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# Evaluation of the Possible Immunomodulatory and Anti-Inflammatory Effects of Gamma-Irradiated Basil, Ocimum basilicum Against Arsenic Toxicity in Rats

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### **ABSTRACT**

In view of the widespread incidence of arsenic poisoning around the world, it was necessary to study this phenomenon and analyze it to find out how to treat it through the application of alternative medicine. Gamma irradiation as a phytosanitary treatment of food and herbal materials is increasingly recognized throughout the world by improving their hygienic quality. The aim of this study was to evaluate the therapeutic effect of raw or irradiated basil on rats exposed to arsenic toxicity. Basil was irradiated by gamma rays at dose 10 KGy. Forty-eight adult Wistar albino rats were divided into six groups as follows group-1: Control group, group-2: received 400 mg/ kg of aqueous extract of basil, group-3: received 400 mg/ kg of aqueous extract of irradiated basil, group-4: received 10 mg/kg of sodium arsenate solution, group- 5: received 10 mg/kg of sodium arsenate solution and 400mg/kg of aqueous extract of basil, and group- 6: received 10 mg/kg of sodium arsenate solution and 400mg/ kg of aqueous extract of irradiated basil. At the end of the experiment (5 weeks), the rats were sacrificed, blood and brain tissue samples were subjected to estimate of the following: CBC, inflammatory markers (CRP, TNF- $\alpha$ , and IL6), immunoglobulin markers (IgA, IgG, IgM), and the levels of oxidative stress and antioxidant in brain tissue (MDA, CAT, SOD, and GSH). The results showed a rise in the antioxidant of basil after the irradiation process. The arsenic caused a significant decreased in levels of HB, RBC, LYM, NEU and PLT and increased levels of WBC and reticulocyte count as compared to the control group. Also, the rats exposed to arsenic showed a significant increase in inflammatory markers in serum, and oxidative stress in brain accompanied with significant decreased in immunoglobulin markers in serum and antioxidant in the brain. In contrast, the administration of raw basil or irradiated basil extract with arsenic was a helpful factor in alleviation of these side effects. In conclusion, our findings showed that the irradiation process enhanced antioxidants in basil plants which attenuated arsenic toxicity.

KEY WORDS: BASIL; GAMMA IRRADIATION; ARSENIC; HEMATOLOGY; INFLAMMATION; IMMUNE SYSTEM; BRAIN.

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#### INTRODUCTION

Arsenic is a toxic substance that occurs naturally and is found in water, rock soil and many foods (Duker et al., 2018). Arsenic toxicity affects millions of people in different parts of the world through the ingestion of arsenic-contaminated drinking water and food (Huq et al., 2006; Flanagan et al., 2012). In recent years, arsenic toxicity has created significant public concern (Duker et al., 2018).It has no taste or smell, which makes it particularly hazardous, so one can be exposed to it without knowing it (ATSDR, 2017). The increasing presence of variable amounts of arsenic in the environment presents major risks, as exposure through inhalation, ingestion and dermal contact can cause various adverse effects on health systems (Vimercati et al., 2017, Chiocchetti et al., 2018 Mochizuki et al., 2019, Tutkun et al., 2019), (Zhao et al., 2019).

The National Center for CAM (complementary and alternative medicine) was reported as a category of healthcare and medical systems (NCCIH, 2018). A systemic review conducted by Eardley et al. (2012) found that, for many reasons, people are using CAM, including its availability, perceived health, and disease prevention. As one of the main forms of CAM is herbal medicines, which uses parts of a plant or whole plants to avoid and cure diseases (Bent, 2008; Pan et al., 2014). The herbal medicinal products are characterized by the presence of complex chemical compounds responsible for the pharmacological activities that contribute to health benefits (Bent, 2008). Basil plants considered from herbal medicine that has eminence value as a cure for various diseases (Patel et al., 2018).

