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Antioxidant properties of *Pelargonium graveolens* L'Her essential oil on the reproductive damage induced by deltamethrin in mice as compared to alpha-tocopherol

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Abstract

Background: Exposure to the pyrethroid pesticide deltamethrin has been demonstrated to exert a wide range of effects on non-targeted organisms. The beneficial effects of geranium essential oil (EO) as an antioxidant has been assessed in deltamethrin (DL) orally administered mice by studying whether the reprotoxicity caused by deltamethrin can be effectively combated with the geranium oil and the effects were compared to vitamin E, as the standard reference drug.

Result: Sixty male albino mice were divided into six equal groups: a control group, a group of mice was given deltamethrin (5 mg/kg b.w.), two groups were administered deltamethrin after having given geranium essential oil (67 mg/kg b.w.) or vitamin E (Vit E) (100 mg/kg b.w.), and two groups received only EO of geranium or Vit E. When compared to control, a dose of deltamethrin 5 mg/kg/day causes a decrease in the epididymal sperm count motility and viability and an increase in the number of abnormal morphology in spermatozoa. DL-exposed mice showed a significant increase of lipid peroxidation (LPP) in the testis compared to control animals.

Conclusion: Essential oil of geranium prevented testicular oxidative damage explored by reduced LPP and improved total sperm motility, viability and morphology in mice spermatozoa. Our study showed a positive influence of geranium essential oil in the animal male reproductive system similar than that of Vit E.

Keywords: Deltamethrin, Oxidative damages, Geranium oil, Antioxidants

Background

The increasing release of chemicals into the environment dictates attention to a better understanding of their toxicity in human and ecotoxicological effects. Many studies suggest that environmental contaminants disrupt male reproduction and play an important role in the declination of the quality and quantity of human semen [1]. Several currently used pesticides are known to adversely impair reproductive competence of males under laboratory, field, clinical or occupational setting. Published studies reported that pyrethrinoid can impair fertility, deteriorate semen

quality, and cause testicular degeneration, male reproductive failure and malformations in the fetuses in the rodents following repeated exposure [2,3]. Deltamethrin [(R, S)] is a type-II pyrethroid synthetic insecticide, which has been widely used to control noxious insects in agriculture, forestry and horticulture. However, a number of studies have demonstrated genotoxic and tumorigenic effects of deltamethrin in mammalian and non-mammalian species [4,5]. For many pesticides, induction of oxidative stress is one of the main mechanisms of their action. The chronic exposure to deltamethrin caused also haemolysis, hepatic, renal toxicities [6] and neurodegeneration [7]. The mechanism of such pathological facts may be prompted by the free radical release and the lipid peroxidation that it induces. During the past few years, estimation of free radical

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generation and antioxidants defense has become an important aspect of investigation in mammals for the protection of cells against oxidative damage due to pesticides [8], heavy metals [9] and chemotherapeutic agent toxicities that they generated probably oxygen-reactive species (ROS) which led to oxidative stress.

Rose-scented geranium (*Pelargonium graveolens* L'Hér.) is widely known as one of the medicinal herbs with the highest antioxidant activity. Essential oils are a folk medicine and recently their use has expanded worldwide to include therapy against various kinds of inflammatory diseases. A great number of new drugs discovered in the last few decades are originate from natural sources [10]. Natural products have been increasingly used for the prevention and treatment of various conditions [11,12].

Geranium essential oil has historically been used in the treatment of dysentery, hemorrhoids, inflammation, heavy menstrual flows and even cancer [13]. The French medicinal community currently treats diabetes, diarrhoea, gallbladder problems, gastric ulcers, liver problems, sterility and urinary stones with this oil [14,15]. In Chinese homeopathy, the geranium essential oil is known to open up the liver charka and promote the expulsion of toxins that prohibit the achievement of balance within the body [16]. The aim of the current study was to determine the components of *Pelargonium graveolens* essential oil and to evaluate its protective effects against deltamethrin-induced reprotoxicity in male mice.

Material and methods

Chemicals

DL is a synthetic pyrethroid insecticide ($C_{23}H_{19}Br_{23}NO_3$). CAS chemical name is a (1R,3R)-3-(2,2-dibromovinyl)-2,2-dimethyl-cyclopropane carboxylate de (S)- α -cyano-3-phénoxybenzyle, CAS registry number 52918-63-5. A commercial formulation of DL, named "Decis EC25" (Bayer CropScience) was used in the experiments. All other chemical products used in this study were purchased from Sigma Chemical Co. (St Louis, France).

