Effect of irradiated almond and hazelnut enricheddiet on biochemical aspects in rats

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Abstract

The present studv aims to evaluate effect irradiationatdoses 6 kGy and 10 kGy on chemical and biochemical parameters of nuts including almond (Prunusamygdalus) and hazelnut (Corylusavellana).70 albinoratsused in this study, the rats divided into 7 groups as following: **Group** (1): fed on basal diet, as a control group. Group (2): fed on basal diet containing un-irradiated almond. Groups (3 and 4): fed on diets containing the irradiated almond on 6 and 10kGy, respectively. Group (5): fed on basal diet containing unirradiated hazelnut. Groups (6 and 7): fed on diets containing the irradiated hazelnut on 6 and 10 kGy, respectively for 8 weeks. The statistical analysis of irradiated almond and hazelnutat dose 6 and 10 kGy at zero time and storage for 6 months on its approximate analysis showed no significant difference in protein, moisture, ash and lipid. The biochemical performance showed non-significant effect on organs weight, high density lipoprotein-cholesterol (HDL-c), low

density lipoprotein-cholesterol (LDL-c), blood hemoglobin, total cholesterol, triacylglycerol's, glucose, serum alanine amino transferase (ALT), serum aspertate amino transferase (AST), liver superoxide dismutase (SOD), liver glutathione peroxidase (GPX) and liver catalase in groups applied doses of non-irradiated or irradiated almond and hazelnut, when compared with control group which fed on basal diet, whileserum alanine amino transferase (ALT) which showed increased in untreated almond and decreased at untreated hazelnut and there were significant increase in antioxidant enzymes at all groups.

Introduction

Almond (*Prunusamygdalus*Batsch.) belongs to the family *Rosaceae*, which also includes apples, pears, peaches, and raspberries (*Jahanban et al., 2010*). Almonds (*Prunusamigdalis*) are a good dietary source of vitamin E, sterols, and flavonoids, each of which has been suggested to play a role in the promotion of health. In particular, increased consumption of flavonoids has been associated with an anti-obesity effect in women and reduced risk of stroke, cardiovascular disease, and some forms of cancer (*Yochum et al., 1999*). The composition of almond is 25.23% moisture, 5.00% ash; 32.73% lipid; 33.66% crude fibre; 3.11% crude protein and 25.47% carbohydrate (*Akpabio, 2012*).

Omer et al., (2011) reported that, the amounts of oleic acid, linoleic acid, palmitic acid and stearic acid in Picantili, Ferraduel, were (57.46 to 68.65%, 11.77 to 25.15%, 5.06 to 7.26% and 1.26 to 2.41%), respectively. Alisonand Oliver (2012) given their favorable fatty acid composition and high fiber content, the U.S. Food and Drug

Administration (FDA) released a health claim recognizing that almonds can help maintain a healthy cholesterol level, particularly in patients with hypercholesterolemia. Nut consumption is not associated with a higher body math index (BMI) among free-living individuals. Almonds are low in available CHO, have a healthy fatty acid profile, and are high in vegetable protein, fiber, and magnesium.

Almonds with skin, hull, and shell show a powerful capacity in collecting free radicals and this activity can be related to the presence of unsaturated fats, terpenoids, flavonoids, phenolic acids, and other beneficial compounds including fiber (*Jahanban et al.*, 2010). Gamma irradiation could increase phenolic content in almond skin(*Harrison and Were*, 2007).

Hazelnuts (*Corylusavellana*) play a major role in human nutrition and health, because of their special composition of fat, protein, carbohydrate, vitamins, minerals and nutrients antioxidant (*Alasalvar et al., 2009*). Hazelnuts contain fat 60.8%, protein 15.0% and fiber10.4%(*Salas-Salvadó., et al., 2006*). On the other hand, (*Alasalvar et al., 2006*) reported that, hazelnut plays a major role in human nutrition and health because of its special composition of fat (around 60%), most of which is highly rich in Monounsaturated Fatty Acids (MUFA) mainly oleic acid.

Hazelnut is a very rich source of energy providing approximately 6 to 6.5 kcals/g fresh seed. Fatty acids composition in Hazelnut play an important role in human health, in this respect (*Connor, 2000, Aronson et al., 2001 and Iso et al., 2002*) reported that, the consumption of specific fatty acids such as omega- 3 fatty acids and oleic acid may provide health benefits.

Dietary compounds other than antioxidant vitamins may provide a critical role in protecting against Radical Oxygen Species ROS induced free radical injury. Phenolic compounds are found in virtually all plant foods and many phenols can act as powerful antioxidants that may reduce free radical damage. Individual phenolic compounds with known antioxidant activities have been identified in hazelnut including gallic acid, 4-OH benzoic acid, p-cou-maricsinaptic acid, and guercetin Hazelnut extracted with 80% Ethanol OH has also demonstrated antioxidant activities. hydrogen peroxide. superoxide, DPPH, and β-carotene linoleate system. The hazelnut extract also inhibited human LDL oxidation and DNA scission. The oxygen radical absorbance capacity (ORAC) assav physiologically relevant antioxidant test because it measures the ability of an antioxidant system to inhibit the oxidative damage to susceptible molecules of peroxyl radicals Haiwen and John (2011) and Shahidi et al., (2007).

