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A single nucleotide polymorphism in the acetylcholinesterase gene of the predatory mite *Kampimodromus aberrans* (Acari: Phytoseiidae) is associated with chlorpyrifos resistance / Cassanelli, Stefano; Ahmad, Shakeel; Duso, Carlo; Tirello, Paola; Pozzebon, Alberto. - In: *BIOLOGICAL CONTROL*. - ISSN 1049-9644. - STAMPA. - 90:(2015), pp. 75-82. [[10.1016/j.biocontrol.2015.05.015](https://doi.org/10.1016/j.biocontrol.2015.05.015)]

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22/09/2024 05:12

(Article begins on next page)

1 **A single nucleotide polymorphism in the acetylcholinesterase gene of**  
2 **the predatory mite *Kampimodromus aberrans* (Acari Phytoseiidae) is**  
3 **associated with chlorpyrifos resistance**

4  
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16  
17 **ABSTRACT**

18  
19 The predatory mite *Kampimodromus aberrans* (Oudemans) (Acari Phytoseiidae) is one of  
20 the most important biocontrol agents for herbivorous mites in European perennial crops.  
21 The use of pesticides, such as organophosphate insecticides (OP), is a major threat to the  
22 success of biocontrol strategies based on predatory mites in these cropping systems.  
23 However, resistance to OP in *K. aberrans* has recently been reported. The present study  
24 investigated the target site resistance mechanisms that are potentially involved in OP  
25 insensitivity. In the herbivorous mite *Tetranychus urticae*, resistance to OP is due to a  
26 modified and insensitive acetylcholinesterase (AChE) that bears amino acid substitution  
27 F331W (AChE Torpedo numbering). To determine whether the predators and prey had  
28 developed analogous molecular mechanisms to withstand the same selective pressure,  
29 the AChE cDNA from a putative orthologous gene was cloned and sequenced from  
30 susceptible and resistant strains of *K. aberrans*. No synonymous mutation coding for a  
31 G119S substitution was determined to be strongly associated with the resistant phenotype  
32 instead of the alternative F331W. Because the same mutation in *T. urticae* AChE was not  
33 associated with comparable levels of chlorpyrifos resistance, the role of the G119S  
34 substitution in defining insensitive AChE in *K. aberrans* remains unclear. G119S AChE  
35 genotyping can be useful in ecological studies that trace the fate of resistant strains after

36 field release or in marker-assisted selection of improved populations of *K. aberrans* to  
37 achieve multiple resistance phenotypes through gene pyramiding. The latent complexity of  
38 the target site resistance in *K. aberrans* vs. that of *T. urticae* is also discussed by exploiting  
39 data from the genome project of the predatory mite *Metaseiulus occidentalis* (Nesbitt).

40

## 41 **1. Introduction**

42 *Kampimodromus aberrans* (Oudemans) (Acari: Phytoseiidae) is a predatory mite that  
43 occurs in various European cropping systems, such as grapevines, apples and hazelnuts  
44 (Ivancich Gambaro, 1973; El Borolossy and Fischer-Colbrie, 1989; Tsolakis et al., 2000;  
45 Ozman-Sullivan, 2006). This predatory mite is also common on the uncultivated plants that  
46 surround crops and represent a potential reservoir for biocontrol agents (Tixier et al., 2002,  
47 2006). *K. aberrans* is considered to be generalist predator (McMurtry and Croft, 1997;  
48 Kreiter et al., 2002; Broufas et al., 2007; Lorenzon et al., 2012) and an effective biocontrol  
49 agent of tetranychid and eriophyoid mites in European vineyards (Duso 1989; Girolami et  
50 al., 1992; Duso and Pasqualetto, 1993; Duso et al., 2012). In addition to several ecological  
51 factors, insecticide and fungicide applications strongly affect naturally occurring and  
52 artificially introduced *K. aberrans* populations (Ivancich Gambaro 1973; Girolami, 1987;  
53 Pozzebon et al., 2002; Peverieri et al., 2009). However, a *K. aberrans* strain has been  
54 detected in North Italian vineyards under conditions of integrated pest management  
55 strategies (IPM) that rely on common ethylene-bis-dithiocarbamate (EBCD) fungicides and  
56 organophosphate (OP) insecticides (Posenato 1994). This strain was also successfully  
57 released in other vineyards and apple orchards following organic or IPM strategies (Duso  
58 et al., 2007; Duso et al., 2009; Ahmad et al., 2013). Recently, laboratory studies have  
59 confirmed significant levels of chlorpyrifos resistance in this same strain (Tirello et al.,  
60 2012). The biochemical basis of OP resistance in phytoseiid mites depends on the active  
61 ingredients involved in the selective pressure and on species/strain-specific genetic  
62 backgrounds. The resistance phenotype might rely on high detoxifying enzyme activities  
63 and/or on a modified and insensitive target AChE. For example, laboratory selection with  
64 methidathion in *Amblyseius womersleyi* (Schicha) leads to increases in monooxygenase  
65 activity and CYP4-d isoform overexpression (Sato et al., 2001, 2006, 2007), while in  
66 *Phytoseiulus persimilis* (Athias-Henriot), this selection results in an enhancement of  
67 glutathione transferase (Fournier et al., 1987). High rate *in-vitro* degradation of azinphos-  
68 methyl has been observed under both polygenic and monogenic control in OP-resistant  
69 strains of *Amblyseius fallacis* (Garman) (Motoyama et al., 1971; Croft et al., 1976). Target

70 site resistance to OP has also been detected biochemically either in isolation or  
71 combination with enhanced OP detoxifying pathways. Resistance to certain OP and  
72 carbamate compounds, such as parathion and propoxur, in a Dutch strain of  
73 *Typhlodromus pyri* Scheuten has been found to be under monogenic control and to be  
74 associated with an insensitive target AChE (Overmeer and van Zon, 1983). In a paraoxon-  
75 resistant strain of *Amblyseius andersoni*, (Chant) the resistant phenotype has been  
76 revealed to be due to an insensitive AChE coupled with modified carboxylesterases (Anber  
77 et al., 1988, 1989).

