Pregabalin induces pathological changes in cerebrum and cerebellum of Albino Rats

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Abstract

Background: Pregabalin (PGB) used as analgesic in remedy of neurogenic pains of persistent diseases, is taken into consideration as one of the maximum abused anti-epileptic pills and it's been proved that it induces addictive behaviors. The current study investigated the impact of PGB management on cerebral cortex and cerebellar cortex, in each acute and chronic toxicity. Seventy- male and non-pregnant female albino rats' 6- to 8-week-vintage divided into three principal corporations of 24 rats each had been studied. Group 1 was the control group, group 2 represented the acute toxicity group, and group three represented the hronic toxicity group. Rats had been sacrificed and examined.

Result: Cerebral cortex tissue of acute toxicity group displayed astrocytosis and dystrophic changes, while the chronic toxicity group showed degeneration, necrosis and inflammatory cell infiltrates. The cerebellum of chronic toxicity group showed affection of Purkinje cells.

Conclusion: Acute and chronic intoxication with pregabalin adversely altered the structure and thus the function of the cerebral cortex and cerebellum.

Introduction

Anti-epileptic drugs are one of the maximum abused drugs worldwide, specifically pregabalin (PGB) and gabapentin [1]. PGB is extensively used as analgesic in remedy of neurogenic ache in diabetic neuropathy [2,3]. It is likewise permitted for the remedy of generalized anxiety disorder, post herpetic neuralgia, post-traumatic strain disorder, seizures, migraine and fibromyalgia [4]. Studies have proven that PGB can also be used in treatment of benzodiazepine dependence and alcohol dependence [5]. PGB is taken into consideration an "perfect psychotropic drug" to have unique effects, which includes sedation, euphoria, dissociation, and psychoactive effects [6, 7]. Thus, PBG is considered one of the most regularly abused substance [8]. Pregabalin is classified as a CNS depressant and its mechanism of toxicity is meant to be CNS depression with respiratory failure and cardiac complications [9]. Cases with cerebellar disorder associated with PGB intoxication had been also reported [10,11]. The mechanism for PGB-prompted cerebellar disorder stays unclear, but a few research said that PGB could result in reduced excitatory inputs from the brainstem [12]. This study was carried out to assess the histopathological changes of PGB on cerebrum and cerebellum in a rat model.

Materials And Methods

Drugs and chemicals:

All drugs and chemical purchased were listed in supplementary table 1.

Test animals:

This experimental work was carried out on 72 male and non-pregnant female adult Albino rats 6- to 8-week-old weighing 100–150 grams. Rats were housed in plastic cages under standard pathogen free conditions and quite non-stressful special room. Every five rats of same sex were placed in each cage, so that stress of isolation or overcrowding was avoided. The rats are divided into 3 main groups of 24 rats each: **Group I (Control group; n=24)**: which is further subdivided into subgroup IA (control group for acute toxicity of Pregabalin; n=12) & IB (ontrol group for chronic toxicity of Pregabalin; n=12). **Group II (Acute toxicity group; n=24)** were given a single dose of PGB (5000 mg/kg body weight) orally by gavage. After 24 hours, rats were sacrificed and subjected to histopathological investigations. **Group III (Chronic toxicity group; n=24)** received chronic doses of PGB as following: 500 mg/kg orally gavage (equivalent 1/10 LD50) for 12 weeks [13]. The drug solution was given with a volume of 1-2 ml. using a stomach tube gavage.

Histopathological evaluation:

The organs were isolated and washed with normal saline; small segments of tissues from the normal and experimental rats were preserved in 10% formalin solution and dehydrated in ascending grades of alcohol. The specimens were then embedded in paraffin blocks, sectioned into 4-5 µm thick and stained with hematoxylin and eosin (H&E) staining [14].

Results

General observations of signs of toxicity

The animals were observed for any behavioral changes, food intake and body weight.

Control groups

The control groups were clinically normal during the period of experiment.

Acute toxicity group:

After oral administration of PGB with dose of 5000 mg/kg (LD50), rats showed reduction in both of motor activity and food consumption, while there was increase in sleep duration. The clinical signs of toxicity appeared within 2-6 hours after PGB admiration and included repetitive circling and rearing, followed by convulsions and they expelled froth from mouth during the convulsion episode (**Figure 1**).

Chronic toxicity groups:

In chronic toxicity groups, rats showed reduction in both motor activity and food consumption and increase in sleep duration. **Abnormal** movements were observed in rats' necks in the form of cervical dystonia. In addition, repetitive circling and convulsions occurred during sample collection and on exposure of the rats to minor stimulus (**Figure 1**).

