The Effect of Selenium and Lycopene on Oxidative Stress in Bone Tissue in Rats Exposed to Cadmium

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Abstract

This study examined the effect of selenium (Se) or lycopene (Lyc) and their combination on oxidative stress in bone following cadmium (Cd) exposure in vivo. Cd exposure enhanced accumulation of Cd in femur with subsequent increase in LPO and PC and decreased antioxidative enzyme activities in rat femur, along with significant decreases in body and femur bone weights. Subcutaneous (s.c.) injection of Se or Lyc reduced Cd accumulation in bone and increased body and femur weights, when given individually or concomitantly. Antioxidant enzyme activities maintained and/or increased following Se and Lyc supplementation when given individually or concomitantly. However, lycopene dose of 10 mg/kg of body weight alone or selenium alone provided better protection to bone tissues of rats against oxidative stress compared to treatment with their combination. Selenium and lycopene could be used as a dietary supplement to protect against bone oxidative stress in areas exposed to Cd pollution.

Keywords

Selenium, Oxidative Stress, Lycopene, Cadmium

1. Introduction

Industrial development in Saudi Arabia has led to various environmental pollutions especially with heavy metals. 

Badr et al. [1] indicated that pollution with heavy metals including cadmium increased in Jeddah, Rabeg and

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Yunbou (West of Saudi Arabia). Moreover, the pollution of rice consumed in Saudi Arabia has increased due to contamination by cadmium [2]. Study conducted by Al-Farraj and Al-Wabel [3] confirmed the increase in cadmium contamination in soils of Saudi Arabia. They pointed out that contamination of soil with cadmium in Mahd Alzahab (South East of the city of Madinah, Saudi Arabia) comprised 88% of the total heavy metals contamination.

Cadmium is dangerous to humans since it affects many organs in the human body particularly bone [4]. Cadmium may affect bone tissue through disorders due to oxidative/antioxidative balance which leads to oxidative stress and antioxidants deficiency can have a negative impact on bone health, whereas antioxidants may prevent the disorders impacted by the oxidative damage induced by reactive oxygen species (ROS) in the bone tissue [5]. Daily selenium intake of 77.8% breast-fed infants in Saudi Arabia is low [6] compared to US recommendation [7]. Selenium is a known antioxidant essential element [8]. Lycopene is also an antioxidant which is found to decrease the risk of bone resorption markers [9]. The combination of selenium and lycopene could have even more effects compared to the effects of each one alone. Therefore, the aim of this research was to evaluate the effect of selenium and lycopene alone or in combination on oxidative stress in bone tissue in rats exposed to cadmium.

2. Materials and Methods

2.1. Animals and Experimental Protocol

Fifty-four Sprague Dawley male rats (7 weeks old) with an approximate weight of 200 - 250 gm. were utilized in this study. The rodents were housed at King Faisal Specialist Hospital in plastic cages in a temperature-controlled (21°C ± 2°C) room with 12 h light/12 h dark cycle. A water and pellet laboratory animal diet was available ad libitum (Table 1). Initial weights and weekly gain in weight of rats were recorded. The rats were randomly divided into six groups with nine rats per group. Three groups were designated as “control groups” and 3 were designated “treated groups”. The control groups were subcutaneously (s.c.) injected with saline solution or olive oil or Cd (3 mg/kg body weight), respectively. The first treated group was s.c. injected with Se (selenium sodium selenite 99% (Na₂SeO₃)) (3.5 mg/kg) + (s.c.) injected with Cd (3 mg/kg body weight), while the second group was s.c. injected with 10 mg Lyc/kg + (s.c.) injected with Cd (3 mg/kg body weight). The third group was s.c. injected with Se (3.5 mg/kg) and 10 mg Lyc/kg + s.c. injected with Cd (3 mg/kg body weight). The injection of Se or Lyc was 1 h prior to Cd injection. Administration of all doses was daily for the first four weeks; however, due to changes in the skin of the treated rats, doses were given for four days weekly for the remaining eight weeks of the study. Guidelines for animal care approved by the ethics committee of KFSHRS were followed (RAC# 2110019). All animal experiments were maintained in accordance with guidelines for the care and use of laboratory animals of the research centre at King Faisal Specialist Hospital & Research Centre, Riyadh, KSA. Cd and Se were prepared by dissolving in sodium chloride (saline, 0.9% NaCl). Lyc was dissolved in olive oil before it was s.c. injected. The rats were euthanized at the end of the twelfth week using ketamine (80 mg/kg) and xylazine (5 mg/kg) through intramuscular injection. The right femurs were dissected out and the surrounding muscles and tissues were removed [10]. All femur samples were weighed and preserved at −80°C.

