

Susceptibility of Helicoverpa zea (Lepidoptera: Noctuidae) Neonates to Diamide Insecticides in the Midsouthern and Southeastern United States

Authors: Adams, A., Gore, J., Catchot, A., Musser, F., Cook, D., et al.

Source: Journal of Economic Entomology, 109(5): 2205-2209

Published By: Entomological Society of America

URL: https://doi.org/10.1093/jee/tow175

The BioOne Digital Library (https://bioone.org/) provides worldwide distribution for more than 580 journals and eBooks from BioOne's community of over 150 nonprofit societies, research institutions, and university presses in the biological, ecological, and environmental sciences. The BioOne Digital Library encompasses the flagship aggregation BioOne Complete (https://bioone.org/subscribe), the BioOne Complete Archive (https://bioone.org/archive), and the BioOne eBooks program offerings ESA eBook Collection (https://bioone.org/esa-ebooks) and CSIRO Publishing BioSelect Collection (https://bioone.org/esa-ebooks)

Your use of this PDF, the BioOne Digital Library, and all posted and associated content indicates your acceptance of BioOne's Terms of Use, available at www.bioone.org/terms-of-use.

Usage of BioOne Digital Library content is strictly limited to personal, educational, and non-commmercial use. Commercial inquiries or rights and permissions requests should be directed to the individual publisher as copyright holder.

BioOne is an innovative nonprofit that sees sustainable scholarly publishing as an inherently collaborative enterprise connecting authors, nonprofit publishers, academic institutions, research libraries, and research funders in the common goal of maximizing access to critical research.



Insecticide Resistance and Resistance Management

Susceptibility of *Helicoverpa zea* (Lepidoptera: Noctuidae) Neonates to Diamide Insecticides in the Midsouthern and Southeastern United States

A. Adams, J. Gore, A. Catchot, F. Musser, D. Cook, N. Krishnan, and T. Irby

¹Department of Biochemistry, Molecular Biology, Entomology and Plant Pathology, Mississippi State University, MS 39762 (caa109@msstate.edu; acatchot@entomology.msstate.edu; Fmusser@entomology.msstate.edu; nkrishnan@entomology.msstate.edu),
²Department of Biochemistry, Molecular Biology, Entomology and Plant Pathology, Delta Research and Extension Center (DREC),
Stoneville, MS 38776 (jgore@drec.msstate.edu; dcook@drec.msstate.edu),
³Corresponding author, e-mail: jgore@drec.msstate.edu
and
⁴Department of Plant and Soil Sciences, Mississippi State University, MS 3976 (tirby@pss.msstate.edu)

Received 15 February 2016; Accepted 15 July 2016

Abstract

Corn earworm, *Helicoverpa zea* (Boddie), is a significant pest of agroecosystems in the midsouthern and southeastern regions of the United States. These insects have developed resistance to, or inconsistent control has occurred with, most insecticide classes. With their unique mode of action, insecticides in the diamide class have become a key component in management of agriculturally important lepidopteran pests. In this study, field populations of H. zea were collected in the southern United States and compared to susceptible laboratory colonies to generate baseline concentration–mortality data. LC_{50} and LC_{90} values were generated for flubendiamide and chlorantraniliprole using neonates. To achieve equivalent levels of mortality, a higher concentration of flubendiamide was required compared to chlorantraniliprole. Flubendiamide LC_{50} values for H. zea ranged from 16.45 to 30.74 ng/ml, with a mean of 23.53 ng/ml. Chlorantraniliprole LC_{50} values for H. zea ranged from 2.94 to 4.22 ng/ml, with a mean of 3.66 ng/ml. Significant differences were observed for some field populations relative to the laboratory colony. For flubendiamide, five populations had greater LC_{50} values and two populations had lower LC_{50} values compared to the laboratory colony. For chlorantraniliprole, three populations had greater LC_{50} values and three populations had lower LC_{50} values compared to the laboratory colony. The response of these populations most likely represents natural variability among populations and does not indicate a significant shift in susceptibility of this species.

