Effect of cadmium chloride on nucleus preopticus in *Heteropneustes fossilis* and its recovery by herbal compound, Ashawagandha

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ABSTRACT

The effect of cadmium chloride on nucleus preopticus in *Heteropneustes fossilis* was analysed histologically. Fish when treated with cadmium chloride of 0.5 ppm for 7, 14, and 21 days exhibited degeneration, hypertrophy, reduced neurosecretory material and vacuolization in NPO. Their size was significantly (p<0.001) increased after cadmium chloride exposure. In ashawagandha recovery group, these nuclei displayed a gradual reorganization in structural detail and size. Recovery by a herbal compound ashawagandha exhibited reduced hypertrophy and vacuolization in NPO.

KEY WORDS: ASHWAGANDHA, CADMIUM CHLORIDE, HETEROPNEUSTUS FOSSILIS, NUCLEUS PREOPTICUS

INTRODUCTION

Heavy metals occur naturally in the environment and are found in varying levels in ground and surface water. Heavy metals are reported as pollutants which caused the metabolic, physiological and structural alterations in fish (Jiraungkoorakul et al., 2007; 2008; 2009). Among heavy metal cadmium has been shown to be responsible for a number of reproductive abnormalities in fish (Sharma et al.,2013 ). The pituitary gland is one of the most important endocrine organs of fish. The histology of the pituitary of teleost fish has been described by a number of authors (Balci et al., 2006; Ozen and Timur, 1993; Hibiya, 1982). The heavy metals (lead, cadmium, mercury etc.) are known to interfere with the endocrine system of model organisms such as mammals, fish etc. and lead to a disturbance in hormonal metabolism, hormone-regulated cellular of physiological processes(Comborn and Clement,1992 ; Kavlock et al.,1996). However, metal deterioration of hypothalamic
nuclei in fish are not well illustrated. Since only in *Heteropneustus fossilis* (Shukla and Pandey, 1984; Pandit and Bhattacharya, 2013)

**MATERIAL AND METHODS**

*Heteropneustus fossilis* measuring about length 12±5 cm and weight 25±5 gm were used in the present study. Cadmium was used for present study in the form of cadmium chloride (CdCl₂). The dose of cadmium chloride was decided after determination of LC 50 value. It was found to be 0.5 ppm. The herbal compound Ashawagandha (*Withania somnifera*) is used as recovery agent of damaged tissue. Fishes were acclimatized to laboratory condition for 7 days before the commencement of the experiment and were treated with 0.01 KMnO₄ solution to remove dermal infection. Fishes were fed with chopped prawn twice a day. The 72 hrs LC50 value of cadmium chloride was found to be 0.5 ppm in *H. fossilis*.

The fishes were divided in three groups having 36 fishes in each one.

<table>
<thead>
<tr>
<th>S. No.</th>
<th>Group</th>
<th>Treatment</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control</td>
<td>Water without CdCl₂ + plain food</td>
</tr>
<tr>
<td>2</td>
<td>Treated</td>
<td>Exposed to 0.5 ppm CdCl₂ + plain food</td>
</tr>
<tr>
<td>3</td>
<td>Recovery by Ashawagandha</td>
<td>Exposed to 0.5 ppm CdCl₂ + Ashawagandha with food</td>
</tr>
</tbody>
</table>

Fishes of all experiment groups (control, treated and recovery) were sacrificed each after 7, 14, and 21 days. Pituitary with brain were fixed in aqueous Bouin’s solution for 24 hours. The material was washed with water, dehydrated and cleared through graded alcohol and xylene respectively after filtration in paraffin blocks were prepared and section of 5 μ thickness were cut and stretched on albuminized slides. The slides were stained with Chrome Alum Haematoxyline Phloxine (CAHP) (Gomori, 1941) stains and mount in DPX for histological observation. All the data and results for final observation were processed in the form of microphotographs and table. The diameter of NPO were recorded and difference if any were compared by statistical analysis using student ‘t’ test (Bancroft, 1966).

**RESULTS AND DISCUSSION**

The hypothalamo-hypophysial neurosecretory system is a unique endocrine apparatus consisting of the cells of nucleus preopticus (NPO) in neurohypophysis. The nerve fibers of these hypothalamic nuclei terminate in the neurohypophysis. Nucleus preopticus is well developed in *H. fossilis*. These cells were spherical in shape with evenly distributed cytoplasm and their nuclear material. The nucleus usually contains a single nucleolus but sometime more than two nuclei were also observed. The perikarya of NPO cells were laden with neurosecretory material. The neurosecretory cells of NPO were positive to AF and CAHP. Cellular differentiation was more marked after using CAHP technique. The hypothalamus of control fish exhibited the NPO neurons presented in their active secreting stage. The hypothalamic nuclei (NPO) exhibited strong affinity to CAHP and AF stain. (Fig. 1, 3 and 5)

**TREATED GROUPS**

The hypothalamus of cadmium chloride treated fish after 7 days duration exhibited the NPO neurons were present in necrotic condition. The NPO neurons exhibited thick cell boundaries and clumping of cytoplasm. The cell as well as nuclei appeared turgid. The neurosecretory material in the perikarya of the cells was coarse. The cadmium chloride treatment induced deformities in these neuronal cells (Fig. 2).

