

# Resistance Potential of *Chrysoperla carnea* (Stephens) to Insecticides Used Against Sucking Complex of Cotton

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**Abstract:** The reported high loss mortality rate of green lacewing, (*Chrysoperla carnea*) have been attributed to diverse factors including unattended use of insecticides. Since chemical control is one of a significant practice to manage insect pest in cotton. However, this kind of practice may impair the natural control provided by generalist predator *C. carnea*. Although, natural control adoption is limited in crops, area and season due to wide-spread use of insecticides but presence of resistance potential in *C. carnea* may improve the design of solid IPM strategies. Herein, we aimed to assess the toxicity of four insecticides to two strains of *C. carnea* (viz. laboratory reared and field collected adults) and to evaluate their resistance potential by calculating their resistance ratio. LC<sub>50</sub> was calculated at 24 h following topical application administered when the adults were 3 days old. Control adult mortalities were less than 10% at 24 h. The LC<sub>50</sub> values (μl mL<sup>-1</sup>) for laboratory reared strains of each tested insecticide were: acetamiprid, 0.0064; bifenthrin, 3.75; chlorpyrifos, 0.067; and profenofos, 0.052. The LC<sub>50</sub> values for field collected strains were 0.096 (acetamiprid), 34.8 (bifenthrin), 0.21 (chlorpyrifos) and 0.44 (profenofos). The toxicity of the test insecticide to *C. carnea* from more to least toxic was acetamiprid > profenofos > chlorpyrifos > bifenthrin. Field collected strain possessed 15 (acetamiprid)-, 9.28 (bifenthrin)-, 3.13 (chlorpyrifos)-, and 8.5 (profenofos)-fold more resistance than the susceptible population. These results are pretty worthwhile for integration of *C. carnea* in IPM programs, impairing with insecticides.

**Keywords:** Green Lacewing, *Chrysoperla*, Resistance Potential, Insecticides, Toxicity

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## 1. Introduction

Conservation natural control, in which beneficial fauna are preserved in the agroecosystem, has been considered as an element progressively important in Integrated Pest Management (IPM) programs [1]. It is the only way to maintain and enhance the survival, reproduction, and efficiency of natural enemies, which are directly involved in regulating populations of various agricultural and forest insect pests. Approaches to the conservation of these natural enemies involve the adoption of practices that benefit them, as well as avoidance of practices that could be harmful [2].

Among natural enemies in Asia, a large number of predators can control insect pest in cotton (*Gossypium hirsutum* L.) such as Coccinellids, Chrysopids, Anthocoris and Spiders [3]. The green lacewing *Chrysoperla carnea* (Stephens) (Neuroptera: Chrysopidae) stand out as important predator, especially of the sucking complex of cotton [4]. The larvae of *C. carnea* are generalist predator which can also feed on eggs and small larvae of lepidoptera, scales, aphids, psilids and whiteflies [5]. It is worth mentioning that the use of chrysopids in IPM as increased since last few years,

mainly due to its adaptability, voracity and relative wide tolerance to several insecticides [6, 7].

Regardless of advances in insect management technologies, use of insecticides is still a common practice worldwide for the control of various insect pest populations [8]. However, insecticides have been also reported with numerous demerits, such as the emergence of secondary pest populations, resurgence of various pests, producing resistant pest populations, and lethal effects on the natural enemies cohabiting in agroecosystem [7, 9]. Based on the importance of *C. carnea* eggs and pupae, tolerance to insecticides may play a vital role to eliminate the pests which are usually escaped from chemical control [1]. One of the main agenda of IPM is the combination of selective insecticides with natural enemies. Therefore, the evaluation of insecticidal effects on biological control agents is essential prior to the execution of and IPM program [10, 11]. Theoretically, few studies have been addressed that all developmental stages (eggs, larva, pupa, and adult) of green lacewing are susceptible to insecticides, the exposure may lead to lethal and sub-lethal effects [7, 12]. Most of the studies highlighted lethal and sub-lethal effects of selective insecticides on larvae and adult lacewing [13-15], and some have focused on eggs and pupae that can be considered more tolerant to insecticides [6, 16, 17].

