Provided by International Institute for Science, Technology and Education (IISTE): E-Journa

Journal of Natural Sciences Research ISSN 2224-3186 (Paper) ISSN 2225-0921 (Online) Vol.5, No.1, 2015



Effect of Alcoholic Leaf Extract of Hellotropium Earopiam on the Levels of Testosterone , Prolactin ,TSH , T_3 and T_4 Hormones in Blood of Male Albino Rats Exposed to Oxidative Stress by H_2O_2

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Abstract

The current study aimed to compare the effects of alcoholic leaf extract of Heliotropium europiam and Vit. C as antioxidants on levels of Testosterone (testo), Prolactin(PRL), TSH, T₃,T₄ hormones in blood of adult albino male rats exposed to oxidativ stress by hydrogen peroxide (0.5%) in the drinking water throughout (30) days. For the purpose of the study preliminary experiments were carred out yo find the most effective dose of plant extracts and it was found that (300mg / kg).. Rats (3-4 months)age and (250±285g) weights were distributed randomly into five groups each group included five rats as follows: control group, H₂O₂ group, H₂O₂+ Heliotropium europiam alcoholic leaf extract group, H₂O₂+ Vit. C group, H₂O₂+ Heliotropium europiam alcoholic leaf extract+Vit.C group. The plant extract of leaves as (300) mg / kg and Vit.C (250) mgkg of body weight in a single daily dose. The results of the statistical analysis shwed that the animals exposed to stress oxidative mediated by hydrogen peroxide led to a significant decrease (p \leq 0.05) in the levels of testo. and RRLand T₃ and non significant difference in the TSH and T₄ levels when compared with control. The treatment with H₂O₂+ plant extract shwed asignificant increase in the testo. level and a significant decrease in T₃ and T₄ levels, and lack of significant difference in the levels of PRL and TSH, while the treatmen of group H₂O₂+Vit.C led to asignificant increase in testo., PRLand TSH and a significant decrease in T₃ and T₄. While the group exposed to stress and treated with plant extract and Vit.C showed asignificant increase in testo. And PRL levels and significant decrease in TSH and T₄ with non significant difference in the T₃ level when compared with hydrogen peroxide group.

Keywords: Hellotropium earopiam, H₂O₂, Free racicals, oxidative stress Testosterone, Prolactin, TSH.

Introduction

Free radicals (FR)are atoms or molecules which contain a single electron or more than one electron in their external orbit, they were produced in living cells especially in mitochondria and they are either reactive oxygen species or eactive nitrogen species(Cooper *et al*; 2002), while oxidative stress it represents a situation that increases the production of free radicals and oxidants in cells so that exceeds the ability of antioxidants to remove or minimize their effects, leading to the formation of oxidative damage to cells and tissues and thus lead to initiation of various diseases in humans, such as cancer, diabetes, cardiovascular diseases, arthritis and disorder in hormones level (Seung *et al.*, 2009). And for reducing to free radicals activity and damages caused by them there are complex systems of multiple models for antioxidants maintained by man and animals, some of them are exogenous and some of them are endogenous and they are a reducing substances their activity represent by inhibition or ending reactions that electrons moved from a certain material to oxidant factor and that leading to production of intermediate free radicals(Venluri& Venluri, 2007)

Human have used plants surrounding it during the thousands of years that he lived on earth in search of food and heal their pains that's where the use of herbs and medicinal plants are an important tool in the treatment of many diseases as most currently known drugs are from plant sources(Ghers *et al.*,2005) Recently, The interest has increased by using some plant extracts in the treatment of many diseases such as cancer, diabetes, arthritis and the hormonal and nervous system disorders because they give better results than manufactured chemicals and lacking of their harmful side effects(Sangameswaran *et al.*,2009).

The Plants of Borginaceae family such as *Heliotropium europium* which have a great biological importance because they contain a many of active compounds, include alkaloids as pyrrolizidine and Acetyl lasiocarpine in the entire plant (Sharma *et al.*,2009) and Alkaloids in leafs as lasiocarpine, heliotrine (Youssef *et al.*,1978; AL-Blesh,2012).and indicine-N-Oxide wich acts as anticancer and antioxidants(Kumardas,2011) and contain types of Glycosides as Qurencetin, Isorhamntine, heliotrope and Isorhamntin (Dnidl,2012) in addition to Flavonoids as Narigrnin (Modak *et al.*, 2009).

