Evaluation of histopathological and ultrastructural changes in the testis of tadalafil treated adult male albino rats

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Abstract

**Background:** Male infertility is a private problem. There was no ideal medication for treatment of the infertile males. PDE5 inhibitors are line of management for male infertility. Tadalafil is efficient and selective PDE5 Inhibitor. It is known to cause less visual manifestations in contrast to the other drugs.

**Aim of the work:** This study was designed to evaluate the effect of tadalafil on the histological structure and ultrastructure of adult male albino rats testis.

**Material and methods:** Twenty-four adult male albino rats were divided into 4 groups; control and 3 treated groups (T1, T2, T3). Tadalafil was given orally to the treated groups in 3 different doses for 30 days. At the end of the experiment, testis was dissected for histological and electron microscopic study.

**Results:** Light microscopic examination of T1 revealed normal structure of tubules with intercellular vacuoles. T2 showed thin irregular thin basement membrane, more vacuoles and congested blood vessels. T3 showed acidophilic exudate in the lumen of the tubules with degenerated germinal cells. The electron microscopic examination of T1 showed normal germinal cells structure with some dilated mitochondria. T2 revealed irregular nuclear membrane of both Sertoli and spermatogonia cells. T3 showed interrupted inter-Sertoli junctional specialization, irregular nuclear membrane in sertoli and Leydig cells. Round spermatid showed loss of acrosomal cap.

**Conclusion:** It can be concluded that, long-term daily use of tadalafil can lead to noticeable pathological effect in the testis which may be implicated in male fertility.

**Keywords:** Tadalafil, testis, sertoli cells, seminefrous tubules, leydig cells

Introduction

Male infertility is a private problem. However the assisted reproductive technology (ART) is a line of management for infertility due to male etiology, significant percent of infertile males still infertile after intrauterine insemination or *in vitro* fertilization (IVF) techniques. There was no ideal medication for treatment of the infertile males. So, it appears that modern medication with a positive effect on male infertility is essential. PDE5 (PhosphoDiestrase Enzyme) inhibitors are line of management for male infertility [1].

It is known that inhibition of the enzyme phosphodiesterase (PDE) type 5 can maintain nitric oxide induced smooth muscle relaxation in the corpus cavernosum of penis and so cause penile erection [2,3]. PDE5 is well expressed in the smooth muscle cells of the penile corpus cavernosum and the smooth muscle cells in blood vessels in the lung, so PDE5 inhibitors can manage erectile dysfunction (ED) [4] and pulmonary hypertension [5]. Although the most of the data about PDE5 inhibitors and ED comes from studies on sildenafil [6], two new PDE5 inhibitors; tadalafil [7] and vardenafil [8] are recently available for the treatment of ED.

The three PDE-5 inhibitors have same mode of action, but their selectivity differs for PDE-5. Sildenafil and vardenafil less selective for PDE5 than tadalafil [9].

Tadalafil is efficient, reversible and selective PDE5 Inhibitor. It is used as an oral therapy in ED caused by psychological, organic or mixed factor [7]. It is a long-acting therapy [10]. Its half-life is about 17.5 hour, high-fat foods do not interfere...
Tadalafil is also known to cause less visual manifestations like abnormal color vision in contrast to the other two drugs [9]. This study was designed to evaluate the effect of tadalafil on the histological structure and ultrastructure of adult male albino rats testis.

Materials and methods

Experimental animals
Twenty-four adult male albino rats (4-6 months old, 200-350gm weight) were used. The use of the animals was prospectively approved by the Committee at Mansoura University, Faculty of Medicine. The rats were housed in the Animal Care Centre of Mansoura faculty of Pharmacy. The rats were provided with fresh food and water daily and inspected for any possible signs of infection (e.g., redness or ulceration of skin).

Experimental design
Animals were divided into four groups (6 rats each). The first group used as a control group. The other three groups were used as treated groups.

Tadalafil administration
Group 1 (Control group): received oral saline solution via orogastric feeding tube for 4 weeks.
Group 2 (treated): 9 mg/kg tadalafil dissolved in distilled water daily for 30 days, through orogastric feeding tube (equivalent to 10 mg adult human dose).
Group 3 (treated): received 1.8mg/kg tadalafil dissolved in distilled water daily for 30 days, through orogastric feeding tube(equivalent to 20 mg adult human dose) [12].
Group 4 (treated): received 2.6mg/kg body weight of tadalafil dissolved in distilled water daily for 30 days, through orogastric feeding tube. The giving doses were similar to the recommended human oral doses 40 mg. Tadalafil was supplied from Pfizer Inc. (Pfizer, Egypt), stored at 2-4°C and protected from sunlight.