78 Although reductions in chlorpyrifos susceptibility have been reported in other predatory  
79 mites, e.g., *T. pyri* (Fitzgerald and Solomon 1999; Cross and Berrie 1994; Bonafos et al.,  
80 2008), little is known about the underlying molecular mechanisms. Among the Acari, high  
81 levels of chlorpyrifos resistance in *Tetranychus urticae* Koch have been found to be due to  
82 a F331W amino acid substitution in the target enzyme acetylcholinesterase (AChE)  
83 (Khajehali et al., 2010). Knowledge of a genetic marker associated with chlorpyrifos  
84 insensitivity in *K. aberrans* could be useful for understanding the amplitude of this  
85 phenomenon and managing predatory mite populations with IPM strategies. Therefore, we  
86 report the cloning and sequencing of a *T. urticae*-like acetylcholinesterase cDNA in *K.*  
87 *aberrans* and its genotyping in chlorpyrifos-susceptible and resistant strains. The potential  
88 complexity of the target site resistance that occurs in predatory mites was also inferred by  
89 inspecting the annotated genome of *M. occidentalis*.

90

## 91 **2. Materials and Methods**

92

### 93 *2.1. Kampimodromus aberrans* populations

94

95 This study was performed on seven *K. aberrans* strains collected in North-eastern Italy  
96 (Veneto Region). Four strains were collected from commercial vineyards, and three strains  
97 were collected from untreated European nettle trees (*Celtis australis* L.) (Table 1).

98 All strains were reared without insecticide exposure in separate rearing units at the  
99 Department of Agronomy, Food, Natural Resources, Animals and the Environment of the  
100 University of Padova, Italy. Grapevine leaves on pads of wet cotton were used as a  
101 substrate for the predatory mites, and small pieces of PVC were placed for shelter and  
102 oviposition. *Typha latifolia* L. pollen was provided as food (Lorenzon et al., 2012).

103 Information about the effects of OP was available for only two strains; specifically, the PO  
104 strain is resistant to chlorpyrifos, and the LE strain is highly susceptible to this insecticide  
105 (Tirello et al., 2012).

106

## 107 *2.2. Insecticide bioassays*

108

109 Laboratory bioassays were conducted for the preliminary screening for resistant and  
110 susceptible phenotypes. The bioassays were performed with a commercial formulate  
111 (Dursban® 75WG, 75% a.i., Dow AgroSciences). The discriminant concentration for the  
112 resistant and susceptible phenotypes was set at 70 g/hl of formulate, which is the  
113 recommended field dose for use in vineyards against grape berry moths and leafhoppers.  
114 The pesticide formulate was diluted in distilled water before the toxicological test  
115 procedures (Tirello et al., 2013). The latter procedures were performed using rectangular  
116 leaf sections (approximately 6 cm<sup>2</sup>). The sections were immersed in the insecticide  
117 solution for 30 s, and distilled water was used in the control treatments. When the  
118 pesticide residues completely dried out, the leaf sections were placed on wet cotton pads,  
119 and cotton barriers were created along their perimeters to prevent predatory mite escape.  
120 Two 12-d-old *K. aberrans* females were gently transferred to each leaf section, and fresh  
121 pollen was provided as food. The experimental units were maintained in a climate  
122 chamber at 25 ± 2° C and 70 ± 10% relative humidity with a 16L:8D photoperiod. Female  
123 mortality was assessed 72 h after the treatments. The females that drowned or escaped  
124 were removed from the initial test number. In total, we assessed 40-45 females per strain.  
125 The corrected mortalities (Abbott, 1925) were calculated.

126

## 127 *2.3. Primer design for cloning AChE cDNA in K. aberrans*

128

129 The annotated version of the genome assembly (release Mocc\_1.0, March 2012) of the  
130 predatory mite *Metaseiulus occidentalis* (Nesbitt) (WOPM genome project) was used to  
131 search for putative AChE-like proteins with the tBlastn algorithm using the AChE sequence  
132 that was amplified from the susceptible strain of *T. urticae* (GenBank accession n.  
133 ADK12697.1) as the query sequence.

134 Transcripts predicted to code for putative AChE-like proteins were extracted from the  
135 scaffolds, and their open reading frames (ORFs) were compared to the *T. urticae* AChE

136 protein using Lasergene sequence analysis tools EditSeq and MegAlign 5.0 (DNASTAR,  
137 Inc., Madison, WI, USA).

138 Degenerate primers were designed by manual inspection of the conserved domains after  
139 the alignments of *T. urticae* AChE and putative orthologous AChE-like proteins in *M.*  
140 *occidentalis*. The resulting primers were used to amplify the cDNA core fragments of the  
141 orthologous AChE in *K. aberrans*. To complete the cloning, walking steps and 3'-5' RACEs,  
142 were performed using no degenerate primers and outlined with PrimerSelect 5.0  
143 (DNASTAR, Inc., Madison, WI, USA).

144

#### 145 2.4. mRNA extraction and AChE cDNA cloning

146

147 Total RNA was extracted by homogenising 200 adults in 500 µl Tri-Reagent (Sigma),  
148 according to the manufacturer's instructions. The sample integrities were examined by  
149 electrophoresis in 1.2% agarose and 2.2 M formamide/formaldehyde denaturing gel.  
150 Quality and quantity assessments of the extracted RNA were performed in a Nanodrop  
151 ND-1000 Spectrophotometer (NanoDrop, Fisher Thermo, Wilmington, DE, USA). First-  
152 strand cDNA was synthesised according to the protocol recommended by the supplier  
153 using Improm-II reverse transcriptase (Promega) and random primers. Amplification of a  
154 cDNA fragment for a putative AChE in *K. aberrans* was achieved through two consecutive  
155 rounds of reverse-transcription PCR (RT-PCR) with degenerate primers. The PCR  
156 mixtures (25 µl) contained GoTaq Flexi 1x buffer, 1.5 mM MgCl<sub>2</sub>, 0.2 mM dNTPs, 30 pmol  
157 forward and reverse degenerate primers, 1 U GoTaq and 2.5 µl of cDNA. The degenerate  
158 primers were based on partially conserved functional domains of homologous AChEs  
159 proteins in *T. urticae* and *M. occidentalis*. For the first RT-PCR round, the forward and  
160 reverse primers were KaAChEF1d (GIPYAKP domain) and KaAChER1d or KaAChER2d  
161 (WVYGGSF motif) (Table 2). The PCR product was then diluted 10-fold and used as the  
162 template for a second RT-PCR in which the KaAChEF1 primer was replaced with the more  
163 internal primer KaAChEF2d, which was designed based on the PYAKPP domain (Table 2).  
164 The two PCR rounds shared the following profile: an initial denaturation step of 3 min at  
165 94 °C; 5 cycles at 94 °C for 30 s, 45 °C for 30 s and 72°C for 60 s; 5 cycles at 94 °C for 30  
166 s, 45 °C plus +1 °C/cycle and 72°C for 60 s; 25 cycles at 94 °C for 30 s, 50 °C for 30 s and  
167 72°C for 60 s; and a final extension step at 72 °C for 10 min. PCR products of the  
168 expected size (approximately 300 bp) were purified from 1% (w/v) agarose/TBE 1x gel  
169 using a EuroGOLD Gel Extraction Kit (Euroclone) and cloned using a pGEM-T easy vector

