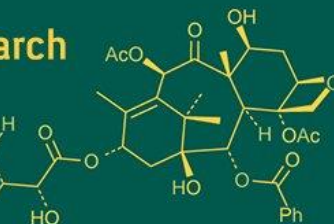
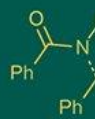


International Journal of Advanced Biochemistry Research



ISSN Print: 2617-4693
ISSN Online: 2617-4707
NAAS Rating (2025): 5.29
IJABR 2025; SP-9(10): 458-463
www.biochemjournal.com
Received: 13-07-2025
Accepted: 16-08-2025

Nahida Afreen
Chandra Sekhar Azad
University of Agriculture and
Technology, Kanpur, Uttar
Pradesh, India

AN Singh
ICAR-Indian Institute of
Vegetable Research, Varanasi,
Uttar Pradesh, India

CL Maurya
Chandra Sekhar Azad
University of Agriculture and
Technology, Kanpur, Uttar
Pradesh, India

SK Biswas
Chandra Sekhar Azad
University of Agriculture and
Technology, Kanpur, Uttar
Pradesh, India

Naushad Khan
Chandra Sekhar Azad
University of Agriculture and
Technology, Kanpur, Uttar
Pradesh, India

Corresponding Author:
Nahida Afreen
Chandra Sekhar Azad
University of Agriculture and
Technology, Kanpur, Uttar
Pradesh, India

Lethal and sub-lethal effects of fipronil on growth and development of okra shoot and fruit borer (*Earias vitella*: Noctuidae: Lepidoptera)

Nahida Afreen, AN Singh, CL Maurya, SK Biswas and Naushad Khan

DOI: <https://www.doi.org/10.33545/26174693.2025.v9.i10Sf.5873>

Abstract

The shoot and fruit borer *Earias vitella* (Fabricius) is a major pest affecting okra (*Abelmoschus esculentus*), causing significant yield and quality losses across all crop stages. Chemical control remains a primary strategy, with fipronil, a phenyl pyrazole insecticide, being widely used. This study evaluated the lethal (LC₅₀) and sub-lethal (LC₁₅ and LC₁₀) effects of fipronil on the growth and development of *E. vitella* under laboratory conditions at ICAR-IIVR, Varanasi, using diet incorporation bioassays. The LC₅₀ value was determined to be 4.844 ppm, while LC₁₅ and LC₁₀ were 0.068 ppm and 0.025 ppm, respectively. Sub-lethal doses significantly influenced key biological parameters, including prolonged larval and pupal durations, reduced larval and pupal weight and length, increased mortality, reduced pupation and adult emergence rates, and decreased fecundity. Growth index and adult longevity were also notably reduced compared to the control. These findings suggest that, beyond direct lethality, sub-lethal exposures to fipronil can impair developmental and reproductive fitness in *E. vitella*, potentially offering extended pest suppression benefits. The study highlights the importance of incorporating sub-lethal effects into insecticide efficacy assessments for integrated pest management (IPM) strategies in okra cultivation.

Keywords: *Earias vitella*, LC₅₀, okra, Sub lethal

Introduction

Vegetables are fresh, edible parts of herbaceous plants. They are a vital food that is very helpful in preserving health and preventing illness. They include beneficial food components that the body may effectively use to strengthen and heal itself. Vegetables are beneficial for preserving the body's alkaline reserves. Their high levels of carbohydrates, vitamins, and minerals make them valuable. Vegetables come in several varieties. They could be edible seeds, fruits, stems, leaves, or roots. Every vegetable makes a unique contribution to diet (Robinson, 1990) [3]. Due to the nation's increased vegetable output, the number of vegetables available per person increased from 264 grams per day in 2004-05 to 355 grams per person per day in 2011-16.

India ranks first in the production of peas (*Pisum sativum*) and okra (*Abelmoschus esculentus*) and second in the production of tomatoes (*Solanum lycopersicum*), cauliflower (*Brassica oleracea*), potatoes (*Solanum tuberosum*), onions (*Allium cepa*), and brinjal (*Solanum melongena*). The nation's primary cereals can be supplemented with vegetables, which are a rich and relatively inexpensive source of proteins, carbs, vitamins, and minerals. Vegetables provide us with a variety of chemicals that our bodies need for development, reproduction, and health maintenance.

