Functional characterization and fitness cost of spinosad-resistant alleles in *Ceratitis capitata*

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Abstract
The sustainability of control programs for the Mediterranean fruit fly, *Ceratitis capitata*, for citrus crops in Spain has been threatened by the development of resistance to malathion and lambda-cyhalothrin in recent years. Spinosad is widely used without apparent loss of efficacy. However, a highly resistant strain, JW-100s, has been obtained after laboratory selection. Spinosad resistance in JW-100s has been associated with different mutant alleles of the α6 subunit of the nicotinic acetylcholine receptor (*Ccα6*) including an isoform-specific truncation allele, *Ccα6*\(^{3aQ68*}\). Using the GAL4 > UAS system in *Drosophila melanogaster* to demonstrate expression of this truncated α6 subunit, in a *dα6* loss-of-function genetic background, does not rescue susceptibility to spinosad, while the expression of *Ccα6* wild-type isoforms does. We have also generated *C. capitata* isolines from JW-100s homozygous for: (1) the *Ccα6*\(^{3aQ68*-Δ3b-4}\) allele, which contains the mutation 3aQ68*, and (2) the *Ccα6*\(^{3aQ68*-K352*}\) allele, which contains the mutations 3aQ68* and K352*. Neither of these produce complete *Ccα6* transcripts. The frequency of resistant alleles declined when in competition with individuals carrying the wild-type allele. Through extensive testing of both biological and behavioral fitness traits, we identified a reduced ability of *Ccα6*\(^{3aQ68*-Δ3b-4}\) males to detect the parapheromone and to mate with females carrying the *Ccα6*\(^{3aQ68*-K352*}\) allele in competition experiments. Thus, not only the potential for spontaneous resistant mutations to arise in *Ccα6* but also their fitness costs must be considered when planning resistance management strategies for *C. capitata*.

Keywords Medfly · nAChR · GAL4 > UAS · Fitness traits · Behavior

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Key message

- The sustainability of medfly control programs is threatened by insecticide resistance.
- Our results highlight the plasticity of medfly to generate mutations in nAChR α6 that can lead to loss-of-function phenotypes conferring spinosad resistance.
- Resistant alleles have a fitness cost, which could be the reason for the absence of resistance events in the field.
- The results of this study will be useful in designing resistance management strategies for medfly.

Introduction

The Mediterranean fruit fly (medfly) *Ceratitis capitata* (Wiedemann) (*Diptera: Tephritidae*) is one of the main pests for citrus and other fruit trees. It causes significant...
economic losses because of direct damage to fruit as well as restricting the capacity of growers to export to medfly-free areas due to costs related to harvesting surveillance and control actions. While the sterile insect technique is being implemented in some area-wide integrated pest management programs, C. capitata control still relies heavily on the use of insecticides (Reddy and Rashmi 2016). Malathion was the most widely used insecticide in Spain from the 1990s until its withdrawal from use in the EU in 2009. Since then, spinosad and lambda-cyhalothrin in bait sprays and deltamethrin in mass trapping and lure and kill devices (Navarro-Llopis et al. 2015) have been the most widely used insecticides for medfly control in citrus crops (MAPA 2019).

The sustainability of medfly control programs is threatened by insecticide resistance (Vontas et al. 2011). Spanish field populations of C. capitata have developed resistance to malathion (Magaña et al. 2007, 2008) and lambda-cyhalothrin (Arouri et al. 2015) due to the intensive use of these products. Cross-resistance to deltamethrin has been observed in a laboratory lambda-cyhalothrin-resistant strain (Arouri et al. 2015). There is still no evidence of field resistance to spinosad in Spain, but a highly resistant strain, JW-100s, has been obtained after laboratory selection (Ureña et al. 2019). Since then, spinosad and lambda-cyhalothrin in bait sprays and deltamethrin in mass trapping and lure and kill devices (Navarro-Llopis et al. 2015) have been the most widely used insecticides for medfly control in citrus crops (MAPA 2019).

More than 250 cases of spinosad resistance have been documented for different pest species (IRAC 2019). Target site-based resistance has been reported for a number of insects, including Plutella xylostella (Baxter et al. 2010; Rinkevich et al. 2010; Wang et al. 2016), Baculovora dorsalis (Hsu et al. 2012), Frankliniella occidentalis (Puinean et al. 2013), Thrips palmi (Bao et al. 2014) and Tuta absoluta (Berger et al. 2016; Silva et al. 2016). In most cases, it has been associated with mutations that result in premature stop codons (Baxter et al. 2010; Rinkevich et al. 2010; Hsu et al. 2012), exon-skipping events (Berger et al. 2016) and deletions (Wang et al. 2016) in the a6 gene, which presumably result in non-functional proteins. However, the point mutations P146S and G275E in the a6 gene that do not disrupt transcription (Puinean et al. 2013; Bao et al. 2014; Somers et al. 2015; Silva et al. 2016) and enhanced metabolism mediated by P450 and esterase enzymes (Geng et al. 2013) have also been associated with spinosad resistance in some insects.

Reversion of resistance in the absence of spinosad selection has been reported for Musca domestica (Khan et al. 2014), Liriomyza trifolii (Ferguson 2004), Spodoptera litura (Rehan and Freed 2014), P. xylostella (Sayyed et al. 2008). T. absoluta (Campos et al. 2014) and F. occidentalis (Bielza et al. 2008b), indicating the existence of a fitness cost associated with spinosad resistance. There is evidence that the knockout of a6 function in D. melanogaster does not cause lethality and no fitness effects have yet been reported in this species (Perry et al. 2007). However, when different trade-offs related to growth and developmental traits were analyzed in other species, fitness costs ranged from absent or minor (Wyss et al. 2003; Bielza et al. 2008a; Abbas et al. 2014) to substantial (Li et al. 2007; Wang et al. 2010; Okuma et al. 2018). The need to investigate fitness, behavioral and life-history traits that require the function of receptors being targeted by insecticides has been highlighted (Perry and Batterham 2018). While nAChRs may have many roles in behavior, the known roles for the different subunits identified in D. melanogaster currently include sleep (Dx1, Dx3 and Dx4—Shi et al. 2014; Somers et al. 2017; Wu et al. 2014), courtship (Dx1—Somers et al. 2017) and escape response (Dx7—Fayyazuddin et al. 2006). Only a few studies have considered how spinosad resistance affects behavior and the impact that loss-of-function or truncated a6 transcripts could have. Changes to feeding behavior in F. occidentalis (Li et al. 2017) and reduced competitive mating in Heliothis virescens (Wyss et al. 2003) have been reported, but the molecular basis of resistance was unknown in both cases.

