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Acute and residual concentrationdependent toxicities of some selected insecticides to adult *Bactrocera invadens* Drew, Tsuruta and White (Diptera: Tephritidae)



G. Abdullahi^{1*}, D. Obeng-Ofori², K. Afreh-Nuamah³ and M. K. Billah⁴

Abstract

Background: The high yield loss reported from the infestations of fruits by *Bactrocera invadens* Drew, Tsuruta and White in West Africa can impact negatively on the growth of horticulture in the region. Similarly, the shortage of insecticides specifically registered for the control of this pest in Ghana makes decision by farmers on which insecticide to use against this pest uncertain and risky.

Results: The result for the acute lethal toxicity shows that the lethal concentration (LC) values of the tested insecticides are 0.59 (diazinon (Diazol)), 0.81 (chlorpyrifos (Pyrinex)), 1.21 (cypermethrin+dimethoate (Cydim super)), and 3.12 (deltamethrin (Deltapaz)) ml L $^{-1}$ of water. The result for the residual toxicities of three concentrations for each of the tested insecticides shows that the lethal times LT (LT $_{50}$ and $_{90}$) for the adult *B. invadens* differed significantly for all the ages of the residues. Some insecticides exhibited low lethal time even on panels that are up to 6 days old after treatment.

Conclusion: We concluded that the acute lethal toxicity of the insecticides from most-to-least toxic are diazinon (Diazol) > chlorpyrifos (Pyrinex) > cypermethrin + dimethoate (Cydim super) > deltamethrin (Deltapaz). The residual toxicity of the label rates of all the insecticides are equal to those of the higher doses but better than those of the least doses. The implication of these findings for the management of *B. invadens* has been discussed.

Keywords: Lethal concentration, Management, Rearing, Bioassay, Tephritid fruit fly

Introduction

Tephritid fruit flies are pest of major significance in most fruit and fruity vegetable crops (Vayssières et al., 2011; Vayssières, Sanogo, & Noussourou, 2007; White & Elson-Harris, 1992; Wih & Billah, 2012). They have been reported to cause colossal economic destruction and losses in important tropical crops like citrus, mangoes, papaya, and guava (Badii, Billah, Afreh-Nuamah, & Obeng-Ofori, 2015; Ekesi & Billah, 2006; Umeh, Garcia, & De Meyer, 2008; Vayssières et al., 2007, 2011). Several

species of fruit flies including *Bactrocera invadens* Drew, Tsuruta and White, *Ceratitis capitata* (Wiedemann), *B. dorsalis* (Hendel), and *B. curcubitacea* (Coquillett) among others have been implicated globally in causing the afore mentioned damage leading to vast economic loss (Badii et al., 2015; Ekesi, Nderitu, & Rwomushana, 2006; Rwomushana, Ekesi, Gordon, & Ogal, 2008).

In west Africa, *B. invadens* have been reported to cause damage in the range of 17–73% on total fruits yield in Mango (Vayssières et al., 2007, 2011). This damage regime will have monumental effects on the sustainable productions of fruits (especially mango) in the region and other similar ecologies known to be within the geographical range of this pest (Copeland, 2006).

Full list of author information is available at the end of the article



^{*} Correspondence: gatsaranyi@yahoo.com

¹Department of Crop Protection, Modibbo Adama University of Technology, Yola, Nigeria

Sequel to this, exploring effective and appropriate management strategies to protect plantation from tephritid fruit flies pressure has become a matter of great urgency (Vayssières et al., 2011).

Despite the existence of other control methods against B. invadens, the use of synthetic insecticides still plays a central role in adult population suppressions programmes (Bateman, 1982; Vayssières et al., 2011), as the eggs are normally deposited inside the fruit and hatch therein into a larvae that feeds and develop to the last instars within the fruits, creating galleries that permits the opportunist micro-organism invasion with consequent decay of tissues around such galleries (Ekesi & Billah, 2006). Likewise, given the array of insecticidal products used by farmers for the control of fruit flies in Ghana, there are limited products that have been specifically registered for this purpose (Badii, Billah, Afreh-Nuamah, & Obeng-Ofori, 2012; EPA, 2008). Also, information on both the acute and residual toxicities of these products against especially B. invadens are lacking in the literature. Hence, the choice of the mango farmers of any of these insecticides depends on recommendations of such insecticides for control of other pests on mango and other fruits, a practice which does not work well for most pest problems (Raga & Sato, 2006; Vayssières et al., 2011). This study was therefore carried out with the sole aim of generating information on the acute and dosedependent residual toxicities of some selected insecticides commonly used by mango farmers in Ghana against B. invadens.

