

# Assessment of testis histological changes and sperm parameters in experimentally-induced unilateral and bilateral cryptorchid mouse model

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## Abstract

**Background:** In most mammals, the testis is always maintained at a lower temperature than that in the abdomen, and exposure of the testis to body temperature causes degeneration of germ cells.

**Objective:** In this research, the long effect of heat exposure on sperm parameters and microstructure of mouse testis were investigated. Cryptorchid mouse were induced by exposure to abdominal heat.

**Materials and Methods:** Immature mice were anesthetized and a small incision was made in the abdominal skin, then fat pad at the upper end of testis was sutured to peritoneum. Weight of testis, spermatogenic cell numbers, tubular ectasis (rate of tubular lumen comparing to the thickness of germinal epithelium) as well as epididymal sperm parameters were measured.

**Results:** The results showed that spermatogenesis was arrested and testicular weights, seminiferous tubular diameters and epididymal sperm parameters were significantly reduced in the bilateral undescended testis compared with unilateral undescended testis and the control mice. However, complete depletion of seminiferous tubules and absence of germ cells was not found in the animals.

**Conclusion:** In general, high temperature caused a decreased in all analyzed parameters except spermatogonial cell number probably due to the apoptosis and these changes significantly increase in bilateral groups compared with unilateral groups. We believe that the present model is a suitable tool for enrichment of spermatogonial stem cells, also it is useful for treatment of cryptorchidism and further biological research on spermatogenesis.

**Key words:** Cryptorchidism, Testis, Spermatogonial stem cells.

## Introduction

It was shown that mammalian testis must descend from the abdominal cavity for normal development to occur, and that the elevated testicular temperature disrupts spermatogenesis and causes infertility (1).

Testicular maldescent is a common urological disorder. In full-term infants, the incidence is 3.4%

to 5.8% and at the time of puberty, 0.8%- 1.8% of boys remain cryptorchid (2).

The abdominal positioning of cryptorchid testis is associated with disruption of germinal epithelium (3). In addition to degeneration of the seminiferous epithelium, accumulation of lipid in sertoli cells, local dilations of the intercellular spaces between sertoli cells junctions and modifications in the interstitial tissue have been reported (4).

Damber *et al* showed that blood flow to the cryptorchid testis is reduced despite the fact that the cryptorchid testis contains a greater percentage and number of blood vessels than the normal testis (5).

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Leydig cell hypertrophy and hyperplasia are accompanied by an increase in the cytoplasmic organelles involved in steroidogenesis, such as smooth endoplasmic reticulum and mitochondria (6). By contrast, Hall *et al* reported that steroidogenic functions of the somatic cells (Leydig and Sertoli cells) appear to be normal when exposed to the core body temperature (7). The elevated testicular temperature affects the normal functioning of all major cell types (Germ, Leydig and Sertoli cells) to some extent (8). Also heat-induced germ cell degeneration is usually accompanied by alteration in Sertoli cell morphology and function, a decrease in testes fluid, and increase in serum FSH with no changes in serum Testosterone and LH levels (9). More recently, a number of studies have documented that high temperatures induced an increased synthesis of several proteins and a decreased synthesis of many others.

The effect of temperature on testicular function, especially sperm motility and sperm viability are unclear however, several enzymatic activities are affected and the lipid composition of plasma membranes is modified (10). Besides, testicular stem cell enrichment has been widely used to investigate the biology of these cells in addition to the production of transgenic animals, germ cell transplantation and restoration of fertility in various species. The potential model for enrichment is cryptorchidism (11). In this study, we are reporting the effects of prolonged cryptorchidism on sperm parameters and testis microstructure of unilateral and bilateral experimental cryptorchidism. We believe that the present model is a suitable tool for enrichment of spermatogonial stem cells also it is useful for the treatment of cryptorchidism and further biological research in to spermatogenesis.

## Materials and methods

### Animals

Immature NMRI mice at under 2 months of age (Weigh 7-12 gr) were purchased from Razi Vaccine & Serum Research Institute (Karaj, Iran). Mice were anesthetized with an injection of 1.6ml/kg of a mixture of Ketamine and xylezine. A cut was made along the skin in the upper abdominal region and the adipose tissue of the caput epididymis was sutured to the inner peritoneal wall, pushing the testes in to the abdomen.

The first experimental group mice were unilaterally cryptorchidized and in the second

experimental group, mice were bilaterally cryptorchidized. Experimental and control animals were sacrificed by cervical dislocation, 2, 4, 6 and 8 weeks after surgery. Following cervical dislocation left testis of unilateral groups and both testes of bilateral groups and epididymis were removed, the testes were weighed and fixed in Bouin's fixative, dehydrated and embedded in paraffin.