Insecticide resistance in insect pest has adverse consequences but can be used as a positive attribute for biological control agents as a valuable tool in pest management [18]. Most studies on insecticide resistance in biological control agents try to establish the degree of compatibility using only a population, without considering the natural variation in susceptibility of insecticides. However, variation in response to insecticides among the populations of natural enemies is similar to the response in any other insect pest [19]. So, the knowledge of the potential resistance of natural enemies to insecticides may improve strong IPM approaches.

In the present study, the toxicity of insecticides used in cotton cultivation, commonly against sucking complex were investigated on laboratory reared and field collected strain of *C. carnea*. The information generated through the present study would be a better understanding of insecticide resistance in natural enemies that will guide us to improve the integration of chemical and biological tools in IPM programs.

## 2. Materials and Methods

### 2.1. Insect Collection

*C. carnea* adults (about 2700) were collected from different fields (cotton, brassica etc) of Tandojam (location: 25.4203°N, 68.5445°E) in the year 2017 by using specialized LED-based light traps which can easily target lacewing trichromatic vision [20]. Each trap was equipped with four strips (each strip contain 12 LEDs) able to emit UV the first (250 mcd and 395 nm wavelength) and flashing white light

second (11,000 mcd and 455 nm wavelength). Traps were provided with a 12V battery for 7Ah and left the switch on from 7 pm to 10 pm. Capturing was also done by hand sweeping net around the traps.

### 2.2. Insect Rearing

Collected adults (field strain) and laboratory reared (G-273) strains were kept in rearing cage (24.5 x 24.5 x 24.5 cm) provisioned with an artificial diet consisting sugar, honey, yeast and distilled water (2:1:1:2) in the laboratory [21]. Black linen cloth was provided at the top of cages for egg laying and removed on each alternate day to harvest the eggs. The eggs were placed in black cloth (cover cloth) and fed on frozen Angoumois grain moth (*Sitotroga cerealella* Olivier) eggs [9]. After hatching (48 - 60 hrs), individual 1<sup>st</sup> instar larvae were sealed in 2 inches polypropylene transparent straw with three pin aeration holes to avoid the cannibalism. The culture was maintained in the Biological Control Laboratory, Nuclear Institute of Agriculture at 25 ± 1°C and 60 ± 5% RH and 15:9 (L:D) till pupation. Tubes were cut out and small open pieces of the tube along with pupae and placed on glass petri plates (9 cm diameter and 1.5 cm height) for emergence. The population of *C. carnea* reared for 10 years in the mentioned laboratory without exposure, was designated as Susceptible strain [22, 23].

### 2.3. Insecticides Formulation

Commercial insecticides were used for bioassays. The four insecticides (Acetamiprid: Acelan<sup>®</sup> 20 SL, FMC, expiry date March 2019; Chlorpyrifos: Cordelia<sup>®</sup> 40 EC, FMC, expiry date July 2018; Bifenthrin: Talstar<sup>®</sup> 10 EC, FMC, expiry date March 2019; and Profenofos: Curacron<sup>®</sup> 50 EC, Syngenta, expiry date November 2018) used under study were purchased their registered dealers. Firstly determined the correction factor (CF) to have a 100% stock solution. Pilot studies were conducted to determine the suitable concentration range to use for each insecticide. On these basis, following concentrations were evaluated to determine the LC<sub>50</sub> for susceptible strain: Acetamiprid: 0.001, 0.002, 0.005, 0.01, 0.02, 0.05 and 0.1 µl mL<sup>-1</sup>; Bifenthrin: 0.9, 1.2, 1.5, 2, 5 and 10 µl mL<sup>-1</sup>; Chlorpyrifos: 0.0075, 0.01, 0.02, 0.05, 0.1, 0.15, 0.2 and 0.5 µl mL<sup>-1</sup>; and Profenofos: 0.0075, 0.01, 0.02, 0.05, 0.07, 0.08, 0.1 and 0.2 µl mL<sup>-1</sup>. The following concentrations were used to determine LC<sub>50</sub> for field strain: Acetamiprid: 0.001, 0.002, 0.005, 0.01, 0.05, 0.1, 0.2 and 0.4 µl mL<sup>-1</sup>; Bifenthrin: 5, 10, 15, 30, 50, 60 and 70 µl mL<sup>-1</sup>; Chlorpyrifos: 0.01, 0.02, 0.03, 0.06, 0.12, 0.24, 0.48 and 0.96 µl mL<sup>-1</sup>; and Profenofos: 0.01, 0.03, 0.05, 0.1, 0.25, 0.5, 1 and 1.5 µl mL<sup>-1</sup>. All insecticides were dissolved in acetone to prepare stock solution.