Food irradiation has been recognized and regulated as an effective food processing technology in many countries being able to destroy or reduce ubiquitous pests and pathogens that contaminate raw foods (*Diehl*, 1981).Radiation processing is well established as a physical, non-thermal method to preserve various food products that involves the exposure of food products (raw or processed) to ionizing or non-ionizing radiation (*Antonio et al.*, 2012).Irradiation can also influence the levels of antioxidants/phytochemicals in plant products (*Behgar et al.*, 2011).

Gamma radiation, more energetic than X-rays, is used from sources of radioactive isotopes, cesium-137 or cobalt-60, and it is identified by the World Health Organization as a food preservation

technique that improves food safety without altering the toxicological, biological or nutritional quality of the food *(Farkas andMohácsi-Farkas2011)*. Therefore, the present study was carried out to assess the effects of diet containing untreated and treated almond and hazelnut on healthy rats.

Material and methods

Material:

Almond (*Prunusamigdalis*), Hazelnut (*Corylusavellana*), sucrose, starch and corn oil were obtained from the local market, Cairo, Egypt.

Casein, all vitamins, minerals, cellulose, L -Cystine and choline chloride were obtained from El–Gomhoriya company, Cairo, Egypt.

Kits: kits used to determine hemoglobin, glucose, AST, ALT, cholesterol, triglycerides; HDL-c andantioxidant enzymes including (malonaldehyed, GPX, Catalase and SOD) were obtained from Gama tread Company, Cairo, Egypt.

Methods:

y Irradiation treatment:

Almond (*Prunusamigdalis*) and Hazelnut (*Corylusavellana*) were packed in polyethylene bags, and sealed by heat. Each bag contained about 200 g. they were subjectedγ Irradiation from Co⁶⁰ at National Center for Radiation Research and Technology at Nasr City, Cairo Egypt. The facility used was Gamma Chamber 400 A, Co-60 facility of India. The doses applied were 6 and 10 kGy delivered at dose rate of 1.606 kGy /h as calibrated using small pieces of radiochromic film *(Maclaughlin et al., 1985)*at the time of experimentation. The samples were stored at 5°C until used.

Chemical composition of almond and hazelnut:

Moisture content, crude fat, crude protein, ash content were determined according to the method described by the (A.O.A.C. 2003). The fatty acids compositionwere determined as the method described by (Hamilton and Hamilton, 1993).

Diet preparation:

The diets were prepared by using untreated and treated (almond and hazelnut) with irradiation. The diets were prepared according to (*Reeves et al., 1993*), the salt mixture was prepared according to (*Hegested al., 1941*) and the vitamin mixture was prepared according to (*A.O.A.C. 2003*).

Experimental rats:

Seventy male albino rats, Sprague-Dawley strain, with an initial weight of about 80 ±5 g were used in this study. They were obtained from animal house of National Center for Radiation Research and Technology, Atomic Energy Authority, Cairo, Egypt. The animals were housed individually in plastic cages with wire mesh bottoms at a room temperature of 22-25°C and 60±5% relative humidity, with a photoperiod of 12h and water for eight week. Groups of ten rats were then assigned to receive one of seven experimental diets (i.e. control, untreated nuts and treated nuts with 6 and 10 kGy irradiation) almond and hazelnut, alongside casein diet. All animals will be free access to feed and water on based diet for one week for adaptation. After this week, the rats divided into seven groups as the following: group 1 fed on basal diet as a control group, group 2, 3 and 4 were feed on untreated almond and treated almond with 6 and 10 kGy, respectively. Group 5, 6 and 7 were feed on untreated hazelnut and treated hazelnut with 6 and 10 kGy, respectively.

During the experimental period (8 week), the diets consumed and body weights were recorded every week. At the end of the experiment period, the animals were fasted overnight, then the rats were anaesthetized and sacrificed, and blood samples were collected from the aorta. The blood samples were centrifuged and serum was separated to estimate some biochemical parameters, i.e. Serum cholesterol (*Richmond, 1973*), triglycerides (*Stein, 1987*),high density lipoprotein-cholesterol HDL-c (*Firdewald et al., 1972*), low and very low density lipoprotein-cholesterol LDL-c and VLDL-c (*Firdewald et al, 1972*), glucose(*Young, 2001*), aspartate amino transferase (AST) and alanine amino transferase (ALT) (*Young, 1990*). Hemoglobinwere estimated according to (*Dacia, and Lewis. 1985*). Liver and kidney were separated from each rat and weighted to calculate organs to body weight %.

Determination of antioxidant enzymes:

Superoxide dismutases (SODs) are metabolioenzymes that catalyze the dismutation of the superoxide anion to molecular oxygen and hydrogen peroxide and thus forma crucial part of the cellular antioxidant defense mechanism (*Nishikimiet al.*, 1972).

Cellular glutathione peroxidase (GPs) is a member of a family of GPx enzyme whose function is to detoxify peroxides in the cell. The GPx enzymes play a critical role in protecting the cell from free radical damage, particularly lipid peroxidation (*Pagliaand Valentine*1967).

Catalase is antioxidant enzyme that is present in the most aerobic cells. Its serves as one of the body's defense systems against H_2O_2 a strong oxidant that can cause intracellular damage (*Aebi*, 1984).

The data obtained was analyzed statistically for standard deviation and one way ANOVA test according to *(Duncan, 1955)*.

