Comparative Potential of Zinc Sulfate, L-Carnitine, Lycopene, and Coenzyme Q10 on Cadmium-Induced Male Infertility

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The human exposure to toxic chemicals and heavy metals is one of the main predisposing factors contributing to male infertility. Acute exposure to cadmium chloride results in testicular damage and infertility. The purpose of the present study was to investigate and compare the curative effect of coenzyme Q10 (CoQ10), lycopene, L-carnitine (LC), and zinc sulfate against the cadmium-induced infertility in male Wistar rats. Cadmium chloride (0.4 mg/kg/day) was orally administered to rats for three consecutive days. Then, oral administration of different treatments (i.e., LC 100 mg/kg, CoQ10 20 mg/kg, lycopene 4 mg/kg, zinc sulfate 6 mg/kg, and a combination LC-CoQ10 at 500/50 mg/kg) was carried out for 30 days. The impact of different treatments on semen parameters, such as sperm count and motility, testicular antioxidants, and serum testosterone, was determined. Furthermore, the morphometry of epididymis sperms and histopathology of rat testes were also assessed. Cadmium exposure decreased the sperm count, progressive sperm motility, testosterone, superoxide dismutase (SOD), and catalase and reduced glutathione (GSH). It also caused banana sperm tail, bent sperm head, vacuolization of seminiferous tubules, and oligospermia in rat testes. All treatments with nutraceuticals improved sperm count, sperm morphology, serum testosterone, vacuolization of seminiferous tubules, and oligospermia in diseased rats. Treatment with lycopene, LC, and LC-CoQ10 improved progressive sperm motility and other parameters and increased SOD, GSH, and CAT in the rat testes. CoQ10 also increased SOD activity in rat testes’ tissue homogenates. It is concluded from the current study that all nutraceuticals partially improved reproductive toxicity of cadmium. The administration of lycopene and a high-dose combination of LC-CoQ10 were more efficacious in treating cadmium-induced infertility than other treatments. Treatment of cadmium-exposed rats with lycopene, LC, CoQ10, and LC-CoQ10 improved sperm count and motility through reduction of testicular oxidative stress and improving serum testosterone.
1. Introduction

It is estimated that the infertility affects about 15% couples globally and this frequency is gradually increasing [1]. The male infertility is traditionally caused by varicocele, cryptorchidism, infections, obstructive lesions, trauma, tumors, and hormonal and environmental factors. Oxidative stress due to excessive reactive oxygen species (ROS) raises numerous problems related to male infertility, like sperm damage and sperm deformity [2]. The pathophysiological origins of infertility include hypogonadotropic hypogonadism (HH), coital disorders, erectile dysfunction, and ejaculation disorder. In male HH, testes fail to produce androgen and sperms, resulting in congenital or acquired diseases, affecting hypothalamus and/or the pituitary gland. Spermatogenesis and androgen production are maintained and induced in HH by exogenous regulation of gonadotropins, permitting natural fertility in several cases [3]. Clomiphene citrate and selective estrogen receptor modulators (SERM) are mainly used to treat HH [4–6].

Several heavy metals cause the male infertility, such as cadmium, lead, and mercury. Acute cadmium poisoning can lead to hepatic and testicular injury. Ototoxicity, severe testicular hemorrhage, edema, and necrosis have resulted from chronic exposure to cadmium. Cadmium 21 is the analogue of cadmium, which is toxic to humans and animals [7]. Cadmium absorption occurs in human by ingestion or inhalation of the food containing cadmium. Mostly, cadmium present in terrestrial foods is absorbed into the body. Cadmium is used in batteries, metal plating, pigments, and in plastics and alloy industries. Human exposure to cadmium occurs ecologically or by occupational ways. Ingestion of food, drinking water, contaminated soil and dust, cigarette, smoking, and dietary consumption are the main causes of cadmium exposure to human [8]. Cadmium inactivates the enzyme containing sulfhydryl groups. It also causes the uncoupling of oxidative phosphorylation in mitochondria of the cell. Moreover, it competes with other metals (e.g., Zn and selenium) for attachment to metalloenzymes and with calcium for binding sites (e.g., calmodulin on regulatory proteins). Cadmium develops the oxidative stress in the different organs by acute poisoning and causes severe degenerative changes in most tissues leading to osteomalacia, hepatotoxicity, renal toxicity, neurotoxicity, infertility, and cancer [9].