Earias vitella (Fabricius) (Lepidoptera: Noctuidae) is a shoot and fruit borer that is a major pest of lepidopterans that is found throughout North Africa, India, Pakistan, and several other countries. It mostly affects crops in the Malvaceae family, including cotton (Nada *et al.* 2010; Kumar *et al.* 2014) [4, 5] and okra (Aziz *et al.*, 2011) [2]. It has a wide host range. These insects directly harm the delicate shoots and fruits of okra, causing the afflicted plants to produce smaller, malformed fruits (Rahman *et al.*, 2013) [6]. Okra is attacked by shoot and fruit borers at every stage of growth, which seriously impairs the crop's quantity and quality (Shah *et al.*, 2001) [7]. Fruit loss from *Earias spp.* infestations ranges from 21.00% to 91.58% in different okra varieties. They damage the crop throughout vegetative and the reproductive

stages, thus causing ample decline in yield (Kataria and Singh 2021)^[8].

Fipronil, which is a novel insecticide of the phenyl pyrazole class, is 5-amino-1-(2,6-dichloro- α,α,α -trifluoro-p-tolyl)-4-trifluoromethylsulfinyl pyrazole-3-carbonitrile. Rhone-Poulenc Ag Company (now Bayer Crop Science) produced it for the first time in 1987, and the product was introduced in 1993 (Tomlin, 2000). It inhibits the normal activity of GABA by targeting gamma-aminobutyric acid (GABA) receptors, which are the main neurotransmitters in insects (Sammelson *et al.*, 2004)^[9].

Fipronil is a very effective compound; piercing, sucking, and chewing insect pests that are resistant to other chemicals like pyrethroids, organophosphates, and carbamates may often be controlled with just a few grams of the active ingredient per hectare.

While fipronil is commonly used to control various pests, including *E. vitella*, its potential impacts on the pest's biology beyond simple mortality are not well-documented. Understanding the lethal effects of these insecticides is critical for evaluating its direct efficacy in controlling *E. vitella* populations, but equally important is examining the sub-lethal effects, which can influence pest behaviour, reproduction, and overall pest dynamics. Sub-lethal exposure to insecticides can lead to changes in feeding patterns, decreased reproductive success, altered development rates, and increased pest resistance over time, even in the absence of immediate mortality. Given the importance of *E. vitella* as a pest of okra, it is essential to understand how sub-lethal doses of fipronil and novaluron might affect pest survival and fitness in the long term. This study will provide crucial insights into the broader impact of insecticide application, helping to inform future pest control

practices and contribute to the sustainable management of *E. vitella* in okra cultivation.

Materials & Method

Collection and rearing of fruit and shoot borer

Larvae of *Earias vitella* were collected from unsprayed okra fields of ICAR- Indian Institute of Vegetable Research, Varanasi and maintained in the laboratory under controlled conditions in transparent plastic jars (25 ± 2 °C, $60 \pm 5\%$ relative humidity). The larvae were provided with fresh okra fruit until they pupated. Once pupation occurred, the pupae were transferred to a plastic box lined with tissue paper. Upon emergence, the adult moths were fed a 10% sugar solution. Butter paper strips were placed in the rearing cage for egg laying. The eggs were collected and transferred to separate plastic jars (1 kg capacity). After hatching, the second instar larvae of *E. vitella* were used for bioassays and biological studies. Precautions: The utmost hygienic conditions were kept in rearing facilities.

Artificial diet composition and preparation

Earias vitella were reared on artificial diet. An artificial diet was developed for mass rearing of the *Earias vitella* in environmentally controlled conditions. The ingredients for 1l of diet ingredients consisting of soybean flour (*Glycine max*) (60.0 g), chickpea flour (*Cicer arietinum* var. *kabuli*) (58.0 g), wheatgerm (16.0 g), dried yeast powder (16.0 g), casein (8.0 g), L-ascorbic acid (2.4 g), cholesterol (0.4 g), methyl-p-hydroxybenzoate (0.4 g), sorbic acid (2.4 g), streptomycin sulphate (0.4 g), agar-agar (16.0 g) and distilled water (820 ml). The insect rearing and mass multiplication was done using the methodology given by (Gupta *et al.*, 2005)^[11].

