Testicular dysfunction induced by aluminum oxide nanoparticle administration in albino rats and the possible protective role of the pumpkin seed oil

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Abstract

Background: Even though the widespread of nanoalumina and their benefits in all fields, its potential impacts on male reproductive system have limited information.

Objective: The present study was conducted to investigate the testicular dysfunction of nanoalumina and the protective role of pumpkin seed oil (PSO) against potential adverse impacts induced by alumina nanoparticles (Al2O3-NPs) in male rat.

Methodology: Al2O3-NPs were administered to the rat orally at a dose of 70 mg/kg body weight once a day for 28 successive days, while pumpkin seed oil was administered to the rat orally at 4 mL/kg b w before administration of Al2O3-NPs, once a day for 28 successive days. After the administration period, sperm concentration, motility, morphology, and DNA damage, as biomarkers of reproductive toxic effects, were evaluated using sperm analysis and comet assays, and histopathological examination of testis was performed. In addition, level of the serum testosterone hormones were estimated, and the levels of oxidative stress biomarkers that take part in the reproductive pathologies such as catalase, glutathione, and malondialdehyde were estimated.

Results: The present results revealed that Al2O3-NPs induced DNA damage in testicular cells, marked histopathological alterations, and caused a significant elevation in MDA in testicular tissue. There was a significant decline in GSH and CAT activities. Furthermore, there was a significant decline in serum testosterone level in the testicular tissue of Al2O3-NP-administered rats. In contrast, pumpkin seed oil co-administration alleviated DNA damage and improved the histopathological alterations in the testicular tissues. Moreover, pumpkin seed oil co-administration significantly reduced MDA and improved the antioxidant defenses in testicular tissue.

Conclusion: The current study concluded that Al2O3-NPs caused testicular dysfunction by generating oxidative injury. Otherwise, PSO co-administration successfully attenuated the adverse impacts of Al2O3-NPs via suppression of oxidative stress and apoptosis as well as enhancement of the antioxidant defense system.

Keywords: Al2O3 nanoparticles, Pumpkin seed oil, Lipid peroxidation, Testicular dysfunction
Introduction
Alumina nanoparticles (Al$_2$O$_3$-NPs) are among the most widely used nanoparticle owing to their distinct physical, thermal, chemical, and biological properties (Arul Prakash, Dushendhra Babu, Lavanya, Vidhya, & Devasena, 2011; Shirai, Watanabe, Fuji, & Takahashi, 2009). Al$_2$O$_3$-NPs are extensively used in many fields of everyday life, including the modification of polymers, functionalization of textiles, aerospace, electronics, pharmaceutical and cosmetics (Bartsch, Saruhan, Schmücker, & Schneider, 1999; Feng et al., 2015; Zhang et al., 2007), heat transfer fluids, treatment of wastewater, biosensors (Tarlani, Isari, Khazraeli, & Eslami Moghadam, 2017), biofiltration, drug delivery (Borbane, Pande, Vibhute, Kendre, & Dange, 2015), and antigen delivery for immunization purposes (Beyzay, Zavaran Hosseini, & Soudi, 2017; Frey, Dange, 2015), and hepato- and nephrotoxicity (Morsy, Abou El-Ala, & Ali, 2016a, 2016b, 2016c; Shirai, Watanabe, Fuji, & Takahashi, 2009). Al$_2$O$_3$-NPs are extensively used in many fields of everyday life, including the modification of polymers, functionalization of textiles, aerospace, electronics, pharmaceutical and cosmetics (Bartsch, Saruhan, Schmücker, & Schneider, 1999; Feng et al., 2015; Zhang et al., 2007), heat transfer fluids, treatment of wastewater, biosensors (Tarlani, Isari, Khazraeli, & Eslami Moghadam, 2017), biofiltration, drug delivery (Borbane, Pande, Vibhute, Kendre, & Dange, 2015), and antigen delivery for immunization purposes (Beyzay, Zavaran Hosseini, & Soudi, 2017; Frey, Neutra, & Robey, 1997). This excessive utilization of Al$_2$O$_3$-NPs makes it widely distributed in our environment. Aluminum oxide nanoparticles may possibly enter the food chain and be responsible for toxicity in animals (Rastogi et al., 2017). NPs have greater toxicity than the same conventional sized particles; hence, their potential adverse effects on the health of humans and wildlife have great concerns (Qin et al., 2017; Serra et al., 2019; Tang et al., 2015).