Spinosad resistance in the laboratory selected JW-100s strain has been associated with three different mutant alleles of the a6 nAChR gene of medfly (Ccα6): (1) the Ccα6 3aAG>AT allele, with the AG > AT mutation in the
5' splice site of exon 3a, which produces full-length transcripts carrying exon 3b and incomplete transcripts that skip exon 3a and join exons 2 and 4; (2) the Cca63aQ68* allele, with 3aQ68* mutation giving rise to truncated transcripts at exon 3a and full-length transcripts with exon 3b; and (3) the Cca63aQ68*–K352* allele with mutations 3aQ68* and K352*, which lead to truncated transcripts at exon 3a, and to 3b transcripts truncated at exon 10 (Ureña et al. 2019). None of the Cca6 transcripts from the Cca63aQ68*–K352* allele could generate complete Cca6 protein products. This mutant would therefore be expected to be resistant to spinosad (Perry et al. 2007). However, why individuals homozygous for the Cca63aQ68* or Cca63aAG>AT alleles are resistant is not known, as these alleles only affect isoforms containing exon 3a while generating wild-type full-length Cca6 isoforms with exon 3b. The analysis of the frequency of the three resistant alleles during the selection process for JW-100s revealed fluctuations. In brief, alleles Cca63aAG>AT and Cca63aQ68* were most frequent at generation F29. However, after generation F45, the allele Cca63aAG>AT was not found and Cca63aQ68*–K352*, not detected in previous generations, had the highest frequency. Finally, at generation F85, the population mainly consisted of individuals homozygous for Cca63aQ68*–K352* (92%). These data could be explained by a combination of differences in resistance levels and fitness costs for the three alleles. Further, the spontaneous generation of mutations during the laboratory selection process may also have been a factor (Ureña et al. 2019). Resistance to spinosad in the JW-100s strain is recessive (Ureña et al. 2019), consistent with other reported examples of α6-based resistance (Bielza et al. 2007; Perry et al. 2007; Baxter et al. 2010; Wang et al. 2016).

The goal of this study was to determine whether or not the relationship between spinosad resistance and Cca63aQ68* and Cca63aQ68*–K352* alleles is causal. To achieve this goal, we have created transgenic D. melanogaster lines that express mutated and wild-type isoforms of Cca6 using the GAL4 > UAS expression system and also generated C. capitata isolines homozygous for these two mutant alleles. In addition, we have assessed the stability of spinosad resistance in C. capitata linked to these two alleles in the absence of spinosad selection. This along with the analysis of biological (developmental time, adult weight) and behavioral (parapheromone detection by wind tunnel and electrophysiological assays, and mating success) traits in the mutants allows us to examine whether there are fitness costs associated with these resistant alleles. The results of this study will be useful in designing resistance management strategies for medfly.

**Materials and methods**

**C. capitata strains and generation of isolines**

The susceptible laboratory strain of *C. capitata*, C, was established from medfly individuals from non-treated experimental fields of the Instituto Valenciano de Investigaciones Agrarias (IVIA, Valencia, Spain) in 2001. Since then, it has been reared in the laboratory without any exposure to insecticides.

The *C. capitata* isolines described here were derived from the spinosad-resistant laboratory strain JW-100s (Ureña et al. 2019). To obtain isolines homozygous for the mutant alleles, a process of crossing and genotyping was performed. All of the crosses were initiated with 50 virgin females (isolated as pupae) and ten males under standard rearing conditions (16:8 h light: dark photoperiod and a temperature of 26 ± 2 °C (light) and 22 ± 2 °C (dark) with water and food *ad libitum*). After 7–10 days, individual females were placed in tubes containing around 20 g of larval rearing medium (standard rearing conditions) for oviposition. When the progeny of these females had developed into larvae and pupae, their female parents were genotyped and the progeny of females carrying the allele of interest were selected for the next cross. The genotyping was done according to two PCR protocols described in Ureña et al. (2019), a PCR–RFLP for the detection of the K352* mutation in exon 10, identified via the presence of an *Msel* restriction site in the mutant allele, and a multiplex (“multiplex PCR-2” in Ureña et al. (2019)) to detect the Q68* mutation in exon 3a. Both protocols allow mutant- and wild-type homozygotes and heterozygotes to be distinguished. Homozygous lines were sequenced to ensure that all individuals were of the expected genotype. The sequencing was performed by Secugen (Madrid, Spain) on a 3730 XL DNA Analyzer (Applied Biosystems) using BigDye® Terminator v3.1 reagents. The DNA sequences produced were analyzed using Geneious 11.0.5 (https://www.geneious.com).

In order to obtain the Q68*–K352* isolate, a cross between JW-100s females and C males, homozygous for the wild allele (*Cca6*/Cca6+), was performed. F1 progeny which female parent carried the *Cca6*3aQ68*–K352*/ Cca63aQ68*–K352* genotype were selected and inter-crossed. The progeny F2 embryos were placed on larval rearing medium (stock prepared by mixing 300 g of sugar, 11.2 g of N-propyl-p-hydroxybenzoate (Sigma-Aldrich, Japan), 11.2 g of methyl-4-hydroxybenzoate (Sigma-Aldrich, India), 10 g of benzoic acid (Merck, China), 145.2 g of dry beer yeast (Difco Laboratories, Erembodegem, Belgium), 1000 g of wheat bran and 2400 ml of distilled water) containing 5 ppm of spinosad to eliminate the *Cca6*/Cca6+...
and Cca6+/Cca63aQ68+/K352s genotypes. F2 survivors were used to establish the homozygous Q68s–K352s isolate (Cca63aQ68+/K352s & Cca63aQ68+/K352s) (Online Resource 3a).

To obtain Q68s isolate, an initial cross was done between JW-100s females and males of the Q68s–K352s isolate. F1 females carrying the Cca63aQ68s+ allele were selected for crossing to C (Cca6+/Cca6+) males, and the same process was repeated with F2 females. F3 individuals carrying the Cca6+/Cca63aQ68s+ genotype were inter-crossed, and the resulting F3 embryos were placed on larval rearing medium containing 5 ppm spinosad to eliminate Ccα63aQ68sΔ3bs homozygous for a new allele (see Results).

**Genomic DNA extraction, PCR of intronic/exonic boundaries and sequencing of Cca6**

Genomic DNA (gDNA) extraction from single flies from Q68s and Q68s–K352s isolines and JW-100s and C strains of C. capitata was carried out with tissue homogenization and DNA isolation using SpeedTools Tissue DNA Extraction (Biotools, Madrid, Spain), according to the manufacturer’s instructions. gDNA was quantified using Nanodrop ND-1000 spectrophotometer (Thermo Fisher Scientific, Waltham, MA, USA).

The boundaries between intronic regions and exons 2, 3a, 3b, 4 and 5 of Cca6 were PCR amplified. The reactions were performed in a volume of 10 μl using 0.4 μM of each oligonucleotide (Online Resource 1), 0.5 U of AmpliTaq Gold DNA Polymerase (Thermo Fisher Scientific, Austin, USA), 10× PCR Buffer II, 1.5 mM MgCl2, 0.2 mM dNTPs (Thermo Fisher Scientific, Austin, USA) and 100–200 ng of the corresponding template. PCR conditions were as follows: an initial step at 95 °C for 3 min; 45 cycles of 95 °C for 10 s and 60 °C for 30 s; 50 °C for 3 min and a melting ramp of 50–99 °C raising by 1 °C and waiting for 5 s in each step. Actin-5C, glycerol-3-phosphate dehydrogenase (GPDH) and α-tubulin were used as reference genes to normalize the data.