Basil (Ocimum basilicum L.) is an herbaceous and aromatic plant worldwide cultivated which belongs to the Lamiaceae family (Jakovljevic et al., 2016). It has several therapeutic properties including antioxidant, anti-aging, anticancer, antiviral, antimicrobial, antigenotoxic, and anti-inflammatory(Sakr and Al-Amoudi, 2012; Shirazi et al., 2014). Studies have shown many pharmacological effects for basil in several diseases such as liver fibrosis(Alomar and Al-Attar, 2019), diabetes mellitus (Widjaja and Rusdiana, 2019), asthma (Eftekhar et al., 2019), anemia (Zangeneh et al., 2019), and cerebral injury (Singh, Krishan and Shri, 2018). Based on the pharmacological and therapeutic properties, basil has played an important role in both traditional pharmaceutical products and contemporary pharmacological and clinical science(Shirazi et al., 2014). But there may be concerns about the use of herbs, which is that herbs are susceptible often to contamination during processing or storage by micro-organisms and insect pests. This leads to shortens their shelf life and triggering serious illness in some cases, particularly if the herbs contaminated with Salmonella and Staphylococcus aureus (Chatterjee et al., 2016). Thus, herbs should be subjected to sterilization or microbial treatment before use.

There are various techniques for the decontamination of medicinal plants, including irradiation (Garg and Gupta, 2016). This is a physical process that applies high-energy ionizing radiation to the plants in order to enhance their safety and shelf-life (Byun et al., 1999; SádECká, 2007; Alothman, Bhat and Karim, 2009). In particular, gamma irradiation seemed to be the best way to decontaminate herbs from microbes without triggering quality changes(Lee et al., 2005). As the absorbed energy of radiation can break the bonds of DNA molecules in microorganisms present in the product and inactivates certain enzymes, this greatly reduced its damaging impact on products. Not only microorganisms are destroyed, but even gametes, insects and parasites are prevented from reproducing, resulting in various preservative effects, (Farkas, 2006).

Moreover, irradiation serves as a safe technique in food processing supported by many internationally recognized organizations, where joint (Food and Agriculture Organization/ International Atomic Energy Agency/ World Health Organization)Expert Committee on the Wholesomeness of Irradiation of Food has ruled that food subject to low irradiation dosage (up to 10 kGy) is safe and not need any testing of toxicology (Wen et al., 2006). The present study was therefore designed to evaluate the possible immunomodulatory and anti-inflammatory effects of raw and irradiated basil (*Ocimum basilicum*) against arsenic toxicity in rats.

#### MATERIAL AND METHODS

Chemicals and kits: Sodium arsenate was obtained from British Drug Houses (BDH) chemical company, England, UK. Diethyl ether was obtained from Sigma-Aldrich company, Louis, USA. Kits for assays of immunoglobulin G (IgG), immunoglobulin M (IgM), immunoglobulin A (IgA), tumor necrosis factor-alpha (TNFα), interleukin (IL-6), C- reactive protein (CRP), superoxide dismutase (SOD), glutathione (GSH), catalase (CAT) and malondialdehyde (MDA) were obtained from Abcam Chemical Company, Cambridge, UK. kits for assay Total capacity antioxidant were obtained from Cell Biolabs company, California, USA. Kits assay protein concentration in tissues (Pierce TM BCA Protein Kit) was obtained from Thermo Fisher Scientific Company, Waltham, US.

**Plant material and preparation of extract:** The basil (*Ocimum basilicum*) leaves were purchased from the local traditional market in Jeddah, Saudi Arabia. The water extracts of raw or irradiated dried basil leaves were prepared according to the method described by Ghazwani et al. (2020). The total antioxidant capacity was measured in basil extracts using the Total Capacity Assay kit with CAT# STA-360.

**Gamma Irradiation treatment:** The samples of dry basil powder were irradiated with 10 KGy of gamma rays after the leaves were transferred into polyethylene bags, using a Cobalt-60 source at a dose rate of 4.75 KGy/h at the National Centre for Radiation Research and Technology (NCRRT), Nasr City, Cairo, Egypt.