Materials and methods

Rearing of the Adult B. invadens used

Adult fruit flies used in this experiment were reared out of infested fruits collected from the field and processed as described by Copeland (2006) and Rwomushana et al. (2008). Fruits were placed on large plastic racks with trays of pre-sterilized moistened sand under the container to collect exiting larvae (Fig. 1a) or incubated in plastic bowls lined with sand (Fig. 1b). The sand was heat-sterilized at 100 °C for 12 h. Fruits were dissected after 2 weeks of incubation to remove any remaining larvae that may prefer to remain inside the rotting fruit if the outside environment is not conducive (Billah, personal communication). After 3 days, the sand was sieved to recollect the puparia, which were placed in petri dishes lined with moist filter paper. Daily check for emerged insect was done, and the emerged insects were transferred into holding cages (Fig. 1c). Adult insects were allowed to stay for 3-4 days to achieve full adult characteristics to facilitate proper identification. They were provided with water-soaked cotton wool and fed on a diet containing pure baker's yeast and sugar (1 yeast to 3 sugar or natural honey, v/v combination).

Adult fruit flies were identified using literature and keys of Billah and Mansell (2006); De Meyer (1998, 2000); Drew, Tsuruta, and White (2005) and White and Elson-Harris (1992). Where there was doubt, flies were sent to Dr. M. K. Billah (Department Animal biology and conservation studies, University of Ghana) for confirmation.



Insecticides products used

Four commercial insecticidal products: cypermethrin+dimethoate (Cydim super[™]), chlorpyrifos (Pyrinex 48 EC[™]), diazinon (Diazol 50 EW[™]), and deltamethrin (Deltapaz[™]) were evaluated during the study. Cydim super is an emulsifiable concentrate containing 36 g cypermethrin and 400 g dimethoate per liter. It is manufactured by Iprochem Company limited (Shenzen, China) and distributed in Ghana by Agrimat Ltd (Agrimat House, Madina). Pyrinex 48 EC™ contains chlorpyrifos 480 g/L. It is manufactured by Makhteshim Chemical Works Ltd (Beer Sheva, Israel) and distributed in Ghana by Dizengoff (Gh) Ltd. Diazol 50 EW contains diazinon 500 g/L. It is manufactured by Makhteshim chemical works Ltd (Beer Sheva, Israel) and distributed in Ghana by Dizengoff (Gh) Ltd. Deltamethrin (Deltapaz) contains deltamethrin 12.5 g/L, manufactured by Makhteshim chemical works Ltd, Beer sheva, Israel, and distributed in Ghana by Dizengoff (Ghana) Ltd.

Determination of acute toxicities of the insecticidal product

To determine the lethal concentration (LC) values of the products, the commercial grade of each product was diluted as follows: cypermethrin + dimethoate, 0.90, 1.60, 2.30, and 3.00 ml L⁻¹; chlorpyrifos, 0.80, 0.90, 1.00, and 1.10 ml L⁻¹; diazol, 1.20, 1.80, 2.50, and 3.20 ml L⁻¹, and detamethrin, 2.00, 2.60, 3.40, and 4.00 ml L⁻¹. The hard cardboard panels were dipped into the solution of the respective concentration for each product and allowed to dry for 10 min under the shade in the lab. The bioassays were conducted in 1-L transparent bottles fitted with yellow caps, from which treated cardboard panels (4 × 4 cm) were hanged (Fig. 2). A hundred and fifty

(150) holes of 1.5 mm diameter were made on the sides of the bottles for ventilation during the bioassay (Cheng, Cho, & Li, 2009). A hole of 1.5 mm in diameter was made at the center of the bottles' yellow cap to hang the treated panels with the aid of a string attached to the panels (Fig. 2). Ten adult *B. invadens* of mixed sexes in three replicates for each product were added to the treated panels in transparent bottles. Observations for mortalities were done every 2 min for 30 min. Data on mortalities were recorded at each interval and used to determine the LC values.

