

## **Histopathological Changes Induced by Specific Pesticides on Some Tissues of the Fresh Water Snail, *Bellamya dissimilis* Müller**

P. R. Jonnalagadda,<sup>1</sup>B. P. Rao M.<sup>2</sup>

<sup>1</sup>Food and Drug Toxicology Research Centre, National Institute of Nutrition, Jamai-Osmania, Hyderabad 500 007, India

<sup>2</sup>Department of Zoology, Nagarjuna University, Guntur 522 510, India

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Pathological and biochemical disturbances in aquatic organisms due to pesticide toxicity is well documented (Muley and Mane 1990). These disturbances can be structural and functional at the cellular and subcellular levels in organisms (Rodriguez et al., 1994). The action of the toxins on the liver and kidney might alter the enzyme activity (Dikshit et al. 1975). Histopathological changes were mostly confined to organs directly involved in their metabolism and detoxification (Rashatwar and Ilyas 1984).

The present study has been carried out to study the effect of the pesticides endosulfan, methylparathion, quinalphos and nuvan (DDVP) on the fresh water snail, *Bellamya dissimilis* Müller. This type of snail is consumed by the population in and around Kolleru lake, which is one of the largest natural freshwater lakes in India. It is also used as a duck bait.

### **MATERIALS AND METHODS**

A total of 50 adult fresh water snails (*B. dissimilis*) Müller were collected from Kolleru lake and divided into five groups of 10 each. The first group consisted of control snails while the other four groups were exposed to 96-hr LC<sub>50</sub> concentrations of different pesticides (Table 1). The 96-hr LC<sub>50</sub> values for the four pesticides were determined by employing the unweighted regression method of probit-analysis (Roberts and Boyce 1972). The fiducial limits of the LC<sub>50</sub> values were derived from the probit transformation at 95% confidence level (Snedcor & Cochran 1967). The digestive gland, mantle and foot of the control group and those of experimental groups that survived (50%) at the end of 96-hr exposure were removed. The tissues were fixed in alcoholic Bouin's fluid and processed (Weesner 1968). The sections (8 µm) were prepared from paraffin embedded tissues and stained with routine haemotoxylin and eosin preparations.

Correspondence to: P. R. Jonnalagadda

**Table 1. 96- hr LC<sub>50</sub> concentrations of different pesticides**

| Group | Pesticides  | Indian supplier                                      | 96-hr LC <sub>50</sub> conc. g/l (10 <sup>-3</sup> ) |
|-------|---|--|--|
| I     | Control unexposed   |  |  |
| II    | Endosulfan 35% EC<br>6,7,8,9,10 hexachloro,<br>1,5,5a,6,6,9,9a-<br>hexahydro<br>6,9 methano-2,4,3-<br>benzo(e)dioxothiepin-3<br>oxide | A.P. Agro<br>Industries<br>Develop<br>Corp. Kurnool. | 1.8<br>(1.65 to 2.11)                                |
| III   | Metacid or Methyl-<br>parathion 50% EC<br>O,O,Dimethyl<br>O-P nitrophenyl<br>Phosphorothioate   | Bayer India<br>Ltd., Bombay.                         | 4.7<br>(3.59 to 6.36)                                |
| IV    | Quinalphos 25% EC<br>Dimethyl O-quinoxalin<br>2yl phosphate   | Premier<br>Pesticides Pvt.<br>Ltd. Cochin.           | 1.9<br>(1.52 to 2.38)                                |
| V     | Nuvan 36 % EC<br>2-dichloroethenyl<br>O-O-dimethyl Phosphoric<br>acid ester   | Hindustan<br>Ciba Geigy Ltd.<br>Bombay.              | 20.89<br>(19.96 to 23.43)                            |

Numbers in parenthesis indicate 95% confidence limits.

## RESULTS AND DISCUSSION

The histopathological studies have shown that the four pesticides have affected the digestive gland, mantle and foot of B. dissimilis Müller when compared with their respective controls (Figs. 1a, 2a and 2e). The lobules, interlobular connective tissue, muscle fibers surrounding the lobules were also affected but the degree of damage varied with different pesticides. The lumen of the digestive gland tubule was shrunken, hypertrophied with conspicuous vacuolations in endosulfan (Fig. 1b) exposed snails. In methyl parathion (Fig. 1c) exposed snails, interlobular tissue was broken and large vacuoles appeared in the hepatic cells and in the connective tissues. Vacuoles increased in inter lobular connective tissue which lost its original shape and the secretory cells became irregular in quinalphos (Fig.1d) exposed snails. In nuvan exposed snails (Fig. 1e) damage to the digestive gland tubules was extensive resulting in deformity of the hepatic cells, or complete destruction of the tubules, hypertrophy of muscles and vacuolation of hepatic tissue. Hepatic necrosis was conspicuous.