The most important materials induced oxidative stress and cause damage in living cells is Hydrogen Peroxide which attack many living systems as polyunsaturated fatty acids of cellular membranes due to lipid peroxidation which cause initiation of many diseases as Diabetes , Cancer and Cardiovascular diseases (Atawodi;2005) in addition to its role in breaking peptide bonds , denaturation of proteins and damage of DNA leading to intiation of genetic mutation (Valacchi&Davis,2008) . Therefore, the current study aimed to determine the effect of alcoholic extract of *Heliotropium europium* and Vitamin C as wel known antioxidant on the levels



of some hormones: Testosterone, Prolactin, Triiodothyronine (T_3) , Tetraiodothyronine (T_4) and Thyroid Stimulating Hormone (TSH) in the blood of male adult albino rats exposed to oxidative stress.

Materials and Methods

This study used male albino rats weights $(250\pm285)g$ and age ranges between (12-16)weeks obtained from Faculty Medicine / University of Mosul and housed in animal house of Biology Department in Tikrit University where placed in metal cages at (3 ± 22) with normal light cycle (12 h light and 12 h dark), Animals were fed with standard laboratory diet(Ashokkumar *et al.*, 2005) and water was given *ad libitum* during the experiment period from November 2011 to February 2012.

Dtermination the effect dose of plant extract

For the purpose of the study preliminary experiments were carred out yo find the most effective dose of plant extracts and it was found that (300mg / kg).

Experimental Design

Rats were divided randomly into six groups, each group contain five animals as follows:

- 1-Control group (untreated group): were given food and water continuously throughout the duration of the experiment.
- 2-Hydrogen Peroxide group: Given (0.5%) of hydrogen peroxide in drinking water for 30 days.
- 3- Hydrogen Peroxide and plant extract group: given hydrogen peroxide as in the second group with dosage with alcoholic extract of *Heliotropium europium* leafs (300mg/kg)for 30 days.
- 4- Hydrogen Peroxide and vitamin C :this group was given hydrogen peroxide as in the second treatment with the dosage of vitamin C (250mg/kg)for 30 days .
- 5-Hydrogen Peroxide and plant extract with Vitamin C: this group were given hydrogen peroxide as in the second treatment with dosage by alcoholic extract of *Heliotropium europium* leafs (300mg/kg) and Vitamin C (250mg/kg) for 30 days.

Blood sampling

At the end of 30 days of treatment and administration period, the animals where fasted overnight after which they were painlessly sacrificed by decapitation under chloroform anaesthesia and the blood (6-8mls) collected by cardiac puncture ,blood put in tubes without anticoagulant and left in the laboratory temperature for (15) minutes then it was separated by centrifuge and save serum at degree (-20) C^0 until do the tests.

Hormonal assay

- 1- Determination of Testosterone concentration.: the concention of testosterone was estimated depending on the steps came with kits of American Monobind company and as instruction of the manufacturer's own technology ELISA (Tietz, 1995).
- 2- Determination of Prolactin (PRL) concentration: the concention of Prolactin was estimated depending on the steps came with kits of American Monobind company and as instruction of the manufacturer's own technology ELISA (Tietz,1995).
- 3- Determination of TSH concentration: the concention of TSH was estimated depending on the steps came with kits of American Monobind company and as instruction of the manufacturer's own technology ELISA (Fesher1996).
- 4- Determination of T_3 concetraion: the concention of T_3 was estimated depending on the steps came with kits of American Monobind company and as instruction of the manufacturer's own technology ELISA(Braverman, 1996).
- 5-Determination of T_4 concentration: the concention of T_4 was estimated depending on the steps—came with kits of American Monobind company and as 1998).

Statistical analysis

The significant differences between the studying groups were found by using ready statistical software (SAS,2001), and statistical means was tested using polynomial Duncan test at significance level ($p \le 0.05$) to determine the significant differences between groups (Duncan,1955).

Results and Discussion

Figure (1) illustrated that there was a significant decreasing ($p \le 0.05$) in Testosterone concentration in Hydrogen Peroxide group in comparison with control group, this result corresponding with results of AL-Hiali (2010) study who noted to a significant decreasing in hormone concentration in treatment of male albino rats with hydrogen peroxide with drinking water for 30 days. The reasons may be due to oxidative stress which lead to



inhibition of hypothalamus-pituitary axis (Knol , 1991) or due to reactive oxygen species effects in leydig cells which it works on the inhibition of steriodogenic acute regulatory protein replication found in mitochondrial membrane in leydig cells which represent the main center of testosterone synthesis and this protein simplify the entry of cholesterol into the cell , it also work on inhibition cytochrome P_{450} side chain cleavage enzyme which has a major role in hormone biosynthesis in leydig cells (Allen *et al* .,2000).

