Ameliorative effects of melatonin and zinc oxide nanoparticles treatment against adverse effects of busulfan induced infertility in male albino mice

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Abstract: Testicular damage is one of the most hazardous effects as it’s associated with azoospermia. Busulfan (Bu) is a highly toxic chemotherapeutic drug that affects testis. Thirty male Swiss albino mice divided into six groups of 5 animals each. Control (oral 0.9% saline daily for 75 days); Mel (20 mg/kg/day orally for 30 days); ZnO NPs (5 mg/kg/day i.p. for 30 days); BU (single i.p. injection of 40 mg/kg and then left for 45 days); BU + Mel (single 40 mg/kg dose of BU and left for 45 days followed by 20 mg/kg/day Mel for 30 days); BU + ZnO NPs (single dose of 40 mg/kg of BU and left for 45 days, then 5 mg/kg/day ZnO NPs for 30 days). Preparation and Characterization of ZnO NPs. Specimens from testis prepared for ultrastructural investigations using TEM after Masson's trichrome and toluidine blue staining. BU induced histological and ultrastructural damage of the testis. Moreover, the present results could be concluded that Mel or ZnO NPs can protect the testicular tissue against ultrastructural alterations induced by BU by its antioxidant and anti-apoptotic effects.

Introduction

Infertility is a serious health problem around the world (Inhorn and Patrizio, 2015). Infertility can be caused by many medications such as chemotherapies and fertility is often impaired after chemotherapy (Delessard et al., 2020; Poganitsch-Korhonen et al., 2017). Deleterious in germ cells was detected due to the exposure to alkylating agents, such as cyclophosphamide, procarbazine, and busulfan (Delessard et al., 2020).

Busulfan (BU) is an alkylating agent that induces deoxyribonucleic acid (DNA) damage impairing DNA replication (Li et al., 2018). Therefore, BU is a highly toxic chemotherapeutic agent that results in adverse effects on gonads (Mirhoseini et al., 2014). After one or two intraperitoneal injections, BU can effectively consume most spermatagonia. Thus, it is important to use antioxidants to treat infertility related to BU (Brinster et al., 2003).

Melatonin (Mel.) is a highly conserved molecule synthesized in all organisms. It is the chief secretory product of the pineal gland and was recently reported to be a powerful antioxidant (Gurunathan et al., 2020; Haghi-Aminjan et al., 2018). In both in vitro and in vivo studies, Mel protected tissues against oxidative damage caused by various free radicals. One of the protective mechanisms that might be related to Mel is anti-apoptotic effects (Haghi-Aminjan et al., 2018).

Melatonin (N-acetyl-5-methoxytryptamine), the chief secretory product of the pineal gland in the brain, is well known for its functional versatility. In hundreds of investigations, melatonin has been documented as a direct free radical scavenger and an indirect antioxidant, as well as an important immunomodulatory agent. The radical scavenging ability of melatonin is believed to work via electron donation to detoxify a variety of reactive oxygen and nitrogen species, including the highly toxic hydroxyl radical. It has long been recognized that the damaging effects of ionizing radiation are brought about by both direct and indirect mechanisms. The direct action produces disruption of sensitive molecules in the cells, whereas the indirect effects (70%) result from its interaction with water molecules, which results in the production of highly reactive free radicals such as ·OH, ·H, and eaq− and their subsequent action on subcellular structures. The hydroxyl radical scavenging ability of melatonin was used as a rationale to determine its radioprotective efficiency (Reiter et al., 2004).

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Melatonin is a hormone produced naturally by the testes, retina, ovary, skin, and intestine (Chen et al., 2016). Melatonin is involved in the regulation of various biological pathways, including hormone secretion and modulation, reproduction, and circadian rhythms, according to previous research (Kim et al., 2012; Wang et al., 2012; Maroufi et al., 2020). Furthermore, melatonin has been demonstrated to have an effective anticancer effect on a variety of cancer cells with no adverse effects (Wang et al., 2012; Maroufi et al., 2020). Melatonin’s actions are mediated by receptor-dependent or receptor-independent mechanisms (Maroufi et al., 2020; Witt-Endersby et al., 2006). Melatonin receptors MT1 and MT2 suppress many signaling pathways involved in cancer growth, including phosphoinositide-3-kinases (PI3K) and mitogen-activated protein kinase (MAPK) pathways, as well as induce apoptotic pathways in cancer cells (Kong et al., 2008; Mao et al., 2010). Furthermore, when taken alone or in conjunction with therapeutic drugs, melatonin has been shown to have anticancer effect in hormone-dependent cancers (Menéndez-Menéndez and Martínez-Campa, 2018; Naja et al., 2019). Toxicity, cellular DNA breakage, and lipid peroxidation are all reduced by melatonin. Melatonin’s ability to scavenge free radicals implies that it could be employed as a dependable protective medication in radiotherapy (Tan et al., 2008).