170 (Promega). The plasmids were purified with a EuroGOLD Plasmid Miniprep Kit  
171 (Euroclone) and sent for sequencing at BMR genomics (Padua, Italy). The sequences  
172 were assembled and analysed using SeqMan 5.0 (DNASTAR, Inc., Madison, WI, USA).  
173 Identification of the AChE-like sequences was performed via a BLASTX search in  
174 GenBank (<http://www.ncbi.nlm.nih.gov>) using the ORFs deduced from the cloned cDNA  
175 fragments. The cDNA clones were further extended in the 3' direction by performing an  
176 RT-PCR that used a forward primer that was designed based on the first cloned cDNA  
177 fragment in *K. aberrans* (KaAChEF3) and a reverse primer (KaAChER3) that was  
178 designed based on the sequence coding for the conserved domain CAFWKNFL in both of  
179 the AChE transcripts found in *M. occidentalis* without any primer degeneration (Table 2).  
180 The RT-PCR mixture had the same composition described above except that the primer  
181 concentration was reduced to 15 pmol. The PCR was performed as follows: 1 cycle of  
182 94 °C for 2 min; 5 cycles that included the three steps of 94°C for 30 s, 50 °C for 30 s and  
183 72 °C for 60 s; 5 cycles of 94 °C for 30 s, 50 °C for 30 s (+ 1 °C/cycle) and 72 °C for 60 s;  
184 20 cycles of 94°C for 30 s, 55 °C for 30 s and 72 °C for 60 s; and a final extension step at  
185 72 °C for 10 min. The PCR product was purified, sequenced and analysed as described  
186 above. Three prime and 5' rapid amplification of cDNA ends reactions (RACEs) were  
187 performed to complete the *AChE* cDNA sequences. In the RACE reactions, the first strand  
188 cDNAs were synthesised using total RNA and polyT-adaptor primer for 3' RACE or  
189 KaAChE-R4 for 5' RACE (Table 2) according to the manufacturer's protocol (5' RACE  
190 System for Rapid Amplification of cDNA Ends, Invitrogen). The 3' RACE product spanning  
191 across the unknown 3'-end of the *AChE* cDNA was amplified in two consecutive PCR  
192 rounds with the KaAChEF4-Adaptor1 and KaAChEF5-Adaptor2 primer pairs. To obtain  
193 the 5' end of the AChE transcript, the cDNA was subjected to polyC-tailing of its 3'-end  
194 with terminal deoxynucleotidyl-transferase (TdT) following the protocol of the kit (5' RACE  
195 System for Rapid Amplification of cDNA Ends, Invitrogen). The upstream cDNA sequence  
196 encompassing the 5' untranslated region was amplified with two PCR rounds using the  
197 coupled primers KaAChER5-TS-primer and KaAChER6-TS-PCR (Table 2). The first 5'  
198 RACE round was performed as follows: 94 °C for 2 min (1 cycle); 5 cycles at 94 °C, 56 °C  
199 for 30 s and 72 °C for 60 s; 5 cycles at 94 °C, 57 °C for 30 s and 72 °C for 60 s; and 20  
200 cycles at 94 °C for 30 s, 58 °C for 20 s and 72 °C for 60 s. The second 5' RACE round  
201 consisted of the following: 1 cycle at 94 °C for 2 min; and 30 cycles at 94 °C for 30 s,  
202 55 °C for 30 s and 72 °C for 60 s. The 5' RACE fragment was purified from the agarose gel  
203 and sequenced as previously described.

204

## 205 *2.5. Full length AChE cDNA sequencing*

206

207 Total RNA was extracted from adults of both sensitive (LE) and resistant (PO) strains with  
208 TRI-Reagent described for the cDNA cloning. First-strand cDNA was synthesised from  
209 total RNA with Improm-II reverse transcriptase (Promega) and random primers as  
210 indicated by manufacturer's protocol. To sequence the ORF of the cloned cDNA, three RT-  
211 PCR fragments that partially overlapped were generated using the following primer  
212 couples: KaAChEF6-R7, KaAChEF7-R8, and KaAChEF8-R9 (Table 2). The PCR reaction  
213 (25 µl) included 2 µl of cDNA, a final concentration of GoTaq Flexi 1x buffer, 1.5 mM MgCl<sub>2</sub>,  
214 0.2 mM dNTPs, 0.6 µM of each primer and 0.625 U/µl GoTaq (Promega). The thermal  
215 profile adopted was as follows: 94°C for 2 min (1 cycle); 30 cycles of 94 °C for 30 s, 56 °C  
216 for 30 s and 72 °C 60 s; and a final extension step at 72 °C for 10 min. The PCR products  
217 were checked by electrophoresis on 1% agarose in TBE 0.5x buffer, purified with the  
218 EuroGOLD Cycle-Pure Kit (Euroclone) and sent to BMR genomics (Padua, Italy) for  
219 sequencing. To this aim, the same primers used for the RT-PCR amplifications and new  
220 internal primers (KaAChEF9, R10, and F10) were used (Table 2). Chromatograms were  
221 assembled with SeqMan tools (DNASTAR, Lasergene), and the alignments of the cDNA  
222 consensus sequences from sensitive and resistant strains were manually inspected for  
223 non-synonymous SNPs with the MegAlign program (DNASTAR, Lasergene).