Table 1. Composition of artificial diet for rearing *Earias vitella* larvae

Groups	Ingredients	Quantity (g)
Fraction A	Soybean flour (<i>Glycine max</i>)	60.0
	Chickpea flour (<i>Cicer arietinum</i> var. <i>kabuli</i>)	58.0
	Wheatgerm (Wheatgerm, Bakers, India Ltd.)	16.0
Fraction B	Dried yeast powder (Tower Brand, The Indian Yeast Co. Ltd.)	16.0
	Casein (Central Drug House, Ltd.)	8.0
	L-Ascorbic acid (SDS, s.d. Fine Chem Ltd.)	2.4
	Cholesterol (SDS, s.d. Fine Chem Ltd.)	0.4
Fraction C	Methyl-p-hydroxybenzoate (LOBA Chemie)	0.4
	Sorbic acid (SDS, s.d. Fine Chem Ltd.)	2.4
	Streptomycin sulphate (Indian Drugs and Pharmaceuticals Ltd.)	0.4
Fraction D	Agar-agar (Qualigens Fine Chemicals)	16.0
	Distilled water	820 ml

The ingredients for the artificial diet of *E. vitella* are provided in Table 3.3 The components of fraction 'A' were thoroughly mixed in a blender with 400 ml of lukewarm distilled water. Agar-agar (fraction 'D') was dissolved in 420 ml of lukewarm water, then boiled in a Borosil beaker. Once cooled to approximately 50° C, it was added to fraction 'A' in the blender. This mixture was blended at high speed for about 1 minute. Next, the ingredients of fractions 'B' and 'C' were added, and the mixture was well stirred. The resulting warm, homogeneous diet mixture was poured into sterilized 15 x 2.5 cm Petri dishes and allowed to cool. After solidifying at room temperature (25-30 °C), the diet was refrigerated.

Toxicity Bioassay

To find out LC₅₀ bioassay experiments were performed using the diet incorporation method. Different concentration of insecticides were added to the artificially prepared diet in the bioassay tray on which insects were reared and evaluated further. The experiment was replicated three times. Mortality was noted after 24 hours after treatment. Lethal and sublethal effects against *E. vitella* larvae was evaluated using a lethal and sublethal concentration calculated from the toxicity experiment (Gulzar *et al.* 2019)^[12].

Statistical analysis

Mortality counts were taken 24 hours after treatment. The moribund insects were considered as dead. The mortality data so obtained were converted into corrected per cent mortality using the Abbott's formula (Abbott, 1925) as given below:

$$P'' = P' - C / 100 - C \times 100$$

Where,

P'' = Corrected per cent mortality in the test insect

P' = Observed per cent mortality in the test insect

C = Per cent mortality in the control

The corrected per cent mortality data thus obtained were subjected to the probit analysis (Finney, 1971) ^[13] to compute LC_{50} value for each insecticide.

Assessing effect of sub lethal dose

The lethal and sublethal effects of fipronil and novaluron on *E. vitella* larvae were assessed using lethal and sublethal concentrations of fipronil and novaluron determined from the toxicity experiment using diet incorporation method.

To study the lethal and sub-lethal effect on growth, lethal and sub-lethal concentration of fipronil and novaluron, were tested against *E. vitella*. The sublethal dose (LC_{10} and LC_{15}) of each insecticide was applied using the diet incorporation method. Second instar larvae were carefully transferred to bioassay tray containing toxicant incorporated artificial diet using a camel hair brush. Three replications of each treatment were maintained, with 15 larvae per replication. The bioassay trays were placed in a BOD incubator set at 28 ± 2 °C with $70 \pm 5\%$ relative humidity. After 24 hours of

exposure, the larvae were transferred to jars containing fresh okra fruits, and observations were made until the completion of the life cycle.

