EFFECTS OF TAURINE AGAINST HISTOMORPHOLOGICAL AND ULTRASTRUCTURAL CHANGES IN THE TESTES OF MICE EXPOSED TO ALUMINIUM CHLORIDE

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The aim of this study was to investigate the protective effects of taurine against histomorphological and ultrastructural changes in the testes of Swiss albino mice caused by acute in vivo exposure to AlCl₃. Light microscopy revealed that a single intraperitoneal (i.p.) dose of AlCl₃ (25 mg kg⁻¹ Al³⁺) was associated with sloughing, tubular atrophy, germ-cell degeneration, and foci of Leydig cell hyperplasia. In addition, transmission electron microscopy showed a destruction of inter-Sertoli cell tight junctions, apoptotic cell death of spermatogonia and primary spermatocytes, various types of abnormalities in spermatid morphology, accumulation of lipid droplets, reduction of the smooth endoplasmic reticulum (sER), and mitochondrial damage in Leydig cells. Taurine post-treatment at i.p. dose of 1 g kg⁻¹ diminished these changes and significantly reduced the number of affected tubules compared to Al-poisoned mice. This is the first study to evidence that taurine protects against pathological changes in the testicular tissue of Al-treated mice.

KEY WORDS: histopathology, male reproductive system, spermatogenesis, toxic metal, ultrastructure
acid and that it is a beta-amino acid as opposed to alpha-amino acid. Taurine is known to participate in several metabolic actions, including formation of bile salts in the liver, membrane stabilization, osmoregulation, and modulation of neurotransmission and intracellular calcium influx (18). Several studies (19-22) have demonstrated the therapeutic effects of taurine against heavy metals and some drugs (23-24), but its action against Al-induced testicular damage has not been studied so far. The aim of this study was to fill in this gap by looking into the therapeutic effects of taurine against Al overdose through histopathological and ultrastructural changes in testis cells of mice.

Figure 1 Photomicrographs of mice testicular cells stained with haematoxylin and eosin. (A) Control group showing regularly arranged seminiferous tubules and interstitium. (B-D) Al-treated group: note tubular atrophy (asterisk, in B), arrows point to sloughing of degenerated germ cells into tubule lumen, LC: Leydig cells, V: tubule vacuolisation (in C), focal hyperplasia of Leydig cells (LC) is seen adjacent to damaged tubule (in D). (E) Al+TAU group showing almost normal spermatogenesis. Arrow illustrates sloughing of seminiferous epithelium. (F) TAU alone group is comparable to control. Scale bar: 200 μm (A-F).

MATERIALS AND METHODS

Chemicals

AlCl₃, taurine, and other reagents used in this experiment were supplied by Sigma-Aldrich Corporation (St. Louis, MO, USA).

Animals and treatments

Male Swiss albino mice weighing 15 g to 20 g were procured from the College of Veterinary Medicine, King Faisal University, Saudi Arabia. The animals were housed in polypropylene cages and
maintained under controlled temperature (25±2 °C), relative humidity (50±15) %, and 12/12-hour light/dark cycle. The animals had free access to standard pellet diet and drinking water. After a 10-day acclimatisation, the animals were randomly assigned to four groups with five animals each. The control group received a single i.p. injection of physiological saline. The Al group received an i.p. injection containing 25 mg kg⁻¹ of Al³⁺. In addition to Al³⁺, the Al+TAU group received a single i.p. dose of 1 g kg⁻¹ of taurine, shortly after AlCl₃ injection. The TAU group received a single i.p. injection of taurine alone (1 g kg⁻¹). The doses were selected based on earlier in vivo studies (19, 25). Mice from each group were killed by cervical decapitation under light ether anaesthesia 24 h after injection. Experimental design conformed to the NIH standards for the care and use of laboratory animals.