**Experimental animals and Design:** The study was carried out using adult female Wistar rats weighing (150-200g); they were obtained from faculty of pharmacy at King Abdulaziz University. The animals were housed in cages and received normal rat chow and tap water in a constant environment (room temperature  $28 \pm 2^{\circ}$ C, room humidity 60+5%) in a 12 h light and 12 h dark cycle. Rats were kept under supervision for two weeks before the experiments started and during all stages of the whole experiment. Animal's procedures were performed in accordance with the Ethics Committee of the King Fahad Medical Research Center and in accordance with the recommendations for the proper care and use of laboratory animals. In the experiment, 48 rats were divided into six groups each of 8 rats as follows:

**Group 1:** normal control rats were given only distilled water. Group 2: rats received 400 mg/ kg of aqueous extract of raw basil (Ezeani et al., 2017). Group 3: rats received 400 mg/kg of aqueous extract of irradiated basil (Ezeani et al., 2017). Group 4: rats received 10 mg/kg of sodium arsenate solution (Firdaus et al., 2018). Group 5: rats received 10 mg/kg of sodium arsenate solution and 400mg/kg of aqueous extract of basil. Group 6: rats received 10 mg/kg of sodium arsenate solution and 400mg/ kg of aqueous extract of irradiated basil. All doses were given through an oral gastric tube daily for five weeks. At the end of the experiment, rats fasted overnight for scarification.

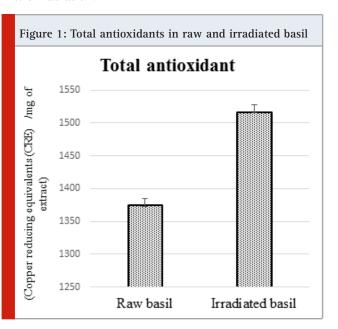
Blood samples were withdrawn by a heparinized capillary tube from the retro-orbital plexus of each rat under anesthesia with diethyl ether, it is put into two tubes, one is ethylenediamine tetra-acetic acid (EDTA) tube and the other is a serum-separating tube. The EDTA tube immediately turned to lab analysis while serumseparating tube centrifuged at 3000 rpm for 15 min to separate serum and then stored at -40° C until the biochemical analysis is done. Directly after preparing the blood sample, rats sacrificed and the brain was kept in ice for homogenate preparation.

**Biochemical analysis:** The biochemical analysis for complete blood count (CBC) was done in Kingdoms Labs. The blood serum IgG, IgM and IgA, TNFα, IL-6 and CRP were identified using the kits with CAT# ab189578, ab157735, ab157738, ab46070, ab119548, and ab 108827, respectively. For tissue analysis, the brain was homogenized and estimated each of SOD, GSH, CAT and MDA according to the kits with CAT# ab65354, ab138881, ab83464, and ab118970, respectively, while the protein concentration estimated in the brain was used Kits with CAT# 23225.

**Statistical analysis:** The data of each group were analyzed using MegaStat 9.4 (add-in for Excel). The data were expressed as arithmetic mean and standard deviation of the mean (SD). Differences between groups were analyzed for parametric parameters using oneway variance analysis (ANOVA), the least significant difference equation (LSD). A P value below or equal to 0.05 was considered significant.

#### RESULTS AND DISCUSSION

Antioxidant in irradiated plant leaves: The effect of irradiation on antioxidants is showed in figure 1. Where leaves treated with gamma irradiation (10kGy) showed a change, a 10.3% increase in the content of total antioxidants compared to raw leaves not treated with radiation.



**Complete blood count (CBC):** The effects of aqueous extracts of raw basil or irradiated basil for 5 weeks on the levels of hemoglobin (HB), red blood cells(RBC), hematocrit (HCT), platelets (PLT), White blood cells (WBC), WBC differential and reticulocyte count in rats exposed to arsenic are present in Tables 1 and 2. No effect of raw or irradiated basil administration in normal animals on all previous parameters was noted. Rats, which were exposed to arsenic showed a significant decrease in levels of HB, RBC, PLT, neutrophils (NEU), and lymphocytes (LYM) compared to the control group. While there was a marked increase in the levels of HCT, WBC, monocytes (MON) and reticulocyte count compared to the control group. Administration of raw and irradiated basil along with arsenic significantly reduced arsenic toxicity effect by the improvement levels of these parameters.