Then, 5- $\mu$ m serial microscopic sections were prepared and at least 5 slides from each testis stained with hematoxylin and eosin for histological assessment.

In each experiment, at least 5 animals were prepared and analyzed (13). All animal experimentation protocols were approved by the Institutional Animal Care and Use Committee of Tarbiat Modarres University (Tehran, Iran).

### Sperm parameters assessment

The epididymis was placed in 1 ml PBS (Ph 7.4) and minced into small pieces before being incubated at 37°C for 30min. Sperm parameters were monitored by light microscopy. Sperm viability was assessed by determining the percentage of sperms excluding vital dye (25% eosin solution). Briefly, 7  $\mu$ l of eosin solution was added to 20  $\mu$ l of cell suspension after incubation and mixed thoroughly. Motility of sperm was also assessed by determining the percentage of motile sperm (14). Finally sperm numbers was also calculated with a hemocytometer count and compared in unilateral, bilateral, sham and control groups.

### Morphometrical analysis of testis

For each testis, in 100 randomly selected tubular profiles that were round or nearly round, the diameters of tubules and epithelium thickness were measured under light microscopy. Volume density of spermatogonial cells, spermatocytes and spermatids in seminiferous tubules were determined.

The location and morphology of the cells within the seminiferous tubules were used to identify them. An estimate of each parameter was performed by examining 20 fields in 5 histological sections from each testis (13).

### Statistical analysis

The results were analyzed by performing ANOVA and Tukey's tests, with  $p < 0.05$  considered as statistically significant. The mean and standard deviation (SD) was also calculated for each value.

## Results

### Analyses of sperm parameters in epididymis following creating unilateral and bilateral cryptorchidism mouse model

There was no significant difference between the result of sham and control groups. As the results show, there was a significant decline in sperm viability and motility rates (Table I and II) as well as a decline in sperms counts (Table III) following surgery. The viability of sperms decreased in all weeks after surgery, however, only in 2<sup>nd</sup> and 4<sup>th</sup> week after surgery there was significant difference between unilateral and bilateral groups. Sperm viability was not observed after 4<sup>th</sup> week in bilateral group but this situation occurs in unilateral group after 8<sup>th</sup> week (Table I). Similar observation was seen on sperm motility after surgery. However, the effect of heat on sperm motility was more severe. The significant difference between unilateral and bilateral groups only observed 2 weeks after surgery (Table II). The number of sperms per ml aspirated from one epididym was determined by hemocytometer counts. As presented in Table III, in all times after surgery the number of sperms decreased significantly in epididymis of experimental groups compared with control group but there was no significant difference between unilateral and bilateral groups (Table III). In addition, we showed decline in all sperm parameters after heat exposure especially in bilateral group.

### Analyses of morphological changes of mouse testis following creating unilateral and bilateral cryptorchid mouse model

As the data show, the values of the testicular weight in unilateral and bilateral groups were significantly less than those in the control group. These decrease observed from 2 week in both experimental groups but significant difference between these groups only observed in 4<sup>th</sup> week after surgery (figure 1). Histological examination of the testis in control and experimentally groups revealed that tubular ectasis (rate of seminiferous tubules diameter comparing to the seminiferous epithelium thickness) was increased in experimental groups in time dependent manner. This difference started from 2 week after surgery between control and experimental groups (figure 2). The testes were histologically examined by H&E staining (figure 3). Compared to the testes in the control group, in cryptorchid model, testicular tissues were degenerated. Spermatogonia, spermatocyte and round spermatids were observed,

**Table I.** Viability rates (%) of sperms aspirated from epididymis after creating unilateral and bilateral mouse model.

Sperm parameters Time of assessment	Viability rate (mean ± SD)			
	2 <sup>nd</sup> w	4 <sup>th</sup> w	6 <sup>th</sup> w	8 <sup>th</sup> w
Control	75.3 ± 2.6	78 ± 1.7	77.3 ± 1.4	78.6 ± 1.4
Unilateral	28.3 ± 3.1 <sup>a</sup>	12.33 ± 1.4 <sup>a</sup>	3.66 ± 1.2 <sup>a</sup>	0 <sup>a</sup>
Bilateral	10 ± 2.8 <sup>a,b</sup>	0 <sup>a,b</sup>	0 <sup>a</sup>	0 <sup>a</sup>

The replication of experiences was at least 3 times for all groups.

a: Significant difference versus control group in the same column (p<0.05)

b: Significant difference versus unilateral group in the same column (p<0.05)

Abbreviation: SD: Standard deviation

**Table II.** Mortality rates (%) of sperms aspirated from epididymis after creating unilateral and bilateral mouse model.

