on the basis of their higher bioavailability as compared to inorganic mineral salts [10, 11, 12 and 13]. For mineral proteinates, it had been shown that, beside higher bioavailability, they are also more bioactive [14].

It is known that Mn possesses ability to change its oxidative state and to generate reactive oxygen species (ROS). Dose-dependent increase in formation of ROS was found after in vitro exposure to MnCl2 in the rat brain [15]. Contrary
to this, Mn has important role as a constituent of the antioxidant enzyme MnSOD, which catalyzes the dismutation of superoxide anion radical (O₂•−). Also, high intracellular manganese has been known to suppress oxidative stress [16,17]. AsMn may play a role in maintaining a healthy balance between oxidants and antioxidants [18], in the present study, the effects of supplementation with Mn proteinates on the antioxidant defense components: CuZnSOD, MnSOD, CAT, GSH-Px, glutathione reductase (GR), glutathione S-transferase (GST), and the amount of glutathione (GSH), in the blood, liver, brain, heart and small intestine of the rats, were examined.

Materials and methods

Male rats (Mill Hill hybrid hooded, age 4 months) were divided into two groups (six animals within each group). Body weight in control group was 370±12 g and in Mn-supplemented group 395±13 g. All animals were housed in individual cages with woodshop bedding at 20-25°C and fed the same pelleted rat diet. The control group had food and water ad libitum, and the test group was supplemented with the organic form of manganese (chemically bound to amino acids and small peptides as a chelating agents) – Bioplex Mn liquid (Alltech Inc., USA) through drinking water for 45 days.

At the end of experiments the intake of Mn was calculated. The rats received trough drinking water 0.323 mg Mn/animal/day, which for 34% above the standard amount in the control diet. Expressed per body weights, animals received 0.851±0.045 mg Mn/kg. The experimental protocol was approved by the Ethical Committee for the Treatment of Experimental Animals of the Institute for Biological Research, Belgrade.

The rats were sacrificed by decapitation and fresh heparinized blood was collected. The erythrocytes and plasma were separated by centrifugation. Erythrocytes were washed three times with saline and lysed with cold water. The liver was perfused with a cold physiological saline solution and samples of all organs (liver, heart, brain and small intestine) were dissected within 3 min after death, and thoroughly rinsed with saline to wash out trace of blood. The obtained samples were homogenized (Janke and Kunkel Ka/Werke Ultra/Turrax homogenizer, 0-4°C) in a solution containing 0.25 mM ethylenediaminetetraacetic acid (EDTA) and 0.05 M Tris-HCl buffer, pH 7.4. The homogenates were sonicated as suggested by Takada et al. [19]. SOD activity was examined by a modified method of Misra and Fridovich [20], and expressed in U g⁻¹ Hb for erythrocytes and U mg⁻¹ protein for the respective tissues. SOD units are defined as the amount of the enzyme inhibiting epinephrine oxidation by 50% under the appropriate reaction conditions. Total specific SOD activity and CuZnSOD activity after inhibition with 4mM KCN, were measured, and then MnSOD activity was calculated. CAT was assayed as suggested by supplier (SIGMA Chemicals, St. Luis, MO, USA) and activity expressed as nmol H₂O₂ min⁻¹ g⁻¹ Hb for erythrocytes, and nmol H₂O₂ min⁻¹mg⁻¹ protein of the tissue. GSH-Px was determined using t-buthlyhydroperoxide as a substrate [21] and the experiment as expressed as nmol of NADPH oxidized min⁻¹ per mg Hb for erythrocytes, and per mg protein for tissue. GR was assayed as suggested by Glatzle et al. [22] and expressed as nmol NADPH min⁻¹mg⁻¹protein for the tissue. GST was determined by the procedure of Habig et al. [23] and the activity was expressed as nmol GSH min⁻¹mg⁻¹ protein. The quantity of GSH was examined in the plasma and in the tissues after deproteinization with 10% sulfosalicylic acid. The GSH was determined by using the method of Griffith [24] and results were expressed as nM GSH L⁻¹ plasma and nM GSH g⁻¹ tissue.

Protein content was estimated by the method of Lowry et al. [25].

Student’s t-test was used for data comparison between different groups according to Hoel [26]. The 0.05 level was selected as the point of minimal acceptable statistical significance.