Result and discussion

Effect of radiation processing with storage on chemical composition of nuts (Almond and Hazelnut).

The effect of radiation processing at (6 and 10 kGy) on the mean values of moisture, ash, crude protein, and crude fat of almond and hazelnut in zero time and after storage for 6 months presented in tables (1 and 2).

1-1 Almond:

The variation of moisture, ash, crude protein, and crude fat of almonds treated with irradiation at zero time and after storage for 6 months were studied, the results are presented in table (1). The mean values ofmoisture, protein, ash and crude fat in zero time does not change significantly in irradiated almonds at 6 kGy, except ash which showed significant decrease, ascompared to un-irradiated almonds (control). On the other hand, irradiated almonds with 10 kGy caused significant decrease in moisture and ash, while the other nutrients showednon-significantchanges, as compared to un-irradiated almonds.

Results in this table indicated that, non-significant changed in moisture, crude protein, ash and crude fat in almonds treated with irradiation (6 and 10 kGy) after storage for 6 months, as compared to the control sample. Several authors agree with our previous mentioned results among them (*Bhatti et al., 2013*) reported that gamma irradiation exerted no considerable effect on the proximate

almond seed composition. In addition to, *Mahfouz, (2015)* reported that the moisture, protein, lipid, and ash of almond nut were not substantially affected by gamma irradiation. Similar finding showed that gamma irradiation with doses up to 10 kGy had no real effect on moisture, protein, lipid, fiber and ash of almond. During storage at room temperature, the total protein and total fat of almond nut also significantly (p<0.05) increased in both irradiated and non-irradiated samples. (*Bela et al., 2008 and Maityet al. 2009*) reported that gamma irradiation could change the seed protein quantitatively, not the qualitatively.

HazeInut:

The variation of moisture, ash, crude protein, and crude fat of hazelnut treated with irradiation at zero time and after storage for 6 months presented in tables (2). The data showed that there was a different in the mean values of moisture between the control sample and treated sample with (6 and 10 kGy). Total lipid also decreased significantlyin hazelnutwhich treated with irradiation at doses (6 and 10 kGy), as compared to un-treated sample.

At the storage (6 moths), the results in the same table indicated that, non-significant increase was observed in the moisture content of hazelnut which treated with irradiation at (6 kGy), while increased significantly with irradiation at (10 kGy), as compared to the control sample. This increase may be due to the humidity of atmosphere. Ash, protein and lipidcontent of hazelnut did not changed significantly with irradiation at (6 and 10 kGy) after storage period for 6 months, as compared to the control sample.

In this respect, (Saadetkoc and Ahmet2017) reported that, After irradiation, 0.5, 1 and 1.5 kGy doses of gamma irradiation

significantly increased the total fat values (p < 0.05) and this value decreased during the storage. (Aycaet al., 2007) indicated that the low dose irradiation treatment slightly increased the total lipid content whereas it dramatically decreased for high dose treatment.

Fatty acids composition of un-treated and treated almond and hazelnut with two doses of irradiation at zero time and after storage for 6 months:

The fatty acids composition of two types of nuts sample exposed to different doses of irradiation are presented in table (3 and 4).

Almond:

The data in table (3) indicated that, at zero time the untreated almond (control) contains 7.42% saturated fatty acids (SFA) {palmitic acid} that was the major saturated fatty acid.

Treating almond sample with irradiation-increasedpalmitic acid by 2.4% and 28.0% at 6 and 10 kGy, respectively. Monounsaturated fatty acid (MUFA) {Oliec acid} of control sample was 73.74% that was the major (MUFA). After treated almonds with irradiation (6 and 10 kGy) oleic acid decreased by about (- 9.68% and - 2.7%), respectively.

Polyunsaturated fatty acids (PUFA) {linoleic acid} of raw sample was 18.8 % that wasthe major (PUFA). The data indicated that the linoleic acid increased by (+ 36.64) at 6kGy and decreased by (-21.8%) at 10 kGy.

After storage almonds for 6 months, palmitic acid in almond which treated with 6 kGyincreased by about 13.6%, oliec acid decreased by 10.49%, and archidic was decreased by 27.1%, while

linoliec acid did not appeared. On the other hand, palmitic acid in irradiated almonds with 10 kGy increased by 33.96%, oliec acid decreased by 24.23%, linoliec acid did not appeared and archidic increased by 34.45%.

Hazelnut:

The data in table (4) indicated that, at zero time the raw hazelnut (control) contain 6.58% saturated fatty acids (SFA) {palmitic acid} that was the major saturated fatty acid. Treating hazelnut sample with irradiation decreasedpalmitic acid by 8.8% at 6kGy and increased by 6% at 10 kGy.Monounsaturated fatty acid (MUFA) (oliec acid) of control sample was 81.28% that was the major (MUFA). Oleic acid decreased by (- 8.5%) at 6kGyand by (- 2%) at 10 kGy.Polyunsaturated fatty acids (PUFA) {linoleic acid} of control sample was 8.19% that the major (PUFA). The data indicated that the linoleic acid change by3.6% at 6 kGy and by 5.6% at 10 kGy.