### 2.4. Concentration Response Bioassay

Concentration response bioassays of insecticides were conducted for 48 – 60 hours old *C. carnea* adults as described previously [21]. All above mentioned

concentrations were made as serial dilutions, each concentration was replicated three times for each bioassay. 1 µl of each serial concentration was applied on the thorax of each individual adult with Arnold Micro-applicator (type LV.65. Burkard, UK) as described by Mansoor et al., [21]. Each replication contained thirty adults, and a total of ninety adults was exposed to each concentration of insecticide. Treated adults were provided with artificial diet as described earlier. Mortality was assessed 24 h after exposure to insecticides. Mortality was determined by viewing the adult's movement and activity of antennae by touching.

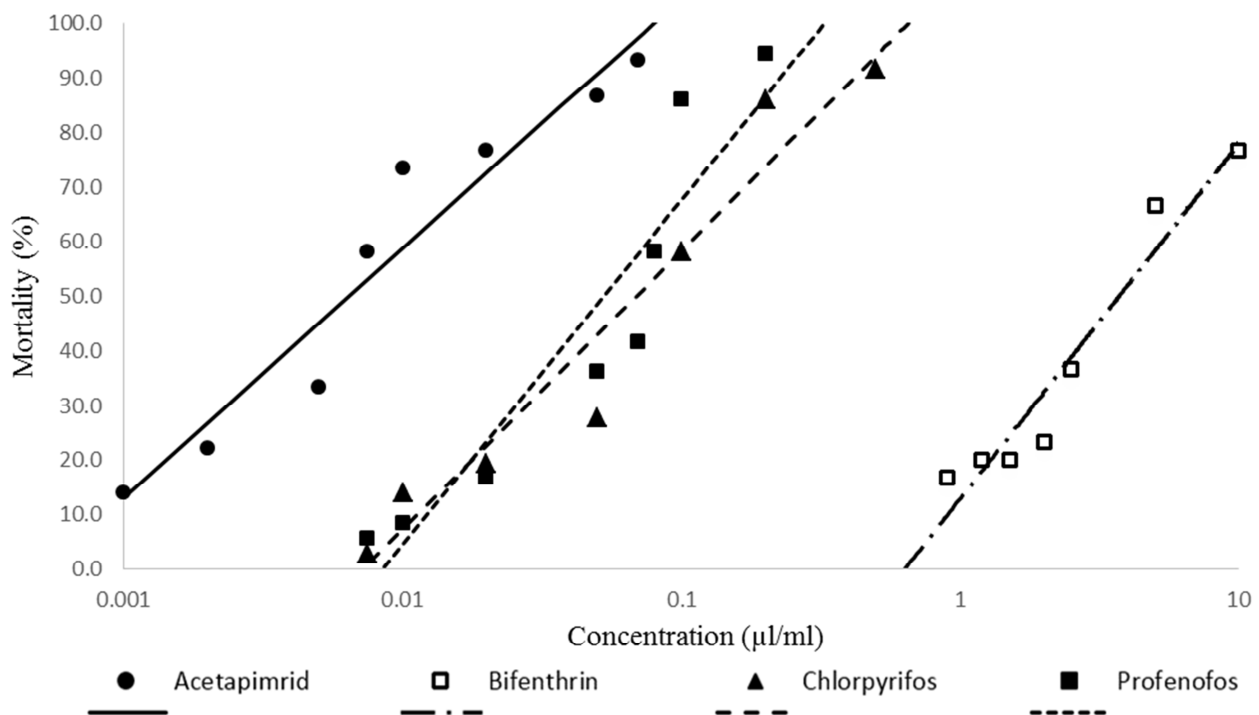
### 2.5. Statistical Analysis

Mortality ratio of *C. carnea* was corrected using solvent control mortality via the Abbott formula. Concentration-response data were analyzed with analytical software POLO [24]. The log concentration-response curves allowed determination of LC<sub>50</sub>, LC<sub>90</sub>, Chi-square ( $\chi^2$ ) and slopes  $\pm$  SE values for the adult Bioassay according to probit analysis [25]. The 95% confidence limits for the range of LC<sub>50</sub> values

were calculated by least-regression analyses against the logarithm of insecticide concentration. The lethal concentration values were considered similar if their 95% confidence limits overlapped [26].

