Effect of irradiated pistachio and walnuts enriched-diet on biochemical aspects in rats

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Abstract

The effect present study aims to evaluate of irradiationatdoses 6 kGy and 10 kGy on chemical and biochemical parameters of sometypes ofnuts including pistachio (Pistachiavera L.) and walnuts (JuglansregiaL.) 70albinorats used 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 unirradiated pistachio. Groups (3 and 4): fed on diets containing the irradiated pistachio on 6 and 10kGy, respectively. Group (5): fed on basal diet containing un-irradiated walnut. Groups (6 and 7): fed on diets containing the irradiated walnut on 6 and 10 kGy, respectively for 8 weeks. The statistical analysis of irradiated pistachio and walnut at dose 6 and 10 kGy at zero time and storage for 6 months showed no significant difference in protein, moisture, ash and lipid. The biochemical performance showed non-significant effect on organs weight, total cholesterol, triacylglycerol's, high density lipoproteincholesterol (HDL-c), low density lipoprotein-cholesterol (LDL-c), blood hemoglobin, glucose, serum alanine amino transferase (ALT), serum

aspartate amino transferase (AST), liver superoxide dismutase (SOD), liver glutathione peroxidase (GPX) and liver catalase in groups applied doses of non-irradiated or irradiated pistachio and walnut, as compared to control group which fed on basal diet , while serum alanine amino transferase (ALT) which showed increased in untreatedpistachio and there were significant increase in antioxidant enzymes at all groups.

Introduction

Pistachio (*Pistachiavera L.*) belongs to Anacardiacea family being one of the most important edible nuts(*Tomaino et al.,2010*). The tendency toward consumption of pistachio nuts has been increased due to their nutrient contents such as sterols, vitamins, minerals, fatty acids, phenolic compounds, protein and dietary fiber. Pistachio nuts contain about 50% oil, an oil rich in oleicand linoleic acids, which provides important therapeutically effects. Oleic acid is an important monounsaturated fatty acid that helps in reducing triglycerides, low-density lipoproteins (LDL), total cholesterol and glycemic index. In addition, oleic acid is responsible for the increase of stability and reduction of oxidation in vegetable oils (*Kamazani et al., 2015*).

During the drying process, nuts can undergo undesirable reactions (especially rancidity) which cause degradation of quality, because of the odd colors and flavors formed; the major oxidative reactions in dried foods are due to peroxidation of lipids. Lipid oxidation in foods is associated almost exclusively with unsaturated fatty acids and it is often autocatalytic, with oxidation products themselves catalyzing the reaction so that the rate increases. The pistachio is a nut with a high lipid content and very rich in unsaturated fatty acids; this makes pistachio nuts very sensitive product owing to rancidity (*Kashani et al., 2008*).

Pistachio green hull (PGH) is a rich source of phenolic compounds with high antioxidant activity. Gallic acid is a predominate compound of (PGH) and can scavenge free radicals, chelating metals and reducing tocopherol radicals (*Aliet al., 2018*).

Walnuts (*JuglansregiaL.*) are widely distributed all over the world. Walnuts are receiving increasing interest as a healthy foodstuff because their regular consumption has been reported to decrease the risk of coronary heart disease. Therefore, walnuts can be utilized as ingredients of many foodstuffs such as bakery products to enhance the nutrition value and sensory properties of the final product(*Xiaoyingand Yufei 2012*).

Many beneficial effects on human health have been related to walnut consumption and attributed largely to oil composition, such as prevention of cardiovascular diseases, type 2 diabetes in women, and cholesterol increases. Walnut oil is predominantly composed of polyunsaturated fatty acids (PUFA), mainly omega6 linoleic acid, and omega3 linolenic acid in contrast to other nuts that they have high content of monounsaturated fatty acids (MUFA)*(Miltiadisand Eleni 2015).*

English walnuts have been reported to be an excellent source of antioxidants (*Jacki et al., 2011*). Walnut is rich in polyphenols and contains numerous ellagitannins, walnut polyphenol not only has antioxidant and anti-inflammatory activity but improves interneuronal signaling and increases neurogenesis (*Dandan et al., 2014*).

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 and Mohácsi-Farkas2011).*

Therefore, the present study was carried out to assess the effects of diet containing untreated and treated pistachio and walnuts on healthy rats.