After storage for6 months the fatty acid arachidic was appeared at 6 kGy (2.4%). Palmitic acid increased by 72% at 6 kGy, Oliec acid decreased by 2.1% and linoliec acid did not appeared, but at 10 kGy (Palmiticacid decreased by 20.3%, Oliec acid decreased by 31.8%, linoliec acid decreased by 62.0%. The effects of gamma irradiation doses and storage duration interaction on fatty acid composition of natural hazelnut kernels were not found to be significant (p>0.05) (Koc et al., 2017).

Mahfouz, (2014) stated that the fatty acid profile slightly changed due to irradiation. The major change in fatty acid composition was the decrease in the quantity of fatty acids (C16:0, C18:0 and C18:1) and the increase in the quantity of fatty acid (C18:2). There were fluctuating changes (either increasing or decreasing) in fatty acids throughout the storage duration.

Biological analysis for rats feed on diet contain untreated and treated nuts:

The relative organs weight of rats feed on diet containing untreated and treated nuts with irradiation:

The data in table (5& 6) showed the mean value of relative organs weight of rats fed on diets containing un-treated and treated almond and hazelnut with irradiation at dose of 6 and 10 kGy. The mean values of heart, liver, spleen, testes and kidney showed non-significant changes between the groups fed on diets containing untreated almonds or hazelnut, as compared to control group fed on basal diet. On the other hand, feeding rats on diet containing almonds or hazelnut, which treated with irradiation at dose (6 and 10 kGy), caused non-significant differences in all organs weights, as compared to the group fed on diet containing un-treated almond except heart and spleen in almonds & spleen in hazelnut groups.

Effect of diet containing untreated and treated almond and hazelnutwith irradiation, onsome biochemical analysis of healthy rats.

Almond:

The table (7) showed that the mean values of hemoglobin, glucose and Alanin Amino Transferase ALT did not affect with diet containing untreated almonds, while Aspartate Amino Transferase AST increased significant, as compared to control group (fed on control diet). Treating almonds with irradiation at dose (6 kGy) showed non-significant changes in all parameters, except AST enzyme, as compared to the group fed on diet containing un-treated almonds. On the other hand, the high dose of irradiation of almonds

led to significant increase in hemoglobin and glucose, as compared to the group fed on diet containing un-treated almonds

Hazelnut:

Un-treated hazelnut led to non-significant changes in glucose and AST enzyme, while caused significant decrease in hemoglobin and ALT enzyme were observed, as compared to the control group fed on basal diet. The data presented in Table (8) revealed that, treated hazelnut with (6 and 10 kGy) increased the mean value of hemoglobin significantly, while the other parameters showed non-significant changes, as compared to the control group fed on untreated hazelnut.

In this respect (*Jiang et al., 2002*) reported that, the Nurses' Health Study showed in 83,818 healthy women that eating 140 g of nuts per week was related to a significant lower DMT2-risk compared to non-consumers. This result was inter alia attributed to the low glycemic index of nuts and their high fiber and magnesium content. In addition, recent studies with 135,956 women confirmed an association between increased walnut consumption (> 56 g/week) and a lower incidence (15 %) for DMT2 (*Pan and Manson 2013*). Contrary to that, *Kochar et al.* (2010) observed no effect of nut consumption on the DMT2-risk in 20,224 male subjects of the Physician's Health Study.

The responsible mechanisms mediated by nut consumption which cause a reduction of the DMT2-riskare not yet fully understood. A modulation of the adiponectin concentration appears conceivable (*Aronis et al., 2012*). The protein, formed by fat-laden adipocytes, is involved in the regulation of appetite and inverse associated with the DMT2-risk (*Heidemann et al., 2008*).

It is also possible that an increase in insulin sensitivity results from the arginine and zinc content of the nuts, which stimulate both insulin secretion and the receptor tyrosine kinase and thereby increase the insulin sensitivity of the cells. In addition, a reduced postprandial glycemic response mediated by nut consumption and a significantly higher release of satiety hormones may also contribute to the prevention of DMT2 (*Reis et al.*, 2012).

Nuts are highly nutritious foods rich in unsaturated fatty acids, fiber, vitamins, minerals and some bioactive substances, such as phenolic antioxidants and phytosterols (*Bao et al., 2013*) and due to these wholesome benefits, individuals living with liver disease are usually advised to include nuts in their diet (*Han et al., 2014*).

Wien et al., (2003) found that a hypocaloric, almond-enriched diet led to greater reduction in weight andbody mass index BMI. This study results were in accordance with the findings of most previous investigations in this field, supporting that weight loss through lifestyle modification, medication or bariatric surgery is associated with a decrease in liver enzymes (Wang et al., 2003, Straznicky et al., 2012 and Clark 2006).

The improvement in liver enzymes in treated diabetic rats with some nuts may be related to the antioxidant properties of these nuts, which have scavenge free radicals and thereby may protect cells from oxidative stress. Almonds (Amygdaluscommunis L.) are a rich source of nutrients and phytochemicals such as vitamin E also polyphenols (Alasalvar et al., 2006 and Yang et al., 2009) Ithat is known as antioxidants and had strong free radical scavenging (Choi et al., 2002).

Effect of diet containing untreated and treated almond and hazelnut with irradiationon lipid profile of healthy rats: Almond:

Table (9) showed the levels of serum cholesterol, triacylglycerol's, high density lipoprotein-cholesterol HDL-c and low density lipoprotein-cholesterol LDL-c in serum rats which were fed on diets containing untreated almonds and treated almonds with irradiation at levels (6 and 10 kGy).