<table>
<thead>
<tr>
<th>Dietary Components</th>
<th>% of Laboratory Animal Diet (Net Weight 50 kg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Protein</td>
<td>20%</td>
</tr>
<tr>
<td>Fat</td>
<td>4%</td>
</tr>
<tr>
<td>Fiber</td>
<td>3.05%</td>
</tr>
<tr>
<td>Energy</td>
<td>2770 kcal/kg</td>
</tr>
<tr>
<td>Calcium</td>
<td>0%</td>
</tr>
<tr>
<td>Phosphorus</td>
<td>0.60%</td>
</tr>
<tr>
<td>Vitamin A</td>
<td>20 μg</td>
</tr>
<tr>
<td>Vitamin D</td>
<td>2.20 μg</td>
</tr>
</tbody>
</table>

*Silos and flour mills organization, KSA.
2.2. Preparation of Homogenised Bone Tissue Samples

Known weight slice of femurs were washed out in ice-cold saline (0.9% NaCl) to eliminate the bone marrow. The homogenised samples were prepared as previously described [11] [12]. RIPA buffer, as opposed to potassium phosphate buffer, was used to obtain a 20% bone concentration, with an addition of butyl-hydroxytoluene (BHT; Sigma-Aldrich) and protease inhibitors. The homogenates were centrifuged at 10,000 × g for 20 min at 4°C and the supernatants were collected and used for the measurements of oxidative stress indices and antioxidant status. Analysis was performed in duplicate for all the analyses measured in this study.

2.3. Measurement of Cd Concentration

Slices of the femurs were dried at 64°C for 48 hours, and then digested and pulverised in nitric acid, according to Kingston and Walter [13]. Cd concentration levels were then determined with MS inductively ICP couple plasma mass spectrometry (ICP-MS).

2.4. Measurement of Antioxidants and Oxidation Levels in Bone Tissue

2.4.1. Catalase (CAT) Activity

CAT enzyme activity was assayed according to the method of Aebi [14] using ready-made enzymatic preparations supplied by Sigma Aldrich, USA. The CAT enzyme activity was further determined by measuring the rate of transformation of hydrogen peroxide to water by employing an optical spectrometer (Absorbance Microplate Reader SpectraMax Plus 384, Molecular California Devices, USA) at a wavelength of 520 nm.

2.4.2. Cu-Zn Superoxide Dismutase (Cu-Zn SOD)

Cu-Zn SOD activity was determined according to the method of Kakkar et al. [15] by applying ready-made enzymatic solutions supplied by Cayman Chemical Company (Anne Arbour, MI, USA). An optical spectrometer was used for the analysis at a wavelength of 520 nm.

2.4.3. Cellular Glutathione Peroxidase (GPx)

Cellular GPx was determined using the method described by Rotruck et al. [16] using pre-prepared enzymatic solutions (BioxytechGPx kit 340) supplied by Oix Research Chemical Company, USA. An optical spectrometer was used for the analysis at a wavelength of 340 nm.

2.5. Lipid Peroxidation (LPO)

Malondialdehyde and 4-hydroxyalkenal were used as indicators of oxidation of polyunsaturated fatty acids. The estimated fat oxidation was assayed by colouration of pre-prepared enzymatic solutions supplied by Oix Chemical CO, USA (Biotech LPO-586 kit), and using a spectrometer at a wavelength of 586 nm [17].

2.6. Protein Carbonyl Content (PC)

PC were estimated as previously described [18] using the pre-prepared enzymatic solution (Protein Carbonyl Flurometric Assay Kit, Cyman Chemical Company) and a Spectral Scanning Microplate Reader (Thermo-Lab-systems Varioskan Flash Multimode, Thermo Fisher Scientific Inc., USA).

2.7. Statistical Analysis

Data were analysed using one-way ANOVA and SAS program. Duncan’s multiple range test was used to analyse data and to determine significant differences among treatment means. Values were considered statistically significant when P < 0.05. We determined that a sample size of 8 would detect a difference in bone weight following Cd administration of =25% at P = 0.05 with 80% power. However, a sample size of n = 9 per group was used to account for any animal loss or bad sample collection.