Material and methods

Material

Pistachio (*Pistachiavera L.*), Walnuts (*Juglansregia*L.), 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, serumaspartate amino transferase (AST)serum alanine amino transferase (ALT).cholesterol, triglycerides; HDL-c and antioxidant enzymes

including (malonaldehyed, liver glutathione peroxidase (GPX), Catalase andliver superoxide dismutase (SOD) were obtained from Gama tread Company, Cairo, Egypt.

Methods

γ Irradiation treatment

Pistachio (*Pistachiavera L.*) and Walnuts (*JuglansregiaL.*) were packed in polyethylene bags, and sealed by heat. Each bag contained about 200 g. they were subjected γ 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 pistachio and walnuts

Moisture content, crude fat, crude protein, ash contentofpistachio and walnuts, were determined according to the method described by the (*A.O.A.C., 2003*).

The fatty acids composition were determined as the method described by (*Hamilton and Hamilton, 1993*).

Diet preparation:

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

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Experimental rats

male albino rats, Sprague-Dawley strain, with an Seventy 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) pistachio and walnut, 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 pistachio and treated pistachio with 6 and 10 kGy, respectively. Group 5, 6 and 7 were feed on untreated walnut and treated walnut with 6 and 10 kGy, respectively.

During the experimental period (6 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*), highdensity 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*).Hemoglobin were 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 metabolio enzymes 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)*.

Results and Discussion

Effect of radiation processing with storage on chemical composition of Pistachio and Walnut.

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

Pistachio

The variation of moisture, ash, crude protein and crude fat of pistachio treated with irradiation at zero time and after storage for six months were studied and the results are presented in Tables (1). The mean values ofmoisture content of pistachio showed non-significant changes after treated with irradiation at dose (6 and 10 kGy). Ash content of pistachio, which treated with irradiation at dose (10kGy), showed significant decreasethan the control sample (at zero time). The other nutrients (protein and lipid) recorded non-significant changes with the two dosage of irradiation, as compared to the control sample at zero time.

Results in this Table indicated that, non-significant changes in the chemical composition of pistachio which treated with irradiation at two doses with storage, while the content of ash in pistachio, which treated with irradiation at dose 10 kGy, decreased, as compared to the control sample. (*Mahfouz, 2014*) resulted that irradiation doses of 1, 2 and 3 kGy of gamma irradiation showed not substantially affected in contents of moisture, proteins, lipid, and ash, with respect to the pistachio control samples.

Walnut

The mean values of moisture, ash, crude protein and crude fat of walnut treated with irradiation with doses 6 and 10 kGy at zero time and after storage for six months were studied and the results are presented in Table (2).The data showed that, at zero time, nonsignificant differences were observed in moisture, ash and protein in un-treated walnut and treated with irradiation at (6 and 10 kGy), while total lipid in this type of nuts which treated at dose 10 kGy, it increased, as compared to the control sample.Storage for 6 months

presented untreated and treated walnut with irradiation at (6 and10 kGy) induced non-significant changes in moisture, ash and protein, while total lipid increased significantly in walnut which treated with 6 and 10 kGy, as compared to the control sample.Generally, there was non-significant effect in moisture at zero time but there was significant effect in moisture at storage period may be due to the humidity from storage condition.

The average values for protein and fat were18.1% and 58.2% in walnut respectively *(Ferhad et al., 2010)* and *(Sza-Tao and sathe ,2000)* reported that, walnut contained 16.66% protein and 66.90% lipids on a dry weight basis. Two walnut cultivars were examined as fresh samples and under storage at 8 °C for two months. The oil content is the lowest in fresh samples of all the cultivars. At this stage, the oil content of 'Chandler' was lower than of the others. The results show significant difference between the two ways of drying for the Hungarian cultivars (P<0.05) *(Konya et al., 2015)*.