Several studies have reported that Al$_2$O$_3$-NPs can cross biological barriers and deposit in different body organs and tissues; however, their accumulation at body sites can lead to adverse effects, such as genotoxicity (Balasubramaniam et al., 2009), inflammatory response (Oesterling et al., 2008), carcinogenicity (Dey et al., 2008), cytotoxicity, ROS generation and mitochondrial dysfunction (Chen, Yokel, Hennig, & Toborek, 2008), neurotoxicity (Chen et al., 2008; Li et al., 2009; Prabhakar et al., 2012; Shrivastava, Raza, Yadav, Kushwaha, & Flora, 2014; Zhang et al., 2011), and hepato- and nephrotoxicity (Morsy, Abou El-Ala, & Ali, 2016a, 2016b, 2016c; Shrivastava et al., 2014; Yousef, Mutar, & Kamel, 2019). Therefore, the potential risks to reproductive health should be investigated, especially in those who are occupationally exposed to Al$_2$O$_3$-NPs.

Pumpkin seed oil has been widely used not only as edible oil but also as a nutraceutical. PSO is rich in various antioxidant vitamins as α- and γ-tocopherol, β carotene, and vitamin E (Stevenson et al., 2007). Ninety-eight percent of pumpkin seed oil constituents are fatty acids (mainly linoleic acid, oleic acid, palmitic acid, and stearic acid). In addition, PSO contains also phenolic compounds such as vanillic acid, tyrosol, and vanillin and high levels of selenium and lutein (Andjelkov, Camp, Trawka, & Verhe, 2010). Furthermore, it contains squalene which is a carbon organic compound which has several commercial uses (Nyam, Tan, Lai, Long, & Che Man, 2009). In addition, it has been shown to contain high levels of phytosterols and proteins (Ryan, Galvin, O’Connor, Maguire, & O’Brien, 2007). Pumpkin seed oil has many benefits to human health, due to the presence of all these constituents. Nowadays, it is used in treatment of several diseases as diabetes (Caili, Huan, & Quanhong, 2006), hypertension, hypercholesterolemia (Al-Zuhair, Abd el-Fattah, & Abd el Latif, 1997), symptomatic micturition disorders (Fruhwirth & Hermetter, 2007), and arthritis (Fahim, Abd-el Fattah, Agha, & Gad, 1995). Pumpkin has anti-mutagenic activities as reported by Elfiy, Elelaimy, Hassan, Ibrahim, and Elsayad (2012).

The toxicological impacts of Al$_2$O$_3$-NPs on male re-productive system have limited information. Therefore, it is important to assess the possible testicular toxicity induced by Al$_2$O$_3$-NPs administration and the possible protective role of pumpkin seed oil in prevention and/or reduction of toxicity associated with oxidative stress induced by alumina nanoparticle.

Materials and methods
Animals
Twenty adult male Wister rats (Rattus norvegicus; 7–9 weeks old; 180–200 g body weight) were purchased from the animal house of the Faculty of Veterinary Medicine, Cairo University, Egypt. Animals were placed in the laboratory for a week under the laboratory conditions. The current study is related to the protocol that was approved by the Institutional Animal Care and Use Committee (CU-IACUC) of Cairo University (No. CU/I/F/61/19). This study was carried out in accordance with the Guide for the Care and Use of Laboratory Animals 8th Edition 2011.

Materials
Nanopowder, white, solid Al$_2$O$_3$-NPs (≤ 50 nm) were obtained from Sigma-Aldrich (St. Louis, MO, USA, product number 544833). Characterization of aluminum oxide nanoparticles was characterized using Transmission Electron Microscope (HR-TEM, FEL, Tecnia G20); X-ray diffraction (XRD, PanAnalytical, X pert Pro) was done in my previous work (not published) which confirmed the very thin particles of Al2O3-NPs (nanopowder, < 50 nm).

Pumpkin seed oil (PSO), 100% natural cold pressed pumpkin seed oil, was obtained from Intenan Health Shop, Egypt.

Mouse Negative Control Serum (932B) was obtained from Sigma-Aldrich.