Observations were made on larval length, weight of mature larvae, larval longevity, and mortality. Similar data were also collected for pupal length, weight, longevity, and mortality. Pupae were weighed at two days of age to ensure they had hardened sufficiently to avoid damage during handling. Each individual pupa was properly labelled and placed in small jars for further observation, including the time required for adult emergence, percentage of adult emergence, and adult longevity. The sex of the *E. vitella* pupae was determined by examining the cocoon, as described by Gupta (1978) ^[14], who noted that male cocoons have a distinct knob at the antero-dorsal end, which is absent in females. Moths that emerged from larvae exposed to sublethal doses of fipronil were observed to record the longevity of both males and females. These moths were placed in an egg-laying chamber to assess the fecundity of individual females. The results were compared to the control group.

Results and Discussion

Calculation of LC_{50} , LC_{15} and LC_{10} of fipronil against *Earias vitella*.

The toxicity of fipronil and novaluron by bio-assaying second instar larvae of *Earias vitella* was determined in laboratory by artificial diet incorporation method using Mekapogu, A.R (2021) Finney's probit analysis spreadsheet calculator (Version 2021). LC_{50} value of fipronil was found to be 4.844 ppm. The value of LC_{15} and LC_{10} of fipronil were calculated 0.068 ppm and 0.025 ppm respectively (Table 2).

Table 2; Comparative toxicity of fipronil and novaluron against second instar larvae of *E. vitella* by diet incorporation method

Insecticides	Regression equation	LC_{50} (ppm)	Fiducial limit	LC_{15} (ppm)	Fiducial limit	LC_{10} (ppm)	Fiducial limit	χ^2 (df)	Slope
Fipronil	$0.562x + 4.6339$	4.844	1.239-18.939	0.068	0.017- 0.266	0.025	0.006-0.097	0.996(4)	0.0571

Y= probit kill; X=log (concentration in per cent $\times 10^{-3}$); LC_{50} = concentration (parts per million) calculated to give 50 per cent mortality.

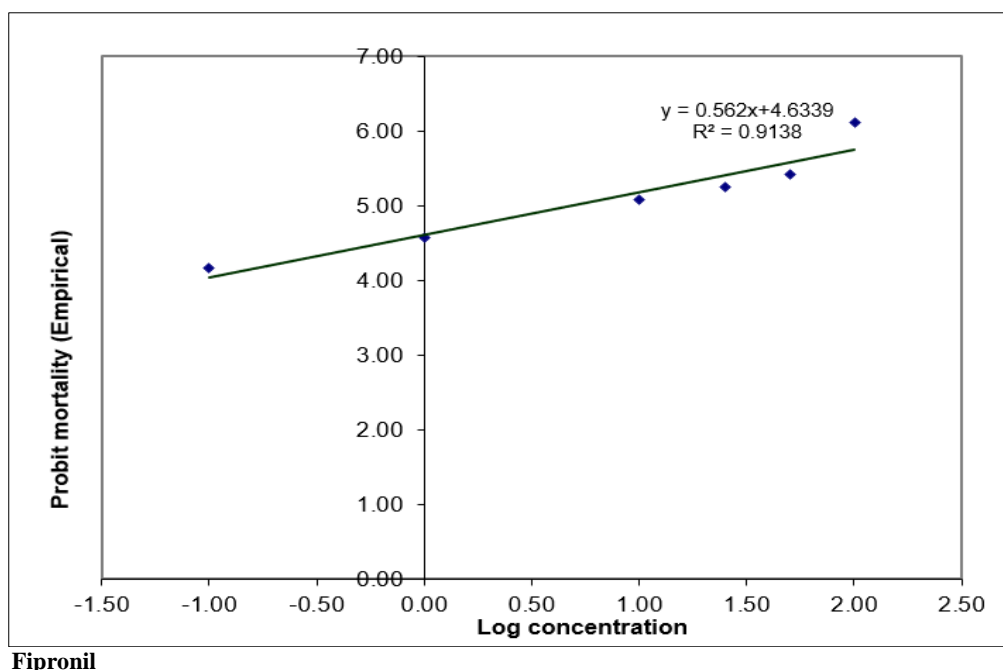


Fig 1: Standard doses mortality curve for second instar larvae of *E.vitella* (Artificial diet incorporation method)

Effect of lethal and Sub-lethal doses on growth and development of *Earias vitella*.