Key words: corn earworm, chlorantraniliprole, flubendiamide

Lepidoptera is the most damaging insect order in soybean, Glycine max (L.) Merr., production in the southern United States (Fitt 1989, Musser et al. 2015). Corn earworm, Helicoverpa zea (Boddie); soybean looper, Chrysodeixis includens (Walker); beet armyworm, Spodoptera exigua (Hübner); and fall armyworm, Spodoptera frugiperda (Smith) are widely distributed polyphagous pests of numerous cultivated crops throughout the midsouthern and southeastern United States. In 2014, these insects resulted in a combined US\$138,874,796 economic loss across Alabama, Arkansas, Louisiana, Mississippi, North Carolina, Tennessee, and Virginia in soybean (Musser et al. 2015). Foliar applications of synthetic insecticides are instrumental in the management of lepidopteran pests in the southern United States. The widespread use of synthetic insecticides has led to resistance and/ or inconsistent control with most insecticide classes, including chlorinated hydrocarbons, organophosphates, carbamates, pyrethroids, and benzoylphenylureas (Sparks 1981, Brown et al. 1998, Temple et al. 2006, Jacobson et al. 2009, Lai and Su 2011).

The diamide class of insecticides was introduced in 2008 and is the newest major class of insecticides (EPA 2008). This class has a novel mode of action and is classified as ryanodine receptor modulators (MoA Group 28) (IRAC 2014). Ryanodine receptors (RyR) are intracellular calcium channels located in the sarcoplasmic reticulum that specialize in the rapid and massive release of calcium from intracellular stores, which is necessary for excitation contraction coupling in striated muscle (Ebbinghaus-Kintscher et al. 2006). Calcium serves as the primary physiological regulator of insect ryanodine receptors (Xu et al. 2000, Scott-Ward et al. 2001). Diamide insecticides bind to the ryanodine receptor complex, prompting the prolonged release of intracellular calcium stores, resulting in cessation of feeding and uncoordinated muscle contraction of intoxicated insects, eventually causing mortality (Ebbinghaus-Kintscher et al. 2006, Nauen et al. 2007, Hannig et al. 2009, Roditakis et al. 2015). The diamide insecticides are characterized by their low mammalian toxicity and are effective against a large number of lepidopteran species (Sattelle et al. 2008, Tohnishi et al. 2005, Lahm et al. 2009, Teixeira and Andaloro 2013, Qi et al. 2014). Two representatives from this class of insecticides are flubendiamide (Belt, Bayer CropScience, Raleigh, NC), a pthalic acid diamide, and chlorantraniliprole (Prevathon, DuPont Crop Protection, Newark, DE), an anthranilic diamide (Lahm et al. 2009). Although they are structurally independent, these insecticides share the same target site (Lahm et al. 2009, Teixeira and Andaloro 2013). Eight years after their introduction to the global market, these two active ingredients comprise 7% of global insecticide use (Sparks 2013). Large global market shares result from the favorable biological, ecological, and toxicological attributes of this insecticide class (Teixeira and Andaloro 2013). It is perceived that the use of this insecticide class will continue to increase globally on a wide variety of crops (Teixeira and Andaloro 2013, Roditakis et al. 2015).

Repeated field applications of the diamide insecticides has resulted in numerous reports of resistance development for several lepidopteran species (Roditakis et al. 2015). To date, cross resistance between chlorantraniliprole and flubendiamide has been documented for diamondback moth, Plutella xylostella (L), smaller tea tortrix, Adoxophyes honmai (Yasuda), and tomato borer, Tuta absoluta (Meyrick) (Wang and Wu 2012, Uchiyama and Ozawa 2014, Roditakis et al. 2015). Also, resistance to chlorantraniliprole has been documented for rice stem borer, Chilo suppressalis (Walker), cutworm, Spodoptera litura (F), and S. exigua (Su et al. 2012, Che et al. 2013, Gao et al. 2013). Furthermore, resistance to flubendiamide has been documented in C. suppressalis (Wu et al. 2014). Because resistance has occurred among a number of pest species globally, monitoring susceptibility levels of target pest species is important for pest management and resistance management efforts. Furthermore, development of successful insecticide resistance management strategies requires the establishment of baseline susceptibility levels of target pest species while resistant allele frequencies are low (Ffrench-Constant and Roush 1990, Cook et al. 2005). Baseline responses of laboratory and field strains of target pest populations to novel modes of action act as a historical reference, and are necessary to mitigate resistance development, prolonging the effectiveness of novel modes of action. Although the diamide class has been used commercially since 2008, no field control failures have been documented for *H. zea* in the southern United States. The primary objective of the present study was to generate dose-mortality responses of *H. zea* to flubendiamide and chlorantraniliprole after several years of commercial use for future resistance monitoring efforts, development of resistance management strategies, and development of a more rapid assay method.