The mantle of the exposed snails showed disorganisation of the tissue especially in the lamellae. Interlamellar space was filled with hyperplastic epithelium and lamellar shape and structure were lost due to desquamation, and this was more conspicuous in methyl parathion (Fig.2c) and Quinalphos when compared with endosulfan exposure (Fig.2b). Cytoplasmic vacuolation and the nucleus were either karyorrhectic or pyknotic. Hemorrhage and infiltration were also observed. Hyperplasia was so

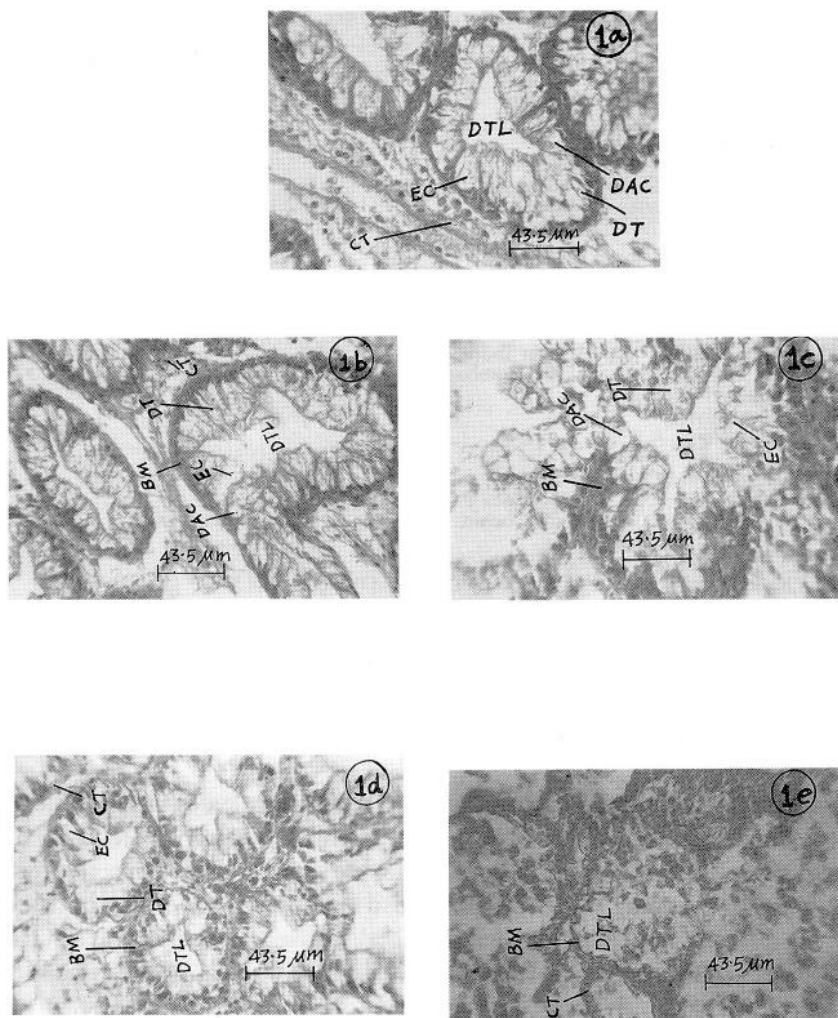


Figure 1. Transverse section of digestive gland (Figs 1a to 1e) of (a) control (unexposed) and after 96-hr exposure to 96-hr  $LC_{50}$  concentrations of (b) endosulfan, (c) methyl parathion, (d) quinalphos and (e) nuvan (X250)

DTL - Digestive Gland Lumen; DAC - Digestive Absorptive Cell;  
DT - Digestive Tubule; BM - Basement Membrane; CT - Connective  
Tissue; EC - Excretory cell.

acute in methyl parathion (Fig.2c) and nuvan (Fig.2d) exposed snails that the lamellae were transferred into globular or balloon-like structures.

The main targets of toxic action were epithelium, mucous glands and muscle fibers of the foot. The epidermis was broken and the mucocytes were concentrated. The central connective tissue and columnar muscle fibers were disrupted in endosulfan exposed snails (Fig.2f). In methyl parathion (Fig.2g) and quinalphos exposed snails, desquamation of the hyperplastic epithelium was noticed. The subepidermal cells were concentrated and muscular layers were damaged. Damage to the foot tissue in nuvan (Fig. 2h) exposed snails was extensive. In addition to the disruption of the muscle fibers, spaces and gaps appeared between fibers and the vesicular cells, and the mucous cells were concentrated and lost their shape.

The present study has shown several degeneration changes in the histological structure of the digestive gland, mantle, and foot of B. dissimilis exposed to 96-hr LC<sub>50</sub> concentrations of the four pesticides. Mane et al. (1979) have noted similar changes while investigating the histopathological changes in the hepatopancreas of the bivalve, Indonaiia cauruleus exposed to DDT, malathion, endrin and thiometon.

There seems to be scanty information on the histopathological effects of pesticides on the foot of molluscs. Swelling of epithelium, reduction in the number and size of mucous glands in parenchyma and disruption of muscle fibers are noted in B. dissimilis.