**Light microscopy**

Testicular tissue samples were immediately fixed in 10 % buffered formalin, dehydrated in ascending grades of ethanol, and embedded in paraffin. Five to six micrometer thick sections were sliced using a rotary microtome, stained with haematoxylin and eosin, and examined under a light microscope (Olympus America Inc., Center Valley, PA, USA). Quantitative histopathological analyses of seminiferous tubules was blinded according to the method of Sayim (26) and included 100 tubules per animal to be categorised as follows: “normal”, “sloughing” with broken cellular association, “atrophic” with scarce or no germ cells, and “germ-cell degeneration” with multinucleated giant cells.

**Electron microscopy**

Small pieces of 1 mm thick testes samples were fixed in 2.5 % glutaraldehyde in sodium cacodylate buffer (pH 7.2) for (24 to 48) h, postfixed for 2 h in 1 % osmium tetroxide solution at 4 °C, dehydrated in graded ethanol series, and embedded in Epon-araldite mixture (Embed-812). Ultra-thin sections (40 nm to 60 nm thick) were sliced with a diamond knives using a Leica EM UC6 ultramicrotome (Leica Co., Austria), transferred to 200 mesh copper grids, and double-stained with 2 % uranyl acetate and lead citrate. Grids were viewed on a transmission electron microscope (Jeol JEM 1011, Jeol Ltd., Japan) operated at 80 kV.

**Statistics**

Data obtained with histopathological analyses were evaluated with the Kruskal-Wallis variance analysis and post-hoc Mann-Whitney U-test using SPSS software (version 17.0; SPSS Inc., Chicago, IL, USA). A p value <0.05 was considered significant.

**Figure 2 Transmission electron micrographs of control mice testicular cells.** (A) Sertoli cell (Sc), spermatogonia type B (SgB), primary spermatocyte (1ry Sp) with a well-developed Golgi complex (Gx), and round spermatids (RS). Tight junctions between adjacent Sertoli cells are indicated by arrow. (B) Round spermatids (RS) and elongated spermatids (ES) in the adluminal compartment. (C) Leydig cell with few lipid droplets (L), numerous mitochondria (M) and abundant smooth ER (sER). N: nucleus. Scale bars: 10 μm (A), 5 μm (B), 2 μm (C).
Figure 3 Transmission electron micrographs of the testicular cells in Al-treated mice. (A) Spermatogonia type B (SgB) displays morphological characteristics of apoptosis, including cell shrinkage, blebbing or swelling of nuclear envelope, and excessive chromatin condensation. Also note damaged primary spermatocytes (1ry Sp) with notable swelling of the nuclear envelope (Ne) and dilatations of endoplasmic reticulum (ER). Inter-Sertoli cell tight junctions show numerous membrane discontinuities (arrow). Sc: Sertoli cell, arrowhead points to a myelin figure, (*) indicates phagocytic vesicle, SgA: spermatogonia type A. (B) Apoptotic primary spermatocyte (1ry Sp) presenting large bodies of condensed chromatin (arrow), swelling of nuclear envelope (Ne), and enlargement of ER membranes. (C) Round spermatids with abnormal positioning of many acrosomal vesicles (arrows). Elongated spermatids (arrowheads) can be seen embedded in the Sertoli cell (Sc) cytoplasm, which also contains swollen mitochondria with degenerated or missing cristae (M). (D) Early-stage spermatids with deleterious nuclear changes. Note intense chromatin density in spermatid nucleus (arrow). 1ry Sp: primary spermatocyte, RS: round spermatids. (E) Elongated spermatids at the initial stage of chromatin condensation. Note various nuclear invaginations in acrosome-deficient spermatids (thick arrows), wide subacrosomal space (thin arrow), and sharply pointed acrosome (arrowhead). 1ry Sp: primary spermatocyte, Sc: Sertoli cell, M: mitochondria, Ly: lysosomal elements. (F) Elongated spermatid (ES) near the tubule lumen still shows markedly dilated ER. Sc: Sertoli cell; (G) Leydig cell showing accumulation of large lipid droplets (L). Note disintegrated mitochondria (M), large cytoplasmic vacuoles (V), and nucleus (N) with perinuclear chromatin condensation. Scale bars: 5 μm (A, D, E, G, H), 2 μm (B, F), 10 μm (C).
containing spermatic cells at different stages of maturation. Leydig cells presented as scattered clusters in the interstitial tissue (Figure 1A). In the Al-treated mice, the seminiferous tubules were mostly irregular in shape, and some of the tubules were atrophied with moderate to severe loss of spermatogenic cell layers (Figure 1B). Degenerated cellular material with darkly stained nuclei was sloughed into the lumen of tubules, indicating a breach in the blood-testis barrier (Figure 1C). The seminiferous tubules displayed focal areas of vacuolar degenerative changes in the cytoplasm of spermatogenic cells and in the Sertoli cells. Foci of Leydig cell hyperplasia were also observed (Figure 1D). Acute lesions caused by Al were substantially remedied by taurine in the Al+TAU group (Figure 1E). The number of affected tubules was significantly lower than in the Al group (Table 1). Tubular atrophy and cellular degeneration were not seen in the seminiferous tubules, but a mild detachment (i.e., sloughing) of germ cells was still noted. The testicular histology in mice treated with taurine alone was normal and without changes in germ cell shape and organisation (Figure 1F).

