How the Pregabalin (Lyrica) Administration as a Abuse Drug Can affects the Reproductive Health of male Wistar Rat

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ABSTRACT
Drugs addiction considered a massive problem persist in diverse populations everywhere the world. Our work aimed to illustrate the correlation between drug addiction and reproductive function in an animal model, also the potential impact of pregabalin (Lyrica) intake on spermatozoa formation process and sexual hormone levels. In this study, we used 14 adult healthy rats that allocated into two groups (n = 7) as control and treated that orally administrated with pregabalin (23.7 mg/kg) for 30 subsequent days. Reproductive hormones, spermatozoa parameters (motility and morphology), lipid peroxidation (MDA), nitric oxide, total antioxidant activity, DNA damage and histopathological investigation were performed. The results revealed that pregabalin addiction had a harmful impact on the hypothalamus-pituitary-gonadal axis of male rats through hindered hormones secretion, raised reactive oxygen species, affect antioxidant enzymes, triggered DNA damage and distorted testicular histology. Finally, we found that addiction of Lyrica caused adverse impact on the male reproductive health and subsequently affect fertility.

Keywords: Pregabalin (Lyrica), Abuse drug, Male, Wistar rat, Reproductive toxicity.

INTRODUCTION
The genitourinary organs can be influenced by drugs, many studies investigate the impacts of drugs on this structure is of extraordinary significance. Several experimental works have shown that oral opium can diminish luteinizing hormone (LH), dihydrotestosterone, and follicle stimulating hormone (FSH), and can cause hypogonadism in 89% of users. The predominance of erectile brokenness and diminished libido is additionally significantly higher in sedate abusers 1.

Pregabalin is an alkylated simple of c-aminobutyric acid (GABA) and basically linked to gabapentin 2. In expansion, GABA mimetic properties have been appeared in rats 3. A study by Grosshans et al. found that unlawful utilize of pregabalin was communal among opioid-addicted persons 4, 5. Lyrica is not a narcotic or an opioid. Lyrica is in a kind of medicines named anticonvulsants.

Various components can disrupt spermatogenesis progression and decrease sperm's property and amount. Effective spermatogenesis based on a multifaceted relation between endocrine, paracrine and autocrine components 6, 7. Obviously, many inherited illnesses might impede the spermatozoa formation mechanism. The non-

genetic reasons of male infertility, oxidative stress (OS) coming about from overstated generation of reactive oxygen species (ROS) is maybe the foremost identified issue.

ROS are required for capacitation, the acrosome reaction and eventually fertilization; be that as it may, diminished removal and overproduction are capable to initiate DNA harm and imperfect membrane reliability of spermatozoa, in this manner coming about in decreased fertility capacity 8, 9. An extraordinary generation of reactive oxygen species (ROS) can be harming to the sperm cells. The spermatozoa plasma membrane contains enormous quantities of unsaturated fatty acids. Hence, it is vulnerable to peroxidative damage. The lipid peroxidation abolishes the constitute of lipid ground of spermatozoa membranes and distracts sperm’s motility 9, 10.

The present study aimed to investigate the potential harmful effect of pregabalin (Lyrica) addiction on the male reproductive health and try to illustrate the exact mechanism through which the used drug can caused infertility and affect sperm function.

MATERIALS AND METHODS

Animals
14 male Wistar with the weight range of 200–220 g was obtained from National organization for drug control and research (NODCAR). The animals were reserved in an animal house for one week under well-ordered laboratory settings at temperature 22°C ± 2°C, humidity (50%–60%) and 12 h in light and 12 h in dark with permitted admission to water and food.
Experimental design
The practical work was agreed by the Cairo University, Faculty of Science Institutional Animal Care and Use Committee (IACUC) (Egypt), (CU/I/5/3/20). The rats were allocated into two groups as follows: group one (control), received distilled water and group two (treated), and administrated orally with abuse dose of pregabalin (Lyrica) 23.7 mg/kg for one month according to World Health Organization 2018 11 for four weeks day after day.

Sperm collection
After one month, the rats were sacrificed using an intraperitoneal injection of sodium pentobarbital (100 mg/kg). Then, testes, epididymis and seminal vesicle were collected, washed with saline, dried and weighted. The cauda part of the epididymis was used to evaluate sperm parameters. In each animal, right testes fixed in neutral 10% formalin for histopathological analysis and left one freeze at -20°C for further investigation. Cauda epididymis were minced, and protected in a warm Petri dish containing 5 ml physiological saline solution (Ph 7.4) at 37°C. The spermatozoa were permitted to disperse into the buffer 12.