Experiment design
Seventy milligrams per kilograms body weight of Al$_2$O$_3$-NPs was suspended in 1-ml distilled water (aqueous suspension) according to Yousef et al. (2019); the suspensions were ultrasonicated before they were orally
administered to rats to avoid aggregation and provide an optimum size distribution for dispersed particles. Rats were randomly designed into four groups that had five rats each group. Doses were administered once daily via gastric tube for 28 successive days. Group 1: Control group, untreated rats. Group 2: Pumpkin Seed Oil group, rats were administered with 4 ml/kg oil according to Sayed (2014). Group 3: nanoalumina group. Group 4: combined group—rats were co-administered with pumpkin seed oil and Al₂O₃-NPs aqueous suspension in the same doses in 2nd and 3rd groups. Rats were sacrificed under anesthesia using 50 mg/kg of sodium pentobarbital. Blood samples were collected for hormonal analysis (testosterone) from the right ventricle of the rats by syringe, and the testicular and epididymal tissues were removed for further assays.

### Sperm quality analysis

Concentration, motility, and viability of sperm are the three parameters used to determine sperm quality. In a Petri dish containing 2 ml phosphate-buffered saline (PBS) at 37 °C, the cauda epididymis was removed and cut into small portions to prepare sperm suspension. The method of hemocytometer was used to analyze concentration and motility of sperm (WHO, 1999). One drop of sperm suspensions was mixed with two drops of 1% eosin Y (to determine the viability of the sperm); a total of 1000 sperms were counted to determine the proportion of malformed sperm.

### Comet assay

The alkaline comet assay was carried out according to the method of Nandhakumar et al. (2011). It was performed using testicular tissue of male rats from all the experimental groups. The fluorescent microscope (Carl Zeiss Axiosplan with epifluorescence using filter 15 BP546/12, FT580, and LP590) was used to examine the slides. For each sample, the extent of DNA migration was determined by capture of simultaneous image and scoring of 50 cells at magnification × 400 using the Comet 5 image analysis software developed by Kinetic Imaging, Ltd. (Liverpool, UK). By using a closed-circuit digital (CCD) camera, the comet images were captured.

### Serum testosterone assay

By using commercial ELISA kits according to Tietz (1995), serological testosterone hormone was estimated.

### Estimation of the markers of oxidative stress and antioxidants

#### Tissue homogenate preparation

Testicular tissues of all groups were removed, washed in 0.9% saline, and then dried on filter paper. One hundred milligrams of tissue was homogenized in 1 ml of PBS and stored at −20 °C overnight. The homogenates were centrifuged at 300 rpm for 15 min. Supernatants were used to evaluate malondialdehyde (MDA) according to Ohkawa, Ohishi, and Yagi (1979); glutathione (GSH) according to Beutler, Duron, and Kelly (1963); and catalase (CAT) according to the method of Aebi (1984).

### Histopathological assay

Testicular tissues were fixed in 10% neutral formalin for 72 h. Testicular tissues were dehydrated in alcohol, cleared in Xylene, and embedded in paraffin wax. Three to 5-μm sections were cut by microtome. Hematoxylin and Eosin were used to stain the sections (Bancroft & Gamble 2008). By using light microscope at magnification × 400, the histological changes were investigated.

### Immunohistochemistry for caspase-3

Immunohistochemical staining of anti-caspase-3 antibodies was performed with streptavidin-biotin (Duan et al., 2003). The sections of 3–5 μm thick were incubated with fresh 0.3% hydrogen peroxide in methanol for 30 min at room temperature. The specimens were then incubated with anti-caspase-3 antibodies as the primary antibody at a 1:100 dilution. After the incubation, the reaction product was visualized with (freshly prepared) diaminobenzidine solution by incubating for 60 min. Finally, sections were counterstained with Mayer’s hematoxylin, mounted with mounting medium, and examined with the Olympus BH2 photo-light microscope (Olympus America Inc., Center Valley, PA, USA). Negative controls were prepared by substituting normal mouse serum for each primary antibody.

### Statistical analysis

Data were represented as mean ± standard error (SE). Statistical analysis of data was performed using GraphPad Prism 5. Data were analyzed for statistical significance by the one-way analysis of variance, followed by the Tukey’s multiple comparison test. The data at \( p < 0.05 \) was considered significant.