On the other hand, zinc (Zn) exists at high concentrations in testes of vertebrates comparable to the liver and kidney (Fallah et al., 2018). In addition, Zn could decrease testicular damage caused by many chemotherapeutic agents such as cyclophosphamide (Torabi et al., 2017). During early spermatogenesis, Zn is collected in the testes, and this plays a significant role in the meiosis division of germinal cells (Elgazar et al., 2005). Additionally, Zinc Oxide Nanoparticles (ZnO NPs) is a new type of material with outstanding biological properties and no toxicity (Suresh et al., 2018). It seems to have great potential for effective cell and molecular selection through certain obstacles in many diseases (Keerthana and Kumar, 2020). The important characteristics of ZnO NPs are antioxidant, antimicrobial, anticancer, anti-inflammatory activities (Madhumitha et al., 2016; Umar et al., 2019). Therefore, the present study evaluated the protective role of Mel or ZnO NPs has antioxidant effects on the alterations of testis induced by BU in the adult male mice as ultrastructure study.

Materials and Methods

Chemicals
Myleran (busulfan) (1, 4-butanediol dimethanesulfonate), used as tablets (2 mg), was manufactured by Excella GmbH (Feucht, Germany). The BU dosage was prepared in normal saline after being dissolved in DMSO and then was administrated in a single (i.p) dose of 40 mg/kg (Mirhoseini et al., 2014).

Melatonin was purchased as 10 mg tablets from Puritan’s Pride (Holbrook, NY, USA). Mel dosage prepared in water and was administrated orally at doses of 20 mg/kg/day (Mirhoseini et al., 2014).

Zinc oxide nanoparticles were white powdered with particle size 10–30 nm (Catalog Number US 677450, Sigma-Aldrich, Saint Louis, MO, USA). The ZnO NP dosage was prepared in standered saline solution and administrated intraperitoneally at 5 mg/kg/day (Torabi et al., 2017).

ZnO nanoparticle preparation and characterization
ZnO nanoparticle powder was autoclaved and UV-sterilized overnight. The sterilized ZnO nanoparticles were suspended and ultra-sonicated (Branson ultrasonic cleaner 3210, 120W, Branson Ultrasonic Corporation, Danbury, CT, USA) for 15 min in pure ethanol (99.9%, Samchun Chemicals, Seoul, Korea). The ZnO nanoparticles of Transmission electron microscopy images were observed in a Jeol JSM-6510 L.V SEM microscope using 30 KV at EM Unit, Mansoura University, Egypt (Fig. 1).

Experimental animals
Thirty healthy male Swiss albino mice weighting 25–32 g were obtained from the Animal House of the Faculty of Pharmacy, Mansoura University, Egypt. They were housed in an artificially illuminated (12 h light–12 h dark cycle), air-conditioned and temperature-controlled room (22–25°C); the commercial laboratory chow and water were available for the animal’s ad libitum. The experiment started after two weeks of acclimatization. All animals received humane care and compliance-with the guidelines of the Animal Care and Use Committee of Mansoura University. Code No. ZD180015.

Experimental design
After the acclimatization period, the animals were randomly assigned into six groups of five mice each.

- **Group 1** (control): mice received orally 0.9% saline solution for 75 days.
- **Group 2** (Mel): mice received oral doses of 20 mg/kg/day of Mel for 30 days.
- **Group 3** (ZnO NPs): mice injected daily i.p with 5 mg/kg/day of ZnO NPs for 30 days.
- **Group 4** (BU): mice injected i.p with a single dose of Bu 40 mg/kg and left for 45 days.
- **Group 5** (BU + Mel): mice injected with a single dose of BU followed by Mel after 45 days of BU injection for 30 days.
- **Group 6** (BU+ ZnO NPs): mice injected with a single dose of BU followed by ZnO NPs after 45 days of BU injection for 30 days.