Sperm parameters Time of assessment	Motility rate (mean ± SD)			
	2 <sup>nd</sup> w	4 <sup>th</sup> w	6 <sup>th</sup> w	8 <sup>th</sup> w
Control	75 ± 2.5	72.66 ± 3.2	75.33 ± 3.4	75.66 ± 3.4
Unilateral	12.33 ± 1.4 <sup>a</sup>	5.33 ± 0.8 <sup>a</sup>	1.6 ± 0.8 <sup>a</sup>	0 <sup>a</sup>
Bilateral	3.33 ± 0.6 <sup>a,b</sup>	0 <sup>a,b</sup>	0 <sup>a,b</sup>	0 <sup>a,b</sup>

The replication of experiences was at least 3 times for all groups.

a: Significant difference versus control group in the same column (p<0.05)

b: Significant difference versus unilateral group in the same column (p<0.05)

Abbreviation: SD: Standard deviation

**Table III.** Sperm count per ml in epididymis after creating unilateral and bilateral mouse model.

Sperm parameters Time of assessment	Viability rate (mean ± SD)			
	2 <sup>nd</sup> w	4 <sup>th</sup> w	6 <sup>th</sup> w	8 <sup>th</sup> w
Control	4.4 ± 0.1	4.3 ± 0.2	4.1 ± 0.2	4.4 ± 0.1
Unilateral	0.1 ± 0.01 <sup>a</sup>	0.1 ± 0.01 <sup>a</sup>	0.7 ± 0.02 <sup>a</sup>	0.3 ± 0.8 <sup>a</sup>
Bilateral	0.01 ± 0.001 <sup>a</sup>	0.06 ± 0.008 <sup>a</sup>	0.03 ± 0.01 <sup>a</sup>	0 <sup>a,b</sup>

The replication of experiences was at least 3 times for all groups.

a: Significant difference versus control group in the same column (p<0.05)

b: Significant difference versus unilateral group in the same column (p<0.05)

Abbreviation: SD: Standard deviation

but the differentiated cells after them, elongated spermatids or spermatozoa were not detected in the experimental groups. Numbers of spermatogonia increased after 8<sup>th</sup> week post surgery in bilateral group, while absence or sloughing of germ cells were not found in all experimental animals (Table IV). Based on these results the bilateral cryptorchid model is a useful tool for enrichment of spermatogonial stem cells.

**Table IV** .Cell count per (mm)<sup>3</sup> in seminiferous tubules (mean  $\pm$ SD) $\times 10^4$ .

		Control	Unilateral	Bilateral
Time of extract	2 <sup>nd</sup> week	Spg 4.4 $\pm$ 0.7	7.5 $\pm$ 1.4 <sup>a</sup>	11.7 $\pm$ 2.4 <sup>a,b</sup>
		Spc 14.9 $\pm$ 2.1	19.5 $\pm$ 1.9 <sup>b</sup>	8 $\pm$ 1.1 <sup>a</sup>
		Spt 16.4 $\pm$ 1.2	2.2 $\pm$ 1.2 <sup>a</sup>	0.9 $\pm$ 0.2 <sup>a,b</sup>
	4 <sup>th</sup> week	Spg 3.8 $\pm$ 0.6	11.7 $\pm$ 2.4 <sup>a</sup>	11 $\pm$ 2.7 <sup>a</sup>
		Spc 15.5 $\pm$ 2.3	20 $\pm$ 2.6 <sup>a</sup>	5 $\pm$ 0.3 <sup>a,b</sup>
		Spt 21.4 $\pm$ 2.5	4.3 $\pm$ 0.4 <sup>a</sup>	0.1 <sup>a,b</sup>
	6 <sup>th</sup> week	Spg 3.7 $\pm$ 0.9	12.2 $\pm$ 1.3 <sup>a</sup>	11.7 $\pm$ 1.2 <sup>a</sup>
		Spc 13.7 $\pm$ 1.3	9.1 $\pm$ 1.0 <sup>a</sup>	4.6 $\pm$ 0.1 <sup>a,b</sup>
		Spt 26.1 $\pm$ 2	2.5 $\pm$ 0.2 <sup>a</sup>	0.1 <sup>a</sup>
	8 <sup>th</sup> week	Spg 2.5 $\pm$ 0.6	10.8 $\pm$ 2 <sup>a</sup>	12.2 $\pm$ 2.1 <sup>a</sup>
		Spc 12.2 $\pm$ 1.6	5.1 $\pm$ 0.4 <sup>a</sup>	1.7 $\pm$ 0.1 <sup>a,b</sup>
		Spt 23.1 $\pm$ 2.1	0 <sup>a</sup>	0 <sup>a</sup>

