



MORTALITY STUDY OF VIVIPARUS BENGALENSIS AFTER EXPOSURE TO THE PHYTOTOXIN FROM AZADIRACHTA INDICA

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Abstract:

It is common practice to use synthetic chemicals in the control of pesticides but such use of pesticides cause damage to the environment and the living organisms. Hence natural phytotoxins are used in the control of pesticides. The phytotoxin *Azadirachta indica* is used to study its effect on the biochemical contents of the gills of snail *Viviparus bengalensis*. The snails were exposed to the phytotoxin from *A.indica* (neem oil) with sub lethal concentration 148.90 ppm for 24, 48, 72, 96 and 120 hrs. to study its effect on the glycogen, proteins and lipids in the gills of the snail *V. bengalensis*. It is observed that glycogen, proteins and lipid contents of gills were reduced. The results were discussed on the basis of metabolism of snails.

Keywords:- *Azadirachta indica*, *Viviparus bengalensis*, phytotoxin, snails.

Introduction:

Although synthetic chemicals are more effective they are not trustworthy due their bad effects on the aquatic and sub aquatic animals and their environment. They also destroy the food of organisms such as zooplanktons and phytoplanktons, in addition due to the residual effects of the synthetic chemicals and heavy metals affect the food chain and the water bodies get affected for prolonged period. They may even result in mutation and these changes become prominent after few generations. The chemical pesticides also kill friendly insects. Hence scientists had thought about the alternative way of using plant toxins which are easily degradable, comparatively less expensive and cannot create prolonged hazards in the water. Some of the plant species shows properties like specific toxicity, feeding inhibition, growth regulation, and anti-fertility effects. (Grainge and Ahmed;1985 and Parmer and Devkumar, 1993)

The present work is concerned with study of the one of the important vital activity i.e. respiratory activity of the gastropod, *Viviparus bengalensis* after its exposure to the phytotoxins from *Azadirachta indica*. The effect of the phytotoxins on the vital activities of the molluscs is being undertaken for the detailed investigation.

In the present investigation it was decided to carry out detailed analysis of biochemical contents of gills.

Material and Methods:-

Neem oil from *Azadirachta indica* used for present study is market product Neem Shastra (having emulsifier itself) was obtained from Siemen's Agrineed Industries 298/295

village Angapur (Vandan), Satara (Maharashtra). 1 ml of Neem Shastra was dissolved in 100ml 95% ethyl alcohol; this was a stock solution. This solution further diluted into acetone and desired concentrations were prepared (100 to 500 ppm). The sublethal concentration 148.90 of this oil is used for study of mortality.

Observations:

Observation on mortality of *V. bengalensis* after its exposure to phytotoxin from *A. indica*. - Fresh water snail, *V. bengalensis* was exposed to the different concentrations such as 100, 200, 300, 400 and 500 ppm of alcoholic extract of *A. indica* (Neem oil) for different time intervals such as 24, 48, 72, 96 and 120 hrs. respectively.

The per cent mortality of *V. bengalensis* to the different concentrations of phytotoxin from *A. indica* was recorded in the Table No. 1 and graphically represented in figure No.1. No mortality was observed in the control set. Mortality was initiated in 100 ppm concentration after 48 hrs.of exposure period. The maximum mortality was observed after 120 hrs. of exposure *Azadirachta indica* on mortality snail, *Viviparus bengalensis*

period in each concentration. 15%, 30%, 55% 65% and 80% mortality was observed respectively in the concentrations 100, 200, 300, 400 and 500 ppm respectively after 120 hrs. of exposure period.

Different values such as LnXY , ΣLnXY , LnX , ΣLnX , LnX^2 , ΣLnX^2 were calculated with the help of percentage, mortality, probit (Y) and concentrations in ppm

By using numerical data, the values of 'b' and 'a' were calculated. Then the regression

equations and LC₅₀ were calculated and have been recorded in Table No. 2.

Table. 1:- Per cent mortality of *Viviparus bengalensis* after exposure to molluscicide *Azadirachta indica* for different concentrations

Conc. in ppm	Per cent mortality				
	24 hrs	48 hrs	72 hrs	96 hrs	120 hrs
control	00.00	00.00	00.00	00.00	00.00
100	00.00	05.00	10.00	10.00	15.00
200	05.00	15.00	20.00	25.00	30.00
300	15.00	35.00	45.00	50.00	55.00
400	35.00	50.00	60.00	75.00	65.00
500	55.00	65.00	75.00	80.00	80.00

Table. 2:- Regression equation with LC₅₀ values of *Azadirachta indica* neem oil on snail, *Viviparus bengalensis* at different time intervals.