### Results

#### Sperm quality analysis

Table 1 presented the parameters of epididymal spermatozoa of Al₂O₃-NPs and/or PSO groups. PSO group did not differ significantly from the control in sperm motility, sperm viability, and abnormal sperm rates. Concentration, motility, and viability of spermatozoa were significantly low in Al₂O₃-NP-administrated group compared to the control group. Significant elevation in abnormalities was in the Al₂O₃-NP-administrated group compared to the control group as seen in Table 1 and Fig. 1. Co-administration of PSO significantly alleviated the decrease in concentration, motility, and viability of
spermatozoa and significantly lowered the percentage of abnormal spermatozoa compared to Al₂O₃-NP-administrated group. PSO had a remarkably protective role against DNA damage caused by Al₂O₃-NPs.

**Comet assay**

Figure 2 represented the comet cells in all groups. Administration of Al₂O₃-NPs for 28 days for male rats revealed an elevated DNA amount in the comet tail in testicular cells. Significant increment (\( p < .05 \)) in all comet parameters (% DNA damage, tail length, and tail moment) in testicular cells of Al₂O₃-NP-administrated rats was compared to control (Table 2). In contrast, there was a significant decline in the comet parameters of PSO and Al₂O₃-NPs co-administrated rats, thus PSO alleviating Al₂O₃-NPs toxicity.

### Effect of PSO and/or Al₂O₃-NPs on serum levels of testosterone

Serum levels of testosterone significantly declined (\( p < .05 \)) in Al₂O₃-NP-administrated rats when compared to control rats. Furthermore, serum testosterone concentration in PSO and Al₂O₃-NPs co-administrated rats was significantly elevated (\( p < .05 \)), compared to Al₂O₃-NP-administrated rat (Fig. 3).

### Effect of PSO on Al₂O₃-NP-induced oxidative stress in the testicular tissue

Al₂O₃-NP-administrated rat showed a marked increase (\( p < .05 \)) of MDA and a decline in GSH content and CAT activities in the testicular homogenates, as compared to control rat. PSO-administrated rat revealed a significant change in the levels of the aforementioned

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**Table 1** Effect of pumpkin seed oil on the percentage of sperm parameters in rat administrated with Al₂O₃-NPs

| Parameter                  | Control         | PSO             | Al₂O₃-NPs       | PSO + Al₂O₃-NPs  |
|----------------------------|-----------------|-----------------|----------------|------------------|
| Sperm concentration (× 10⁶ ml⁻¹) | 79.05 ± 0.3069 | 80.85 ± 0.3617  | 40.13 ± 0.2394ª | 59.4 ± 0.3109ª²  |
| Sperm motility (%)          | 85.38 ± 0.4479  | 86.13 ± 0.225    | 56.8 ± 2.478ª²  | 70.05 ± 0.3571ª² |
| Sperm viability (%)         | 84.18 ± 0.4131  | 85.85 ± 0.3524   | 59.63 ± 0.2394ª | 69.98 ± 0.225ª² |
| Sperm abnormal forms(%)     | 6 ± 0.4082      | 5.5 ± 0.2887     | 21.5 ± 0.6455ª² | 12.25 ± 0.25ª²   |

ªRefers to a significant change from the control rat  
²Refers to a significant change from the Al₂O₃-NP-administrated rat

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\[\text{Fig. 1} \quad \text{Photomicrograph showing sperm morphology from the cauda epididymis of male albino rats. (N) normal sperm; (a) bent head; (b) banana head; (c) coiled tail; (d) detached head; (e) headless sperm. Eosin stain}\]
oxidative stress markers. Meanwhile, PSO in combination with Al₂O₃-NPs reestablished the oxidant/antioxidant balance in the homogenates of the testes of Al₂O₃-NP-administered rat (Figs. 4, 5, and 6).

**Histopathological examination**

Figure 7 showed the photomicrographs of the testes from the different groups. The seminiferous tubules with normal structure showed regularly arranged rows and complete set of germinal epithelium in both control and pumpkin seed oil groups as seen in Fig. 7a and b. Otherwise, seminiferous tubules of Al₂O₃-NP-administered rats showed vacuolization and degenerative and necrotic changes specially in spermatogonial cell layer and in some areas of the germinal epithelium as well as interstitial edema and congestion of blood vessels (Fig. 7c). In PSO and Al₂O₃-NP co-administered rats, amelioration in histopathological examination was seen in testicular tissues (Fig. 7d), where examined sections revealed no histopathological alterations.

