

Effect of exposure and withdrawal of pyrethroid and herbal based mosquito vaporizers on histology of spinal cord of male wistar albino rats

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Abstract

Introduction: Pyrethroids, organochlorines, organophosphates, and carbamates constitute important active ingredients of available mosquito repellents in the Indian market, in which pyrethroids are found very effective and safe. The natural compounds produced by plants can interfere major metabolic pathways, thus causing death of the insects. The aim of present study was to know the effect of exposure and withdrawal of transfluthrin and herbal based mosquito vaporizers on spinal cord of male albino rats.

Materials and Methods: We have taken spinal cord of 26 well preserved male albino wistar rats and divided in to three groups.

Control Group: Six rats, notexposed to any mosquito repellents.

Experimental Group 1: 12 rats: divided in subgroup 1A, the pyrethroid Exposure group; and subgroup1B, the Pyrethroid withdrawal group.

Experimental Group 2: Eight rats: divided in subgroup 2A, the herbal Exposure group and 2B, the herbal withdrawal group.

Results: Spinal cord of pyrethroid vaporizer exposed group showed gross destruction in their histology than the herbal based mosquito vaporizer exposure group. Also, some of the normal histological features were recovered after withdrawal of chemicals and reversal was better in herbal based mosquito vaporizer.

Conclusion: Histological findings of present study suggests that pyrethroid (Transfluthrin) based mosquito repellents are harmful to the spinal cord. Herbal based mosquito repellent also showed damage though the severity was mild as compared to pyrethroid. Reversal changes were much better in herbal withdrawal group than the pyrethroid withdrawal group.

Keywords: Pyrethroid, Transfluthrin, Herbal, Mosquito repellent, Spinal cord, Withdrawal.

Introduction

In the race of better quality of life, knowingly or unknowingly we are exposing ourselves to many harmful chemicals in our daily routine, and protecting ourselves from mosquito bites is one such example. Mosquito borne diseases are burden on community health. There are many chemical mosquito repellents in the form of Mosquito Coils, Liquid Vaporizers are available in the market as personal protective measures from mosquito bite. Pyrethroids, organochlorines, organophosphates, and carbamates constitute important active ingredients of available mosquito repellents in the Indian market, in which pyrethroids are found very effective and safe.¹

There are two categories of pyrethroids. Type I pyrethroids (like allethrin, resmethrin, and permethrin) does not contain an alfaciano group, while type II pyrethroid (like deltamethrin, cypermethrin, and fenvalerate) contains an alfaciano group. Type I usually causes T-syndrome characterized by tremors, whereas type II pyrethroids usually present its adverse effect in the form of CS-syndrome.² The type I pyrethroid transfluthrin (TFL) is synthesized by tetrafluorobenzyl alcohol and trans-2,2-dimethyl-3-(2,2-dichlorovinyl) cyclopropanecarboxylate.³ Since plant produces a large number of secondary compounds like alkaloids, terpenoids, phenolics, flavonoids, chromenes which possess physiological activities against insects so this may provide a new source of natural insecticides. These natural compounds produced by plants can interfere major metabolic pathways, thus causing death of the insects.⁴

There is a lack of studies on effect of herbal based mosquito vaporizers on histology of organs so the aim of present study was to know the effect of exposure and withdrawal of trans fluthrin and herbal based mosquito vaporizers on spinal cord of male albino rats.

Material and Methods

The present study was conducted in the Department of Anatomy, King George's Medical University, UP, Lucknow during August 2016 to September 2017. Ethical clearance was obtained from Institutional Animal Ethics committee (IAEC), King George's Medical University. The proposed study was carried out on various formalin fixed viscera from 26 male wistar albino rats.

Inclusion Criteria

Spinal cord from well preserved male wistar albino rats exposed to pyrethroid and herbal based mosquito vaporizers were taken for study.

Exclusion Criteria

Spinal cord of male albino wistar rats not well preserved with distorted morphology were excluded from the present study.