Oxidative stress is implicated in several human diseases including but not limited to atherosclerosis, cancer, diabetes, rheumatoid arthritis, inflammatory bowel disease, Parkinson’s disease, and infertility [10]. Although small amount of ROS is beneficial for fertilization, the excessive amounts of ROS and nitric oxide (NO) promote capacitation and acrosomal reaction, thus adversely affecting spermatogenesis. Even the lipid peroxidation resulting from low level of ROS causes the plasma membrane modification and facilitates the adhesion of sperm-oocyte. These prooxidants are then shifted to the semen and vaginal secretions, which ultimately cause the infertility [11]. Morphologically abnormal spermatozoas are produced by the ROS in the infertile men having low capacity of the antioxidants and less sperm motility. In the seminal plasma of asthenozoospermic men, the ROS activity is associated with the chain-breaking antioxidants [12]. Dietary and endogenous antioxidants including low level of chain-breaking antioxidants (CBA) are indicated to treat reduced spermatogenesis and testicular stress [13].

Coenzyme Q10 (CoQ10) protects from thrombolysis, congestive heart failure (CHF), essential hypertension, renovascular hypertension, and ventricular arrhythmia in addition to stroke, retinopathy, and muscular dystrophy. Moreover, CoQ10 is widely distributed to different body organs and exhibits a long half-life [14]. Lycopene belongs to carotenoids, which are the main sources of vitamin A. Lycopene possesses the antioxidant properties and is naturally found in fruits and vegetables. Lycopene inhibits various human cancer cells, prevents DNA damage, and is more potent than a- and b-carotenes [15].

L-Carnitine (LC) belongs to carnitines, which are highly polar compounds, broadly distributed in nature. Within the male genital tract, carnitines are concentrated in the epididymis and spermatozoa [12]. Carnitines are also frequently used in the treatment of male infertility, chronic renal failure, and erythropoietin resistant anemia [16]. Zinc (Zn) is an essential trace element that acts as a cofactor involved in deoxyribonucleic acid (DNA) profiling and protein synthesis. Furthermore, Zn has also antiapoptotic and antioxidant properties. Zn is useful in the treatment of asthenozoospermic male infertility. It improves the sperm integrity, conception, and pregnancy rates. Zn also plays a pivotal role in lipid metabolic flexibility and stabilization of sperm membrane [17]. Lack of Zn is related to the compromised sperm functioning and enhancement of oxidative stress in seminal plasma. Moreover, Zn is also important in the regulation of various processes, for example, capacitation and the acrosome reaction of sperm for conception and embryonic implantation [18]. The investigation reason behind this study was to assess and compare the efficacy of lycopene, LC, ZnSO4, and CoQ10 against cadmium-induced infertility in male rats.

2. Materials and Methods

Diethyl ether, phosphate buffer solution (PBS), paraformaldehyde, ethanol, xylene, paraffin, hematoxylin, 10% tricyclic acid (TCA), diethiobis-2-nitrobenzoic acid (DTNB), pyrogallol solution, potassium phosphate, hydrogen peroxide (H2O2), and cadmium chloride were obtained from Sigma Aldrich (Germany). LC and zinc sulfate (ZnSO4) were obtained from Selmore Pharmaceuticals (Pakistan). CoQ10 and lycopene were acquired from Bionext Pharmaceuticals, Pakistan. Eosin-Nigrosin stain was obtained from Hitech Specialties Solutions, India. Centrifuge (800D–4000 rpm), microtome, compound microscope (Germany), Eppendorf tubes, UV-visible spectrophotometer (UV-1601, Shimadzu), and computer-assisted sperm analysis (CASA) system (Minitube®, Germany) were used in the study.