**Immunohistochemistry for caspase-3**

Figure 8 showed the photomicrographs of the testes from the different groups. The intensity of activated caspase-3 immunostaining (brown) is predominant on seminiferous tubules and Leydig cells of Al₂O₃-NP-administered rats. In PSO and Al₂O₃-NP co-administered rats, the immunostaining was mitigated by PSO.

**Table 2** Effect of pumpkin seed oil on the percentage of DNA damage (%DNA), tail length (TL), and tail moment (TM), in the testicular tissue of rat administrated with Al₂O₃-NPs

| Parameter            | Experimental groups |
|----------------------|---------------------|
|                      | Control | PSO     | Al₂O₃-NPs | PSO + Al₂O₃-NPs |
| DNA% tail            | 15.59 ± 0.5653      | 15.33 ± 0.2822 | 34.12 ± 2.491ª | 22.93 ± 1.257b |
| Tail length (μm)     | 5.113 ± 0.3702      | 4.943 ± 0.0348 | 24.31 ± 0.8417ª | 11.01 ± 0.3405b |
| Tail moment (μm)     | 1.083 ± 0.3346      | 1.031 ± 0.2924 | 5.827 ± 0.4737ª | 2.936 ± 0.2135b |

ªRefers to a significant change from the control rat
bRefers to a significant change from the Al₂O₃-NP-administered rat
Discussion
NPs possess the ability to cross the blood–testis barrier (BTB) that protects reproductive tissue (Bai et al., 2010; Gao et al., 2013). Hussein, Ali, Saadeldin, and Ahmed (2016) reported that many NPs can cross the BTB and exert their toxic impacts on spermatogenesis. Regarding accumulation and translocation, the capacity of NPs to penetrate both the BTB and the blood–brain barrier is very important (Araujo, Sheppard, Löwenberg, & Kreuter, 1999; Kim et al., 2006).

Lin et al. (2007) showed that SiO2 NPs decreased sperm count and the sperm mobility rate and significantly increased the malformation rate, in the testis of rat after intraperitoneal injection. Guo et al. (2009, 2010) observed that TiO2 NPs and/or ZnO NPs significantly reduced sperm density and motility, and increased sperm abnormality and germ cell apoptosis in the testis of male mouse after intraperitoneal injection. Zhou, Yue, Li, Zhou, and Liu (2019) showed that PbSe-NP administration led to reduction in the quantity and quality of
sperm; accumulation of PbSe-NPs caused oxidative stress, disorder of normal serum sex hormones, endoplasmic reticulum, and mitochondria-mediated cell apoptosis which caused a significant decline in fertility.

Brezina, Yunus, and Zhao (2012) reported that male infertility can be evaluated by using semen parameters as concentration, motility, and spermatozoa morphology. In the current work, reproductive toxicity of Al₂O₃-NPs was characterized by a significant decrease in the concentration and normal morphology of spermatozoa, significant increase in DNA damage, histopathological alterations, and increase in intensity of caspase-3 expression. MDA is a lipid peroxidation by-product, which could indirectly indicate the severity of injured tissues (Sreelatha, Padma, & Umadevi, 2009; Yu et al., 2011). To confirm oxidative injury caused by the accumulation of Al₂O₃-NPs in the testes, MDA levels were determined. There was a significant increment in
Fig. 7 Photomicrographs of testicular sections of different groups, hematoxylin, and eosin. 

- **a, b** Testicular sections from control and PSO groups, respectively, showing normal histological features of seminiferous tubules.
- **c** Testicular section from Al₂O₃-administrated rat showing the degenerative, vacuolization, and necrotic changes specially in spermatogonial cell layer and in some areas of the germinal epithelium as well as interstitial edema and congestion of blood vessels (arrows), interstitial edema, (asterisk)
- **d** Testicular section from the co-administrated group with PSO and Al₂O₃ showing an obvious preservation of spermatogenic epithelium in most seminiferous tubules. SZ, spermatozoa, IS, interstitial cells

Fig. 8 Immunohistochemical staining of activated caspase-3 in rat testis. 