PCR for the coding regions was performed in a volume of 25 μl using 0.8 μM of each oligonucleotide (Online Resource 1), 0.5 U of AmpliTaq Gold DNA Polymerase (Thermo Fisher Scientific, Austin, USA), 10×PCR Buffer II, 1.5 mM MgCl2, 0.2 mM dNTPs (Thermo Fisher Scientific, Austin, USA) and 5 μl of the corresponding cDNA as template. PCR conditions are described in Sect. 2.2., and then, PCR products were purified and sequenced.

**GAL4 > UAS targeted expression of Cca6 isoforms in D. melanogaster**

* D. melanogaster transgenic flies with an insertion of wild-type Cca6 isoforms 3a8b (MK251469) (strain UAS-Cca63a8b) and 3b8a (MK251470) (strain UAS-Cca63b8a) and the mRNA isoform carrying 3aQ68s mutation (strain UAS-Cca63aQ68sΔ3bs) of the Cca6 gene were created in the da6s background (as per Perry et al. 2015) to specifically express Cca6 isoforms using the GAL4 > UAS method (Brand and Perrimon 1993).

**Cloning of wild-type and mutant mRNA isoforms of Cca6**

Mutated and wild-type sequences of Cca6 were obtained by conventional PCR from JW-100s resistant and C wild-type individuals, respectively. Samples were acquired by RNA extraction of single flies, and cDNA synthesis was performed as described above. The oligonucleotides used (Sigma-Aldrich, Spain) are listed in Online Resource 2.
The mutant allele was amplified through PCR in a volume of 25 µl using 0.4 µM of forward FnAch6ex1 and reverse RnAch6Q68 oligonucleotides, with reagents and conditions as mentioned above. Wild-type isoforms were amplified in a volume of 50 µl using 0.5 µM of forward FnAch6ex1nest and reverse RnAch6ex12 oligonucleotides, 2 U of iTaqTM DNA Polymerase (Bio-Rad, California, USA), 5× HF Buffer, 0.2 mM dNTPs and 8 µl of the corresponding cDNA diluted 100-fold as template. PCR conditions were as follows: an initial denaturation step at 98 °C for 30 s; 35 cycles of 98 °C for 10 s, 60 °C for 30 s and 72 °C for 1.5 min; and a final step of 72 °C for 7 min for full extension.

PCR products were cloned into the pGEM-T-easy vector (Promega Corporation, Madison, USA) following manufacturer instructions. Transformation was performed by electroporation with a 2.5 V electric pulse for 4.6 ms using a MicroPulser™ electroporator (Bio-Rad, USA). Minipreps were performed using the alkaline lysis method (Engbercht et al. 1991). The amount of plasmid DNA was quantified using a NanoDrop (Thermo Fisher Scientific, Massachusetts, USA) and sequenced (Secugen, Madrid, Spain) to identify the isoform present and to detect any mutations introduced in the cloning process.

**Construction of vectors for transformation of D. melanogaster**

Clones were digested with NotI (New England Biolabs, NEB) and ligated into digested pUASattB (Bischof et al. 2007) using 1U DNA T4 ligase (Promega Corporation, Madison, USA) and 2× buffer. Transformation was performed using a standard heat shock protocol. Clones with proper orientation were selected and cultures established. Plasmids were purified (Wizard® Plus SV Minipreps DNA Purification System, Promega, USA) and sequenced (Macrogen, South Korea) with T7 and T3 oligonucleotides (Online Resource 2).

**Microinjection of Cca6 constructs to generate transgenic lines**

Transgenic UAS-Cca6 lines of D. melanogaster were produced by embryonic microinjection into 786G strain, which lacked expression of the nAChα6 gene (da6ns) (Perry et al. 2015). Plasmid DNA (100–250 mg/µl) was injected into the posterior pole of dechorionated 0–30-min-old embryos using a Femtojet microinjector (Eppendorf, USA) with a FemtoII needle (Eppendorf, Hamburg, Germany). Single-pair crosses were established with survivors that developed to the adult stage, and then, F1 individuals carrying the transgene (distinguishable by their eye color) were crossed to obtain homozygous lines.

To drive the expression of UAS constructs, homozygous transgenic males were crossed to females homozygous for the X-linked Da6 > GAL4 which drives expression in cells that normally express Da6 (Perry et al. 2015). This was conducted in the da6ns background.

**Susceptibility bioassays with spinosad**

**Susceptibility of C. capitata**

The susceptibility of C. capitata Q68* and Q68*–K352* isolines to spinosad was tested with a topical application assay with increasing doses of this insecticide (Dow AgroSciences 88% p/p, Indianapolis, USA). Dilutions were prepared using acetone as a solvent for spinosad and for the non-treated control. A volume of 0.5 µl of each solution was applied to each fly with an automated microapplicator 900X (Burkard Manufacturing Co., Hertfordshire, UK). Four replicates of each dose were performed with 15 flies per replicate. After the insecticide treatment, flies were confined in ventilated plastic dishes (89 mm in diameter and 23 mm in height) with water and the rearing diet (4:1 sugar/yeast) ad libitum, in an environmentally controlled chamber (Sanyo MLR-350-H, Sanyo, Japan), at 25 ± 1 °C and 16 h light and 8 h dark photoperiod (standard conditions). Mortality was recorded after 48 h. Flies were considered dead if they were ataxic.

**Susceptibility of D. melanogaster**

The D. melanogaster transgenic strains were also tested for susceptibility to spinosad. The resistant 786G strain (carrying da6ns) and da6ns;UAS-Da6-3a3b flies (that carries a susceptible nAChα6 transcript from D. melanogaster, isoform 3a3b) were used as positive and negative controls, respectively. Bioassays were performed by placing F1 first-instar larvae in vials containing food with increasing doses of spinosad (0, 0.05, 0.5 and 5 ppm). Food was prepared using Instant Drosophila Medium (Carolina™, Burlington, North Carolina, USA). The experiment included five replicates of each spinosad concentration and five vials with non-treated food, with 50 larvae per vial. Vials were kept in the environmentally controlled chamber in constant darkness at 25 ± 1 °C. The number of flies that had emerged after 16–19 days was recorded.