Fatty acids composition of Pistachio and Walnut treated with irradiation at zero time and after storage period:

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

Pistachio

The data in Table (3) indicated that, at zero time the control pistachio contain 13.09% saturated fatty acids (SFA) (palmitic acid) that was the major saturated fatty acid. Treating pistachio sample with irradiation led to decrease in palmitic acid by about 17.2% at 6kGy and decreased at 10 kGy by 20.01%. Monounsaturated fatty

acid (MUFA) (oliec acid) of raw sample was 69.97% that was the major (MUFA). The data in this table showed that there is change in oleic acid at 6kGy increased by 1.7% and at 10 kGydecreased by 0.7%.Polyunsaturated fatty acids (PUFA) (linoleic acid) of control sample was 16.92 % that the major (PUFA). The data indicated that the linoleic acid increased by about (6.6% and 12.29%) at 6 kGy and 10 kGy, respectively, than that of that control. After 6 months palmitic acid increased by about 5.4%;oliec acid decreased by 0.4%, and linoliec acid were increased by 3.3%, when treated pistachio with irradiation at 6 kGy, while at 10kGy the palmitic acid, olic acid and linoliec acid decreased by about (1.0%, 9.0% and 16.7%), respectively in pistachio, as compared to the control sample.

*Mahfouz(2014)*indicated that the highest used dose (3kGy) to pistachio samples slightly decreased the fatty acid content.

Walnut

The data in Table (4) indicated that, at zero time the control walnut contain 15.7% saturated fatty acids (SFA) (palmitic acid and stearic acid) the major saturated fatty acid was palmitic acid (11.59%)followed by and stearic acid was (4.13%). Treating walnut sample with irradiation decrease in palmitic acid change by 35.97% at 6kGy and increased at 10 kGy by 9.4%).Monounsaturated fatty acid (MUFA) (Oliec acid) of raw sample was 23.33% that was the major (MUFA). Oleic acid decreased by 31.67% at 6 kGy and by 0.8% at 10 kGy.Polyunsaturated fatty acids (PUFA) (linoleic acid) of control sample was 50.16 % that the major (PUFA). The data indicated that the linoleic acid increased by 28.42% at 6kGy and decreased by 2.3% at 10 kGy.

After storage for 6months saturated fatty acids decreased in walnut with irradiation, while unsaturated fatty acids increased. **Stamatios and Michael (2009)** determined treated and untreated walnuts with irradiation doses at 1.0, 1.5, 3.0, 5.0 and 7.0 kGy. The fatty acids determined, stearic and palmitic acids concentration increased while oleic acid decreased with irradiation dose. Polyunsaturated fatty acids were unaffected by irradiation.

*Umit et al., (2011)*The concentration of total saturated fatty acids increased while total mono unsaturated and total polyunsaturated fatty acids decreased in hazelnut, walnut almonds and pistachio nut which treated with 1, 3, 5 and 7 kGy.

Vassilia et al., (2015)GC-FID results showed that SFA increased and MUFA and PUFA decreased with the increase of irradiation dose (5, 10, 13 KGy dose). Moreover, MUFA/SFA and PUFA/SFA ratios decreased (P< 0.05) compared to control samples.

Biological analysis for rats feed on diet contain untreated and treated pistachio and walnutwith irradiation

Relative organs weight of rats Pistachio

The mean value ±SD of relative organs weights of rats fed on diets containing un-treated and treated pistachio with irradiation at dose (6 & 10 kGy) showed in Table (5). The results indicated that, all organs weights, recorded non-significant changes in-group of rats fed on diet containing untreated and treated pistachio with two dosage of irradiation, as compared to the control group, which fed on control diet onlyexcept the mean value of spleen, which recorded significant increase.

Walnut

The mean value ±SD of relative organs weight of rats fed on diets containing un-treated and treated walnut with irradiation at dose (6 & 10 kGy) showed in Table (6). The mean values of heart and liver weight showed non-significant changes in the group fed on untreated walnut, while the weight oftests and kidney recorded significant increase, as compared to the control (fed on basal diet only). The two treated groups with irradiation with dose (6 and 10 kGy) showed significant increase in heart, spleen and kidney weights, while liver and tests recorded non-significant differences, as compared to the group fed on untreated walnut.

Effect of diet containing untreated and treated pistachio and walnut on somebiochemical analysis of healthy rats. Pistachio

The effect of enriched diet with untreated and treated pistachio with irradiation at (6 & 10 kGy) on hemoglobin, glucose, AST and ALT of healthy rats presented in Table (7). The results in this Table showed significant differences in the mean values of hemoglobin, glucose, AST and ALT between the groups fed on diet containing untreated pistachio, as compared to control group. Treated pistachiowith irradiation at dose 6 and 10 kGy, led to significant decrease in the mean value of hemoglobin and glucose levels, as compared to untreated pistachio. The data in this Table showed significant decrease in AST and ALT enzymes with high dose from irradiation, as compared to the control group (fed on untreated pistachio).