Materials and Methods

Insects

The H. zea susceptible colony was a laboratory colony maintained at the Mississippi State University Department of Entomology, Mississippi State, MS insect rearing facility. This colony originated from non-Bt corn in 2006 and wild individuals collected from non-Bt corn were incorporated into the colony on a yearly basis. Prior to the initiation of this experiment the susceptible colony was not known to be exposed to insecticides. Field-derived populations for this study were composed of 15 H. zea colonies collected during 2013 and 2014 (Table 1). Each collection consisted of at least 300 third instars. Larvae were placed in 36-ml Solo cups (Bio-Serv, Frenchtown, NJ) containing Stonefly Heliothis Diet (Product No. 38-0600, Ward's Natural Science, Rochester, NY) with matching lids. At pupation, ~50 pupae were placed in 3.79-liter cardboard containers with matching lids with the corresponding colony and generation information labeled on the outside of each bucket. Adults were fed a 10% sugar-water solution that was changed daily. For the purpose of egg collection for bioassays, the cardboard containers were lined with Reynolds Cut-Rite Wax Paper (Reynolds Consumer Products, Lake Forest, IL). The center of each lid was removed so that only the rim remained. Cotton cloth was placed over each bucket and kept in place by the lid to serve as an oviposition substrate. Eggs were collected daily and new cloths and wax paper were applied to every bucket. Collected egg sheets and wax paper from each colony were kept in 3.79-liter Ziploc (S.C. Johnson & Johnson, Inc., Racine, WI) bags until larvae hatched for use in bioassays. The laboratory susceptible colonies and field derived

Table 1. Description of *H. zea* field-derived populations and laboratory colonies by identification code, collection host, month collected, and collection location

Colony	Species	Collection host ^a	Month	Location	
LAB	H. zea	Z. mays		MSSTATE, MS	
AR14	H. zea	Z. mays	July	Lonoke County, AR	
GA14	H. zea	Z. mays	July	Tift County, GA	
LA14	H. zea	Z. mays	July	Franklin Parish, LA	
MSKIL14	H. zea	T. incarnatum	May	Montgomery County, MS	
MSLEL14	H. zea	T. incarnatum	May	Washington County, MS	
MSNAT14	H. zea	T. incarnatum	May	Adams County, MS	
MSSTARK13-1	H. zea	T. incarnatum	May	Oktibbeha County, MS	
MSSTARK14-1	H. zea	Z. mays	July	Oktibbeha County, MS	
MSSTONE13-1	H. zea	Z. mays	June	Washington County, MS	
MSSTONE13-2	H. zea	S. bicolor	July	Washington County, MS	
MSSTONE13-3	H. zea	C. arietinum	August	Washington County, MS	
MSVIC14	H. zea	T. incarnatum	May	Warren County, MS	
MSYAZ14	H. zea	T. incarnatum	May	Yazoo County, MS	
NC14	H. zea	Z. mays	July	Washington County, NC	
SC14	H. zea	Z. mays	July	Barnwell County, SC	
TN14-1	H. zea	Z. mays	June	Madison County, TN	
TN14-2	H. zea	Z. mays	July	Madison County, TN	

^a All collections from Z. mays were from non-Bt.

populations of each species were reared at the Mississippi State University insect rearing facility under the following conditions: 25 °C, 80% relative humidity (RH), and a photoperiod of 16:8 (L:D) h. All assays were conducted on first- and/or second-generation progeny of field-collected colonies.

Bioassays

Concentration-mortality bioassays were conducted to determine the susceptibility of H. zea to commercial formulations of flubendiamide (Belt; Bayer CropScience, Raleigh, NC) and chlorantraniliprole (Prevathon; DuPont Crop Protection, Newark, DE). Preparation of insecticide-treated diet was similar to Temple et al. (2009). Dilutions of flubendiamide and chlorantraniliprole in distilled water were made from a stock solution with a concentration of 1,000 ng/ml and 500 ng/ml, respectively, to yield eight concentrations ranging from 0 to 35 ng/ml for flubendiamide and 0 to 6.8 ng/ ml for chlorantraniliprole. Aliquots from these solutions were combined with Stonefly Heliothis Diet to yield 400 g of insecticidetreated diet for each concentration. Insecticide treated diet was stored in 0.95-liter Ziploc bags and refrigerated. All diet was used or disposed of within 7 d of preparation. Insecticide-treated diet for each concentration was dispensed into 16 wells of a 128-well bioassay tray (Product No. BAW128, Frontier Agricultural Sciences, Newark, DE) in 0.5-ml aliquots. Each well was infested with one neonate (<24h after hatching). Cells were covered with perforated, clear 16-well lids (P.E. film, Bio-Serv, Frenchtown, NJ). Infested assay trays were labeled and placed in a rearing chamber maintained at 25 °C, 80% RH, and a photoperiod of 16:8 (L:D) h. All bioassays were replicated at least four times over different days based on date of oviposition. Insect mortality was measured 7 d later. Ingestion of the diamides results in feeding cessation (Nauen et al. 2007, Hannig et al. 2009). Typically the ability of larvae to right themselves after being flipped onto their dorsal surface is considered an appropriate criterion for determining mortality with intoxicated larvae (Temple et al. 2009). Based on preliminary data of four H. zea colonies (data not presented), it was observed that intoxicated larvae, though severely stunted, could still right themselves when flipped onto their dorsal surface. Given the mobility of intoxicated larvae, severe growth inhibition was a more reliable indication of mortality. Therefore, the criteria for mortality was defined as larvae that had not molted to the second instar and weighed <10 mg after 7 d (Siegfried et al. 2000).