Investigated parameters

- **Light microscopic study**
In all experimental groups, the specimens of right side testis were removed and immediately washed in normal saline and fixed in 10% neutral buffered formalin. After fixation, the samples were dehydrated with an ascending ethanol series, cleared in xylene, embedded in paraffin wax and sectioned at 5 µm thickness. Sections were stained with Masson’s trichrome to demonstrate collagen fibers and examined under a bright-field Olympus microscope.

- **Toluidine blue stain**
Semithin sections were cut with a diamond knife on an LKB microtome and mounted on slides, stained with toluidine blue and examined by a light microscope.
Transmission electron microscopic investigations (TEM)
The ultrastructural investigations of the present work were mainly focused on the testis of male mice. Small pieces of the right testis were immediately separated and fixed in 4% glutaraldehyde in Dulbecco’s modified phosphate buffered saline. Samples were washing and fixed in 1% osmium tetroxide for 1 h at room temperature. After that, they were dehydrated in an ethanol gradient, treated via propylene oxide, then embedded in Epon resin (Epon 812; Fluka Chemie, Switzerland) for TEM investigations. Ultrathin sections (60–70 nm) were cut with a diamond knife on an LKB microtome and mounted on slides, stained with uranyl acetate and lead citrate (Hall et al., 2012). The sections were then examined and photographed at 80 kV using a JEOL 2100 Transmission Electron Microscope in the Electron Microscopic Unit at Faculty of Agriculture, Mansoura University, Mansoura, Egypt.

Results
Characterization of ZnO nanoparticles
Transmission electron microscopy (TEM) images showed that ZnO nanoparticle had a sub-100 nm size on average after the sterilisation process.

Masson trichrome observations
The effect of control, Mel or ZnO NPs on testis was further assessed by demonstrating the expression of Masson on the testis. Masson trichrome stained sections of control, Mel or ZnO NPs groups showed minor expression of collagen fibers at the boundary of seminiferous tubules and in between the tubules (Figs. 2A–2C). A section in the testis of the BU group showed intense positive expression of collagen fibers at the boundary of seminiferous tubules and in between the tubules. Busulfan (BU) was reported as cytotoxic drug associated with pulmonary toxicity, acute lung injury, chronic intestinal fibrosis that defined as the overgrowth, hardening and deposition of extracellular matrix components including collagen (Fig. 2D). Mel or ZnO NPs supplemented mice after BU treatment showed a mild expression of Masson trichrome staining (Figs. 2E and 2F).

Toluidine blue observations
Semithin sections in the testis of control, Mel and ZnO NPs groups stained with toluidine blue, showed seminiferous tubules lined by germinal epithelium, including several types of spermatogenic cells at different stages of spermatogenesis. Sertoli cells were resting on regular basement membranes ensheathed by flat myoid cells and nearly similar to the control. Normal spermatids and an increased number of sperms in the lumen of seminiferous tubule were also noticed (Figs. 3A–3C). The adjacent tubules showed wide intercellular spaces separating the germ cells in the BU-treated group. Degenerated Leydig cells in the interstitial spaces and congested blood vessel were observed. Few number and abnormal shape of spermatids were also seen. The lumen of seminiferous tubules of the BU treated group showed devoid of sperms (Fig. 3D). BU injected mice and supplemented with Mel or ZnO NPs groups improved the general histological picture of the testis. The architecture of seminiferous tubules showed nearly normal and observed the lining of germinal epithelium with well-organized spermatogenic layers at different stages of spermatogenesis. Sertoli cells were nearly normal in their structure. Moreover, seminiferous tubules with much sperms in their lumen and some Leydig cells in the interstitial spaces were also seen in-between (Figs. 3E and 3F).

Electron microscopic observations
The ultrathin sections of control testis revealed that seminiferous tubules (S.T) were surrounded by thin basal lamina that contains flat myoid cells and lined by spermatogenic cells. The epithelium of spermatogenic cells showed the sequences of spermatogonia, spermatocytes, spermatids, Sertoli cells and germ cells in active spermatogenesis. Sertoli cells showed a distinct nucleus with prominent nucleoli and a cytoplasm containing numerous mitochondria, endoplasmic reticulum, lysosomes and lipid droplets (organelles) in an active state. It is located between spermatogenic cells and rested on the basement membrane of the seminiferous tubule. The spermatogonia rested on the basal lamina of the tubule, and had a rounded nucleus with prominent nucleoli and numerous mitochondria were seen in their cytoplasm (Fig. 4A). The primary spermatocytes were round in shape with central large rounded nuclei containing peripheral euchromatin condensation and heterochromatin; numerous mitochondria and lysosomes were also seen (Fig. 4B).