Walnut

The effect of enriched diet with untreated and treated walnut with irradiation at (6 & 10 kGy) on hemoglobin, glucose, AST and ALT of basil diet rats presented in Table (8). Non-significant changes in all tested parameters was observed between the groups fed on diet containing (un-treated or treated walnut with irradiation at dose 6 and 10 kGy), except AST and ALT enzymes in the group of rats which treated with 10 kGy, as compared to the control group fed onbasal diet only.

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*).

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 adiponectine 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 (PYY) 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). 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.

*Nazirogluet al., (1999)*reported that, administered vitamin E has protective effects against CCl₄-induced chronic liver damage and cirrhosis as evidenced by biochemical data and conventional histological examination.

Effect of diet containing untreated and treated pistachio and walnut on lipid profile of healthy rats: Pistachio

The effect of feeding rats ondiet containing untreated and treated pistachio, on lipid profile of healthy rats presented in Table (9).The results indicated that, non-significant changes in all parameters of lipid profile in rats fed on diet containing untreated

pistachio, as compared to healthy rats fed on basal diet, except triacylglycerol which showed significant increase. On the other hand, feeding rats on diets containing pistachio, which treated with irradiation (6 & 10 kGy) caused non-significant differences in serum cholesterol, triglyceride, HDL-c and LDL-c, as compared to the group fed on diet containing untreated pistachio. In this respect, *(Nur et al,. 2007)* reported that, consumption of pistachio as 20% of daily caloric intake increased high-density lipoprotein (HDL) levels and decreased the ration of total cholesterol/ high density lipoprotein TC/HDL, compared with those not taken pistachio.

In general, all types untreated and treated nuts did not changed the lipid profile as compared to the control group, or improved the most parameters of lipid profile. In this respect, Clinical trials up to 2009 have already shown that regular nut consumption can have beneficial effects on cholesterol and triglyceride levels Subsequent studies have confirmed these effects. Several intervention studies obtained with different types of nuts (pistachios, Brazil nuts, walnuts. almonds. hazelnuts or peanuts) and consumption patterns (30-80 g/d) in healthy individuals and subjects with hyperlipidemia showed both a reduction in total cholesterol (TC) by an average of 10.1 % as well as a reduction in LDL-cholesterol (LDL-C) by an average of 8.6 %(Damasceno et al., 2011 and McKiernan et al., 2010).

These effects were often attributed to the favorable fatty acid profile and the dietary fiber content of nuts. Some studies also showed an increase in HDL-cholesterol (HDL-C) as well as a decrease in triglycerides (TG) and apolipoprotein B (Apo B) (*Sabate et al., 2003 and Tey et al., 2011*) those results were particularly evident in people already suffering from hyperlipidemia.

Another intervention study even achieved in obese participants an improvement of blood-lipid parameters, but the level of significance for this effect was only reached in subjects with hyperlipidemia (*McKiernan et al., 2010*).

Walnut

The result of serum cholesterol, Triacylglycerol, HDL-c and LDL-c in rats fed on basal diet, diet containing untreated walnut and diets containing irradiated walnut with (6 kGy and 10kGy) presented in Table (10).Feeding healthy rats on diet containing untreated walnut showed non-significant differences in all lipid profile, except triglyceride, which showed significant increase, and LDL-c which recorded significant decrease, as compared to healthy rats fed on basal diet. Treating healthy rats with irradiated walnut (6 and 10 kGy) recorded non-significant changes in total cholesterol, triglyceride, HDL-c and LDL-c, except the mean value of serum cholesterol in treated rats with irradiated walnut with 10 kGy and LDL-c in the group which treated with irradiated walnut. *Jacki et al., (2011)* reported that, walnut consumption has been associated with improvements in serum lipid profile.

The effect of untreated and treated pistachio and walnut with irradiation on antioxidant enzymes in 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*).