## 3. Results

Solvent control mortality was lower than 10% at 24 h after treatment. The toxicity of insecticides tested are reported in Table 1 and 2 and have shown graphically in Figure 1 and 2. The LC<sub>50</sub> values of acetamiprid, bifenthrin, chlorpyrifos, and profenofos for laboratory reared susceptible strain of *C. carnea* were 0.0064, 3.75, 0.067 and 0.052 µl mL<sup>-1</sup>, and LC<sub>50</sub> values for field collected strains were 0.096, 34.8, 0.21 and 0.44, respectively. Toxicity of the test insecticides to *C. carnea* adults form most to least toxic was acetamiprid > profenofos > chlorpyrifos > bifenthrin. The 95% confidence intervals for chlorpyrifos and profenofos were overlapped, suggesting no significant difference in the toxicities of these two insecticides to *C. carnea* adults.



**Figure 1.** The mortality response of susceptible (G-273) strain of *C. carnea* after exposure to Acetamiprid, Bifenthrin, Chlorpyrifos, and Profenofos. The concentration response line of each population was drawn using a probit linear model  $y = ax + \beta$  in which  $a$  and  $\beta$  are the slope and intercept, respectively.  $x$  is the log-transformation concentration (µl/ml) and  $y$  is the percent mortality.

**Table 1.** Toxicity of insecticides to the susceptible strain (G-273) of *C. carnea*.

Insecticides	LC <sub>50</sub> (95% CI) [µl mL <sup>-1</sup> ]	LC <sub>90</sub> (95% CI) [µl mL <sup>-1</sup> ]	The fit of probit line			
			Slope ( $\pm$ SE)	df	N <sup>a</sup>	$\chi^2$
Acetamiprid	0.0064 (0.004 $\pm$ 0.011)	0.059 (0.035 $\pm$ 0.1)	1.34 ( $\pm$ 0.12)	6	750	0.98
Bifenthrin	3.75 (2.49 $\pm$ 5.64)	19.39 (12.88 $\pm$ 29.62)	1.81 ( $\pm$ 0.09)	5	660	0.97
Chlorpyrifos	0.067 (0.04 $\pm$ 0.11)	0.38 (0.25 $\pm$ 0.59)	1.75 ( $\pm$ 0.1)	6	750	0.78
Profenofos	0.052 (0.04 $\pm$ 0.07)	0.22 (0.15 $\pm$ 0.31)	2.14 ( $\pm$ 0.08)	6	750	0.87