Plant material

Samples of *P.graveolens* (*geraniaceae*; known as Geranium); were collected from Sfax, South of Tunisia in March 2011. A voucher specimen was deposited at the laboratory of Natural substances chemistry, Faculty of Science of Sfax, Tunisia.

Extraction of essential oil

A bout 200 g of the fresh leaves of *P.graveolens* was completely immersed in 1000 ml distilled water and hydro-distilled for 4 hours in Clevenger-type apparatus until 250 ml of the water-oil layer was obtained. The oily layer obtained on the top of the aqueous distillate was extracted

three times with diethyl ether. The recovered organic solvent was dried over anhydrous sodium sulfate until the last traces of water were removed and evaporated by vacuum distillation at room temperature to yield greenish yellow oil. The obtained essential oil was stored in a dark glass bottle at 4°C until tested and analyzed. Yield based on dried weight of the sample was calculated.

Gas chromatography-mass spectrometry (GC-MS)

The analysis of the essential oil was performed on a GC-MS HP model 5975B inert MSD (Agilent Technologies, J&W Scientific Products, Palo Alto, CA, USA), equipped with an Agilent Technologies capillary DB-5MS column (30 m length; 0.25 mm i.d.; 0.25 μ m film thickness), and coupled to a mass selective detector (MSD5975B, ionization voltage 70 eV; all Agilent, Santa Clara, CA). The carrier gas was He and was used at 1 ml/min flow rate. The oven temperature program was as follows: 1 min at 100°C ramped from 100 to 260°C at 4°C min⁻¹ and 10 min at 260°C. The chromatograph was equipped with a split/splitless injector used in the split mode. The split ratio was 1:100. Identification of components was assigned by matching their mass spectra with Wiley and NIST library data, standards of the main components and comparing their Kovats Retention Indices (KRI) with reference libraries and from the literature [17,18]. The component concentration was obtained using semi-quantification by peak area integration from GC peaks and by applying the correction factors.

Animals care

Adult, healthy and virgin Swiss Albino male mice weighing 27 ± 3.0 g were obtained from the Centre of Veterinary Research (Sfax, Tunisia). Animals were examined for health status and acclimated to the laboratory environment for 1 week *prior* to use. All animals were housed in stainless steel cages and maintained on a 12 h light /dark cycle. The animal room was designed to maintain temperature at $23 \pm 2^\circ\text{C}$ and relative humidity at approximately 50%. Food and water were provided *ad libitum*.

Experimental design

The animals were randomly divided into six groups of ten mice each and they were treated as follows: group 1: control mice received 2.5% diluted DMSO by gavages (C), and served as negative control; group 2: received DL (5 mg/kg b.w./day ($\approx \frac{1}{4}$ DL50 for mouse)); group 3: received DL(5 mg/kg b.w./day) and essential oil of *P. graveolens* at (67 mg/kg b.w./day) after 120 min of DL administration; group 4: received DL(5 mg/kg body weight/day) and Vitamin E (Vit E) at 100 mg/kg b.w./day after 120 min of DL administration; group 5: received only essential oil of *P.graveolens* (EO) at 67 mg/kg b.w./day; group 6: received only Vitamin E (Vit E) at

Table 1 Chemical composition of *P. graveolens* essential oil

N°	KI ^a	Compound	Composition (%)	TR ^b
1	939	α-pinene	0.54	8.67
2	1031	β-phellandrene	0.10	11.55
3	1040	cis-ocimene	0.08	12.03
4	1098	linalool I	10.42	13.82
5	1127	rose oxide-trans	0.6	14.62
6	1154	menthone	7.34	15.22
7	1189	α-terpineol	2.87	16.05
8	1233	β-citronellol	29.30	17.5
9	1255	geraniol	10.53	18.33
10	1275	citronellyl formate	9.54	18.81
11	1290	Thymol	7.84	19.02
12	1418	Trans-caryophyllene	1.2	20.5
13	1439	aromadendrene	0.91	23.56
14	1480	germacrene-d	3.46	29.98
15	1513	γ-cadinene	1.20	24.91
16	1514	leden	1.63	25.08
17	1524	δ-cadinene	1.25	25.33
18	1530	epizonaren	0.21	25.42
Identified components (%)		89.04%		
Monoterpenes hydrocarbons		0.72		
Oxygenated monoterpenes		68.9		
Sesquiterpenes hydrocarbons		9.88		
Oxygenated sesquiterpenes		-		
Other compounds		9.54		

TR^b : Retention Time (min) KI^a : Retention Index (kovats).

dose 100 mg/kg body weight, and served as positive control for antioxidant treatment. The essential oil and Vitamin E were diluted to 2.5% by 2.5% dimethyl sulfoxide (DMSO) and then we added 12.5 μl/ml of Tween20 [19].