Histopathological changes with PBG toxicity

Examination of sections of the **cerebral cortex** (CC) grey and white matter of both control groups showed normal structures of the layers of CC and its cells. The grey matter was well organized and regularly arranged in six layers from outside inwards: the outer molecular layer was a fibrous layer with few neuroglia and neurons cells, the granular layers consisted of stellate-shaped nerve cells with large round nuclei, the inner and outer pyramidal cell layers were formed of medium and large sized cells, respectively, the innermost layer constitutes the pleomorphic layer and was formed of a variety of nerve cells. Pyramidal cells were seen with large central vesicular nuclei, containing one or more nucleoli and basophilic cytoplasm. Granular cells are seen with large rounded vesicular nuclei while glial cells seen as small dense nuclei. The fibrillary eosinophilic background substance constituted the neuropil, with glial cells and blood vessels within. **(Fig. 2A).**

Examination of sections of the **cerebellum** of both control groups showed normal appearance of cerebellar folia: each folium formed of outer grey cortex and inner white matter. The cerebellar cortex showed three layers from outside inwards; the molecular layer, the Purkinje cell layer and the granular cell layer. The molecular layer was formed of 2 types of neurons: stellate and basket cells. The Purkinje cells layer arranged in one row between the granular and molecular layers, showed large flask shaped cells with vesicular rounded nuclei with apparent nucleoli and basophilic Nissel's granules in their cytoplasm. The granular layer consisted of closely packed numerous, oval granular cells with deeply stained rounded nuclei **(Fig. 3A).**

Effect of acute pregabalin toxicity on cerebral cortex (Fig.2B) and cerebellum (Fig.3B)

In acutely intoxicated group II, examination of sections of CC tissue of both male and female groups showed increase in cellularity and irregular distribution of the astrocytes, with hypertrophy and enlarged vesicular nuclei cells. Dystrophic changes in the form of red neurons were seen with shrinkage of the cell body, pyknosis of the nucleus, disappearance of the nucleolus and intense eosinophilia of the cytoplasm. No pathological changes were seen regarding neuropil, blood vessels and layers of CC.

Examination of sections of cerebellar tissue of both male and female groups showed preserved architecture of cerebellar cortex layers with few shrunken Purkinje cells. The neuropil had few vacuoles of variable sizes and pericellular haloes. The molecular and granular layers have apparently normal architecture.

Effect of chronic pregabalin toxicity on cerebral cortex (Fig.2C) and cerebellum (Fig.3C)

In chronic intoxicated group, the CC showed multifocal histological changes in all layers of the CC of both male and female groups as compared to the control group. Most of pyramidal cells were shrunken and showed deeply stained nuclei with loss of their processes, and pericellular halos. Many vacuoles of variable sizes were found in neuropil and inside most of cells in all layers. Small focal area of inflammatory infiltrate and necrosis was seen. The CC showed dilated congested blood vessel surrounded by inflammatory infiltrate, area of intracerebral hemorrhage and encapsulated necrosis.

Similarly, cerebellar cortex of both male and female groups showed shrinkage of most of Purkinje cells with loss of their processes, and pericellular halos. Many vacuoles of variable sizes were seen in neuropil and inside most of cells in all layers.

Discussion

Pregabalin toxicity has been related to overdose consumption resulting in suicide, recreation and accidental overdose. Moreover, PGB toxicity may occur after its synergistic depressant effect with other CNS depressant drugs and following accumulation of PGB in the setting of renal impairment [15-17].

Accordingly, the present study investigated possible toxic effects of acute and chronic PGB administration on cerebral cortex and cerebellum of adult albino rats.

The brain histopathological changes in the present study showed significant degenerative changes in neurons in both cerebral cortex and cerebellum with hypertrophy and hyperplasia of astrocytes with acute intoxication with PGB. These changes were attributed to the direct effect of PGB on the brain. These results were similar to previous reports of PGB causing encephalopathy and EEG changes [18-20].

In addition, the present study showed degenerative changes in the cerebellar cortex of all chronic PGB intoxicated groups, affecting mainly Purkinje cells, which became shrunken with ill-defined cytoplasmic processes and some of them had vacuolated cytoplasm with absence of Nissel's granules. The molecular and granular layers were also affected, being less in thickness and had multiple vacuolated areas. Such findings agreed with previous reports [21,22], which reported the chronic effect of administration of 600mg/kg PGB to maternal Swiss albino mice and resulted in degenerative effects on developing cerebellum of fetus, in the form of disruption of the architecture of the Purkinje cell layer, mild vacuolization in the granular layer, necrosis and hemorrhage in focal area of white matter.

The cerebellar cortex in chronically treated rats with 500 mg/kg PGB showed degenerative changes of both Purkinje and granular cells. Purkinje cell became shrunken, disarranged and there was cell loss. Purkinje cells play a crucial role in motor coordination and alteration in structure or functions of these cells may cause changes in motor coordination function [23, 24]. There has been some evidence of cerebellar dysfunction in pregabalin intoxication in the form of nystagmus, in-coordination, myoclonus and ataxia [25-28].

