Diazinon-Induced Ovarian Toxicity and Protection by Vitamins E

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Received: 03.03.2014
Accepted: 23.04.2014

ABSTRACT

Background: DZN (diazinon) is an organophosphate insecticide that had been used in agriculture and for domestic and veterinary use for several years and caused many negative effects on plants and animal species, especially on human. The aim of present study was to evaluate the effects of DZN on MDA (malondialdehyde) and GSH (glutathione) levels in female rat reproductive tissue (ovary) and to assess the protective role of vitamin E.

Methods: A total of 30 adult female Wistar rats were divided into five groups: control group (without any intervention), sham group (received only pure corn oil, as solvent), experimental group 1 (DZN+corn oil, 60 mg/kg), experimental group 2 (vitamin E, 200 mg/kg), and experimental group 3 (DZN+vitamin E, the same dosage). All drugs were injected intraperitoneally, except vitamin E which was administrated by gavage. The animals were scarified after two weeks and MDA as a marker of lipid peroxidation and GSH content were measured in ovarian tissue.

Results: DZN reduced GSH content and increased MDA level in ovary compared with the control group (P<0.001). Vitamin E plus DZN increased GSH content but decreased DZN-induced MDA elevation in rat ovarian tissue.

Conclusion: Oxidative stress contributes to DZN-induced ovarian toxicity. The results of this study suggested that vitamin E may have a protective effect on DZN-induced ovarian toxicity.

Keywords: Diazinon, Glutathione, Malondialdehyde, Ovary, Vitamin E.

INTRODUCTION

Pesticides and herbicides are persistent and dangerous chemical compounds that are used extensively in agriculture to enhance food production. Among them, pesticides organophosphates are commonly used as insecticides. They are generally the most toxic pesticides for animal species, especially vertebrate animals [1]. Organophosphorus compounds have been utilized in agriculture, industry, medicine, farming, animal keeping, and households to kill insects, worms, nematodes, fungi, and weeds for five decades [2-5].

Unfortunately, the improper use of these products damages the environment- plants, animals, soil, and water- and creates resistance to them which is an increasing concern in this field [6,7]. Residual amounts of organophosphate insecticides are detectable in soil, tissues of organisms, vegetables, grains, and other foods products [7]. On the other hand, it has been reported that organophosphate insecticides negatively affect different tissues, including liver, kidney, immune system, pancreas, and cardiac and vascular walls, and induce hematological and biochemical changes. One of the affected organs is the ovary, which has a key role in reproductive function by synthesizing hormones and producing oocyte [8, 9]. DZN (0,0-diethyl-0-[2-isopropyl-6-methyl-4-pyrimidinyl] phosphorothioate) is a pesticide with a wide range of action which inhibits acetylcholinesterase (AchE) activity. It has been widely used throughout the world with applications in agriculture and horticulture for controlling insects in crops, ornamentals plants, lawns, fruits, vegetables, and food products. It is also used as a pesticide in household and agricultural public health, and for veterinary purposes [3, 7, 10-12]. Research has shown that DZN can be absorbed through the digestive system, skin, or via the respiratory mucosa when inhaled [13]. In addition to its inhibitory effects.
on AChE, it can increase formation of free radicals and so induce oxidative stress and tissue lipid peroxidation in mammals and other organisms [14,15].

Reactive Oxygen Species (ROSs) are part of normal oxidative metabolism, but they can cause tissue damage as lipid peroxidation when produced in large amounts [16]. MDA is a major oxidation product of peroxidized polyunsaturated fatty acids and elevated MDA content is a significant indicator of lipid peroxidation [7,14]. Thiol groups are sensitive to oxidative damage and normally decrease in response to oxidative stress [17]. GSH is one of the thiol groups that has an essential role in protecting cells from damage induced by oxidative stress [18].

Antioxidants, on the other hand, are scavengers that detoxify excessive ROS and play an important role in maintaining oxidant/antioxidant balance in the body. Antioxidants are of two types: enzymatic and nonenzymatic [15]. Vitamin E (α-tocopherol) is a family of lipid-soluble vitamins and acts as an antioxidant in cells by protecting cellular membranes and lipoproteins from peroxidation. In addition, several studies have shown that α-tocopherol inhibits free radical formation and so decreases lipid peroxidation in biological systems [5]. The aim of this study was to investigate the effects of DZN, an organophosphate insecticide, on MDA and glutathione levels and evaluate the protective role of vitamin E in rat ovarian tissue.