The experimental was continued for 35 days. The proper doses of treatment for each animal were placed into a syringe that was inserted orally with the help of plastic tube directly into the oropharyngeal region. The experimental protocol was performed according to the

European convention for the protection of vertebrate animals used for experimental and other scientific purposes (Council of Europe No123, Strasbourg, 1985) and approved by the ethics committee for research on laboratory animal use of our institution.

Sperm quality

Epididymal sperm were collected by cutting the one epididymis into small pieces in 2 ml of physiological saline at 32°C. Sperm evaluation included cell concentration, viability, progressive motility, and normal morphology according to World Health Organization guidelines [20].

Sperm motility and count

Sperm motility was determined by scoring the number of all progressive sperm and then the non-progressive and the immotile sperm in the same field. On each slide, at least 100 sperm cells were counted. Sperm motility assessment was repeated in a new preparation of the same sample. First, the percentage of sperm cells in each motility group was calculated and then the average in each group for the two assessments of the two slides was assayed according to Kvist and Björndahl [21]. From this solution, 20 μl aliquots were placed on the Neubauer Hemacytometer for counting the number of spermatozoa/epididymis.

Sperm viability

Sperm viability study was assessed using the Eosin/Nigrosin stain. The staining was performed with 1 drop of fresh semen into 2 drops of staining solution on a microscope slide. Using another slide, a smear was made and allowed to dry. Unstained (intact) and red-colored (with damaged membranes) spermatozoa were counted under the microscope using ×1000 objectives and oil immersion. Sperm viability was defined as the percentage of intact cells, as per the procedures described in the WHO Laboratory Manual [22].

Sperm morphology

To evaluate the spermatozoa abnormalities, sperm suspension was stained with Eosin; smears were made on slides, air-dried and made permanent. The spermatozoa were classified according to the Wyrobek and Bruce

Table 2 Effect of different treatments on cell's density, viability, motility and morphology in control group (C) and mice treated with deltamethrin (DL), essential oil of *P. graveolens* (EO) and Vitamin E (Vit E) or their combination

	Control	DL	DL + EO	DL + Vit E	EO	Vit E
Spermatozoa count Per epididymis (×10 ⁶)	5.90 ± 0.55	1.96 ± 0.25***	5.09 ± 0.35***	5.11 ± 0.31***	5.77 ± 0.62***	5.76 ± 0.71***
Sperm parameters Motility (%)	78.44 ± 5.85	39.33 ± 6.82***	76.11 ± 10.01***	66.11 ± 18.6***	75.00 ± 8.87***	88.67 ± 2.74***
Viability (%)	89.33 ± 3.97	72.11 ± 3.14***	83.22 ± 1.92***	83.00 ± 4.61***	85.89 ± 6.13***	89.00 ± 4.27***
Morphology (%)	7.00 ± 1.58	22.00 ± 3.94***	15.53 ± 3.59**	16.88 ± 7.12**	6.25 ± 3.28***	7.88 ± 3.00***

Data are presented as mean ± SD. All groups vs. control group: * P < 0.05; ** P < 0.01; *** P < 0.001. All groups vs. DL group: # P < 0.05; ## P < 0.01; ### P < 0.001.

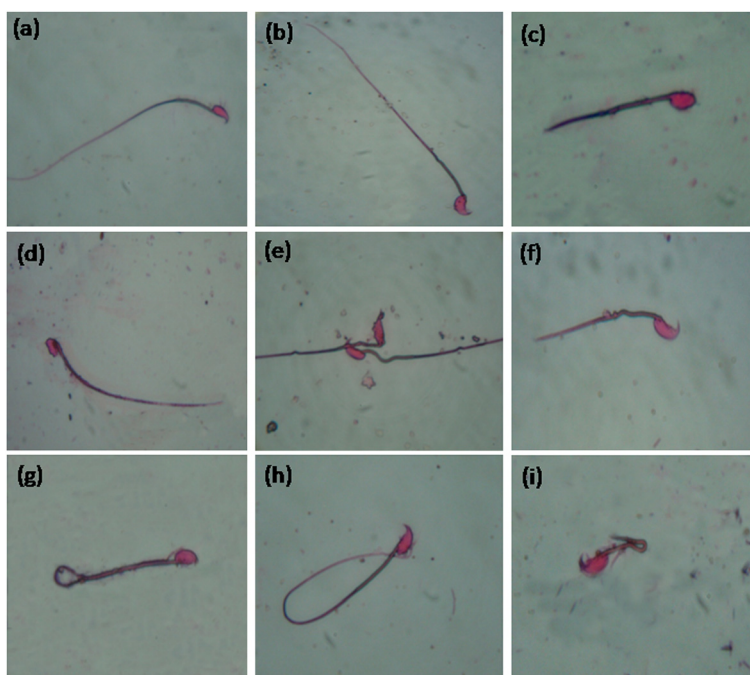


Figure 1 Eosin-stained spermatozoa of mice, (a-b) Control; (c-d) head abnormalities of the spermatozoa; (e-f) mid-piece abnormalities and (g-i) tail abnormalities.