Data Analysis

Data were corrected for control mortality using Abbott's formula (Abbott 1925). Corrected data were analyzed with probit analysis to calculate slope, LC_{50} , LC_{90} , and confidence intervals (PROC PROBIT, SAS Institute 2012). Goodness of fit tests (P > 0.10) were evaluated to ensure the trend line fit the model. LC_{50} and LC_{90} values were considered different when 95 percent confidence intervals did not overlap.

Results and Discussion

Significant differences in LC_{50} and LC_{90} values were observed among populations of H. zea for both chlorantraniliprole (Table 2) and flubendiamide (Table 3). Overall, mean LC_{50} and LC_{90} data suggest that H. zea were \sim 6.4-fold (23.43 vs. 3.65 ng/ml) to 5.4-fold (30.51 vs. 5.65 ng/ml) more tolerant to flubendiamide than chlorantraniliprole, respectively. A similar study suggested that S. frugiperda was 13.67-fold more tolerant to flubendiamide compared with chlorantraniliprole (Hardke et al. 2011). Based on these results and those reported previously, lepidopteran larvae appear to be more sensitive to chlorantraniliprole than flubendiamide at lower concentrations.

The LC₅₀ values for chlorantraniliprole ranged from 2.94 to 4.22 ng/ml (1.44-fold), with a mean of 3.66 ng/ml (Table 2). The LC₉₀ values for chlorantraniliprole ranged from 4.52 to 9.17 ng/ml (2.02 fold), with a mean of 5.68 ng/ml. The LC₅₀ values for flubendiamide ranged from 16.45 to 30.74 ng/ml (1.86 fold), with a mean of 23.53 ng/ml (Table 3). The LC₉₀ values ranged from 21.22 to 35.33 ng/ml (1.66 fold), with a mean of 30.59 ng/ml. Overall, three field populations had LC₅₀ values that were significantly lower than the laboratory colony and three field populations had LC₅₀ values that were significantly greater than the laboratory colony for chlorantraniliprole (Table 2). Similarly, two field populations had LC₅₀

Table 2. Comparative susceptibility of *H. zea* neonates to chlorantraniliprole in dose–mortality curves generated with concentration–mortality bioassays with insecticide-treated diet

Colony	Species	N^a	Slope (±SE)	LC ₅₀ (95% CL) ^b (ng/ml)	$LC_{90}~(95\%~CL)^b~(ng/ml)$	χ^2 (df)	P^{c}
LAB	H. zea	991	3.35 (±0.24)	3.58 (3.41–3.73)	5.25 (5.01–5.55)	6.13 (5)	0.2935
AR14	H. zea	256	$2.83 (\pm 0.48)$	3.38 (2.85-3.76)	5.32 (4.79-6.23)	3.86 (5)	0.5700
GA14	H. zea	384	$3.37 (\pm 0.43)$	3.09 (2.81-3.32)	4.52 (4.20-5.00)	6.27 (5)	0.2808
LA14	H. zea	288	$3.38 (\pm 0.94)$	3.84 (2.85-4.25)	5.61 (5.17-6.88)	5.25(3)	0.1541
MSKIL14	H. zea	634	$3.82 (\pm 0.37)$	4.21 (4.02-4.39)	5.86 (5.56-6.28)	8.57 (5)	0.1275
MSLEL14	H. zea	448	$1.84 (\pm 0.23)$	3.79 (3.44-4.14)	7.61 (6.57-9.54)	5.73 (4)	0.2202
MSSTARK13	H. zea	384	$1.60 (\pm 0.21)$	4.11 (3.50-4.77)	9.17 (7.53-12.32)	3.89 (5)	0.5659
MSSTARK14	H. zea	512	$4.02 (\pm 0.64)$	3.62 (3.42-3.84)	4.98 (4.54-5.86)	1.56(5)	0.9059
MSSTONE13-1	H. zea	881	$3.13 (\pm 0.40)$	2.94 (2.75-3.11)	4.43 (4.07-5.03)	4.93 (4)	0.2947
MSSTONE13-2	H. zea	1643	$2.76 (\pm 0.23)$	3.21 (3.05-3.36)	5.10 (4.74-5.61)	4.44 (5)	0.4882
MSSTONE13-3	H. zea	1166	$4.51 (\pm 0.64)$	3.52 (3.36-3.68)	4.68 (4.35-5.27)	6.09 (5)	0.2974
MSYAZ 14	H. zea	128	$2.12 (\pm 0.43)$	4.09 (3.25-4.84)	7.47 (6.15–10.72)	8.43 (5)	0.1340
NC14	H. zea	256	$5.54 (\pm 1.01)$	4.22 (3.86-4.49)	5.32 (4.98-5.91)	6.38 (5)	0.2702
SC14	H. zea	384	$5.72 (\pm 0.93)$	4.05 (3.76-4.26)	5.06 (4.8-5.49)	9.07 (5)	0.1064
TN14-1	H. zea	256	$5.21 (\pm 0.96)$	3.72 (3.37-3.98)	4.76 (4.44-5.32)	0.43 (5)	0.9946
TN 14-2	H. zea	256	$2.49 (\pm 0.61)$	3.15 (2.19–3.67)	5.26 (4.65–6.53)	6.83 (5)	0.2336