Pistachio

Feeding rats on diet containing treated and untreated pistachiowithirradiation on antioxidant enzymes including malonaldehyed and antioxidant enzymesof healthy rats presented in Table (11). The table showed decrease malonaldehyed significantly in the group feed with untreated pistachio, while non-significant changes in GPX, Catalase and SOD were observed, as compared to healthy rats, fed on basal diet.Treating pistachio with the two doses from irradiation (6 and 10 kGy) showed significant decrease in malonaldehyed, while the other antioxidant enzymes recorded significant increase with irradiation, as compared to the group feed with untreated pistachio.

Sari and Bagic (2010) who reported that, substituted saturated fat with pistachio (comprising about 20% of daily caloric intake) for a period of 4 weeks. The pistachio diet led to lower glucose, lipids, total oxidant status, and malondialdehyde, and increased superoxide dismutase.

Walnut

Data in Tables (12) revealed that, the mean values of malonaldehyed decreased significantly in the group fed on untreated walnut, while GPX and SOD showed non-significant changes, as compared to the control group. On the other hand, the mean value of catalase increased significantly in healthy rats fed on diet containing untreated walnut, as compared to control group fed on basal diet. The mean value of malonaldehyed in healthy groups, which were feed with irradiated walnut with 6 and 10 kGy, showed significant

differences; on the other hand, the other antioxidant enzymes antioxidant enzymesrecorded significant increase, as compared to the group feed with untreated walnut. In this respect, (*Dandan et al., 2014*) stated that, walnut polyphenol (WP) significantly decreased serum total triglycerides, cholesterol and malondialdehyde (MDA) level and increased superoxide dismutase (SOD) activity. Administration of WP significantly decreased MDA level and increased SOD activity in brain tissues. The treatment of rats with alcohol and alcohol + walnut containing diet supplementation caused changes in the level of serum enzymes and (MDA) content (*Bedia et al., 2015*).

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

Radiation		Zero	time		Storage (6 months)			
dose kGy	Moisture	Ash	Protein	Lipid	Moisture	Ash	Protein	Lipid
Control	7.73 [⊳]	2.66ª	28.10 ^a	52.65ª	6.63ª	3.16ª	21.10ª	60.47ª
	±0.088	±0.033	±0	±1.012	±0.328	±0.33	±0	±5.567
6 kGy	6.86 ^b	2.50 ^{ab}	28.70 ^a	52.03 ^a	3.91ª	3.50 ^a	19.60 ^a	54.67 ^a
	±0.120	±0.000	±0	±0.328	±1.000	±0.300	±0	0.338
10 40 4	6.36 ^b	2.47b	27.50 ^a	52.19 ^a	5.93ª	2.40 ^b	18.20ª	54.90 ^a
IU KOY	±0.239	±0.088	±0	±0.634	±1.517	±0.230	±0	±0.000

*Data represented mean ± standard error.

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

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

Radiation		Zero	time		Storage (6 months)			
dose kGy	Moisture	Ash	Protein	Lipid	Moisture	Ash	Protein	Lipid
Control	6.25ª	1.60ª	21.10 ^a	67.97ª	10.43ª	1.83ª	21.80 ^a	60.17 ^c
	±1.134	±0.100	±0	±0.233	±0.219	±0.939	±0	±0.233
6 kGy	4.80ª	1.73ª	22.60 ^a	64.73 ^a	11.60ª	2.00 ^a	20.30 ^a	66.70 ^b
6 кGy	±0.058	±0.033	±0	±1.524	±0.602	±0.306	±0	±0.000
10 kGy	4.33ª	1.80ª	22.50ª	64.13⁵	11.77ª	1.43ª	22.60 ^a	69.80 ^a
	±0.333	±0.057	±0	±0.677	±0.426	±0.290	±0	±0.000

*Data represented mean ± standard error.

