

Mechanisms of pyrethroid resistance in *Haematobia irritans* (Muscidae) from Mato Grosso do Sul state, Brazil

Mecanismos de resistência da *Haematobia irritans* (Muscidae) a piretróides em Mato Grosso do Sul, Brasil

Antonio Thadeu Medeiros Barros^{1*}; Teresinha Tizu Sato Schumaker²; Wilson Werner Koller³;
Guilherme Marcondes Klafke⁴; Thais Aguiar de Albuquerque²; Rodrigo Gonzalez²

¹Embrapa Pantanal, Corumbá, MS, Brasil

²Departamento de Parasitologia, Instituto de Ciências Biomédicas – ICB, Universidade de São Paulo – USP, São Paulo, SP, Brasil

³Embrapa Gado de Corte, Campo Grande, MS, Brasil

⁴Instituto de Pesquisas Veterinárias Desidério Finamor – Fepagro/IPVDF, Eldorado do Sul, RS, Brasil

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Abstract

Horn fly resistance to pyrethroid insecticides occurs throughout Brazil, but knowledge about the involved mechanisms is still in an incipient stage. This survey was aimed to identify the mechanisms of horn fly resistance to cypermethrin in Mato Grosso do Sul state, Brazil. Impregnated filter paper bioassays using cypermethrin, synergized or not with piperonyl butoxide (PBO) and triphenyl phosphite (TPP), were conducted from March 2004 to June 2005 in horn fly populations (n = 33) from all over the state. All populations were highly resistant to cypermethrin, with resistance factors (RF) ranging from 89.4 to 1,020.6. Polymerase chain reaction (PCR) assays to detect the knockdown resistance (*kdr*) mutation also were performed in 16 samples. The *kdr* mutation was found in 75% of the tested populations, mostly with relatively low frequencies (<20%), and was absent in some highly resistant populations. Addition of TPP did not significantly reduce the LC₅₀ in any population. However, PBO reduced LC₅₀s above 40-fold in all tested populations, resulting in RFs ≤ 10 in most cases. Horn fly resistance to cypermethrin is widespread in the state, being primarily caused by an enhanced activity of P450 mono-oxygenases and secondarily by reduced target site sensitivity.

Keywords: Horn fly, insecticide resistance, metabolic resistance, *kdr*.

Resumo

Resistência da mosca-dos-chifres a inseticidas piretróides ocorre em todo o país, entretanto, o conhecimento sobre os mecanismos envolvidos é ainda incipiente. Este estudo objetivou identificar os mecanismos de resistência desta mosca à cipermetrina em Mato Grosso do Sul. Bioensaios utilizando papéis impregnados com cipermetrina, isoladamente ou sinergizada por butóxido de piperonila (PBO) ou trifetil fosfato (TPP), foram realizados de março/2004 a junho/2005 em 33 populações. Todas as populações apresentaram elevada resistência à cipermetrina, com fatores de resistência (FR) variando de 89,4 a 1.020,6. Ensaios de reação em cadeia da polimerase (PCR) visando a detecção de *kdr* (“knockdown resistance”) foram realizados em 16 amostras. A mutação *kdr* foi detectada em 75% das populações, geralmente em baixas frequências (<20%) e ausente em algumas populações resistentes. A adição de TPP não reduziu significativamente a CL₅₀ em nenhuma população. Entretanto, o PBO reduziu em mais de 40 vezes a CL₅₀ de todas as populações testadas, resultando em FR ≤ 10 na maioria dos casos. Resistência da mosca-dos-chifres à cipermetrina encontra-se disseminada no estado, sendo causada primariamente por um aumento da atividade de P450 mono-oxigenases e secundariamente pela redução da sensibilidade do sítio de ação do inseticida.

Palavras-chave: Mosca-dos-chifres, resistência a inseticidas, resistência metabólica, *kdr*.

*Corresponding author: Antonio Thadeu Medeiros Barros
Embrapa Pantanal, Rua 21 de Setembro, 1880, Nossa Senhora de Fátima,
CP 109, CEP 79320-900, Corumbá, MS, Brasil
e-mail: thadeu.barros@embrapa.br

Introduction

It took about a century for the horn fly, *Haematobia irritans irritans*, to cross the American continent after its introduction in the United States during the 1880's (SLINGERLAND, 1891) until it reached Southern Cone countries (VALÉRIO; GUIMARÃES, 1983; LUZURIAGA et al., 1991; CARBALLO; MARTÍNEZ, 1991).

