

Control Of Cotton Pest (*Mylabris Cichorii*) Using *Azadirachta Indica* Leaf And Stem Bark Extracts

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ABSTRACT

The present study was investigated important medicinal plant extract *A. indica* stem bark aqueous extract against the fourth instars larvae of cotton Pest, During the experimental period insects' weight were significantly decreased from 24, 48, 72 and 96 hrs of exposure time of leaves and stem bark aqueous extract of *A. indica* treated against *M. cichorii*. The high percent mortality presents in the 800-ppm concentration of *T. arjuna* at 96 hrs exposure time of leaves extract followed by 600, 400, and 200 ppm, respectively. The high percentage mortality was observed in the 800-ppm concentration of *A. indica* at 96 hrs exposure time of leaves extract followed by 600, 400, and 200 ppm, respectively. The high percentage mortality was observed in the 400-ppm concentration of *A. indica* at 96 hrs exposure time of stem bark extracts followed by 800, 600, and 200 ppm, respectively. These results have been suggested that the *A. indica* plant leaves and stem bark aqueous extracts were used as the integral pest management studies of the cotton pest, *M. cichorii*.

Key words: Biocidal activity, cotton pest management, *M. pustulata*, *A. indica*, mortality rate, weight of the insect.

INTRODUCTION

India is gifted with varied agro-climatic conditions for growing an array of vegetable crops to combat the present ill-balanced diet. The eggplant/brinjal belongs to the family Solanaceae and to the genus Solanum, species melongena. The egg plant, Guinea squash are some synonyms to brinjal. It is a hardy crop and does best in warm and moist climate. The crop is generally grown twice or thrice in a year. The edible part, fruit is of high nutritive value and can well be compared with tomatoes. Insect pest constitutes the major limiting factors in the successful production of eggplant. In the sub-Himalayan region of north east India eggplant is cultivated at a commercial scale but insect and mite pest damage constitutes a limiting factor for its successful production (Ghosh 1999). In the sub-Himalayan region of north east India major vegetables like eggplant, ladyfinger cabbage, cauliflower etc are cultivated at a commercial scale but insect and mite pest damage constitutes a limiting factor for its successful production (Ghosh et al., 1999; Ghosh et al., 2000; Ghosh and Senapati 2001a; Ghosh and Senapati 2001b; Chaudhury et al., 2001). Hadda/ spotted beetle (*Epilachne vigintioctopunctata*), aphid (*Aphis gossypii*), leafhopper (*Amrasca biguttula biguttula*), thrips (*Thrips tabaci*), spider mite (*Tetranychus* spp.), blister beetle (*Mylabris pustulata* Thunberg) and white fly (*Bemisia tabaci*) are the important pests of eggplant causes heavy damage (Ghosh, 1999). Pest complex of eggplant is very high and blister beetle causes heavy damage to the crop. These beetles are brightly colored insects that secrete a compound containing cantharidin when disturbed. Cantharidin, a terpenoid, produces blisters on human skin upon contact and hence these beetles are called "blister" beetles. In general, this is a minor pest, although occasional outbreaks can occur.

MATERIALS AND METHODS

Collection and preparation of stem bark and leaves of *A. indica*

The barks and leaves of *A. indica* were collected from our college campus, The barks of the plant were washed thoroughly with distilled water to remove dust and other particles. The washed plant part is then dried at room temperature. The collected *A. indica* barks and leaves were cut into small pieces. The plant parts were dried in an incubator for 7 days at 40°C, crushed in an electrical grinder and then the powder was separated.

Extraction Methods

For extraction of secondary metabolites cold percolation methods (Thillairajasekar *et al.*, 2009; Das *et al.*, 2010) was followed.

Cold percolation method

For cold percolation method, 250 gm of the powdered stem bark of *A. indica* material was soaked with 750 ml (1:3 w/v) of selected aqueous in an aspirator bottle for 48 hrs at room temperature. The extract was filtered through a funnel with Whatman number 1 filter paper. After filtration, the filtered extract was poured into the distillation unit at 20° C for separation of the solvent and the secondary metabolites residue (10 mL) was evaporated and dried over sodium sulphate in a desiccator under vacuum. The crude extracts were stored in the refrigerator (LG, India) for further use. The extraction rate was calculated by weighing the crude extracts obtained from 100 g of dry plant material after extraction with respective solvent.

Pest collection and rearing

Nymphs and adults of *M. cichorii*, were collected from cotton fields. The collected pests were maintained in the insectory under laboratory conditions (temperature 28 ± 2°C, 70 ± 5 % RH and a photoperiod of 11L: 13D hrs) in transparent plastic containers (8cm height × 6.5cm diameter) containing a layer of sterile coarse sand (4cm thick). Insects were fed with its natural host cotton flower and also cotton bolls. Insects were maintained at least for 2 generations. The laboratory emerged 6-12 hrs old third stadium *M. cichorii* were used for this experiment.