**Antibodies:** The extracts of raw and irradiated basil had distinct effects on antibodies in rats exposed to arsenic that are showed in Table 3. The treatment with raw or irradiated basil to normal animals not producing an effect on levels of serum IgG, IgM and IgA. The arsenic group showed a significant decrease in IgG, IgM and IgA levels compared to the control group. The treatment with basil extract as raw or irradiated along with arsenic dose was showed a significant increase in those antibodies' levels. compared with the group of rats given only arsenic.

**Inflammatory markers:** The level of markers of inflammation in rats exposed to arsenic which received raw or irradiated basil as a treatment are showed in Table 4. In normal animals, the administration of raw

or irradiated basil has not had an effect on the levels of serum TNF, IL-6 and CPR. The rats exposed to arsenic showed a significant increase in levels of TNF, IL-6, and CPR as compared to the control group. Giving raw or irradiated basil extract significantly reduced inflammatory markers induced by arsenic.

Table 1. Effect of basil extracts on hematological parameters of rats exposed to arsenic				
Test				
Group	HB (g/dl)	RBC (10^6/mL)	HCT (%)	PLT (10^3/mL)
С	12.661 ±1.258	4.450± 0.575	29.635± 1.870	263.13± 52.58
В	12.725 ± 0.902 bbb	4.570± 0.491 bbb	29.286± 0.516 bbb	274.50± 19.17 bbb
IB	12.810 ± 0.756 bbb	4.568 ± 0.487 bbb	29.518± 0.908 bbb	277.25± 13.63 bbb
A	8.748 ± 0.590 aaa	3.100 ± 0.524 aaa	39.058± 0.655 aaa	143.25± 7.96 aaa
B + A	10.918 ± 0.71 aaa bbb	4.066 ± 0.463 bbb	30.978± 1.748 bbb	188.38± 10.89 aaa bb
IB + A	11.249 ± 0.290 aabbb	4.151± 0.164 bbb	32.330± 1.863 aaa bbb	177.88± 10.47 aaa bb

Values are the mean of 8 observation  $\pm$  SD, Significant different from C value at P <  $0.05^a$ ,  $0.01^{aa}$ ,  $0.001^{aaa}$ , Significant different from A at value at P <  $0.05^b$ ,  $0.01^{bb}$ ,  $0.001^{bbb}$ 

Table 2. E	Table 2. Effect of basil extracts on hematological parameters of rats exposed to arsenic					
Test						
Group	WBC WBC differential (10^ 3 / mL)			Reticulocyte Count (%)		
	(-1 - 7)	NEU (%)	LYM (%)	MON (%)		
С	5.661 ± 0.508	62.403 ±1.869	28.16 ±2.658	4.113±0.631	1.413± 0.416	
В	5.255 ± 0.532	63.388 ±1.886	27.61±4.022	4.250±0.460	1.100± 0.185	
	bbb	bbb	bb	bbb	bbb	
IB	5.134 ± 0.604	64.538 ±1.577	29.46 ±4.530	4.064±0.261	1.263± 0.424	
	bbb	bbb	bbb	bbb	bbb	
A	9.761± 0.893	38.051 ±2.361	20.00 ±2.624	7.150±0.493	2.350± 0.540	
	aaa	aaa	aaa	aaa	aaa	
B + A	6.160± 0.825	54.175 ±5.389	24.95±5.687	5.163±0.460	1.011±0.110	
	bbb	aaa	b	aaa	bbb	
		bbb		bbb		
IB + A	5.978± 0.610	51.850±5.011	25.73±5.035	5.113±0.500	1.163±0.518	
	bbb	aaa	b	bbb	bbb	
				aaa	bbb	
	Values are the mean of 8 observation $\pm$ SD, Significant different from C value at P < 0.05a, 0.01aa, 0.001aaa, Significant different from A at value at P < 0.05b, 0.01bb, 0.001bb					

