Introduction

Smoking has been frequently associated with male infertility (1, 2). It reduces sperm density in ardent smokers compared with non-smokers (3). The percentages of motile and normal sperms are also reported to be lower among male smokers because of the increased generation of reactive oxygen species (ROS) (4). The effect of nicotine on the male reproductive system is a rising concern, as a previous study indicated that nicotine caused a decrease in the number of spermatocytes and spermatids (5). The testes are sensitive to cigarette smoke, and long-term exposure to it alters the morphology of spermatogenic cells and sperm

Abstract

**Background:** Nicotine is a major toxic and hazardous component of cigarette smoke, and it has been widely used in nicotine replacement therapy (NRT). This study was aimed to investigate the effects of chronic low-dose nicotine on sperm characteristics and reproductive organ integrity in adolescent male Sprague–Dawley rats.

**Methods:** Twelve rats were equally divided into two groups. Group I received normal saline, and group II received 0.6 mg/kg body weight nicotine intraperitoneally for 28 consecutive days. At the end of the experimental period, sperm was collected for sperm characteristic evaluation, and the testes and prostate were isolated for biochemical and morphological analysis. The effects of nicotine on the body and reproductive organ weights of the animals were evaluated.

**Results:** Chronic nicotine treatment significantly (P < 0.05) altered the sperm count, motility, viability, and morphology, and remarkably increased the malondialdehyde (P < 0.001) and advanced oxidation protein product (P < 0.05) levels in the testes and prostate of nicotine-treated group compared to control group. Moreover, nicotine caused a significant decrease (P < 0.05) in the superoxide dismutase activity of the testes. No significant differences were observed in the reduced glutathione level in both of the testes and prostate of nicotine group compared with control group. Nicotine also induced histopathological alteration in the testes.

**Conclusion:** A low-dose nicotine exposure at 0.6 mg/kg caused detrimental effects on sperm characteristics and induced oxidative stress in the testes and prostate.

**Keywords:** infertility, prostate, testes, sperm, male, reproductive
At the end of the study, all rats were anesthetised with sodium thiopental and dissected. The weights of the prostate, seminal vesicle, cauda epididymis, and testes were recorded. Sperm was collected from the left and right cauda epididymis by immediately placing the cauda epididymis in 2 mL of Hank’s Buffered Salt Solution enriched with 0.5 % bovine serum albumin (Sigma Chemicals, St. Louis, USA) and pre-warmed at 37 °C. Both cauda epididymis were minced and centrifuged at 1000 rpm at 4 °C for 3 minutes. The testes and the prostate were homogenised in 1.15% potassium chloride solution and centrifuged at 4 °C for 20 minutes. The homogenates were used for further biochemical analysis.

**Biochemical Assays**

The levels of lipid malondialdehyde (MDA) were determined by the method of Stocks and Dormandy (13), advanced oxidation protein product (AOPP) by the method by Witko et al. (14), superoxide dismutase (SOD) activity by the method of Beyer and Fridovich (15), and reduced glutathione (GSH) levels by the method of Ellman (16).

**Sperm Characteristic Analysis**

Sperm count was calculated by a Makler Counting Chamber. About 10 µL of sperm suspension was taken and calculated for all grids under a 10x magnification of a light microscope. Sperm count was expressed as x 10^6 cells/mL. Sperm motility was evaluated by counting the motile and the non-motile sperms and assessed up to 200 sperms under a 10x magnification of a light microscope. Data are shown as the percentage of sperm motility. Sperm viability was determined by a routine gold standard method suggested by the World Health Organisation (17).

**Morphological Analysis of Male Reproductive Organs**

The testes and the prostate were fixed with 10% formalin solution, dehydrated in graded ethanol, and embedded in paraffin wax. Both the testes and the prostate tissues were sectioned at 5 µm and stained with hematoxylin and eosin. The slides were evaluated under light microscopy at 10x and 40x magnifications.

**Statistical Analysis**

Statistical analysis was performed using the Statistical Package for Social Sciences version 19.0. The differences between the control group
and the treated group were statistically evaluated using independent \( t \)-test. All of the results were expressed as the mean (SD).

**Results**

**Weight of the Body and Reproductive Organs**

Table 1 indicates that treatment with nicotine did not cause any significant changes \((P > 0.05)\) in the weight of the body and reproductive organs, such as the prostate, seminal vesicle, caudal epididymis, and testes.

**Sperm Characteristics**

Table 2 shows the significant decline of the sperm count, sperm motility, and sperm viability \((P < 0.05)\) in the nicotine-treated groups compared with the control group. A significant increase in the abnormal sperm morphology was found in the nicotine group \((P < 0.001)\).