^a Total number of insects tested.

^b Confidence limits.

^c Goodness of Fit test (P > 0.10).

6.1	Species	> 7/2	Cl (+ CE)	LC ₅₀ (95% CL) ^b (ng/ml)	LC ₉₀ (95% CL) ^b (ng/ml)	χ^2 (df)	P^c
Colony		N^a	Slope (±SE)				
LAB	H. zea	447	4.54 (±0.53)	21.96 (20.36-23.21)	29.12 (27.71–30.95)	5.39 (4)	0.2497
AR14	H. zea	256	$4.25 (\pm 0.81)$	21.88 (19.23-56.58)	29.59 (27.58-33.19)	6.29 (5)	0.2789
GA14	H. zea	256	$5.66 (\pm 1.06)$	27.75 (26.23-28.99)	34.79 (32.67-39.24)	3.35 (5)	0.6459
LA14	H. zea	256	$4.80 (\pm 0.60)$	23.34 (21.85-24.66)	30.47 (28.65-33.15)	7.27 (5)	0.2010
MSKIL14	H. zea	398	$6.89 (\pm 1.13)$	29.34 (27.58-30.48)	35.33 (34.07-37.38)	0.70(2)	0.7042
MSLEL14	H. zea	384	$4.32 (\pm 0.90)$	24.43 (21.82-25.90)	32.87 (30.86-37.31)	9.06 (5)	0.1068
MSNAT 14	H. zea	192	$5.22 (\pm 1.16)$	24.04 (19.94-26.28)	30.76 (28.57-33.74)	2.68(3)	0.4439
MSSTARK13-1	H. zea	406	$5.23 (\pm 0.61)$	17.02 (16.19-17.91)	21.75 (20.37-23.88)	1.25(3)	0.5354
MSSTARK14-1	H. zea	256	$5.04 (\pm 1.03)$	16.45 (15.25–17.77)	21.22 (19.29-25.73)	1.56(5)	0.9060
MSSTONE13-1	H. zea	947	$2.58 (\pm 0.20)$	19.72 (18.54-20.83)	32.38 (30.28-35.21)	3.28(3)	0.1938
MSSTONE13-2	H. zea	672	$2.75 (\pm 0.76)$	21.23 (15.30-23.73)	33.82 (30.75-43.90)	4.60(3)	0.2035
MSVIC14	H. zea	320	$8.04 (\pm 1.43)$	25.19 (22.51-26.86)	29.55 (27.91-31.13)	1.62(2)	0.4443
MSYAZ14	H. zea	560	$6.46 (\pm 0.72)$	30.74 (29.99-31.52)	37.49 (35.95-39.87)	1.03(3)	0.7948
NC14	H. zea	288	$9.07 (\pm 1.46)$	24.47 (23.22-25.36)	28.19 (27.23-29.60)	2.60(3)	0.4572
SC14	H. zea	896	$7.37 (\pm 0.60)$	25.70 (25.14-26.22)	30.58 (29.79-31.61)	9.23 (5)	0.1002
TN14-1	H. zea	256	$3.72 (\pm 0.63)$	22.82 (20.37-24.64)	32.20 (29.66-36.70)	3.85 (5)	0.5718
TN14-2	H. zea	256	$5.19 (\pm 0.90)$	22.30 (20.24-23.80)	28.55 (26.74-31.49)	4.34 (5)	0.5009