The data in this Table revealed that, feeding rats on diet containing untreated almond showed non-significant changes in the mean value of serum cholesterol and HDL-c, while serum LDL-c and triglyceride changed significantly, as compared to healthy rats fed on basal diet. The mean value of serum triacylglycerol's increased significantly in serumrats treated with untreated almonds, as compared to the control group.

Treating healthy rats with diets containing irradiated almond (6 and 10 kGy) led to non-significant differences in the mean values of serum cholesterol, triacylglycerol's, HDL-c and LDL-c, as compared to the group of rats fed on diet containing untreated almond. Increase the intake of monounsaturated fat in the diet has associated with reduced of LDL been susceptibility oxidation (David et al., 2008), almonds consists of a number of components that may reduce LDL oxidation. On the other hand, (Jennifer et al. 2002) indicated that HDL cholesterol was significantly lower with the almond diets and suggested that nuts can reduce heart disease risk in healthy persons.

The National Cholesterol Education Program demonstrated that, almond consumption fits well with current American Heart

Association guidelines to replace saturated fats with unsaturated fats and with the National Cholesterol Education Program (NCEP) guidelines to liberalize total fat intake, specifically from monounsaturated fat (MUFA), (Garg et al., 1988) related to its ability to increase HDL cholesterol. Hyson et al., (2002) revealed that the consumption of almond kernels with or without brawn skin is effective in reducing the level of total cholesterol and LDL.

HazeInut:

The result of lipid profile including (serum cholesterol, triglyceride, HDL-c and LDL-c) in healthy rats fed on basal diet, diet containing untreated hazelnut and diets containing irradiated hazelnut with (6 kGy and 10kGy) presented in Table (10).

Feeding healthy rats on diet containing untreated hazelnut showed non-significant differences in serum cholesterol and HDL-c, as compared to healthy rats fed on basal diet. Untreated hazelnut caused significant increase in the mean value of triglycerides and significant decrease in LDL-c in healthy rats, as compared to healthy rats fed on basal diet. Treating healthy rats with irradiated hazelnut (6 and 10 kGy) recorded non-significant changes in total cholesterol, triglycerides, HDL-c and LDL-c, as compared to basil rats fed on untreated hazelnut, except LDL-c in the group treated with irradiation hazelnut with (10 kGy). In this respect, (Simone et al., 2016) who reported that Hazelnut-enriched diet is associated with a decrease of LDL and total cholesterol, while HDL cholesterol, triglycerides and body mass index (BMI) remain substantially unchanged. Mercanligilet al., (2007) reported that, hazelnut-enriched diet decreased the mean value of the ratios of total cholesterol/HDL cholesterol and LDL-c/HDL-c.

The effect of untreated and treated nuts with irradiation on antioxidant enzymes inhealthy rats:

Important determinants of cellular antioxidant enzyme are the enzymes Sodium Oxide Dismutase (SOD), Catalase (CAT) and Glutathion Peroxidase (GPx), which are responsible for the elimination of Reactive Oxygen Species ROS. Because these enzymes act sequentially to remove ROS, the balance of the activity of these enzymes may be as critical in the defense against ROS as the activity of the enzymes alone (*Boateng et al., 2016*).

Almond:

Data in table (11) showed the effect of untreated and treated almonds with irradiation (6 kGy and 10 kGy) on malonaldehyedand antioxidant enzymes of healthy rats. Feeding rats on diet containing untreated almonds showed significant decrease in the mean value of malonaldehyed, while the other parameters recorded significant increase, as compared to control group. On the other hand treating rats with irradiationat doses (6 and 10 kGy)showed non-significant changes in malonaldehyed, as compared to the group fed on diet containing untreatedalmond. On the other hand, the mean value of antioxidant enzymes including "catalase activity and superoxide dismutase activity in liver tissue" increased significantly in rats which treated with irradiated almonds, as compared to the control groups.

In this respect, **Soheil** et al., (2015) resulted that after treatment with almond hull, the serum level of MDA in all test groups significantly decreased. **David** et al., (2008) the full-dose almonds reduced serum Concentrations of malondial dehyde. **Li** et al., (2010) reported that feeding diabetic rats on diets containing irradiated almond with 6 and 10 kGy showed non-significant differences in

malonaldehyed, as compared to the group fed on diet containing untreated almond.

HazeInut:

Table (12) showed the effect of untreated and treated hazelnut with irradiation (6 kGy and 10 kGy) on malonaldehyed and antioxidant enzymes of healthy rats. The mean value of the malonaldehyeddecreased significantly in the groups treated with irradiated and un irradiated hazelnut, as compared to the control group. The mean values of GPX and SOD in healthy rats fed on diet containing untreated hazelnut showed non-significant changes, while catalase increased significantly, as compared to healthy rats fed on based diet.

Treated healthy groups with diets containing irradiated hazelnut with (6 and 10 kGy) showed non-significant differences in all antioxidant enzymes (Malonaldehyed, GPX, Catalase and SOD), as compared to the group of rats, which was fed on diet containing untreated hazelnut.

Aydan et al., (2004) indicate that hazelnut oil reduced oxidative stress and cholesterol accumulation in the aortas of rabbits fed on high cholesterol diet.

(Table 1): Approximate analysis of un-treated and treated almonds with irradiation.