224

## 225 *2.6. DNA extraction and exon-intron junction amplification*

226

227 DNA extraction was performed according to the methods described by Tixier et al. (2008)  
228 while scaling up the reagents. Two hundred frozen adults of each strain were  
229 homogenised in 150 µl of extraction buffer (2% CTAB, 1.4 M NaCl, 0.2% 2-mercaptoethanol,  
230 100 mM EDTA, and 100 mM Tris-HCl at pH 8.0) using a micro tissue grinder (Wheaton,  
231 Millville, NJ). The homogenate was transferred to a 0.5-ml test tube and incubated for one  
232 hour at 65°C with periodic hand mixing. One hundred and fifty microliters of a  
233 chloroform:isoamyl alcohol mixture (24:1) was added, the solution was mixed by inversion,  
234 and tubes were centrifuged at 6°C for 5 min at 1000 g. The aqueous solution was collected  
235 in a new test tube, and 80 µl of isopropanol was added to the decanted aqueous phase,  
236 which was then chilled at -20 °C for 20 min for DNA precipitation. After centrifugation (15  
237 min, 6 °C, 1000 g), the pellet was suspended in 100 µl of 96% alcohol at 4 °C. After a final

238 centrifugation of 10 min (6 °C, 1000 g), the dried pellet was suspended in 30 µl of de-  
239 ionised water. The quality and quantity of the extracted DNA were assayed by  
240 spectrophotometric analyses with a Nanodrop ND-1000, and the integrities were verified  
241 through electrophoresis on 1% agarose/TBE 0.5x gel. Exon-intron boundary predictions  
242 were made by aligning the AChE cDNA sequence cloned in *K. aberrans* with the scaffold  
243 form the *M. occidentalis* genome project from which the transcript XR\_145413 had been  
244 predicted (Genbank accession n. AFFJ01003151.1). Relying on hypothetical gene  
245 structure conservation, the primers were designed on the exon sequences to generate  
246 partially overlapping PCR fragments that encompassed the putative introns in the *K.*  
247 *aberrans* AChE gene. The PCR products were purified and sequenced as described for  
248 the cDNA sequencing using the same primers that were employed for the DNA  
249 amplifications.

250

### 251 **3. Results**

252

#### 253 *3.1. Insecticide bioassays*

254

255 Laboratory trials confirmed the findings reported by Tirello et al. (2012). At the discriminant  
256 dose, 100% corrected mortalities were observed for the LE, PA and PD strains, which  
257 originated from untreated nettles, and high survival rates were observed for the PO, ME,  
258 SF and BX strains (4.08%, 9.57%, 7.69% and 4.26% corrected mortalities, respectively),  
259 which were collected from commercial vineyards.

260

#### 261 *3.2. T. urticae AChE-like gene in the M. occidentalis genome*

262

263 The tBlastn search on the annotated genome of the predator mite *M. occidentalis* using  
264 the AChE cloned from *T. urticae* as the query sequence resulted in two predicted mRNAs  
265 that codes for putative AChEs and had sequences that were significant similar to that of  
266 the query (XR\_145413 and XR\_145279; identity 54%, positive 69%, e-value 0.0). These  
267 mRNAs originated from genes in partially overlapping contigs (AFFJ01003151 and  
268 AFFJ01002402). The corresponding open reading frames differed primarily in the amino  
269 terminal due to a diverse prediction of the first splicing site, while they shared the  
270 remaining five, which resulted in only 8 mismatches out of the 593 conserved amino acid  
271 residues. These mismatches arose from indels in the coding regions of the two genes,

272 which did not differ in the intronic sequences with the exception of the first intron that  
273 originated from alternative splicing paths. Because the algorithms used for automatic  
274 splicing site predictions often fail to identify splicing sites at the 5' end of putative  
275 transcripts and because of the low level of sequence divergence between the two genes, it  
276 was unclear whether there were two copies per genome or if they were derived from *in*  
277 *silico* mis-assembling of the high-throughput sequencing reads. In any case, when the *M.*  
278 *occidentalis* transcriptome shotgun assembly was interrogated with Blastn with the two  
279 putative transcripts, a pair of cDNA fragments were retrieved that covered both mRNAs  
280 (JL046593.1 and JL050556.1; identities 99% and 98%), which confirmed that they were  
281 actually transcribed. Altogether, these findings suggested that the two very similar  
282 predicted mRNAs could be informative for cloning *T. urticae*-like AChE cDNA in *K.*  
283 *aberrans*.

284

### 285 3.3. AChE cDNA in the susceptible strains of *K. aberrans*

286

287 cDNA of 2329 was isolated from the susceptible LE strain (Genbank accession number:  
288 HF934042). The deduced precursor was composed of 655 amino acids (Fig. 1) with a  
289 signal peptide that was predicted to encompass the first 32 amino acids from the amino  
290 terminal (Shen et al. 2007). The cloned KaAChE displayed most of the amino acids  
291 responsible for the functional integrity of the enzyme that are typically well conserved both  
292 in insect and mite AChEs; i.e., the KaAChE residues involved in the intramolecular  
293 disulphide bonds (C139, C166, C325, C336, C471, and C593), the catalytic triad (S271,  
294 E395, and H509), the anionic subsite (W156), the oxianion-hole (G189, G190, and A274),  
295 and the acyl pocket (W304, F360, and F399) (Fig. 1). The highest identity (> 93%) was  
296 observed for the AChE that was annotated in *M. occidentalis* from the transcript  
297 XR\_145413 because the first splicing path was consistent with that predicted in this  
298 putative mRNA. No alternative cDNA sequence similar to the *M. occidentalis* transcript  
299 XR\_145279 was detected in *K. aberrans*. As expected, the greatest divergences in the  
300 amino acid sequences between the KaAChE and XR\_145413 predicted AChEs were  
301 restricted to the amino and carboxy terminals of the protein outside of the functional  
302 domains. The amino acid identities with the other cloned and predicted AChEs in the Acari  
303 genomes that carry multiple AChE loci ranged from 61% (*Ixodes scapularis* putative AChE,  
304 XP\_002413212) to 33% (*Rhipicephalus microplus*, AChE3, AAP92139). The amino acid  
305 identity was 52% between the AChEs coded by single copy genes in the *T. urticae* and *T.*

306 *evansi* that carry mutations associated with reduced chlorpyrifos sensitivity (GQ461344,  
307 ADK12694, and AFS60097) This divergence was compatible with that observed in the  
308 AChEs from different species of Acari and even between AChEs from multiple loci in the *I.*  
309 *scapularis* or *R. microplus* genomes. AChEs of insects are divided in two groups, i.e.,  
310 those orthologous and those paralogous to the *D. melanogaster* AChE (Kim et al., 2012),  
311 and KaAChE exhibited a high level of similarity to the paralogous AChEs found in  
312 *Nephotettix cincticeps* (Hemiptera: Deltocephalidae) and *Blattella germanica* (Blattodea:  
313 Blattellidae) with an amino acid identity of approximately 57% (ADZ15146; ABB89946).