**Antioxidant Status and Oxidative Damage Markers in the Reproductive Organs**

Table 3 shows that treatment with nicotine caused a significant \((P < 0.05)\) decrease in the SOD activity in the testes and an insignificant decrease in the prostate compared with that in the control group. The MDA levels and AOPP in the prostate \((P < 0.05)\) and the testes \((P < 0.001)\) were augmented in the nicotine-treated rats compared with the non-nicotine treated rats. No significant changes were observed in the GSH levels in the testes and the prostate.

**Histopathology of the Reproductive Organs**

**Testes**

Figure 1A and Figure 1B illustrate the normal architecture of the seminiferous tubules and the spermatogenesis process. The seminiferous tubules contain abundant spermatogenic cells, which are surrounded by peritubular myoid cells, and the Leydig cells are located in the interstitial tissue between the seminiferous tubules (1B). Figure 1C and Figure 1D show that the spermatogenesis process is reduced in the lumen of the seminiferous tubules. Some of the seminiferous tubules exhibit a complete absence of spermatogenic cells, acinus, and fibromuscular stroma with normal secretory epithelium. The architecture of the seminiferous tubules is markedly shrunken.

**Table 1.** Body and reproductive organ weight in normal and nicotine rats for 28 days of treatment

<table>
<thead>
<tr>
<th>Organs (g)</th>
<th>Experimental groups</th>
<th>( t )-statistic (df = 12)</th>
<th>( P )-value&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control group&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Nicotine group&lt;sup&gt;a&lt;/sup&gt;</td>
<td></td>
</tr>
<tr>
<td>Final body weight</td>
<td>270.83 (37.80)</td>
<td>279.20 (19.71)</td>
<td>3.817</td>
</tr>
<tr>
<td>Prostate</td>
<td>0.57 (0.08)</td>
<td>0.49 (0.15)</td>
<td>1.418</td>
</tr>
<tr>
<td>Seminal vesicle</td>
<td>0.74 (0.23)</td>
<td>0.65 (0.15)</td>
<td>2.173</td>
</tr>
<tr>
<td>Cauda epididymis</td>
<td>0.48 (0.07)</td>
<td>0.46 (0.11)</td>
<td>1.307</td>
</tr>
<tr>
<td>Testes</td>
<td>2.58 (0.22)</td>
<td>2.67 (0.49)</td>
<td>1.037</td>
</tr>
</tbody>
</table>

<sup>a</sup>Mean (SD). <sup>b</sup>Independent \( t \)-test. \( n \): Number of sample. \( t \): \( t \)-statistics. df: Degree of freedom.

**Table 2.** Sperm characteristics in normal and nicotine rats for 28 days of treatment

<table>
<thead>
<tr>
<th>Sperm characteristics</th>
<th>Experimental groups</th>
<th>( t )-statistic (df = 12)</th>
<th>( P )-value&lt;sup&gt;b&lt;/sup&gt;</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control group&lt;sup&gt;a&lt;/sup&gt;</td>
<td>Nicotine group&lt;sup&gt;a&lt;/sup&gt;</td>
<td></td>
</tr>
<tr>
<td>Sperm count (X10&lt;sup&gt;6&lt;/sup&gt;/ml)</td>
<td>15.28 (0.52)</td>
<td>13.05 (0.35)</td>
<td>3.986</td>
</tr>
<tr>
<td>Sperm motility (%)</td>
<td>58.67 (2.37)</td>
<td>44.13 (1.27)</td>
<td>3.632</td>
</tr>
<tr>
<td>Sperm viability (%)</td>
<td>65.58 (3.14)</td>
<td>34.50 (1.95)</td>
<td>2.273</td>
</tr>
<tr>
<td>Abnormal sperm morphology (%)</td>
<td>3.17 (0.40)</td>
<td>8.92 (0.51)</td>
<td>3.093</td>
</tr>
</tbody>
</table>

<sup>a</sup>Mean (SD). <sup>b</sup>Independent \( t \)-test. \( n \): Number of sample. \( t \): \( t \)-statistics. df: Degree of freedom.
Table 3. Oxidative stress markers in reproductive organs in normal and nicotine rats for 28 days of treatment.