Radiation		Zero	time			Storage (6 months)	
	Moisture	Ash	Protein	Lipid	Moisture	Ash	Protein	Lipid
Dose kGy	g/100 g	g/100g	g/100 g	g/100 g	g/100 g	g/100 g	g/100 g	g/100 g
Control	6.17 ^a ±0.186	3.03° ±0.03	28.1ª ±0.12	52.4ª ±0.2	6.53 ^a ±3.274	4.36 ^a ±0.0	23.5° ±0.12	54.6 a ±0.7
6 kGy	5.99 ^a ±0.521	2.83 ^b ±0.03	28.7ª ±0.15	54.2ª ±4.8	6.28 ^a ±0.643	4.46 ^a ±1.6	22.3 ^a ±0.15	54.2 a ±0.2
10 kGy	5.26 ^b ±0.066	2.73 ^b ±0.09	27.5 ^a ±0.12	52.9ª ±0.7	6.52 ^a ±1.574	4.30 ^a ±0.5	21.90 ^a ±0.12	54.1 a ±1.1

^{*}Data represented mean ± standard error.

(Table 2): Approximate analysis of un-treated and treated hazelnut with irradiation.

Radiation		Zero time				Storage (6 months)			
	Moisture	Ash	Protein	Lipid	Moisture	Ash	Protein	Lipid	
Dose kGy	g/100 g	g/100g	g/100 g						
Control	4.73 ^a	2.33ª	15.00 ^a	62.49 ^a	14.60 ^b	2.53 ^a	21.00 ^a	56.96ª	
Control	±0.176	±0.06	±0.12	±0.499	±0.200	±0.033	±0.12	±3.996	
6 kGv	4.20 ^b	1.83ª	15.20 ^a	59.07 ^b	15.13 ^b	2.03ª	19.90 ^a	54.80 ^a	
6 кСу	±0.058	±0.328	±0.15	±0.067	±0.570	±0.333	±0.14	±3.468	
10 kGy	4.10 ^b	1.71 ^a	16.80 ^a	58.13 ^b	16.70 ^a	2.10 ^a	20.10 ^a	51.56 ^a	
	±0.100	±0.00	±0.12	±1.10	±0.058	±0.05	±0.15	±1.14	

^{*} Data represented mean ± standard error.

^{*}Values at the same column with different letters are significant at P<0.05.

^{*}Values at the same column with different letters are significant at P<0.05.

Table (3): The percentage of fatty acids in control and irradiated almond at zero time and after storage for 6 months.

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	Almond at Zero time								
Fatty acid	Carbon no.	control	6 kGy	Change%	10kGy	Change%			
Palmitic acid	C 16:0	7.42	7.6	+ 2.4	9.5	+ 28.0			
Oleic acid	C 18:1	73.74	66.6	- 9.68	71.7	- 2.7			
Linoliec	C 18:2	18.8	25.69	+ 36.64	14.7	- 21.8			
	Alı	mond after sto	rage for s	x months					
Palmitic acid	C 16:0	10.6	12.05	+ 13.6	14.2	+ 33.96			
Oleic acid	C 18:1	76.42	68.40	- 10.49	57.9	- 24.23			
Linoliec	C 18:2	-	-	-	-	-			
Arachidic	C20:0	3.57	2.6	- 27.1	4.8	+ 34.45			

Table (4): The percentage of fatty acids in control and irradiated Hazelnut at zero time and after storage for 6 months.

•									
	Hazelnut at zero time								
Fatty acid	Carbon no.	control	6 kGy	Change%	10 kGy	Change%			
Palmitic acid	C 16:0	6.58	6.00	- 8.8	6.98	+ 6.0			
Oleic acid	C 18:1	81.28	74.3	- 8.5	79.61	- 2.0			
Linoliec	C 18:2	8.19	7.6	- 3.6	7.7	- 5.9			
	Ha	zelnut after s	torage for	six months					
Palmitic acid	C 16:0	5.9	10.2	+72	4.7	- 20.3			
Oleic acid	C 18:1	79.71	78.0	-2.1	54.3	- 31.8			
Linoliec	C 18:2	8.3	-	-	3.1	- 62.0			
Arachidic	C20:0	-	2.4		-				

Table (5): Relative organs weight of rats fed on diet containing untreated and treated almond with irradiation.

	Organs	Organs weight / body weight %					
Groups		Heart	Liver	Spleen	Testes	Kidney	
Basil die	nt .	0.7 ^a	5.02 ^a	0.56 ^b	1.34 ^b	1.13 ^a	
Dasii uit	5 1	±0.01	±0.14	±0.04	±0.08	±0.07	
Un-treated almond		0.68ª	4.79 ^a	0.67 ^{ab}	1.83ª	1.13 ^a	
UII-II ea	ieu aimonu	±0.09	±0.48	±0.11	±0.20	±0.09	
ed with	6 KGv	0.79 ^b	4.77 ^a	0.61 ^b	1.86ª	1.13 ^a	
	o Noy	±0.08	±0.37	±0.06	±0.07	±0.03	
Treat	<u>e</u> <u>10</u>	0.77 ^b	5.11ª	0.91°	1.88ª	1.13 ^a	
alu	KGy	0.09	±0.60	±0.10	±0.21	±0.19	
	P. Value	0.0001	0.0001	0.0001	0.0001	0.0001	

^{*}Values are expressed as means ±SD.