Ultrastructural changes

In control mice, electron micrographs showed the normal structure of seminiferous tubules (Figure 2A-B). Sertoli cells were identified by their large indented nuclei, numerous mitochondria, rough endoplasmic reticulum (rER), and lipid inclusions. These Sertoli cells, closely connected by many tight junctions, separated spermatogonia from primary spermatocytes. Two types of spermatogonia could be distinguished: type A spermatogonia, with ovoid and lightly stained nuclei, and type B spermatogonia, with smaller, spherical nuclei of a more electron-dense nucleoplasmic matrix. In primary spermatocytes, the nuclei had fine chromatin accumulation and the cytoplasm was copious with a well-developed Golgi complex and aggregated mitochondrial structures. The adluminal compartment of seminiferous tubules contained round spermatids in the Golgi phase and in different stages of acrosomal cap formation, elongations of condensed nuclei, and normal tail formation during spermiogenesis. Leydig cells in the control group (Figure 2C) had spherical euchromatic nuclei, many mitochondria with tubular cristae, abundant smooth endoplasmic reticulum (sER), and a few lipid droplets.

RESULTS

Histopathology

The testes of control mice had a normal histological pattern, with well-organised seminiferous tubules containing spermatic cells at different stages of maturation. Leydig cells presented as scattered clusters in the interstitial tissue (Figure 1A). In the Al-treated mice, the seminiferous tubules were mostly irregular in shape, and some of the tubules were atrophied with moderate to severe loss of spermatogenic cell layers (Figure 1B). Degenerated cellular material with darkly stained nuclei was sloughed into the lumen of tubules, indicating a breach in the blood-testis barrier (Figure 1C). The seminiferous tubules displayed focal areas of vacuolar degenerative changes in the cytoplasm of spermatogenic cells and in the Sertoli cells. Foci of Leydig cell hyperplasia were also observed (Figure 1D). Acute lesions caused by Al were substantially remedied by taurine in the Al+TAU group (Figure 1E). The number of affected tubules was significantly lower than in the Al group (Table 1). Tubular atrophy and cellular degeneration were not seen in the seminiferous tubules, but a mild detachment (i.e., sloughing) of germ cells was still noted. The testicular histological architecture in mice treated with taurine alone was normal and without changes in germ cell shape and organisation (Figure 1F).
collapsed cristae, and their cytoplasm contained many myelin figures and large phagosomes. Inter-Sertoli cell junctions were completely lacking or partly deleted (Figure 3A). Spermatogonia and spermatocytes showed morphological changes typical of apoptosis, including cell shrinkage, membrane blebbing, and excessive chromatin condensation (Figure 3A-B). Distortions of developing spermatids were also common after Al treatment. In some slides, the acrosomal vesicle or cap was dislocated into spermatid nucleus (Figure 3C). Round spermatid nuclei had more dense patches of heterochromatin, and in cases of severe structural damage chromatin was fully condensed (Figure 3D). A variety of spermatids, including round and elongated cells, had oddly shaped nuclei with acrosome defects (Figure 3E). This was seldom seen in control spermatids. Near to the tubule lumen, deformed spermatids had a markedly dilated endoplasmic reticulum (ER) (Figure 3F). In the interstitium, the cytoplasm of Leydig cells (Figure 3G) was almost filled by lipid droplets, whereas smooth endoplasmic reticulum (sER) decreased. Several Leydig cells frequently presented condensed chromatin at the nuclear margin, vacuolisation of the cytoplasm, and swollen, vacuolated mitochondria with disarranged or discontinuous cristae.