3. Results

3.1. Effects of Cadmium, Selenium and Lycopene on Body Weight

The effect of cadmium on the body weight of rats is shown in Table 2. The body weight of the cadmium treated
Table 2. Body weight and body weight gain in control saline C and rats exposed for 12 weeks to cadmium (Cd), cadmium + selenium (Cd + Se), cadmium + lycopene (Cd + Lyc), cadmium + lycopene + Se (Cd + Lyc + Se).

<table>
<thead>
<tr>
<th></th>
<th>C</th>
<th>Cd</th>
<th>Cd + Se</th>
<th>Cd + Lyc</th>
<th>Cd + Lyc + Se</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial body weight (g)</td>
<td>240 ± 63A</td>
<td>237 ± 4.5A</td>
<td>242 ± 3.3A</td>
<td>235 ± 3.8A</td>
<td>238 ± 3.0A</td>
</tr>
<tr>
<td>Final body weight (g)</td>
<td>349 ± 64B</td>
<td>265 ± 12.2A</td>
<td>285 ± 11B</td>
<td>323 ± 4.0B</td>
<td>294.8 ± 5.4B</td>
</tr>
<tr>
<td>% Body weight gain (12 weeks)</td>
<td>46 ± 2.9a</td>
<td>9 ± 1.3d</td>
<td>24 ± 3.6c</td>
<td>38 ± 1.9b</td>
<td>24 ± 2.01c</td>
</tr>
</tbody>
</table>

Results are presented as mean ± SE; Means not sharing a common superscript a big letters in column are significantly different at P < 0.05; Means not sharing a common superscript a small letters in row are significantly different at P < 0.05 as assessed by Duncan’s multiple-range test.

Rats increased slightly (9%) but the body weight of the other two control treated rats increased significantly (P ≤ 0.05) at the end of the experimental period. Chen et al. [19] indicated a decrease in the body weight of male Sprague Dawley rats treated with cadmium at a dose of 1.5 mg/kg body weight. Rabbits treated with oral dose of 5 mg cadmium/kg b.w. showed significant (P ≤ 0.05) decrease in body weight after four months when compared to the control group. On the other hand, treatment of the male rats with selenium and lycopene either alone or in combination caused statistically (P ≤ 0.05) increase in body weight (Table 2). The increase in body weight was higher in the rats treated with 10 mg lycopene/kg b.w. than the other treated groups as well as the cadmium treated one. These results agree with results reported by Messaoudi et al. [20], who reported that selenium increased the body weight of Wistar rats exposed to cadmium. The increase in body weight of rats in this study also agree with the finding of Rencuzogullari and Erdogan [21], who showed an increase in the body weight of rats exposed orally to cadmium and given a dose of 10 mg lycopene/kg b.w.

3.2. Effects of Selenium and Lycopene on Cadmium Concentration in Femoral Bone Tissue and on Its Weight

Injection of the rats with dose of 3 mg cadmium/kg b.w. for 12 weeks caused accumulation of this element in femoral bone tissue. Cadmium concentration increased significantly (P ≤ 0.05) in these rats (about 574 folds) compared to the other two control groups given saline solution and olive oil, respectively (Table 3). Results of this study agree with the finding of Brzoska et al. [5], who indicated similar increase in bone tissue of rats given orally a dose of 2.5 mg cadmium/kg b.w. for six months.

A significant (P ≤ 0.05) decrease in cadmium concentration in femoral bone tissue was observed in rats administered selenium and lycopene either alone or in combination. However, selenium and lycopene alone had the greatest effect than their combinations. Data are lacking in the literature (to the best of our knowledge) on the effects of selenium and lycopene on cadmium concentration on bone tissue and this study report such data for the first time. Due to the lack of such data comparison in this study was made with studies evaluated their effects on other tissues. Brzoska et al. [10] studied the effect of zinc (60 mg/L of drinking water) on cadmium concentration of the femoral bone tissue and found that zinc reduced the cadmium concentration by 18%. Alhazza’s study [22] also indicated a reduction in cadmium concentration on the testes of rats exposed to cadmium and injected by selenium (0.35 mg/kg b.w.).