Table (6): Relative organs weight of rats fed on diet containing untreated and treated hazelnut with irradiation.

	Organs		Organs weight / body weight %				
Groups		Heart	Liver	Spleen	Testes	Kidney	
Basil diet		0.70 ^a	5.0 ^a	0.56ª	1.34 ^b	1.13 ^a	
Dasii ulet		±0.01	±0.14	±0.04	±0.08	±0.07	
Untreated		0.7 ^a	4.47 ^a	1.05 ^b	1.74 ^a	1.25 ^a	
hazelnut		±0.09	±0.57	±0.21	±0.16	±0.18	
_ +	6 KGv	0.71 ^a	4.95 ^a	0.98 ^{ab}	1.79ª	1.38ª	
Treated hazelnut with	6 KGy	±0.08	±0.46	±0.06	±0.20	±0.12	
reate azeln with	10KGy	0.65 ^a	4.70 ^a	0.82°	1.68ª	1.39 ^a	
Г <u>с</u>	TUNGY	±0.07	±0.44	±0.12	±0.14	±0.12	
P.Va	alue	0.0001	0.0001	0.0001	0.0001	0.0001	

^{*}Values are expressed as means ±SD.

^{*}Values at the same column with different letters are significant at P<0.05.

^{*}Values at the same column with different letters are significant at P<0.05.

Table (7): Effect of diet containing untreated and treated almond with irradiation on blood hemoglobin, serum glucose and liver enzymes of healthy rats.

	Parameters	Hemoglobin	Glucose	AST	ALT
groups		g/dl	mg/dl	U/L	U/L
Basil diet		15.73 ^a	74.22 ^a	83.75 ^{ab}	22.16 ^b
Dasii ület		±1.5	±7.0	±7.3	±1.10
Untreated	almond	15.72 ^a	83.46 ^{ab}	94.00°	26.83 ^{ab}
Uniteated	allilollu	±0.84	±5.3	±0	±0.83
	6 kGy	14.10 ^a	73.87 ^a	89.16ª	26.00 ^a
Treated almonds with	бкбу	±1.46	±5.4	±3.08	±0.110
rea Imc wi	10 kGy	16.90°	91.38°	88.80 ^a	26.00ª
р в	10 kGy	±3.87	±14.36	±3.32	±0.042
F	γ	0.8515	0.4517	0.3165	0.0014

^{*}Values are expressed as means ±SD.

Table (8): Effect of diets containing untreated and treated hazelnut with irradiation on blood hemoglobin, serum glucose and liver enzymes of healthy rats.

Para	Parameters		Glucose	AST	ALT
groups		g/l	mg/l	U/L	U/L
Basil diet		15.73⁵	74.22 ^a	83.75ª	22.16 ^a
Dasii ület		±1.5	±7.0	±7.3	±1.10
Untreated h	azalnut	13.17°	80.51 ^{ab}	94.00 ^a	13.83 ^b
Officeated fi	azemut	±1.03	±6.17	±0	±0.94
	6 kCv	16.59 ^a	72.72 ^{ab}	94.00 ^a	16.00 ^{ab}
Treated hazelnut with	6 kGy	±1.51	±16.03	±1.11	±0.81
rea aze wi	10	16.68 ^a	78.34 ^{ac}	94.00 ^a	15.00 ^b
ר ב	kGy	±1.71	±3.03	±0	±1.09
PV		0.326	0.932	0.179	0.0065

^{*}Values are expressed as means ±SD.

^{*}Values at the same column with different letters are significant at P<0.05.

^{*}Values at the same column with different letters are significant at P<0.05.

Table (9): Effect of diet containing untreated and treated almond with irradiation on lipid profile of healthy rats.

P	arameters	Cholesterol	Triacylglycerol	HDL-c	LDL-c
Groups		mg/dl	mg/dl	mg/dl	mg/dl
Basil diet		139.00 ^b	79.37 ^a	61.56 ^a	61.57ª
Dasii ulet		±4.35	±17.18	±1.14	±7.10
Untreated	almond	138.33 ^{ab}	155.66 ^b	63.46 ^a	44.70 ^b
Uniteated	alliloriu	±7.53	±4.70	±0.50	±7.77
	6 kGy	135.00 ^a	149.00 ^b	61.64ª	44.64 ^b
Treated almonds with	0 KGy	±3.57	±6.80	±1.43	±0.27
reate Imono with	10 kGy	135.33 ^a	146.66 ^b	61.75 ^a	44.25 ^b
г а		±1.66	±25.71	±1.81	±4.54

^{*}Values are expressed as means ±SD.

Table (10): Effect of diet containing untreated and treated hazelnut with irradiation on lipid profile of healthy rats.

Pa	rameters	Cholesterol	Triacylglycerol	HDL-c	LDL-c
Groups		mg/dl	mg/dl	mg/dl	mg/dl
Basil die		139 ^b	79.33°	61.56 ^a	61.57°
Dasii ule	ı	±4.35	±17.18	±1.14	±7.10
Untreated		133.33 ^{ab}	149.66ª	60.42 ^a	38.98ª
hazelnut		±11.78	±15.34	±0.23	±8.83
	6 kGv	131.33ª	161.33 ^{ab}	60.61 ^a	37.45 ^a
Treated hazelnut with	th the department of the depar	±4.66	±8.29	±0.77	±4.36
aze W	10 kGy	131.00ª	172.66 ^{ab}	60.99 ^a	45.47 ^b
- ح	10 kGy	±00	±5.84	±0.87	±1.64

^{*}Values are expressed as means ±SD.