**Antioxidants and oxidative damage:** The results of determining the content of SOD, GSH, CAT and MDA in rats given arsenic along with raw basil or irradiated

basil were shown in Table 5.The normal rats treated with irradiated basil showed an increase in the level of SOD than rats treated with raw basil. The level of GSH was

showed an increase in normal rats of both of raw basil group and irradiated basil group as compared to the control group. Arsenic-exposed rats showed a significant increase in MDA levels accompanied by a significant decrease in SOD, GSH and CAT compared to a control group. The management of raw or irradiated basil along with arsenic alleviated the effects of arsenic and resulted in a significantly decreased MDA with significantly increased SOD, GSH, CAT.

Table 3. Effect of basil extracts on immunoglobulins of rats exposed to arsenic				
Test	IgG (µg/ml)	IgM (µg/ml)	IgA (μg/ml)	
Group				
С	517.16 ± 35.137	0.628 ± 0.084	90.180 ± 6.688	
В	522.490 ± 41.527	0.640 ± 0.053	93.236 ± 7.697	
	bbb	bbb	bbb	
IB	514.663 ± 32.665	0.655 ± 0.064	94.614 ± 5.087	
	bbb	bbb	bbb	
A	202.500 ± 40.818	0.226 ± 0.024	45.779 ± 5.816	
	aaa	aaa	aaa	
B + A	298.911 ± 32.731	0.389 ± 0.015	72.754 ± 4.942	
	aaa	aaa	aaa	
	bbb	bbb	bbb	
IB + A	287.915 ± 35.128	0.408 ± 0.021	76.745 ± 4.229	
	aaa	aaa	aaa	
	bbb	bbb	bbb	
-				

Values are the mean of 8 observation  $\pm$  SD, Significant different from C value at P < 0.05a, 0.01aa, 0.001aaa, Significant different from A at value at  $P < 0.05^b$ , 0.01bb, 0.001bbb

Table 4. Effect of basil extracts on the inflammation markers o rats exposed to arsenic				
Test	TNFα (pg/ml)	IL-6 (pg/ml)	CRP (ng/ml)	
Group	(PB))	(58)	(118/1111)	
С	17.829 ± 1.221	21.048 ± 2.134	0.134 ± 0.039	
В	16.144 ± 1.428	20.753 ± 1.962	0.134± 0.034	
	bbb	bbb	bbb	
IB	16.123 ± 1.216	20.456 ± 2.940	0.140± 0.018	
	bbb	bbb	bbb	
A	40.043 ± 2.453	35.334 ± 3.973	0.990± 0.076	
	aaa	aaa	aaa	
B + A	27.656 ± 2.173	27.413 ± 3.842	0.660± 0.093	
	aaabbb	aaabbb	aaabbb	
IB + A	28.935 ± 2.634	26.876 ± 1.563	0.490± 0.033	
	aaabbb	aaabbb	aaabbb	

Values are the mean of 8 observation  $\pm$  SD, Significant different from C value at P < 0.05a, 0.01aa, 0.001aaa, Significant different from A at value at P < 0.05b, 0.01bb, 0.001bbb

The irradiation treatment can increase the content of some phytochemicals and the plant's antioxidant activity, thereby increased biological value (Zevallos-Concha et al., 2016; Pereira et al., 2018). The results indicated that the extract of irradiated basil showed a highly significant

increase in total antioxidants as a comparison to raw basil. Similar observations were reported in previous studies on basil and some other plants (Khawory et al., 2020; Osman et al., 2020; Rady et al., 2020).