Although resistance to several insecticide classes had been previously reported in U.S. (SPARKS et al., 1985), horn fly populations remained susceptible in Brazil until the mid 1990's, as shown by efficacy (GRISI; SCOTT, 1992; PEREIRA et al., 1994) and bioassay (SCOTT et al., 1994) studies with pyrethroid and organophosphate insecticides. However, continued reliance on commercial pyrethroid products for controlling cattle pests led to development of pyrethroid resistance in horn fly populations and became a major concern throughout the country (BARROS et al., 2012).

Insecticide resistance is an individual mutation-induced reduction in susceptibility to lethal drugs, which may become a population trait through selection by drug exposure, thus impairing the insect control. Individual horn fly resistance to pyrethroids may be phenotypically expressed by changes in penetration and metabolism of those compounds (SPARKS et al., 1990; SHEPPARD, 1995) as well as reduced target site sensitivity (knockdown resistance) (GUERRERO et al., 1997). Knockdown resistance (*kdr*) and enhanced metabolic detoxification have been considered the major mechanisms involved in pyrethroid resistance (BULL et al., 1988; SPARKS et al. 1990; SHEPPARD, 1995).

The *kdr* is already known in Brazilian pyrethroid-resistant horn fly populations (GUERRERO; BARROS, 2006; SABATINI et al., 2009), but little is known about its real importance and the role played by metabolic mechanisms.

This study reports a survey on susceptibility of horn flies to cypermethrin and the search for the mechanisms behind horn fly resistance to pyrethroids in the state of Mato Grosso do Sul, Brazil.

Materials and Methods

The field survey was conducted from March 2004 to June 2005 in the state of Mato Grosso do Sul (Figure 1), located in the Brazilian Mid-West, looking for mechanisms of pyrethroid resistance in horn fly populations. Horn fly field bioassays and sampling were conducted by Embrapa Pantanal and the molecular analyses of fly samples were performed at the University of São Paulo (USP).

1. Field bioassays

Previous selection of cattle ranches for conducting the insecticide bioassays was based on convenience and practical factors (ease of access, owner concurrence, suitable infrastructure, and fly availability), but was random regarding suspicion of insecticide resistance or any other particular situation. Ranchers were previously requested to keep a cattle herd untreated for at least two weeks before testing.

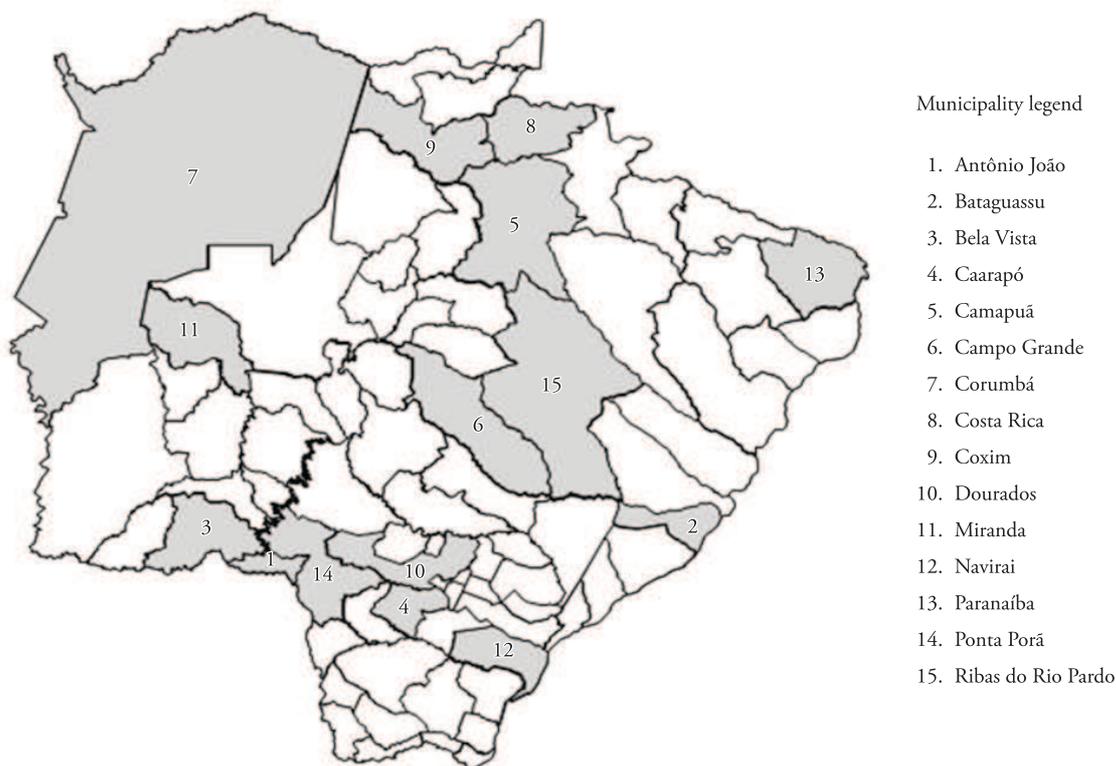


Figure 1. Geographic scope of the survey on horn fly susceptibility and resistance mechanisms to pyrethroids in the state of Mato Grosso do Sul, Brazil (2004-2005).