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

Zero time									
Fatty acid	Carbon no.	control	6 kGy	Change%	10kGy	Change%			
Palmitic acid	C 16:0	13.09	10.77	- 17.2	10.47	- 20.01			
Oleic acid	C 18:1	69.97	71.18	+ 1.7	70.52	+ 0.7			
Linoliec	C 18:2	16.92	18.04	+ 6.6	19.0	+12.29			
	Pis	stachio after sto	rage for 6	months					
Palmitic acid	C 16:0	9.80	10.33	+5.4	9.7	-1.02			
Oleic acid	C 18:1	70.81	70.47	-0.4	64.4	-9.0			
Linoliec	C 18:2	17.3	17.9	-3.3	14.4	-16.7			

Table (4): The percentage of fatty acids in control and irradiatedwalnut with irradiation at zero time and after storage for 6ment

Zero time										
Fatty acid	Carbon no.	Control	Control 6 kG		Ch	ange%	1	0 kGy	Change%	
Palmitic acid	C 16:0	11.59	7.4	2	-	- 35.97		12.68	+ 9.4	
Stearic	C 18:0	4.13						4.17	+ 0.9	
Oleic acid	C 18:1	23.33	15.9	94 - 3		- 31.67		23.14	- 0.8	
Linoliec	C 18:2	50.16	53.9	90 + 2		- 28.42		49.0	- 2.3	
		Six	month	3						
Palmitic acid	C 16:0	13.5		13.	35	- 1.1		8.3	- 38.51	
Stearic	C 18:0	31.96	6	26.	22	- 17.9	5	21.74	- 31.97	
Linoliec	C 18:2	35.6		38.	97	+ 9.4	6	45.8	+28.65	
Linolenic	C 18:3	3.0		3.	6	+ 20		7.9	+ 163	

treated and treated pistachio with irradiation.								
		Heart (g /	Liver (g/	Spleen (g/	Tests (g/	Kidney (g/		
		100 g body	100 g body	100 g body	100 g body	100 g body		
		weight)	weight)	weight)	weight)	weight)		
Basil diat		0.70 ^a	5.02 ^a	0.56 ^c	1.34 ^{ab}	1.13ª		
Basil diet		±0.01	±0.14	±0.04	±0.08	±0.07		
untreated		0.69ª	4.59 ^a	0.67ª	1.75ª	1.03ª		
pistachio		±0.04	±0.14	±0.06	±0.06	±0.04		
0	6	0.65ª	5.02 ^a	0.60 ^b	1.40 ^b	1.14ª		
th th	KGy	±0.08	±0.27	±0.06	±0.04	±0.05		
rea ista	10	0.8ª	4.48 ^a	0.60 ^b	1.51ª	1.03ª		
⊢ <u> </u>	KGy	±0.06	±0.15	±0.05	±010	±0.04		
P. val	ue	0.0001	0.0001	0.0001	0.0001	0.0001		

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

*Values are expressed as means ±SD.

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

Table (6): Relative weight of body organs of rats feed on un-treated and treated walnu twith irradiation

Padia	tion	Heart (g /	Liver (g /	Spleen (g /	Tests (g /	Kidney (g /	
doses		100 g body	100 g body	100 g body	100 g body	100 g body	
		weight)	weight)	weight)	weight)	weight)	
Pooil dio	+	0.70 ^a	5.02 ^a	0.56 ^a	1.34 ^b	1.13°	
Dasii ule	ι	±0.02	±0.14	±0.04	±0.08	±0.07	
untreated		0.69 ^a	4.88ª	0.506 ^b	1.81ª	1.23 ^b	
walnut		±0.0.06	±0.21	±0.03	±0.12	±0.07	
th _	6	0.81 ^b	5.34ª	0.58ª	1.66 ^{ab}	1.30ª	
ated t wi	KGy	±0.06	±0.37	±0.05	±0.09	±0.04	
Trea	10	0.77 ^a	4.89 ^a	0.60ª	1.93ª	1.29 ^a	
L S	KGy	±0.03	±0.26	±0.071	±0.16	±0.05	
P. va	lue	0.0004	0.581	0.580	0.013	0.220	

*Values are expressed as means ±SD.

Table (7): Effect of diet containing untreated and treated pistachio on blood hemoglobin, serum glucose and liver enzymes

Р	arameters		Glucose	A ST 11/1	
Groups		Tiemogiobin mg/i	g/l	A31 0/L	ALT U/L
Bacil diot		15.73 [⊳]	74.22 ^a	90.75 ^a	22.16 ^b
Dasii ulet		±1.5	±7.0	±7.3	±1.10
Untreated		18.58ª	88.27 ^b	89.00 ^b	21.33 ^b
pistachio		±1.14	±15.02	±2.34	±1.47
0	6 kCv	14.44 ^c	64.00 ^b	90.00 ^a	25.33ª
ated Ichic ith	0 KGy	±1.49	±6.97	±0	±0.42
realista	10 40 4	14.68°	71.73 ^a	83.33°	15.50°
	TUKGy	±1.17	±1.17	±2.65	±0.50
PV		0.326	0.579	0.384	0.0001

*Values are expressed as means ±SD.