**Stability of Cca63aQ68Δ3b–4 and Cca63aQ68–K352* alleles in C. capitata**

Pupae from the Q68*(Cca63aQ68Δ3b–4/)Ccα63aQ68Δ3b–4 and Q68*–K352* (Ccaα63aQ68–K352*/Ccα63aQ68–K352*) isolines and C strain (Cca6+/Cca6+) were isolated, and the adults sexed immediately after emergence to guarantee virginity.
Males and females from each strain were placed separately into ventilated plastic dishes and maintained with water and rearing diet for 2 days. The following reciprocal crosses were performed: 100 ♂ Q68* × 100 ♀ C and 100 ♂ Q68* × 100 ♂ C to obtain QC-G0 (genotype Cca6b3aQ68Δ3a/3bCcα6*); 100 ♂ Q68*–K352* × 100 ♀ C and 100 ♀ Q68*–K352* × 100 ♂ C to obtain KC-G0 (genotype Cca6b3aQ68Δ3b*/3aCcα6*); and 100 ♂ Q68*×100 ♀ Q68*–K352* and 100 ♀ Q68*×100 ♀ Q68*–K352* to obtain QK-G0 (genotype Cca6b3aQ68Δ3b*/3aCcα6*). The progeny of reciprocal crosses were pooled at G0. Three replicates of QC and KC crosses were established and maintained for nine generations without insecticide selection pressure. Three replicates of the QK cross were established and maintained without selection until G1. At generation G2, each replica of QK was split into two, one continued without exposure to insecticide (QKn), whereas the other one was exposed to 100 ppm spinosad for 72 h (QKs) every generation for nine generations. Between 20 and 40 flies of each cross and generation were frozen at −20 °C for genotyping.

Fitness of resistant isolines and the susceptible strain of C. capitata

Several assays were performed to analyze biological and behavioral traits in Q68* and Q68–K352* isolines and the C strain.

Developmental time to pupation and adults’ weight

Eggs (corresponding to a volume of 50 µl) were collected and spread on larval rearing medium (160 g approx.) in containers (130×90×25 mm) covered with an aluminum foil to avoid desiccation. Containers were placed in 21 ventilated plastic boxes allowing third-instar larvae that jump from the food container and pupate to be collected. Eggs were kept in the environmentally controlled chamber in constant darkness at 25 ± 1 °C. Three replicates of each strain were prepared. The number of pupae obtained was recorded every day and the pupae removed from the box. Fifty females and 50 males (2–3-day-old adult flies) of each strain were collected and weighed with a precision balance (AM100, Mettler-Toledo, Switzerland).

Attraction to a parapheromone in a wind tunnel

The attraction of males for trimedlure (a parapheromone) was evaluated in a wind tunnel (Jang et al. 1994). A piece of trimedlure dispenser weighing 0.52 g (Aralure, Agrisense, Pontypridd, UK) was suspended from the top of the wind tunnel (63×63×215 cm, Plexiglas flight tunnel, Analytical Research Systems, Gainesville, FL, USA) at a height of 20 cm, next to the air inlet, while 50 male adult flies were released downwind at the other end of the tunnel. A yellow sticky trap (Koppert, Berkel en Rodenrijs, The Netherlands) was located next to the trimedlure so that flies approaching the lure would be stuck. The rate of airflow was 0.2 m/s, the temperature 25 °C and the humidity 40%. Flies were left inside the tunnel for 30 min. The time taken for the first fly of each strain to reach the trap was recorded, as well as the number of caught flies in each 5 min interval. Each isolate/strain was evaluated four times. A control assay (same conditions, flies and yellow trap without trimedlure) was performed to measure any effect of the yellow traps on flies. The number of flies stuck in the presence of trimedlure was corrected by the number of flies stuck without it.

Electrophysiological antennal activity

Electroantennography (EAG) was performed to determine whether males were able to produce a response when stimulated with trimedlure (Cossé et al. 1995). Heads of 3–7-day-old adult flies were removed from the rest of the body and connected to two electrodes of an EAG probe-type PRG-2 (Syntech) coated with a conductive Spectra 360 electrode gel (Parker Laboratories Inc., Fairfield, NJ, USA); one electrode was inserted into the tip of one antenna and the other into the back of the head. A Syntech IDAC 2 acquisition controller and a Syntech CS-55 stimulus controller (OCKENFELS SYNTECH GMBH, Kirchzarten, Germany) were used to evaluate the electroantennographic response. GC-EAD 32 (v 4.3) software was employed to record the data. Serial dilutions of the tested compound were made in re-distilled HPLC-grade hexane. The solution (5 µl) was applied to a filter paper strip located inside a 15-cm-long Pasteur pipette. Control puffs of air (pipette containing a filter paper with no impregnated solution) and a blank (filter paper plus hexane) were performed with each head before the exposure to trimedlure. The response to air was used as a negative control as it showed low EAG activity, very close to 0 mV. The response to hexane was used as a positive control as C. capitata individuals show EAG activity when exposed to this compound. Between 5 and 15 replicates of each strain/isoline were performed. Thus, EAG amplitudes recorded when exposed to trimedlure were corrected by the value of observed with hexane exposure.

Male competition in mating

Mating assays were performed to study the mating success of males of the three strains. Virgin 3–5-day-old males and females were used to make crosses with ten females of the same strain and 12 males of each of the three strains. Five replicates of each cross were established. Flies were located in ventilated methacrylate boxes (12 cm diameter, 5 cm in height) provided with water and food ad libitum. Crosses

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were kept during 6 h in a climatic room, at 26 ± 2 °C during the light hours. Then, individual females were placed in separate tubes containing rearing medium for the laying of the eggs, and under standard conditions until F1 pupae were obtained (approximately 10 days). Three pupae derived from each female were genotyped. The presence of Cca6\(^{3aQ68*–K352*}\), Cca6\(^{3aQ68*3b3b}\) and Cca6\(^{3a}\) alleles allowed the genotype of the male mating with each female to be determined.

**Statistics**

In all experiments, data were statistically analyzed with the Levene and Shapiro–Wilk tests to test the homogeneity and normality of the data, respectively. Developmental time to pupation, weight of adult flies, first fly to reach the source of stimuli in the wind tunnel, competitive mating assays with C. capitata and arsine-transformed susceptibility to spinosad of D. melanogaster transgenic lines were analyzed by ANOVA followed by the Student–Newman–Keuls post hoc test. Expression levels of Cca6 exons were performed on Cq values (log2 transformation of NRQs) using Student's t test (exonic regions 3a–4 and 2–3b) or ANOVA followed by Student–Newman–Keuls post hoc test (exonic regions 2–3a and 7). The attraction to trimedlure of a group of flies in wind tunnel was evaluated by a repeated measures ANOVA, also followed by Student–Newman–Keuls post hoc test. A Kruskal–Wallis test was performed for the analysis of EAG. Chi-square test was completed to test the fit of the data to expected genotypic frequencies in the stability assay. For the analysis of susceptibility to spinosad of C. capitata isolines/strains, mortality data were used to estimate LD\(_{50}\) values (dose needed to cause 50% mortality) and Probit analysis was performed using the program POLO-PC (LeOra Software 14), which corrects exposed samples’ mortality with the control mortality using Abbott's transformation (Abbott 1925). Resistance ratios [RR = LD\(_{50}\) (lab strain/isoline)/LD\(_{50}\) (C strain)] were considered significant if their 95% fiducial limits did not include 1 (Robertson and Preisler 1992).