Impregnated filter paper bioassays (SHEPPARD; HINKLE, 1987) were used to assess susceptibility of horn fly populations to technical grade cypermethrin (89.56% purity) serially diluted in acetone (Merck P.A.). Cypermethrin kits contained three replicates of ten concentrations ranging from 1.6 to 819.2 $\mu\text{g}/\text{cm}^2$, which allowed assessment of variable levels of susceptibility in the field. Control papers were treated with acetone only. Impregnated filter papers were kept in aluminum foil under refrigeration until they were placed in plastic Petri dishes (90 mm diameter) before the bioassay. Each treated paper was used in four bioassays (twice per side).

To investigate the involvement of metabolic mechanisms in resistance, bioassays with cypermethrin synergized with either 5% piperonyl butoxide (PBO, 90% purity, Sigma-Aldrich, USA) or 5% triphenyl phosphate (TPP, +99% purity, Sigma-Aldrich, USA) were also performed at each site, depending on fly availability. After dilution in acetone, the synergist concentration (5%) was kept constant along the insecticide concentrations. After preliminary field bioassays for adjusting insecticide concentration range in the kits, concentrations of cypermethrin-TPP were similar to cypermethrin alone (1.6-819.2 $\mu\text{g}/\text{cm}^2$), while the concentration range of cypermethrin-PBO was much lower and varied from 0.1 to 3.2 $\mu\text{g}/\text{cm}^2$. In each bioassay, the potential toxicity of synergists was evaluated by papers treated with the 5% synergist solution only.

Horn flies were collected from cattle with entomological hand nets and transferred to dishes immediately after an adequate number of flies had been collected. Early fly mortality was assessed immediately after dishes were loaded and dead flies were excluded if present. Actual fly susceptibility to the insecticide was evaluated by assessing fly mortality after a 2-hour exposure; flies unable to walk were considered dead. After insecticide kits were loaded, a sample of that population was transferred to a plastic vial with commercial ethanol for later molecular studies.

Pooled mortality data from the three replications were analyzed by probit analysis using POLO-PC (LEORA SOFTWARE, 1987) to obtain lethal concentration (LC_{50}) and respective fiducial limits for each field population. Bioassays with fly mortality >10% in control dishes were not considered. The insecticide kits produced yearly were tested with a susceptible horn fly colony maintained at the USDA-ARS Knipping-Bushland US Livestock Insects Research Laboratory (Kerrville, TX, USA) to provide reference LC_{50} . Resistance factors (RFs) were calculated by dividing LC_{50} from field populations by the LC_{50} from the susceptible colony. Differences in LC_{50} were assumed to be statistically significant when 95% fiducial limits did not overlap.

2. Molecular analysis

Polymerase chain reaction (PCR) was performed in horn flies from 16 populations sampled in 14 municipalities. Detection of the *kdr* genotype mutation for each population followed the protocol of Guerrero et al. (1998). Alleles from 35 individuals were amplified per sampled population.

3. Extraction of genomic DNA

Genomic DNA was extracted according to Moreira-Ferro et al. (1998), with some modifications. The head of flies were individually homogenized in 50 μL of lysis buffer (100 mM Tris-HCl pH 7.5;

100 mM NaCl; 100 mM EDTA; 1% SDS; 1 mg/mL proteinase K) and incubated for 1h at 60 °C. The sample containing genomic DNA was subjected to extraction with an equal volume of phenol (2 \times), phenol/chloroform (1 \times), and chloroform/isoamylalcohol (1 \times). The DNA contained in the aqueous phase was precipitated by adding 100% ethanol and incubated (20 minutes) in ethanol with dry ice.

After centrifugation (5 minutes, 11,600 g, 4 °C), the pellet was dissolved in 1ml of TE (10 mM Tris; 1 mM EDTA, pH 8.0) containing 10 mg/mL RNase. The DNA was again precipitated by adding 20% (v/v) of 3 M sodium acetate and 1 mL of isopropanol. Next, the suspension was maintained for 15 minutes at room temperature and then centrifuged (30 minutes, 11,600 g, 4 °C). The precipitate obtained was washed with 70% ethanol, dried in a vacuum centrifuge, and then dissolved in 20 μL of bidistilled water.