Femoral bone weight decreased significantly (P ≤ 0.05) in the group of rats exposed to cadmium compared to control one (Table 3). Selenium and lycopene either alone or in combination reversed the effect of cadmium on bone weight. Selenium and lycopene (10 gm/kg b.w.) alone and selenium plus lycopene (15 gm/kg b.w.) had the greatest effect in reversing the loss of bone weight; however, full attainment of bone weight was not accomplished by the use of selenium and lycopene alone or in combination. Brzoska et al. [10] indicated that zinc given to rats exposed to cadmium increased bone weight after 12 months of the treatment. Selenium with vitamin E or C improved also skeletal changes of the bone of the femoral osteoporotic rabbits [23].

3.3. Effects of Selenium and Lycopene on Oxidative Enzymes

The effects of selenium and lycopene either alone or in combination on CAT of male rats treated with cadmium are shown in Table 4. Cadmium administration for 12 weeks reduced CAT activity in rats bone tissue by 60% (from 30 to 18 µmole/min/mg of protein). Selenium and lycopene alone preserved CAT activity completely, whereas their combinations preserved most of this activity (Table 4). Study by Messaoudi et al. [24] revealed that zinc and selenium either alone or in combination preserved CAT activity in the red blood cells of rats ex-
Table 3. Cadmium concentration (Cd conse) in femoral bone and femoral bone weight of control saline C and rats exposed for 12 weeks to cadmium (Cd), cadmium + selenium (Cd + Se), cadmium + lycopene (Cd + Lyc), cadmium + lycopene + Se (Cd + Lyc + Se).

<table>
<thead>
<tr>
<th></th>
<th>C</th>
<th>Cd</th>
<th>Cd + Se</th>
<th>Cd + Lyc</th>
<th>Cd + Lyc + Se</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cd conse mg/kg dry weight</td>
<td>13.3 ± 1.8&lt;sup&gt;a&lt;/sup&gt;</td>
<td>7652 ± 254&lt;sup&gt;b&lt;/sup&gt;</td>
<td>4392 ± 123&lt;sup&gt;c&lt;/sup&gt;</td>
<td>4565 ± 275&lt;sup&gt;d&lt;/sup&gt;</td>
<td>5798 ± 229&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td>Femoral bone weight gm</td>
<td>1.02 ± 0.023&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.69 ± 0.014&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.95 ± 0.020&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.90 ± 0.020&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.81 ± 0.021&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Results are presented as mean ± SE; Means not sharing a common superscript letters in a row are significantly different at P < 0.05, as assessed by Duncan’s multiple-range test.

Table 4. Catalase (CAT), copper zinc superoxide dismutase (Cu-Zn SOD) activities, and glutathion peroxidase (GPx) in bone tissue of control saline C and rats exposed for 12 weeks to cadmium (Cd) cadmium + selenium (Cd + Se), cadmium + lycopene, cadmium + lycopene + Se (Cd + Lyc + Se).

<table>
<thead>
<tr>
<th>Antioxidant activity</th>
<th>C</th>
<th>Cd</th>
<th>Cd + Se</th>
<th>Cd + Lyc</th>
<th>Cd + Lyc + Se</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAT Umol/min/mg</td>
<td>29.6 ± 1.40&lt;sup&gt;a&lt;/sup&gt;</td>
<td>19 ± 0.88&lt;sup&gt;b&lt;/sup&gt;</td>
<td>30 ± 1.13&lt;sup&gt;c&lt;/sup&gt;</td>
<td>30.3 ± 0.73&lt;sup&gt;d&lt;/sup&gt;</td>
<td>22.3 ± 0.70&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td>Cu-Zn SOD U/mg</td>
<td>0.28 ± 0.01&lt;sup&gt;a&lt;/sup&gt;</td>
<td>0.20 ± 0.02&lt;sup&gt;b&lt;/sup&gt;</td>
<td>0.32 ± 0.20&lt;sup&gt;c&lt;/sup&gt;</td>
<td>0.29 ± 0.02&lt;sup&gt;d&lt;/sup&gt;</td>
<td>0.31 ± 0.01&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
<tr>
<td>GPx mu/mg protein</td>
<td>4.11 ± 0.3&lt;sup&gt;a&lt;/sup&gt;</td>
<td>1.9 ± 0.1&lt;sup&gt;b&lt;/sup&gt;</td>
<td>4.5 ± 0.15&lt;sup&gt;c&lt;/sup&gt;</td>
<td>4.4 ± 0.31&lt;sup&gt;d&lt;/sup&gt;</td>
<td>4.9 ± 0.35&lt;sup&gt;e&lt;/sup&gt;</td>
</tr>
</tbody>
</table>

Results are presented as mean ± SE; Means not sharing a common superscript letters in a row are significantly different at P < 0.05, as assessed by Duncan’s multiple-range test.