Sperm count
For calculating the sperm, 500 μL of the sperm suspension was dilut (dilution of 1:10) with formaldehyde fixative (10% formalin in phosphate buffered saline). 10 μL from the diluted solution was placed into a hemocytometer. Hemocytometer was situated in a moist chamber for 7 min. Hemocytometer was placed on the microscope stage. Then, the sperms at the four corners of the central square were counted 13.

Sperm morphology
Eosin/ nigrosin stain was operated to evaluate spermatozoa morphology. One drop of eosin/nigrosin was added to the suspension and slightly mixed. The slides were then examined under the light microscope at ×400. A total of 300 spermatozoa were evaluated on each slide to find the anomalies of the head and tail 14.

Sperm high motility
The sperm motility was measured through the light microscope at ×400. One drop of sperm suspension was set on a glass slide. The number of the sperms with rapid progressive forward movement was computed and the percentages of high motile sperms were achieved 15.

Hormones profile
Blood were harvested from the rats by cardiac puncture in clean tube, centrifuged at 3000 rpm for 15 min to obtain sera and then were kept at -20°C for hormones analysis. Quantitative determination of Testosterone (T), follicle-stimulating hormone (FSH) and luteinizing hormone (LH) level were measured using ELISA kits specific for rats (SunLong Biotech Co., LTD).

Redox status
Reactive oxygen species evaluated using reagent kits obtained from Bio Diagnostic (Egypt). Freeze testes minced in PBS (1:10 ml). The suspension was centrifuged at 3000 rpm for 20 minutes, and the supernatant was used to test MDA level by means of Satoh 16, NO concentration based on Montgomery and Dymock 17 and total antioxidant capacity as designated by Koracevic et al. 18.

Comet assay
DNA injury was estimated using the single-cell gel electrophoresis technique 19. The DNA damage were calculated through comet score software, in which the DNA % in the tail, the tail length and the tail moment might be acquired.

Histological examination
For light microscopical investigation, the testes of two groups were fixed by immersion in 10% formalin. All samples were relocated in 70 % ethanol and then dehydrated in an ascending run of ethanol, cleared in xylol and embedded in paraffin. Five μm thick sections were yielded using a rotary microtome. Histological staining was performed with Ehrlich’s hematoxylin and counterstained with aqueous eosin 20. Microscopical examination and photographing of the histological sections were implemented with AmScope microscope.

Statistical analysis
Statistical analysis was achieved by the independent t-test to reveal mean and standard error mean of all examined parameters using Statistical Package for the Social Sciences (SPSS). The differences between control and treated group at significance level of 0.05 (P<0.05).

RESULTS
Effect of Lyrica on tissues weight
The absolute and relative reproductive organs weights of different treated groups are showed in Table 1. Lyrica exposure triggered a reduction in in testes, epididymis and seminal vesicle weights.

| Parameters                       | Treated     | Control    |
|----------------------------------|-------------|------------|
| Absolute right testes weight (g) | 1.2±0.136   | 1.36±0.08  |
| Relative right testes weight (%) | 0.496±0.49  | 0.508±0.032|
| Absolute left testes weight (g)  | 1.16±0.181  | 1.33±0.098 |

Table 1: Showing effect of Lyrica on testes weight.
Relative left testes weight (%)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.48±0.071          | 0.491±0.031 |

Absolute right epididymis weight (g)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.175±0.017         | 0.183±0.016 |

Relative right epididymis weight (%)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.072±0.006         | 0.065±0.005 |

Absolute left epididymis weight (g)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.175±0.017         | 0.183±0.016 |

Relative left epididymis weight (%)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.072±0.006         | 0.065±0.005 |

Absolute seminal vesicle weight (g)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.316±0.072         | 1±0.143     |

Relative seminal vesicle weight (%)

| Group               | Control     | Treated     |
|---------------------|-------------|-------------|
| 0.316±0.072         | 1±0.217     |

Effect of Lyrica on spermatozoa

Lyrica exposure resulted in non-significant decrease in sperm motility with significant rise in the total abnormalities as banana head, abnormal tail, hookless sperm compared with the control group (Table 4 & Fig. 2).

Table 2: Showing effect of Lyrica on sperm analysis profile.

| Parameters          | Control     | Treated     |
|---------------------|-------------|-------------|
| Motility            | 41.66±2.47  | 40.83±5.06  |
| Abnormality %       | 3.33±0.954  | 12.5±1.945* |
| Banana Head         | 0.66±0.33   | 1.83±0.60   |
| Abnormal Tail       | 1.66±0.614  | 5.50±1.62   |
| Hook Less           | 0.66±0.333  | 1.83±0.307* |

Values are expressed as Mean ± SEM. The statistical differences were analyzed by independent samples t-test. * = P <0.05 compared with control.