4. *Kdr* detection

The *kdr* gene was amplified from total genomic DNA according to Guerrero et al. (1998). The PCR was conducted with 25 ng of genomic DNA, 20 pmol of each primer (FG-129, FG-138, and/or FG-130 or FG-134) (GUERRERO et al., 1998), 10 mM Tris (hydroxymethyl) aminomethane hydrochloride pH 8.3, 50 mM KCl, 0.05 mM each dNTP, 3.5 mM MgCl_2 , and 0.1 μL of 1:1 (vol:vol) mix of *AmpliTaq* DNA polymerase (5 units per microliter of stock solution (ss)) and TaqStart Antibody (1.1 $\mu\text{L}/\mu\text{L}$ ss). PCR was conducted as follows: 96 °C for 2 minutes, 35 cycles (94 °C for 1 minute, 62 °C for 1 minute, 72 °C for 1 minute) and 72 °C for 7 minutes. The PCR product was visualized on 1.4% agarose gel stained with ethidium bromide after electrophoresis.

Results and Discussion

A total of 87 bioassays were conducted on 33 cattle ranches located in 15 municipalities from all ten microregions of the state (Table 1). High levels of resistance to cypermethrin were detected in all populations, with RF ranging from 89.4 to 1,020.6 (Table 2). Except for one population, the RFs were above 100. All resistance levels were much higher than necessary to reduce efficacy of cypermethrin-based products for controlling horn flies (GUGLIELMONE et al., 1998). Therefore, failure of horn fly control was expected to occur in those ranches, as informed during visits.

Table 1. Survey on horn fly susceptibility and resistance mechanisms to pyrethroids in the state of Mato Grosso do Sul, Brazil (2004-2005).

Mesoregions	Microregions	Municipalities
Mid-North	Alto Taquari	Camapuã, Coxim
	Campo Grande	Campo Grande
East	Cassilândia	Costa Rica
	Nova Andradina	Bataguassu
	Paranaíba	Paranaíba
	Três Lagoas	Ribas do Rio Pardo
Pantaneis	Baixo Pantanal	Corumbá
	Aquidauana	Miranda
Southwest	Bodoquena	Bela Vista, Ponta Porã
	Dourados	Antônio João, Caarapó, Dourados, Naviraí

Table 2. Bioassay dosage-mortality results to cypermethrin with and without synergist in horn fly populations from the state of Mato Grosso do Sul, Brazil (2004–2005).

