

# Acute high dose reprotoxic effects of bisphenol-A on the testicles of adult rodents

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## Author's Contribution

<sup>1</sup>Substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work.

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## ABSTRACT

**Objectives:** To evaluate the acute effects of high dose of bisphenol-A (BPA) induced in the testicles of adult rats.

**Methodology:** An Experimental study was conducted at Anatomy Department, University of Health Sciences, Lahore, from January 2015 to December 2015.

Twelve adult Wistar albino rats were weighed and divided into three groups of four rats each. Group A (control) was given corn oil (2ml/kg/day). Group B (BPA-100) was given BPA 100mg/kg/day dissolved in corn oil (2ml/kg/day). Group C (BPA-200) was given BPA 200mg/kg/day dissolved in corn oil (2ml/kg/day). The animals were sacrificed on the 8th day after weighing. Testis were weighed and fixed in Buoin's solution. Johnson scoring was done in H&E stained slides. Basement membrane integrity was observed in PAS stained slides.

**Results:** Group A rats showed normal parameters. Group B (BPA-100) showed a significant decrease in Johnson score and an increase in basement membrane disruption while the changes in body weight of animals and paired weight of testis was statistically insignificant. Group C (BPA 200) rats died within 3-4 days and the experiment was limited to two groups.

**Conclusion:** The results of the present study indicate that a high dosage of Bisphenol A in rats causes significant disruption of the testicular histology and proves to be a significant reprotoxin even with a short duration of exposure.

**Key Words:** Bisphenol A, Testicles, Johnson score, Basement membrane.

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## Introduction

Endocrine disrupting chemicals (EDCs) are becoming an uprising environmental factor leading to an increase in deteriorative reproductive health effects and infertility.<sup>1</sup> The xeno-estrogenic effects of EDCs are mainly associated with reproductive health problems and involve the estrogen signaling pathways.<sup>2</sup> Interference with androgen signaling pathways is another mechanism associated with male reproductive developmental and functional health problems<sup>3</sup>. Testicular dysgenesis syndrome includes a range of male reproductive problems such as cryptorchidism, poor semen quality and

testicular carcinoma that are increasing partially due to the increased exposure of environmental pollutants.<sup>4</sup>

Bisphenol A (BPA) is a plasticizer and one of the EDCs with production at high volumes worldwide annually along with increasing levels in the environment.<sup>5</sup> Its estrogen mimicking property was first reported in 1936.<sup>6</sup> It is employed in the production of polycarbonate plastic and resins, which are further used for making water bottles, dental sealants, the lining of food and beverage cans. BPA monomers tend to migrate from its lining resins of food and beverage cans into food especially in temperate environments.<sup>7</sup>

Male reproductive health is targeted and deteriorated by this pollutant.<sup>8</sup> BPA has a wide range of adverse male reproductive health effects in the developmental period as well as adulthood by changing hormonal balance or direct dystrophic effects on testicular tissue.<sup>9</sup> It also affects Leydig cells directly and via hypophysial-gonadal axis to decrease serum testosterone production and release.<sup>10</sup>

BPA has weak estrogenic effects and it acts both on the surface and nuclear estrogenic receptors. It also causes changes in the expression of hormone binding proteins.<sup>11</sup> Oxidative stress is another mechanism by which it has deteriorative effects on the process of spermatogenesis even at low doses.<sup>12</sup>

Chronic low dose exposure of this pollutant has been well studied but acute high dose effects have been well investigated. This study has been conducted to determine the acute high dose effects of BPA on testicular tissue and to determine the tolerance of rodents to high oral doses of this compound.

## Methodology

Twelve rats of age 6 to 8 weeks and weight 180-200gm were procured from the animal house of the University of Health Sciences, Lahore. The initial body weight was noted while marking the animals with numbers and randomly dividing them into three groups of four rats each. The animals were allowed to acclimatize for 2 days before experiments. Free access to water and food was provided at room temperature ( $24^{\circ}\pm 5^{\circ}$ ) and normal conditions of humidity (45%  $\pm 5\%$ ).

The dosage was administered at 10 am daily.

**Group A** (control) was given corn oil 5ml/kg/day orally for 7 days

**Group B** (BPA-100) was given BPA 100mg/kg/day in corn oil orally for 7 days

**Group C** (BPA-200) was given BPA 200mg/kg/day in corn oil orally

At the end of the experimental duration, the animals were weighed before sacrifice. The pair of testes of each rat was dissected out, weighed and fixed in Buoin's solution.