314

#### 315 3.4. Organisation of the clone AChE locus in *K. aberrans*

316

317 The intron-spanning amplifications of the *K. aberrans* AChE locus confirmed the exon-  
318 intron junctions that were predicted *in silico* in the *M. occidentalis* genome scaffold  
319 AFFJ01003151, which lead to XR\_145413 transcript annotation and coding for a putative  
320 *M. occidentalis* AChE (MoAChE). However, the first 106 nucleotides of the 5' UTR region  
321 of the KaAChE cDNA did not match with any portion of the scaffold sequence  
322 AFFJ01003151. In contrast, the unmatched 5'UTR portion of the KaAChE cDNA exhibited  
323 an 81% identity with segments of two partially overlapping scaffolds in the *M. occidentalis*  
324 genome (Genbank accession n. AFFJ01002403 and AFFJ01002403). The GT-AG  
325 consensus rule for donor and acceptor splice sites was also respected using the KaAChE  
326 cDNA sequence to guide the joining of the putative and still unannotated 5'UTR portion of  
327 MoAChE on the scaffolds AFFJ01002403 and AFFJ01002403 to the 5' end of the  
328 remaining open reading frame relying on the AFFJ01003151 scaffold. Because the  
329 AFFJ01002403 and AFFJ01002403 scaffolds do not overlap with AFFJ01003151, a long  
330 intronic sequence has to be envisaged in the MoAChE locus and is likely excluded from  
331 the assembly step. Assuming intron size conservation between the two phytoseiids, this  
332 hypothesis was supported by the unsuccessful amplification of this intron in the *K.*  
333 *aberrans* AChE locus. Although we were able to characterise 5 introns experimentally and  
334 an additional putative splicing site bioinformatically, we suggest that the KaAChE gene  
335 includes seven exons (I-VII) that are separated by 6 introns (Table 3, (Genbank accession  
336 n. HG328327). Exon I is non-coding, whereas exon II contains the initiation codon (ATG),  
337 which is similar to the observations of the majority of the AChE gene loci that have been  
338 annotated in insects and mites. Exons III-VI formed the catalytic domain and exhibited  
339 partially amino acid conservation across the AChEs that were cloned from the mites. Exon

340 exon seven contains the stop codon (TAG) and the 3' UTR region. All intron-exon  
341 boundaries followed the GT-AG rule (Breathnach et al., 1978); furthermore, these  
342 boundaries contained the YTNAN consensus sequence for lariat formation at the branch  
343 point close to the 5' end of the acceptor-splicing site. In addition to the positions, the  
344 lengths of the amplifiable introns were also conserved in the homologous AChE loci from  
345 the two phytoseiidae species with the exception of the third intron, which was slightly  
346 longer in the *K. aberrans* than in the *M. occidentalis* AChE gene (1162 bp vs. 936 base  
347 pairs, respectively). Sequence inspection of the third intron in the *K. aberrans* AChE locus  
348 revealed the presence of short microsatellite repeats and a long inverted repeat (LIR)  
349 (Wang et al., 2006). These nucleotide motifs can cause sliding of the intron sequences  
350 during DNA replication and might account for the different sizes of the third intron in the  
351 KaAChE gene.

352

### 353 3.5. Comparison the AChE cDNA sequences across different strains

354

355 Full-length sequencing of the KaAChE cDNA of the susceptible (LE) and resistant (PO)  
356 strains revealed a non-synonymous G to A mutation at position 687 that led to a G191S  
357 substitution in the protein sequence (G119S AChE *Torpedo* numbering; Fig. 1). This  
358 residue is involved in the oxianion hole, which is one of the functional domains of AChE  
359 activity (Zhang et al. 2002). The susceptible and resistant strains also differed in another  
360 single nucleotide polymorphism (SNP) at position 1499 of the cloned cDNA; this C to T  
361 transition did not affect the codon for the D461 residue. The resistant strain was  
362 homozygous at this site and carried only the T allele, while the sensitive strain exhibited  
363 both SNPs with a preference for C over T as indicated by the electropherograms. The  
364 phenylalanine residue (F339) that was replaced by a tryptophan in the chlorpyrifos-  
365 resistant stains of *T. urticae* (F439W mutation, or F331W AChE, *Torpedo* numbering) was  
366 still conserved both in the susceptible and resistant strains of *K. aberrans*. The same was  
367 true for the glycine residue (G336) that was found to be replaced with alanine (G328A) in  
368 the F331W-bearing strains of *T. urticae*. The cDNA KaAChE sequences of two additional  
369 susceptible (PA and PD) and three resistant (ME, SF and BX) strains of *K. aberrans* were  
370 also examined. The resistant strains were all homozygous for the G191S substitution,  
371 while the susceptible strains carried only the G191 allele. The resistant BX strain  
372 sequence differed from the other strains in a SNP in the 3' UTR that consisted of a G to A  
373 substitution.