Figure 1: Showing effect of pregabalin on testes weight.

Effect of Lyrica on reproductive hormones

Lyrica administration revealed significant imbalance in the reproductive hormones' levels in comparison by control group. The testosterone, LH and FSH levels were decreased in the treated rats when compared with control group. Also treated group revealed non-significant increase in alkaline acid phosphate (ACP) in comparison with the control one (Table 3).

Table 3: Showing effect of Lyrica on reproductive hormones.

| Parameters          | Control     | Treated     |
|---------------------|-------------|-------------|
| Testosterone (ng/ml)| 0.456±0.044 | 0.22±0.036* |
| FSH (mIU/ml)        | 4.35±0.375  | 3.05±0.028* |
| LH (mIU/ml)         | 3.47±0.306  | 2.55±0.028* |

Values are expressed as Mean ± SEM. The statistical differences were analyzed by independent samples t-test. * = P <0.05 compared with control.

Effect of Lyrica on redox status

Lyrica exposure increase the malonaldehyde (MDA) level non-significantly in comparison with the control group. Similarly, addiction of lyrica create a significant rise in content of nitric oxide (NO). Also, lyrica taken elevate the total antioxidant activity non-significantly as compared with the control group in all examined tissues (Table 4).

Table 4: Showing effect of Lyrica on ROS markers.

| Parameters          | Control     | Treated     |
|---------------------|-------------|-------------|
| MDA (nmol/g. tissue)| 1126.26±70.94 | 1225.42±55.77 |
| NO (μmol / L)       | 7.23±0.294  | 13.95±0.196* |
| TAC (mM / L)        | 0.7488±0.118 | 0.9840±0.008 |

Values are expressed as Mean ± SEM. The statistical differences were analyzed by independent samples t-test. * = P <0.05 compared with control.

Figure 2: Showing effect of pregabalin on sperm morphology. A& B: control. C-F: Amorphous head. G-J&M: bent tail. K-M: sperm with Hook defect.
Table 5: Showing effect of Lyrica DNA damage index.

| Groups       | Control          | Treated          |
|--------------|------------------|------------------|
| Comet %      | 8.65±0.375       | 3.45±0.837*      |
| Tail Length  | 5.07±0.268       | 12.61±0.827*     |
| Tail DNA %   | 7.48±0.144       | 8.04±0.575       |
| Tail Moment  | 0.44±0.065       | 1.012±0.024*     |
| Olive Moment | 0.687±0.022      | 1.32±0.28*       |

Values are expressed as Mean ± SEM. The statistical differences were analyzed by independent samples t-test. * = P <0.05 compared with control.

Effect of Lyrica on DNA damage

We used the comet assay (single cell gel electrophoresis) to inspect the influence of Lyrica on DNA by evaluating the DNA tail, tail length and tail moment in the examined tissue (Fig. 3 and Table 5). The treated group exposed to Lyrica displayed a significant change in the comet parameters compared with the control rats except for Tail DNA % showed insignificant increase as compared with the control group.

Figure 3: Showing effect of pregabalin on DNA. (A) Control and (B) Treated.

Histopathological results

Histological investigation of testis of control rat stained with H&E presented standard form of seminiferous tubules and interstitial tissue. Each tubule was lined with stratified epithelium (germinal cells) and supportive Sertoli cells. The germinal cells were organized in several layers (spermatogenic cells). Leydig cells were found between stroma connective tissue. Animals taken Lyrica displayed different histopathological alterations. The seminiferous tubules have irregular outline, degenerated tubules were noticed with absence of spermatozoa. Most of the seminiferous tubules revealed injured and disordered spermatogenesis cells and the impaired germ cells were exfoliated in the lumen. Furthermore, nonexistence of spermatozoa was obviously documented. Plentiful spermatocytes seemed with pyknotic nuclei and many vacuoles were observed in the seminiferous tubules. Some seminiferous tubules be seen cracked (Fig. 4).