<table>
<thead>
<tr>
<th></th>
<th>Testis Experimental groups</th>
<th>Prostate Experimental groups</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control group (n = 7)</td>
<td>Nicotine group (n = 7)</td>
</tr>
<tr>
<td>SOD (U/min/mg protein)</td>
<td>2.16 (0.44)</td>
<td>1.44 (0.40)</td>
</tr>
<tr>
<td>GSH (nmol/mg protein)</td>
<td>106.52 (13.91)</td>
<td>111.52 (7.41)</td>
</tr>
<tr>
<td>MDA (nmol/g protein)</td>
<td>0.68 (0.11)</td>
<td>1.57 (0.18)</td>
</tr>
<tr>
<td>AOPP (µmol/g protein)</td>
<td>48.23 (2.23)</td>
<td>57.04 (2.85)</td>
</tr>
</tbody>
</table>

SOD: Superoxide dismutase. GSH = Glutathione. MDA = Malondialdehyde. AOPP = Advanced oxidation protein products.

*S*Mean (SD). **Independent t-test. n: Number of sample. t: t-statistics. df: Degree of freedom.

Figure 1. Testes: Histology of the testes of control and nicotine groups (H&E staining) (Magnification A and C ~ 10×; Magnification B and D ~ 40×). Figure 1A shows that normal architecture of the seminiferous tubules and spermatogenesis process. Figure 1B shows that the seminiferous tubules contain of abundant spermatogenic cells (G) and spermatozoa (S), which has surrounded by peritubular myoid cells (MY) and leydig cells (L) were located in the interstitial tissue between the seminiferous tubules. Figure 1C shows that architecture of seminiferous tubules were markedly shrunken and distorted (DT) and spermatozoa production was reduced in the lumen of seminiferous tubules (AS). Figure 1D shows that some of the seminiferous tubules show complete absence of spermatogenic cells and spermatozoa (AS).
and distorted. Some of the seminiferous tubules exhibit a complete absence of spermatogenic cells (Figure 1D).

Prostate

Figure 2A and Figure 2B illustrate the normal prostate gland, which consists of the acinus and fibromuscular stroma with normal secretory epithelium, from control group. The prostate exhibits no change in histology in the nicotine-treated group (Figure 2C and Figure 2D).

Discussion

Researchers have reported the detrimental effect of cigarette smoking on male fertility (18, 19, 20). Nicotine exposure reduces the weight of the testis and the number of spermatocytes and spermatids. Nicotine also affects the testosterone level, pituitary gonadotropins, and testicular antioxidant status. Moreover, nicotine contributes to ROS generation in the testis (21).

SOD protects tissues against oxygen free radicals by catalysing the deletion of superoxide radical (O$_2^-$), which harms the membrane and biological structures (22). GSH is an important inhibitor of free radical mediated lipid peroxidation. It is involved in several reactions in the body and is one of the most prominent non-enzymatic antioxidant (23). SOD and GSH are important in scavenging the enzyme that eliminates the toxic free radicals in vivo (24). Unlike in the control group, the SOD

![Figure 2. Prostate: Histology of the prostate of control and nicotine groups (H&E staining) (Magnification A and C ~ 10×; Magnification B and D ~ 40×). Figure 2A and Figure 2B illustrated that normal prostate of the control group. Figure 2B shows that prostate gland consists of acinus (A) and fibromuscular stroma (FS), secretory epithelium (SE). Figure 2C and Figure 2D show that normal acinus, fibromuscular stroma and secretory epithelium from the nicotine treated group. No changes were found in the prostate of nicotine treated group.](image-url)
Spermatozoa are susceptible to oxidative stress-induced damage because of the presence of large quantities of PUFA in their membrane, especially diPUFA (phospholipids esterified with two PUFAS), which is especially found in the retina and certain parts of the brain aside from the sperm (30). The abnormal morphology of the sperm structure can be linked to the lipid peroxidation damage to the PUFA component caused by free radicals (31). Abnormal sperm production may be due to the disruption of spermatogenesis by nicotine action (19).

The testes and the prostate are closely related to sperm quality and play an important role in male fertility. The oxidative damage and the morphological alterations of the testes are associated with the reduction in sperm quality (32). The prostate secretes fluids that make up the semen and provides nutrient to the sperm. Lipid peroxidation of the prostate may affect the secretion of semen and the sperm characteristics. According to Zhang et al. (3), when the semen volume decreases, sperm density and sperm viability also decrease. Therefore, any alteration in the structure and function of these reproductive organs can lead to male infertility by affecting the sperm quality.

Conclusion

Chronic nicotine treatment at a dose of 0.6 mg/kg has detrimental effect on sperm characteristics and can cause oxidative damage to the testes and the prostate.

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Authors’ Contributions

Conception and design: SBB, SZ, ESL, JM
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Drafting of the article: FFJ, ESL
Critical revision of the article for important intellectual content: IST, ESL, JM
Final approval of the article: SBB
References


Chronic nicotine treatment induces male reproductive damage