2.1. Experiment Design. Seventy male Wistar rats of 10 weeks old and weighing 150 ± 20 g were used in the study.
The animals were acquired from Riphah Institute of Veterinary Sciences, Lahore, Pakistan, and then acclimatized for two weeks in well-ventilated animal house of RIPS, Lahore, Pakistan. The animal study was approved and conducted according to the guidelines of the Institutional Ethical Committee of Riphah International University (Reference no. REC/RIPS-LHR/2017044). Animals were kept at a temperature of 26 to 30°C and humidity of 30 to 70%, maintained according to the National Institute of Health (NIH) guidelines. These rats were randomly divided into seven groups, with each group comprising of ten animals. All animal groups were separately placed in steel cages. To demonstrate the curative effect of different nutraceuticals against male infertility, the LC, CoQ10, lycopene, and ZnSO4 were orally given to rats for 30 days previously treated with cadmium chloride. Afterwards, the sperm, biochemical, hormonal, and histological parameters were determined. Different treatments given to various groups are given in Table 1.

Cadmium was administered orally to rats for three times on consecutive days. Then, different treatments with nutraceuticals were orally administered for 30 days. 24h after administration of last doses, the rats were anesthetized with diethyl ether, the blood samples were collected from heart puncture, and their peritoneal cavities were opened through a low transverse incision to remove the testes of rats from the control and treatment groups immediately. The blood taken in plain tubes was centrifuged (3000 rpm) at 4°C for 10 min to prepare the serum from each animal and stored it at −20°C until assayed. The harvested testes specimens were preserved in paraformaldehyde for histological analysis [23].

2.2. Epididymis Sperm Count, Viability, and Motility. Determination of the motility was done by the CASA system. The contralateral epididymis was transferred to another petri dish containing PBS followed by the diffusion method in which sperms diffused from epididymis tubes. The layers of connective tissue were removed at the juncture between the proximal and distal regions of the cauda epididymis using Jeweler’s forceps. Approximately within the 30 S, the study samples were picked up to avoid variations that occur during late samples. The sperm viability was determined by staining with Eosin-Nigrosin [24]. Sperm motility parameters such as total motility, progressive motility of sperms (PMS), progressive fast motility (PFM), progressive slow motility (PSM), progressive circular motility (PCM), and linear motility of sperms (LMS) were determined.

2.3. Morphological Studies. Abnormal spermatozoa in sperm cells were evaluated after obtaining from the cauda epididymis and calculated according to the Wyrobk method by counting and identifying 1000 sperm heads in three samples per animal. The abnormalities observed in spermatozoa were categorized into four groups according to their frequency of appearance.

Spermatozoa were observed for flagellum abnormalities such as dual, bifid, tangled or wrong insertion of the flagellum, abnormalities in head morphology such as macrocytoplasm, spermatozoa having macrocephaly; acephalous, strange head shape, and alterations in acrosome [25].

2.4. Histological Studies. Tissues of testes were preserved in a solution of 4% paraformaldehyde in 0.1 M phosphate buffer. Then, tissues were detached, dried by ethanol and then sanitized with xylene, and lastly embedded in paraffin wax. Microtome was used for preparing tissue section of 5 μm thickness. Hematoxylin-eosin staining was used for countermarking of tissues, which were observed under light microscope for examination and photomicrographs [26].

2.5. Hormonal Assay. The serum obtained from the collected blood was used to determine the hormone profile. Testosterone level was measured using commercially available immunnoassay enzyme linked immunoassay (Rocky Mountain Diagnostics, Inc., USA, according to manufacturer’s specs).

2.6. In Vivo Antioxidant Activity. Enzymatic antioxidants such as glutathione (GSH), catalase (CAT), and superoxide dismutase (SOD) were assessed in rat testes by the methods described previously. To prepare tissue homogenate, harvested testicular tissues were washed with ice-cold normal saline; 1 g tissue was minced with 2-3 ml PBS via tissue homogenizer and then diluted with PBS to prepare 10% w/v tissue homogenate. The resultant mixture was centrifuged at 4000 rpm for 20 min and the supernatant was drawn into a sterile tube and oxidative stress markers were determined immediately. Moreover, protein content in testicular tissue homogenate was determined by Lowry methods [27].

2.7. GSH Level. For GSH determination, a previously described method was followed [27]. A 1 ml of tissue homogenate supernatant was taken and added to 1 ml of 10%
TCA, 4 ml of sodium phosphate solution, and 0.5 ml DTNB. The absorbance of resultant solution was taken at 412 nm with UV-Vis spectrophotometer.