Histological analysis
At the end of treatment, rats of each group were anaesthetized with Ketamine (60 mg/kg i.p.) and perfused intracardially with 0.9% NaCl, followed by 4% paraformaldehyde in 0.1 M phosphate buffer (PB) pH 7.2. The testis was carefully dissected and removed. Both testis were taken and weighted. One of them was immersed in 0% paraformaldehyde solution for 24 hours. The specimens were processed for paraffin sections of 5 μm-thickness and stained with H&E for histological examination.

Tissue preparation for ultrastructural study
The other testis was cut into small pieces 1mm³ each. Specimens were immediately fixed in cold 3.5% glutaraldehyde and washed in 0.1 M phosphate buffer (PH 7.2). Then, postfixed with 1% osmium tetroxide (OsO₄), dehydrated and embedded in epoxy resin. Semithin sections (1μm thick) were cut on an LKB ultratome and stained with toluidine blue stain and examined by light microscope. Ultrathin sections (50-70nm thick) were cut, mounted on copper grids, stained with uranyl acetate and lead citrate [13]. Sections were examined by JEOL-100SX transmission electron microscope provided with a digital cameral the Electron Microscopic Unit of Tanta Faculty of Medicine.

Quantitative analysis
The transverse and longitudinal diameters of the seminiferous tubules were measured. The number of cells in tests were measured in a fixed field in serial sections by LeicaQuin 500” image analyzer computer system (England). The cells were counted in five fixed non-overlapping microscopic fields using a 100×objective lens.

Statistical analysis
Statistical analysis was done using computer software SPSS program (statistical package for social science) version 10. All data were expressed as the mean±SD. The significance level considered was P≤0.05.

Results

Light microscopic examination

Group 1 (control group)
Histological examination of H&E and toluidine blue stained sections in the testis of the control group showed multiple rounded seminiferous tubules with regular outlines. They were lined by 4-6 layers of germinal epithelium at different stages of spermatogenesis. The flagella of mature sperms were seen in the lumen of the tubules. The lining epithelium consisted of Sertoli cells and other germinal cells. Sertoli cells appeared pyramidal in shape and resting on the basement membrane. The germinal epithelium consisted of spermatogonia, 1ry spermatocytes, rounded spermatids and elongated spermatids. The interstitial spaces in-between the tubules contained Leydig cells and some blood vessels (Figure 1).

Group 2 (treated-T1)
L/M examination of testicular sections of this group showed seminiferous tubules with germinal epithelium. The flagella of sperms were obvious in the lumen. Sertoli cells were seen resting on the basement membrane. The spermatogenic cells were noticed; 1ry spermatocyte, round spermatid and elongated sperms. Areas of intercellular vacuoles were detected. The interstitial space showed some Leydig cells separated by congested blood vessels (Figure 2).

Group 3 (treated-T2)
L/M examination of testicular sections of this group showed some seminiferous tubules with irregular outline. The flagella of sperms were obvious in the lumen. Sertoli cells were seen resting on the thin, irregular basement membrane. The spermatogenic cells were noticed; 1ry spermatocyte, round...
vacuoles were detected. The interstitial space showed some Leydig cells separated by congested blood vessels (Figure 3).

**Group 4 (treated-T3)**

L/M examination of testicular sections of this group showed seminiferous tubules with irregular outline with acidophilic exudate in their lumen. Some spermatogenic cells showed signs of degeneration, vacuolated cytoplasm and pyknotic nuclei. Increased areas of cellular loss and vacuolation. Congested blood vessels between tubules (Figure 4).

**Transmission electron microscopic results (TEM)**

**Group 1 (control group)**

The ultrastructure of the control testis showed Sertoli cells were resting on the basement membrane with adjacent myoid cell. The nucleus appeared triangular and euchromatic with well-developed nucleolus. The cytoplasm showed some organelles like mitochondria, and ribosomes. The inter-Sertoli junctional specialization with adjacent cells was apparent (Figure 5A). The germinal epithelium showed different stages of maturation. The primary spermatocytes appeared as large rounded cells,
The spermatogonia appeared large rounded cells, with irregular nuclear membrane, and acidophilic exudate in their lumen and interstitial space (*). Interstitial cells of Leydig (Ly) and congested blood vessel (bv) are noticed (H&E x100). B. Some spermatogenic cells showed signs of degeneration, vacuolated cytoplasm and pyknotic nuclei (arrows). Increased areas of cellular loss and vacuolation. And acidophilic exudate in the lumen (*) (H&E x400). C. Seminiferous tubules with intercellular vacuolation (*) (Toulidine blue x100). D. Higher magnification of C, showing Sertoli cells (S), rounded spermatids (Sd), elongated spermatid (arrows) Areas of intercellular vacuoles are detected (*). Degenerated cells with pyknotic nuclei (arrow heads) (Toulidine blue x400).