**Table 1: Independent-sample t Test showing two-tailed significance of difference between mean Final body weights of animals in groups A and B.**

Parameter of Study	Groups	Mean $\pm$ SD	Standard Error of Mean (SEM)	Mean Difference	p-value
Final Body Weight of Rats (gm)	A	190.875 $\pm$ 4.028	2.014	3.900	0.475
	B	186.975 $\pm$ 9.416	4.708		

\*p-value  $\leq 0.05$  is considered statistically significant

Tissue processing was done and 4 $\mu$ m thick sections were cut. H&E stained sections were examined to assess the Johnson score of tubules. PAS staining was done to examine the basement membranes of seminiferous tubules.

The data were entered and analyzed using SPSS version 21. Quantitative parameters like body weights paired weight of testes and Johnson scoring was analyzed using the Independent-sample t test and two-tailed significance was calculated between both groups. The qualitative parameters like basement membrane integrity was analyzed using Pearson Chi-square test and association of the histological parameter and groups was assessed. Frequencies and percentages were given for the parameter of basement membrane and mean with standard deviation and standard error of means were given for the body weight, testicular weight and Johnson score. P-value  $\geq 0.05$  was considered statistically significant.

## Results

Animals of group A were healthy. Group B and C rats showed irritability and decreased appetite which was more marked in the higher dose group C. Group C rats developed marked fatigue and loss of appetite before they expired after 3-4 days of intervention and investigation proceeded with two groups. The rats of group C showed signs of diarrhea, muscles paralysis and difficult respiration and died within 2 to 3 hours of development of these symptoms.

The initial body weight of animals in both groups started from the same level. The mean initial weight of group A rats was  $182\text{gm} \pm 3.162$  and that of group B was  $185\text{gm} \pm 4.242$  ( $p=0.300$ ). There was no significant difference in mean final body weight of both groups ( $p=0.475$ ). Mean final body weight of group A was  $190.875\text{gm} \pm 4.028$  and that of group B was  $186.975\text{gm} \pm 9.416$  (Table I).

There was no statistical difference between both groups regarding this parameter ( $p=0.499$ ). Mean paired weight of gonads in group A was  $2.445\text{gm} \pm 0.171$  and that of group B was  $2.367\text{gm} \pm 0.130$  (Table II).

**Table II: Independent-sample t Test showing two-tailed significance of difference between mean Paired weight of testes of animals in groups A and B.**

Parameter of Study	Groups	Mean $\pm$ SD	Standard Error of Mean (SEM)	Mean Difference	p-value
Paired Weight of Testes (gm)	A	2.445 $\pm$ 0.171	0.086	0.775	0.499
	B	2.367 $\pm$ 0.130	0.065		

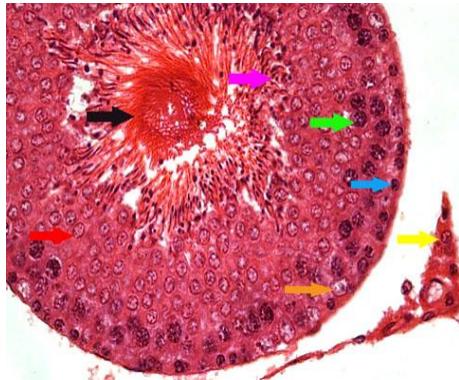
\*p-value  $\leq 0.05$  is considered statistically significant

**Table III: Independent-sample t Test showing two-tailed significance of difference between mean Johnson score of seminiferous tubules in groups A and B.**

Parameter of Study	Groups	Mean $\pm$ SD	Standard Error of Mean (SEM)	Mean Difference	p-value
Johnson Score of Seminiferous tubules	A	9.612 $\pm$ 0.281	0.140	3.297	0.001*
	B	6.315 $\pm$ 0.177	0.088		

\*p-value  $\leq 0.05$  is considered statistically significant

Statistically significant difference was observed in this parameter (p=0.001). Mean Johnson score of group A was  $9.612 \pm 0.281$  and that of group B was  $6.315 \pm 0.177$  (Table III, Figure 1-2)



**Figure 1.** Photomicrograph of seminiferous tubule of group A with Johnson score 10. Sertoli cells (orange arrow), Leydig cells (yellow arrow), Spermatogonia (blue arrow), Primary spermatocytes (green arrow), Round spermatids (red arrow), Elongated spermatids (pink arrow) and Spermatozoa in lumen (black arrow) are identified clearly. H&E. X400



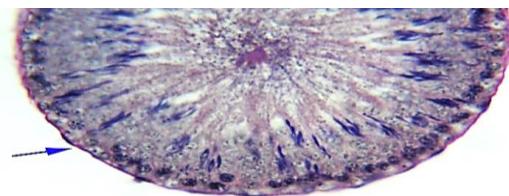
**Figure 2.** Photomicrograph of seminiferous tubule of group B with Johnson score of 6. Sertoli cells (orange arrow), Leydig cells (yellow arrow), Spermatogonia (blue arrow), Primary spermatocytes (green arrow) and desquamating spermatids (red arrow) are identifiable. H&E. X400