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

Table (8):	Effect c	of diet	containing	untreated	and	treated	walnut	on
hemoglobin, glucose and liver enzymes								

Par	rameters	Hemoglobin mg/l	Glucose g/l	AST U/L	ALT U/L
Groups					
Bacil diot		15.73 ^b	74.22 ^{ab}	91.75 [⊳]	22.16°
Dasii ulet		±1.52	±7.03	±7.30	±1.10
		17.46 ^a	81.23 ^b	81.25ª	17.66 ^{ab}
Uniteateu	wannut	±1.15	±10.50	±1.03	±1.17
6 kGy	6 kGy	17.40 ^a	75.52°	83.00 ^{ab}	18.33 ^b
	0 KGy	±0.59	±3.37	±2.00	±2.71
val val	10	17.14ª	88.99ª	80.75ª	17.16 ^b
⊢	kGy	±1.58	±4.07	±4.98	±1.77
PV		0.748	0.449	0.268	0.227

*Values are expressed as means ±SD.

	on lipid profile of healthy rats.							
Para	ameters	Cholesterol	Triacylglycerol	HDL-c	LDL-c			
Groups	<u> </u>	mg/dL	mg/dL	mg/dL	mg/dL			
Basil dia		139 ^{ab}	79.33 ^b	61.56ª	61.57 ^{ab}			
Dasii ulei	L	±4.35	±17.18	±1.14	±7.10			
Untreated		138.33ª	151.00ª	62.32ª	48.48 ^b			
pistachio		±2.33	±10.50	±1.81	±11.10			
	6 kGy	135ª	112.33ª	61.42ª	69.11ª			
be oir	₽ .9 b kGy	±5.68	±29.31	±1.43	±6.40			
Treate pistach with ADA	134.00 ^{ab} ±12.5	178.50ª ±39.50	61.13ª ±0.85	51.66 ^b ±21.25				

 Table (9): Effect of diet containing untreated and treated pistachio, on lipid profile of healthy rats.

*Values are expressed as means ±SD.

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

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

Pa	rameters	Cholesterol	Triacylglycerol	HDL-c	LDL-c
Groups		mg/dL	mg/dL	mg/dL	mg/dL
Basil dia	+	139.00 ^b	79.33 ^b	61.56ª	61.57°
Dasii ule	L	±4.3	±17.18	±1.14	±7.10
Untreated walnut		139.33 ^b	151.33ª	58.33ª	50.73 ^b
		±8.08	±18.67	±3.61	±6.02
_	6 kGy	136.66 ^{ba}	170.66ª	58.90ª	44.63ª
tth nut	±5.36	±6.43	±1.05	±6.27	
rea wal	10 kGy	134.33ª	127.00ª	51.68ª	56.25 ^{ab}
	10 KGy	±5.33	±11.15	±5.11	±7.56

*Values are expressed as means ±SD.

Table (11): Effect of diet containing untreated and treated almond or	ſ
malonaldehyed and antioxidant enzymes of healthy rats.	

Rarameters		Malonaldehyed	GPX	Catalase	SOD
Groups		ng/gT	U/gT	U/gT	U/gT
Basil diet		5.45°	64.87 ^b	70.65 ^b	20.59 ^b
		±0.18	±0.49	±0.19	±0.07
Untreated pistachio		3.90 ^b	^b 66.06	86.48 ^b	21.46 ^b
		±0.49	±0.58	±5.19	±0.72
Treated pistachio with	6 KGy	3.32 ^b ±0.30	69.62ª ±0.75	93.18ª ±0.75	27.77ª ±0.189
	10 KGy	2.18 ^a ±0.18	^a 69.29 ±0.82	94.68ª ±1.12	27.46 ^a ±1.23

*Values are expressed as means ±SD.