Study Design

Well-preserved spinal cord of 26 Male albino rats with, were divided into 3 groups:

Control Group

Spinal cord of 6 male albino rats, those were not exposed to mosquito vaporizer fumes.

Experimental Group I

Spinal cord of 12 male albino rats (divided in subgroup I A and subgroup I B; each subgroup contains spinal cord of 6 male albino rats).

Subgroup I A

Contains spinal cord of male albino rats exposed to chemical-based mosquito vaporizer (Transfluthrin 0.88% w/w) for 8 hours /day, 6 days in a week for 12 weeks.

Subgroup I B

Contains spinal cord of male albino rats those were given exposure of chemical-based mosquito vaporizer (Transfluthrin 0.88% w/w) for 8 hours/day, 6 days in a week for 12 weeks and further kept for 4 weeks to see the reversal changes.

Experimental Group II

Spinal cord of 8 male albino rats (divided in subgroup II A and subgroup II B; each subgroup contains organs of 4 male albino rats).

Subgroup II A

Spinal cord of male albino rats exposed to herbal based mosquito vaporizer for 8 hours/day, 6 days in a week for 12 weeks.

Subgroup II B

Contains spinal cord of male albino rats those were exposed to herbal based mosquito vaporizer for 8 hours/day, 6 days in a week for 12 weeks and further kept for 4 weeks to see the reversal changes.

Products detail

Chemical Based Mosquito Vaporizer

Good Knight Refill (Vaporizer containing 0.88% transfluthrin, Reg. No. CIR-48,420/2004; manufactured by Godrej Consumer Products Ltd.) and herbal based Mosquito Vaporizer Mosrelief Refill Vaporizer (Vaporizer containing Cymbopogon citrius*, Bhutika, oil:6%, Azadirachtaindica**, Nimbaoil:6%, Eucalyptus Globulus, Tailaparna, oil:6%, Excipients: OS, *API, **AFI; Mff. Licence No. AUS920; manufactured by Strategi, Plot No.:50, Part 1, 4th Phase, KIADB industrial area, Mular-563130, Kolardist., Karnataka, India.) were used in our study.

Slides were prepared for histopathological observation, by routine Histological procedure i.e. fixation, clearing, paraffin embedding and sections of 5 µm thick were cut by rotary microtome and stained with Hematoxylin and Eosin stain.

Light microscopic examination of sections of spinal cord was done to evaluate and compare the histopathological changes. Observation of slides was done

under 10X, 40X, by Kyowa Trinocular Research Microscope. Microphotograph of different tissues were taken for documentation by digital camera (DSC-WX80/RC E32, manufactured in China by Sony corporation, 1-7-1 konan, Minto-KU, Tokyo 108-0075, Japan, month of import April 2014).

Observations

Histological findings of spinal cord

Control group

At 10X magnification the histological section of spinal cord at cervical region showed intact ciliated cuboidal ependymal linings of central canal. Both anterior and posterior grey horns were evident with their multipolar neurons. A large central nucleus, a prominent nucleolus, and several radiating cell processes were seen in the multipolar motor neurons of the spinal cord at higher magnification (40X). A single, thin axon was seen arising from a cone-shaped, clear area of the neuron; axon hillock. Smaller cells with basophilic nuclei (neuroglia) were seen between spaces between the neurons (Fig. 1).

Group IA

Haematoxy line and Eosine stained section of spinal cord of pyrethroid exposed group showed discontinuity in ependymal linings of central canal (10X). There was extensive neuronal degeneration and perineuronal vacuolations of grey matter. Few neuroglial cells were seen (Fig.1.1). There was capillary dilatation and haemorrhage at some places in grey matter. The vacuolations of white matter was also seen at some places. The extensive oedema and loss of inter neuronal contacts leading to wide separation of grey and white matter was clearly visible (Fig.1.1).