The effect of lethal and sublethal doses (LC₅₀, LC₁₅, LC₁₀) of insecticides viz. fipronil and novaluron was assessed on the basis of their effect on the larval period, larval weight, larval length, larval mortality, per cent pupation, pupal period, pupal weight, pupal length, pupal mortality, per cent adult emergence, adult longevity, fecundity and growth index.

Effect of lethal and Sub-lethal doses on Larvae Larval longevity

Fipronil

Effect of lethal and sub lethal doses of fipronil on larval longevity presented in Table 4.15. The longest larval period was 16.06 days in T₁ (LC₅₀), followed by 13.4 days in T₃ (LC₁₀), 12.66 days in T₄ (control), and 12.43 days in T₂ (LC₁₅). Both T₃ (LC₁₀) and T₁ (LC₅₀) had significantly longer larval periods compared to T₂ (LC₁₅) and T₄ (control).

Table 3: Effect of lethal and Sub-lethal doses on Larval longevity (Days)

Larval longevity (days)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	16.06	-26.84
T ₂ (LC ₁₅)	12.43	-5.78
T ₃ (LC ₁₀)	13.4	1.84
T ₄ (Control)	12.66	0
C.D.	0.428	
SE(m)	0.129	

Larval weight

Fipronil

Effect of lethal and sub lethal doses of fipronil on larval weight presented in Table 4.16. The highest mean weight of 0.089 g was recorded in T₄ (control) which is reduced to 0.079 g in T₃ (LC₁₀), 0.077 g in T₂ (LC₁₅) and 0.074 g in T₁ (LC₅₀). T₁ (LC₅₀), T₂ (LC₁₅) and T₃ (LC₁₀) were significantly at par with each other.

Table 4: Effect of lethal and Sub-lethal doses on Larval weight (g)

Larval weight (g)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	0.074	16.85
T ₂ (LC ₁₅)	0.077	13.48
T ₃ (LC ₁₀)	0.079	11.23
T ₄ (Control)	0.089	0
C.D.	0.008	
SE(m)	0.003	

Larval length

Fipronil

Effect of lethal and sub lethal doses of fipronil on larval length presented in Table 4.17. In T₁ (LC₅₀) and T₂ (LC₁₅) there is significant reduction in length 14.98 mm and 15.98 mm respectively, whereas there was no significant reduction 15.96 mm in T₃ (LC₁₀) as compared to 17.07 mm in T₄ (control).

Table 5: Effect of lethal and Sub-lethal doses on Larval length (mm)

Larval length (mm)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	14.98	12.24
T ₂ (LC ₁₅)	15.98	6.36
T ₃ (LC ₁₀)	16.24	4.84
T ₄ (Control)	17.07	0
C.D.	0.991	
SE(m)	0.299	

Larval mortality

Fipronil

Effect of lethal and sub lethal doses of fipronil on larval mortality presented in Table 4.18. The maximum mortality is 41.66 per cent in T₁ (LC₅₀), followed by 38.33 per cent in T₂ (LC₁₅), 36.66 per cent in T₃ (LC₁₀). Significantly lower mortality was found 6.66 per cent in T₄ (control). It was observed that 41.66 per cent in T₁ (LC₅₀), 38.33 per cent in T₂ (LC₁₅) and 36.66 per cent T₃ (LC₁₀) were significantly at par.

Table 6: Effect of lethal and Sub-lethal doses on Larval mortality (%)

Larval mortality (%)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	41.66	-524.97
T ₂ (LC ₁₅)	38.33	-474.96
T ₃ (LC ₁₀)	36.66	-449
T ₄ (Control)	6.66	0
C.D.	14.603	
SE(m)	4.41	

Effect of lethal and Sub-lethal doses on Pupa

Effect on pupation

Fipronil

Effect of lethal and sub lethal doses of fipronil on pupation percentage presented in Table 4.19. Fipronil doses cause significant reduction in percent pupation when compared to control. Maximum pupation percent was found 90.54 per cent in T₄ (control) followed by 60.47 per cent in T₃ (LC₁₀), 51.94 per cent in T₂ (LC₁₅) and minimum 51.25 per cent in T₁ (LC₅₀). It was observed that 51.94 per cent in T₁ (LC₅₀) and 51.25 per cent T₂ (LC₁₅) were significantly at par.