374

#### 375 **4. Discussion**

376

377 Resistance to pesticides can be a desirable feature in *K. aberrans* because this predatory  
378 mite is an effective biocontrol agent for spider mites in perennial crops. Indeed, strains of  
379 this predatory mite that are apparently resistant to OP have successfully been released in  
380 vineyards and apple orchards in which the pest control strategies included chlorpyrifos and  
381 many other pesticides (Duso et al. 2009; Duso et al., 2012; Ahmad et al., 2013). The  
382 resistance to chlorpyrifos of these strains has been definitively demonstrated (Tirello et al.,  
383 2012), but the underlying molecular mechanisms remain poorly understood. An initial clue  
384 about this issue came from the chlorpyrifos-resistant strain of *T. pyri* that exhibited a lower  
385 level of AChE activity than the susceptible strain, which suggests that the reduced  
386 substrate affinity observed in the biochemical assay might be associated with a modified  
387 AChE (Fitzgerald and Solomon, 1999). In Acari, target site resistance due to a modified  
388 AChE that confers high levels of insensitivity to OPs, including chlorpyrifos, has been  
389 described in *T. urticae* and *Tetranychus kanzawai* Kishida (Aiki et al., 2004; Van Leeuwen  
390 et al. 2010; Khajehali et al., 2010). A G119S substitution (AChE torpedo numbering) in the  
391 single copy AChE gene has been associated with the moderate decreases in chlorpyrifos  
392 susceptibility between the resistant compared to the sensitive strains of *T. urticae*  
393 (resistance ratio at LD<sub>50</sub>, RR<sub>50</sub> = 31), and a greater resistance ratio has been detected in  
394 cases of F331W replacement (RR<sub>50</sub> > 400). *In vitro* expression of the AChE isoforms of *T.*  
395 *urticae* that carry F331W and/or G119S substitutions has revealed a reduction in  
396 sensitivity to another organophosphate (monocrotophos) and a decrease in the catalytic  
397 efficiency of the enzyme (Kwon et al., 2012); however, no data had been reported for  
398 chlorpyrifos. Although these features appeared much more evident in the F331W-mutated  
399 AChE, the two substitutions acted synergistically when they were associated *in vitro* and  
400 thus were favourably co-selected in the resistant strains *in vivo* (Kwon et al., 2010b; Ilias et

401 al., 2014). In mosquitoes, two acetylcholinesterase genes are present, both substitutions  
402 affect the paralogous AChE in the highly OP- and carbamate-resistant strains of *Culex*  
403 *pipiens* L. and *Anopheles gambiae* Giles (G119S; Weill et al., 2004a) and in *Culex*  
404 *tritaeniorhynchus* Giles (F331W; Alout et al., 2007). The role of these mutations in  
405 reducing the AChE sensitivity to OP was confirmed via inhibition analysis of the expression  
406 of AChE from mutated mosquito S2 cells (Weill et al., 2003; Oh et al., 2006). The F331W  
407 substitution has also been detected in AChE1 of a chlorpyrifos-resistant strain of the sweet  
408 potato whitefly *Bemisia tabaci* Gennadius (Alon et al., 2008). The *K. aberrans* strain with  
409 the highest level of insensitivity to chlorpyrifos (PO strain) described by Tirello et al. (2012)  
410 has a  $RR_{50} = 539,602$ , and this ratio is even higher than that found in the *T. urticae* and *B.*  
411 *tabaci* populations with the F331W AChE genotype. There a target site resistance might  
412 be present in that strain. Because no AChE sequences for *K. aberrans* are stored in  
413 databases, the annotated genome project of the predatory mite *M. occidentalis* was  
414 inspected. More than a dozen AChEs-like sequences were predicted by the curators of the  
415 *M. occidentalis* genome project using an automated computational analysis, although  
416 some of the transcripts represented uncompleted open reading frames or differed only in  
417 their splicing paths. To identify a suitable AChE candidate that is potentially responsible for  
418 target site resistance in *K. aberrans*, the AChE protein sequence from *T. urticae* was used  
419 to probe probing the annotated genome of *M. occidentalis*. Once a putative homologous  
420 AChE in *M. occidentalis* was found, its sequence was used to speed up the cloning of the  
421 corresponding AChE cDNA in *K. aberrans*. Full sequencing of the cloned AChE cDNA  
422 revealed that the resistant strain (PO) differed from the susceptible strain (LE) in terms of  
423 non-synonymous G to A mutation that introduced a G191S substitution in the AChE open  
424 reading frame. That mutation corresponds to the aforementioned G119S substitution in  
425 AChE Torpedo numbering. Strangely, the corresponding amino acid position in the  
426 homologous AChE that was found in the annotated genome of *M. occidentalis* is occupied

427 by a serine. Unfortunately no information is available concerning the chlorpyrifos  
428 susceptibilities of the *M. occidentalis* strains employed for the genome project. Notably, the  
429 G119 in the *K. aberrans* AChE is encoded by a GGC codon, which could easily be  
430 converted to the AGC codon for serine. This substitution in the mosquito paralogous  
431 AChE seems not to be neutral under the selective pressure produced by organophosphate  
432 and carbamate treatments (Weill et al. 2004b). Indeed, when KaAChE cDNA from an  
433 additional two chlorpyrifos-susceptible and three chlorpyrifos-resistant unrelated strains of  
434 *K. aberrans*, the G191S substitution was absent only in the resistant strains in the in  
435 homozygous condition. The silent nucleotide polymorphisms found in the KaAChE cDNA  
436 of the resistant strains might may also suggest that different G119S mutation events  
437 occurred independently. Although, in *Culex quinquefasciatus* Say, a chlorpyrifos inhibition  
438 study of a paralogous AChE bearing the G119S substitution revealed a reduced sensitivity  
439 to the insecticide (Liu et al., 2005) that likely resulted from the reduced accessibility of the  
440 catalytic site (Weill et al., 2004a). *In vivo*, *T. urticae* strains with the same mutated AChE  
441 genotype display only a moderate resistance to chlorpyrifos. In contrast, all examined  
442 resistant strains of *K. aberrans* are highly resistant to the insecticide (Tirello et al., 2012).  
443 Thus, the role of the G119S remains unclear, although the possibility that the same  
444 mutation has different effects on chlorpyrifos AChE sensitivity in the predatory mite cannot  
445 be ruled out. Nevertheless, the non-silent nucleotide polymorphism responsible for the  
446 G119S substitution represents a reliable molecular marker that is associated with the  
447 detected resistant phenotype. Alternative target site resistance mechanisms not explored  
448 in this study might also rely on the amplification of modified AChEs as occurs in *T. urticae*  
449 and *T. evansi* (Kwon et al., 2010a; Carvalho et al., 2012) or on mutations that affect  
450 multiple AChE loci with additive effects such as has been observed in *R. microplus*  
451 (Temeyer et al., 2009, 2010, 2012). In either case, the co-expression of sensitive and  
452 insensitive AChEs might contribute to reducing the fitness costs associated with OP