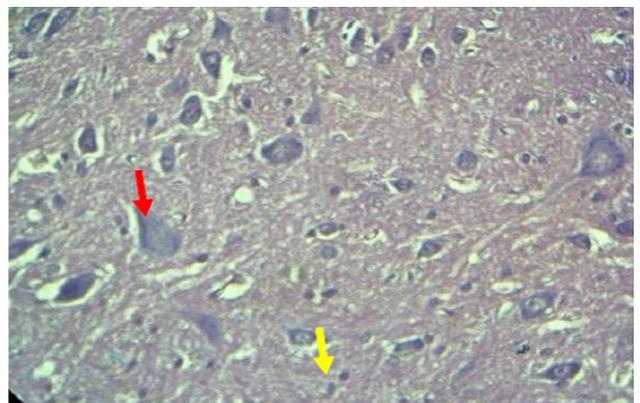


Fig. 1: (40X) Photograph of spinal cord of control group showing multipolar motor neuron (red arrow) and neuroglial cells (yellow arrow).

Group IB

The histological sections of group IB showed disruption of ependymal lining of central canal. Degeneration of neurons and perineuronal vacuolation of the grey matter was seen. There was loss of neuroglial cells (Fig.1.2) but it was less marked as compared to group 1A. Capillary dilatations

and haemorrhage at some places in grey matter was observed. The vacuolations of white matter were less marked as compared to group IA. There was extensive oedema and loss of inter neuronal contacts resulted into separation of grey and white matter (Fig. 1.2).

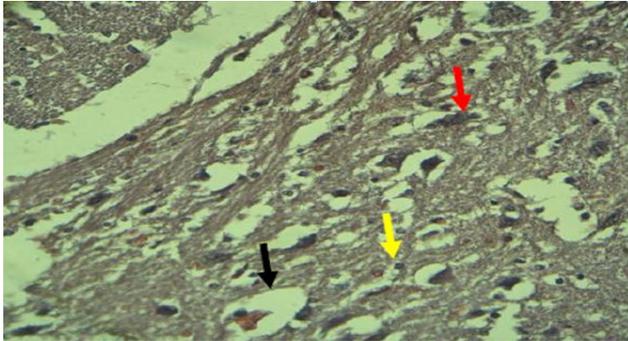


Fig. 1.1 (40X) Group IA showing neuronal degeneration (red arrow), perineuronal vacuolation (black arrow) and few neuroglial cells (yellow arrow).

Group IIA

Disruption of central canal lining and oedema was well evident in grey matter of spinal cord of Group IIA. Degeneration of neurons and loss of neuroglia of grey matter of spinal cord was also observed (Fig.1.3). There was sparsely placed axons and loss of neuroglia from the white matter (Fig.1.3).

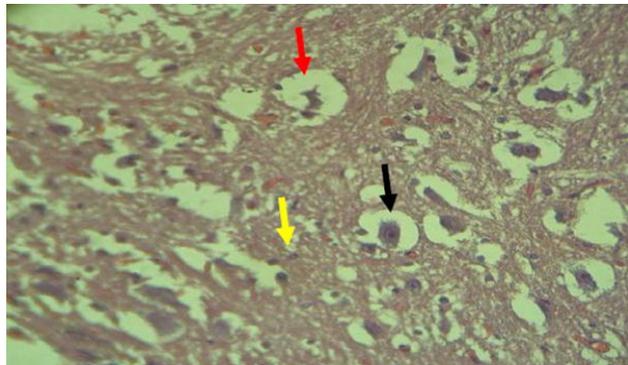


Fig. 1.2: (40X) Group IB showing neuronal degeneration (red arrow), perineuronal vacuolation (black arrow) and more neuroglial cells (yellow arrow).

