



# Potential Use of Predatory Bug *Eocanthecona furcellata* for Biological Control of *Plutella xylostella* in Chinese Kale Production

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**Abstract** Predatory bug *Eocanthecona furcellata* was investigated for the potential of biological control agent to *Plutella xylostella* in the laboratory and greenhouse condition. Predation rate of 3<sup>rd</sup> to 4<sup>th</sup> instar nymphs *E. furcellata* on 2<sup>nd</sup>, 3<sup>rd</sup> and 4<sup>th</sup> instar larvae *P. xylostella* were studied in the laboratory. The results showed that predation rate of 4<sup>th</sup> and 5<sup>th</sup> nymphal instars of *E. furcellata* were higher than 3<sup>rd</sup> instar. Feeding rate were greater when they feed on 2<sup>nd</sup> and 3<sup>rd</sup> instar larvae than feed on 4<sup>th</sup> instar larvae of *P. xylostella*. The efficiency of *E. furcellata* in controlling *P. xylostella* under greenhouse conditions was conducted. When *P. xylostella* population reached to economic threshold (ET), 20 of *E. furcellata* were released to chinese kale greenhouse (18 m<sup>3</sup>), Results showed that pest population was reduced 10.32% after 5 days of release when compared to control. Additionally, percentage of plant damage from released predatory bug greenhouse was lower than non-released greenhouse. Therefore, it is possible to release the predatory bug *E. furcellata* in vegetable production as a biological control agent.

**Keywords** predation, biological control, diamondback moth

## INTRODUCTION

*Plutella xylostella* (L.) is an important pest to Brassicaceae including Chinese kale (*Brassica oleracea* var. *alboglabra*). The first instar larvae feed on mesophyll tissues inside the leaf, whereas, other instar larvae are surface feeders. They consume leaves, buds, flowers, siliques, and the green outer layer of stems (Sarfracz et al., 2005). The damage varies, depending on plant growth stage, plant varieties, larval densities and size. If larvae are numerous, they can consume the entire leaf except the veins. *P. xylostella* has been a serious pest of vegetable crops for many decades. The annual cost for its global control is estimated at US\$ 4-5 billion per year (Zalucki et al., 2012). The short generation time, high fecundity and broad host usage within Brassicaceae host plant of *P.*

*xylostella* are the important factors for insecticide resistance for example chlorantraniliprole, Spinosad, chlorfenapyr and lufenuron, an insect growth regulator (IGR) (Arruda et al., 2020).

A method to delay the evolution of insecticide resistance is to integrate the use of control methods such as chemical and biological controls (Mccord and Yu, 1987). *Eocanthecona furcellata* Wolff (Hemiptera: Pentatomidae) is a common predatory stink bug in Southeast Asia (Tuan et al., 2016). It has been mass-reared and used as a biological control agent in Thailand (Suasa-ard, 2010). It has been reported for controlling many caterpillars worm such as *Mythimna separata*, *Helicoverpa armigera*, and *Heliothis assulta* etc. Predation of *E. furcellata* gradually increased from the 2<sup>nd</sup> instar to 5<sup>th</sup> instar nymph when fed on *Maruca vitrata* (Pillai and Agnihotri, 2013) and *Spodoptera litura* (Tuan et al., 2016). Even though, there are some studies of feeding efficiency and utilization of *E. furcellata* for many insect pests, there are not many studies of *E. furcellata* fed on *P. xylostella*. Tuan et al. (2016) revealed predation rate of each stage and life table of *E. furcellata* when fed on 4<sup>th</sup> larval instar but not all stage. Therefore, this study revealed the predation rate for 3<sup>rd</sup> -5<sup>th</sup> nymphal stages of *E. furcellata* when fed on 2<sup>nd</sup>-4<sup>th</sup> larval stages of *P. xylostella* for finding suitable stages for biological control. Additionally, *E. furcellata* was evaluated for its efficiency as a biological control agent in the greenhouse.

## OBJECTIVE

The objectives of this study were to examine predation rate of the *E. furcellata* in predation life stages and to evaluate its efficiency as a biological control agent in the chinese kale greenhouse.