Insecticide concentration-dependent efficacy bioassay

Insecticide concentration-dependent efficacy was determined by calculating the lethal time (LT₅₀ and LT ₉₀) values for the field strain of adult B. invadens for each candidate insecticide. All bioassays were conducted in the Research Laboratory at the African Regional Postgraduate Programme in Insect Science (ARPPIS) building, University of Ghana, Legon, under 28 ± 2 °C, RH 68-81% and photoperiod of 12:12 D to L hours. Bioassays were conducted in a 1-L transparent bottle as described in Section 2.3 above with modification on the mode of treatment of hard cardboard panels. Here, the hardboard panels were dipped into a solution of the test insecticides at three different concentrations for each insecticidal product for 2 min. The rates used to get the respective solutions are 10 ml below the manufacturers' recommended rate (lowest concentration), the manufacturers' recommended field label rate (median concentration), and 10 ml above the manufacturers' recommended rate (highest concentration) (Moore and Miller 2006). Thus, 1.6, 2.3, and 3.0 ml of cypermethrin+dimethoate (Cydim super) per liter of water chlorpyrifos (Pyrinex)



Fig. 2 Insecticides efficacy bioassay system

48 EC were used at the rates of 0.9, 1.0, and 1.10 ml L⁻¹ of water, diazinon (Diazol) 50 EW was used at the rates of 1.8, 2.5, and 3.2 ml L⁻¹ of water, and deltamethrin (Deltapaz) was used at the rate of 2.6, 3.4, and 4.0 ml L⁻¹ of water for the concentration-dependent and residual toxicity bioassay. Prior to introduction of the insects, the treated panels were allowed to dry under laboratory condition for 20 min. Then, ten adult *B.invadens* of mixed sexes (five males and five females) aged 4–12 days were collected from the rearing cages using a 1.5 cm diameter vial and introduced into the bioassay bottles containing the treated panels. The lids of the bottles were replaced and keep in an upright position (Fig. 2).

Observations for mortalities were made at least once every 5 min. Adult flies that were knocked down and could not move any appendages for 30 s after responding to the insecticides were considered dead. The respective mortalities and times taken (lethal time (LT)) for 50 and 90% of the introduced flies to die were recorded and used to determine LT50 and LT90, for a treatment and concentration level using probit analysis. Three experiments were conducted based on the age (in days) of the treated panels to test the residual toxicities of the insecticides for each of concentration. Thus, (1) freshly treated panels (20 min after dipping), (2) 3-dayold treated panels (72 hours after dipping), and (3) 6day-old treated panels (144 h after dipping) (Urbaneja et al. 2009). In each experiment, all treatments were replicated three (3) times. A similar panel treated with only water before exposure to insects was used as a control for the different doses of the four insecticides.

Statistical analysis

Insecticide acute toxicities were determined within 30 min and expressed in LC values, and their 95% confidence limits and slope of regression lines determined by probit analysis (EPA probit analysis program, version 1.5). Data on lethal times were subjected to one-way ANOVA in GENSTAT statistical packages (GenStat Release 9.2 (PC/Windows) Genstat, 2007, Lawes Agricultural Trust (Rothamsted Experimental Station) 9th Edition). Means were separated using the least

significant difference (LSD; P < 0.05). Prior to analysis, mortalities exceeding 5% in the controls were corrected for natural mortality using Abbotts' formula (Abbott, 1925).

Results

Acute toxicities of the selected insecticidal products

The acute toxicities of the insecticidal products tested are shown in Table 1. The LC₅₀ values of all tested products are low, ranging from 0.59 (diazinon (Diazol)) to 3.10 (deltamethrin (Deltapaz)) ml L^{-1} . The toxicity ranges from most-to-least toxic are diazinon (Diazol) > chlorpyrifos (Pyrinex) > cypermethrin+dimethoate (Cydim super) > deltamethrin (Deltapaz), respectively. The LC₉₀ values of the tested products ranged from 1.03 chlorpyrifos (Pyrinex) to 5.23 deltamethrin (Deltapaz) ml L⁻¹; hence, the toxicity ranges from most-to-least toxic are chlorpyrifos (Pyrinex) > diazinon (Diazol) > cypermethrin+dimethoate (Cydim super) and deltamethrin (Deltapaz). Similarly, the 95% fiducial limits (FL) (0.55-0.89 ml L⁻¹) for chlorpyrifos (Pyrinex) and cypermethrin+dimethoate (Cydim super) (LC₅₀ 0.4–1.68 ml L⁻¹) for the LC₅₀ overlapped. Also, the 95% FL for deltamethrin (Deltapaz) (4.01–18.49 ml L⁻¹) and cypermethrin+dimethoate (Cydim super) (2.11–15.56 ml L⁻¹) overlapped at their LC_{90} values (Table 1).