Round spermatid appeared with regular plasma membrane surrounded by acrosomal vesicle with normal

**FIGURE 1.** (A & B): Characterization of the sterilized ZnO nanoparticle; A and B transmission electron microscopy (TEM) images of the ZnO nanoparticle.
round nucleus and nucleolus. Its cytoplasm contains well-developed mitochondria, ribosomes and Golgi apparatus (Fig. 4C). The tail of mature sperm in the middle piece showed that it is formed of central axoneme, dense outer fibers and mitochondrial sheath were around the dense fibers. The mitochondrial sheath was surrounded by the plasma membrane (Fig. 4D). The head of mature spermatozoa was elongated, containing an acrosomal cap covering the nucleus (Fig. 4E). The Leydig cell showed a regular plasma membrane with euchromatic nucleus and prominent nucleolus; its cytoplasm had many normal mitochondria and lysosomes (Fig. 4F).

On the other hand, transmission electron microscopic examination of the testicular tissues in BU-treated group revealed many alterations manifested by irregular thickened basal lamina and thick collagen fibers between myoid cells and basal lamina in focal area, the thickness of basal lamina causes low rate of sperm production. Some tubules showed reduced spermatogenesis (Figs. 5A–5C). Sertoli cells appeared with irregular plasma membrane and nucleus with disintegrated nucleoli and separated from basal lamina. The nucleus of Sertoli cell was irregular in shape and small in size (Fig. 5A). In this group, some seminiferous tubules showed impairment in the growth of spermatogenic cells, and the basal lamina surrounding the seminiferous tubules was irregular and tortuous. The presence of vacuoles was evident in the cytoplasm of Sertoli cells, spermatocytes and spermatids. In some germ cells, the membranes of the nuclei were swollen and vacuolated, including apoptosis in some germ cells. Spermatogonia and primary spermatocyte cells appeared abnormal architecture with degenerated nuclei in some cells and pyknotic in another (Figs. 5A and 5B). The cytoplasm of some germ cells contains clumped organelles with degenerated mitochondria and vacuoles (Fig. 5B). Round spermatids had distorted and degenerated nucleus, cytoplasm revealed deformed and vacuolated mitochondria (Fig. 5C). Some abnormalities of the acrosome of the elongated spermatids were seen, separation of the acrosome from the nucleus, a wide separation between the cells and many vacuoles were also observed (Fig. 5D). Also, a wide separation between the germ cells was observed (Fig. 5E). The BU-treated group showed damage of Leydig cell with degenerated nucleus, irregular plasma membrane. The cytoplasm showed degenerated mitochondria, many vacuoles and many lysosomes (Fig. 5F).

FIGURE 2. (A–F) Photomicrographs of mice testes in the control and different experimental groups. Masson trichrome staining. (A, B & C) control, Mel and ZnO NPs showing minor collagen fibers (arrowhead); (D) BU group showing dense collagen fibers (arrowhead); (E) BU + Mel group showing minor collagen fiber (arrowhead); (F) BU + ZnO NPs group showing minor collagen fibers (arrowhead).
Electron micrographs of the group treated with BU and supplemented with Mel showed normal regular and thin basal lamina, revealing that Mel reduced the thickness of basal lamina in mice treated with BU. The standard architecture of Sertoli cells resting on the basal lamina with normal nucleus were also seen. All types of spermatogenic cells such as spermatogonia, primary spermatocytes and spermatids were normal structure and cytoplasm contains many mitochondria (Figs. 6A and 6B). Thin and regular basal lamina, normal spermatogonia with normal nucleus, primary spermatocytes and spermatids were normal structure also noticed (Fig. 6C). Round spermatids had normal oval nucleus, the nuclear membrane was thickened from one side forming acrosomal vesicles and minor cells has pyknotic nuclei. The lumen of seminiferous tubules showed increase number of normal sperms with dense stained nuclei covered by acrosome (Figs. 6D and 6E). Leydig cells had regular plasma membrane, nuclei with prominent nucleoli and the cytoplasm showed numerous mitochondria, rough endoplasmic reticulum and lysosomes (Fig. 6F).