Location (Year)	Cypermethrin			Cypermethrin+PBO			Cypermethrin+TPP		
	LC ₅₀ (95% FL)	Slope (SE)	RF	LC ₅₀ (95% FL)	Slope (SE)	RF	LC ₅₀ (95% FL)	Slope (SE)	RF
Antônio João #1 (2004)	18.77 (15.91-22.10) ^a	4.17 (0.36)	89.4	-	-	-	57.45 (26.18-171.21) ^{ab}	2.15 (0.18)	-
Bataguassu #1 (2005)	31.32 (27.17-35.62) ^a	2.88 (0.27)	195.8	0.43 (0.35-0.52) ^{ab}	2.74 (0.21)	10.8	55.46 (31.52-127.04) ^a	1.24 (0.15)	383.0
Bataguassu #2 (2005)	18.98 (16.88-21.43) ^a	4.49 (0.69)	118.6	0.45 (0.39-0.54) ^{ab}	2.82 (0.66)	11.3	17.34 (8.48-25.15) ^a	3.12 (0.38)	462.2
Bataguassu #3 (2005)	36.71 (30.89-43.16) ^a	1.92 (0.15)	229.4	0.37 (0.17-0.49) ^{ab}	3.30 (0.35)	9.3	27.05 (15.30-43.09) ^a	1.67 (0.11)	144.5
Bela Vista #1 (2004)	46.15 (35.47-61.53) ^a	2.49 (0.19)	219.8	0.87 (0.61-1.03) ^{ab}	5.53 (0.71)	4.4	45.56 (26.63-78.78) ^a	2.58 (0.18)	225.4
Caarapó #1 (2005)	33.85 (29.63-38.62) ^a	2.35 (0.16)	211.6	-	-	-	53.17 (47.46-59.58) ^{ab}	3.77 (0.41)	303.7
Camapuã #1 (2005)	22.42 (15.63-30.08) ^a	2.41 (0.20)	140.1	0.18 (0.16-0.21) ^{ab}	2.69 (0.26)	4.5	17.31 (11.45-24.78) ^a	3.02 (0.29)	443.1
C. Grande #1 (2005)	163.29 (111.14-212.77) ^a	1.36 (0.23)	1020.6	1.13 (0.81-1.69) ^{ab}	1.04 (0.15)	28.3	106.15 (56.35-296.50) ^a	0.93 (0.11)	144.3
C. Grande #2 (2005)	52.89 (27.96-88.34) ^a	1.61 (0.14)	330.6	0.40 (0.29-0.53) ^{ab}	1.78 (0.16)	10.0	22.88 (11.25-36.34) ^a	2.09 (0.21)	884.6
Corumbá #1 (2004)	24.68 (13.24-41.95) ^a	2.09 (0.17)	117.5	-	-	-	24.18 (20.75-28.09) ^a	2.08 (0.18)	190.7
Costa Rica #1 (2005)	31.36 (22.09-43.57) ^a	2.76 (0.20)	196.0	0.23 (0.15-0.32) ^{ab}	3.49 (0.32)	5.8	19.17 (10.50-29.77) ^a	1.75 (0.14)	161.2
Costa Rica #2 (2005)	75.58 (59.12-97.21) ^a	1.76 (0.13)	472.4	0.38 (0.32-0.44) ^{ab}	2.79 (0.18)	9.5	72.21 (60.06-86.87) ^a	1.28 (0.08)	159.8
Costa Rica #3 (2005)	31.03 (27.32-35.08) ^a	3.91 (0.49)	193.9	0.20 (0.15-0.25) ^{ab}	2.54 (0.22)	5.0	-	-	601.8
Coxim #1 (2005)	24.75 (17.10-32.81) ^a	2.89 (0.26)	154.7	-	-	-	25.66 (13.00-34.24) ^a	3.92 (0.58)	-
Coxim #2 (2005)	30.24 (18.20-40.69) ^a	3.62 (0.40)	189.0	0.21 (0.16-0.25) ^{ab}	3.29 (0.26)	5.3	17.58 (15.11-20.14) ^a	2.87 (0.30)	213.8
Coxim #3 (2005)	18.25 (16.33-20.14) ^a	3.68 (0.35)	114.1	0.11 (0.08-0.13) ^{ab}	2.54 (0.27)	2.8	18.50 (14.73-23.19) ^a	2.29 (0.13)	146.5
Dourados #1 (2005)	27.15 (16.04-42.27) ^a	3.66 (0.30)	169.7	0.19 (0.12-0.24) ^{ab}	2.57 (0.23)	4.8	41.22 (36.07-47.10) ^a	2.25 (0.16)	154.2
Dourados #2 (2005)	42.77 (32.55-59.44) ^a	2.97 (0.22)	267.3	0.30 (0.16-0.41) ^{ab}	3.35 (0.35)	7.5	87.66 (60.29-128.97) ^{ab}	1.49 (0.09)	343.5
Miranda #1 (2005)	28.46 (15.80-39.82) ^a	2.91 (0.33)	177.9	0.28 (0.18-0.36) ^{ab}	2.62 (0.25)	7.0	37.45 (31.72-43.20) ^a	2.86 (0.31)	730.5
Miranda #2 (2005)	23.87 (15.59-35.48) ^a	3.08 (0.25)	149.2	0.29 (0.16-0.38) ^{ab}	3.65 (0.47)	7.3	17.32 (15.40-19.38) ^a	3.76 (0.36)	312.1
Miranda #3 (2005)	22.72 (15.52-32.53) ^a	3.04 (0.21)	142.0	0.11 (0.04-0.16) ^{ab}	2.44 (0.22)	2.8	18.06 (16.11-20.11) ^a	3.22 (0.27)	144.3
Navirai #1 (2005)	62.16 (55.37-70.70) ^a	2.80 (0.25)	388.5	-	-	-	74.72 (56.12-100.75) ^a	1.01 (0.19)	150.5
Navirai #2 (2005)	28.82 (16.79-46.49) ^a	2.26 (0.16)	180.1	0.21 (0.13-0.28) ^{ab}	2.08 (0.18)	5.3	23.21 (15.35-37.06) ^a	3.35 (0.27)	622.7
Navirai #3 (2005)	31.18 (19.90-44.16) ^a	3.78 (0.34)	194.9	0.16 (0.06-0.23) ^{ab}	3.14 (0.29)	4.0	30.59 (16.36-50.37) ^a	3.35 (0.27)	193.4
Paranaíba #1 (2005)	30.73 (27.61-33.93) ^a	5.08 (0.65)	192.1	0.63 (0.41-0.86) ^{ab}	2.78 (0.21)	15.8	34.65 (26.33-58.90) ^a	1.98 (0.16)	254.9
Ponta Porã #1 (2004)	32.94 (19.51-53.91) ^a	2.65 (0.19)	156.9	-	-	-	27.51 (19.25-38.82) ^a	2.10 (0.40)	288.8
Ponta Porã #2 (2005)	37.73 (27.97-49.93) ^a	2.47 (0.18)	235.8	0.13 (0.04-0.23) ^b	1.56 (0.16)	3.3	32.32 (24.86-41.44) ^a	2.60 (0.20)	183.4
Ponta Porã #3 (2004)	35.94 (26.68-56.81) ^a	1.78 (0.23)	171.1	-	-	-	32.23 (26.86-38.00) ^a	1.74 (0.12)	269.3
Ponta Porã #4 (2005)	76.34 (55.99-122.01) ^a	1.13 (0.17)	477.1	-	-	-	50.21 (36.39-70.66) ^a	2.03 (0.16)	214.9
Ribas do RP #1 (2005)	34.88 (26.27-45.11) ^a	3.06 (0.26)	218.0	-	-	-	36.65 (25.34-50.64) ^a	1.43 (0.10)	418.4
Ribas do RP #2 (2005)	39.55 (35.00-44.50) ^a	2.51 (0.18)	247.2	-	-	-	71.75 (52.56-95.64) ^{ab}	2.92 (0.28)	305.4
Ribas do RP #3 (2005)	26.37 (18.12-39.51) ^a	2.85 (0.22)	164.8	0.28 (0.25-0.31) ^{ab}	2.82 (0.24)	7.0	23.69 (19.93-28.12) ^a	1.55 (0.11)	597.9
Ribas do RP #4 (2005)	33.95 (29.48-38.68) ^a	2.83 (0.27)	212.2	-	-	-	29.62 (18.63-45.27) ^a	1.87 (0.16)	197.4
Kerrville colony (2004)	0.21 (0.16-0.30)	4.07 (0.48)	-	0.20 (0.17-0.24)	5.81 (0.60)	-	0.15 (0.14-0.17)	2.44 (0.20)	246.8
Kerrville colony (2005)	0.16 (0.14-0.18)	7.64 (0.92)	-	0.04 (0.02-0.05)	5.33 (1.02)	-	0.12 (0.10-0.14)	5.64 (0.55)	-