- **a** Testis of control rats.
- **b** Testis of PSO-administered rats.
- **c** Testis of Al₂O₃-administrated rats. The intensity of activated caspase-3 immunostaining (brown color) is pre-dominant on spermatogonia and Leydig cells in Al₂O₃-administered group.
- **d** Testis of PSO+ Al₂O₃ co-administered rats.
testicular MDA when compared with the control group. Furthermore, the administration of Al2O3-NPs resulted in a significant decline in testicular GSH and CAT. It is suggested that Al2O3-NPs could decrease the quantity/quality of sperm through cellular impairment of testicular tissue, which is caused by the generation of ROS. These data are in agreement with former results (Morsy et al., 2016a, 2016b, 2016c), administration of Al2O3-NPs generates a state of oxidative stress in the brain, liver, and kidney tissues of rats, shown by lipid peroxidation increment and antioxidant defense system compromised.

Ema, Kobayashi, Naya, Hanai, and Nakanishi (2010) indicated that in vitro studies revealed that silver, aluminum, and molybdenum trioxide NPs injured spermatogonia stem cells; TiO2 and black carbon nanoparticles (CB NPs) impacted Leydig cell viability, and gold NPs reduced human sperm motility. Yoshida et al. (2009) observed that black carbon nanoparticles have adverse impacts on male reproductive function in mice characterized by elevated serum testosterone (T) levels and histological alterations in both the seminiferous tubules and Sertoli cells such as vacuolation and degenerative changes and Leydig cell dysfunction.

Lan and Yang (2012) stated that the Leydig cells secrete testosterone, an essential hormone for spermatogenesis. Leydig and Sertoli cell apoptosis through DNA damage was reported in vitro after incubation with ZnO NPs (Han et al., 2016) and in vivo after exposure to AgNPs (Castellini et al., 2014; Layali, Tahmasbpour, & Jorsaraei, 2016). Lan and Yang (2012) reported that NP exposure may induce a generalized inflammation in the host and affect Leydig cells, causing decline in testosterone serum levels, which then impaired the integrity of the BTB. In conformity with former findings (Castellini et al., 2014; Ema et al., 2010; Lan & Yang, 2012; Layali et al., 2016), administration of Al2O3-NPs caused the reduction in serum testosterone level and histological alterations such as vacuolation and degenerative changes in the seminiferous tubules.

In conformity with former work (Guo et al., 2009 & 2010; Zhou et al., 2019), administration of Al2O3-NPs caused increase in intensity of caspase-3 expression which is ultimately responsible for the majority of the cell apoptotic process.

Liu et al. (2016) suggested the reproductive toxicity of ZnO NPs using a mouse Sertoli cell line (TM-4) and spermatocyte cell line (GC2-spd) characterized by generating ROS and DNA lesions in germ cells. ZnO NPs could exert their toxicity by downregulating the gap junction protein expression and disrupting the Sertoli cell mitochondrial outer membrane and cell membrane, potentially leading to BTB disruption and further destroying BTB integrity. Furthermore, ROS and the cytokine secretion play important roles in BTB disruption, significant increment in the status of oxidative stress (i.e., leading to ROS and MDA increment and GSH decline), and increasing the cytokine (TNF-α) levels in Sertoli cells.

The induction of sperm abnormalities could be attributed to oxidative stress and lipid peroxidation (Agarwal & Saleh, 2002). Plasma membrane of the spermatozoa is liable to peroxidation due to the high extent of unsaturated fatty acids and low antioxidant capacity. The lipid peroxidation disrupts the structure of the lipid matrix in the membrane of spermatozoa, causing the motility loss and the membrane integrity defects (Vernet, Aitken, & Drevet, 2004).

Morsy et al. (2016a, 2016b) showed that nanoalumina injection for male rats induced the oxidative stress (MDA increment and GSH, and CAT decline) in the brain, hepatic, and renal tissues, leading to DNA damage and histopathological alterations.

It is believed that NPs are absorbed in the testis in their ionic form. Accumulation of Al in the testis can partially contribute to the toxicity of Al2O3-NPs. Human and animal exposure to Al-containing products will increase the concentration of this metallic element in their organs and damage their tissues (including the testicular tissues). Yousef, Kamel, El-Guendi, and El-Demerdash (2007) and Guo, Lin, Yeh, and Hsu (2005, b) reported sperm viability and motility reduction due to the high levels of Al in spermatozoa and seminal plasma. Sharma et al. (2003), Guo et al. (2005, b), and Cheraghi, Golkar, Roshanaei, and Alani (2017) showed accumulation of Al in testicular tissue, necrosis of spermatocytes/spermatids, and a significant decline in fertility in both male rats and mice. Al may exert its toxicity in reproductive system of male through various mechanisms such as generating oxidative stress, interfering with spermatogenesis and steroidogenesis, impairing cell signaling, BTB, and disrupting the endocrine system (Pandey & Jain, 2013).