Moreover, ultrathin sections of testicular tissues of mice treated with BU and supplemented with ZnO NPs showed marked improvement of spermatogonia, primary spermatocytes, Sertoli cells and spermatids. Thin and regular basal lamina and normal architecture of Sertoli cells resting on the basal lamina with normal nucleus were also seen (Figs. 7A and 7B). The round spermatids had normal nucleus and nucleolus with regular plasma membrane and covered with acrosomal vesicles, cytoplasm contains abundant mitochondria, ribosomes and lysosomes (Figs. 7C and 7E). The lumen of seminiferous tubules showed numerous of normal mature spermatids with normal acrosome and nucleus, cytoplasm contains many

**FIGURE 3.** (A–F) Toluidine blue stained semithin sections in the testes of control and experimental groups. (A, B & C) control, Mel and ZnO NPs showing well organized seminiferous tubules and arrangement of Sertoli cell resting on basement membrane spermogonia, rest on intact basement membrane, large primary spermatocytes, round and elongated spermatids and numerous mature sperms are present in the lumen, normal Leydig cells; (D) BU-treated group showing thin wall of seminiferous tubules, wide intertubular spaces with degenerated interstitial cells and reduced number of Leydig cells, no sperms found in lumen, destroyed most spermogonia and spermatocytes cells; (E & F) Mel or ZnO NPs supplemented after BU group showing recover seminiferous tubules surrounded by regular basement membrane, improve Leydig cells, spermogonia, Sertoli cells, and increased volume of sperms in the lumen.

**Abbreviations and symbols:** Lumen (L), Leydig cells (LC), spermogonia (SG), spermatozoa (SZ), separation of spermatogenic cell from the basement membrane (white arrowhead), Sertoli cells (S) and spermatocyte (SC), spermatid (SD), increased interstitial spaces (arrow), degenerated Leydig cells (curved arrows), no sperm in the lumen (*) and no spermatogenic process (arrowhead).
mitochondria and lysosome (Figs. 7D). Leydig cells had regular plasma membrane, nuclei with prominent nucleoli and the cytoplasm showed numerus mitochondria and lysosomes (Fig. 7F).

**Discussion**

Infertility is a disease of the reproductive system defined by the failure to achieve a clinical pregnancy after one year of regular unprotected sexual practice (Bashir et al., 2009). Male infertility risk factor such as age, smoking, obesity, exercise, diet, electronic devices, stress, scrotal temperature, reactive oxygen species and therapeutic drugs (Olayemi, 2010). Some therapeutic drugs such as alkylating agents (cyclophosphamide and busulfan) alter fertility, change organ structure, sex hormones, function and life quality (Tsuyoshi et al., 2015).

Busulfan (BU) is one of the alkylating agents widely used treating various cancers, leukemia, malignant lymphomas and autoimmune diseases (Gabrielsen and Tănăkut, 2016). Despite the many clinical applications of BU, it has critical complications as spermatogenesis disorder and infertility (Juenke et al., 2011).

Antioxidant agents neutralize free radicals and oxidative stress resulting from chemotherapeutic agents (Simone et al., 2007). Melatonin (Mel) is a free radical scavenger that regulates various molecular pathways, such as inflammation, proliferation, apoptosis, and metastasis during different pathological conditions (Cui et al., 2017). Zinc possesses important properties, including its use in many applications and its flexibility in preparing it in different shapes and properties, making it a magic material (Padmavathy and Vijayaraghavan, 2008). It is an antioxidant and a vital trace element necessary to maintain germ cells. The high Zn concentrations in testis and accessory gonadal glands indicate a critical role in reproduction (Tuncer et al., 2011). ZnO NPs have potent antioxidants and can ameliorate reproductive function (Mohammadi et al., 2017; Torabi et al., 2017).

In the present results, Masson trichrome staining in BU-treated group showed dense collagen fibers at the boundary of seminiferous tubules and in between the tubules due to Busulfan (BU) was the first cytotoxic drug associated with pulmonary toxicity, acute lung injury, chronic intestinal fibrosis that characterized by overgrowth, hardening and is attributed to excess deposition of extracellular matrix components including collagen (Oliner et al., 1961; TA Wynn, 2008). These results were in agreement with Hussein et al. (2015). It could be explained by the released factors from the macrophages and T lymphocytes, stimulating collagen deposition. Moreover, activated macrophages could
stimulate the epidermal growth factor (EGF) which induces granulation tissue formation and fibrosis (Murray and Wynn, 2011). In the present results, Mel. supplemented group after BU, resulted in minor collagen fibers. This consistent with Ilbey et al. (2009); Mel improved histopathological changes and this due to its antioxidant and anti-apoptotic properties (Bejarano et al., 2009; Mirhoseini et al., 2014). Also, the present results showed few of collagen fibers in ZnO NPs supplemented group after BU. These go parallel with Meligy et al., 2019; and this protection due to its antioxidant and anti-apoptotic properties of Zn (Alfi et al., 2015; Maremanda et al., 2016).