Table 3. Comparative susceptibility of *H. zea* neonates to flubendiamide in dose–mortality curves generated with concentration–mortality bioassays with insecticide-treated diet

values lower than the laboratory colony and five field populations had LC_{50} values that were greater than the laboratory colony for flubendiamide (Table 3). Statistical differences among populations were minimal and most likely represent natural variation among field populations. Although it cannot be ruled out because these insecticides have been used commercially since 2008, this does not appear to represent a major shift in H. zea susceptibility to this class of insecticides. Flubendiamide and chlorantraniliprole have been used extensively in soybean and grain sorghum to manage H. zea across the southern United States (Catchot et al. 2015) because of their high level of efficacy and long residual control (Tohnishi et al. 2005). As a result, H. zea populations in the southern United States have likely been exposed to some level of selection pressure to these active ingredients prior to being tested.

Concentration-mortality values of chlorantraniliprole for H. zea neonates in the current studies are considerably lower than those previously reported. Temple et al. (2009) reported mean LC₅₀ values that were 15-fold greater than the results reported here (56 vs 3.6 ng/ml). These differences are most likely because Temple et al. (2009) tested third instars compared to neonates tested in the current study. The susceptibility of multiple lepidopteran species to several insecticides has been shown to increase at later instars (Yu 1983). Although direct comparisons cannot be made to previous research, the overall purpose of this study was to develop a more rapid method for testing H. zea susceptibility to diamide insecticides. In many cases, growers and consultants need to respond to a control failure and need an answer as quickly as possible to determine if resistance is the cause of poor control. Rearing larvae to the third instar increases the time needed to confirm resistance and will negatively impact the ability of growers to respond in a timely manner.

Baseline susceptibility to the diamide insecticide class was generated in Louisiana by Hardke et al. (2011) for *S. frugiperda* and Temple et al. (2009) for *H. zea.* However, data have not been produced for these species in Mississippi or other states included in this study. Furthermore, differences observed between this study and previous studies can be attributed to differences in growth stages of

larvae tested. The long residual efficacy of the diamide insecticides may potentially expose multiple generations of the same species to the insecticide. Adams (2016) observed up to 90% and 60% control of H. zea 32 d after treatment in soybean field studies with both chlorantraniliprole and flubendiamide, respectively. This falls well within the generation time frame for H. zea during the warm summer months. Therefore, neonates were used in this study to account for subsequent populations that could potentially be exposed to the diamide insecticides as the growing season progresses and to develop a more rapid method for confirming resistance. Nevertheless, it is critical to document the variability in the response of field populations prior to the occurrence of control failures in the field. Resistance is defined as "a heritable change in the sensitivity of a pest population that is reflected in the repeated failure of a product to achieve the expected level of control when used according to the label recommendation for that pest species" (IRAC 2007). This study generated data that can serve as a reference point for future monitoring programs associated with H. zea, aiding in the detection of resistance alleles prior to field control failures. Future monitoring programs will aid resistance management efforts, allowing the diamide insecticide class to continue to play an important role in crop protection strategies.

Acknowledgments

We wish to thank all the faculty and staff at the Delta Research and Extension Center, the R.R. Foil Plant and Soil Sciences Center, the Mississippi State University Extension Service, and the Mississippi Soybean Promotion Board for their generous funding of this project. We also thank Dominic Reisig, Jeremy Greene, Gus Lorenz, Scott Stewart, Phillip Roberts, David Kerns, John Adamczyk, and Maribel Portilla for contributing populations to be tested. This publication is a contribution of the Mississippi Agricultural and Forestry Experiment Station. This material is based upon work that is supported by the National Institute of Food and Agriculture, U.S. Department of Agriculture, Hatch project under accession number 223813, MIS-721140.

^a Total number of insects tested.

^b Confidence limits.

^c Goodness of Fit test (P > 0.10).