Exposed to cadmium but the effect of the combination of these two elements was less than that observed with either of them alone. On the other hand, El Heni et al. [25] found that exposure to cadmium caused a decrease of 58.5% in CAT activity in the liver of rats; however, selenium and lycopene neither alone nor in combinations had significant effects in the activity of this enzyme.

Treatment of male rats with cadmium also reduced significantly (P ≤ 0.05) the activity of SOD in bone tissue, whereas the treatment of these rats with selenium and lycopene either alone or in combination increased significantly (P ≤ 0.05) the activity of this enzyme (Table 4). These results do not agree with the findings of El Heni et al. [25], who stated that selenium had no effect in SOD activity of rats exposed to cadmium. However, these researchers used low dose of selenium (0.1 mg/L of drinking water). Moreover, selenium was given orally in their study.

GPx activity also reduced significantly (P ≤ 0.05) in the bone tissue of rats treated with cadmium and treatments with selenium and lycopene either alone or in combination increased significantly (P ≤ 0.05) the activity of this enzyme (Table 4). Despite the beneficial effects of the all treatment with lycopene and selenium in increasing the activity of GPx, it seems that combination of both of them had higher but not significant affect compared to the effect of each of them alone (Table 4). Moreover, increasing the dose of lycopene to 15 mg/kg b. w. had no extra beneficial effect on the activity of antioxidant enzymes.

Ognjanovic et al. [26] studied the effect of selenium in the kidney and liver of Wistar rats exposed to cadmium and found that selenium increased significantly (P ≤ 0.05) the activity of GPx in these two tissues.

3.4. Effects of Selenium and Lycopene on Oxidative Status

Measuring the activity of antioxidant enzymes as well as indicators of oxidative stress such as Cu-Zn SOD/GPx, malondialdehyde and protein carbonyl concentration are important to know the balance between the first and the second stage of the function of antioxidant enzymes [27]. This means the balance between ROS production and their destruction [28].

Lipid peroxidation was increased significantly (P ≤ 0.05) by 79% in rats exposed to cadmium compared to rats in the control group; however, both selenium and lycopene either alone or in combination prevented the effect of cadmium and the lipid peroxidation level in the treated rats maintained to its normal level as in the control group (Table 5). This result agrees with the finding of several researchers [5] [21] [29].

Cadmium not only oxidized lipid but also protein and this is indicated by the increase in protein carbonyl concentration in bone tissue of rats exposed to this element (Table 5). Such affect indicates the breakdown of protein in bone especially collagen [5]. Again selenium and lycopene either alone or in combination prevented such effect; however, selenium and lycopene (10 mg/kg b.w.) had the greatest effect compared to the other treatments. These results agree with the finding of Brzoska et al. [5].
Table 5. Malondialdehyde (MDA), protein carboxyl (PC) concentration in bone tissue of control saline C and rats exposed for 12 weeks to cadmium (Cd) cadmium + selenium (Cd + Se), cadmium + lycopene of body weight (Cd + Lyc), cadmium + lycopene of body weight.

<table>
<thead>
<tr>
<th>Oxidative status</th>
<th>C</th>
<th>Cd</th>
<th>Cd + Se</th>
<th>Cd + Lyc</th>
<th>Cd + Lyc + Se</th>
</tr>
</thead>
<tbody>
<tr>
<td>LPO µm/mg protein</td>
<td>0.24 ± 0.01b</td>
<td>0.43 ± 0.02a</td>
<td>0.21 ± 0.01b</td>
<td>0.25 ± 0.013b</td>
<td>0.22 ± 0.02a</td>
</tr>
<tr>
<td>PC nmol/mg protein</td>
<td>1.6 ± 0.14c</td>
<td>2.9 ± 0.23a</td>
<td>1.6 ± 0.12c</td>
<td>1.3 ± 0.13c</td>
<td>2.2 ± 0.20b</td>
</tr>
</tbody>
</table>

Results are presented as mean ± SE; Means not sharing a common superscript letters in a row are significantly different at P < 0.05, as assessed by Duncan’s multiple-range test.