Basement membrane integrity was disturbed at a significant level in the treatment group (p=0.014). Out of 64 sections of seminiferous tubules assessed in group A, 53 were intact (82.8%) and 11 were disrupted (17.2%). In group B, 41 sections showed intact basement membranes (64.1%) and 23 were disrupted (35.9%) from 64 sections observed Table IV (Figure 3-4)

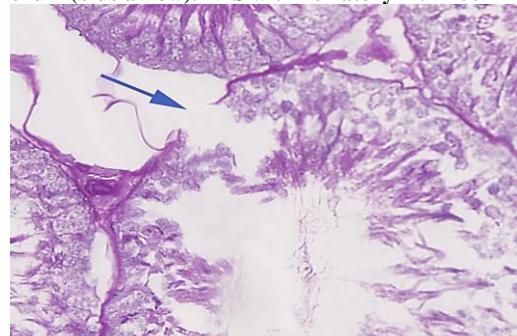
**Table IV: Pearson Chi Square Test showing number and percentages of intact and disrupted basement membranes in groups A and B.**

Groups	Basement membrane n/%age within groups		Total	p-value
	Intact	Disrupted		
A	53/ 82.8%	11/ 17.2%	64/ 100%	0.014*
B	41/ 64.1%	23/ 35.9%	64/ 100%	

\*p-value  $\leq 0.05$  is considered statistically significant



**Figure 3:** Photomicrograph of seminiferous tubule of group A showing Intact basement membrane and normal seminiferous epithelium (blue arrow). PAS with hematoxylin. X400



**Figure 4.** Photomicrograph of seminiferous tubule of group B showing Disrupted basement membrane along with sloughing of seminiferous epithelium (blue arrow). PAS with hematoxylin. X400

## Discussion

Male infertility has increased worldwide.<sup>13</sup> BPA, a synthetic carbon based plasticizer is now well known for its hazardous effects on reproductive health in humans as well as animals.<sup>14</sup> Male reproductive organs are mainly affected and disruption of testicular steroidogenesis and spermatogenesis are the primary adverse effects of this contaminant.<sup>15</sup>

Previous studies have shown that the xenoestrogenic effects of BPA are attributed to its affinity to the estrogenic as well as androgenic receptors. Adverse effects at low and high doses, on spermatogenesis and Leydig cells, have also been documented via increased oxidative stress<sup>16</sup>. The current study was designed to assess the acute high dose effects of BPA on testicular tissue of rodents.

In the present study, BPA had a negative effect on the behavior and appetite but the bodyweight of animals treated insignificantly decreased in the short period of intervention. Morgan et al (2014) also reported no change in body weight when they treated rats with BPA (25mg/kg/day).<sup>17</sup> Low (0.05mg/l) and high (5mg/l) doses of BPA given in drinking water for a longer duration showed decreased body and testicular weights in male prepubertal rats.<sup>18</sup>

There was insignificant decrease in the paired weight of testis that is also attributed to the short duration of the experiment. A statistically significant decrease was observed in animals after exposure to BPA (25mg/kg/day) for a month. BPA decreased the testosterone levels by decreasing the number of Leydig cells while it also caused a decrease in tubular diameter, both factors contributing to the decrease in testicular weight.<sup>19</sup>

Spermatogenesis is a sensitive process affected by exposure to drugs, radiation and pollutants.<sup>20</sup> In the present study, Johnson score of the BPA treated animals significantly decreased as compared to control. Studies have shown that doses of BPA as low as 20 µg/kg/day and 200µg/kg/day have been reported to cause negative changes in seminiferous epithelium and disruption of Blood Testis Barrier (BTB). BPA has been shown to directly target the junctions of Sertoli cells, making cells of different stages of spermatogenesis vulnerable to the toxic effects.<sup>21</sup>

Increased oxidative stress has been attributed to the mechanism of toxicity produced by pollutants like

bisphenols<sup>22</sup>. The present study showed significant disruption of basement membranes in rats exposed to BPA. Basement membrane damage results from protein oxidation and lipid peroxidation during high oxidative stress state.<sup>23</sup> The same mechanism may be attributed to the results of our study.

In this study, a high dose of BPA has shown significant toxic effects on testicular histology. As the plastic and paper industries are significantly growing in developed as well as developing countries, there are increased environmental concentrations of synthetic xenoestrogenic compounds like bisphenols and phthalates. Occupational exposure leads to higher levels of contamination through all routes.<sup>24</sup> The data regarding BPA exposure and adverse effects is still deficient and needs more investigation.

## Conclusion

The testicular histology was markedly deteriorated by exposure to the high dose of BPA as shown by a significant decrease in Johnson score and increased disruption in basement membranes of the seminiferous tubules. Hence, it is concluded that high dose exposure of BPA given for short duration causes testicular toxicity in albino rats.

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