The testicular ultrastructure of animals treated with Al+TAU showed a significant improvement over the Al group. There were normal Sertoli cells, spermatogonia, spermatocytes, round spermatids with intact acrosomal cap, and elongated spermatids in the tail formation stages (Figure 4A-B). In addition, most Leydig cells appeared normal, similar to control (Figure 4C).

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### Table 1 Percentage of tubular histopathologies in the testes of mice exposed to different treatment conditions

<table>
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<tr>
<th>Parameters</th>
<th>Group</th>
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<tbody>
<tr>
<td></td>
<td>Control 25 mg Al&lt;sup&gt;3+&lt;/sup&gt;</td>
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<tr>
<td>Intact</td>
<td>97.3±0.3</td>
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<tr>
<td>Sloughing</td>
<td>2.7±0.3</td>
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<tr>
<td>Tubular atrophy</td>
<td>–</td>
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<td>Germ-cell degeneration</td>
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Statistical significance of data obtained in histopathological analyses was evaluated by Kruskal-Wallis variance analysis and post-hoc Mann-Whitney U-test.

*<sup>p</sup>< 0.05 vs the control group.

*<sup>p</sup>< 0.05 vs the Al+TAU group.

*<sup>p</sup>< 0.05 vs the TAU group.

**DISCUSSION**

Many toxic metals damage testicular germinal epithelium (27-29). Our main findings in Al-treated mice were deformations of the Sertoli cells, epithelial sloughing, tubular atrophy, and abnormal germ cells. These observations confirm and extend previous reports on gonadotoxic effects of Al in male animal models (30-33). According to Hess and Nakai (34), sloughing of immature germ cells is caused by a disruption of microtubules and intermediate filaments of the Sertoli cells. Kim et al. (35) reported that downregulation of cell adhesion proteins such as cadherin in the Sertoli cells increased sloughing of seminiferous epithelial cells, which is likely to lead to tubular atrophy (36).

The morphological hallmarks of apoptosis, which have been detected in spermatogonia and primary spermatocytes after Al administration, might primarily result from microtubule targeting and mitotic arrest, as described before (37, 38). Al treatment also elicited broad cytotoxic effects in the Leydig cells of mice. The presence of multiple lipid droplets in these cells could eventually lower secretory activity, probably by decreasing the use of free cholesterol for steroidogenesis (39-41). Two recent studies (42, 43) have shown that intact mitochondria with active respiration are essential for LH-induced Leydig cell steroidogenesis. According to Kumar et al. (44) reported large bioenergetic deficits, mitochondrial dysfunction, and depletion of cellular ATP in different regions of rat brain after chronic Al exposure. In theory, the
mitochondrial cytopathy observed in our study by electron microscopy of Leydig cells could lead or contribute to low testosterone secretion by these cells. Besides, the state of microtubule polymerisation and/or tubulin can influence steroidogenesis, as observed in rat Leydig cells (45, 46).