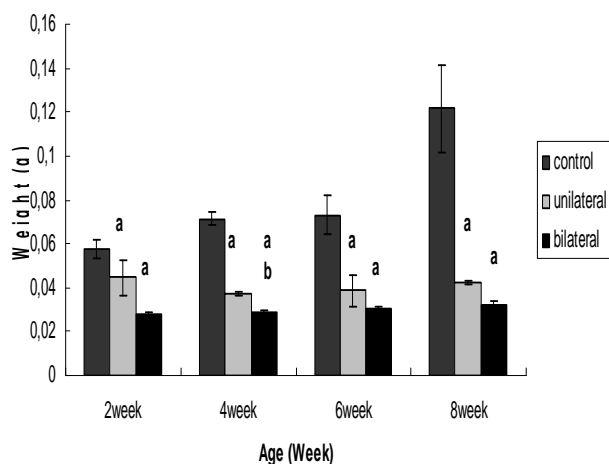
Spg: Spermatogonia, Spc: Spermatocyte, Spt: Spermatid.

The replication of experiences was at least 3 times for all groups.

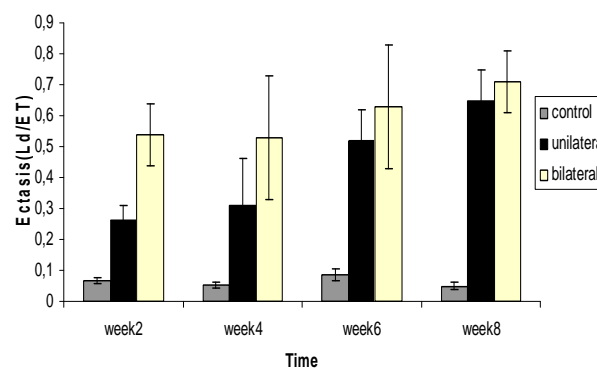
a: Significant difference versus control group in the same column ( $p < 0.05$ ).

b: Significant difference versus unilateral group in the same column ( $p < 0.05$ ).

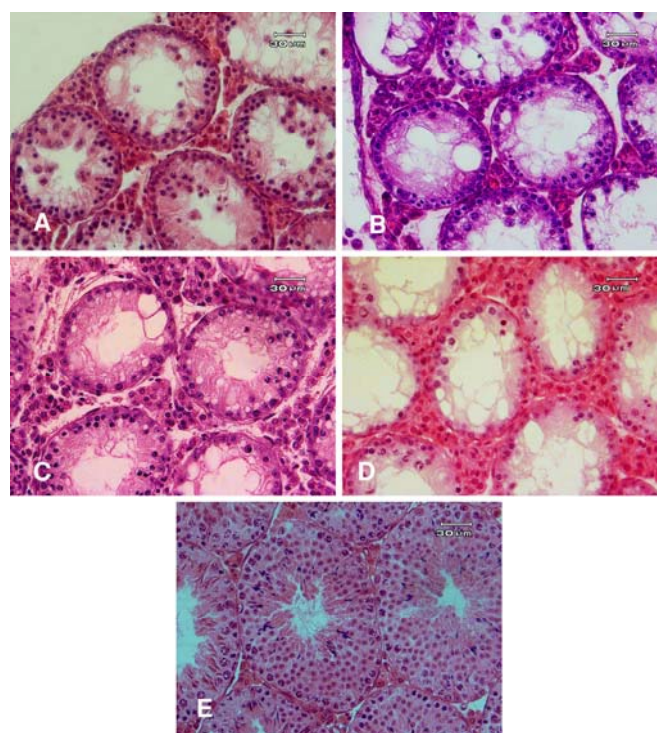
SD: Standard deviation



**Figure 1.** Testicular weights (gr) in control and experimental groups. a: Significant difference versus control group ( $p < 0.05$ ) b: Significant difference with unilateral group ( $p < 0.05$ ).



**Figure 2.** Seminiferous tubular ectasis ( $\mu$ m) of the mouse testis in the control and experimental groups. a: Significant difference versus control group ( $p < 0.05$ ) b: Significant difference with unilateral group ( $p < 0.05$ ).