References Cited

- Abbott, W. S. 1925. A method for computing the effectiveness of an insecticide. J. Econ. Entomol. 18: 265–267.
- Adams, A. 2016. Baseline susceptibility of selected lepidopteran pests to diamides and use strategies in Mississippi soybean. Ph.D. Dissertation, Mississippi State University, Mississippi State, MS.
- Brown, T. M., P. K. Bryson, D. S. Brickle, S. Pimprale, F. Arnette, M. E. Roof, J. T. Walker, and M. J. Sullivan. 1998. Pyrethroid–resistant Helicoverpa zea and transgenic cotton in South Carolina. Crop Prot. 17: 441–445.
- Catchot, A., J. Gore, and D. Cook. 2015. Bollworms showing up in soybean. Mississippi Crop Situation, July 17, 2015. (http://www.mississippi-crops.com/ 2015/07/17/bollworms-showing-up-in-soybeans/) (accessed June 13, 2016).
- Che, W., T. Shi, Y. Wu, and Y. Yang. 2013. Insecticide resistance status of field populations of *Spodoptera exigua* (Lepidoptera: Noctuidae) from China. J. Econ. Entomol. 106: 1855–1862.
- Cook, D. R., B. R. Leonard, J. Gore, and J. H. Temple. 2005. Baseline responses of bollworm, *Helicoverpa zea* (Boddie), and Tobacco Budworm, *Heliothis virescens* (F.), to indoxacarb and pyridalyl. J. Agric. Urban Entomol. 22: 100–109.
- Ebbinghaus-Kintscher, U., P. Luemmen, N. Lobitz, T. Schulte, C. Funke, R. Fischer, T. Masaki, N. Yasokawa, and M. Tohnishi. 2006. Phthalic acid diamides activate ryanodine-sensitive Ca2+ release channels in insects. Cell Calcium 39: 21–33.
- EPA. 2008. Pesticide Fact Sheet, Chlorantraniliprole, (http://www3.epa.gov/pesticides/chem_search/reg_actions/registration/fs_PC-090100_01-Apr-08.pdf) (accessed January 2016).
- Ffrench-Constant, R. H., and R. T. Roush. 1990. Resistance detection and documentation: the relative roles of pesticidal and biochemical assays, pp 4–38. *In R. T. Roush and B. E. Tabashnik [eds.]*, Pesticide Resistance in Arthropods. Chapman and Hall, New York, New York and London, United Kingdom.
- Fitt, G. P. 1989. The ecology of Heliothis species in relation to agroecosystems, Ann. Rev. Entomol. 34: 17–52.
- Gao, C., R. Yao, Z. Zhang, M. Wu, S. Zhang, and J. Su. 2013. Susceptibility baseline and chlorantraniliprole resistance monitoring in *Chilo suppressalis* (Lepidoptera: Pyralidae). J. Econ. Entomol. 106: 2190–2194.
- Hannig, G. T., M. Ziegler, and P. G. Marcon. 2009. Feeding cessation effects of chlorantraniliprole, a new anthranilic diamide insecticide, in comparison with several insecticides in distinct chemical classes and mode-of-action groups. Pest Manag. Sci. 65: 969–974.
- Hardke, J. T., J. H. Temple, B. R. Leonard, and R. E. Jackson. 2011. Laboratory toxicity and field efficacy of selected insecticides against fall armyworm (Lepidoptera: Noctuidae). Fla. Entomol. 94: 272–278.
- IRAC. 2007. Insecticide Resistance Action Committee, Resistance Management for Sustainable Agriculture and Improved Public Health. (http://www.irac-online.org/documents/irac-croplife-irm-booklet/? ext=pdf). (accessed January 2016).
- IRAC. 2014. IRAC MoA Classification Scheme (Version 7.3.1). (http://www.irac-online.org) (accessed October 2015).
- Jacobson, A., R. Foster, C. Krupke, W. Hutchison, B. Pittendrigh, and R. Weinzierl. 2009. Resistance to pyrethroids insecticides in *Helicoverpa zea* (Lepidoptera: Noctuidae) in Indiana and Illinois. J. Econ. Entomol. 102: 2289–2295.
- Lahm, G. P., D. Cordova, and J. D. Barry. 2009. New and selective ryanodine receptor activators for insect control. Bioorganic Med. Chem. 17: 4127–4133.
- Lai, T., and J. Su. 2011. Assessment of resistance risk in *Spodoptera exigua* (hübner) (Lepidoptera: Noctuidae) to chlorantraniliprole. Pest Manag. Sci. 67: 1468–1472.
- Musser, F. R., A. L. Catchot, Jr., J. A. Davis, D. A. Herbert, Jr., G. M. Lorenz, T. Reed, D. D. Reisig, and S. D. Stewart. 2015. 2014 Soybean Insect Losses in the Southern US. Midsouth Entomol. 8: 35–48.