Figure 4: Showing effect of pregabalin on histology. Photomicrographs of a section in the testis of control albino rats (A&B); the normal histological structure, multiple number of seminiferous tubules (ST) lined by many kinds of germinal epithelium (spermatogonia 1, spermatocyte 2, spermatid 3). Mature sperms in the lumen (4) and the interstitial tissue (*) in between the tubules, which contain Leydig cells (LC). Treated group; (C) degenerated and disorganized seminiferous tubule (ST) without spermatozoa and vacuoles between interstitial tissue (*). (D-F) pyknotic spermatogonia (arrow) and primary spermatocytes with karyolysis signs (arrowhead), wide lumen devoid of sperms.

DISCUSSION

In this study, we investigated the correlation between drug abuse and sexual performance also the impending influence of pregabalin (Lyrica) intake on spermatozoa formation and reproductive hormones activity.

Drugs addiction considered a massive problem persist in diverse populations everywhere the world. Lyrica is not a narcotic or an opioid. Lyrica is in a kind of medicines named anticonvulsants.

The outcomes showed that pregabalin (Lyrica) reduced sperm motility, normal sperm morphology, augmented testicular DNA damage, and prompted histopathological
alterations in testicular tissue. These harmful impacts have been complemented by provoked oxidative stress in testicular tissue and the change of serum hormone levels that participate a part in the spermatogenesis progression.

Some findings have implied that the drug misuse have destructively influence male fertility, with an effect on hypothalamic-pituitary-gonadal axis, spermatogenesis, and sperm function, Leydig cells, Sertoli cells and in testicular tissues.7,21

Organ weights are susceptible indicators to reveal toxicity after chemical exposure.22 Tissue weight change mirror the distractions of the reproductive system functions.23 In our study, significant decreasing in absolute and relative testis and epididymis weights after pregabalin (Lyrica) intake were not detected.

The testicular tissue weights are related with the Sertoli cells number and spermatozoa formation consequently the testis size is indication of the germinal cells number in the testis.12 This might be as a result of free radicals’ creation and ROS by pregabalin (Lyrica) and effect of these detrimental elements on testis susceptible cells. Spermatozoa motility, and morphology are pointers used to estimate semen quality, testicular function and verify reproductive toxicity.23 A decline in sperm motility and sperm morphology abnormalities are a significant parameter of chemical caused infertility.24,25

Regulatory authorities like EPA, FDA, OECD, WHO, and ICH underline the significance of the sperm head, sperm midpiece and tail abnormalities especially twisted tail and bent/spiral tail are related to infertility.25,26

FSH, LH, and testosterone have parts in male reproductive functions conservation and so, hormone levels determination is essential in reproductive toxicity reports.25 It is proven that LH and FSH are released below the regulator hypothalamic gonadotropin-releasing hormone from the anterior pituitary. LH motivates the testosterone secretion from Leydig cells, and testosterone is mandatory for secondary sexual appearances and spermatogenesis. FSH controls the spermatozoa production in Sertoli cells.25,27 The hypothalamic-pituitary gonadal axis can be influenced by several agents. Chemicals comprising drugs can lessen fertility and cause infertility by disordering the ordinary function of this axis.28 It has been revealed that antiepileptic drugs affect hypothalamic pituitary gonadal axis and trigger reproductive malfunction.29,30 In our study, fallen serum FSH, LH, and testosterone levels were detected after drug administration. Previous studies presented that LH and FSH levels were not changed.25,31

In addition, a reduction of testosterone level is associated to the spermatogenic cell damage and spermatogenesis distraction, and subsequently initiate reproductive power disordered.25,27,28

Opioids play on the hypothalamic-pituitary axis by hindering the GnRH discharge, that suppress FSH and LH release accordingly cause spermatogenesis impairment and decline testosterone concentrations.32 Vuong et al.33 stated that opioid- convinced hypogonadism. Other articles imply that sperm concentration and quality are damaged in opioid users, augmented DNA fragmentation level and lowered catalase (CAT) and superoxide dismutase (SOD) activity were seen in addict men competed to healthy persons.7,34

Pregabalin drug decreases serotonin discharge in the synaptic cleft. Meanwhile serotonin is the required mediator in melatonin creation, thus causing melatonin reduction. Melatonin is an antioxidant and has an important role in defending the testicular tissue against damage induced by ROS. Low melatonin activity causing lessening in testosterone synthesis and secretion by lessening the glutathione peroxidase (GPx) enzyme.35,36

Prior paper presented that the pituitary gonadotropins, serum FSH, LH, and PRL hormone levels reduced using two doses of pregabalin, these clarifications are arrangement with our study, and is due to the antiepileptics (ADEs) may have an effect on the gonadal level.37,38, Pregabalin (PGB) has the ability to hinder the central nervous system activity, that controls physiological and behavioral consequences correlated with normal reproductive performance through hormonal signals.38,39

