Modulatory Effects of Vitamin C and E on Cypermethrin-Induced Cardiac and Hepatic Damage in Female Wistar Rats

Oladele JO1,3, Adewale OO2, Oywelo OI*, Oyeleke OM3, Ilori OT1, and Olayinka OE1

1Phytomedicine and Toxicology Laboratories, Department of Biochemistry, Faculty of Basic and Applied Sciences, Osun State University, Osogbo, Nigeria
2Cancer Research and Molecular Toxicology Laboratories, Department of Biochemistry, Faculty of Basic and Applied Sciences, Osun State University, Osogbo, Nigeria
3Biochemistry Unit, Department of Chemical Sciences, Faculty of Sciences, Kings University, Ode-Omu, Nigeria

*Corresponding author: Oywelo OI, Phytomedicine and Toxicology Laboratories, Department of Biochemistry, Faculty of Basic and Applied Sciences, Osun State University, Osogbo, Nigeria; E-mail: ioluoye@yahoo.com, jo.oladele@kingsuniversity.edu.ng

Abstract

The rationale behind this study was to determine the possible ameliorating effect of Vitamin C and E on Cypermethrin induced toxicity in heart and liver function of female albino rats. Twenty-eight female albino rats were sorted into four groups of seven rats per groups were used in this study. Group A serves as the control and received distilled water orally. Group B, C and D were administered 25mg/kg body weight cypermethrin orally. Group C and D were treated daily with 40mg/kg body weight vitamin C and 20mg/kg body weight vitamin E respectively by oral administration while group B was left untreated for 14 days. Cypermethrin significantly (P<0.05) induced cardiac and hepatic damage as characterized by elevated levels of serum alanine aminotransferase (ALT), aspartate aminotransferase (AST), hepatic bilirubin and decrease in cardiac albumin, globulin, total protein, ALP, LDH and GGT. However, treatment with vitamin C and E significantly ameliorate the adverse effects of Cypermethrin. Histological examination of the cardiac and hepatic tissues correlates with the biochemical results indicating marked distortions in the architecture of heart and liver of rats administered Cypermethrin alone but treatment with vitamin C and E attenuated these alterations. In conclusion, vitamin C and E effectively suppress the toxicity effects of Cypermethrin in heart and liver of rats by scavenging the free radicals and protecting the membrane integrity of the tissues.

Keywords: Cypermethrin; Vitamins; Transferase; Cardiac Damage; Hepatic Damage

Introduction

The deleterious effect of agrochemicals and pollutants has its impact on all living organism. The water pollutants include industrial, agricultural and urban discharges that have hazardous effect on aquatic organisms especially when these contaminants are slight decomposers and exhibit high potential for accumulation with synergistic effect [1]. Cypermethrin has become one of the most important insecticides in wide scale use. It has wide uses in cotton, cereals, vegetables and fruit, for food storage, in public health and in animal husbandry.

Cypermethrin, an analogue of pyrethrins is classified by the World Health Organization (WHO) as moderately hazardous (class II). It interacts with the sodium channels in nerve cells through which sodium enters the cell in order to transmit a nerve signal. These channels can remain open for up to seconds compared to the normal period of a few milliseconds, after a signal has been transmitted.
Advances in Clinical Toxicology

The animals were divided into four groups of seven rats per groups. Group A serves as the control and received distilled water orally. Group B, C and D were administered 25mg/kg body weight cypermethrin orally. Group C and D were treated daily with 40mg/kg body weight vitamin C and 20mg/kg body weight vitamin E respectively by oral administration while group B was left untreated. Animals were kept at optimum temperature with a 12 hours light/dark cycle and given rat feed in form of pellet and water ad libitum. The cages were cleaned twice daily. The period of administration lasted for 14 days.

Preparation of Serum

The rats were sacrificed 24 hrs after the last treatment by cervical dislocation. The jugular vein was cut and blood sample collected into clean, dry centrifuge tube. The blood was left for 10 min at room temperature to clot after which it was centrifuged at 4,000 rpm in an MSC (Essex, UK) bench centrifuge. The clear supernatant (serum) was aspirated using a Pasteur pipette into clean, dry sample bottles and then stored at -4°C for biochemical analyses.

Preparation of Tissue Homogenates

The heart and liver were quickly excised from the rat and immediately placed on a blotting paper to remove blood stains. The tissues were then rinsed in 1.15% KCl to remove haemoglobin followed by homogenization in 4 volumes of ice-cold 0.01 M potassium phosphate buffer (pH 7.4) using Teflon homogenizer. The homogenates were centrifuged at 12,500 g for 20 min at 4°C to obtain supernatants (post-mitochondrial fractions) which were stored till required for assay.

Determination of Biochemical Parameters

Colorimetric determination of protein concentration was done using bovine serum albumin as standard [9] while globulin was estimated using the method of Mokady, et al. [10]. Albumin concentration in the serum was measured using the bromocresol green method [11]. Serum ALT and AST activities were determined using based on the principle described by Reitman and Frankel [12]. Alkaline phosphatase (ALP) was measured by the method of Wright, et al [13]. Lactate dehydrogenase (LDH) was measured by method of Vanderlinde [14]. Bilirubin and GGT were measuring based on the instruction of the manufacturer (randox) kit.