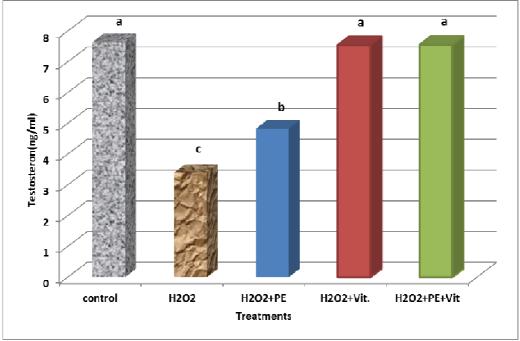


Figure (1): effect of different treatment on the concentration of testosterone in male albino rats exposed to oxidative stress

The treatment of hydrogen peroxide group with alcoholic extract or vitamin C or alcoholic extract with vitamin C lead to a significant increasing of testosterone concentration in comparison with hydrogen peroxide group.

The reason of this increasing due to the containing of alcoholic extract of *Heliotropium europium* leafs on many of antioxidant molecules as Glycosides and Flavonoids which act to reduce or prevent the damage of free radicals and thus reduced oxidative stress resulted from hydrogen peroxide treatment and these molecules listed on work on stimulating selenoprotein enzyme which contain selenocysteine and the last one has a major role in the transport of Selenium and protection tissues from Reactive Oxygen Species (ROS) effects and his work is similar to Glutathione Peroxidase and has a relationship with Testosterone secretion in leydig cells (Nishimura *et al.*, 2001). As for the role of Vitamin C as antioxidant is known to protect cells and tissues from oxidative damage and also has an important role in stimulating the pituitary gland to secrete Interstitial cell stimulating hormone (ICSH), Which stimulates the leydig cells to production of Testosterone, We note from figure (1) that the treatment of rats exposed to hydrogen peroxide with alcoholic extract of plant with vitamin C was better than the treatment of alcoholic extract of plant alone and this result may be due to synergism and cooperation of Vitamin C with compounds found in plant and thus leading to best results.

It is noted from Figure (2) a significant decreasing($p \le 0.05$) in the concentration of prolactin in the group treated with hydrogen peroxide as compared with control group and the reason may be due to the effect of oxidative stress on the axis of the hypothalamus - pituitary which leads to inhibition of this axis and thus reduce the hormones secreted from pituitary gland, including the prolactin and the treatment of animals exposed to oxidative stress with vitamin C and plant extract with vitamin C together have led to a significant increasing in prolactin concentration as compared to hydrogen peroxide group. The reason may be due to the presence of Qurencetein in plant which consider as a powerful antioxidant and reduce from free radicals generation thus leading to prevent or reduce their effect on hypothalamus-pitutary axis and thus return pituitary gland to secrete their hormone , including prolactin(Dnidl ,2012),As for Vitamin C as antioxidant , it reduce from free radicals effects on this axis and stimulate the secretion of pituitary gland .



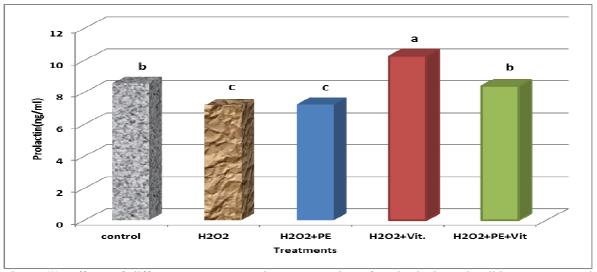


Figure (2): effects of different treatment on the concentration of prolactin in male albino rats exposed to oxidative stress.

It was Shown from (Figure 3)that there is no significant difference in the concentration of the Thyroid Stimulating Hormone (TSH) in hydrogen peroxide group as compared with the control group, These findings were compatible with the results of AL-Douri, (2012) in male rabbits exposed to oxidative stress that did not find any significant difference in TSH concentration between the group exposed to oxidative stress and the control group when treated rabbits for 30 days and studies also shown that there was nonsignificant differences in TSH concentration and oxidative stress resulted from diabetes (Celani *et al* .,2004; Kedari , 2001). It is clear from Figure (3) presence of a significant increasing in hydrogen peroxide with vitamin C group as compared to hydrogen peroxide group only and this is attributable to stimulate the vitamin C content of the pituitary gland to secrete TSH as a result of the low concentration of T3 (Figure 4)by negative feedback mechanism, or may be attributed to the depletion of iodine thereby increasing the responsiveness to TSH(AL-Douri, 2012). or to the inhibition of Thyroperoxidase .As for group exposed to oxidative stress and group treated with alcoholic extract to the leaves of the plant did not lead to a significant difference in the concentration of TSH compared with the group with findings (AL-Rubaei and AL-Musawi, 2011) Because I have noticed no change in the concentration of TSH in the case of oxidative stress, which in turn is reflected on the levels of thyroid hormones circulating in the blood.