Harden and Pennell40 indicated that ADEs may intermingle with gonads. Since PGB drug blockade calcium channels, that analogous neurochemical mechanisms are involved in the interaction of these drugs with hypothalamic neurohormones synthesis as gonadotropin-releasing hormone (GnRH), so PGB may be affects hypothalamus because GnRH discharge from neurons is based on the depolarization - caused influx of extracellular calcium and because PGB effects on Ca2+ channels by blockading or inhibiting Ca2+ influx on hypothalamus that is susceptible to change the GnRH pulsations and trigger reduction in LH and FSH levels.38,41

Findings demonstrated that testosterone clearly affects the Sertoli cells. Sertoli cells providing nutrients for dividing spermatogenic cells. They produce many growth factors and transferring proteins which has a critical role in cell division and spermatozoa formation.36 Concerning the testosterone role on spermatogenesis, reduction in this hormone secretion induced sperm density decrease. Mammalian spermatozoa has high unsaturated fatty acids quantity that are main substrates in oxidation. In normal situations antioxidant mechanisms are participate in reproductive tissues and inhibit oxidative injure in different gonadal cells and developed sperm.42 The previous reports revealed that free radical’s creation directly disturbs sperm proliferation, activity and fertility.36

Hypothalamus release Luteinizing hormone releasing hormone (LHRH) that initiates the secretion of LH from pituitary gland then testosterone releasing from the testes. We did not identify if the decreased LH was because of pituitary malfunction or the decreasing LHRH release in...
the treated group. It is probable that LHRH, LH, and testosterone were distressed by the PGB.1,43.

The oxidative stress induce lipid peroxidation affects semen quality, and inducing male infertility.25,44,45. Oxidative stress induced a reduction in intracellular ATP levels, apoptotic factors creation resulting in mitochondrial membrane disturbance, protein phosphorylation disorder, rise in membrane permeability, and spermicidal molecules formation, impairment of acrosome membrane so reduces semen quality, concentration, motility and morphology 44. Moreover, antioxidant defense mechanism deficiency participates to the sperm vulnerability.25,46. In our study, reduced total antioxidant activity levels and increased MDA and NO levels were observed following PGB administration signifying that PGB induced oxidative stress in testicular tissue.

Nitric oxide has an essential role in sperm physiology and have many undesirable effects on hypothalamic–pituitary–testicular axis.47. There is a correlation between nitric oxide and sperm acrosome and tail. Nitric oxide can reduce the sperm motility by decreasing ATP level.48. Nitric oxide can damage sperm mitochondrial membrane, thus releasing C chromosome, initiating caspase cascade activity and promoting apoptosis.12,15.

In the study of Daniel et al., severe morphologic variations of the sperms were seen in microscopic analysis of the semen of addicts, which verifies the outcomes of the existing study12,49.

Our results showed significant initiation of oxidative stress after pregabalin exposure. This comes in accordance with the results of Kamel.50 who explored the effect of chronic oral pregabalin administration for 90 days on the rat brains and stated significant decline in SOD and CAT in pregabalin administered groups.51.

Sperm DNA integrity is a clue of the sperm reproductive power.52. So, the DNA’s structural integrity was examined to estimate sperm function. The neutral comet assay is simpler, sensitive and accurate method in clarifying DNA damage double-strand breaks in human sperm.25,53.

In this study tail length, DNA percentage in tail and olive tail moment were recorded to evaluate genotoxic impact of the PGB. These parameters were particularly crucial to determine the DNA damage severity after exposure to genotoxic environmental agent.54. Many studies stated that tail moment is a clearer value to evaluate DNA damage.25,55.

Thus, PGB generated DNA damage seen in our study may be due to oxidative stress, promoting DNA histone modification. Additionally, sperm head morphology is an indirect mark of mutagenic effects induced by chemical exposure. The preceding report found a confident association between sperm head abnormalities and DNA damage, and it was discussed that imperfections in sperm head morphology developed from genetic material damage.25,56. Here sperm head abnormalities boosted by PGB intake reflected DNA damaging.

CONCLUSION

Based on the preceding elucidation we imply that Lyrica may be create imbalance redox status by producing reactive oxygen species and nitrogen with low antioxidant power that initiating cell death through interaction with the lipid of cell membranes, nucleic acid and proteins which influences cells signaling pathways that controlling programmed cell death (apoptosis and necrosis) and cell proliferation finally causing reproductive hormones levels and histopathological alternation.

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