^{*}Values at the same column with different letters are significant at P<0.05.

^{*}Values at the same column with different letters are significant at P<0.05.

Table (11): Effect of diet containing untreated and treated almond with irradiation on malonaldehyed and antioxidant enzymes of healthy rats.

		Malanaldahuadaa/aT	GPX	Catalase	SOD
		Malonaldehyedng/gT	U/gT	U/gT	U/gT
Basil diet		5.45ª	64.87°	70.65°	20.59 ^d
Dasii ület		±0.18	±0.79	±0.19	±0.07
Untreated a	lmond	3.32 ^b	66.79 ^b	82.08 ^b	23.03°
Officeated a	imona	±0.7	±0.25	±4.79	±0.16
	6 KGv	3.6 ^b	67.94 ^{ab}	93.21ª	26.07 ^a
ted ond	6 KGy	±0.301	±0.21	±0.76	±0.22
Treated almond with	10	3.8 ^b	68.72 ^a	94.63ª	24.85 ^b
F '	KGy	±0.401	±0.74	±1.94	±0.63
	KGy	±0.401	±0.74	±1.94	±0.63

^{*}Values are expressed as means ±SD.

Table (12): Effect of diet containing untreated and treated hazelnut on antioxidant enzymes of healthy rats.

		Malanaldahyadaa/aT	GPX	Catalase	SOD
		Malonaldehyedng/gT	U/gT	U/gT	U/gT
Basil diet		5.45° ±0.18	64.87 ^b	70.65°	20.59 ^b
Basii diet		5.45°±0.16	±0.49	±0.19	±0.07
Untreated		4.65 ^b ±0.75	65.70 ^{ab}	95.89 ^{ab}	21.93 ^{ab}
hazelnut		4.03 ±0.73	±1.27	±6.04	±1.12
	6	4.65 ^b ±0.50	66.72 ^{ab}	105.11 ^{ab}	22.47 ^a
nut ad	KGy	4.05° ±0.50	±0.77	±3.25	±0.25
Treated hazelnut with	10 KGy	4.64 ^b ±0.311	71.32 ^a ±3.26	110.76 ^a ±0.93	22.70 ^a ±0.16

^{*}Values are expressed as means ±SD.

^{*}Values at the same column with different letters are significant at P<0.05.

^{*}Values at the same column with different letters are significant at P<0.05.

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تأثير اللوز والبندق المشع والمدعم للوجبات على بعض الخصائص البيوكيميائيه في الفئران

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المستخلص العربى

تهدف هذه الدراسة إلى تقييم تأثير الإشعاع عند الجرعات Γ كيلوجرايو Γ كيلوجراي على بعض التقديرات الكيميائية والكيميائية الحيوية للمكسرات المتضمنة اللوز والبندق. استخدم في هذه الدراسة Γ من جرذان الالبينو، قسمت هذه الجرذان إلى Γ مجموعة على النحو التالي: المجموعة Γ المجموعة (Γ): تتغذى على الغذائي الاساسي، واستخدمتكمجموعة ضابطة. المجموعة Γ تتغذى على تغذيتها على غذائي أساسي يحتوي على لوز غير مشعع. المجموعات (Γ و Γ): تتغذى على وجبات غذائية محتوية على اللوز المشعع Γ و Γ كيلو جراي، على التوالي. المجموعة (Γ): تتغذى على نتغذى على غذاءاساسي يحتوي على البندق غير المشعع. المجموعات (Γ و Γ): تتغذى على الوجبات الغذائية المحتوية على البندق المشعع Γ و Γ كيلوجراي على التوالي لمدة Γ أسابيع.

أظهر التحليل الإحصائي لللوز والبندق المعامل بجرعات ٦ و١٠ كيلو جراي عند بدء التجربة وبعد التخزين لمدة ٦ أشهر في التحليل الكيميائي عدم وجود فرق كبير في البروتين والرطوبة والرماد والدهون باستثناء بعض المكسرات المعاملة بالجرعات الإشعاعيةالمختلفة. ولم

تظهر التحاليل الكيمائية الحيوية وجود تأثير معنوي على وزن الأعضاء ، وكولسترولالليبوبروتينات منخفضه الكثافة (HDL-c) ، وكولسترولالليبوبروتينات منخفضه الكثافة (LDL-c) ، وهيموجلوبين الدم ، والكولسترول الكلي ، و الجليسريداتالثلاثية ، والجلوكوز ، وألانين أمينترانسفيراز (ALT) ، سوبراوكسيديسمايوتيز (SOD) ، جلوتاثيونبيروكسيديز و كتاليز الكبد في مجموعات الفئران التي عوملت باللوز والبندق غير المشعع أو المشعع ، بالمقارنة بالمجموعة الضابطة التي تغذت على الغذاء الاساسي، باستثناء الانين أمينو ترانسفيراز (ALT) التي أظهرت زيادة في اللوز الغير معامل وانخفضت في البندق الغير معامل ، وكان هناك زيادة كبيرة في الانزيمات المضادة للأكسدة في جميع العينات.