criteria [23]. Sperm cells with normal morphology and cells presenting abnormalities in head, mid-piece and tail were assessed. At least 200 sperm cells were observed in each slide, under 1000× magnification.

Biochemical assays

Measurement of Lipid Peroxidation and Protein Carbonyl Levels in testis

LPP process is determined by the thiobarbituric acid (TBA) method which estimates the malondialdehyde formation (MDA) [24]. Briefly, 0.5 ml of testis homogenate was mixed with 1 ml of trichloroacetic acid solution and centrifuged at 2500 g for 10 min. One millilitre of a solution containing 0.67% thiobarbituric acid (TBA) and 0.5 ml of supernatant were incubated for 15 min at 90°C and cooled. Absorbance of TBA-MDA complex was measured at 532 nm. Lipid peroxidation is expressed as nmoles MDA/g tissue.

Protein oxidation was determined based on the reaction of the carbonyl groups with 2, 4-dinitrophenylhydrazine

(DNPH) to form 2, 4 dinitrophenylhydrazone, as described by Reznick and Packer [25]. Samples were read at 370 nm and carbonyl content was calculated using the molar absorption coefficient for aliphatic hydrazones ($22,000 \text{ M}^{-1} \text{ cm}^{-1}$) and expressed as $\mu\text{mole carbonyl/mg protein}$.

Determination of testis enzymatic antioxidants

Catalase (CAT) was assayed by the decomposition of hydrogen peroxide and converts it to water and molecular oxygen according to the method of Aebi [26]. Decrease in absorbance due to H_2O_2 degradations was monitored at 240 nm for 1 min and the enzyme activity was expressed as $\mu\text{mol H}_2\text{O}_2$ consumed /min/mg protein.

Superoxide dismutase (SOD) activity was estimated according to Beauchamp and Fridovich [27]. The reaction mixture contained 50 mM of testis tissue homogenates in potassium phosphate buffer (pH = 7.4), 0.1 mM methionine, 2 μM riboflavine and 75 μM Nitro Bleu Tetrazolium

Table 3 Results of sperm morphology assessment in control group (C) and mice treated with deltamethrin (DL), essential oil of *P. graveolens* (EO) and Vitamin E (Vit E) or their combination

		Control	DL	DL + EO	DL + Vit E	EO	Vit E
Categories of sperm abnormality in semen samples	Head-abnormalities	-	++++	+	+	-	-
	Mid-piece abnormalities	-	++++	-	-	-	-
	Tail abnormalities	+	++++	+	+	+	+

+++ Severe intensity of the anomaly, + moderate to low intensity of the anomaly, - very low intensity of the anomaly.

Table 4 Effect of different treatments on oxidative stress parameters: lipid peroxidation (MDA) and protein carbonyl content (PCO) in testis of controls (C) and mice treated with deltamethrin (DL), essential oil of *P. graveolens* (EO) and Vitamin E (Vit E) or their combination

	C	DL	DL + EO	DL + Vit E	EO	Vit E
MDA ^a	299.26 ± 11.64	470.93 ± 11.02***	303.8 ± 13.88###	297.46 ± 5.57###	242.20 ± 8.72###	284.53 ± 12.10###
PCO ^b	0.30 ± 0.10	0.92 ± 0.25***	0.46 ± 0.10##	0.47 ± 0.08##	0.23 ± 0.13###	0.53 ± 0.08###

^a (nmoles/g); ^b (μmoles carbonyl/mg protein). Values are expressed as means ± SD of ten mice in each group. All groups vs. control group: * P < 0.05; ** P < 0.01; *** P < 0.001. All groups vs. DL group: # P < 0.05; ## P < 0.01; ### P < 0.001.