Table 5. Effect of basil extracts on the antioxidants and oxidative damage in the brain of rats exposed to arsenic

Test	SOD (Inhibition rate %/	GSH (µM/gram tissue)	CAT (µM/mg of protein)	MDA (µM/mg of mg of protein)
Group	protein)	(1)	(12.14)	g p,
С	105.406 ±7.342	90.743 ± 7.044	0.020 ± 0.003	2.496± 0.451
В	107.548 ±9.384	$96.709 \pm 3.420$	$0.019 \pm 0.004$	2.400± 0.422
	bbb	abbb	bbb	bbb
IB	114.655 ±8.165	95.425 ± 4.426	0.021 ± 0.004	2.441± 0.433
	abbb	abbb	bbb	bbb
A	77.977 ±10.370	60.248 ± 2.657	0.009 ± 0.000	6.005± 0.126
	aaa	aaa	aaa	aaa
B + A	82.049 ±8.953	84.259 ± 4.487	0.012 ± 0.001	3.479± 0.610
	aaabbb	aabbb	aaa	aaabbb
IB + A	89.249 ±9.381	82.533 ± 3.153	0.014 ± 0.003	3.165± 0.484
	aaabbb	aabbb	aabb	aabbb

Values are the mean of 8 observation  $\pm$  SD, Significant different from C value at P < 0.05<sup>a</sup>, 0.01<sup>aa</sup>, 0.001<sup>aaa</sup>, Significant different from A at value at P < 0.05<sup>b</sup>, 0.01<sup>bb</sup>, 0.001<sup>bbb</sup>

Basil contains phenolic compounds and flavonoids (Bahcesular et al., 2020), that are considered as natural antioxidants. These biomolecules exhibit their activity through various mechanisms, including inhibiting enzymes that inducing free radical produce, increasing endogenous antioxidants, removing free radicals, and inducing the expression of the numerous genes responsible for enzyme synthesis that inhibit oxidative stress (Primiano, Sutter and Kensler, 1997). Ghazwani, Osman and Balamash (2020) have recently reported that the Fourier-transform infrared (FTIR) analysis indicated to increase the content of phenolic acids and flavonoids in basil leaves after treated with 10 kGy of gamma-ray. Moreover, Maraei, Khaled and Elsawy (2017)reported that the gamma irradiation-induced the biosynthesis of certain phenolic compounds.

Also, it seems that gamma irradiation with 10 kGy might stimulate some chemical reactions in basil, which perhaps increase in phenolic content by the breakdown of covalence bonds among phenolic components and, free phenolic components with low molecular weight are increasing (Jamshidi, Barzegar and Sahari, 2014). A complete blood count test is a blood test used to assess general health and detect a range of disorders in hematological parameters. A complete blood count test measures many blood components and features, including red blood cells that carry oxygen, white blood cells that fight infection, hemoglobin that oxygen-carrying protein in red blood cells, hematocrit that indicate to the ratio of red blood cells to the liquid or plasma component of the blood and platelets that help blood clot, and that

any change whether an abnormal rise or decrease in the census, indicate the incidence of diseases or disorders requiring medical procedures (Clinic, 2018).

In our study, it has been observed that the levels of HB, RBC, PLT, NEU, and LYM are decreased significantly with a marked increase in the HCT, WBC, MON and reticulocyte count in rats exposed to arsenic compared to the control group. The results are consistent with some studies (Kajiguchi et al., 2005; Bhattacharya and Haldar, 2012; Sumedha and Miltonprabu, 2013; Lemaire et al., 2015; Ghosh et al., 2017; Su et al., 2018). This effect of Arsenic exposure on the hematopoietic system may be attributed to the mechanisms of arsenic toxicity which may induce hemolysis and erythrophagocytosis through increased oxidation of sulfhydryl groups in hemoglobin and decreased oxygen intake by cells as a result of decreased intracellular glutathione, which decreases the lifespan of erythrocytes(Abdul et al., 2015). Moreover, arsenic exposure can also cause a range of changes, such as increasing ceramide formation, membrane disintegration, cytosolic calcium levels, besides decreasing in adenosine triphosphate (ATP) levels, cell membrane integrity affecting erythrocyte lifespan (Abdul et al., 2015).