In fish exposed to cadmium chloride for 14 days duration, the hypothalamic nuclei (NPO and NLT) exhibited cytoplasmic and nuclear abnormalities and depletion of Neurosecretory material. The NPO cells showed degeneration in their cytoplasmic contents. Their cell boundaries were disappeared (Fig. 4). After the treatment of cadmium chloride of 21 days duration these neurons presented exhausted condition due to cadmium stress. Some of the NPO cells showed hypertrophy. Degenerated cytoplasm and disappeared cell boundaries were observed in these cells. They had vacuolated cytoplasm (Fig. 6). Similar results were found by (Shukla and Pandey, 1984) in hypothalamic nuclei of *Sarotherodon mossambicus* exposed to DDT. While Katti and Sathaynesan (1986) reported degeneration of the NPO neurons in *Clarias batrachus* exposed to lead nitrate. Our findings corroborates with study of Ram and Joy (1988). The observed that the neurons containing less quantity of neurosecretory material and exhibited various degrees of degeneration and nuclear necrosis in *Channa punctatus* exposed to HgCl₂ and methyl mercury chloride of the dose.

Fig-1. Control group (7 days): Showing NPO cells stained brightly and neurosecretory material on the periphery of NPO cells visible.

Fig-2. Treated group (7 days): Showing fusion of NPO cells and atrophic condition, among neurons visible.

Fig-3. Control group (14 days): Showing NPO cell bodies with evenly distributed cytoplasm and clearly visible neurosecretory material.
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PLATE 1. Microphotographs of sections through hypothalamus showing NPO of Heteropneustes fossilis in control and treated group (7, 14 21 days, CAHP X1000)

Fig- 4. Treated group (14 days) : Showing deformed NPO cell bodies and their hypertrophied nature.

Fig-5. Control group (21 days): Showing NPO cells with their normal structural configuration.

Fig-6. Treated group (21days): Showing hypertrophied nature and vacuolization of cytoplasm in cell.

Recovery group: After 21 days treatment of cadmium chloride the fishes were administered with Ashawagandha.

In 7 days: After 7 days duration in Ashawagandha group the NPO exhibited reformed condition. Vacuolization was still present in their cytoplasm. Some of them were appeared with regenerated cytoplasm and their

Table 1. Diameter of Nucleus preopticus of Heteropneustes fossilis in control and experimental group.

<table>
<thead>
<tr>
<th>S. No</th>
<th>Experimental group</th>
<th>Days of exposure</th>
<th>7 days</th>
<th>14 days</th>
<th>21 days</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Control</td>
<td></td>
<td>10.4±0.06</td>
<td>10.9±0.05</td>
<td>11.2±0.03</td>
</tr>
<tr>
<td>2</td>
<td>Treated</td>
<td></td>
<td>13.6±0.07**</td>
<td>15.2±0.06***</td>
<td>17.9±0.03***</td>
</tr>
<tr>
<td>3</td>
<td>Recovery Ashawagandha</td>
<td></td>
<td>11.0±0.02**</td>
<td>12.2±0.04**</td>
<td>12.3±0.01**</td>
</tr>
</tbody>
</table>

All values are expressed in Mean ± SEM, Total no. of samples for each observation: 10, Significant level (**p < 0.05, ***p < 0.001).
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Fig-7. Ashawagandha Recovery Group (7 days): Showing accumulation of neurosecretory material in NPO cells and vacuolization in their cytoplasm still persist.

Fig-8. Ashawagndha Recovery Group (14 days): Showing still exhibited hypertrophied condition in a few NPO cells, most of them in normal appearance.

Fig-9. Ashawagndha Recovery Group (21 days): Showing regenerated cytoplasm and nuclear contents of cell body of NPO.

ABBREVIATIONS

C-Cytoplasm, NPO-Nucleus preopticus
NSM-Neurosecretory material DN-Degenerated neurons
HN-Hypertrophied neurons, VN-Vacuolized neurons
RH-Reduced hypertrophy, RN-Regenerated neurons
RV-Reduced vacuolization

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nuclear contents (Fig. 7) In 14 days In Ashawagndha recovery group the NPO cells showed reduced hypertrophy and increased population with some extent of neurosecretory material (Fig. 8) In 21 days After 21 days duration the fishes of Ashawagndha recovery group showed signs of reconstitution in the neurons. In recovery group cellular hypertrophy still persisted in a few of NPO neurons. Most of them were in normal appearance. They had regenerated cytoplasm and nuclear contents (Fig. 9)

These observations were in good agreement with the study of Shukla and Pandey (1984). They postulated decreased size of hypothalamic nuclei in S. mos-sambicus exposed to DDT and BHC when kept in plain water. Pandit and Bhattacharya (2013) observed after Hgcl2 exposure NPO size were increased in the spirulina and chorella were effective to recover the histological changes in the NPO of H. fossilis.


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