Table 7: Effect of lethal and Sub-lethal doses on Pupation (%)

Pupation (%)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	51.25	43.39
T ₂ (LC ₁₅)	51.94	42.62
T ₃ (LC ₁₀)	60.47	33.21
T ₄ (Control)	90.54	0
C.D.	3.11	
SE(m)	0.94	

Pupal longevity

Fipronil

Effect of lethal and sub lethal doses of fipronil on pupal longevity presented in Table 4.20. Pupal longevity differs from each other with maximum pupal period 11.57 days in T₁ (LC₅₀) followed by 9.59 days in T₂ (LC₁₅), 9.39 days in T₃ (LC₁₀) and minimum 7.68 days in T₄ (control). It was observed that 9.59 days in T₂ (LC₁₅) and 9.39 days in T₃ (LC₁₀) were significantly at par with each other.

Table 8: Effect of lethal and Sub-lethal doses on Pupal longevity (Days)

Pupal longevity (Days)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	11.57	-50.63
T ₂ (LC ₁₅)	9.59	-24.86
T ₃ (LC ₁₀)	9.39	-22.25
T ₄ (Control)	7.68	0
C.D.	1.952	
SE(m)	0.589	

Pupal weight**Fipronil**

Effect of lethal and sub lethal doses of fipronil on pupal weight presented in Table 4.21. The lowest pupal weight was found 0.057 g in T₁ (LC₅₀) followed by 0.059 g in T₂ (LC₁₅), 0.066 g in T₃ (LC₁₀) and 0.069 in T₄ (control). There is significant reduction in 0.057 g in T₁ (LC₅₀) and 0.059 g in T₂ (LC₁₅) over control. It was observed that 0.057g in T₁ (LC₅₀), 0.059g in T₂ (LC₁₅) and 0.066g T₃ (LC₁₀) were significantly at par with each other.

Table 9: Effect of lethal and Sub-lethal doses on Pupal weight (g)

Pupal weight (g)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	0.057	17.39
T ₂ (LC ₁₅)	0.059	14.49
T ₃ (LC ₁₀)	0.066	4.32
T ₄ (Control)	0.069	0
C.D.	0.009	
SE(m)	0.003	

Pupal length**Fipronil**

Effect of lethal and sub lethal doses of fipronil on pupal length presented in Table 4.22. The pupal length reduced significantly in treatments when compared to 10.68 mm in T₄ (control). Minimum pupal length was observed in 8.39 mm T₁ (LC₅₀) followed by 9.44 mm in T₂ (LC₁₅) and 9.8 mm in T₃ (LC₁₀). It was observed that 9.44 mm in T₂ (LC₁₅) and 9.8 mm in T₃ (LC₁₀) were significantly at par with each other.

Table 10: Effect of lethal and Sub-lethal doses on Pupal length (mm)

Pupal length (mm)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	8.39	21.41
T ₂ (LC ₁₅)	9.44	11.61
T ₃ (LC ₁₀)	9.8	8.23
T ₄ (Control)	10.68	0
C.D.	0.669	
SE(m)	0.202	

Pupal mortality**Fipronil**

Effect of lethal and sub lethal doses of fipronil on pupal mortality presented in Table 4.23. The highest mortality was found 24.74 per cent in T₁ (LC₅₀) followed by 22.11 per cent in T₂ (LC₁₅) and 22.01 per cent in T₃ (LC₁₀). All these treatments recorded significantly higher mortality than 12.58 percent in T₄ (control).

Table 11: Effect of lethal and Sub-lethal doses on Pupal mortality (%)

Pupal mortality (%)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	24.74	-75.75
T ₂ (LC ₁₅)	22.11	-74.96
T ₃ (LC ₁₀)	22.01	-96.68
T ₄ (Control)	12.58	0
C.D.	2.316	
SE(m)	0.699	

Effect of lethal and Sub-lethal doses on adult**Effect on adult emergence****Fipronil**

Effect of lethal and sub lethal doses of fipronil on adult emergence presented in Table 4.24. The lowest adult emergence was found 72.73 in T₁ (LC₅₀) followed by 73.91 per cent in T₂ (LC₁₅), 77.73 per cent in T₃ (LC₁₀) which are significantly lower than 86.14 per cent T₄ (control). T₁ (LC₅₀) 72.73 per cent and T₂ (LC₁₅) 73.91 per cent were significantly at par with each other.