2.8. Nymphicidal activity bioassay

Bioassay studies were carried out using uniform sized (24.7±0.4 mg weight), 6-12 hrs old third stadium *M. cichorii*. which was selected randomly from the stock culture. Five insects were placed in a transparent plastic container (8 cm height × 6.5 cm diameter). Different concentrations of *A. indica* stem bark aqueous extracts [200, 400, 600 and 800ppm (4mg extract in 5ml diet- 800 ppm)] mixed in aqueous. Experimental animals were allowed to feed for 96 hrs continuously. 200µL of aqueous extracts of stem bark was pour into the small cotton flowers and provided to the insects. The food was changed every day. Three replications were maintained for each concentration. Mortality was recorded every 24 hrs, till 96 hrs. The mortality was corrected using Abbott's formula (Abbott, 1925), if more than 10% mortality was observed in control category. The corrected mortality data was subjected to probit analysis (Finney, 1971) to find out the LC₅₀, Chi square, df and p values. After 96 hrs, live nymphs were provided with water-soaked cotton flowers till their death.

RESULTS

During the experimental time the weight of the insect was significantly decreased in 800 ppm (t=3.5; df= 5; p=0.057), 400 ppm (t=10.07; df=5; p=0.000) and 600 ppm (t=18.31; df=5; p=0.000) at 24 hrs of exposure time of stem bark aqueous extract compare to control at 200 ppm the weight was increased when compare to control (Figure 1). During the experimental time the weight of the insect was significantly decreased in 800 ppm (t=5.5; df= 5; p=0.057), 200 ppm (t=10.07; df=5; p=0.000), 400 ppm (t=17.31; df=5; p=0.000) and 600 ppm (t=10.07; df=5; p=0.000) and 800 ppm (t=16.07; df=5; p=0.000) at 48 hrs of exposure time of stem

bark aqueous extract when compare to control (Figure 2). During the experimental time the weight of the insect was significantly decreased in 400 ppm ($t=32.55$; $df=5$; $p=0.000$), 200 ppm ($t=19.20$; $df=5$; $p=0.000$), 600 ppm ($t=16.10$; $df=5$; $p=0.000$) and 800 ppm ($t=17.10$; $df=5$; $p=0.000$) at 72 hrs of exposure time of stem bark aqueous extract when compare to control (Figure 3). During the experimental time the weight of the insect was significantly decreased in 800 ppm ($t=26.55$; $df=5$; $p=0.000$), 200 ppm ($t=19.10$; $df=5$; $p=0.000$), 600 ppm ($t=15.00$; $df=5$; $p=0.000$) and 400 ppm all the animals are dead at 96 hrs of exposure time of stem bark aqueous extract when compare to control. The high percentage mortality was observed in the 400-ppm concentration of *A. indica* at 96 hrs exposure time of stem bark extracts followed by 800, 600, and 200 ppm, respectively (Figure 4).

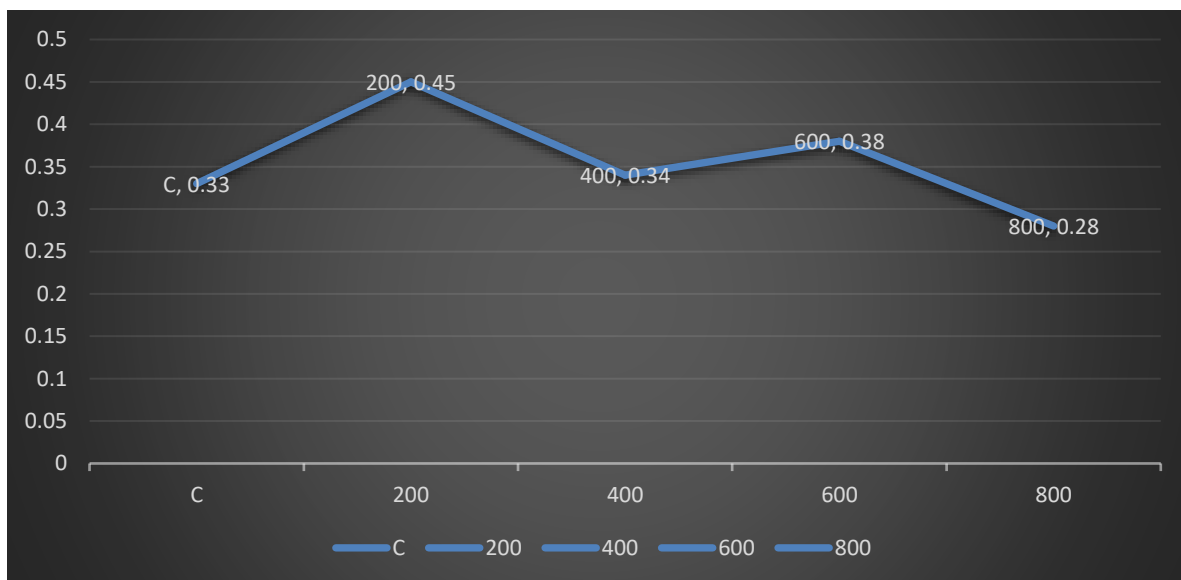


Figure 1. Effect of *A. indica* stem bark aqueous extract on whole body wet weight (g) of *M. cichorii* nymphs at 24 hrs