- Nauen, R., S. Konanz, T. Hirooka, T. Nishimatsu, and H. Kodama. 2007. Flubendiamide: A unique tool in resistance management tactics for pest lepidoptera difficult to control. Pflanzenschutz-Nachrichten Bayer 60: 247–262.
- Qi, S., P. Lummen, R. Nauen, and J. E. Casida. 2014. Diamide insecticide target site specificity in the Heliothis and Musca ryanodine receptors relative to toxicity. J. Agric. Food Chem. 62: 4077–4082.
- Roditakis, E., E. Vasakis, M. Grispou, M. Stavrakaki, R. Nauen, M. Gravouil, and A. Bassi. 2015. First report of *Tuta absoluta* resistance to diamide insecticides. J. Pest Sci. 88: 9–16.
- SAS Institute. 2012. SAS 9.3 for windows. SAS Institute, Cary, NC.
- Sattelle, D. B., D. Cordova, and T. R. Cheek. 2008. Insect ryanodine receptors: Molecular targets for novel pest control chemicals. Invert. Neurosci. 8: 107–119.
- Scott-Ward, T. S., S. J. Dunbar, J. D. Windass, and A. J. Williams. 2001. Characterization of the ryanodine receptor-Ca2+ release channel from the thoracic tissues of the lepidopteran insect *Heliothis virescens*. J. Membr. Biol. 179: 127–141.
- Siegfried, B. D., T. Spencer, and J. Nearman. 2000. Baseline susceptibility of the corn earworm (Lepidoptera: Noctuidae) to the Cry1Ab toxin from Bacillus thuringiensis. J. Econ. Entomol. 93: 1265–1268.
- Sparks, T. C. 1981. Development of insecticide resistance in *Heliothis zea* and *Heliothis virescens* in North America. Bull. Entomol. Soc. Am. 27: 186–192.
- Sparks, T. C. 2013. Insecticide discovery: An evaluation and analysis. Pest. Biochem. Physiol. 107: 8–17.
- Su, J., T. Lai, and J. Li. 2012. Susceptibility of field populations of *Spodoptera litura* (Fabricius) (Lepidoptera: Noctuidae) in China to chlorantraniliprole and the activities of detoxification enzymes. Crop Prot. 42: 217–222.
- Teixeira, L. A., and J. T. Andaloro. 2013. Diamide insecticides: Global efforts to address insect resistance stewardship challenges. Pestic. Biochem. Physiol. 106: 76–78.
- Temple, J. H., D. R. Cook, P. L. Bommireddy, S. Micinski, W. Waltman, A. M. Stewart, B. Garber, and B. R. Leonard. 2006. Monitoring cypermethrin susceptability in Louisiana bollworm 2004-2005, pp. 1236–1240. Proc. Beltwide Cotton Conf., San Antonio, TX. 3-6 Jan. 2006. Natl. Cotton Counc. Am., Memphis TN.
- Temple, J. H., P. L. Bommireddy, D. R. Cook, P. Marcon, and B. R. Leonard. 2009. Susceptibility of selected lepidopteran pests to rynaxypyr, a novel insecticides. J. Cotton Sci. 13: 23–31.
- Tohnishi, M., H. T. Nakao, T. Furuya, A. Seo, H. Kodama, K. Tsubata, S. Fujioka, H. Kodama, T. Hirooka, and T. Nishimatsu. 2005. Flubendiamide, a novel insecticide highly active against lepidopterous insect pests. J. Pestic. Sci. 30: 354–360.
- Uchiyama, T., and A. Ozawa. 2014. Rapid development of resistance to diamide insecticides in the smaller tea tortrix, *Adoxophyes honmai* (Lepidoptera: Tortricidae), in the tea fields of Shizuoka Prefecture, Japan. Appl. Entomol. Zool. 49: 1–6.
- Wang, X., and Y. Wu. 2012. High levels of resistance to chlorantraniliprole evolved in field populations of *Plutella xylostella*. J. Econ. Entomol. 105: 1019–1023
- Wu, M., S. Zhang, R. Yao, S. Wu, J. Su, and C. Gao. 2014. Susceptibility of the rice stem borer, *Chilo suppressalis* (Lepidoptera: Crambidae), to flubendiamide in China. J. Econ. Entomol. 107: 1250–1255.
- Xu, X., M. B. Bhat, M. Nishi, H. Takeshima, and J. Ma. 2000. Molecular cloning of cDNA encoding a drosophila ryanodine receptor and functional studies of the carboxyl-terminal calcium release channel. J. Biophys. 78: 1270–1281.
- Yu, S. J. 1983. Age variation in insecticide susceptibility and detoxification capacity of fall armyworm (Lepidoptera: Noctuidae) larva. J. Econ. Entomol. 76: 219–222.