Regarded the change in platelet count, this confirmed that arsenic inhibited platelet differentiation within the hematopoietic system of bone marrow, leading to reduced platelet production Wu et al., 2014. The white blood cell level was increased in arsenic feed groups be due to the impact of arsenic which induced apoptotic

effect on plasma cells as noted by Rousselot et al. (2004). WBC may generally be divided into five classes, based on their function, morphology and origin: LYM, MON and NEU(Villa et al., 2003). The changes in the LYM and NEU populations present in this study may be due to arsenic caused immune inhibition in rats (Taheri et al., 2016).

Our findings demonstrated that extracts of basil caused improved the disorders that occur in CBC. This is in agreement with the results of previous researches (Ofem et al., 2012; Zangeneh et al., 2019). This effect may due to basil which contains a proportion of iron (Nworgu, Yekini and Oduola, 2013), that contributes to improving the level of HB in the blood and has the ability to stimulate production and increase of RBC to treat deficiency caused by arsenic. Furthermore, in normal, the lack of oxygen in the local tissue appears to lead to the production of glycoprotein known as erythropoietin, which induces increased erythrocyte output(Bowman and Rand, 1980).

Basil leaves extract is very likely to contain erythropoietinlike agents that are responsible for increased erythrocyte production (Ofem, Ani and Eno, 2012). Saha et al. (2012) reported that secondary metabolites of basil, consisting of important elements include essential oil geraniol, a monoterpene and citral, play a role as the modulator in hematological abnormalities (Ofem et al., 2012). Moreover, in results about the increased lymphocyte count after basil administration, it has been reported that Ocimum basilicum modulates the cell-mediated as well as a humoral immune response that could be due to the presence of flavonoids and terpenoids (Mediratta et al., 2002).

Antibodies also are known as immunoglobulins, are substances made by the body's immune system in response to foreign substances. Antibodies bind to these foreign substances and they can be killed by the immune system. IgG, IgM and IgA from the major types of antibodies. If reduced levels of antibodies produced by the immune system, it leads to more likely to develop repeated infections(Staff, 2019). The results of our study showed that in the arsenic group of rats (10 mg/kg BW) a highly significant reduction was observed in IgG, IgA and IgM levels compared to the control group. The results are consistent with some studies (Institoris et al., 2001; Sankar et al., 2013). The effect on immunoglobulin levels associated with arsenic exposure can attribute to the arsenic disrupts glucocorticoid regulation, responsible for immune function, (Kaltreider et al., 2001).

Furthermore, the apoptosis caused by arsenic may result in decreased immune responses(Harrison and McCoy, 2001). The administration of water extract from raw or irradiated basil extract demonstrated a protective effect against arsenic toxicity in rats, by increasing antibodies that decreased as a result of arsenic poisoning. These results are in line with the findings of Mohammed, Kadhim and Taher (2017) and Jahejo et al. (2019). Jeba, Vaidyanathan and Rameshkumar (2011)showed that aqueous extract of basil stimulated the antibody

production in rats. The flavonoids present in the basil leaves are mainly responsible for the immunomodulatory effect (Ravindran, 2017).

Inflammation is a biological response of the immune system which can be induced by damaged cells, toxic compounds or pathogens (Medzhitov, 2010). It is part of the body's defense mechanism. one of the major aims of inflammation is to bring immune cells to the area of concern as well as to inactivate or destroy any injurious stimuli and to also begin the repair(Ferrero-Miliani et al., 2007; Medzhitov, 2010). The inflammation response is caused by specific immune factors released from the damaged cells. Where, the damaged cells release cytokines, including interleukins, such as IL-6, IL-8, and tumor necrosis factor-α, that are responsible for communication between white blood cells. Interleukins also stimulate the production and release of CRP from the liver; an important component of the innate immune system (Sinclair et al., 2012).