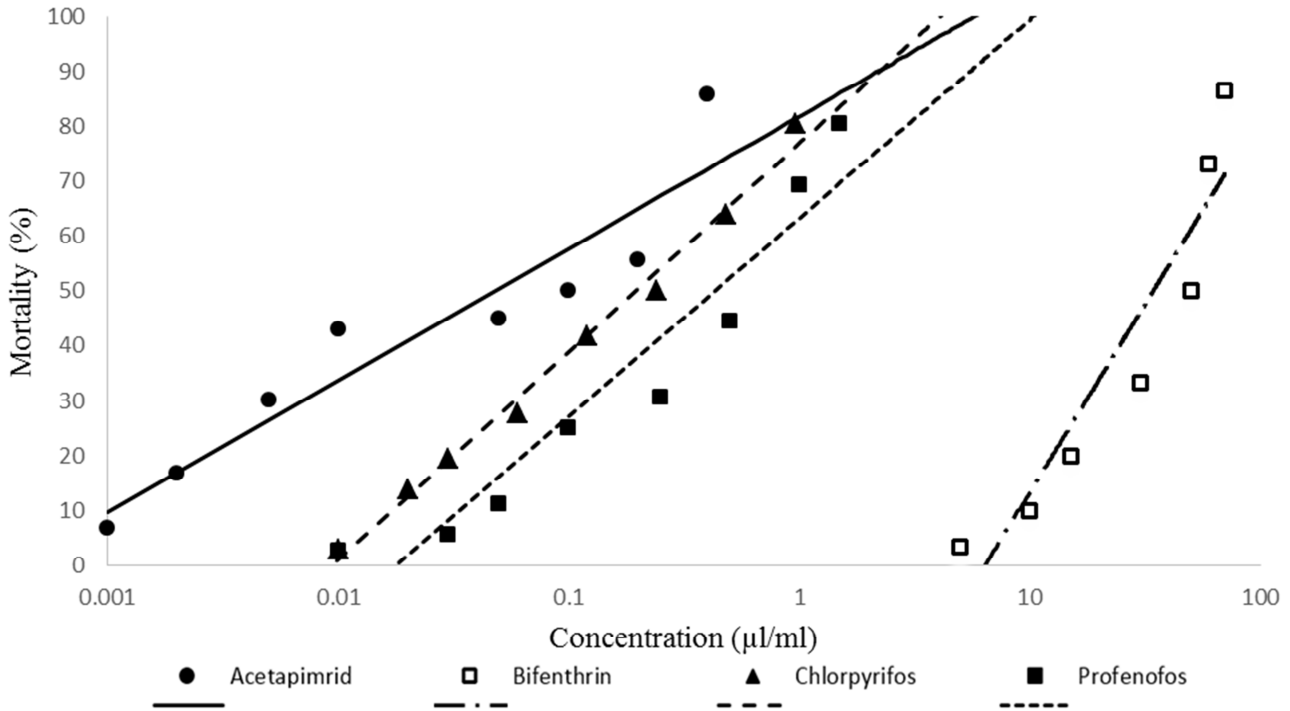
<sup>a</sup> Number of *C. carnea* individuals used in bioassay and control.

**Table 2.** Toxicity of insecticides to the field collected strain of *C. carnea*.

Insecticides	LC <sub>50</sub> (95% CI)[ $\mu\text{l mL}^{-1}$ ]	LC <sub>90</sub> (95% CI)[ $\mu\text{l mL}^{-1}$ ]	The fit of probit line				
			Slope ( $\pm$ SE)	df	N <sup>a</sup>	$\chi^2$	RR <sup>b</sup>
Acetamiprid	0.096 (0.04 $\pm$ 0.31)	18.4 (5.72 $\pm$ 59.31)	0.58 ( $\pm$ 0.3)	6	510	0.79	15
Bifenthrin	34.8 (25.4 $\pm$ 49.9)	130.1 (92.7 $\pm$ 182.2)	2.32 ( $\pm$ 0.08)	5	450	0.96	9.28
Chlorpyrifos	0.21 (0.11 $\pm$ 0.36)	2.28 (1.27 $\pm$ 4.09)	1.24 ( $\pm$ 0.1)	6	510	0.97	3.13
Profenofos	0.44 (0.24 $\pm$ 0.81)	4.79 (2.64 $\pm$ 8.68)	1.25 ( $\pm$ 0.1)	6	510	0.98	8.5

<sup>a</sup> Number of *C. carnea* individuals used in bioassay and control.

<sup>b</sup> Resistance ratio, LC<sub>50</sub> of the field strains / LC<sub>50</sub> of susceptible (G-273) strains



**Figure 2.** The mortality response of field collected strains of *C. carnea* after exposure to Acetamiprid, Bifenthrin, Chlorpyrifos, and Profenofos. The concentration response line of each population was drawn using a probit linear model  $y = ax + \beta$  in which  $a$  and  $\beta$  are the slope and intercept, respectively.  $x$  is the log-transformation concentration ( $\mu\text{l/ml}$ ) and  $y$  is the percent mortality.

## 4. Discussion

The role of a generalist predator, *C. carnea* is obvious and well known, however, their presence in the field is dependent upon the lack of disruption due to different insecticides [21]. Therefore, there is a need to study the evolution of insecticides resistance in *C. carnea*, which are regularly used for the management of various insect pests. The *in-vitro* studies demonstrated its worth for evaluating the toxicity of insecticides on adults *C. carnea* by providing quantitative data with high accuracy and reproducibility and avoiding environmental variation [27]. The techniques discussed herein demonstrated the effectiveness of a reliable acute toxicity assay for laboratory reared adults and their difference to the field collected strain. The LC<sub>50</sub> values of acetamiprid, bifenthrin, chlorpyrifos, and profenofos for laboratory reared susceptible strain and field collected strains of *C. carnea* were evaluated.