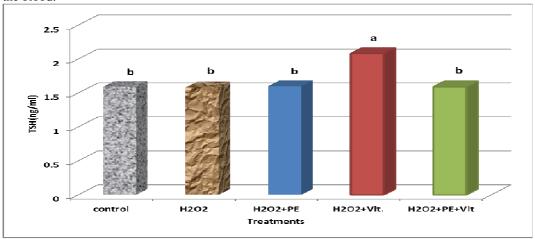


Figure (3):effect of different treatment on the concentration of Prolactin in male albino rats exposed to oxidative stress

It was showed from figures (4and 5) that group exposed of oxidative stress has lead to a significant increasing in T3 and T4 concentrations and this reason may related to inhibition of 5-monodeiodinase which controls of the transfer of T4 to T3 (Kedari,2011).

According to Hydrogen peroxide with Vitamin C group there was a significant decreasing in concentration of T3 and T4 as compared with control group and this reflected on the significant increasing on the concentration of TSH in this group as compared with control group and this reason in decreasing of the levels of two hormones may be due to roles of Vitamin C as antioxidant which works on the inhibition of antithyroid



compounds and these compounds metabolize in the body to Thyocyanate and Isothyocynate and finally to sulfur containing compounds. Rouzaud *et al.*,(2004) noted that Cyanogenic glycosides convert in the body to Thyocyanate by Clucosidase or Salfer transferase found in tissues of plants or animals.

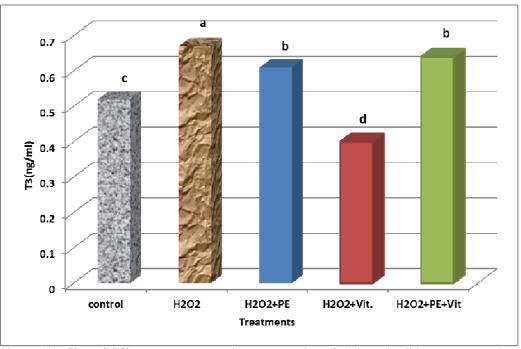


Figure (4): effect of different treatment on the concentration of T3 in male albino rats exposed to oxidative stress

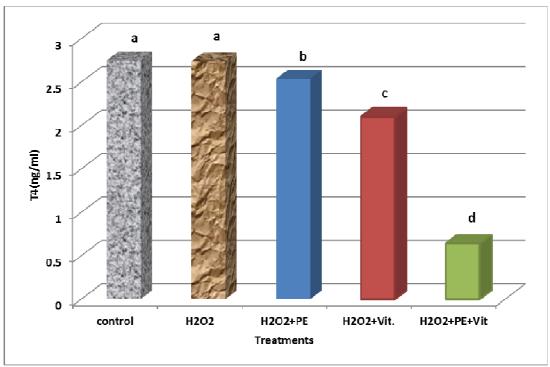


Figure (5):effect of different treatment on the concentration of T4 in male albino rats exposed to oxidative stress. The high concentrations of Thyocyanate inhibit the combination of iodine with insoluble thyroglobuline (Ermans and Bourdoax, 1989), While the treatment of hydrogen peroxide exposed rats of alcoholic extract and vitamin C didn't lead to any significant differences in the concentration of T4 but led to a significant increasing in T3 concentration as compared with hydrogen peroxide group and the reason may be due to reducing of the linkage of T3 with proteins carrying it and finally leading to decreasing the concentration of T3 in blood.



Conlution

There are a lot of data wich show that free radicals and reactive oxygen species increased damage done the compounds of living cells the effect of alcoholic leaf extract of *Hellotropium earopiam* acts as antioxidant supplement wich is decrease harmful effect of the free radicals through some improvement was observed in the levels of studed hormones.

Acknowlegdments

The author is grateful to thre college of science-Tikrit Unniv.and woulk like to thank MSc. Safa Amed for her supports, Dr.Prof. Zaid M. Mubark for kind advice, Assist.Prof.Arkan B.Mqhamad for his assistance in the prepration of this parper.

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