Long Term Administration of Nicotine in Albino Rat Cerebellum - Histopathological Study

S.Lokanadham1, UshaKothandaraman2, R.Chidamabaram3, WMS.Johnson4, K.Prabhu5
1 Research Scholar, Department of Anatomy, Bharath University, Chennai, Tamilnadu, India.
2Professor and Head, Department of Anatomy, ESIC Medical College & PGIMSR, Chennai, Tamilnadu, India.
3R & D Director, Centre for Research, Sri Lakshmi Narayana Institute of Medical Sciences, Puducherry, India.
4Professor, Department of Anatomy, Sri Balaji Medical College, Chennai, Tamilnadu, India.
5Associate Professor, Department of Anatomy, Sri Balaji Medical College, Chennai, Tamilnadu, India.
*Corresponding author’s E-mail: loka.anatomy@yahoo.com

Accepted on: 20-08-2015; Finalized on: 30-09-2015.

ABSTRACT
The actions of nicotine upon cerebellar function are poorly understood. Small doses of nicotine have a stimulating action on the central nervous system whereas large doses depress. 16 adult male albino rats of wistar strain weighing 135-150 grams were used for the present study. The rats were divided in to four groups each consists of 4 animals. The control group was given 0.9% saline as vehicle and the experimental groups (drug treated- I, II, III, IV) were given intraperitoneal injection of Nicotine a dose of 0.5 mg/kg, 0.75 mg/kg and 1mg/kg body weight dissolved 1:1 ratio in normal saline respectively for a period of 28 days. Cortical layers were intact in the entire drug treated group. White matter with few vacuoles was observed in 0.5 mg/kg nicotine induced group. Disruption of Purkinje cell layer and granular cells was observed in 0.75 mg/kg induced group. Dose of 1mg/kg nicotine induced group animals shown complete disruption of Purkinje cell layer, cellular disruption in granular cell layer. Extended core of white matter into the folia of the cerebellum cellular microcystic change with interstitial oedema was observed in the same group. The loss of white core, Purkinje cell layer, and granular layer cellular disruption give better knowledge in understanding long term nicotine exposure. Our study reveals the histopathological changes of nicotine on long term administration in adolescent rat cerebellum with their clinical significance.

Keywords: cerebellum, disruption, nicotine, white core.

INTRODUCTION
Recent studies on neurophysicochemical aspects also suggested that the cerebellum also could participate in excitotoxicity. Dysfunction of the cerebellum causes problems such as loss of balance and motor coordination, ataxia, dystonia through several genetic mutations and alterations in the biochemical pathways of neuroprotection; however, a large number of patients remain undiagnosed. Many researchers have studied the effect of nicotine on cerebellum and found significant deficits. Nicotinic receptor binding sites and subunits are expressed in rat cerebellum from embryonic stage through adult hood, with the highest expression levels seen during the development of the cerebellar cortex. Nicotine acts as neurodegenerative agent on higher doses, recent studies stating that nicotine can use as protective agent due to its positive effects in the same area but still its action of mechanism unknown.

MATERIALS AND METHODS
A total of 16 adult male albino rats of wistar strain weighing 135-150 grams were used for the present study. The animals were maintained under controlled conditions and in room temperature (23±2° C), humidity (50±5%) and a 12 h light and dark cycle. The animals were housed in sanitized polypropylene cages. The animals were fed with standard rat pellet diet commercially available manufactured by Kamathenu Pvt Ltd., Bangalore and clean drinking water ad libitum. The rats were divided in to four groups each consists of 4 animals. The control group was given 0.9% saline as vehicle and the experimental groups (drug treated-I,II,III,IV) were given an intra peritoneal injection dose of 0.5 mg/kg, 0.75 mg /kg, 1mg/kg body weight Nicotine dissolved in normal saline (0.9% Nacl) respectively for a period of 28 days. All the animals were sacrificed after a 12 hrs of fasting from the last day of experimental study and preserved in the 10% formalin for 48 hours. The specimens were processed for routine paraffin technique to study the histopathological changes in cerebellum on administration of nicotine.

All the animals were maintained as per the national guidelines and protocols approved by Institutional animal ethical committee of Sugen Life sciences (SLS/COM/IAEC/02/2013-2014).

RESULTS
Molecular layer with basket and stellate cells, single Purkinjee cell layer lies between the molecular and granular cell layer was observed in control group. We have observed Column of nerve fibers in the white matter of cerebellum. Granular cell layer with granule cells (pink stained) glomeruli (light stained) were also observed in control group (Figure - 1).
In 0.5 mg /kg nicotine treated Group (II); there was vacuoles in the white matter and all the layers of cortex were observed (Figure - 2).

Disruption of purkinje cells layer and granular cells was observed in 0.75 mg/kg nicotine treated group (III). We were also observed few purkinje cells in the same group (Figure-3).

Vacuoles in the white matter and disruption of cells in the granular layer under high magnification were observed (Figure-4).

In 1mg /kg nicotine treated group (IV), there was complete disruption of Purkinje cell layer along with a loss of Purkinje cell layer (very few cells only present) and also cellular disruptions in granular cells were observed (Figure-5).

Under high magnification we have observed the extended core of white matter into the folia of the cerebellum, cellular microcystic change with interstitial oedema of
white matter and significant loss of Purkinje cells with disruption of cellular layers were observed (Figure - 6).

**CONCLUSION**

The loss of white core, Purkinje cell layer, and granular layer cellular disruption give better knowledge in understanding long term nicotine exposure in animal models.

**Acknowledgement:** Authors acknowledge the great help received from DR. ChandraMouli during this work.

The authors are also grateful to authors, editors and publishers of all those articles, journals and books from where the literature for this article has been reviewed and discussed.

**REFERENCES**


**DISCUSSION**

The interaction of nicotine and the Alpha subunits of nicotine Ac receptor in white core may subsequently trigger apoptotic process that leads to loss of white matter. In our study there was a loss of white core of cerebellum on long term administration of nicotine. Adolescent nicotine exposure has deleterious effects on cell development particularly in Purkinje cell layer in cerebellum of albino rats characterized by reduced number of Purkinje cell number. Exposure to nicotine during the brain growth spurt period in rats has been associated with significant reduction of the Purkinje cell numbers in the cerebellar vermis. Belluzzi and co-workers reported that the brain in early adolescent, late adolescent and adult rats is differentially sensitive to the nicotine administration. The adolescent or adult brain could be more sensitive to acute and also to chronic administration of nicotine. Neuro-pathologically, reduction in Purkinje cell numbers in cerebellar vermis and hemispheres have been reported in nearly all mentally retarded autistic cases. The loss of Purkinje cells combined with loss of granule cells and white matter of cerebellum in two mentally retarded cases suggested that cerebellar hyperplasia. On higher doses of nicotine for long term administration we have found Purkinje cell layer and granular cells disruption and loss which were in agreement with previous literatures. One-dose application had no significant effect, but the chronic treatment was accompanied with a reduction in locomotor activity in adult rats. Administration of nicotine specifically inhibits cerebellar Purkinje cells which could result in the reduction of Purkinje cells, and consequently, the middle Purkinje cell layer of the cerebellum.

The results and observations of our study were in agreement with literature.


Source of Support: Nil, Conflict of Interest: None.