**MATERIALS AND METHODS**

**Animals**

A total of 30 adult female Wistar rats were obtained from the Animal Lab of Mashhad University of Medical Sciences, Mashhad, Iran. The animals were divided into five groups: control (without any intervention), sham group (received only pure olive oil, as solvent), experimental group 1 (DZN+ olive oil, 60 mg/kg), experimental group 2 (Vitamin E, 200 mg/kg), and experimental group 3 (DZN+vitamin E, with the same dose). In this study, olive oil was used as solvent. DZN and solvent were administrated by intraperitoneal injection and Vitamin E was given by gavage. All of these animals were scarified after two weeks and MDA, as a marker of lipid peroxidation, and GSH content were determined in ovarian tissues.

**Chemicals**

DZN was diluted in olive oil. Malondialdehyde, thiobarbituric acid, Malondialdehyde tetrabutylammonium, reduced glutathione (GSH), DTNB [5,5′-Dithiobis(2-nitrobenzoic acid)] and vitamin E (α-tocopherol acetate) were purchased from Sigma Co.

**Lipid Peroxidation Test**

The amount of lipid peroxidation was assessed through the measurement of MDA levels in ovarian tissues. MDA reacts with thiobarbituric acid (TBA) and produces a pink colored complex which has the maximum absorbance at 532nm. Initially, 3 ml of phosphoric acid (1%) and 1 ml of TBA (0.6%) were added to 10% homogenized tissue in KCl and, then, the compound was heated for 45 min in a boiling water bath. After cooling the compound, 4 ml of n-butanol was added to it and vortex-mixed was used for 1 min followed by centrifugation at 3000 g for 10 min. After that, the organic layers were removed and transferred to other tubes and absorbance level was read at 532 nm [19]. A calibration curve was designed using Malondialdehyde tetrabutylammonium. MDA levels were expressed by nmol/g tissue.

**Reduced Glutathione (GSH) Examination**

GSH was evaluated in ovarian tissue through the method employed by Moron et al. [20]. The basis of this was the formation of yellow color after adding DTNB [5,5′-Dithiobis(2-nitrobenzoic acid)] to compounds containing sulphydryl groups. For this purpose, 300 μl of homogenates tissues was blended with 300μl of 10% tricloroacetic acid (TCA) and vortexed. After centrifugation at 2500 g for 10 min, the upper layers were removed and blended with reaction mixtures containing 2 ml phosphate buffer (pH: 8) and 500μl DTNB. After 10 min, the absorbance was evaluated at 412 nm using a spectrophotometer (Jenway 6105 uv/vis, UK). At the end of this process, the amount of GSH was determined based on a standard curve drawn with commercially available GSH and GSH Levels were expressed by nmol/g tissue.

**Statistical Analysis**

Results are expressed as mean±SD. Statistical analysis was performed with ANOVA.
followed by Tukey–Kramer test to compare the differences between means. Differences were considered statistically significant at P<0.05.

RESULTS

Effect of Vitamin E on Ovary Lipid Peroxidation Induced by Diazinon

The results of the present study indicated that MDA level significantly increased in the DZN treated group compared to the control group (P<0.001). MDA level decreased significantly in vitamin E+DZN-treated group compared with the DZN group. Administering vitamin E (200 mg/kg) alone also significantly decreased the level of MDA compared with DZN treated rats (P<0.001). Administration of vitamin E (200 mg/kg) and oil (60 mg/kg) alone had no significant effects on the MDA content compared to control group (Figure 1).

Effect of Vitamin E on GSH Content in Ovarian Tissue Following Exposure to Diazinon

The results also revealed significant decreases in GSH content in the DZN-treated group compared with the control group (P<0.001). GSH content increased significantly in the vitamin E+DZN-treated group compared with the DZN group (P<0.01). Administering vitamin E (200 mg/kg) alone significantly increased the GSH content in comparison with the DZN treated rats (P<0.001). Administration of vitamin E (200 mg/kg) and oil (60 mg/kg) alone had no significant effects on the GSH content compared to the control group (Figure 2).

Figure 1. Effects of DZN and vit E on MDA levels in groups. 
*** P< 0.001 vs. control, ### P< 0.001 vs DZN

Figure 2. Effects of DZN and vit E on GSH levels in groups.
*** P< 0.001 vs. control, ### P< 0.001 vs DZN, ## P< 0.01 vs DZN
DISCUSSION