Concentration-dependent lethal times residual toxicities of the insecticidal products

Cypermethrin+dimethoate (Cydim super)

The results of the concentration-dependent toxicity level comparisons for cypermethrin+dimethoate (Cydim super) across the residue ages are shown in Table 2. As expected, freshly treated panels with 3.0 ml L⁻¹ cypermethrin+dimethoate (Cydim super) was more toxic to the flies and gave the least mortality times (LT₅₀ = 29.67 \pm 13.68 and LT₉₀ = 73.67 \pm 41.67 min). However, the differences between the mean LT₅₀ values for all concentration levels were not significant except for the control that was highly significantly (P < 0.001) higher than all. However, the difference between LT₉₀ values of the least and the highest concentration (1.6 and 3.0 ml L⁻¹) were highly significant (P < 0.001) (LT₉₀ = 128.00 \pm 20.81 and 73.67 \pm 41.67 min, respectively).

Table 1 Probit analysis of mortality of adult B. invadens to insecticides

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Insecticides	No. exposed	Slope ± SE	LC ₅₀ (FL) ml/L ⁻¹	LC ₉₀ (FL) ml/L ⁻¹	χ^2
Cypermethrin+dimethoate	30	3.18 ± 1.15	1.20 (0.4–1.68)	3.04(2.11–15.56)	2.29
Chlorpyrifos (Pyrinex)	30	12.67 ± 0.93	0.81 (0.55–0.89)	1.03 (0.93–1.58)	1.04
Diazinon (Diazol)	30	3.92 ± 2.33	0.59 (*)	1.25 (*)	0.54
Deltamethrin (Deltapaz)	30	5.66 ± 1.98	3.10 (2.53-4.09)	5.23(4.01-18.49)	0.41

Legend: FL fiducial limits

^{*}Fuducial limit could not be established

Table 2 Mean lethal time (LT) for different concentrations of cypermethrin+dimethoate (Cydim super) and age of residues

		Mean ± S. E. lethal time (min)	
Age of residue	Concentration (ml L^{-1})	LT ₅₀	LT ₉₀
FTP	0.0	> 2880.00 ± 0.00	> 2880.00 ± 0.00
	1.6	66.33 ± 8.78	128.00 ± 2081
	2.3	62.00 ± 15.52	103.00 ± 14.18
	3.00	29.67 ± 13.68	73.67 ± 41.67
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	50.22 10238.18 < .001	56.24 3257.96 < .001
3-DOTP	0.0	$> 2880.00 \pm 0.00$	$> 2880.00 \pm 0.00$
	1.6	127.00 ± 54.20	194.00 ± 72.00
	2.3	66.00 ± 17.40	105.33 ± 27.51
	3.0	44.67 ± 14.66	60.33 ± 18.46
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	96.00 2270.89 < .001	129.19 1217.66 < .001
6-DOTP	0.0	$> 2880.00 \pm 0.00$	$> 2880.00 \pm 0.00$
	1.6	144.33 ± 55.47	199.67 ± 129.66
	2.3	74.33 ± 17.32	94.33 ± 21.30
	3.0	42.67 ± 9.74	62.00 ± 17.62
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	96.10 2248.90 < .001	114.30 1555.47 < .001

Legend: FTP freshly treated panels, 3-DOTP 3-day-old-treated panels, 6-DOTP 6-day-old-treated panels

In the 3-day-old treated panels experiment, the highest concentration (3.0 ml $\rm L^{-1})$ was more toxic to the flies (LT₅₀ = 44.67 \pm 14.66 min; LT₉₀ = 60.33 \pm 18.46 min), but was only highly significantly (P < 0.001) more toxic than the least concentration level (1.6 ml $\rm L^{-1})$ (LT₅₀ = 127 \pm 54.20 min; LT₉₀ = 194 \pm 72.00 min). All concentration levels were significantly more toxic (P < 0.001) than the control (LT₉₀ > 2880.0 min).

Mean toxicity difference among concentration levels in the 6-day-old treated panels was the same as the 3-day-old treated panels. Difference between the LT of the flies on the panels treated with 2.3 and 3.0 ml L⁻¹were not significant. Both concentration levels were, however, significantly more toxic (P < 0.001) to the adult $B.\ invadens$ than the controls panels.

Chlorpyrifos (Pyrinex 48 EC)

Table 3 summarizes the within treatment comparisons for the different doses of chlorpyrifos (Pyrinex) over the residues ages. The result for the freshly treated panel experiment indicated that all concentrations were highly significantly (P < 0.001) more toxic to the adult *B. invadens* than the controls. The median concentration (1.0 ml L⁻¹) was more toxic (LT₅₀ = 39.33 ± 9.39 and LT₉₀ = 56.67 ± 2.40 min) than the least concentration, but these mean mortality times were not significantly different

from the highest (1.1 ml L^{-1}) and the lowest concentration (0.9 ml L^{-1}) .