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

Table (12):	Effect	of diet	containing	untreated	and	treated	walnut	on
	malona	aldehye	d and antic	xidant enz	ymes	s of hea	Ithy rate	5.

Parameters		Malonaldehyed	GPX	Catalase	SOD
Groups		ng/gT	U/gT	U/gT	U/gT
Basil diet		5.45ª	64.87 ^{bc}	70.65°	20.59 ^b
		±0.18	±0.49	±0.19	±0.07
Untreated walnut		3.90 ^b	64.48°	82.42 ^b	21.31⁵
		±0.49	±1.09	±0.93	±0.63
Treated Walnut with	6 KGy	4.10 ^b ±0.60	67.37 ^{ab} ±0.56	98.01ª ±1.57	29.06 ^a ±0.37
	10 KGy	4.01 ^b ±1.05	68.84 ^a ±0.86	99.98 ^a ±0.46	26.07 ^a ±3.92

*Values are expressed as means ±SD.

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تأثير الفستق وعين الجمل المشع والمدعم للوجبات علي بعض الخصائص التي البيوكيميائيه في الفئران

آيات لطفي خضر سعودي (... هنيه فتحي غريب النيلي (أشرف عبدالعزيز عبد المجيد ٢

اقسم بحوث تشعيع الأغذية – المركز القومى لبحوث وتكنولوجيا الاشعاع هيئة الطاقة الذرية – القاهرة – جمهورية مصر العربية تقسم التغذية وعلوم الأطعمة – كلية الاقتصاد المنزلى - جامعة حلوان– القاهرة جمهورية مصر العربية

المستخلص العربى

تهدف هذه الدراسة إلى تقييم تأثير الإشعاع عند الجرعات ٦ كيلوجري و ١٠ كيلوجراي في المعاملات الكيميائية الحيوية للمكسرات بما في ذلك الفستق وعين الجملاستخدم في هذه الدراسه ٢٠ جرذان ألبينو ، وتنقسم هذه الجرذان إلى ٢ مجموعة على النحو التالي: المجموعة (١): تتغذى على النظام الغذائي الاساسي ، كمجموعة ضابطة. المجموعة (٢): تتغذى على نظام غذائي أساسي يحتوي على فستق غير مشعع. المجموعات (٣ و ٤): تتغذى على الوجبات الغذائية التي تحتوي على عين الجمل المشعع بـ ٦ و ١ KGy ، على التوالي. المجموعة (٥): تتغذى على النظام الغذائي الاساسي يحتوي على عين الجمل غير المشعع. المجموعات (٣ و ٢). المجموعة (٥): الغذائية النظام الغذائي الاساسي يحتوي على عين الجمل المشعع بـ ٦ و ١ ماميع. المجموعات (٦ و ٢) أساسي على التوالي. المجموعة (٥): المدم النظام الغذائي الاساسي يحتوي على عين الجمل غير المشعع بـ ٦ و ١ محموعات (٦ و ٢). المجموعة (٢)

أظهر التحليل الإحصائي للفستق المشعع و عين الجمل المشعع عند الجرعة ٦ و ١٠ كيلوجراي عند بدء التجربه و بعد التخزين لمدة ٦ أشهر على تحليله الكيميائي عدم وجود فرق كبير في البروتين والرطوبة والرماد والدهون. ولم تظهر التحاليل البيوكيميائيه وجود تأثير معنوي على وزن الأعضاء، وكولسترول الليبوبروتينات عاليه الكثافة (HDL-C) و كولسترول الليبوبروتينات منخفضه الكثافة (LDL-C) ، هيموجلوبين الدم ، الكوليسترول الكلي ، الجليسريدات الثلاثية، الجلوكوز ، وألانين أمين ترانسفيراز (ALT) والاسبرتات امين ترانسفيراز (AST) ، سوبر

اوكسيد ديسموتيز(SOD) ، جلوتاثيون بيروكسيديزوكاتالز الكبد في مجموعات عينات الفستق وعين الجمل غير المشعع أو المشعع ، عند مقارنتها مع المجموعه الضابطه التي تتغذى على النظام الغذائي الاساسي ، في حين أن ألانين الأميني ترانسفيراز (ALT) أظهر زيادة في الفستق غير المعالج وكانت هناك زيادة كبيرة في إنزيمات مضادات الأكسدة في جميع العينات.