4. Discussions

Cd is an environmental pollutant that causes harm to the overall health of human beings. It adversely affects many of the body’s tissues, including bone [30]-[32]. The importance of nutritional supplements such as phytochemicals on lowering the harmful effects of Cd on the liver and kidney had been reported [29] [33]. These studies are consistent with the research of Brzoska et al. [5], who concluded that oxidative stress in bone tissue is mainly from Cd exposure and may be responsible for the increase in the amount of people afflicted with bone diseases. Therefore, the aim of this study was to find ways to reduce the oxidative stress levels in bone tissue resulting from Cd exposure; specifically, to study the individual and concomitant effects of Se and Lyc on oxidative stress levels in the bone tissue of laboratory rats that had been exposed to Cd.

The measurement of body weight and the rate of weight gain are important in toxicological studies [34], as they are considered indicators of the extent of deterioration of health. A decrease in these factors is also an indicator of the deficiency in the biological processes in the body [35]. In the present study, exposure to Cd did not lead to a significant increase in body weight gain after 12 weeks from the commencement of the study. This indicates a disruption in the growth of the Cd-exposed rats. These results are consistent with the findings of Chen et al. [19], which indicated a decrease in the body weight of Sprague Dawley rats that were subcutaneously injected with a dosage of 5 mg/kg of Cd. This study also confirms the findings of Yousef et al. which demonstrated a significant reduction in the body weight of laboratory rabbits that were administered a dosage of 5 mg/kg of oral Cd for one month [35].

The reduction in body weight in the Cd-treated group observed could be due to a lack of appetite caused by the high dosage levels. However, the findings could also be attributed to the impact that Cd exposure had on weight of femurs and other tissues [10] [36] [37]. Interestingly, the reduction in the weight of bone tissue has been associated with Cd as a result of its effect on several body mechanisms resulting in the generation of free radicals [38]. For this reason, treatment with Se and Lyc, when administered separately or concomitantly, decreased the oxidative stress and enhanced or maintained antioxidant enzyme activities in the Cd treated rats, thus improving rat food palatability, food intake and (eventually) their body weight gain (Table 2).

The impact of Cd on bone tissue is dependent on the time of exposure as well as the age of the animal. The present study demonstrated that exposure of rats to Cd with an age from seven weeks to five months led to the deposition of high concentrations of Cd in the femur. Brzoska et al. [39] found that exposure to Cd increased its concentration levels in the bone only during the most intensive phase of skeletal growth, that is, up to the 4th month of the animal’s life. These results are consistent with other studies which demonstrated higher concentrations of Cd were found in the bones of younger rats than in older ones [40] [41]. This highlights the seriousness of exposure to Cd at an early age, which is an important stage in the development of bone. Exposure to Cd may impede bone development, causing damage with advancing age. One of the possible reasons for the increased Cd concentration in bone could be attributed to its interference in the formation of hydroxyapatite (Ca_{10}(PO_{4})_{6}(OH)_{2}) which is responsible for the strengthening of the bone tissue, by replacing calcium ions with Cd ions [42]. A link may exist between Cd and various important proteins in the body in that Cd plays a role in raising the protein concentration of bone tissue. Cd binds to ions that compete with existing thiols to bind with some proteins. When the synthesis of these proteins decreased, the free Cd ions in tissue increased, and started to affect bones and other defense systems in the body [43] [44]. Thus, the increase in ions may be the result of the high concentration of Cd in the bone tissue. Therefore, Se and Lyc, either separately or concomitantly, played a significant role in reducing the rate of deposition of Cd (Table 3). Furthermore, these antioxidants may contribute to the reduction of Cd ions that bond with important bone components. Bone tissue has defense mechanisms that protect it from the effect of reactive oxygen species (ROS). These defense mechanisms include...
antioxidant enzymes such as CAT, SOD, and GPx. GPx represents the body's main defense system that activates cellular antioxidant enzymes in the bone tissue. It is responsible for restricting LPO, and interacts with CAT in converting hydrogen peroxide (H₂O₂) to oxygen (O₂), whereas SOD is accountable for converting O₂•⁻ to O₂ and H₂O₂ [45].