DZN is not only used in pest control of fruits and plants, but it is also used as an ectoparasiticide for cattle in veterinary applications [21]. DZN is absorbed from the gastrointestinal tract and rapidly metabolized [8]. Organophosphate insecticides induce biochemical and histopathological changes in several organs, such as liver, kidney, immune system, pancreas, and cardiac and vascular walls [22,23]. One of the targeted organs is the ovary, which has a major role in reproductive function by synthesizing hormones and producing oocyte [24, 9]. This study was done to investigate the effects of DZN, an organophosphate insecticide, on MDA and GSH levels and assess the protective role of vitamin E in rat ovarian tissues. One of the mechanisms of toxicity is increasing oxidative stress by the generation of free radicals and producing tissue lipid peroxidation. The intensity of lipid peroxidation was assessed through the measurement of MDA levels in ovarian tissues. The increased MDA content is a significant indicator of lipid peroxidation. This study showed that exposure to DZN (60 mg/kg) for two weeks significantly increased MDA level as compared with the control group (P<0.001). Ogutcu et al. showed that DZN raises MDA level in heart tissues of rats, which is an indicator of free radical formation in this organ [7]. Koc et al. confirmed that endosulfan and malathion exposure increase MDA levels in ovarian tissues of female rats which is an indicator of free radicals formation and lipid peroxidation during metabolism of these insecticides [25,26]. Jahromi et al. found that malathion which is one of the organophosphorus insecticides also may increase MDA level in ovary which is a definite reaction caused by released radicals from body and lipids metabolisms [4]. Previous studies have shown that DZN raised the lipid peroxidation level in erythrocytes and pancreas tissues of rats [7,13,27]. Several studies have also indicated that organophosphate insecticides increase MDA level in hepatic, cerebral, pulmonary, pancreatic, intestinal, and cardiac tissues and these changes are accompanied by increasing antioxidant system activities in some tissues [28,29].

In the present study, exposure to DZN (60 mg/kg) significantly decreased GSH content in ovary compared to the control group (P<0.001).

Manna et al. reported that reduction of GSH content is an important indicator of oxidative damage in heart tissues [30]. Razavi et al. reported that exposure to DZN significantly increases MDA levels and decreases GSH contents in heart tissues [31]. The previous studies have also indicated that exposure to fenthion, an organophosphate insecticide, decreases GSH level and increases MDA level in erythrocytes of mice and rats [29,32]. Shah et al. reported that DZN administration induces lipid peroxidation in a dose-dependent manner in rats. He also demonstrated that DZN treatment decreases renal GSH, reduces the activities of antioxidant enzymes, including the enzymes involved in GSH metabolism and large production of oxidants which is associated with renal damage; all of which are involved in the cascade of events leading to DZN-mediated renal oxidative stress and toxicity [1]. Cells have several ways to reduce the effects of oxidative stress, through repairing damage (damaged nucleotides and lipid peroxidation created by products) or reducing oxidative damage by enzymatic and non-enzymatic antioxidants [7]. Vitamin E, a non-enzymatic antioxidant, is a lipid-soluble vitamin present in biological membrane [33,34]. Vitamin E allows free radicals to remove a hydrogen atom from the antioxidant molecule rather than from polyunsaturated fatty acids; thus, it breaks the chain of free radical reactions; the resulting radical is relatively inactive [35]. Minerals and vitamins are essential for the normal functioning of the reproductive system [36]. In the present study, vitamin E was administrated via gavage for 14 days. The experimental group 2 received vitamin E plus DZN while experimental group 3 received vitamin E alone. Vitamin E (200 mg/kg) plus DZN significantly decreased the level of MDA compared to DZN treated rats (P<0.001) and increased the GSH content compared to DZN treated rats (P<0.01). Sutcu et al. have also shown that DZN increased lipid peroxidation in erythrocytes of rat and combinations of vitamin C and E reduce lipid peroxidation [37]. Yilmaz et al. reported that oxidative stress contributes to DZN-induced brain toxicity and vitamins E plus C combination may have a protective effect on this toxicity [38]. Kalender et al. observed that endosulfan administration increases MDA level in rats and
vitamin E protects cells and cellular structures from oxidative damage by reducing the MDA level [39]. A different study reported that vitamins C and E prevent hepatotoxicity induced by methyl parathion in rats [34]. Ogutcu et al. found that vitamin E reduces MDA level in DZN-induced heart tissue toxicity but does not prevent toxicity completely [7].

CONCLUSION

Based on the findings of this study, it can be concluded that DZN causes toxicity in ovarian tissues by inducing oxidative stress, reducing GSH content, and increasing MDA level. Vitamin E can protect ovarian tissues against this toxicity by reducing MDA level and increasing GSH content.

ACKNOWLEDGMENTS

The present study was part of MSc thesis in anatomical research. This study was conducted under contract NO. 911097 supported by a grant from the Research Council of Mashhad University of Medical Sciences. The authors wish to declare their gratitude to Research the Council of Mashhad University of Medical Science and Mrs. Motajadded and Mrs. Ziaee.

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