Pesticides in general induce histopathological changes in the tissues of fish. However, in other short term studies no marked histological changes were observed in the tissues of experimental animals (rabbits and rats) but only increased or decreased weight was reported (Gupta and Gupta 1977; Wayman et al. 1978).

Of the three other parameters (toxicological, metabolic and biochemical) studied, it was clear that endosulfan was more toxic to B. dissimilis than the three other pesticides mentioned (Padmaja et al. 1988, Padmaja and Rao 1991; Padmaja and Rao 1994). In the present study it was interesting to note that histopathological changes induced by nuvan were more intense when compared to endosulfan. Indeed, while finding the effect of endosulfan on Labeo rohita, it was observed that the death of the fish exposed to acute doses is the result of direct toxic action of the pesticide rather than due to destruction of cells and interference with cellular metabolism (Rao et al. 1980). Cellular destruction may perhaps accelerate death by way of affecting the biochemical pathways and energy reserves (Padmaja and Rao 1994).

Most of the pesticides, particularly the organochlorines have been reported to be hazardous to non-target species (Matthiessen et al. 1982). It is interesting to note that in the present study, the organophosphorous pesticide, such as nuvan was more toxic than organochlorine, endosulfan, when various parameters were considered

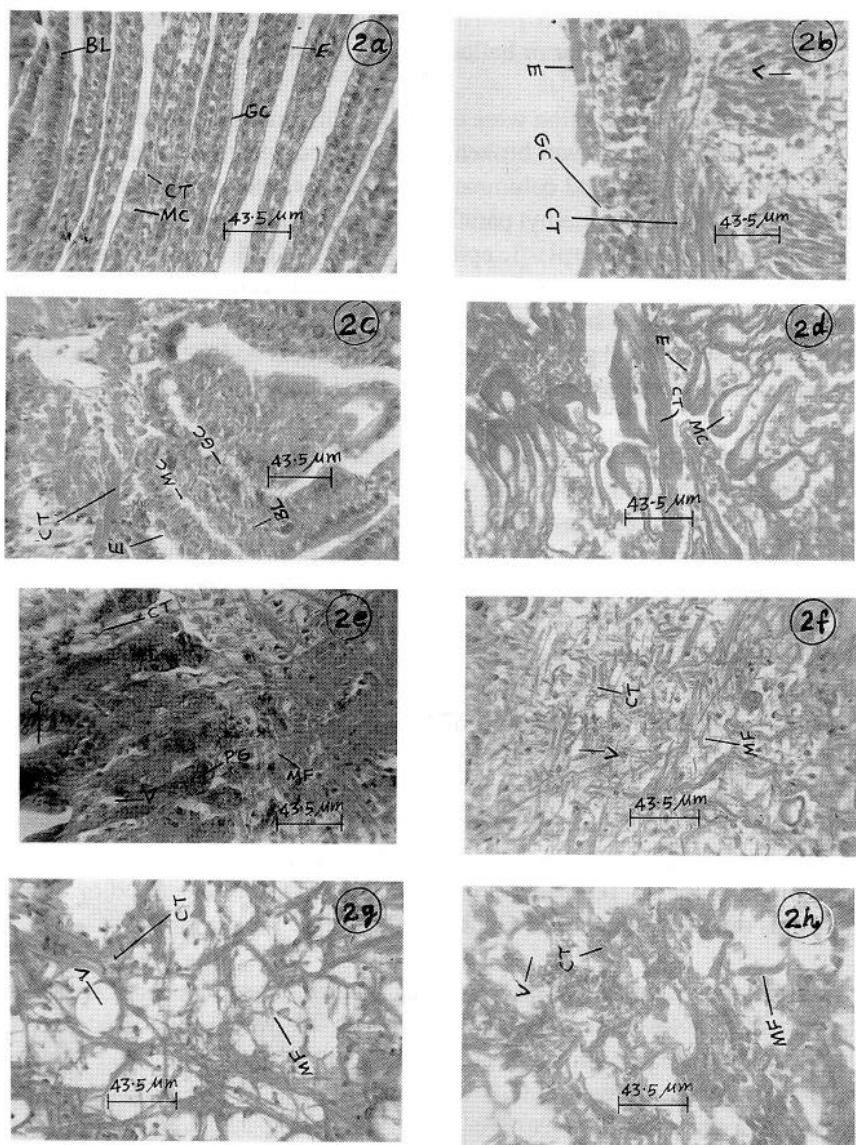


Figure 2. Transverse sections of mantle (Figs 2a to 2d) and foot (Figs 2e to 2h) of (a and e) control (unexposed) and after 96-hr exposure to 96-hr LC<sub>50</sub> concentrations of (b and f) endosulfan, (c and g) methyl parathion, (d and h) nuvan (X250)

E - Epidermis; GC - Gland Cell; CT - Connective Tissue;  
 MC - Mucocyte; BL - Blood Lacuna; MP - Muscle Fibers;  
 V - Vacuole.

together. Hence, it is essential to study the toxicological effects, changes in metabolism and biochemical constituents before arriving at a definite conclusion in the toxicity of the compound.

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