**Figure 3.** Cross sections of mouse testes staining with H&E A: Four weeks after surgery (unilateral group) B: Four weeks after surgery (bilateral group) C: Eight weeks after surgery (unilateral group) D: Eight weeks after surgery (Bilateral group) E: Cross section of normal testis.

## Discussion

Most mammals have a scrotum, and it is well established that the mild hyperthermia leads to infertility (15,16). As a sequence, the effects of local heating on the animal testis have been widely studied (17). In the current study we are reporting the effects of prolong heating caused by surgery on sperm parameters and testis microstructure of mouse testis also we compared this effects in

unilateral and bilateral experimental cryptorchidism mouse model.

One of the findings in this study was the decrease in testis weight and in the number of spermatozoa in testis and epididymis in experimental mouse toward the end of experiment, associated with a significant deterioration in the histology. This finding indicates that degeneration of spermatocytes and spermatids occurs after cryptorchidism is in agreement with Chowdhury and Steinberger (18,19) also by Collis and Lacy (20). By 2 weeks post operation, the mean weight of the testes in the bilateral cryptorchid mouse had diminished about 50% of the weight in the control group. Many studies suggested that germ cell loss after heat exposure caused this decrease in testicular weight (21). Longquan (2006) demonstrated that experimentally induced cryptorchidism in adult male rats caused a decrease in the percentage of motile spermatozoa within 3 days post operation (22). The present study demonstrated that experimentally induced cryptorchidism in immature mouse caused the epididymal percentage of sperm number, motility and viability rates decreased significantly from 2 week postoperation in both experimental groups. Motility and viability rates decline in bilateral experimental group compared with unilateral group.

The seminiferous tubules in the cryptorchid testes were similar in shape in cross section to those of the control testes, but their diameter showed progressive reduction with the increasing duration of cryptorchidism. The volume density measurements showed that the seminiferous tubule lumen is increased after cryptorchidism. Degeneration of germ cells resulted in the increase in the tubule lumen (4,6). We showed tubular ectasis increased after surgery and this subject was more prominent in bilateral groups compared with unilateral groups probably due to rapid degeneration of differentiated germ cells in bilateral groups. Many investigators showed significant decreased diameters of seminiferous tubules and thickness of germinal epithelium in cryptorchid testis (23).

During spermatogenesis, the spermatogonial stem cells (SSC) which undergo self-renewal throughout adult life of the animal and produces progeny cells, undergo the complicated differentiation steps. As in other rare stem cell populations, enrichment strategies are required to study SSC biology. It has been demonstrated that during testis development, the total number of SSC per testis increases 39-fold from newborn to adult

(24). The number of SSC also appears to increase during the regeneration of spermatogenesis following cytotoxic damage (25, 26). Either testis hyperthermia or cryptorchidism may be used to increase the percentage of SSC *in vivo*. Although these experimental conditions cause depletion of more differentiated germ cells, they do not appear to affect SSC survival and biological activity, thus resulting in an increase in the ratio of SSC vs. the total number of germ cells (27). An effect of heat on spermatogonia in mice had already been reported by Reid, who performed studies on the repopulation of seminiferous tubules after irradiation (28). Differentiation of spermatogonia has also been reported to be arrested in cryptorchid mice (29) and heating the testes of sheep caused a reduction in the production of A<sub>1</sub> spermatogonia 20 days later, with no effect on the numbers of A<sub>0</sub> spermatogonia (30). Miralga and Hayashi (31) reported a decrease in the percentage of tubules with spermatogonia from 100 to 89% at day 15 after heating the testes of rats to 43°C for 30 min. We showed spermatocytes and spermatids are major cells which were affected also spermatogonial number increased after 2 week postoperation and this increase is noticeable in bilateral group compared with unilateral group. In addition, experimentally induced bilateral cryptorchidism provides a useful model for enrichment of spermatogonial stem cells.

According to previous study, experimentally induced unilateral cryptorchidism provides a useful model for study of the role of local factors regulating testis cell apoptosis this suggested that apoptotic cell death induced by bilateral cryptorchidism might be affected by changes in systemic factors. The increase of apoptosis in male germ cell after unilateral cryptorchidism is presumably regulated by local testicular factors (32). Also in this research, the differences indicated between unilateral and bilateral groups may arise from this subject.

We believe that the present model is a suitable tool for enrichment of spermatogonial stem cells; also it is useful for treatment of cryptorchidism and further biological research in spermatogenesis.

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