2.8. SOD Activity. An earlier described method was used to determine SOD activity with little modifications [10]. A 100 ml tissue homogenate supernatant was added to 2.8 ml PBS (pH of 7.4) and 100 ml pyrogallol solution. The absorbance of the mixture was taken at 325 nm. One unit of SOD was described as the amount of enzyme responsible for 50% inhibition of pyrogallol autoxidation per ml of the assay solution.

2.9. CAT Activity. The activity of CAT was assessed by H2O2 method. A 50 ml tissue homogenate was added to 1.95 ml potassium phosphate solution and (50 mM) 1 ml of H2O2. Change in absorbance per min was determined at 240 nm to estimate CAT activity [28].

2.10. Statistical Analysis. The obtained results were tabulated and presented as mean ± standard deviation (SD) and evaluated by GraphPad Prism software version 7.01 for one-way analysis of variance (ANOVA) followed by a post hoc test. Bonferroni’s multiple comparison tests showed the significance level of each test. Bonferroni’s multiple comparison tests showed the significance level of each test. Bonferroni’s multiple comparison tests showed the significance level of each test.

3. Results

3.1. Epididymis Sperm Count. After 30 days of dosing, the sperm count was observed in different groups, which indicated that the disease control group had shown significantly reduced sperm count (1.76 ± 0.67 million/ml) as compared to the normal control rats (18.55 ± 0.95 million/ml). Treatment with different nutraceuticals significantly improved the sperm count in cadmium-treated rats. The sperm count in lycopene-treated group (18.33 ± 0.64 million/ml) was the highest among all treated groups which was followed by ZnSO4 (17.95 ± 0.47 million/ml), LC (16.77 ± 0.74 million/ml), LC-CoQ10 (15.24 ± 0.64 million/ml), and CoQ10 (14.58 ± 0.85 million/ml) treated groups. Moreover, the sperm count in all nutraceutical treated rats resulted in a rise as compared to that of normal control rats except CoQ10 treated rats, which showed statistically reduced sperm count in comparison to the normal control group. The effect of nutraceuticals on sperm viability in cadmium exposed rats was also significant. All nutraceuticals increased the sperm viability in diseased rats. The effect of different nutraceuticals on sperm count and viability of cadmium exposed rats is shown in Figure 1.

Results are presented as mean ± SD (n = 10). *** and ## showed statistical significance as compared to disease control and normal control at P < 0.001 and P < 0.01, respectively.

3.2. Motility of Sperms. The CASA analysis revealed the motility of sperms in the cadmium-exposed rats. Treatment with nutraceuticals had variable effect on sperm motility parameters such as total motility, PMS, PFM, PCM, and LMS. The effect of various treatments on cadmium-induced infertility in rats is shown in Figure 2.

It was found that the total motile sperms in the disease control group were significantly declined (P < 0.0001) in comparison to the normal control rats. Administration of lycopene, LC, and LC-CoQ10 increased the percentage of motile sperms in cadmium-treated rats to varying degree. Lycopene and LC-CoQ10 had the most pronounced positive effect on sperm motility followed by the LC.

Administration of cadmium significantly reduced the percentage of PMS in disease control rats as compared to normal control rats. It was revealed that the administration of lycopene, LC, and LC-CoQ10 had a profound positive effect on PMS in rats preexposed to cadmium. Treatment with lycopene had the most prominent effect on PMS in rats while LC and LC-CoQ10 had a comparative effect on PMS of cadmium-exposed rats.

It was found that cadmium exposure in rats significantly decreased the percentage of PFM sperms as compared to normal rats. Administration of CoQ10 and LC-CoQ10 failed to bring about any significant increase in PFM in cadmium-exposed rats. However, treatment with lycopene, LC, and ZnSO4 significantly improved the percentage of PFM in rats. Administration of lycopene exhibited the most pronounced increase of PFM in cadmium-exposed rats as shown in Figure 2. Cadmium exposure also significantly decreased PSM in treated rats in contrast to the normal control group. Treatment with LC-CoQ10 exhibited the most significant increase in PSM in comparison to the disease control group followed by lycopene. However, the administration of CoQ10, LC, and ZnSO4 did not cause any increase in the PSM of cadmium-exposed rats, as shown in Figure 2.