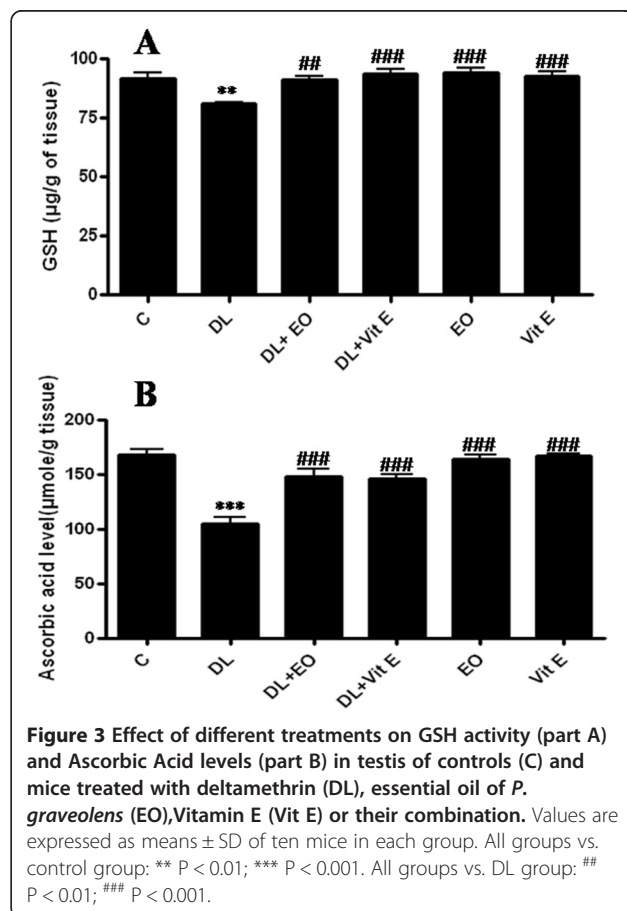
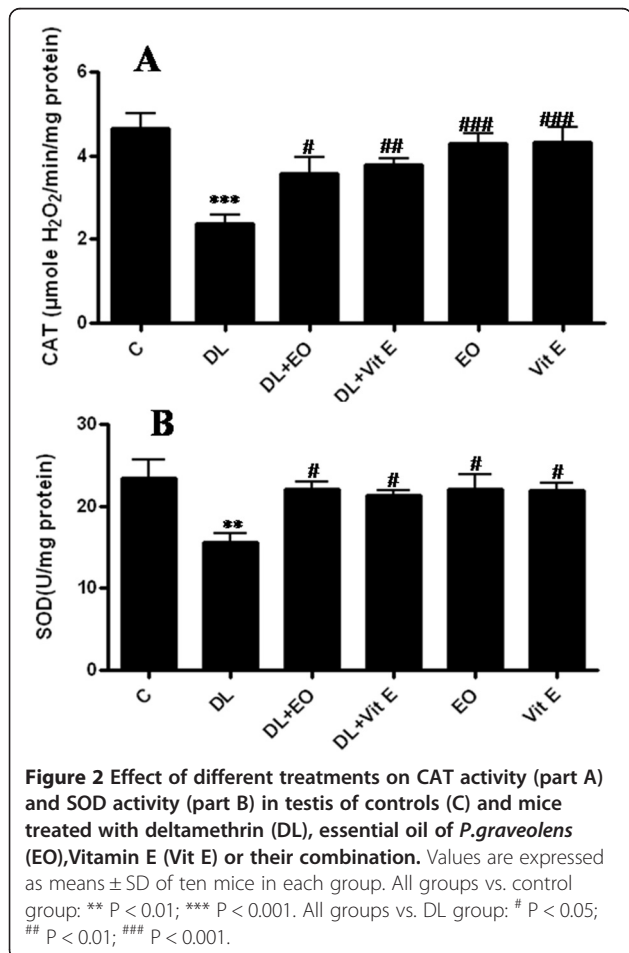
(NBT). The developed blue colour reaction was measured at 560 nm. Units of SOD activity were expressed as the amount of enzyme required to inhibit the reduction of NBT by 50% and the activity was expressed as U/ mg protein.

Determination of testis non-enzymatic antioxidants

GSH in tissues was determined by the method of Ellman [28] modified by Jollow et al [29] based on the development of a yellow colour when 5, 5-dithiobis-2nitrobenzoic acid (DTNB) was added to compounds containing sulfhydryl groups. 500 μl of testis tissue homogenate in phosphate buffer were added to ml of 4% sulfosalicylic acid. The

mixture was centrifuged at 1600 g for 15 min. 500 ml of supernatant were taken and added to Ellman's reagent. The absorbance was measured at 412 nm after 10 min. total GSH content was expressed as μg/mg of tissue.