RESULTS

The effects of Mn proteinate on the activities of antioxidant enzymes: CuZnSOD, MnSOD, CAT, GSH-Px, GST and GR, and the amount of GSH, were

<table>
<thead>
<tr>
<th>ERYTHROCYTES</th>
<th>CuZnSOD</th>
<th>MnSOD</th>
<th>GR</th>
<th>GST</th>
</tr>
</thead>
<tbody>
<tr>
<td>(U g⁻¹ Hb)</td>
<td>control</td>
<td>5314 ± 207</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td></td>
<td>Mn</td>
<td>4020 ± 170  ***</td>
<td>-</td>
<td>-</td>
</tr>
<tr>
<td>LIVER</td>
<td>control</td>
<td>28.0 ± 2.3</td>
<td>4.3 ± 0.1</td>
<td>139 ± 3.1</td>
</tr>
<tr>
<td>(U mg⁻¹ protein)</td>
<td>Mn</td>
<td>32.1 ± 1.8</td>
<td>4.8 ± 0.2 **</td>
<td>151 ± 3.1 ***</td>
</tr>
<tr>
<td>HEART</td>
<td>control</td>
<td>14.0 ± 1.1</td>
<td>9.9 ± 0.2</td>
<td>32.9 ± 0.94</td>
</tr>
<tr>
<td>(U mg⁻¹ protein)</td>
<td>Mn</td>
<td>16.4 ± 1.4 **</td>
<td>8.8 ± 0.8</td>
<td>39.9 ± 1.9 **</td>
</tr>
<tr>
<td>BRAIN</td>
<td>control</td>
<td>32.8 ± 4.1</td>
<td>6.6 ± 0.03</td>
<td>70.0 ± 1.3</td>
</tr>
<tr>
<td>(U mg⁻¹ protein)</td>
<td>Mn</td>
<td>29.4 ± 1.4</td>
<td>6.9 ± 0.2</td>
<td>75.7 ± 2.0</td>
</tr>
<tr>
<td>SMALL INTESTINE</td>
<td>control</td>
<td>12.8 ± 0.9</td>
<td>2.5 ± 0.07</td>
<td>358 ± 10</td>
</tr>
<tr>
<td>(U mg⁻¹ protein)</td>
<td>Mn</td>
<td>11.6 ± 0.7</td>
<td>2.2 ± 0.13</td>
<td>353 ± 5</td>
</tr>
</tbody>
</table>

For definitions of units see the Materials and Methods section. Results are M ± SD, n = 6.

*** P < 0.005; ** P < 0.025

Table 1. Specific activity of CuZnSOD, MnSOD, GR and GST in the erythrocytes, liver, heart, brain and small intestine of the control and Mn-supplemented group.
evaluated in the blood, liver, heart, small intestine and brain of the control and Mn-supplemented rats.

The activities of CuZnSOD, MnSOD, GR and GST in the blood and tissues of control and Mn supplemented rats are presented in Table 1. Activity of CuZnSOD in the erythrocytes was significantly decreased in the Mn-supplemented group (P < 0.005), while in the heart of the same group of animals, treatment with Mn increased CuZnSOD activity, compared with the corresponding control (P < 0.025). Also, MnSOD activity in the liver was increased in this group (P < 0.025). The activity of GR increased after Mn supplementation in the liver (P < 0.005) and heart (P < 0.025). The GR activity in the small intestine was significantly higher (P < 0.005) in animals supplemented with Mn.

GSH-Px activity in the erythrocytes and tissues of rats were depicted in the Figure 1. Decrease in the activity was found after Mn-supplementation in all examined tissues, except in the liver.

CAT activity in rat tissues is presented in the Figure 2. The differences in the CAT activity between the control and Mn-supplemented group were similar than for GSH-Px activity. In all examined tissues the decreases in CAT activity after Mn-supplementation, were found.

The effect of Mn proteinate on GSH content in the plasma and tissues of control and Mn-treated rats is illustrated in the Figure 3. The amount of GSH in all examined tissues of rat (except in small intestine) changed significantly in treated group in comparison to control, but the changes were not in the same direction. In the plasma (P < 0.005) and liver (P < 0.025), of Mn-treated rats, the content of GSH was increased in comparison to the control, but in the heart and the brain (P < 0.005) the significant decrease in GSH content was found.

Discussion

In this study, we have shown that supplementation of rats with Mn proteinates induces tissue-specific changes of antioxidant defense in vivo. Specifically, Mn supplementation increased the activities of CuZnSOD and GR in the heart, MnSOD and GR in the liver, and decreased activity of CuZnSOD in erythrocytes and GST in small intestine. On the other hand, the activities of CAT and GSH-Px decreased in all examined tissues in animals supplemented with Mn. In addition, the content of GSH in the plasma and liver was higher after Mn-supplementation, but lower in the heart and the brain. The effects of observed changes are discussed in more detail below.

In contrast to the number of enzymes activated by Mn, there are few Mn metalloenzymes [27]. One of them is MnSOD, the antioxidant enzyme largely located in the mitochondrial matrix and encoded by nuclear gene. It is essential enzyme for all kind of oxygen-metabolizing cells, and act as an O2•− scavenger. MnSOD expression varies greatly, both among different types of cells and following changes in the environmental conditions. The increased level of MnSOD activity induced by Mn exposure has already been shown [28].