**Results**

**Expression of Cca6\(^{3aQ68*}\), Cca6\(^{3aQ68*–K352*}\) and Cca6\(^{3a}\) mRNA isoforms in D. melanogaster**

In order to study the contribution that the different Cca6 alleles make to spinosad resistance/susceptibility, it is important that they be compared in an identical genetic background. Such experiments are readily performed in D. melanogaster. Thus, we generated UAS transgenic D. melanogaster flies carrying wild-type mRNA isoforms Cca6\(^{3a}\) (UAS-Cca6\(^{3a}\)) and Cca6\(^{3b}\) (UAS-Cca6\(^{3b}\)), and mutant mRNA isoform Cca6\(^{3aQ68*}\) (UAS-Cca6\(^{3aQ68*}\)). We used the methods of Perry et al. (2015) to introduce the mRNA isoforms of Cca6 into the resistant background of 786G strain, carrying the null da6\(^{ns}\) allele.

Bioassays of susceptibility to spinosad were performed with F1 flies from the cross between transgenic UAS-Cca6 strains and the Da6 > GAL4;da6\(^{ns}\) driver strain (Fig. 1a). Flies expressing UAS-Cca6\(^{3a}\) and UAS-Cca6\(^{3b}\) had close to 100% mortality when exposed to 0.5 ppm of spinosad in the food. Those flies expressing UAS-Cca6\(^{3aQ68*}\) only had approximately 20% mortality, even at the highest concentration tested (5 ppm). As expected, the progeny from the control cross (Da6 > GAL4;da6\(^{ns}\)×786G) did not exhibit significant mortality at any of the tested concentrations of spinosad, while mortality of the F1 produced by crossing Da6 > GAL4;da6\(^{ns}\) with da6\(^{ns}\);UAS-Da6-3a3b (the susceptible nACh6 transcript) was greater than 60% at all concentrations tested. We found that UAS-Cca6\(^{3a}\) and UAS-Cca6\(^{3b}\) strains also showed spinosad susceptibility when we tested them without driving the expression of the transgene (Fig. 1b). This could be due to basal expression of the transgenes and suggests that minimal expression levels of Cca6 are enough for spinosad to be toxic. Thus, rescue experiments showed that nACh6 wild-type mRNA isoforms 3a and 3b are capable of rescuing the response to spinosad, while the mRNA isoform carrying the 3aQ68* mutation does not.

**Characterization of Q68* and Q68*–K352* isolines of C. capitata**

To study the contribution of resistant alleles Cca6\(^{3aQ68*}\) and Cca6\(^{3aQ68*–K352*}\) to the resistance phenotype of the laboratory strain JW-100s, we generated two isolines homozygous for each allele and we named them Q68* (Cca6\(^{3aQ68*}\)) and Q68*–K352* (Cca6\(^{3aQ68*–K352*}\)). When obtained, we characterized them molecularly to check whether they carried the expected genotypes.

We first sequenced the cDNA of Cca6 and found that Q68* isoline had transcripts where exon 3a was directly followed by exon 5, and transcripts with exon 2 followed by exon 5, indicating that exons 3b and 4 were not transcribed. In addition, we observed that when exons 3b and 4 were absent, a change in the reading frame of Cca6 occurred, provoking the appearance of a premature stop codon at the beginning of exon 5. Next, we observed that exons 3b and 4 were not present in cDNA of Q68* isolate, while intronic region downstream of exon 3a and upstream of exon 5 were present, indicating that a deletion must had occurred in between of these exons. Conversely, when we checked the sequence of both the gDNA and the cDNA of isolate Q68*–K352*, we found that all exons were present and transcribed. To understand whether the deletion of exons...
had occurred during the generation of Q68* isolate or whether it was already present in JW-100s, we checked the gDNA and cDNA of JW-100s individuals carrying the allele Ccα63aQ68* of generations F29 (30% of individuals homozygous Ccα63aQ68* /Ccα63aQ68*) and F85 (92% of individuals homozygous Ccα63aQ68*–K352*/Ccα63aQ68*–K352* and just 8% of individuals heterozygous Ccα63aQ68*/Ccα63aQ68*–K352*). We found that Ccα63aQ68* allele carried all exons in F29, while exons 3b and 4 were absent in F85, suggesting that the deletion happened at an unknown generation during the selection of JW-100s strain in the laboratory. Furthermore, qPCR analysis to test the expression of exons 3a and 3b in Q68* and Q68*–K352* isolines confirmed that exons 3b and 4 were not expressed in Q68* (Fig. 2). The results also indicated that expression of both 3a and 3b exons was significantly lower in Q68*–K352* isolate than in C strain (Fig. 2). The compendium of results of sequence and expression of Ccα6 allowed to elucidate that Ccα63aQ68* allele had lost exons 3b and 4 during the selection of JW-100s giving rise to a new allele Ccα63aQ68*Δ3b-4, which gives rise to isoforms that truncate at exon 3a, and to isoforms lacking exons 3b and 4 that truncate at exon 5 (Fig. 3). On the other hand, Q68*–K352* isolate kept the allele Ccα63aQ68*–K352* as originally was in JW-100s strain, though the expression of both 3a and 3b isoforms was reduced when compared to the C strain.

We performed bioassays on the isolines in order to compare the levels of resistance in them with that reported for JW-100s. Feeding spinosad at the concentration used in field treatments (260 ppm in bait sprays) did not cause mortality in either isolate. Feeding bioassays could not be performed at higher concentrations because of the limited solubility of spinosad in aqueous solutions. Topical bioassays with a range of doses of spinosad (Table 1) indicated that both Q68* and Q68*–K352* isolines had similar levels of resistance, with resistance ratios (RR) of 3139 and 3602, respectively, when compared to the susceptible C strain. These data

Fig. 1 Mortality percentages obtained from exposure to increasing spinosad concentrations of: a F1 transgenic Drosophila melanogaster flies coming from the cross between the Dα6 > GAL4 and the UAS-Ccα6 lines Ccα63a8b, Ccα63b8a, Ccα6Q68*, 786G and UAS-Dα6-3a3b (with a susceptible nACh6 transcript); and b parental UAS-Ccα6 lines of D. melanogaster flies. All flies used in the crosses were homozygous for the dα6* allele (no Dα6 expression). Different letters within each concentration indicate significant differences (ANOVA, Student–Newman–Keuls post hoc test, p ≤ 0.05. Data were previously arcsin transformed)
indicate that both alleles contribute equally to the resistant phenotype of JW-100s (RR = 1794, Ureña et al. 2019).

**Stability of spinosad-resistant alleles Cca6^{3aQ68*–K352*} and Cca6^{3aQ68*Δ3b–4} in C. capitata**

In order to study the stability of Cca6^{3aQ68*–K352*} and Cca6^{3aQ68*Δ3b–4} alleles in C. capitata, we performed crosses among Q68* and Q68*–K352* isolines and C strain to obtain individuals heterozygous for one of the resistant alleles and the wild-type Cca6^+ allele or heterozygous for both resistant alleles. We then followed the frequency of the different genotypes in the successive generations.