Ascorbic acid content was determined spectrophotometrically by dinitrophenyl-hydrazine method described by Jacques-Silva and others [30]. Briefly, the ascorbic acid in the testes homogenate was oxidized by Cu²⁺ to form dihydro-ascorbic acid, which reacts with acidic 4-dinitrophenylhydrazine to form a red hydrazone. Final color development was achieved with 65% sulfuric acid and then optical density was measured at 540 nm. The concentration of the samples in vitamin c is obtained from a standard range of ascorbic acid from a dilution of 10 mg/ml ascorbic acid. The calibration curve was



prepared using ascorbic acid as standard. Results were expressed as $\mu\text{mole/g}$ of tissue.

Protein determination in testis

The total protein concentration of supernatant was determined according to Lowry et al [31] using bovine serum albumin BSA as a standard.

Histopathology

For light microscopic examination, tissue samples of testis of the experimental mice were fixed in 4% paraformaldehyde solution for 24 h. After routine processing, paraffin-embedded tissue samples were sectioned at 3-5 μm thickness and stained with Mayer's haematoxylin and eosin. For testicular toxicity, approximately 200 circularly sectioned seminiferous tubules for each mouse were assessed microscopically, and then the percentages of tubules with histopathological changes were calculated [32].

Statistical analyses

The results obtained are expressed as mean values \pm SD. Statistical significance was assessed using one-way analysis of variance (ANOVA). Values of $P < 0.05$ were regarded as statistically significant. All statistical analyses were performed using the Graph Pad Prism version 4 Software (Graph Pad Software, Inc. La Jolla, CA, USA).

Results and discussion

The result obtained with GC/MS analysis of the oil is depicted in Table 1. The essential oil was found to contain 18 constituents representing 89.04% of the total essential oil while minor constituents (10.96%) of the oil remained unidentified. The most abundant components ($> 4\%$) of the *P. graveolens* essential oil were β -citronellol (29.3%) followed by geraniol (10.53%), linalool (10.42%) and citronellal formate (9.54%). These results were in agreement with previous studies [33,34] which demonstrated that the geranium leaves essential oil contains high levels of these compounds. Further, Bouaziz and his collaborators were proved that the antioxidant activity of the essential oil could be attributed in part to the presence of compounds such as β -citronellol and geraniol and its ability to decompose free radicals by quenching reactive oxygen species and trapping radicals before reaching their cellular targets [35].

Xenobiotics and environmental contaminants such as pesticides are known to induce a broad spectrum of toxicological effects and biochemical dysfunctions constituting serious hazards to health. The epididymal sperm count motility and viability were significantly decreased ($p \leq 0.001$) after DL administration (Table 2) whereas a significant increase was observed in abnormal morphology of spermatozoa (head anomalies (Figure 1c-d); mid-piece anomalies (Figure 1e-f) and tail anomalies