In the present study, the increase in MnSOD activity after Mn supplementation, was found only in the liver. These differences between examined
After supplementation Paynter [27] found considerably greater changes in Mn concentration in liver in comparison to the heart, but the heart showed the greatest relative change in the activity of MnSOD in respond to changing the concentration of Mn in the diet. In this experiment the Mn were supplemented in the form of MnCl₂. Our data indicate increased CuZnSOD activity only in the heart after supplementation with Mn proteinate, and no changes in the activity of MnSOD. This discrepancy may be the consequence of the different form of supplemented Mn (MnCl₂ versus Mn proteinate) and differences in absorption and organ deposition. It is well known that Mn²⁺ could react with O₂•−, and that the nature of the reaction depends on the ligand of the Mn ions. Thus, Mn may scavenging O₂•− directly, or indirectly playing a role in metal-catalyzed dismutation reaction. [29,30,31,32]. The organic form of Mn used in this experiment is amino acid, peptide chelate and the way of absorption and the bioavailability is different in comparison to inorganic Mn salt. Because of that, it is possible that in our experiment more Mn is transported into the heart. The beneficial effect of Mn treatment in the heart was found in the work of Barandier et al. [33] where suppression of early oxidative burst by Mn-supplementation after myocardial reperfusion injury, was observed. This protective effect of Mn against oxidative damage was explained by antioxidant properties of intracellularly accumulated Mn per se, independently of MnSOD. It is also known that Mn forms complexes with lactic, propionic, succinic, pyruvic and pyridine carboxylic acids, which are efficient O₂•− scavengers [27]. It has been shown that lactate accumulates during ischemia in the cardiomyocytes, accordingly, it was postulated that the observed antioxidant effect of Mn might be mediated, at least in part, by acid complex. Thus, it is shown that Mn complexes in heart have scavenger activity. It is of growing interest to use Mn complexes as pharmaceutical SOD agents (SOD mimics) to aid in reducing cellular damage by O₂•− [34]. In the earlier work we shown [35] increase of CuZnSOD in heart and liver, and decreased in MnSOD activity in heart after supplementation with Cu and Zn chelate. Malecki et al. [36] found that, after parenteral administration, uptake of Mn was higher in the peripheral tissues than in the liver.

In the heart we also found decreased activity of CAT and GSH-Px, and the amount of GSH. As in the papers of Coassin et al. [37], and Stacey & Klaassen [38] the peroxyl radical scavenging capacity of Mn (II) was shown. Decreased CAT and GSH-Px activities in the heart as well as in the blood of supplemented rat in our experiment may be explained by this Mn activity. After rat supplementation with Mn, Scheuhammer & Cherian [39] deduced that increased Mn levels in blood were almost totally accounted by increases in the erythrocyte fraction due to the treatment. At the same time, the concentration of Mn in the whole blood is five to 10-folds greater than in serum [12]. In our experiment, in the plasma of supplemented group, but also in the liver, a significant increase in the level of GSH was annotated. It is known that the level of GSH in plasma sustains the level of GSH in liver [28]. The GSH sulfhydryl-containing compound plays a crucial role in detoxification and cellular defense [40]. Higher level of GSH in the plasma of supplemented rats, may be result of tissue elimination of the accumulated GSH conjugates. Changes in GSH in the heart and the brain were in opposite direction. Many metal ions possess a high affinity for sulfhydryl groups and deplete GSH, due to metal binding to glutathione [41].

The same results, decreased amount of GSH, as well as the activity of CAT and GSH-Px we find in the brain of Mn supplemented rats, Liccione and Maines [42] found in rat striatum exposed to Mn. Also, Hussain and Ali [28, 43] got the reduced GSH content in cerebellum as a consequence of administration of MnCl₂. It is well known that Mn can induce neurological damage. Mn mostly accesses the brain via transferrin-dependent mechanism, but also as a free ion. This uptake may be mediated through an endogenous blood-brain barrier Mn²⁺ transporter, or via channels or transporters that are specific for some other ions [44, 45]. Chronic exposure to Mn causes the degeneration of nigrostriatal dopaminergic neurons and Parkinson's syndrome. GSH depletion and upregulation of γ-glutamil transpeptidase, not accompanied by corresponding increase of GSH disulphide levels in the substantia nigra is also depicted after chronic Mn exposure. [42]. As it may be seen, even the low dose of Mn given in organic form, in our experiment made the changes in the activity and content of antioxidant components in the brain of rats.

The change in the activity of GST after Mn-supplementation was found only in small intestine. The acidic intestinal GST, according to Peters et al. [46] is most probably identical with GST Pi isolated from placenta, but in the hepatic preparation this isofrom is hardly detectable. The difference between the effect of Mn supplementation on GST activity in small intestine and in the other examined tissues may be the consequence of different presence of GST isofroms in these tissues.

Conclusions

Treatment with Mn proteinate increased the activity of MnSOD only in the liver. Decreased activities of the most of examined antioxidant enzymes and the amount of GSH were noticed. Exception is the increased GSH amount in the plasma and liver and the increased activity of CuZnSOD in the heart. The observed changes in antioxidant defense after treatment with organic form of Mn, may be explained by scavenging capacity of Mn for peroxyl and superoxide anion radical. According to all, we can assumed that the alterations in antioxidant defense in Mn-supplemented group is a consequence of new prooxidant - antioxidant equilibrium.

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REFERENCES


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