with rounded nucleus and dispersed granular chromatin. The cytoplasm showed mitochondria, ribosomes, and lysosomes (Figure 5B). The rounded spermatids showed rounded nucleus with fine granular chromatin. Acrosomal vesicle was apparent over the anterior part of the nucleus to form acrosomal cap. The cytoplasm contained mitochondria and ribosomes (Figure 5C). The fusiform spermatids showed elongated pyriform nuclei. Acrosomal cap could be noticed on the anterior part of the nucleus (Figures 5D and 5E). The interstitial cells of Leydig showed nuclei with thin peripheral rim of chromatin. The cytoplasm showed organelles like mitochondria, lysosomes and smooth endoplasmic reticulum (Figure 5F).

Group 2 (treated-T1) 

The ultrastructure of the treated testis revealed normal Sertoli cells, which showed triangular nucleus with prominent nucleolus and fine granular chromatin. The cytoplasm contained some normal and some dilated mitochondria (Figure 6A). The spermatogonia appeared large rounded cell, with irregular nuclear membrane, the cytoplasm contained ballooned mitochondria, lysosomes, vacuoles, ribosomes and smooth endoplasmic reticulum. Try spermatocyte appear with large rounded nucleus (Figures 6B and 6C). Rounded spermatids appeared with spherical nuclei and incomplete acrosomal caps. Cross sections of the middle pieces of numerous spermatozoa could be also seen (Figure 6D). The elongated spermatids showed pyriform shaped nuclei with acrosomal cap (Figure 6E). The interstitial cells of Leydig appeared with slight indentation in the nuclear membrane. The cytoplasm contained mitochondria, lysosomes and vacuoles (Figure 6F).

Group 3 (treated-T2) 

The examination of the treated testis in this group showed abnormal structure of Sertoli cells, its nucleus appeared with irregular nuclear membrane, the cytoplasm contained ballooned mitochondria. The inter-Sertoli junctional specialization with adjacent cells was interrupted (Figure 7A). The spermatogonia showed nucleus with irregular nuclear mitochondria, lysosomes, vacuoles, ribosomes and smooth endoplasmic reticulum.
membrane, the cytoplasm contained ballooned mitochondria, ribosomes, dilated smooth cytoplasmic reticulum and some vaculated areas (Figure 7B). Round spermatid showed rounded nucleus with incomplete acrosomal caps. The cytoplasm contained rounded mitochondria and ribosomes (Figure 7C). The elongated spermatids showed pyriform shaped nucleus with acrosomal cap (AC). The interstitial cells of Leydig appeared with slight indentation in the nuclear membrane (arrows). The cytoplasm contained mitochondria (M), lysosomes (L) and vacuoles (V).

**Group 4 (treated-T3)**

The examination of the treated testis in this group showed abnormal structure of Sertoli cells. It showed triangular nucleus with irregular membrane, prominent nucleolus, the cytoplasm contained large cytoplasmic clear vacuole, dilated smooth endoplasmic reticulum and lysosome. The inter-Sertoli junctional specialization with adjacent cells was disturbed (Figure 8A). The adjoining primary spermatocytes cells exhibited large nucleus with irregular membrane and clumped chromatin (Figure 8B). Round spermatid showed indented nucleus with dense clumped chromatin. There was complete loss of acrosomal caps. The cytoplasm showed vacuoles and ballooned mitochondria. Some abnormal forms of elongated spermatids were also noticed (Figure 8C). The interstitial cells of Leydig appeared with indented nuclei and

Figure 6. Photomicrograph of ultrastructure of the treated testis (T1) A. Sertoli cells, with triangular nucleus with prominent nucleolus (nu) and fine granular chromatin (N). The cytoplasm contains some normal and some dilated mitochondria (M). B&C. The spermatogonia appear large rounded cell, with irregular nuclear membrane (arrows). The cytoplasm with ballooned mitochondria (M), lysosomes (L), vacuoles (V), ribosomes (R), smooth endoplasmic reticulum (sER). D. The elongated spermatids with spherical nuclei and incomplete acrosomal caps (arrow). Cross sections of the middle pieces of spermatooza could be also seen (arrow head). E. The elongated spermatids show pyriform shaped nucleus (N) with acrosomal cap (AC). F. The interstitial cells of Leydig appeared with slight indentation in the nuclear membrane (arrows). The cytoplasm contains mitochondria (M), lysosomes (L) and vacuoles (V).