The vacuolization appeared in brain tissues of rats in the current study could be attributed to cellular degeneration within the cerebellar and cerebral cortex. In addition, shrunken Purkinje cells were surrounded by large prominent pericellular spaces that could be related to the shrinkage of neurons and disintegration of their cyto-skeletal elements [29]. The dark staining of nuclei of degenerated cells could be due to accumulation of denatured proteins and chromatin condensation in cells undergoing apoptosis [30].

The cerebellar dysfunction in pregabalin intoxication could be attributed to action of PGB on $\alpha 2\text{-}\delta$ subunit of VGCC, which are widely distributed throughout the peripheral and central nervous systems. The highest

level of expression of these channels has been found in the cerebellum and in the hippocampus, and their dysfunction or decreased activity level has been linked to ataxia and cognitive impairment [32].

Declarations

Ethical approval

All procedures performed on animals were done in according to the guidelines and approved by committee of ethics in faculty of medicine, Suez Canal university; Egypt (C; No.4521) and in compliance with the ARRIVE guidelines. The staff who dealt with the animals were well trained. Animals were fed appropriately and protected from pathogens. Pain, distress, and suffering were kept to a minimum grade. Adequate food and water were introduced to entire animals and they were assessed for general health and body weight. Animals were anesthetized before scarification. Scarification the animals was done in appropriate manner.

Availability of data and material

All data generated during this study are included in this published article [and its supplementary information files].

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Competing interests

No conflict of interest.

Authors' contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Figures

Figure 1

Clinical effects of PGB toxicity on rats A; Intoxicated rat has tonic clonic convulsions. B; Cervical dystonia (involuntary neck muscle contractions producing abnormal postures) in chronic intoxicated rats. C; Repetitive circling in rat.

A

B

C

Figure 2

Representative figures heamtoxylin and eosin stained sections of cerebral cortex in control, acute and chronic toxicity groups. (Left sided figure; 20x & right sided figure; 40x) A; Control group Section of control rat cerebral cortex showing normal pia matter (black arrow), grey matter layers; molecular layer (1), outer granular layer (2), outer pyramidal layer (3), inner granular layer (4), inner pyramidal layer(5) and the plemorphic layer (6), homogenous neuropil (black star *) and blood vessels (BV). Inset showing

pyramidal cells (P) and granular cells (G), 100x. B; Acute toxicity group Photomicrographs of sections from the cerebral cortex of adult albino rats with acutely intoxicated with pregabalin LD50 (II) for 24 hours, showing astrocytes with hypercellularity and irregular distribution, degenerated neurons (D), homogenous neuropil with slight vacuolation and pericellular halos (arrowheads). In addition, there are intracellular vacuoles (IV), extracellular vacuoles (EV), interstitial edema (thick black arrow) and dystrophic red neurons (R). C; Chronic toxicity group Photomicrographs of sections from the cerebral cortex of adult albino rats with chronic intoxication with pregabalin LD50 (II) for twelve weeks, showing degenerated neurons (D), homogenous neuropil with vacuolation (*), intracellular vacuoles (IV), extracellular vacuoles (EV), interstitial edema and inflammatory infiltrate (Thin black Arrow), pericellular halos (Arrowhead), necrosis (N) and dilated blood vessels (DBV).

Figure 3

Representative figures heamtoxylin and eosin stained sections of cerebellum in control, acute and chronic toxicity groups. (Left sided figure; 20x & right sided figure; 40x) A; Control group Section of control rat cerebellum showing normal folia (F) separated by narrow sulci (S), uniform white matter (W), well-organized three layers of cerebellar cortex; the molecular layer (MCL), the Purkinje cell layer (PCL) and the granular layer (GCL). The cerebellar cortex showing stellate (Sc), basket cells (B), large pyriform Purkinje cells (P), granular cells (G) and surrounding neuropil (*). B; Acute toxicity group Photomicrographs of

sections from the cerebellum of adult albino rats with acutely intoxicated with pregabalin LD50 (II) for 24 hours, showing preserved architecture of cerebellar cortex layers; the molecular layer (MCL), the Purkinje cell layer (PCL) and the granular layer (GCL) with surrounding neuropil (*). Some Purkinje cells (PN) are distorted and shrunken with vacuoles (V). (Incet, 100x) C; Chronic toxicity group Photomicrographs of sections from the cerebellum of adult albino rats with chronic intoxication with pregabalin LD50 (II) for twelve weeks, showing degenerated Purkinje cells (PN), vacuoles (V) and pericellular halos (thick black arrows), with interstitial edema (thin black arrows).

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