PBO (piperonyl butoxide), TPP (triphenyl phosphate); LC₅₀ = lethal concentration to 50% of the population (expressed as µg/cm²) after 2 hours exposure to the insecticide; RF (resistance factor) = LC₅₀ of field population/LC₅₀ of the reference susceptible colony (Kerrville, USA); LC₅₀ statistically significant difference: (a) compared to the Kerrville reference strain, (b) between synergized and non-synergized bioassays.

The widespread resistance of horn flies to cypermethrin found in this survey confirmed previous reports from the state (BARROS et al., 2007) and all over the country (BARROS et al., 2012). Previous cypermethrin RFs detected by Barros et al. (2007) did not exceed 91.3 (most below 60), while in the present study the lowest RF was 89.4 and most were above 140, suggesting that pyrethroid resistance is getting worse in the region. In practice, development of resistance tends to increase insecticide use by producers, by either applying higher amounts of insecticide products and/or reducing the interval between treatments, in an attempt to recover original control levels. As a cyclical process, higher exposure to insecticides tends to increase resistance selection reducing product efficacy and consequently control efficiency, which leads to more insecticide use and higher selection pressure.

The *kdr* mutation, associated with pyrethroid target site insensitivity, was detected in most horn fly populations, confirming previous reports from the state (GUERRERO; BARROS, 2006) and elsewhere in the country (SABATINI et al., 2009). Horn flies with *kdr* genotypes were detected in 75% of sampled populations (Table 3), a higher frequency than previously recorded in the state, which strengthens the apparent worsening trend of pyrethroid resistance in the region. Homozygous (RR) *kdr* flies were detected only in two populations, the ones with the highest frequencies of mutant flies, similarly as found by Guerrero and Barros (2006).