Table 12: Effect of lethal and Sub-lethal doses on adult emergence (%)

Adult emergence (%)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	72.73	15.56
T ₂ (LC ₁₅)	73.91	14.09
T ₃ (LC ₁₀)	77.35	10.19
T ₄ (Control)	86.14	0
C.D.	3.869	
SE(m)	1.168	

Adult longevity**Male longevity****Fipronil**

Effect of lethal and sub lethal doses of fipronil on male longevity presented in Table 4.25. Significantly less male longevity was observed 3.13 days in T₁ (LC₅₀), 3.39 days in T₂ (LC₁₅) and 3.13 days in T₃ (LC₁₀) as compared to 5.35 days in T₄ (control). It was observed that T₁ (LC₅₀), T₂ (LC₁₅) and T₃ (LC₁₀) were significantly at par with each other.

Table 13: Effect of lethal and Sub-lethal doses on Male longevity (Days)

Male longevity (Days)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	3.13	41.47
T ₂ (LC ₁₅)	3.39	36.67
T ₃ (LC ₁₀)	3.62	32.37
T ₄ (Control)	5.35	0
C.D.	1.137	
SE(m)	0.343	

Female longevity**Fipronil**

Effect of lethal and sub lethal doses of fipronil on female longevity presented in Table 4.26. Significantly lower female adult longevity was recorded 7.33 days in T₁ (LC₅₀), 7.75 days in T₂ (LC₁₅) and 7.76 days in T₃ (LC₁₀) when compared to 9.64 days in T₄ (control). It was observed that T₁ (LC₅₀), T₂ (LC₁₅) and T₃ (LC₁₀) were significantly at par with each other.

Table 14: Effect of lethal and Sub-lethal doses on Female longevity (Days)

Female longevity (Days)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	7.33	23.89
T ₂ (LC ₁₅)	7.75	19.53
T ₃ (LC ₁₀)	7.76	19.42
T ₄ (Control)	9.64	0
C.D.	0.513	
SE(m)	0.155	

Fecundity**Fipronil**

Effect of lethal and sub lethal doses of fipronil on fecundity presented in Table 4.27. The adult fecundity significantly reduced to 83.38 egg/ female in T₁ (LC₅₀), 93.33 eggs/ female in T₂ (LC₁₅) and 103.75 eggs/ female in T₃ (LC₁₀) when compared to 149.08 eggs/female in T₄ (control).

Table 15: Effect of lethal and Sub-lethal doses on Fecundity (Eggs/female)

Fecundity (Eggs/female)		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	83.38	44.06
T ₂ (LC ₁₅)	93.33	37.39
T ₃ (LC ₁₀)	103.75	30.40
T ₄ (Control)	149.08	0
C.D.	8.008	
SE(m)	2.418	

Effect on growth index**Fipronil**

The studies on growth and development of *E. vittella* revealed the lower growth index in all the treatments (T₁, T₂ and T₃) compared to T₄ (control). The data presented in Table 4.28 shows higher growth index 7.3 in T₄ (control) as compared to and 3.51 in T₁ (LC₅₀), 4.22 in T₂ (LC₁₅) and 4.32 in T₃ (LC₁₀). It was observed that growth index 4.22 in (T₂) and 4.32 in (T₃) were significantly at par with each other.