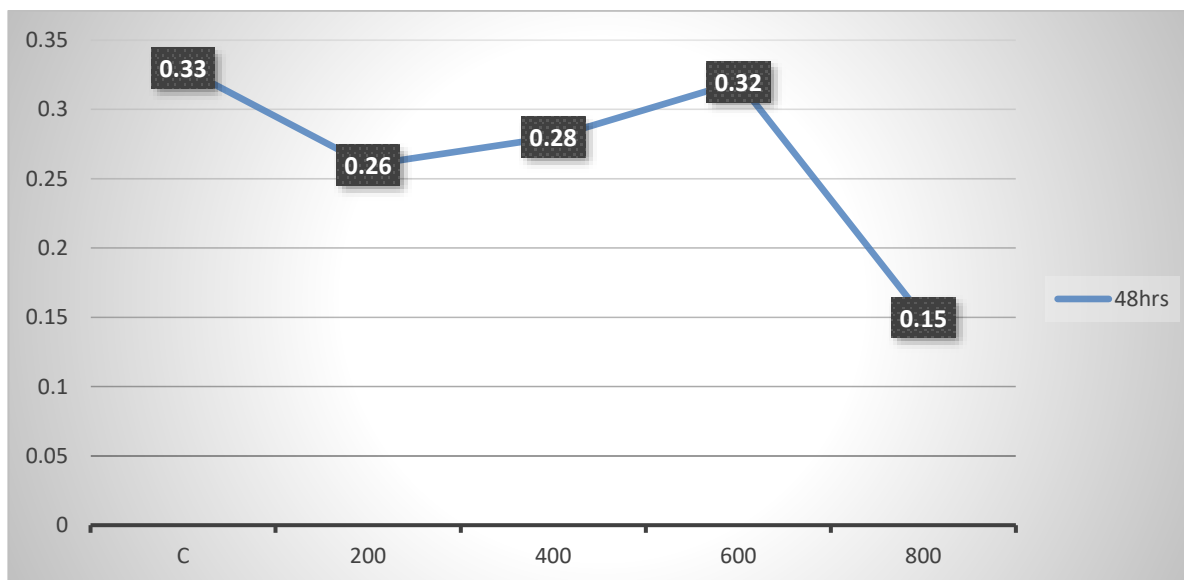


Figure 2. Effect of *A. indica* stem bark aqueous extract on whole body wet weight (g) of *M. cichorii* nymphs at 48 hrs

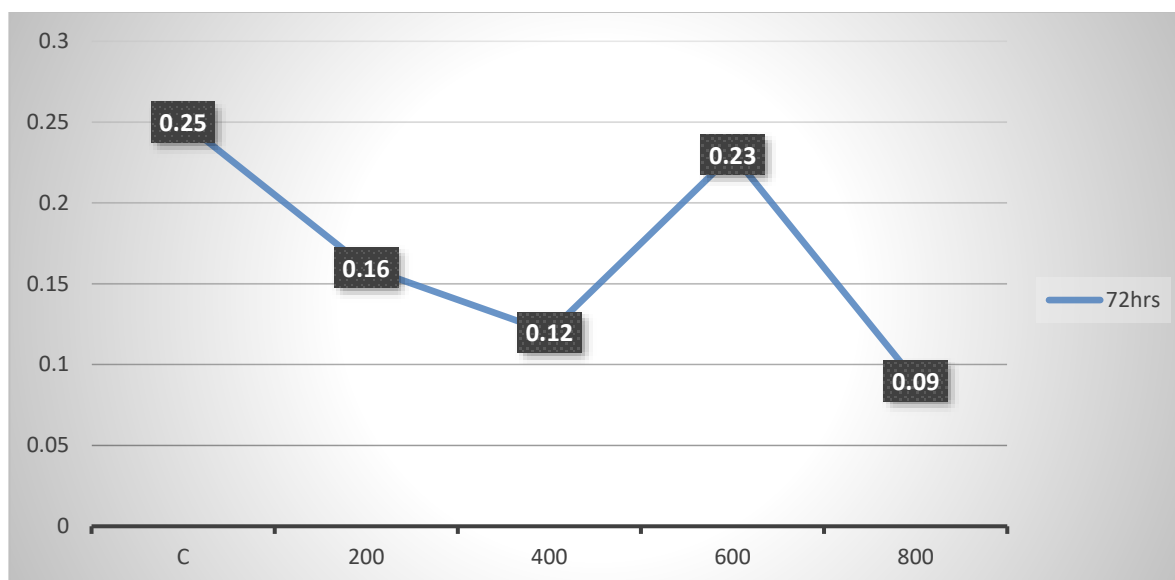


Figure 3. Effect of *A. indica* stem bark aqueous extract on whole body wet weight (g) of *M. cichorii* nymphs at 72 hrs