Although *kdr* was commonly found within populations, the frequency of mutant flies in each population did not exceed 20% (allelic frequency $\leq 10\%$) in 83.33% of the populations (Table 3). Such low frequencies followed a similar trend to that previously reported in the state (GUERRERO; BARROS, 2006; SABATINI et al., 2009) and likely reflected a low to moderate selection pressure associated with routinely poor control practices in most cases. The highest frequency of *kdr* flies (54.3%; allelic frequency = 34.3%) was detected in the top resistant population

(RF = 1,020.6), which seems compatible with the high potential of resistance known for this mechanism (OPPENORTH, 1985). However, inhibition of metabolic mechanisms by PBO dropped that RF to only 28.3, which would be the level of resistance actually attributable to *kdr* in that particular population.

The presence of *kdr* in most populations suggested a significant role played by this mechanism in pyrethroid resistance in the region; however, its absence in some populations as well as its low frequency in most populations cannot account for the high resistance levels found. Actually, the evidences gathered by previous studies (GUERRERO; BARROS, 2006; BARROS et al., 2007; SABATINI et al., 2009) have already suggested that the major mechanism behind pyrethroid resistance in the country would be metabolic, most likely oxidative. This situation contrasted with findings abroad, where higher frequencies of *kdr* have been reported and this mechanism seemed to play a more important role (JAMROZ et al., 1998; LI et al., 2003; OYARZÚN et al., 2011).

No toxicity by 5% TPP was observed to colonized or wild horn flies. Exposure of horn flies to TPP in bioassays did not result in significant reduction of cypermethrin LC₅₀ (Table 2). Increases of cypermethrin susceptibility due to esterase (EST) inhibition by TPP did not exceed 2.3-fold in field populations compared to 1.4-fold for susceptible flies from the reference colony. Guerrero and Barros (2006) failed to detect an esterase-mediated mechanism by using PCR-based assays, but found a significant reduction of cypermethrin LC₅₀ in a single population (with 50% of *kdr* flies) in TPP-synergized bioassays, suggesting the involvement of EST as a mechanism of resistance. However, the effect of a synergist that inhibits a detoxication pathway tends to be much greater in the presence of *kdr*, which increases opportunity for insecticide degradation (OPPENORTH, 1985). Thus, it seems that the actual existence of an EST-based resistance mechanism in Brazilian horn fly populations needs further evidence.

Table 3. Profile of knockdown resistance (*kdr*) genotype in horn fly populations from Mato Grosso do Sul state, Brazil (2004-2005).

Location	Sample size (n)	<i>Kdr</i> genotype ¹			<i>Kdr</i> allelic frequency ²	Frequency of <i>kdr</i> flies ³	Cypermethrin RF ⁴
		SS	SR	RR			
Antônio João #1	35	35	0	0	0.0	0.0	89.4
Bataguassu #1	35	25	5	5	21.4	28.6	195.8
Bela Vista #1	35	32	3	0	4.3	8.6	219.8
Camapuã #1	35	30	5	0	7.1	14.3	140.1
C. Grande #1	35	16	14	5	34.3	54.3	1020.6
Corumbá #1	35	28	7	0	10.0	20.0	117.5
Costa Rica #2	35	32	3	0	4.3	8.6	472.4
Coxim #2	35	35	0	0	0.0	0.0	189.0
Dourados #1	35	35	0	0	0.0	0.0	169.7
Miranda #2	35	34	1	0	1.4	2.9	149.2
Naviraí #2	35	32	3	0	4.3	8.6	180.1
Paranaíba #1	35	35	0	0	0.0	0.0	192.1
Ponta Porã #1	35	33	2	0	2.9	5.7	156.9
Ponta Porã #2	35	34	1	0	1.4	2.9	235.8
Ponta Porã #3	35	31	4	0	5.7	11.4	171.1
Ribas do RP #3	35	34	1	0	1.4	2.9	164.8
Total	560	501	49	10	Avg. 6.2	10.5	–

¹S represents a pyrethroid susceptible-associated allele and R represents a pyrethroid resistance-associated allele; ²*Kdr* allelic frequency – percentage of the *kdr* allele in the total number of alleles in that *locus* (2 alleles per fly); ³Frequency of *kdr* flies – percentage of horn flies with a *kdr* genotype (SR and/or RR) in the population;

⁴RF (resistance factor) = LC₅₀ of field population/LC₅₀ of the reference susceptible colony (Kerrville, USA).

The PBO activity as an inhibitor of microsomal oxidases is well established (CASIDA, 1970; FARNHAM, 1998), being considered as an efficient insecticide synergist and an important tool in studies regarding insecticide metabolism and resistance mechanisms (HODGSON; LEVI, 1998).