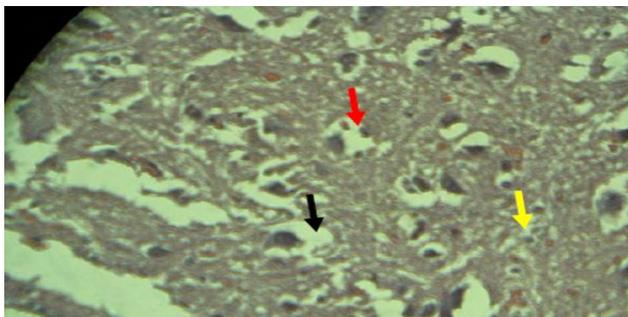


Fig. 1.3: (40X) Group IIA showing neuronal degeneration (red arrow), perineuronal vacuolation (black arrow) and few neuroglial cells (yellow arrow).

Group IIB

Intact central canal was clearly seen. Less oedematous grey matter. Most of the multipolar neurons showed axon hillock along with nissel's granules. There was densely packed nerve fibres and neuroglia almost similar to that of control group, were seen in the white matter (Fig.1.4).

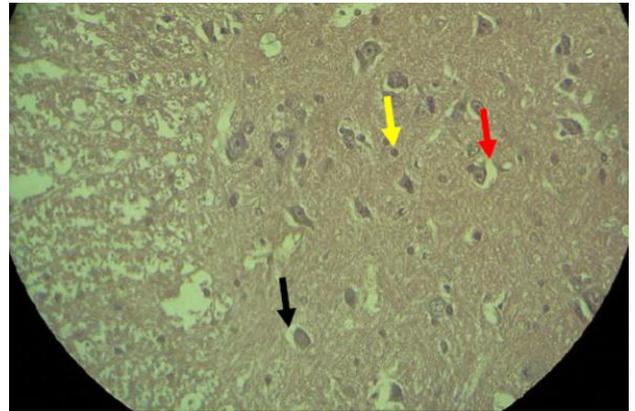


Fig. 1.4: (40X) Group IIB showing less marked neuronal degeneration (red arrow), perineuronal vacuolation (yellow arrow) and abundant neuroglial cells (black arrow).

Discussion

There are many insecticidal agents which cause variable changes in the central nervous system on repeated exposure. Metabolic changes due to hypoxia and hypoglycaemia can lead to cellular injury into the exposed organs.

H&E stained section of spinal cord of pyrethroid exposure group of present study showed extensive neuronal degeneration, perineuronal vacuolation in grey matter and vacuolation of the white matter along with capillary dilatation. There was wide separation of grey and white matter which results due to oedema and loss of interneuronal contacts. Our findings are in accordance to a study of where sub lethal toxicity of cypermethrin in Sprague Dawley rats showed spongiosis and perineuronal degeneration of spinal cord and gliosis, degeneration of neurons and perineuronal vacuolation was also evident in the in the cerebrum of exposure group of rats.⁵ In another study, the rats, those were given orally the sub-lethal doses of pyrethroids showed various histopathological changes in peripheral nerves and spinal cord.⁶ Necrosis and degeneration of neurons of cerebrum was also reported among cypermethrin-intoxicated rats.⁷ Rats those were exposed to cypermethrin showed cellular injury in the form of pyknosis and loss of some cells in the cerebellum.^{8,9} Degeneration and congestive changes were reported in a study which was similar to the present observations.^{10,11,12} Concordant to our findings, haemorrhages were also observed in the brain tissue of rats exposed to cypermethrin.¹²

Conclusion

After evaluation of histological changes, it was seen that Inhalation of pyrethroid and herbal mosquito vaporizers both causes neuronal degeneration and spongiosis in spinal cord. Discontinuation of herbal vaporizer exposure for one month exhibits significant reversal of injury in comparison to withdrawal of pyrethroid based mosquito vaporizer. It can be concluded that toxic effects of these chemicals, whether synthetic or natural, are seen in rats. Therefore, we should limit the use of these vaporizers and further studies should be done to identify other compounds and its related toxicity must be explored as a safer alternative for protection against these vectors.

Conflicts of Interest: None.

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