Histological Examination

The heart and liver were fixed in 10% formalin and embedded in paraffin wax. Thin sections (7–9 mm thickness) of the tissues were cut and dewaxed in xylene, hydrated in decreasing percentage of alcohol and stained with haematoxylin and eosin.
with hematoxylin. They were then dehydrated in increasing percentage of alcohols till 70% and stained with 1% alcoholic eosin. They were differentiated in 90% alcohol and cleared in xylene. These stained sections were observed under the microscope for histopathological analysis.

**Statistical Analysis**

Results obtained were expressed as mean value ± standard deviation (SD). Comparison was done using one-way analysis of variance (ANOVA) between the control and treatment groups. P values <0.05 were considered statistically significant.

**Results**

**The effect of vitamin C and E on cypermethrin-induced changes in cardiac albumin, globulin and total protein concentrations in female wistar rats**

Cypermethrin induced significant reduction in cardiac albumin, globulin and total protein concentrations in the exposed rats when compared to control, individual administration of vitamins C and E significantly raised the concentration of these parameters (Figure 1).

**Figure 1:** The effect of administration of vitamin C and E on cypermethrin-induced changes in cardiac albumin, globulin and total protein in female wistar rats. Values are means ± SD.

**The effect of administration of vitamin C and E on cypermethrin-induced changes in hepatic albumin, globulin and total protein concentrations in female wistar rats in serum bicarbonate and sodium ion concentrations in female wistar rats**

Cypermethrin resulted in significant increase in serum cardiac albumin, globulin and total protein concentrations bicarbonate and sodium ion concentrations in the exposed rats when compared to control. Exposure of these rats individually to vitamins C and E significantly modulated the concentration of these ions (Figure 3).

**Figure 3:** The effect of administration of vitamin C and E on cypermethrin-induced changes in hepatic bilirubin, globulin and albumin in female wistar rats. Values are means ± SD.
The effect of administration of vitamin C and E on cypermethrin-induced changes in the activities of serum AST and ALT in female wistar rats

Activities of AST and ALT were significantly induced in the rats exposed to cypermethrin when compared to control, however, administration of vitamins C and E significantly modulated these activities (Figure 4).

Figure 4: The effect of administration of vitamin C and E on cypermethrin-induced changes in serum AST and ALT in female wistar rats. Values are means ± SD.

Figure 5: Photomicrographs of section of cardiac tissues of rats administered Cypermethrin, vitamin C and E. A: control rats showing normal cardiac muscle fiber architecture with multiple elongated nuclei. The cardiac muscles are branched with less space between the muscle fibers. B: rats administered cypermethrin only showing damage histo architecture of the cardiac muscle characterized by severe vacuolization of cardiac muscle with no clear branching. C: rats administered cypermethrin + Vitamin C showing a fairly normal cardiac muscle branched and multinucleated. Areas of mild vacuolization are seen and enlarge intramuscular fiber space. D: rats administered cypermethrin + Vitamin E showing mild vacuolization and enlarged intramuscular fiber space. Cardiac muscle branched and multinucleated.

Discussions

Biochemical parameters are sensitive index to determine the changes in the body system due to exposure to pesticide toxicity and can constitute important tools in toxicological studies [15]. Therefore, the purpose of this work is to evaluate the effect of cypermethrin on some selected biochemical parameters. Result showed that albumin, globulin, and total protein significantly (P<0.05) decreased in heart and liver of the rats administrated cypermethrin alone. The decrease in the concentrations of these parameters could be attributed to necrosis of the heart and liver by the toxicant which leads to the impairment in the metabolism of these parameters [16]. This result is in tandem with the previous report of Ojutiku, et al [17]. However, treatment with vitamin C and E significantly attenuated the toxic effects of the toxicant. Vitamins C and E individually have been reported for their abilities to ameliorate pathological changes in serum proteins associated with diseased conditions [18-20].

The observed significant decrease in the activities of enzymes (ALP, LDH and GGT) in the heart of rats administered cypermethrin is suggestive of cardiac tissues damage. Enzyme activities in tissues are often used as marker to ascertain early toxic effects of administered foreign compounds to experimental animals [21,22]. Decrease of these enzymes in the organs such as heart might be due to possible membrane damage or due to stress condition resulting into leakage of the enzymes from the heart into the serum. ALP is a marker enzyme for plasma membrane and is required for proper functioning of organs. LDH is an intracellular enzyme present in nearly all metabolizing cells with highest concentration in the heart, skeletal muscle and erythrocytes.

Activities of ALT and AST are routinely measured clinically as diagnostic tools in assessment of hepatocellular injury and health status of the liver [22,23]. Result of this study revealed that administration of Cypermethrin in rats caused a significant increase (P<0.005) in serum concentrations of ALT and AST. This is an indication that Cypermethrin might have caused alteration in biochemical activities in liver and also compromised the integrity of the membrane causing the release of these enzymes into the blood stream. However, treatment with vitamin C and E significantly reverse these alterations.

Photomicrograph of the heart and liver of rats administered cypermethrin show significant histological
distortions in their normal appearance and architecture. The cardiac tissues were characterized by severe vacuolization, deformed nuclei, disarray of myofibres and connective tissue deposits which might probably be due to degeneration of the structural protein in mitochondria of the cytoplasm. The livers of cypermethrin treated rats also indicate acute tubular necrosis, damaged parenchyma and vascular congested cells with degenerated tubules. The hepatocytes were scattered in the hepatic parenchyma instead of radiating in a cord arrangement from the central vein and are separated by a widening sinusoid. These alterations were found to be reduced following the administration of vitamin C and E.

References