On the 3-day-old treated panel experiments, there was a highly significant difference (P < 0.001) among mean lethal times for all treatment levels. The adult flies were more susceptible to the highest concentration (1.1 ml $\rm L^{-1}$) and gave the least lethal times ($\rm LT_{50} = 19.33 \pm 5.28$; $\rm LT_{90} = 31.00 \pm 3.00$ min), acting faster than the same concentration on the freshly treated panels ($\rm LT_{50} = 29.67 \pm 7.13$ and $\rm LT_{90} = 43.66 \pm 6.64$ min). Similarly, the results obtained with the 6-day-old treated panels showed a similar pattern of toxicity to those for the 3-day-old panels treated with chlorpyrifos (Pyrinex).

Diazinon (Diazol 50 EW)

The result for experiments with panels freshly treated with varying concentrations of diazinon (Diazol) is shown in Table 4. It indicated that the highest concentration (3.2 ml $\rm L^{-1}$) was more toxic to the adult *B. invadens* and gave the lowest mean LT values (LT $_{50} = 25.33 \pm 4.41$ and LT $_{90} = 30.67 \pm 6.38$ min). Mean lethal time comparisons across the dosage concentrations were, however, not significantly different from each other, but were highly significantly different (P < 0.001) from the control.

Experiments with 3-day-old treated panels showed that all concentration levels significantly acted faster

Table 3 Mean lethal time (LT) for different concentrations of chlorpyrifos (Pyrinex) and age of Residues

		Mean \pm S. E. lethal time (min)	
Age of residue	Concentration (ml L^{-1})	LT ₅₀	LT ₉₀
	0.9	66.33 ± 16.50	74.67 ± 9.60
	1.0	39.33 ± 9.39	56.67 ± 2.40
	1.1	29.67 ± 7.13	43.66 ± 6.64
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	33.06 19569.51 < .001	19.43 56073.58 < .001
3- DOTP	0.0	> 2880.00 ± 0.00	$> 2880.00 \pm 0.00$
	0.9	38.00 ± 4.58	53.00 ± 9.00
	1.0	33.33 ± 4.41	31.00 ± 3.00
	1.1	19.33 ± 5.28	31.00 ± 3.00
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	13.44 119600.00 < .001	17.06 73617.82 < .001
6-DOTP	0.0	> 2880.00 ± 0.00	$> 2880.00 \pm 0.00$
	0.9	34.33 ± 2.33	48.33 ± 4.10
	1.0	37.33 ± 1.20	50.00 ± 6.11
	1.1	26.33 ± 1.46	35.33 ± 1.33
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	5.00 900800.00 < .001	12.19 143800.00 < .001

Legend: FTP freshly treated panels, 3-DOTP 3-day-old-treated panels, 6-DOTP 6-day-old-treated panels

Table 4 Mean lethal time (LT) for different concentrations of diazinon (Diazol) and age of residues

		Mean ± S. E. lethal time (min)	
Age of residue	Concentration (ml L^{-1})	LT ₅₀	LT ₉₀
FTP	0.0	> 2880.00 ± 0.00	> 2880.00 ± 0.00
	1.8	33.70 ± 16.5	48.88 ± 7.00
	2.5	34.67 ± 3.76	39.00 ± 7.00
	3.2	25.33 ± 4.41	30.67 ± 6.38
	LSD (<i>P</i> < 0.05)	12.38	19.21
	<i>f</i> -value p	140700.00 < .001	58152.10 < .001
3-DOTP	0.0	$> 2880.00 \pm 0.00$	$> 2880.00 \pm 0.00$
	1.8	27.67 ± 3.38	40.00 ± 5.00
	2.5	24.00 ± 3.06	38.33 ± 5.36
	3.2	13.67 ± 2.73	28.67 ± 4.70
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	8.66 289500.00 < .001	14.20 106600.00 < .001
6-DOTP	0.0	$> 2880.00 \pm 0.00$	$> 2880.00 \pm 0.00$
	1.8	24.67 ± 3.38	30.66 ± 3.40
	2.5	19.00 ± 1.16	26.33 ± 6.11
	3.2	14.67 ± 2.33	20.66 ± 1.86
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	7.00 449100.00 < .001	8.37 309300.00 < .001

Legend: FTP freshly treated panels, 3-DOTP 3-day-old-treated panels, 6-DOTP 6-day-old-treated panels

than the control. The highest concentration level (3.2 ml $\rm L^{-1}$) was highly significantly (P < 0.001) more toxic to the flies than the least concentration (1.8 ml $\rm L^{-1}$) (LT₅₀s = 13.67 and 33.7; LT₉₀s = 28.67 and 40 min, respectively).