The fate of the different Cca6 alleles depended on the crosses performed (Table 2). In QC and KC crosses that carried only one of the resistant alleles (Cca6^{3aQ68*Δ3b–4} or Cca6^{3aQ68*–K352*}) and the wild-type allele (Cca6^+), the genotypic frequencies observed in G1 corresponded to those expected under Mendelian inheritance (25% homozygous wild-type, 50% heterozygous and 25% homozygous resistant). After this, we observed a rapid and continuous decline in the frequency of homozygous-resistant genotypes, not observed after G7. Heterozygous individuals remained at low frequency after nine generations. On the other hand, when both Cca6^{3aQ68*–K352*} and Cca6^{3aQ68*Δ3b–4} alleles coexisted in the QK cross, the typical frequencies of the Mendelian inheritance were maintained in Hardy–Weinberg equilibrium until the end of the experiment (G9), in both untreated and spinosad selected conditions. This result indicates that Cca6^{3aQ68*Δ3b–4} and Cca6^{3aQ68*–K352*} resistant alleles have a fitness cost compared to the wild-type allele, and that it is higher in homozygosis than in heterozygosis. However, no difference in fitness cost was observed between both strains.

**Fitness of Q68* and Q68*–K352* isolines of C. capitata**

We analyzed biological and behavioral traits in the Q68* and Q68*–K352* isolines of C. capitata and compared them to C strain in order to associate the existence of any fitness cost with the presence of a mutant allele conferring resistance.

**Biological traits**

We compared the embryo to pupal viability of the isolines and the C strain by analyzing the number of pupae obtained from a volume of 50 µl of eggs. No significant differences were observed (Table 3). The analysis of embryo to pupa development time showed that Q68* isolate developed significantly slower than C strain, with the Q68*–K352* isolate developing even slower (Table 3). We also checked if there were differences in adult weight between these genotypes. It was equivalent for females of all strains (Table 3), but males from the resistant Q68* and Q68*–K352* isolines were significantly heavier than those from the susceptible C strain.

**Behavioral traits**

We performed a wind tunnel assay to study the ability of male flies of C. capitata to get attracted by the synthetic pheromone, trimedlure. We saw that males from Q68* isolate had a lower response to trimedlure than males from C strain and Q68*–K352* isolate, when considered the percentage of males responding to trimedlure during an exposition of 30 min (Fig. 4a) and the time needed by the first fly to reach the pheromone (Fig. 4b). On the contrary, the ability of males of Q68*–K352* isolate to respond to the pheromone resulted similar to that of the wild type.

We also carried out an EAG assay to analyze whether males’ antennal receptors were able to detect the pheromone trimedlure. Among all the concentrations tested, we observed significant differences only with a dose of 5 ng of trimedlure, where the response in Q68*–K352* was higher than in the C males (Fig. 5). We did not observe any significant difference between Q68* and C strain at any of the concentrations tested.

We conducted competitive mating assays to find out whether there were differences in fecundation success when males of the different strain/isolines were put together with a group of females of a specific strain/isoline. We did not observe significant differences in the number of fecundated females when males belonged to C strain or Q68* isolate.
However, Q68*-K352* males were more successful in fecundating Q68*-K352* females than Q68* males (Fig. 6).

The results of the wind tunnel, the EAG and the mating assays indicate that males carrying the Cca6$^{3aQ68*\Delta3b-4}$ allele have impaired ability to detect the pheromone and to mate females carrying the Cca6$^{3aQ68*-K352*}$ allele.

**Discussion**

In a previous study, it was shown that medflies homozygous for Cca6$^{3aQ68*}$ allele were resistant to spinosad, despite generating wild-type full-length Cca6 isoforms with exon 3b (Ureña et al. 2019). Here, we have used the GAL4 > UAS expression system to introduce in D. melanogaster dα6 loss-of-function mutants three different Cca6 isoforms: (1) the isoform carrying 3aQ68* mutation, encoded by Cca6$^{3aQ68*}$ allele; (2) the wild-type mRNA isoform 3b8a, encoded by both Cca6$^{3aQ68*}$ and Cca6$^{+}$ alleles; and (3) the wild-type mRNA isoform 3a8b, encoded by Cca6$^{+}$ allele. As expected, the expression of the isoform carrying mutation 3aQ68* does not rescue susceptibility to spinosad, confirming that the truncated transcripts do not produce a response to spinosad (Baxter et al. 2010; Rinkevich et al. 2010; Hsu et al. 2012). On the contrary, we found that the expression of both wild-type Cca6$^{3b8a}$ and Cca6$^{3b8a}$ isoforms rescued spinosad

Fig. 3 Cca6 gene structure and its mRNA isoforms in: a C strain; b JW-100s strain at generation F29; c Q68* isolate and d Q68*-K352* isolines of Ceratitis capitata. Dark gray structures are truncated isoforms carrying stop codons, light gray means isoforms lacking some exons, and white refers to non-truncated isoforms. Sections a and b have been extracted from Ureña et al. (2019).
susceptibility in the α6-deficient strain, as already reported for Dα6 isoforms from susceptible D. melanogaster strains and of orthologues from susceptible strains of a variety of species (Perry et al. 2011, 2015; Anstead et al. 2015). Ureña et al. (2019) hypothesized that the absence of wild-type isoforms containing exon 3a in individuals homozygous for Ccaα63aQ68* allele could be enough to confer resistance to spinosad in medfly, since it has been suggested that some isoforms could be more responsive to spinosad or better able to form functional receptors than others (Perry et al. 2015). However, our results indicate that medfly 3b8a isoform is enough to mediate spinosad toxicity. We must keep in mind the fact that the GAL4 > UAS system usually confers higher expression levels than that observed under native conditions, which could mean that the susceptible phenotype observed could be due to 3b8a isoform being expressed at abnormally high, non-physiological levels. Nonetheless, we have found that UAS-Ccaα63b8a construct in the absence of GAL4 driver is sufficient to partially rescue the susceptibility to spinosad, which suggests that basal expression of both isoforms is enough for spinosad to be toxic, as observed previously (Perry et al. 2015). Alternative explanations for resistance conferred by Ccaα63aQ68* allele, despite the expression of susceptible 3b isoforms, have been proposed: i) that truncated 3a isoforms may interfere with the correct assembly of nAChR receptors in medfly and ii) the presence of undetected mutations in non-coding regions that could affect the transcript stability or the translation of full-length wild-type 3b isoforms (Ureña et al. 2019). Further studies are required to prove whether the 3b8a isoform encoded by the Ccaα63aQ68* allele has affected its functionality in medfly.