Figure 7. Photomicrograph of ultrastructure of the treated testis (T2) A. Sertoli cell nucleus with irregular nuclear membrane (arrow heads) and fine granular chromatin. The cytoplasm contains ballooned mitochondria (M). The inter-Sertoli junctional specialization with adjacent cells is interrupted (arrow). B. The spermatogonia showed nucleus with irregular nuclear membrane (arrow heads). The cytoplasm shows ballooned mitochondria (M), ribosomes (R), dilated smooth cytoplasmic reticulum (sER) and some vaculated areas (*). C. Round spermatid with rounded nucleus and incomplete acrosomal caps (arrow). The cytoplasm contains rounded mitochondria (M) and ribosomes(R). D. The elongated spermatids with pyriform shaped nuclei (N). Ectoplasmic specialization can be observed between the Sertoli cell and the spermatids. It is made of three layers; Sertoli cell plasma membrane (arrow head), surface cistern of smooth endoplasmic reticulum that lies parallel to the plasma membrane (*), and layer of actin filaments located between the plasma membrane and surface cistern (arrow). E. Some abnormal forms of sperms can be seen (arrow). F. The interstitial cells of Leydig appear with indented nuclei (arrow). The cytoplasm contains lysosomes (L), rounded mitochondria (M), and vacuoles (V).
clumped chromatin. The cytoplasm contained lysosomes, mitochondria, lipid droplets and vacuoles (Figure 8D).

Morphometric results and statistical analysis

Weight of testis
There was no significant change in testicular weight between control and treated groups (Graph 1).

The mean diameter of seminiferous tubules
In comparison with the control group, the mean diameter of seminiferous tubules showed a significant increase in the treated groups (Graph 2).

The number of germ cells
In comparison with the control group, the mean number of germ cells showed a significant decrease in the treated groups (Graph 3).

Discussion
The PDE-5 inhibitors; sildnafile, tadalafil and vardenafil helped to treat the erectile dysfunction. All three drugs are safe and tolerable. Tadalafil improved erectile dysfunction even in males with type 1 or type 2 diabetes [14].

In this study, the control rat testis showed multiple rounded seminiferous tubules with regular outlines. The lining
epithelium consisted of Sertoli cells and other germinai cells. Sertoli cells appeared pyramidal in shape and resting on the basement membrane. The germinal epithelium consisted of spermatogonia, primary spermatocytes, rounded spermatids and elongated spermatids. The interstitial spaces in-between the tubules contained Leydig cells and some blood vessels. The ultrastructure of the Sertoli cell showed that the nucleus appeared triangular and euchromatic with well developed nucleolus. The inter-Sertoli junctional specialization with adjacent cells was apparent. The primary spermatocytes appeared as large rounded cells, with rounded nucleus and dispersed granular chromatin. The rounded spermatids showed rounded nucleus with fine granular chromatin. Acrosomal vesicle was apparent over the anterior part of the nucleus. The fusiform spermatids showed elongatedpyriform nuclei. Acrosomal cap could be noticed on the anterior part of the nucleus. The interstitial cells of Leydig showed nuclei with thin peripheral rim of chromatin. This comes in agreement with the histological and ultrastructure of the control adult rat testis [15].

The treated group testicular sections which received the least dose of tadalafil, 0.9 mg/kg orally, showed seminiferous tubules with germinal epithelium. The flagella of sperms were obvious in the lumen. Areas of intercellular vacuoles were detected. The interstitial space showed some Leydig cells separated by congested blood vessels. The ultrastructure of the treated testis revealed normal Sertoli cells, which showed triangular nucleus with prominent nucleolus and fine granular chromatin. The cytoplasm contained some normal and some dilated mitochondria. The spermatogonia appeared as rounded cell, with irregular nuclear membrane. Rounded spermatids appeared with spherical nuclei and incomplete acrosomal caps. Cross sections of the middle pieces of numerous spermatozoa could be also seen. The elongated spermatids showed pyriform shaped nuclei with acrosomal cap. The interstitial cells of Leydig appeared with slight indentation in the nuclear membrane. The cytoplasm contained mitochondria, lysosomes and vacuoles. The diameter of seminiferous tubule increased in this group when compared with control group. The number of germ cells showed significant decrease in this group when compared with the control group.