On 6-day-old treated panels experiment with diazinon (Diazol), only the mean mortality times of the least concentration (1.8 ml $\,\mathrm{L}^{-1}$) and the highest concentration (3.2 ml $\,\mathrm{L}^{-1}$) were highly significantly different (P < 0.001). Generally, there was an overall surprising decrease in mortality times with increase in the age of residues on panels treated with diazinon (Diazol) over the experimental period (Table 4).

Deltamethrin (Deltapaz)

The results for the mean lethal times (LT) for different concentrations of deltamethrin (Deltapaz) across the ages of residues are presented in Table 5. In the freshly treated panels experiments, the highest concentration was highly significantly (P < 0.001) more toxic than all the treatment levels and the control. There were, however, no significant differences between the least concentration and the median concentration (2.6 ml L⁻¹ and 3.4 ml L⁻¹, respectively).

On the 3-day-old panel experiments, control panels significantly (P < 0.001) exhibited no toxicity (LT > 2880.00 min) to the adult *B. invadens* relative to all the

three concentration levels of deltamethrin (Deltapaz). LT $_{50}$ of the highest concentration (411.67 \pm 243.68 min) was highly significantly (P<0.001) lower than that of the least concentration (1291.00 \pm 38.67 min). The LT $_{90}$, however, indicated that median concentration level was also highly significantly different (P<0.001) from the least doses, but not from the highest concentration. The pattern of toxicity of 6-day-old treated panels was similar to that of the freshly treated panels. Deltamethrin (Deltapaz), however, showed a progressive decrease in toxicity with increase in the age of residues.

Discussion

The pestilence of *B. invadens* on host fruits depends on the mobility of the adult flies and most especially the females that lay the eggs (Vayserres et al. 2008). Laboratory studies provide avenue to estimate the toxicities of insecticides on adult flies by excluding the effects of the environmental factors/variables and their variation and can also raise reproducible quantitative data (Mahmoudvand et al., 2011; Raga & Sato, 2006). Similarly, tephritid fruit flies control relies on adult suppression ((Ekesi & Billah, 2006). Similarly, the LC data can be used to rate the acute insecticides toxicities (Mahmoudvand et al., 2011), from highly toxic LC_{50} 0.001–1.99, moderately toxic 2.0–10.99, and non-toxic >11.0 ml L^{-1} (Sunil, Thippaiah, Jagadish, & Chakravarthy, 2016). Lethal

Table 5 Mean lethal time (LT) for different concentrations of deltamethrin (Deltapaz)

		Mean ± S. E. lethal time (min)	
Age of residues	Concentration (ml L^{-1})	LT ₅₀	LT ₉₀
FTP	0.0	> 2880.00 ± 0.00	> 2880.00 ± 0.00
	2.6	360.30 ± 24.91	547.33 ± 32.27
	3.4	318.33 ± 61.76	462.00 ± 14.76
	4.0	153.00 ± 14.53	431.33 ± 14.76
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	111.20 1464.50 < .001	64.84 3648.17 < .001
3 -DOTP	0.0	> 2880.00 ± 0.00	> 2880.00 ± 0.00
	2.6	1291.00 ± 29.20	1691.00 ± 51.00
	3.4	669.67 ± 136.01	844.67 ± 93.18
	4.0	411.67 ± 243.68	771.67 ± 93.18
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	457.50 62.36 < .001	233.90 187.39 < .001
6- DOTP	0.0	> 2880.00 ± 0.00	> 2880.00 ± 0.00
	2.6	1335.00 ± 33.76	1733.00 ± 210.00
	3.4	1277.33 ± 38.76	1459.00 ± 51.10
	4.0	790.00 ± 147.31	1110.33 ± 205.00
	LSD (<i>P</i> < 0.05) <i>f</i> -value p	352.40 135.06 < .001	485.50 26.51 < .001

concentration (LC) values obtained in our study are, therefore, in the overall low, ranging from 0.59 to 3.10 ml $\,\mathrm{L}^{-1}$. This implies that all the chemical insecticides tested are good candidate products for the containment of *B. invadens* adult population explosions.