It was evident that the exposure to cadmium had not brought about any statistically significant decline in the percentage of PCM in the disease control group. Administration of ZnSO4 resulted in a significant (P < 0.001) increase in PCM in contrast to disease control. However, the treatments with lycopene, CoQ10, LC, and LC-CoQ10 had insignificantly affected the PCM induced by cadmium. Furthermore, cadmium exposure did not cause any significant decline in the percentage of LMS in rats. However, administration of CoQ10 and LC-CoQ10 resulted in a statistically significant increase in the LMS of cadmium-exposed rats, as shown in Figure 2.

Results are presented as mean ± SD (n = 10). ****, ***, * showed statistical significance as compared to disease control at P < 0.0001, 0.001, 0.01, and 0.05, respectively.

3.3. Sperm Morphology. In morphological studies, general observation was made to determine various abnormalities of
spermatozoa in all groups. The abnormalities appeared in the flagellum and head of rat sperm exposed to cadmium. Bent sperm tail and banana head were observed in several sperms of the disease control group. Such abnormalities were not evident in groups treated with lycopene, LC, ZnSO₄, CoQ₁₀, and LC-CoQ₁₀. The effect of different treatments on sperm morphology is shown in Figure 3.

3.4. Effect on Testosterone Level. The serum level of testosterone in normal control group (2.39 ± 0.507 ng/ml) was higher as compared to cadmium-exposed rats (1.46 ± 0.096 ng/ml). All nutraceuticals significantly improved the level of serum testosterone in cadmium-exposed rats. The treatment with lycopene, LC, and LC-CoQ₁₀ resulted in the most pronounced effect on serum testosterone level in cadmium exposed rats and were significantly comparable to a normal control group. However, the treatment with CoQ₁₀ and zinc sulfate resulted in lower serum testosterone level as compared to normal control rats. The effect of different nutraceuticals on the serum testosterone level of cadmium exposed rats is shown in Figure 4.

Results are presented as mean ± SD (n = 10). ***, and * showed statistical significance as compared to disease control and normal control at P < 0.001 and P < 0.01, respectively.

3.5. Oxidative Stress Parameters in Rat Testes. Administration of cadmium was followed by treatment with different nutraceuticals and then rat testes’ tissue homogenates (10% w/v) were tested for antioxidant parameters, such as SOD, CAT, and GSH.

3.6. SOD Activity. In the current study, the activity of SOD was found to be significantly higher in normal control group (26.13 ± 0.78 U/mg of protein) as compared to the disease control group (13.29 ± 1.78 U/mg of protein). Treatment with LC-CoQ₁₀ (22.28 ± 2.255 U/mg of protein) showed the highest SOD activity among all nutraceuticals, which was statistically comparable to the normal control group. Treatment with lycopene and LC also exhibited statistically higher SOD activity than the disease control group. The activity of SOD exhibited by LC-CoQ₁₀ was insignificantly different from the normal control group, as shown in Figure 5(a).

Results are presented as mean ± SD (n = 10). ***, ***, **, and * showed statistical significance as compared to disease control at P < 0.0001, 0.001, 0.01, and 0.05, respectively, while ### and ## showed statistical significance compared to normal control at P < 0.0001 and 0.01, respectively.

3.7. CAT Activity. It was found that the activity of CAT in normal control group (1.46 ± 0.096 U/mg of protein) was statistically higher as compared to the disease control group (0.42 ± 0.013 U/mg of protein). The activities of CAT in lycopene, LC, and LC-CoQ₁₀ groups were significantly elevated than the disease control group. However, LC, lycopene, and LC-CoQ₁₀ failed to normalize the CAT activity. The treatment with ZnSO₄ and CoQ₁₀ did not improve the CAT activity in comparison to the disease control group as shown in Figure 5.