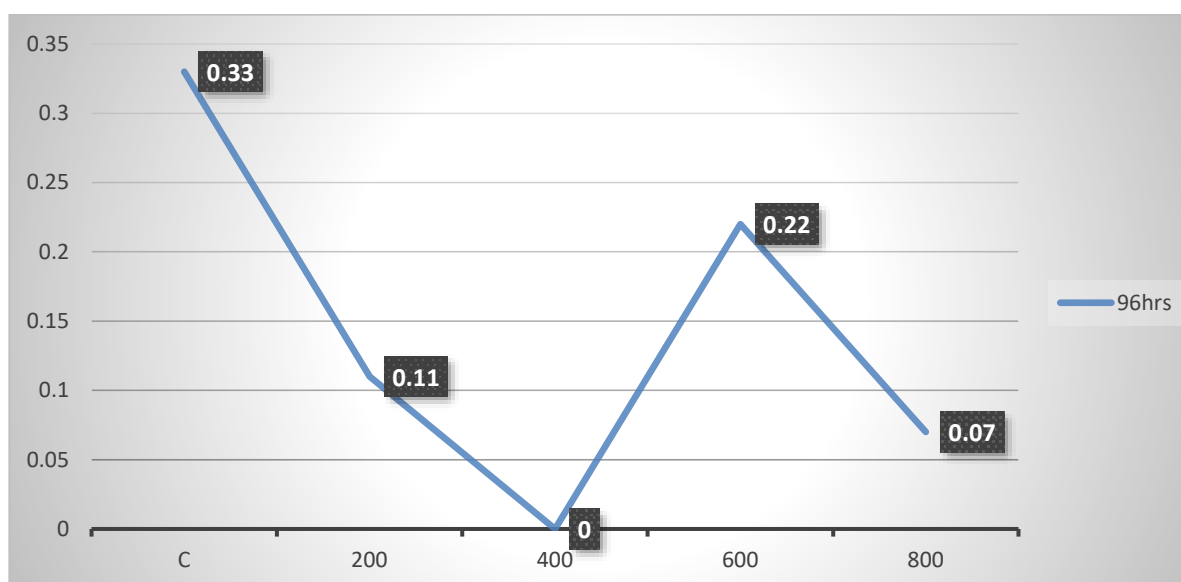


Figure 4. Effect of *A. indica* stem bark water extract on whole body wet weight (g) of *M. cichorii* nymphs at 96 hrs

DISCUSSION

During this study, toxicity and effectiveness of 2 2-tridecanone formulations were assessed against *S. invicta*. In field path, at application rate of 5.28 ml/L and 14 days once mound drench treatment, 100% management was achieved for formulation with piperonyl butoxide (PBO) and 9th management for the formulation while not PBO (Chen, 2016). Signified that antifeedants can be found amongst all the major classes of secondary metabolites, viz., alkaloids, phenolics, and terpenoids which the most probable toxic substances against insects. The ethyl acetate extract of the plant diminished the feeding rate of *S. litura*. The is indicated that the active principles present in the plants retard larval feeding department or make the food unpalatable or the substances directly act on the chemo essentials of the larval resulting in feeding deterrence. Several investigators have already reported that botanicals offer antifeedant action against *S. litura* (Arivoli, 2012). Mendhulkar et al., 2014 investigation that the liquid extracts of *Couroupita guianensis* leaves show high insecticidal impacts on nymphs and adult flies, whereas low effect on the eggs of *Bemisia tabaci* as compared to regulate. Ahmad et al.2013 effectuality the insecticidal activity of *Allium sativum*, *Zingiber officinale*, and *Nigella sativa* extracts against the larvae of *Trogoderma granarium*. The

very best concentration of (6%) *Z. officinale* was found to be relatively a lot of cytotoxic (16.70%) than those of *A. sativum* (10.45%) and *N. sativa* (5.49%) at 96 hrs exposure. In conclusion, nourishment of *T. arjuna* and leaves and bark aqueous extracts were tested against fourth instars larvae of *M. pustulata*, the present investigation suggested that the *T. arjuna* plant leaves and stem bark aqueous extracts were used as the integral pest management studies of the cotton pest, *M. pustulata*.

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