453 resistance (Carvalho et al., 2012, Temeyer et al. 2013a). Alternatively, the overexpression  
454 of distinct AChEs from different loci it is thought to result in to bio-scavenging due to the  
455 supply of excess targets for xenobiotics, including OP and carbamate insecticides (Lee et  
456 al. 2014, in press). While the detection of homozygosity for the G119S substitution argues  
457 against the co-existence of duplicated sensitive and insensitive AChEs in chlorpyrifos-  
458 resistant strains, the role of multiple AChE loci in *K. aberrans* cannot be not excluded.  
459 Intriguingly, the genome of the closely related phytoseiides *M. occidentalis* harbours at  
460 least six AChE-like candidates (GenBank accession numbers XP\_003743019,  
461 XP\_003747509, XP\_003739584, XP\_003738701, XP\_003744479, and XP\_003745369).  
462 Although these putative AChEs have lower amino acid identities (< 34%) to those found in  
463 *T. urticae*, they exhibit conserved functional residues for acetylcholinesterase activity (i.e.,  
464 the form the catalytic triad and the acetylcholine binding pocket), exhibit conserved amino  
465 acid positions, are potentially involved in substitutions that affect AChE sensitivity to  
466 organophosphates and carbamates, have and received support from transcriptomics  
467 analyses (Hoy et al., 2013). Together, these findings suggest that, in predatory mites,  
468 multiple AChEs resemble the composite picture observed in ticks (Temeyer et al. 2013b).  
469 Thus, herbivorous and predatory mites can differ not only in detoxification pathways  
470 (Mullin et al., 1982; Grbic et al., 2001; Dermauw et al., 2012) but also in AChE repertoires,  
471 which potentially offers alternative solutions for the development of target site resistance.  
472 However, chlorpyrifos inhibition assays of AChE activity should be performed to support  
473 this hypothesis in examined OP resistant strains. Although, the contribution of enhanced  
474 detoxifying activities to chlorpyrifos resistance in predatory mites has not yet been reported  
475 as it has for other organophosphates (Sato et al., 2001, Fournier et al., 1987; Motoyama et  
476 al., 1971; Anber et al., 1988), the use of synergists in bioassays should be combined with  
477 detoxification enzyme assays to confirm or deny the involvement of metabolic resistance.

478 In summary, the potential target site resistance to chlorpyrifos in *K. aberrans* has barely  
479 been dissected compared to that in *T. urticae*. The F331W substitution that is responsible  
480 for AChE that is highly insensitive to chlorpyrifos in Tetranychidae was absent in a putative  
481 homologous gene that was cloned from the resistant strain *K. aberrans*. However, a  
482 G119S mutation that was detected in the same gene appeared to be associated with the  
483 resistant phenotype. Because pesticide treatments strongly affect the success of predatory  
484 mites release (Ahmad et al., 2013), this polymorphism might be useful as a molecular  
485 marker for tracing the resistant phenotype in ecological studies or in gene pyramiding and  
486 marker-assisted selection of desirable traits for multiple insecticide resistance.

487

488 **References**

489 Not included

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**Table 1. Strains of *Kampimodromus aberrans* and their origins.**

Strain	Origin
PO	Commercial vineyard at Monteforte d'Alpone (Verona province)
SF	Commercial vineyard at S. Pietro Incariano (Verona province)
BX	Commercial vineyard at Soave (Verona province)
ME	Commercial vineyard at Valdobbiadene (Treviso province)
LE	Untreated European nettle trees ( <i>Celtis australi</i> L.) at Legnaro (Padova province)
PD	Untreated European nettle trees ( <i>Celtis australi</i> L.) at Padova
PA	Untreated European nettle trees ( <i>Celtis australi</i> L.) at Paese (Treviso province)

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**Table 2. Primer sequences**

	Forward primer	Sequence 5' → 3'	Reverse primer	Sequence 5' → 3'
<i>Degenerate primers</i>	KaAChEF1d KaAChEF2d	GGNATHCCNTAYGCNAARCC CCNTAYGCZAARCCNCC	KaAChER1d KaAChER2d	RAANSWNCCNCCRTANACCCA RAARCTNCCNCCRTANACCCA
<i>3' cDNA extension</i>	KaAChEF3	TGGAATGCCAACACTAATA	KaAChER3	AGGAAGTTCTTCCAAAACGCGCA
<i>3' RACE</i>			oligdTadapter	ACAGCAGGTCAGTCAAGCAGTAGC AGCAGTTCGATAAGCGGCCGCAT GGAT <sub>12</sub> DN
<i>5' RACE</i>	KaAChEF4 KaAChEF5	AGGTGGTGAACGAAGCCATCAT TTGAGTACACGGACTGGTTGAACCCT	Adapter1 Adapter2	ACAGCAGGTC AAGTCAAG AGCAGTAGCAGCAGTTCGATA
	TS-Primer	CACCATCGATGTCGACACGCGTCGGG IGGIG	KaAChER4 KaAChER5	CTCCGTAAACCCATAGAAGGA CCACATTGTTGAACCCTCAAAGTCG
<i>cDNA sequencing</i>	TS-PCR	CATCGATGTCGACACGCGTC	KaAChER6	TGTTGAACCCTCAAAGTCGTCG
	KaAChEF6 KaAChEF7 KaAChEF8 KaAChEF9 KaAChEF10	ACCTTGATAAACTGTCGCTGTGGC CAATGCAGGCATGATGGACCAAGT GTTGAACCCTGACGATCCGATCAA AAATGCGATTTTCGACATCCTGTGCC GCGCTATCGGGCAACAACACAAACA	KaAChER7 KaAChER8 KaAChER9 KaAChER10	AAAGTCACGTTGTTCCGGGTTGCCT AGGAAATTCTTCCAGAATGCGCA TCCTAGTTCGCTCCTTCAGTTGGA CCAACAATCTTGTGACGGCATCT
<i>G119S screening</i>	KaAChEF11	AATGCGATTTTCGACATCCTGTGCC	KaAChER11	AAAGTCCGTTGTTCCGGGTTGCC
<i>F331W screening</i>	KaAChEF12	AAACTCGCGGAGGAAGTCAAGTGT	KaAChER12	CCAACAATCTTGTGACGGCATCT

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**Table 3. Genome organisation of the *K. aberrans* ace locus**