This study showed an inverse relationship between antioxidant enzyme activities and Cd concentrations in femoral tissue (Table 4). The significant decline in antioxidant enzymes (CAT, Cu-ZnSOD, and GPx) could be due to Cd bonding with enzymes that contain -SH- functional groups, which are meant to defend tissues leading to reduced activity [46]. This could also be through another mechanism resulting from the impedance of Cd activity by replacing the activity of beneficial substances [47]. Another possibility is that Cd depleted the vital elements necessary for the functioning of these enzymes, such as Se, zinc, copper, manganese and iron. CAT contains iron in its active site [45]. Cd may play a role in decreasing iron levels and, as a result, decreased CAT activity. The possibility that Cd may led to a lack of nutrients such as zinc and copper, which are fundamentals to the formation of SOD, should also be considered [45]. Brzoska et al. indicated that medium and high exposure to Cd reduces the bone tissue’s content of zinc, copper and iron [5] [10]. Se also had a positive impact on the activity of Cu-Zn SOD [48]. Therefore, the decline in the activity of GPx may result from the high concentrations of Cd in the bone tissue and, as a result, this could be the cause of the depletion of Se which is essential for GPx synthesis [45].

The results of this study also showed that Se and Lyc, individually or concomitantly, had a role in the increase in the activity of antioxidant enzymes in the femoral tissue of rats. This may explain the role of each in reducing the concentration of Cd in bone tissue. On the other hand, administration of Se and Lyc alone maintained a normal level of CAT activity compared to the other combination group. This may be the reason why these two groups showed more reduction in Cd concentrations in bone tissue compared to the other treated groups. Se maintained the activity of Cu-Zn SOD and GPx by binding directly with Cd and converting it to an inactive complex compound called Cd complex selenide [50]. Another possibility is that it reduced the Cd link in the active site of these enzymes and weakens their activities. The function of Se is to protect bone tissue from oxidation by protecting antioxidant enzymes. This decreases their vulnerability to Cd and stops the depletion of the elements necessary to build the active site for these enzymes [45]. Se also plays an important role in building seleno-proteins like GPx [51]. The role of Se in increasing GPx activity in the bone tissue that protects DNA from oxidation was evident in this study.

This study also highlighted the role of Lyc, either alone or in combination with Se, in increasing antioxidant enzyme activities and suggests the possibility of Lyc being an antioxidant compound which protects tissues from the effects of free radicals. Additionally, by reducing the concentration of Cd in bone tissue, it may reduce tissue damage and the loss of the elements necessary for the formation of antioxidant enzymes.

In addition, Cd treatment led to an increase in LPO in femur bone tissue (Table 5). This could be attributed to the role of Cd in decreasing the activity of antioxidant enzymes in the tissue responsible for scavenging free radicals. The continued presence of free radicals may be the reason for the oxidation of lipids in the cell walls [52]. The results of this study are consistent with previous results showing that the decline in the activity of GPx was followed by a significant rise in LPO [52]. This supports the role of Se and Lyc in reducing LPO by increasing the activity of GPx and thioredoxin reductase, both of which are important enzymes for the elimination of LPO [29] [51]. Supplementation with Se and Lyc prevented bone lipid from oxidative damage, possibly as a result of their role in maintaining the activity of antioxidant enzymes. This either indicates a reduction in hydrogen peroxide concentration, or that the enzymes became efficient in reducing ROS and, thus, lowering the concentration of LPO. Cd treatment not only led to lipid oxidation, but also to the oxidation of proteins (Table 5). A significant increase was observed in PC concentration in the bone tissue of the Cd treated group. This indicates destruction of the proteins that are present in femoral tissue including collagen [5]. The low PC concentrations in the femoral of rats receiving Se or Lyc could be as a result of their role in maintaining the CAT activity (Table 4) and decreasing Cd accumulation in the femoral tissue (Table 3).

5. Conclusion
This study demonstrated that Se and Lyc supplementation reduced Cd concentrations in femoral tissue and also
increased or maintained the antioxidant enzyme activities by reducing bone oxidative stress. Se and Lyc could be sufficient to reduce the toxicity of Cd in areas exposed to Cd pollution and could be recommended as a dietary supplement to protect against bone oxidative stress. Further studies are needed to confirm the protective effect of bone tissues by Se and Lyc.

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