This study also reveal that diazinon (Diazol) had the highest toxicity (LC₅₀ 0.59 ml L⁻¹) > chlorpyrifos (Pyrinex) (LC₅₀ 0.81 ml L^{-1}) > cypermethrin+dimethoate (Cydim super) (LC₅₀ 1.20 ml L^{-1}) > deltamethrin (Deltapaz) (LC $_{50}$ 3.10 ml L $^{-1}$). This implies that deltamethrin is the least effective against the adult B. invadens. However, our result appears to be in contrast with the findings of Mahmoudvand et al. (2011) and Mehta et al. (2000) who reported that deltamethrin is more effective against Dacus ciliates Loew compared to dimethoate, malathion, endosulfan, and chlorpyrifos, an active ingredients similar to that of cypermethrin+dimethoate (Cydim super) and chlorpyrifos (Pyrinex) used in our studies. Also, contrary to our findings, chlorpyrifos has been reported to be of inferior efficacy when compared with lamba-cyhalothrin a pyrethroids like deltamethrin in our study (Khursheed & Raj, 2012). This disparity in result may be partly attributed to species differential response to insecticidal products or due to variable environmental conditions of the test scenarios or both factors combined (Bradbury & Coats, 1989; Worthing, 1991). Ours was a laboratory trial while Khursheed and Raj (2012) results are from field trials. Also, in a bait study, Mahat and Drew (2015) reported that females of B. tyroni, a close relative of B. invadens (Ekesi & Billah, 2006), were found to be more deterred from feeding of baits containing chlorpyrifos and malathion with a resultant rapid knockdown and higher mortalities. This is in tandem with our findings in this study that chlorpyrifos (chlorpyrifos (Pyrinex) is more effective than deltamethrin (Deltapaz) and cypermethrin+dimethoate (Cydim super). Conversely, an overlap was found between the 95% FL of the tested products with exception to dizinon (FL could not be established) indicating that all the compounds exhibited significant toxicities to the adult B. invadens. Furthermore, the B. invadens adults were more sensitive to diazinon (Diazol) and most tolerant to deltamethrin (Deltapaz). This implies that diazinon (Diazol) is the most efficacious among the entire insecticidal products tested in this study for the suppression of adult population of B. invadens. Other trials on the control of fruit flies using similar products include Khan and Khattak (2000) which tested trichlorfon and malathion as both cover sprays and bait against melon fruit flies Bactrocera cucurbitacea on musk melon Cucumis melon with both products performing much better than the control plots. Similar studies with Trichlorfon, malathion, endosulfan, Dimetheote, and Curbicron reported impressive performance against B. curcubitae on C.

melon (Agarwal, Sharma, & Rehman, 1987; Chughtai & Baloch, 1988; Chughtai, Khan, & Islam, 1984; Hussain & Khan, 1980).

The result of the dose-dependent residual response of the adult *B. invadens* to cypermethrin+dimethoate (Cydim super) concentrations exhibited a variable concentration-dependent residual lethal times (LT $_{50}$ and LT $_{90}$). The highest concentration has shown to exert least mortality times which were only significantly (P < 0.001) lower at LT $_{90}$ for the freshly treated panel experiment. This implies that even concentrations lower than the field recommended rate for this insecticide can still give a good control of adult *B. invadens*. Residual toxicities on 3- and 6-day-old treated panels showed low lethal times for the adult *B. invadens* (LT $_{50}$ 42.67 \pm 7.4–144; LT $_{90}$ 62–199.67 min on 6-day-old treated panels).

Cypermethrin acts as a stomach and contact insecticide. Its structure is based on pyrethrum, a natural insecticide found in chrysanthemum flowers, but it has a higher biological activity and is more stable than its natural model (Anonymous, 1995; Leahey, 1985). The relatively rapid degradation of cypermethrin means that it is not generally found as a residue in food (Anonymous, 1995). Cypermethrin is therefore classified by the World Health Organization (WHO) as "moderately hazardous" (Class II) (FAO/ WHO, 1996). Dimethoate on the other hand is a widely used organophosphorus (OP) insecticide applied to kill mites and insects systemically and on contact (EXTOXNET, 1996). Like all OPs, dimethoate acts by interfering with the activities of cholinesterase, an enzyme essential for the proper functioning of the nervous system of insects and humans (EXTOXNET, 1996). The foregoing coupled with our findings can add up to guarantee the safe use of cypermethrin+dimethoate (Cydim super) for B. invadens control.

chlorpyrifos (Pyrinex) (chlorpyrifos 480 g/L) showed the same trend of result as the freshly treated panels across the concentrations tested. Residual toxicities were found to be more acute on the 3- and 6-day-old treated panels relative to freshly treated panels. Chlorpyrifos (Pyrinex) contains chlorpyrifos 480 g/L. Chlorpyrifos is a broad-spectrum organophosphate insecticide that inhibits acetyl cholinesterase in insect nervous system. Chlorpyrifos acts on pests primarily as a contact poison, with some action as a stomach poison (EXTOXNET, 1993, 1996). Our study equally demonstrates that it can be used successfully for the control of fruit flies with particular emphasis on adult *B. invadens*.