Exon	Position (nt) <sup>a</sup>	Exon size (bp)	Splice junction	Intron size (bp)	phase	amino acid involved
Exon I (inferred)	(-106) - (-11)	106	TCGGAGGACTGGgt ...	unknown	-	5' UTR
Exon II (ATG)	(-10) - 310	320	agATGATTGGATAT ... ACGCATTCCCTCGgt ...	113	I	G104
Exon III	311-876	566	agGTATTCCGTATG ... CTATTTTCTCAAgt ...	1162	0	Q292
Exon IV	877-1302	426	agGCCGTGCTGCAA ... TTTACCTCGCAGgt ...	194	0	Q434
Exon V	1303-1494	192	agGTGGTGAACGAA ... CGCTCGTCTCAGgt ...	402	0	Q498
Exon VI	1495-1822	328	agAATAAATGGCCG ... TCGCGCTATCGGgt ...	145	I	G608
Exon VII (TAA)	1823-2213	391	agGCAACAACACAA ...			3'UTR

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The numbering of nucleotides is based on the *K. aberrans* AChE cDNA in which +1 corresponds to start codon.

1	KaAChE	MRLGVSGGASGLAK-LALCRSMAATVRSLSLTRA\LF AFLIHLWLVVTL CVGRVDARA AHL	58
2	MoAChE	MWLGVS GGASGLAKCKLALCRSMATILRSMSTRTLFAFLINTWLVVTL CVGRVDARA AHL	60
3		* *****: :*:***:*****: *****	
4		▽	
5	KaAChE	LHHHRHRTAGSAQS QGDPLL VHTTKGPV RGITLQASNGKLVDAFLGIPYAKPPV GKMRFR	118
6	MoAChE	LHHHRHRTAGNAQS QGDPLV VHTTKGPV RGITLQASNGKLVDAFLGIPYAKPPV GKMRFR	120
7		*****.*****:*****:*****:*****:*****:*****	
8		1 + 1	
9	KaAChE	HPVPM DPWEKPLNVTERPATCVQVVD TYFDDFEGSTMWNANTNMSEDC LNMLVWVPRPRP	178
10	MoAChE	HPVPM DPWEKPLNVTEPPATCVQVVD TYFDDFEGSTMWNANTNMSEDC LNMLVWVPRPRP	180
11		***** *****	
12		++	
13	KaAChE	TNAAVLLWVYGG <b>G</b> FYSGCATLDVYDGKILASEENVIVVSFN YRVGSLGFLYLDHADAPGN	238
14	MoAChE	TNAAVLLWVYGG <b>S</b> FYSGCATLDVYDGKILASEENVIVVSFN YRVGSLGFLYLDHADAPGN	240
15		*****.*****	
16		+ +	
17	KaAChE	AGMMDQVMALRWVQDNIHLFGGNPNNVTLFGES <b>S</b> AGAVSVAYHLLSPLSRDLFSQAVLQSG	298
18	MoAChE	AGMMDQVMALRWVQDNIHLFGGNPNNVTLFGES <b>S</b> AGAVSVAYHLLSPLSRDLFSQAVLQSG	300
19		*****	
20		+ 2 2	
21	KaAChE	GATVPWGYNERQTAMTNGYKLAEEVKCPTDDVEATVKCLRLQDPDLLVKSEIFATGVVDF	358
22	MoAChE	GATVPWGYNERQTAITNGYKLAEEVKCPTDDVEATIKCLRLQDPDLLVKSEIFATGVVDF	360
23		*****.*****:*****:*****:*****:*****	
24		+ +	
25	KaAChE	SFVPVVDGAFLTERPEDTMNSGNFKKCKILLGNSNRD <b>E</b> GTYFIIYYLTQLFKRDENVYLTR	418
26	MoAChE	SFIPVVDGAFLTERPEDSMSSGNFKKCKILLGNSNRDEGTYFIIYYLTQLFKRDENVYLTR	420
27		** :*****:*.*****	
28		▽	
29	KaAChE	EDFVDAVQALSPFTSQVVNEAII FEYTDWLN PDDPIKNRDAVDKIVGDYFTCPVIDTAH	478
30	MoAChE	EDFVDAVQALSPFTSSVVNEAII FEYTDWLN PDDPIKNRDAVDKIVGDYFTCPVIDMAH	480
31		*****.***** **	
32		▽	
33	KaAChE	YYSSAGLDVYMYYYVYRSSQNKWPEW MGVI <b>H</b> ADEIAYVFGEPLNQ TWSYRQDEQMFSRRI	538
34	MoAChE	YYFISGLDVYMYYYVYRSSQNKWPEW MGVI <b>H</b> ADEIAYVFGEPLNQ TWSYRQDEQMFSRRI	540
35		** :*****	
36		3	
37	KaAChE	MRYWANFARMGNPSLNP DGNWEKTYWPAHTAFGKEFLILDVNSTQVGYGNRAKHCAFWKN	598
38	MoAChE	MRYWANFARMGNPSLNP DGNWEKTYWPAHTAFGKEFLILDVNSTQVGYGNRAKHCAFWKN	600
39		*****	
40		▽ +	
41	KaAChE	FLPNLIALSGNNTNKAE EGRD GASSQSSSIMLLCSLAASIVVTGRILSQPPATAA	655
42	MoAChE	FLPNLIALSGNNTNKAEDESKDGASTQSSSLTLLCSLAASMIVTGRLLSSSTARAA	657
43		*****:*.***:*****: *****:*****:*****:*****	
44			

**Fig. 1** Alignment of the AChEs cloned in from the chlorpyrifos -susceptible strain of *Kampimodromus aberrans* (KaAChE) and predicted from *Metaseiulus occidentalis* transcript XR\_145413 (MoAChE). Identical amino acids are indicated by asterisks, and conservative substitutions are indicated by dots. The cleavage site of signal peptide is indicated by a slash. The mutated residue (G191S) in the chlopyrifos-resistant strain is in reverse in the background, the cysteine residues that form the intramolecular disulphide bonds are numbered and on the light-gray background (C139-C166, C325-C336, C471-C593), the catalytic triad residues are boxed (S271, E395, H509), and the following conserved residues are indicated with plus signs: anionic subsite (W156), oxianion-hole (G189, G190, A274), acyl pocket (W304, F360, F399), and cysteine residue forming

- 1 intermolecular disulphide bond (C634). The inverted triangles indicates the exon
- 2 boundaries in the cDNA open reading frame.