## METHODOLOGY

### Insect Rearing

*Plutella xylostella* was collected from chinese kale production area in Khon Kaen Province, Thailand. Thirty larvae were transferred in an aluminum cage (90x 60x 60 cm) with 15 days old chinese kale in a pot inside the cage. The larvae were maintained in the cage until they became to pupae. Pupae were transferred to the mating cage (10 cm in diameter and 10 cm in height). Newly emerged adults were provided with cotton balls soaked with 20% honey water as food source. Shredded of chinese kale leaves was used as the substrate for oviposition by the diamondback moth and the F3 was then used in the experiment. *E. furcellata* were taken from a colony (10<sup>th</sup> generation) maintained at NBCRC, Upper Northeastern Regional Center. The experimental insect was maintained in the laboratory at room temperature, 28 ± 5°C, relative humidity of 60 ± 10%.

### Bioassay

**Laboratory:** The experiment was examined in the laboratory using Completely Randomized Design (CRD) with 10 replications. A newly hatched of 3<sup>rd</sup>, 4<sup>th</sup> and 5<sup>th</sup> nymphs of *E. furcellata* were test for predation rate. One of each nymph was transferred into a container (7x 9x 4 cm) with cotton balls soaked with water, and a chinese kale leaf. Second larval instar of *P. xylostella* (n = 20) were provided daily as a food until predator change to a new nymphal stage. Then, the experiments were repeated, 3<sup>rd</sup> and 4<sup>th</sup> instar larvae of *P. xylostella* were provided instead as a pray. Predation rate were recorded every day until *E. furcellata* change to new nymphal stage. The data were statistically analyzed using Statistix 10 (Analytical software, 2013) and treatments were compared using Tukey's HSD test (P < 0.05).

**Greenhouse:** The experiment was examined in the 8 chinese kale greenhouses (18 m<sup>3</sup>) at NBCRC. Forty pots of chinese kale were use as the experimental units for each greenhouse. The population of *P. xylostella* was artificial infestation by releasing a cluster of eggs/pot when plants were 30 days old. After eggs hatched to the larvae, the population were recorded. When the population reached to the economic threshold level (ET) (1 larva/plant), Twenty of *E. furcellata* were released into the

greenhouse. While, control greenhouse was not released. The experiment was studied with 4 replications. The data were statistically analyzed using Paired sample t-test using Statistix 10 (Analytical software, 2013). Percentage of leaf area damage was recorded in level of injury (0=no damage, 1 = < 25% damage, 2 = 25-50% damage, 3 = 50-75% damage, 4 = >75% damage). The percent reduction of *P. xylostella* was statistically calculated according to the equation of Henderson and Tilton (1955).

$$\% \text{ reduction} = 100 \times (1 - (Ta \times Cb)/(Tb \times Ca)) \quad (1)$$

Where, Ta = population of insect counts after treatment, Cb = population of untreated insect count before treatment, Tb = population of insect counts before treatment, and Ca = population of untreated insect count after treatment.

## RESULTS AND DISCUSSION

Prey consumption of *E. furcellata* fed on larvae of *P. xylostella* was significantly different. The third nymphal instar of *E. furcellata* fed on 3<sup>rd</sup> and 4<sup>th</sup> instar larvae more than 2<sup>nd</sup> instar larvae, whereas, 4<sup>th</sup> and 5<sup>th</sup> nymphal instar of *E. furcellata* fed on younger more than the older prey (Table 1). Predation rate of *E. furcellata* may involve with a prey species. Tiwari et al. (2017) revealed that when *E. furcellata* fed on *S. litura*, and *M. vitrata*, predation rate of prey consumed slightly increased with increasing prey stage. However, when *E. furcellata* fed on *Spilarctia obliqua* and *S. frugiperda* predation rate of prey consume decreased with increasing prey size (Kumar et al., 2001; Keerthi et al., 2020). Rani and Wakamura 1993 suggested that physical stimuli such as host shape or movement had no influence on acceptability of the prey to *E. furcellata*.