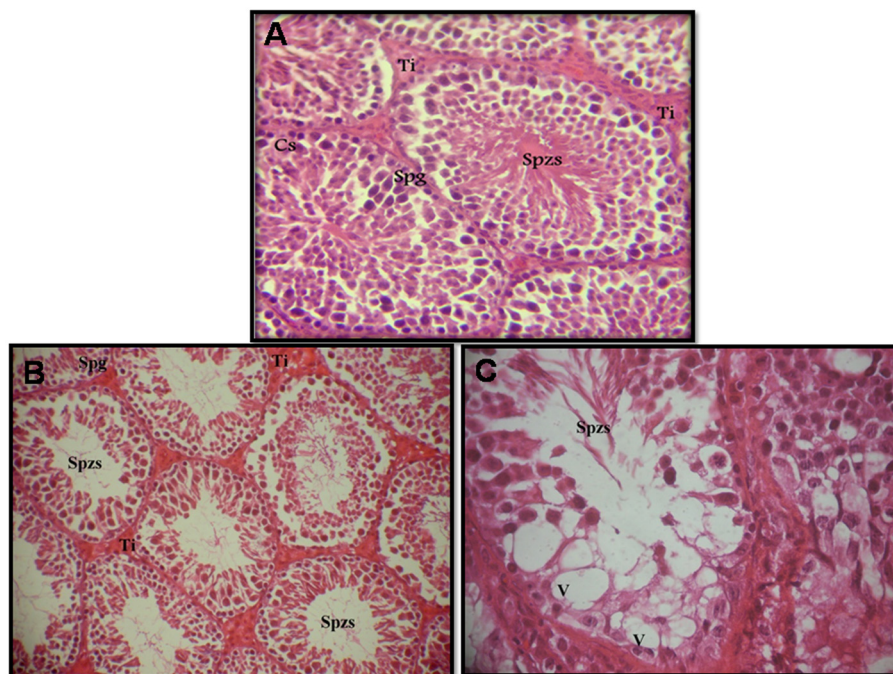


Figure 4 Testicular sections of control mice which show normal spermatogenesis (part A): Note the normal cell arrangement in the seminiferous tubules. The interstitial spaces also appear normal Ti, interstitium; Sg, spermatogonia; Sd, spermatid; Spzs, spermatozoa; CS, Sertoli cell (x 200 H&E). Testicular sections of mice treated with 5 mg/kg/day of DL: Note the atrophic seminiferous tubules (part B) (x400 H&E). Sloughing of germ cells into tubular lumen, V, vacuolization in Sertoli cells and (arrow) multinucleated giant cell (part C) (x1000 H&E).

(Figure 1g-i) compared to the control group (Figure a-b). These anomalies are counted in each group and classified according to their intensities in Table 3. Thus, our results support and extend previous reports suggesting that synthetic pyrethroid generally impairs sperm characteristics [36].

The reduction in sperm count may be due to an adverse effect of DL on spermatogenesis. Lipid peroxidation (LPP) is a marker of oxidative damage. DL-exposed mice showed a significant increase ($p \leq 0.01$) of LPP and protein carbonyls content in the testis compared to control animals (Table 4). Increased lipid peroxidation and altered membrane function can render sperm dysfunctional through impaired metabolism, motility, acrosome reaction reactivity and fusogenic capacity as well as oxidative damage to sperm DNA [37,38]. These results were similar to those of Samah et al [39] who reported the DL induced LPP production in testis rats. The DL induced reduction in testicular CAT activity (Figure 2) may reflect less capacity of testicular mitochondria and microsomes to eliminate H_2O_2 . SOD is considered the first line of defense against deleterious effects of free radicals in the cell by catalyzing the dismutation of superoxide radicals to hydrogen peroxide and molecular oxygen. Testicular SOD activity (Figure 3) declined significantly ($p \leq 0.01$) in the testis of DL-treated animals

when compared with control animals. Thus, our results support and extend previous reports suggesting that pesticide intoxication generally impairs the testicular antioxidant defense system and induces lipid peroxidation in experimental animals and humans [40,41]. Wang et al [42] have also reported that Beta-Cypermethrin decreased the activity of both CAT and SOD. Yet, these activities increased after antioxidants supplementation. We have also investigated the contribution of GSH and vitamin C against oxidative stress. DL treatment led to a significant decrease in GSH ($P < 0.01$) (Figure 4) and ascorbic acid ($P < 0.001$) levels in testis (Figure 5) compared with control group. GSH is an important naturally occurring antioxidant, which prevents free radical damage and helps detoxification by conjugating with chemicals. Under oxidative stress, GSH is consumed by GSH related enzymes to detoxify the peroxides produced due to increased lipid peroxidation [43]. These effects were significantly mitigated by 67 mg/kg EO co-treatment compared with DL-treated group. However, in essential oil supplemented groups the LPP and protein carbonyls declined significantly with relative to DL-exposure group. Our results showed that deltamethrin induced oxidative stress in the mice testis may be successfully treated with the essential oil, geranium, through its antioxidant effects. In this setting, geranium