3.8. GSH Level. It was found that the GSH content in the normal control group (3.02 ± 0.123 nmol/ml) was significantly higher as compared to disease control group (2.39 ± 0.318 mg/g). Administration of lycopene, LC, and LC-CoQ₁₀ significantly increased GSH in diseased rats. The
Figure 2: Effect of lycopene, L-carnitine, coenzyme Q10, and zinc sulfate on sperm motility of cadmium-exposed rats. Results presented as mean ± S.D. (n = 10). ****, ***, **, and * showed statistically significant as compared to disease control at P < 0.0001, 0.001, 0.01, and 0.05 respectively. (a) Total motility. (b) Progressive motility. (c) Progressive fast motility. (d) Progressive slow motility. (e) Progressive circular motility. (f) Local motility.
effect of ZnSO₄ on GSH was negligible in comparison with the disease control group. Moreover, the treatment with CoQ10 significantly reduced the GSH content in cadmium-exposed rats. The effect of treatment with lycopene, L-carnitine, CoQ10, and ZnSO₄ on cadmium-induced reduction in GSH content in rat testicular tissue homogenate is as shown in Figure 5.

3.9. Histological Studies. Photomicrograph of testicular histology of the normal control rats showed intact and normal structure germinal epithelium. However, treatment with cadmium chloride resulted in vacuolization in germinal epithelial cells and oligospernia in the testes of diseased rats. The fissures were observed clearly, and the seminiferous epithelium was incomplete. Treatment with lycopene, CoQ10, LC, ZnSO₄, and LC-CoQ10 resulted in amelioration of cadmium toxicity. All treatments with CoQ10, LC, ZnSO₄, and LC-CoQ10 showed only partial improvement in testicular lesion caused by cadmium chloride. The damage of histological structures in the LC-CoQ10 and L-carnitine treated group was even worse than that in other nutraceutical treatment groups. The effect of treatment with lycopene, L-carnitine, CoQ10, and ZnSO₄ on cadmium-induced histological damage to rat testicles is shown in Figure 6.

4. Discussion
Several drugs, heavy metals, and environmental chemicals have toxic effects on human reproductive system. Human
In morphological study, spermatozoa abnormalities, such as vacuolization and oligospermia, were observed in cadmium-treated rats. Animals treated with lycopene, CoQ10, LC, and ZnSO4 showed no abnormal spermatozoa in contrast to the disease control group. A previous study showed that treatment of cyclophosphamide-induced infertility Sprague-Dawley male adult rats with lycopene normalized the sperm morphology [36]. Previous studies confirm that the lycopene, LC, CoQ10, and ZnSO4 and CoQ10 supplementations reduced morphological anomalies in diseased or injured animals [37].

The current study also showed the remarkable inhibition in serum testosterone level by cadmium exposure. The treatments with lycopene, LC, CoQ10, and ZnSO4 and CoQ10 significantly raised the level of testosterone, sperm count, sperm motility, and antioxidant enzyme concentrations after acute poisoning of cadmium as compared to control group [35]. Testosterone level is an important factor in the determination of male infertility. A previous study conducted on LC showed the similar effect on serum testosterone level [20]. A study conducted on LC showed a significant increase in plasma testosterone level in male adult albino mice that were treated with cadmium chloride at dose 0.35 mg/kg [38]. Similarly, a study conducted on ZnSO4 in improving fertility also supported that Zn had a positive effect on testosterone level and semen quality in infertile 7-week-old Sprague Dawley rats [39]. CoQ10 also reversed the testicular damage associated with cadmium toxicity by enhancing the levels of LH and testosterone in rodents likewise the present study [39].

It is found that the reduction in the amount of glutathione and protein containing sulphydryl group is associated with cadmium exposure, which eventually leads to increased ROS production. This enhanced production of ROS increases the lipid peroxidation, excretion of urinary lipid metabolites, DNA damage, membrane damage, altered gene expression, and apoptosis [31]. The in vivo antioxidants such as SOD, CAT, and GSH were reduced by exposure to cadmium in diseased rats. Treatment with LC, lycopene, ZnSO4, and LC-CoQ10 raised the activity of SOD in rat testes as compared to the disease control group. Moreover, the CAT activity and GSH content were increased by treatment with LC and lycopene and LC-CoQ10 in testicles of cadmium-exposed rats. It is previously found that LC reduced oxidative stress through reduction of malondialdehyde (MDA) and increasing GSH in lipopolysaccharide exposed rats at 500 mg/kg single dose [40]. The in vivo antioxidants (SOD, CAT, and GSH) play a pivotal role in the reduction of oxidative stress and thus decrease the chances of infertility. In case of histopathological study, disease control group showed vacuolization in germinal epithelial cells whereas the administration of LC, lycopene, CoQ10, ZnSO4, and LC-CoQ10 in infertile rats showed no deterioration as evident in previous study [41].