Trends of lethal times among concentrations across the ages of treated panels similar to those of cypermethrin+dimethoate (Cydim super) and chlorpyrifos (Pyrinex) were found for both diazinon (Diazol) and deltamethrin (Deltapaz). Diazinon (Diazol) is a broadspectrum insecticide that contains diazinon and is an

organophosphate and a non-systemic insecticide, effective against a wide range of insect pest on all major agricultural crops. Our study points to its greater propensity of serving as a dependable insecticide against adult B. invadens. Deltamethrin on the one hand is a synthetic insecticide based structurally on natural pyrethrums, which rapidly paralyze the insect nervous system giving a guick knockdown effect (Haug & Hoffman, 1990; Hayes & Laws, 1990). Death of insects exposed to deltamethrin seems to be due to irreversible damage to the nervous system occurring when poisoning lasts more than a few hours (Leahey, 1985). Its poisoning occurs through cuticular penetration or oral uptake (EXTOX-NET, 1995: Hayes & Laws, 1990), with high compatibilities with other common insecticides and fungicides (Thompson 1998). Deltamethrin is also reported to have a very good residual activity for outdoor uses (field crops, cattle dip, and tsetse) as well as indoor uses (mosguitoes, stable flies, horseflies, fleas, cockroaches, stored product insects) (Bradbury & Coats, 1989; Worthing, 1991). It is considered the most powerful of the synthetic pyrethroids. It is up to three orders more active than some pyrethroids (Bradbury & Coats, 1989; Spencer 1981). However, it has shown a lesser residual toxicities in this our study, even when compared to its sister synthetic pyrethroids, cypermethrin on the acute toxicity ratings.

Lastly, this study revealed a general trend of the lowest doses becoming less effective in causing significantly slower mortalities along the ages of the panels. There is, therefore, a general decline of the efficiency of the least doses levels for all the tested products with an increasing age of the residues. Decline in toxicities of insecticides in the field as well as under laboratory conditions against fruit flies with an increasing treatment time has been reported (Mahat & Drew, 2015; Peck & Mcquate, 2000; Reynolds, Osborne, & Barchia, 2017; William, Valle, & Vanuela, 2003)

Conclusion

In conclusion, the results of this study suggest that all the tested insecticidal products can be a potential tool for the control of *B. invadens* in Ghana and similar ecologies. Diazinon is, however, the most effect product owing to the least LC and LT times. However, actual field trials are required to validate our result under the prevailing agro-climatic conditions.

Abbreviations

LC: Lethal concentrationLTLethal timeECEmulsifiable concentratesARPPISAfrican Regional Postgraduate Programme in Insect ScienceFLFiducial limitsLSDLeast significant differenceLtdLimitedANOVAAnalysis of varianceet al.And othersFTPFreshly treated Panels3-DOTP3-Day-old treated panels panels

Acknowledgements

We gratefully acknowledge the German Academic Exchange Service (DAAD) for funding this research as part of the first authors work for his M. Phil in Entomology at African Regional Postgraduate Programme in Insect Science (ARPPIS)-West African sub-regional center, University of Ghana, Legon, Ghana. All facilities provided by ARPPIS and University of Ghana are hereby acknowledged.

Authors' contributions

AG had conceived the idea, performed the practical section, and collected the samples, analyzed the data, and wrote his master thesis (at which this manuscript is based on). MKB had identified the insect and carried out the taxonomic part of this study. DO, KA, and MKB helped in designing the study, supervised the preparation of the experiment, and helped in writing the manuscript. All authors read and approved the final manuscript.

Funding

Funding for the entire Master degree program for the first author was provided by German DAAD.

Availability of data and materials

Data is available on request.

Ethics approval and consent to participate

Not applicable for this kind of research.

Consent for publication

Not applicable

Competing interests

The authors declare that they have no competing interests.

Author details

¹Department of Crop Protection, Modibbo Adama University of Technology, Yola, Nigeria. ²Department of Crop Science, University of Ghana, Legon, Ghana. ³Agricultural Research Centre, Kade, Institute of Agricultural Research, University of Ghana, Legon, Ghana. ⁴Department of Animal Biology and Conservation Sciences, University of Ghana, Legon, Ghana.

Received: 4 October 2019 Accepted: 11 February 2020 Published online: 04 May 2020

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