An alternative way for Al to suppress testosterone is to induce nitric oxide (NO) (11). Inhibition of LH-stimulated steroidogenesis may be reinforced by NO in Leydig cells (47). A well characterised consequence of stress induced by testicular NO is the decrease in steroidogenic enzyme activities (48). Guo et al. (13) showed that AI-induced excessive NO compounds might decrease the production of testicular adenosine 3,5-cyclic monophosphate (cAMP), which helps to transport cholesterol to the inner mitochondrial membrane. The result is lower testosterone release. Reduced steroidogenesis results in altered spermatogenesis and spermatic failure (49, 50). All this suggests that the degenerative changes in germ cells found in this study might be due to hormonal deficiency.

Several studies (14, 16, 31, 32) have demonstrated the role of antioxidants in AI-induced reproductive toxicity. Our findings suggest that taurine counters AI-induced testicular lesions. In an earlier study, Das et al. (20) reported that oral administration of taurine was effective in counteracting arsenic-induced oxidative stress and attenuating testicular damage, and apoptosis of testicular tissue in Wistar rats by controlling the reciprocal regulation of Bcl-2/Bad, phospho-ERK1/2, phospho-p38, phospho-Akt, and NF-κB. Furthermore, Manna et al. (22) reported that taurine pre-treatment could prevent cadmium-induced testicular pathophysiology in mice. Taurine has been shown to stimulate testicular steroidogenesis in vivo and in vitro (51) and to promote spermatogonial proliferation and/or meiosis (52). It is considered a capacitating agent (53, 54), and as a sperm motility factor (55, 56).

In summary, our results suggest that Al may act as an endocrine disruptor in male mice and has the potential to induce adverse changes in Leydig cell ultrastructure. These morphological alterations provide a reasonable explanation why Al lowers the fertility index in males. For the first time, our results have also demonstrated that taurine can antagonise the pathological effects of acute Al poisoning in male genital organs, acting as an antioxidant. However, further investigation is required to pin-point the biochemical and molecular mechanisms of these countereffects.

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Sažetak

DJELOVANJE TAURINA PROTIV HISTOMORFOLOŠKIH I ULTRASTRUKTURNIH PROMJENA
U TESTISIMA MIŠEVA IZLOŽENIH ALUMINIJEVU KLORIDU

Cilj je ovog istraživanja bio proučiti zaštitno djelovanje taurina od histomorfoloških i ultrastrukturnih
promjena u testisima švicarskih albino miševa akutno izloženih AlCl₃. Svjetlosnom je mikroskopijom
utvrđena povezanost između jednokratne intraperitonealne (i.p.) doze AlCl₃ (25 mg kg⁻¹ Al³⁺) i odvajanja
nekrotičnoga tkiva, atrofi je tubula, degeneracije zametnih stanica te žarišta hiperplazije Leydigovih stanica.
Usto su se elektronskom mikroskopijom mogli vidjeti razoreni čvrsti spojevi između Sertolijevih stanica,
apoptoza spermatogonija i primarnih spermatocita, različite morfološke abnormalnosti spermatida,
nakupljanje lipidnih kapi, stanjenje glatkog endoplazmatskog retikuluma (sER) te oštećenje mitohondrija
u Leydigovim stanicama. Naknadna primjena taurina u i.p. dozi od 1 g kg⁻¹ ublažila je ove promjene i
značajno smanjila broj zahvaćenih tubula u odnosu na miševa otrovane aluminijem. Ovo je prvo istraživanje
koje potvrđuje zaštitno djelovanje taurina protiv patoloških promjena na tkivu testisa miševa uzrokovanih
aluminijem.

KLJUČNE RIJEČI: histopatologija, muški reproduktivni sustav, spermatogeneza, toksični metali,
ultrastruktura

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