It is well known that normal Sertoli and Leydig cell functions provide a principle activation and maintenance of spermatogenesis. In the current work, the ultrastructure study of testicular tissue elucidated that BU altered seminiferous tubules and changed the normal structure of Sertoli cells. These results were consistent with Anan et al. (2018) who observed the impairment of Sertoli cells because of the increased oxidative stress in the testes induced by chemotherapeutic agent. Also, the present results revealed that basal lamina surrounded seminiferous tubules plays an important role in spermatogenesis, which was found thickened and irregular and this thickness would cause a decreased rate of sperm production. Our results were consistent with who stated that stimulation of myoid cells that produce more collagen and extracellular matrix leads to thicken and irregular, basal lamina (Sakr et al., 2012). The previous authors reported that the change succeeded thickened basal lamina in its binding properties, impairing spermatogenesis by limiting access of different hormones and paracrine factors regulating sperm production (Aitken and Roman, 2009). These alterations due to BU exposure caused by over production of ROS. Moreover, degeneration of Leydig cells may impair the function of secreting hormone in BU group; this was in agreement with who revealed that chemotherapeutic drugs increased free radicals, damaging the Leydig cells (Colpi et al., 2004; Hutson, 2006).

On the other hand, Mel has protective role against hazardous effects of BU. In Mel supplemented group after BU showed normal spermatogenic cells, Sertoli cells and thin basal lamina. These results parallel those who reported that Mel restores almost normal basal lamina by decreasing the thickness and provided protection for spermatogenesis.
In another study, Espino et al. (2010) demonstrated that Mel also prevents oxidative stress and apoptosis of mitochondrial pathways. Antioxidant could protect living cells from BU damage by improving the deleterious effect of free radicals, so Mel has an ability of scavenging free radicals and regulate gene transcription for some antioxidant enzymes (Bharti et al., 2011). Also, the present results investigated the protective effects of ZnO NPs against hazardous effects of BU showed marked improvement of spermatogonia, spermatocytes, Sertoli cells, Leydig cells, thin and regular basal lamina. These results agreed with those who reported that ZnO NPs improved the adverse effects of chemotherapeutic agent (Anan et al., 2018). It was showed that ZnO NPs have anti-apoptotic activates and Zn supplementation gives a protective effect against molecules apoptosis (Tian et al., 2009). Authors stated that one of potential mechanisms of decreased apoptosis was suppressing caspase enzyme activity. The other explanation of such findings was that ZnO NPs increased mitochondrial functions resulting in decreased apoptosis- inducing factor and cytochrome c then preventing apoptosis (Kaufmann and Hengartner, 2001). Also, in 2017, Torabi et al. (2017) reported that ZnO NPs have powerful antioxidant properties and can improve the reproductive function.

FIGURE 6. (A–F): Electron micrographs of seminiferous tubules of BU-treated mice supplemented with Mel; (A, B & C) thin basal lamina (BL), normal appearance of Sertoli cell resting on basal lamina (Thick arrow), normal spermatogonia (closed arrow), nucleus of Sertoli cell (N), nucleolus (Nu), normal spermatocyte (curved arrow), mitochondria (M); normal spermatid (zigzag arrow); (D) many sperm with nucleus (N) and acrosome (A), spermatid (zigzag arrow); (E) many elongated sperm with nucleus (N) and acrosome (A); many mitochondria (M); (F) nucleus of normal Leydig cell (N), lysosome (Ly), mitochondria (M), rough endoplasmic reticulum (RER).
Conclusion

In conclusion, chemotherapeutic drugs exert harmful effects on the reproductive system, degenerative changes of the testis of albino mice and arrested spermatogenic activity through several pathways. BU induced histological and ultrastructural damage of the testis. Moreover, the present results could be concluded that Mel or ZnO NPs can protect the testicular tissue against ultrastructural alterations induced by BU by its antioxidant and anti-apoptotic effects.

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Data Availability Statement: The data generated or investigated during this study are obtainable from the corresponding author on reasonable request.

Ethics Approval: All experimental procedures were approved and supervised by the Institutional Ethical Committee for the care and use of laboratory animals in the Faculty of Science, Mansoura University, Egypt. All efforts were made to minimize animal suffering.

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