Remarkably, when we tried to obtain an isoline homozygous for the Ccaα66aQ68s allele, we realized that the resulting Q68* isolate was homozygous for a new Ccaα63aQ68sΔ3b-4 allele composed by isoforms that truncate at exon 3a and isoforms lacking exons 3b and 4 as a consequence of a deletion in the gDNA that truncate in exon 5 (Fig. 3). We confirmed that the allele Ccaα65aQ68s expressing full-length wild-type 3b isoforms was present in JW-100s individuals from generation F29, as already reported (Ureña et al. 2019). However, we found that heterozygous Ccaα65aQ68sΔ3b-4/Ccaα65aQ68sΔ3b-4 individuals of JW-100s strain from generation F85, about ten generations previous to the parents that were used to generate the Q68* isoline, already lacked exons 3b and 4. Thus, Ccaα65aQ68sΔ3b-4 allele might have appeared at an undetermined point during the selection of JW-100s with spinosad. It is probable that this new allele had been generated from Ccaα65aQ68s. Nevertheless, we cannot discard that Ccaα65aQ68sΔ3b-4 was already present at very low frequency in JW-100s. Regardless of its origin, it could have been selected instead of Ccaα65aQ68s because of better stability or lower fitness cost. Insect nAChR subunit genes have a complex process of splicing, and different isoforms are usually expressed in natural conditions (Sattelle et al. 2005). Indeed, the existence of nAChR transcripts lacking some exons appears to be common in insects (Jones et al. 2006). This includes α6 transcripts missing both forms of exon 3 that have been identified at low proportion in wild-type populations of Bombyx mori, Tribolium castaneum, T. absoluta and P. xylostella (Shao et al. 2007; Rinkevich and Scott 2009; Berger et al. 2016; Wang et al. 2016). Indeed, Berger et al. (2016) reported that spinosad resistance in T. absoluta was associated with altered nAChR α6 transcripts that lacked exon 3a/3b, though in this case, both exons were present and unaltered at the genomic level, indicating that their absence in mRNA results from exon skipping rather than a deletion in gDNA. It is not known whether nAChR transcripts with stop codons and/or exclusion of exons are translated, and if so, the functional role of the resulting proteins, though some regulatory and modulatory roles have been proposed (Sattelle et al. 2005). In any case, our

| Isoline/strain | N(1) | Slope ± S.E. | LD50(95%FL) | χ2(3) | D.f. | RR (95%FL)(4) |
|---------------|-----|-------------|------------|------|-----|---------------|
| JW-100s(5)   | 419 | 4.74 ± 0.61 | 185 (142–224) | 83.7 | 22  | 1794 (1156–2785)# |
| Q68*–K352*   | 390 | 2.26 ± 0.30 | 371 (284–483) | 35.3 | 21  | 3602 (2253–5757)# |
| Q68*         | 390 | 3.59 ± 0.45 | 323 (266–378) | 26.6*| 21  | 3139 (2000–4927)# |
| C            | 306 | 2.34 ± 0.67 | 0.10 (0.08–0.18) | 15.6*| 18  | 1             |

(1) Number of flies considered in the Probit analysis (including non-treated)
(2) Lethal dose (LD50) in µg/g of insect (fresh weight assuming an average weight of 10 mg) for topical assays at 48 h performed with spinosad 88% technical grade insecticide (a 0.5 µl drop of insecticide solution in acetone or acetone alone was applied to the dorsal thorax of each fly by using an automatic micropipettor)
(3) Good fit (*) of the data to the Probit model (P > 0.05)
(4) Resistance ratio (RR) = [LD50 (selected strain)/LD 50 (C strain)]. The fiducial limits for RR were calculated according to Robertson and Preisler (1992). *RR is significant (P < 0.05) if the 95% FL does not include 1
(5) Data from Ureña et al. (2019)
Table 2 Genotype frequencies (%) in the progeny of the crosses between Q68* and Q68*–K352* isolines and C strain of Ceratitis capitata

| Crosses(1) | Generation | Genotypes (X± SE)(2) | Fit the model (Chi-square test)(3) |
|------------|------------|----------------------|-----------------------------------|
| Q68* × Q68*–K352* (QK) | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ |
| QKnt | 0 ± 0 | 100 ± 0 | 0 ± 0 | * |
| G0 | 40.8 ± 3.8 | 50.9 ± 4.7 | 8.3 ± 8.3 | |
| G1 | 16.7 ± 4.2 | 66.7 ± 4.2 | 16.7 ± 4.2 | * |
| G5 | 46.7 ± 3.3 | 33.3 ± 8.8 | 20 ± 11.6 | |
| G7 | 20 ± 5.8 | 46.7 ± 3.3 | 33.3 ± 3.3 | * |
| G9 | 30 ± 5.8 | 56.7 ± 12 | 13.3 ± 6.7 | * |
| QKs | 25 ± 12.5 | 58.3 ± 18.2 | 16.7 ± 8.3 | * |
| G3 | 23.3 ± 3.3 | 36.7 ± 8.8 | 40 ± 5.8 | * |
| G5 | 40 ± 10 | 36.7 ± 6.7 | 23.3 ± 14.5 | * |
| G7 | 40 ± 10 | 46.7 ± 8.8 | 13.3 ± 3.3 | * |
| Q68* × C (QC) | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ |
| G0 | 0 ± 0 | 100 ± 0 | 0 ± 0 | |
| G1 | 34.4 ± 3.6 | 52.6 ± 4.6 | 13 ± 5 | |
| G3 | 54.2 ± 10.5 | 45.8 ± 10.5 | 0 ± 0 | |
| G5 | 55.6 ± 13.6 | 39.7 ± 8.8 | 4.8 ± 4.8 | |
| G7 | 75.4 ± 17 | 24.6 ± 17 | 0 ± 0 | |
| G9 | 70.6 ± 7.6 | 29.4 ± 7.6 | 0 ± 0 | |
| Q68*–K352* × C (KC) | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ | Ccα6+/Ccα6+ |
| G0 | 0 ± 0 | 100 ± 0 | 0 ± 0 | |
| G1 | 23.3 ± 18.6 | 60 ± 17.3 | 16.7 ± 12 | |
| G3 | 58.3 ± 15 | 33.3 ± 18.2 | 8.3 ± 4.2 | |
| G5 | 52.2 ± 7.8 | 42.2 ± 2.2 | 5.6 ± 5.6 | |
| G7 | 75 ± 25 | 25 ± 25 | 0 | |
| G9 | 80 ± 10 | 20 ± 10 | 0 | |

(1) Individuals of the Q68* and Q68*–K352* isolines were crossed among them (QK) and with individuals of the C strain (QC and KC, respectively). Both QC and KC were maintained without selection pressure for nine generations. QK was split at G2 in two different populations: one maintained without selection pressure (QKnt) and the other one selected with spinosad (QKs).
(2) There were three replicates of each cross. Between 20 and 40 flies of each cross and generation were genotyped.
(3) Good fit (*) of data to the expected genotypic frequencies (0, 100 and 0% in G0, and 25, 50 and 25% for all the other generations for first homozygous, heterozygous and second homozygous, respectively). Chi-square test was performed for data analysis (p > 0.05).

Table 3 Biological parameters of adult flies from C strain and Q68* and Q68*–K352* isolines of Ceratitis capitata

| Parameter | C | Q68* | Q68*–K352* |
|-----------|----|------|------------|
| Number of pupae(1) | 878 ± 102 (a) | 755 ± 52 (a) | 909 ± 45 (a) |
| Days from egg to pupae(2) | 8.77 ± 0.01 (a) | 9.3 ± 0.01 (b) | 9.46 ± 0.01 (c) |
| Females’ weight (g)(3) | 8.27 ± 0.14 (a) | 8.6 ± 0.15 (a) | 8.5 ± 0.15 (a) |
| Males’ weight (g)(3) | 7.34 ± 0.11 (a) | 8.29 ± 0.15 (b) | 7.95 ± 0.13 (b) |

(1) Number of pupae obtained from a volume of eggs of 50 µl. Three replicates were performed with each strain/isoline.
(2) n = number of pupae obtained for each strain/isoline in the table.
(3) n = 50 adults 2–3 days old.