This observation could be attributed to the ability of Al13 and Al2O3-NPs to induce oxidative stress, penetrate the BTB, generate lipid peroxidation, and ultimately injure the testis biological membrane. The low sperm count, motility, and viability, as well as the high morphological abnormality, DNA damage, and histological alterations shown in Al2O3-NP-administered rats, confirm the previous findings.

Pumpkin seed oil is known to increase membrane fluidity and osmosis and allows for intracellular and extracellular gaseous exchange, attributed to the presence of linoleic acid, a polyunsaturated fatty acid (Lovejoy, 2002). In addition, pumpkin diminishes the ability of the testes and epididymis to lipid peroxidation due to the presence of oleic acid, a monounsaturated fatty acid.
Pumpkin seed oil possesses some fundamental constituents (vitamin A, tannins, linoleic acid, oleic acid, and alkaloids) which inhibit lipid peroxidation, hence, attenuating testicular function (Akinbayo, 1997; Bensousan, Morales, & Hermo, 1998; Leat, Northrop, Harrison, & Cox, 1983). From the present data, it can be suggested that pumpkin seed oil may be used as antioxidant against testicular toxicity which might be induced by Al2O3-NPs administration. Akang, Oremosu, Dosumu, Noronha, and Okanlawon (2010) demonstrated that pumpkin seed oil attenuated semen parameter level of testosterone (T), Testosterone; TiO2: Titanium dioxide nanoparticles; ROS: Robot Operating System; Telferia occidentalis; PSO: Pumpkin seed oil; MDA: Malondialdehyde; NPs: Nanoparticles; PBS: Phosphate-buffered saline; DNA: Deoxyribonucleic acid; ELISA: The enzyme-linked immunosorbent assay; GSH: Glutathione; H&E: Hematoxylin and eosin; MDA: Malondialdehyde; NPs: Nanoparticles; PBS: Phosphate-buffered saline; PbSe NPs: Lead selenide nanoparticles; PSO: Pumpkin seed oil; Al: Aluminum; Al2O3 NPs: Aluminum oxide nanoparticles; AgNPs: Silver nanoparticles; Al: Aluminum; Al2O3 NPs: Aluminum oxide nanoparticles; BTB: Blood–testis barrier; CAT: Catalase; CB NPs: Black carbon nanoparticles; DNA: Deoxyribonucleic acid; ELISA: The enzyme-linked immunosorbent assay; GSH: Glutathione; H&E: Hematoxylin and eosin; MDA: Malondialdehyde; NPs: Nanoparticles; PBS: Phosphate-buffered saline; PbSe NPs: Lead selenide nanoparticles; PSO: Pumpkin seed oil; T: Testosterone; TiO2: Titanium dioxide nanoparticles; ROS: Robot Operating System; ZnO NPs: Zinc oxide nanoparticles.

Conclusion
According to the current data, it could be concluded that PSO exhibits its protective role by inhibiting cellular damage and apoptosis occurring as a result of oxidative stress in the spermatogenic cells of seminiferous tubules and Leydig cells induced by the administration of Al2O3 nanoparticles.

Abbreviations
AgNPs: Silver nanoparticles; Al: Aluminum; Al2O3 NPs: Aluminum oxide nanoparticles; BTB: Blood–testis barrier; CAT: Catalase; CB NPs: Black carbon nanoparticles; DNA: Deoxyribonucleic acid; ELISA: The enzyme-linked immunosorbent assay; GSH: Glutathione; H&E: Hematoxylin and eosin; MDA: Malondialdehyde; NPs: Nanoparticles; PBS: Phosphate-buffered saline; PbSe NPs: Lead selenide nanoparticles; PSO: Pumpkin seed oil; T: Testosterone; TiO2: Titanium dioxide nanoparticles; ROS: Robot Operating System; ZnO NPs: Zinc oxide nanoparticles.

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Author’s contributions
The author did all the work and read and approved the final manuscript.

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Availability of data and materials
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Ethics approval and consent to participate
The study was carried out in strict accordance with the Guide for the Care and Use of Laboratory Animals 8th Edition 2011.

Consent for publication
Not applicable.

Competing interests
No potential conflict of interest was reported by the author.

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