**Table 1 Prey consumption of different stages of *Eocanthecona furcellata* fed on larvae of *Plutella xylostella***

Treatment <i>E. furcellata</i>	<i>P. xylostella</i> <sup>1/</sup>		
	2 <sup>nd</sup> instar	3 <sup>rd</sup> instar	4 <sup>th</sup> instar
3 <sup>rd</sup> nymph	28±0.50 B	65±4.50 A	50±1.91 A
4 <sup>th</sup> nymph	71±5.91 A	79±4.72 A	24±0.82 B
5 <sup>th</sup> nymph	71±2.65 A	60±5.35 AB	38±2.08 B
F-test	**	**	*
CV (%)	24.33	30.28	26.08

<sup>1/</sup>Within each column, mean±SD followed by the same capital letter indicate no significantly different ( $P>0.05$ )

However, Kumar et al. (2001) revealed that visualization of the predator and movement of prey increases the predation rate. In this case, the active movement of 2<sup>nd</sup> and 3<sup>rd</sup> instar larvae of *P. xylostella* may be a factor of high predation rate of 4<sup>th</sup> and 5<sup>th</sup> nymphal stage of predator. Contrast to 4<sup>th</sup> and 5<sup>th</sup> nymphal stage, 3<sup>rd</sup> nymphal stage may difficult to handle the active prey movement.

The results of laboratory experiment for predation of *E. furcellata* lead to a decision to release 3<sup>rd</sup> nymphal stage of *E. furcellata* as a biological control in chinese kale greenhouse. Twenty of 3<sup>rd</sup> nymphal stage in a bio-product were released into the greenhouse after diamond back moth reach to 1 larva/plant. The results revealed that the number of *P. xylostella* was not significant different between treatment and control (Table 2). However, population of *P. xylostella* was reduced at 3 days after *E. furcellata* released. The percent reduction of *P. xylostella* at 3-5 day after predator released were 10.32, 21.79 and 18.06%, respectively. During 1-2 days after predator released the population in treatment was not reduced. At that period, most of *P. xylostella* was 2<sup>nd</sup> instar which was not suitable for the predator stage. Two day later, most of pest population was 3<sup>rd</sup> instar, and predator was changing nymphal stage from 3<sup>rd</sup> to 4<sup>th</sup> nymph. Then the population was reduced (Percent reduction was higher). Therefore, in case of a serious outbreak, 4<sup>th</sup> nymphal stage may use

instead. Level of chinese kale leaf damage in control greenhouse increased from level 1 to level 3, whereas, level of chinese kale leaf damage in control greenhouse was level 1 and level 2.

However, applying in open field condition may have some factors such as temperature, various pests, other natural enemies and chemical insecticide which involving the efficiency of predator.

**Table 2 Number of *Plutella xylostella* in chinese kale greenhouse and level of leave damage with released and non-released *Eocanthecona furcellata* as a biological control agent, and percentage reduction.**

Plant age (day)	non-released		released		Reduction (%)
	Insect/plant	Level of leave damage	Insect/plant	Level of leave damage	
36 <sup>2/</sup>	2.32±1.15a <sup>1/</sup>	1	1.44±0.15a	1	na
37	2.21±1.12a	1	1.42±0.14a	1	-3.52
38	2.17±1.09a	1	1.40±0.15a	1	-3.94
39	2.12±1.04a	2	1.18±0.18a	1	10.32
40	2.06±1.00a	2	1.00±0.26a	1	21.79
41	1.75±1.00a	3	0.89±0.24a	2	18.06

<sup>1/</sup>Within each row, mean±SD followed by the same small letter indicate no significantly different ( $P>0.05$ )

<sup>2/</sup> *Eocanthecona furcellata* was released into the greenhouse.

## CONCLUSION

*E. furcellata* is an effective predator of *P. xylostella*, the third nymphal stage consumed 3<sup>rd</sup> and 4<sup>th</sup> larval instar than 2<sup>nd</sup> instar. In contrast, 4<sup>th</sup> and 5<sup>th</sup> nymphal stage consumed 2<sup>nd</sup> and 3<sup>rd</sup> instar more than 4<sup>th</sup> instar. *E. furcellata* is an alternative biological control agent in chinese kale green house. It can reduce pest population and level of leaf damage.

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