No toxicity by 5% PBO was evidenced to both colonized and wild horn flies. Addition of PBO in bioassays dramatically reduced cypermethrin LC_{50} s in all field populations, dropping cypermethrin RFs (between 114.1 and 1,020.6 in non-synergized bioassays) to 10 or less in 81.82% of the populations (Table 2). Synergism ratios exceeded 40 (ranged 42.2 to 290.2) in all field populations, while a maximum reduction of 4-fold was observed in LC_{50} of colonized susceptible flies. Such a sharp fall in cypermethrin RF following PBO exposure indicated the strong involvement of P450 mono-oxygenases in pyrethroid resistance in all studied horn fly populations. A similar effect of PBO, primarily associated with an enhanced metabolic detoxification by oxidases, has been also observed in highly permethrin resistant house flies, despite the presence of other mechanisms (SCOTT; GEORGHIU, 1986). Although *kdr* was detected in more than half of the flies of the top resistant population in the present study, the LC_{50} was reduced above 140-fold (from 163.29 to only 1.13 $\mu\text{g}/\text{cm}^2$) following PBO exposure, confirming that a major mechanism of resistance was oxidative.

Although 5% PBO showed a high synergism, later unpublished studies indicated that lower PBO concentrations provided higher synergism to cypermethrin in impregnated paper bioassays, suggesting that the synergism factors reported here would be probably higher if a lower PBO concentration had been used in this study.

Besides inhibition of oxidases, PBO may interfere with other insecticide-insect processes such as the cuticular penetration of insecticides (SCOTT; GEORGHIU, 1986). However, changes in penetration just provide lower levels of resistance, being of secondary importance (OPPENORTH, 1985), and would not explain the observed resistance levels. In addition, a partial inhibition of resistance-associated EST by PBO has been observed in some insects (GUNNING et al., 1998; YOUNG et al., 2005), but not in horn flies (LI et al., 2007). Although it should not be discounted that the higher susceptibility of horn flies to cypermethrin following PBO exposure may not rely solely on the inhibition of metabolic oxidation, there was no evidence that other resistance mechanism affected by this synergist played a significant role in the pyrethroid resistance showed by those populations. As mentioned earlier, the presence of *kdr* may increase the synergist effect, which is also true for PBO; however, the low frequency of this mechanism in almost all analyzed populations surely reduced the *kdr* importance as a synergist enhancer in this study.

The marked decline in resistance levels of all populations exposed to PBO, the lack of significant increase of susceptibility due to TPP, as well as the absence or low frequency of *kdr* in highly resistant populations to cypermethrin, all together pointed to the conclusion that the primary mechanism of horn fly resistance to pyrethroids in the studied populations was an enhanced oxidative metabolism by P450 mono-oxygenases.

Moreover, it should be pointed out that the LC_{50} s obtained from PBO-synergized bioassays were significantly higher than the susceptible colony (even in populations without detected *kdr*),

which may be explained by incomplete inhibition of oxidases associated with the pyrethroid resistance, presence of *kdr* (detected or not), and/or involvement of other (undetected) mechanism of resistance.

Pyrethroid resistance levels and mechanisms detected in this study ultimately resulted from the local strategies for controlling horn flies on pastured beef cattle, which routinely rely on the inadequate use of insecticide products applied by backpack sprayers and application of product doses well below the technically recommended (BARROS et al., 2007). Although the selection pressure imposed by this common practice has obviously succeeded for selecting the metabolic mechanism, it seemed to be less successful regarding selection of the *kdr* mechanism and its associated fitness disadvantages (SCOTT et al., 1997; YOUNGER, 2011). Nevertheless, the importance of *kdr* may increase quickly if intensity of selection pressure by pyrethroid-dependent control strategies, using more frequent treatments or long lasting formulations, becomes strong enough to overcome the fitness disadvantages showed by *kdr* flies in the absence of pyrethroids.

As a short-term approach, the use of PBO tends to increase pyrethroid toxicity against horn flies, thus improving efficacy of pyrethroid products and extending their use in the field for horn fly control. On the other hand, the multiple resistance mechanisms found in most populations makes efficient fly control and resistance management more complex and difficult to achieve if management strategies focus on a single mechanism, such as the use of synergized insecticides.

A worse scenario regarding horn fly resistance to pyrethroids should be expected if pyrethroid products continue to dominate the market and their indiscriminate use persists in the field. Therefore, adequate horn fly control depends not just on reducing pyrethroid use itself, or simply replacing it by an insecticide from another class, but also on developing and adopting alternative control approaches that reduce chemical dependence and improve control efficiency and sustainability.

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