Data are mean ± standard error. Different letters within each row indicate significant differences (ANOVA, Student–Newman–Keuls post hoc test, p ≤ 0.05).

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results and those from other studies highlight the plasticity of nAChR α6 to generate a tremendous diversity of isoforms, including truncated isoforms potentially unaffected by spinosad. An important factor that will determine the progress of spinosad resistance in field populations is the fitness cost of resistance. We found that both $Cca\alpha6^{\Delta q68^*\Delta3b-4}$ and $Cca\alpha6^{\Delta q68^-K352^*}$ alleles were not stable in the absence of spinosad when individuals from isolines homozygous for these two alleles were crossed with individuals carrying the wild-type allele $Cca\alpha6^+$. Our results indicate that both resistant alleles may impart a competitive disadvantage with respect to the wild-type allele. However, no differences among the stability of $Cca\alpha6^{\Delta q68^*\Delta3b-4}$ and $Cca\alpha6^{\Delta q68^-K352^*}$ alleles were perceived when they competed between them, both in the presence or absence of insecticide. Besides, both resistant alleles conferred similar levels of spinosad resistance, which indicated that differences in susceptibility did not play a relevant role in the competition between resistant alleles in the presence of the insecticide. Fitness cost associated with spinosad resistance has been reported to range from minor (Wyss et al. 2003) or absent (Bielza et al. 2008a; Abbas et al. 2014) to substantial (Li et al. 2007, 2017; Sayyed et al. 2008; Wang et al. 2010; Rehan and Freed 2015; Okuma et al. 2018). However, the genetic mechanism underlying spinosad resistance was only known in two of these studies, and in both cases, it was associated with

Fig. 4 Wind tunnel assay with males of C strain and Q68* and Q68*–K352* isolines of Ceratitis capitata; a percentage of males that respond to trimedlure during 30 min; b time needed by the first fly of each strain/isoline to reach the trimedlure source. Data are mean ± standard error of four replicates with 50 adult males in each one. Different letters within each row indicate significant differences (repeated measures ANOVA in male response during 30 min and ANOVA in first fly attracted, in both cases followed by Student–Newman–Keuls post hoc test, $p \leq 0.05$)

Fig. 5 Electroantennography response (mV) of males of C strain and Q68* and Q68*–K352* isolines of Ceratitis capitata when exposed to increasing doses of trimedlure. Data are mean ± standard error of 5–15 replicates. Different letters within each amount of trimedlure indicate significant differences (ANOVA, Student–Newman–Keuls post hoc test, $p \leq 0.05$)

Fig. 6 Number of males of C strain and Q68* and Q68*–K352* isolines of Ceratitis capitata that, under competitive conditions, mated with females of the three strain/isolines. Data are mean ± standard error of five replicates with ten females of a specific strain/isoline and 12 males of all strain/isolines. Different letters within each group indicate significant differences (ANOVA, Student–Newman–Keuls post hoc test, $p \leq 0.05$)
overexpression of detoxifying enzymes (Sayyed et al. 2008; Rehan and Freed 2014). In order to identify trade-offs associated with spinosad resistance in C. capitata, extensive testing of both biological and behavioral traits was undertaken. We found that the effect on the biological traits tested was moderate, with Q68* (Ccα6Δ3aQ68Δ3b–K352*) and Q68*–K352* (Ccα6Δ3aQ68Δ3b–K352*/Ccα6Δ3aQ68Δ3b–K352*) isolines showing longer developmental time but higher male weight than C strain, whereas females of both resistant isolines and susceptible C strain did not differ in weight. However, mutations that would lead to a loss-of-function phenotype may affect behavioral patterns, as it has been reported for a Dα1 knockout mutant with a significant fitness cost in terms of severe mating behavior defects and an altered sleep phenotype (Somers et al. 2017). Thus, we explored the possibility that behavioral fitness traits (parapheromone detection by wind tunnel and electrophysiological antennal assays, and mating success) could be affected in males of the Q68* and Q68*–K352* isolines. We found that males from Q68* isoline, homozygous for Ccα6Δ3aQ68Δ3b–K352* allele, had impaired ability to find the parapheromone source in a wind tunnel and to mate females, though EAG assays demonstrated that this impediment was not due to a failure in the response of receptors in the antennae. This could point to a failure in the flight tools to properly move inside the tunnel, or to a difficulty to orient themselves when they receive a stimulus, though further studies would be needed to confirm these hypotheses. A study focused on the competitive mating of a spinosad-resistant strain of H. virescens concluded that resistant adults must have a reproductive disadvantage, but they were not able to discern whether it was due to differences in mating success or in fecundity (Wyss et al. 2003). On the contrary, a homozygous Ccα6Δ3aQ68Δ3b–K352* allele in the Q68*–K352* isolate does not imply a handicap for the male’s ability to detect the parapheromone and to mate with females when compared to males of the C strain. Therefore, other fitness traits may be affected by the Ccα6Δ3aQ68Δ3b–K352* allele that make it unstable when in competition with the wild-type allele, as it is possible that different isoforms generated by each resistant allele may interfere differently with the normal function of the receptor and with other nAChR subunits that α6 may also co-assemble with.

Our results highlight the plasticity of C. capitata to generate a wide range of mutations in nAChR α6 that can lead to loss-of-function phenotypes potentially conferring spinosad resistance. However, these resistant alleles have a fitness cost that makes them unstable when in competition with wild-type alleles, which could be the reason for the absence of resistance events in the field (Ureña et al. 2019). In addition, the rate of resistance evolution in field populations is expected to be slow, due to the completely recessive nature of spinosad resistance in this species (Ureña et al. 2019). Nonetheless, once resistance evolved, it would be expected to increase to fixation if the selection pressure eliminated all susceptible insects. Moreover, the selection of compensatory mutations that ameliorate the fitness cost of existing resistant genotypes or the selection of less costly new resistant alleles could accelerate the evolution of resistance (Raymond et al. 2001; ffrench-Constant and Bass 2017). Thus, insect resistance management strategies should be implemented to ensure the sustainability of spinosad.

Author contribution statement

FO, PHC and LS conceived the study. FO, PHC, AGA, EU, LS, TP, PB, VN and ELE participated in the design of the experiments and the interpretation of the results. AGA, EU and ELE performed the experiments. AGA and FO wrote the first draft of the manuscript. All authors read, corrected and approved the manuscript.

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Compliance with ethical standards

Conflict of interest The authors declare that they have no conflict of interest.

Ethical approval All applicable international, national and/or institutional guidelines for the care and use of animals were followed.

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