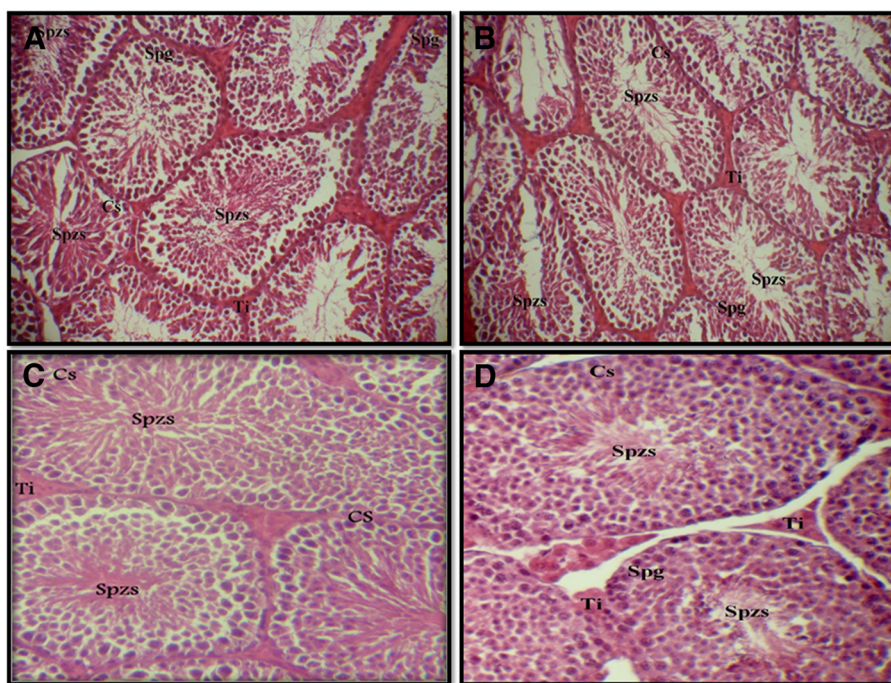


Figure 5 Testicular sections of mice treated with 5 mg/kg/day of DL and 100 mg/kg Vit E (figure A) or DL + EO (part B): Note the increase of germ cells in the seminiferous tubules. The interstitial spaces also appear normal Ti, interstitium; Sg, spermatogonia; Sd, spermatid; Spzs, spermatozoa; CS, sertoli cell (x 400 H&E). Testicular sections of mice treated with Vit E (part C) or EO (part D): Note the normal cell arrangement in the seminiferous tubules. The interstitial spaces also appear normal Ti, interstitium; Sg, spermatogonia; Sd, spermatid; Spzs, spermatozoa; CS, sertoli cell (x 400 H&E).

prevented testicular oxidative damage, reduced lipid peroxidation and improved total sperm motility, viability and morphology in mice spermatozoa. The histological results reported in the current study confirmed the biochemical results. In light microscopic examinations, DL-exposed mice showed severe histopathological changes in the testis such as atrophic seminiferous tubules (Figure 4B). The affected tubules were lined by fewer spermatogenic cells. In addition, appearance of multinucleated giant cells, vacuolization in Sertoli cells and loss of germ cells were also found (Figure 4C). However, the essential oil and vitamin E administration to DL group reduced the appearance of histo-pathological changes in testis (Figure 5 A-D). The testes of control group (Figure 4A) and animals administrated with EO and Vit E (Figure 5 C-D respectively) showed a normal histological structure. In fact, seminiferous tubules presented a complete spermatogenesis and an increase in spermatozoa inside the lumen of seminiferous tubules. Similar histological changes in the testis and epididymis have been documented previously [39]. The administration of essential oil of *P.graveolens* exerts a strong protective effect against oxidative stress damage at the protein level. The essential oil at a dose of 67 mg/kg/day was effective in the reduction of proteins oxidation in testis induced by deltamethrin toxicity. The reduced oxidative stress and lipid peroxidation observed in the geranium essential oils treated animals may be attributed to the important role of the essential oil as antioxidants. The geraniol and β -citronellol may have influenced the geranium essential oil effects as they were major components of the oil (Table 1). In addition, geraniol has been found to have significant antioxidant properties that might be effective for many problems and stabilized and protected the cell membrane against oxidative stress [44]. Numerous studies (in animals and infertile humans) reported a beneficial effect of vitamin E supplementation on sperm quality: reducing the amount of TBARS (marker of lipid peroxidation) in seminal plasma with improved mobility and velocity of sperm, increase concentration and decrease abnormal sperm [45]. Administration of EO at dose of 67 mg/kg bw, 2 hours after DL exposure showed the same effect as vitamin E and has improved the poor quality of sperm induced by deltamethrin which cause higher motility score in the spermatozoa. Their positive effects stem from their ability to inhibit lipid peroxidation, chelate redox- active metals, and attenuating other processes involving reactive oxygen species [46,47].

To the best of our knowledge, this study is the first *in vivo* experimental that investigated the antioxidant effect of *Pelargonium graveolens* on mice sperm kinematic parameters.

The rose-scented geranium *Pelargonium graveolens* L'Her is used for many years in traditional medicine

[48,49]. Recently, Boukhris et al demonstrated that administration of essential oil of *P.graveolens* may be helpful in the prevention of diabetic complications associated with oxidative stress [50]. It seems that many of these pharmacologic features of essential oil of geranium are due to its potent antioxidant actions. In the present study, geranium acts as a natural suppressor of oxidative stress which resulting in sperm promoted quality.

Conclusion

Based on these results, this study showed a positive influence of geranium essential oil in the animal male reproductive system. Further preclinical studies and clinical trials in humans are needed to find out a possible place for it in therapies of fertility disturbances.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

AB; MB and MB designed the experiments, analyzed the data and drafted the manuscript. TB, helped in histopathology. AT helped in data analysis. NA and RG conceived research and approaches and have given final approval of the manuscript to be published. All authors read and approve the final manuscript.

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