Previous study on titanium dioxide and benzopyrene-induced toxicity reported that its testicular toxicity was treated by lycopene through reduction of oxidative stress and suppressing the apoptosis to protect germinal cells [42, 43]. Furthermore, lycopene has demonstrated the anti-
infertility effect against drug-induced testicular damage caused by gentamicin, cisplatin, and deltamethrin mainly through the reduction of lipid peroxidation and oxidative stress and increased production of testosterone [44]. Furthermore, lycopene was useful to improve infertility caused by coal burning related fluorosis through oxidative stress associated with Jun N-terminal kinase (JNK) and extracellular signal-regulated protein kinase (ERK) pathway [45].

Khe LC has also shown improvement of oligospermia against busulfan-induced oligospermia through reduction of oxidative stress [46]. Moreover, LC-ameliorated trazodone induced testicular damage through modulating autophagy, reduction of oxidative stress, and preventing inflammation [47]. LC also corrects testicular perfusion to prevent ischemic injury to testicular tissues [48]. Prevention of apoptosis is also one of the mechanisms through which LC might have demonstrated anti-infertility effect against toxicants through modulation of proapoptotic and anti-apoptotic genes, such as Bcl-2, BAX, and Caspase-3 [49].

Posttreatment of diseased rats with CoQ10 failed to bring

**Figure 5:** Effect of lycopene, L-carnitine, coenzyme Q10, and zinc sulfate on oxidative stress in testicles of cadmium-exposed rats. Results presented as mean ± S.D. (n = 10). **, ****, *** and * showed statistically significant as compared to disease control at P< 0.0001, 0.001, 0.01 and 0.05 respectively, while **** and ## showed statistically significant compared to normal control at P<0.0001 and 0.01 respectively. (a) Superoxide dismutase. (b) Catalase. (c) Reduced glutathione.
out an increase in the antioxidants such as GSH, CAT, and SOD, which is in contrast to a previous study that demonstrated the preventive effect of CoQ10 against cadmium-induced infertility had been mediated through reduction of oxidative stress. This apparent contrast might be due to preventive treatment of cadmium-exposed rats with CoQ10 in the previous study [19]. However, estimation of MDA, an important indicator of lipid peroxidation, must also be carried out in rats’ testes exposed to cadmium as cadmium has been found to increase lipid peroxidation in different animal organs [50].

5. Conclusion

In conclusion, the present study showed that the male infertility caused by exposure to cadmium was associated with increased sperm immobility, reduction in sperm count and viability, damage to testicular histology, and increase of
oxidative stress. Daily administration of LC 100 mg/kg, lycopene 4 mg/kg, ZnSO\textsubscript{4} 6 mg/kg, and LC-CoQ10 500/50 mg/kg improved the sperm parameters, serum testosterone, testicular histology, and oxidative stress in male rats to varying degree. Administration of individual and combination treatment of nutraceuticals to cadmium-exposed rats improved cadmium chloride-induced toxicity through the reduction of oxidative damage and increase serum testosterone. Further studies must be carried out to evaluate the ameliorating effect of these chemicals against other endocrine-disrupting effects of cadmium. These nutraceuticals must also be evaluated individually and in combination with each other against infertility caused by heavy metals and other endocrine disruptors.

Data Availability
The data used to support this study are available from the corresponding author upon request.

Ethical Approval
The animal study was approved and conducted according to the guidelines of the Institutional Ethical Committee of Riphah International University (Reference no. REC/RIPS-LHR/2017-044). The animals were kept and maintained in the experiment according to the National Institute of Health (NIH) guidelines.

Conflicts of Interest
The authors declare that they have no conflicts of interest.

Authors’ Contributions
MFA, AS, and AI planned the study. AI, MFA, AS, MZ, AR, and MHR carried out the experimental work. MFA, MZ, and GMA analyzed the data. All authors contributed to the article writing process and approved the final version of the manuscript.

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