Time of interval (hrs)	Regression equation (Y=a+b X)	LC ₅₀ value in ppm calculated
24	Y = - 10.402 + 5.8084 X	446.8
48	Y = - 2.6503 + 2.9479 X	391.8
72	Y = - 2.1628 + 2.8577 X	318.5
96	Y = - 1.8068 + 2.8668 X	275.7
120	Y = - 1.5131 + 2.6812 X	263.0

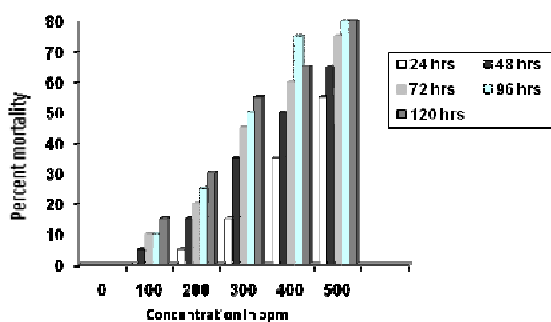


Figure.1 :- Effect of different concentrations of neem oil (Neem Shastra) of

Results and Discussion:

Mortality is most noticeable effect of pollution and it plays an important role in the toxicological studies. It also helps in the determination of LC₅₀ values of toxicant. In the present study of mortality, different concentrations (100, 200, 300, 400 and 500ppm) of the plant toxins from *A.indica* are used to observe the mortality response in the fresh water snail, *V.bengalensis*.

It is observed that mortality of *V.bengalensis* was found altered in different concentrations of phytotoxins at different time intervals.

In the present investigation, it is observed that some snails get died earlier to the effect of plant toxins while some try to survive little more time, when all others are dead. This natural variability in mortality responses is due to the resistance power of animal to each concentration. (Mouirhead - Thomson, 1971)

It is also observed that the mortality of snails under study was observed only in intoxicated sets at definite concentrations of the phytotoxins and not in the control set It is true and noticeable that the pH, temperature,

salinity, hardness and dissolved oxygen of water in control and experimental set is normal even

after addition of the toxicant. This clearly indicated that no factors other than phytotoxins were responsible in this study for the mortality of snail *V- bengalensis*.

In concentration 100 ppm. of the plant toxins from *A. indica* no or less mortality of *V. bengalensis* was observed upto early 24 hrs. because of tolerance, mechanism.

Such type of tolerance phenomenon was observed in case of *T. mossambicus* (Patil, 1988), in *V. bengalensis* from extract of *Eupatorium triplinerve* (Nanaware *et al.*; 2003, and Awati, 2004).

The intoxication data on mortality due to pesticides have been reported by many workers. The pesticide intoxication programme was successfully carried out by some workers by activating the high percentage of mortality as reported in *Lymnae stagnalis* (Bhide 1986 and 1989), in *Pila globosa* (Bhide 1987), *Antheraea agsama* (Bora and Handique 1990), in *Indonaia caeruleus* (Muley 1989) and in *Lymnaea stagnalis* (Bhide 1991).

Percent mortality data recorded in Table No. 1 indicated that the rate of mortality is less in lower concentrations while it is more in higher concentrations of phytotoxins. The percent mortality in the Experimental snails was also seen increased with increase in the exposure period. Therefore percent mortality in the snails under study was found associated with the concentration of toxicant and intoxication period. This also suggests that the rate of mortality in snail, *V. bengalensis* is increased with increase in the concentration of the plant toxins *A. indica* and with increase in the intoxication period also.

Lokhande and Kulkarni (1990) have reported the effect of pesticide toxicity in combination with temperature and pH in *Bellamyia bengalensis*, with similar observations.

According to Malu and Kulkarni (1990) increased mortality is due to increased endosulfan and malathion residue with respect to pesticide exposure time. Higher accumulation of pesticides in mantle than foot and hepatopancreas showed increased mortality in the snail, *Viviparus bengalensis* by (1990) and it was more probably due to pesticide residue accumulation in *Limnaea stagnalis* as reported by Bhide, 1986 and 1989 Similar observations were recorded by Bhide (1987) in *Pila globosa*.

The present investigation provides the data derived from dose concentration and percent mortality for the determination of LC₅₀ values and regression equation by probit analysis. The LC₅₀ values of plant toxins for snail *V. bengalensis* were calculated with respect to the plant toxins derived from *A. indica*. The regression equation is useful for the calculation of lethal concentration. The calculated LC₅₀ value of phytotoxin *A.indica* to the snail *V. Bengalensis* is 263.0 ppm.

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