In contrast to this study, it was reported that administration of single dose of sildenafil (50 mg) in young infertile males resulted in significant increase in the number of spermatozoa [18]. In agreement with this result, it was concluded that tadalafil administration (10 mg) showed a significantly increase in sperm count and motility with increased incidence of abnormal forms. Histological examination showed loosely packed stroma around seminiferous tubules with reduction in number of spermatogenic cells [12]. Also, it was documented that administration of 50 mg sildenafil caused degeneration of somniferous tubules and the spermatogenic cells. The interstitial tissue showed congested blood vessels, degenerated leydig cells. The basement membrane was thin [17]. It was concluded that of sildenafil citrate (50 mg) administration caused significant reduction in the number of sperms [18].

In contrast, it was reported that administration of 10-mg and 20-mg tadalafil in human every day for six months, showed insignificant change in the sperm morphology [19]. Also, it was found that administration of single dose of sildenafil (50 mg) in young infertile males resulted in significant increase in sperm motility, while, tadalafil single dose (20 mg) caused reduction in the motility of the sperms. It was found that impact of sildenafil on the motility of sperms might be due to changes in mitochondria and calcium channels [20].

The Sertoli cell tight junctions is a vital component of the blood-testis barrier which is essential for normal spermatogenesis. The interrupted inter-Sertoli junctional specialization in case of tadalafil administration may be explained by the fact defective testicular blood barrier could cause opening of the testicular blood barrier gate. This disturb the protective and nutritional function of the barrier and impair spermatogenesis [21]. L/M examination of testicular sections of the group received 2.6 mg/kg showed seminiferous tubules with irregular outline with acidophilic exudate in their lumen and congested blood vessels in between. Some spermatogenic cells showed signs of degeneration; vacuolated cytoplasm and pyknotic nuclei. Increased areas of cellular loss and vaculation in the tubules. The ultrastructure of treated testis in this group showed abnormal structure of Sertoli cells. It showed triangular nuclei with irregular membrane, the cytoplasm contained large
cytoplasmic clear vacuoles, dilated smooth endoplasmic reticulum and lysosome. The inter-Sertoli functional specialization with adjacent cells was disturbed. Round spermatid showed indented nucleus with dense clumped chromatin. There was complete loss of acrosomal cap. The cytoplasm showed vacuoles and ballooned mitochondria. Some abnormal forms of elongated spermatids were also noticed with defective ectoplasmic specialization. The interstitial cells of Leydig appeared with indented nuclei, the cytoplasm contained lysosomes, mitochondria, lipid droplets and vacuoles. The diameter of seminiferous tubule increased in this group when compared with control group. The number of germ cells showed significant decrease in this group when compared with the control group.

In agreement with this result, it was documented that administration of 100 mg sildenafil caused degeneration of seminiferous tubules with thin basement membrane and the spermatogenic cells, the interstitial tissue showed congested blood vessels, degenerated Leydig cells [17]. However, it was noticed that 100-mg sildenafil citrate in diabetic patients increased sperm motility and semen volume [22].

In the same time, it was documented that administration of 100-mg singledose of sildenafil had no significant effect on sperm motility, sperm count or morphology in human [23]. Also, single dose of vardenafil (20 mg) had no significant effects on sperm motility, sperm viability, and sperm morphology in males [24].

In this study, round spermatid in treated rats showed incomplete or absent acrosomal cap. Some elongated spermatids showed abnormal nuclei, shape and defective ectoplasmic specialization. According to some authors, the ectoplasmic specialization, which is unique to the testis, have an essential role in spermatid nuclear shaping. F-actin filaments from normal Sertoli cells enhance spermatid head elongation [25]. So, abnormal Sertoli cell may affect spermatid nuclear shaping.

On the other hand, it was reported that PDE5 inhibitors may enhance the capacitation of spermatids [18, 26, 27]. cGMP can open cyclic nucleotide-gated channels for calcium entry inside the spermatooza to initiate the acrosomal reaction. However, PDE5 cause hydrolysis of cGMP, so, inhibition of PDE5 by sildenafil citrate can trigger the effect of cGMP on sperm acrosomal reaction [28].

In this investigation, the diameter of seminiferous tubules increased in treated rats when compared with control rats. Similar result was reported in case of sildenafil and explained by the vasoactive of the drug, which may occur in other PDE5 inhibitors [29].

Conclusion
It can be concluded that, long-term daily use of tadalafil can lead to noticeable pathological effect in the testis which may be implicated in male fertility and this effect is dose dependent, so the effect of tadalafil necessities further investigations.

Competing interests
The author declares that he has no competing interests.

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