Usually, molecular and cellular activities and interactions efficiently alleviate inevitable infection or damage, during acute inflammatory responses. This effect helps restore homeostasis in the tissue and overcome the acute inflammation. Uncontrolled acute inflammation can become chronic, however, and can lead to a number of chronic inflammatory diseases (Zhou et al., 2016). The elevated levels of inflammatory markers are expected to be associated with toxic metals exposure. Results of this study showed that serum IL-6, TNF- $\alpha$  and CRP level in rats exposed to arsenic was highly significantly elevated. Our findings agree with the results of the previous study on the association between arsenic and ability to cause chronic inflammation by demonstrating the increased pro-inflammatory mediators like TNF-α, IL-6 and CRP in the arsenic exposed group in comparison to the control group(Prasad and Sinha, 2017). Inflammation considered to be one of the main arsenic toxicity mechanisms that can be correlated with increasing cellular damages oxidative stress and lipid peroxidation (Bhadauria and Flora, 2007).

In this work also, it was demonstrated that oral treatment with basil extracts diminished inflammations in rats exposed to arsenic. These results in agreement with Aye et al. (2019) and Takeuchi et al. (2020) who found that basil has anti-inflammatory effects. Rodrigues et al. (2016) reported that the basil essential oil was effective in reducing inflammations (acute or chronic) by induced inhibiting of the inflammatory mediator receptors and the migration of cells to stimulus locations. This may be attributed to the contains of basil of rosmarinic acid (Kwon et al., 2019), where this compound has been related to anti-inflammatory activities (Luo et al., 2020).

Antioxidants are considered the enzymes of the body's protection, are able to stabilize free radicals before attacking components of the cell. They work to diminishing free radicals through reducing their energy or donate electrons to them, thus making it stable(Krishnamurthy and Wadhwani, 2012). While,

"oxidative stress, defined as a disturbance in the balance between the production of reactive oxygen species (free radicals) and antioxidant defenses" (Betteridge, 2000). In the present study, observed a very highly significant decrease in GSH, SOD and CAT accompanied by increased MDA of the rat's brain in the arsenic group as compared to the normal control group. This result is similar to the study of Sun et al. (2018) that reported that arsenic caused significantly decreased GSH, SOD and CAT with increased MDA content in the brain tissues of chickens. This may be attributed to the toxic effect of arsenic that may induce oxidative stress by interacting with antioxidants, resulting in the accumulation of free radicals in cells (Bonetto, Villaamil Lepori and Puntarulo, 2014).

In contrast, basil giving a positive effect on the brain by improving the levels of antioxidant antioxidants, and fat oxidation (MDA). These results were in line with recent studies showed that the water extract of gammairradiated basil contributes to improving the oxidative stress induced by arsenic exposure in rats (Ghazwani, Osman and Balamash, 2020; Osman, Ghazwani and Balamash, 2020). Also agree with the results of Khodabakhshi et al. (2017), who proved that the increased level of MDA in the mice brain tissue following seizures was prevented by basil extract. Moreover, Khaki (2016) demonstrated that the basil extract protects brain cells from the harmful effects by regulating the antioxidant enzymes in the serum. This saves the neurons from irreversible cell injury. The antioxidant effect is due primarily to phenolic elements, such as, phenolic acids and flavonoids, which have redox properties and ability to neutralize free radicals (Shahidi et al., 1992).

#### CONCLUSION

In this study, we demonstrated that gamma radiations have the ability to increases antioxidants contents in basil leaves. Moreover, the water extracts obtained from *Ocimum basilicum* can be successful in diminished arsenic toxic effect through the improvement of the Blood homeostasis of CBC and immunoglobins level, reduced inflammatory markers and, enhancement ability of antioxidants to be overcome oxidative stress in the brain.

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**Ethical Clearance Statement:** The Current Research Work Was Ethically Approved by the Institutional Review Board (IRB) of Ethics Committee of the King Fahad Medical Research Center and in accordance with the recommendations for the proper care and use of laboratory animals.

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