Table 16: Effect of lethal and Sub-lethal doses on Growth index

Growth index		
Treatment	Fipronil	Increment or decrement over control (%)
T ₁ (LC ₅₀)	3.51	51.91
T ₂ (LC ₁₅)	4.22	42.09
T ₃ (LC ₁₀)	4.32	40.78
T ₄ (Control)	7.3	0
C.D.	0.462	
SE(m)	0.139	

Kale *et al.*, (2002) evaluated the toxicity of fipronil on neonate (0-10-hour-old) larvae of *Earias vittella* using fruit-dip bioassays. They reported LC₅₀ value of 0.115 ppm for fipronil at 96 hours post-treatment. It differs from, our study determined LC₅₀ value of 4.844 ppm at 24 hours post-treatment for second-instar larvae. The higher LC₅₀ value in our study likely reflects the shorter exposure duration and the use of later-stage larvae, as neonates exhibit greater susceptibility to insecticides. Generally, mortality values decrease with prolonged exposure or when later larval instars are tested, due to cumulative toxic effects and reduced sensitivity in older larvae. Saleem *et al.*, (2024) reported the LC₅₀ values of fipronil against third instar *Spodoptera frugiperda* as 111.8 ppm, 50.75 ppm, and 20.5

ppm after 24, 48, and 72 hours of exposure, respectively. The relatively high LC₅₀ values may be attributed to the comparatively greater resistance of *S. frugiperda* to fipronil than that observed in *Earias vittella*.

References

1. Abbot CG. The solar constant and terrestrial magnetism. Nature. 1925;116(2926):785-785.
2. Aziz MA, Hasan M, Ali A. Impact of abiotic factors on incidence of fruit and shoot damage of spotted bollworms *Earias* spp. on okra (*Abelmoschus esculentus* L). Pakistan J Zool. 2011;43:863-868.
3. Robinson DS. Food biochemistry and nutritional value. New York: Longman Scientific and Technical Publisher; 1990.
4. Nada MS, Singh J, Kadian RS. Population dynamics of major insect pests in cotton and their natural enemies. Ann Plant Prot Sci. 2010;18(2):470-473.
5. Kumar K, Chapman RB. Sublethal effects of insecticides on the diamondback moth *Plutella xylostella* (L.). Pestic Sci. 1984;15(4):344-352.
6. Rahman MM, Islam MN, Ahmed KS, Khalequzzaman M. Incidence and damage assessment of shoot and fruit borer (*Earias vittella*) on okra. Bangladesh J Agric Res. 2013;38(3):491-499.
7. Shah RA, Zaki FA, Wani SH. Seasonal incidence and damage of shoot and fruit borer *Earias vittella* (Fab.) on okra. Appl Biol Res. 2001;3(1-2):76-78.
8. Kataria SK, Singh G. Efficacy of biorational insecticides against spotted bollworm *Earias* spp. in okra. Indian J Entomol. 2021;1-3. doi:10.55446/IJE.2021.84.
9. Sammelson RE, Caboni P, Durkin KA, Casida JE. GABA receptor antagonists and insecticides: common structural features of 4-alkyl-1-phenylpyrazoles and 4-alkyl-1-phenyltrioxabicyclooctanes. Bioorg Med Chem. 2004;12(12):3345-3355.
10. Bobé A, Coste CM, Cooper JF. Factors influencing the adsorption of fipronil on soils. J Agric Food Chem. 1997;45(12):4861-4865.
11. Gupta GP, Rani S, Birah A, Raghuraman M. Mass rearing of the spotted bollworm, *Earias vittella* (Lepidoptera: Noctuidae) on an artificial diet. Int J Trop Insect Sci. 2005;25:134-137.
12. Gulzar A, Ali MM, Tariq M, Bodlah I, Tariq K, Ali A. Lethal and sublethal effects of *Azadirachta indica* seed extract on the development of spotted bollworm *Earias vittella* (Fab.). Gesunde Pflanzen. 2019;71(1):19-24.
13. Finney DJ. Bioassay and the practice of statistical inference. Int Stat Rev. 1979;47(1):1-12.
14. Gupta BD. Differentiation of sex in pupae of spotted bollworm, *Earias fabia* (Stoll) (Lepidoptera: Noctuidae-Erastrinae). 1978.
15. Kale VD, Deth MG, Kale VD. Studies on thiamethoxam, fipronil, abamectin and spinosad against pests of okra. Rahuri: MPKV; 2002.
16. Saleem U, Asrar M, Farhat Jabeen SM. Assessment of selective synthetic insecticides against third instar larvae of *Spodoptera frugiperda* (Lepidoptera: Noctuidae) under laboratory conditions. Pakistan J Zool. 2024;56(1):1-10.