Maximum toxicity [(LD<sub>50</sub> = 0.0064 (0.004  $\pm$  0.011)] was observed in the case of acetamiprid, belong to a new, widely used class of insecticide, the neonicotinoids. With identical

structure to nicotine, they also share agonist activity at ionotropic or nicotinic acetylcholine receptors (nAChRs) [28, 29]. Its toxicity against many insect fauna related to cotton and ornamental plants are well established [30]. Its resistance has previously been reported in *C. carnea* at different locations (Muzaffargarh; 30.0703° N, 71.1933° E) in Pakistan.

The toxicity test in the present study evidently proved the toxicity of organophosphate insecticides (i.e. profenofos and chlorpyrifos) to the *C. carnea* adults. Non-significant difference was found in their toxicity to *C. carnea*. Percent mortality was increased with increase in the concentration of these insecticides. Like other organophosphate insecticides, chlorpyrifos and profenofos kill targeted insects by inhibiting the action of acetylcholinesterase. It is usually responsible for degradation of excitatory neurotransmitter, acetylcholine, thereby termination of nerve impulse transmission at cholinergic synapses [31], which is critical to the functioning of the insect nervous system [29, 32]. Once this enzyme is inhibited, acetylcholine builds up and insect expire from overstimulation of their nervous system [33].

However, among the test insecticides, bifenthrin showed the least toxicity [LC<sub>50</sub> = 3.75 (2.49  $\pm$  5.64)] to *C. carnea*

adults. It is also an insecticide that has been using frequently by the cotton growers of Pakistan against insect pests, also affects natural enemies [34]. The basic mechanism of bifenthrin involves binding to voltage-gated sodium channels, important sites for neurotoxic action and modifying their kinetics, causing disrupt the normal functioning of nerves [35, 36].

The results of our studies, where we applied insecticides directly onto the field collected strain of *C. carnea* suggested the resistance potential is present in their adults. Resistance ratio depicting their folds of resistance to particular insecticide. Field collected population has 15 (acetamiprid)-, 9.28 (bifenthrin)-, 3.13 (chlorpyrifos)-, and 8.5 (profenofos)-fold more resistance than the susceptible population. Although, resistance in insect pest outnumber resistance in the natural enemies especially the predators was more than twenty to one. This attribute most probably indicates limited devotions to resistance in natural enemies as well as biological difference among natural enemies and pests [18]. It is well documented that the natural enemies develop resistance less readily as detoxification enzymes level is lower the predators than in pests or because they suffered due to food limitation insecticide that severely lessens their host or prey [37].

Insecticide resistance mainly credited due to either or all of these mechanisms viz. metabolic, decrease penetration, target site insensitivity and behavioral resistance [38]. Decreased penetration and target site insensitivity are less common and contribute little for resistance in many cases [39]. Enhanced metabolic resistance includes esterases, cytochrome P450 mono-oxygenase and glutathione enzyme complex [38]. In the present study, resistance to the bifenthrin, suggesting the involvement of esterases and mono-oxygenases in the *C. carnea* [21, 40]. Previously resistance to pyrethroids mediated by mono-oxygenases has been reported in *C. carnea* [21, 41]. While, organophosphate are attacked by about three enzyme systems, the polysubstrate monooxygenases (PSMOs), carboxylesterases with phosphatase activity and glutathione transferases [31]. The latter two invariably detoxify the organophosphate by splitting off alkyl or other substituents. However, the performance of cytochrome P450 can result in activation of phosphorothioates if the P=S is changed to P=O by oxidative desulfuration, or in detoxification, the carbon of one of the small alkyl substituents is oxidized [42, 43]. Likewise, P450 mediated detoxification probably acts a substantial part in neonicotinoid (imidachloprid) resistance in most of the insects but there would be some secondary mechanism involved [44].

## 5. Conclusion

*C. carnea* population possesses the resistance potential against the test insecticides and could be used as a positive trait for the improvement in their survival opportunities in the field. This potential may lead *C. carnea* to higher predation rates and fitness advantages as Mansoor *et al.*, [21] reported previously. However, the outcome of our studies would be a better

understanding of insecticide resistance in *C. carnea*, ultimately will allow us to improve the integration of biological and chemical tools simultaneously in IPM programs.

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