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**A Single Amino Acid Substitution in the Third Transmembrane Region  
Has Opposite Impacts on the Selectivity of the Parasiticides Fluralaner  
and Ivermectin for Ligand-Gated Chloride Channels**

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## Running Title Page

Running title: Actions of isoxazoline ectoparasiticide on chloride channels

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**ABBREVIATIONS:** AVM, avermectin; DMSO, dimethyl sulfoxide; GABA,  $\gamma$ -aminobutyric acid; GABA<sub>Cl</sub>, GABA-gated chloride channel; GluCl, glutamate-gated chloride channel; IVM, ivermectin; LGIC, ligand-gated ion channel; SOS, standard oocyte solution; TM, transmembrane segment; TSI, transmembrane subunit interface

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## ABSTRACT

Fluralaner (Bravecto<sup>TM</sup>) is a recently marketed isoxazoline ectoparasiticide. This compound potently inhibits  $\gamma$ -aminobutyric acid (GABA)-gated chloride channels (GABA<sub>Cl</sub>s) and less potently glutamate-gated chloride channels (GluCl<sub>s</sub>) in insects. The mechanism underlying this selectivity is unknown. Therefore, we sought to identify the amino acid residue(s) causing the low potency of fluralaner toward GluCl<sub>s</sub>. We examined the fluralaner sensitivity of mutant housefly (*Musca domestica*) GluCl<sub>s</sub> in which amino acid residues in the transmembrane subunit interface (TSI) were replaced with the positionally equivalent amino acids of *Musca* GABA<sub>Cl</sub>s. Of these amino acids, substitution of an amino acid (Leu315) in the third transmembrane region (TM3) with an aromatic amino acid dramatically enhanced the potency of fluralaner in the GluCl<sub>s</sub>. In stark contrast to the enhancement of fluralaner potency, this mutation eliminated the activation of currents and the potentiation but not the antagonism of glutamate responses that are otherwise all elicited by the macrolide parasiticide ivermectin (IVM). Our findings indicate that the amino acid Leu315 in *Musca* GluCl<sub>s</sub> plays significant roles in determining the selectivity of fluralaner and IVM for these channels. Given the high sequence similarity of TM3, this may hold true more widely for the GluCl<sub>s</sub> and GABA<sub>Cl</sub>s of other insect species.

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## Introduction

Ligand-gated ion channels (LGICs) play vital roles in regulating neuronal excitation and inhibition in animals. These channels are either (1) cation-selective channels, the activation of which depolarizes the postsynaptic membrane towards firing an action potential, or (2) anion-selective channels, the activation of which hyperpolarizes the membrane or suppresses the depolarization generated by cation channels (Smart and Paoletti, 2012). Nicotinic acetylcholine receptors and  $\gamma$ -aminobutyric acid (GABA) receptors, which are members of the Cys-loop family of LGICs, are examples of such cation and anion channels, respectively (Miller and Smart, 2010). Inhibitory glutamate receptors, which are found only in invertebrates, are also Cys-loop LGIC family members. The Cys-loop LGICs are pentamers, the subunits of which are assembled to form a central ion-permeable channel. Each subunit consists of a large N-terminal extracellular domain, four hydrophobic  $\alpha$ -helical transmembrane segments (TMs), an intracellular loop between TM3 and TM4, and a short extracellular C-terminus. The orthosteric agonist-binding site is located at the subunit interface of the extracellular domain. These channels are important targets for drugs and insecticides (Alexander et al., 2015; Ozoe, 2013).

Glutamate-gated chloride channels (GluCl<sub>s</sub>) are the main targets for the insecticidal, acaricidal, and nematocidal macrolides avermectins (AVMs), which allosterically activate and modulate various ion channels, including GABA-gated chloride channels (GABA<sub>A</sub>Cl<sub>s</sub>), glycine-gated chloride channels, pH-gated chloride channels,  $\alpha$ 7 acetylcholine-gated cation channels, ATP-gated P2X receptor cation channels, and G protein-gated inwardly rectifying potassium channels (Chen et al., 2017; Cully et al., 1994; Cully et al., 1996; Dawson et al., 2000; Fuse et al., 2016; Krause et al., 1998; Nakatani et al., 2016; Shan et al., 2001; Silberberg et al., 2007). GluCl<sub>s</sub> are activated by

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nanomolar AVMs, whereas other channels require concentrations in the micromolar range. Abamectin (AVM B<sub>1</sub>) is used to control phytophagous mites and insect pests on agricultural and horticultural crops (Lasota and Dybas, 1991). Ivermectin (IVM) (Fig. 1), C22-23-dihydro AVM B<sub>1</sub>, is widely used to control endo- and ectoparasites in animals and to treat onchocerciasis and lymphatic filariasis caused by parasitic worms in humans (Laing et al., 2017; Ōmura and Crump, 2014; Wolstenholme et al., 2016). GABACls are targets for chlorinated hydrocarbon and phenylpyrazole insecticides, which stabilize the closed conformation of GABACls by interacting with TM2 amino acid residues on the cytoplasmic side within the channel pore (Ozoe, 2013). Isoxazoline and *m*-benzamidobenzamide insecticides are new-generation antagonists that inhibit insect GABACls (Asahi et al., 2015; Gassel et al., 2014; McTier et al., 2016; Nakao and Banba, 2016; Nakao et al., 2013; Ozoe et al., 2010; Ozoe et al., 2013; Shoop et al., 2014). These insecticides inhibited GABACls with mutations conferring insensitivity to the conventional antagonists (Nakao et al., 2013; Asahi et al., 2015). The substitution of a conserved TM3 Gly located in the transmembrane subunit interface (TSI) of *Drosophila* GABACls diminished or eliminated the inhibitory effects of a *m*-benzamidobenzamide (meta-diamide) insecticide on GABA responses, whereas the conventional antagonists remained effective (Nakao et al., 2013). These findings indicate that these insecticides have mode(s) of action distinct from that of the conventional channel-blocking antagonists.

Fluralaner (Bravecto™) (Fig. 1) is a recently marketed isoxazoline ectoparasiticide that is used for flea and tick protection in pets (Taenzler et al., 2014; Wengenmayer et al., 2014). This compound inhibits agonist responses in both GABACls and GluCl<sub>s</sub>, with selectivity for the former over the latter (Ozoe et al., 2010), whereas IVM is a selective

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activator or modulator for GluCl<sub>s</sub> (Fuse et al., 2016). In the present study, we sought to identify the mechanism underlying the difference in fluralaner and IVM sensitivity between GluCl<sub>s</sub> and GABA<sub>A</sub>Cl<sub>s</sub>. We report that the substitution of a single amino acid in the TSI of GluCl<sub>s</sub> with a positionally equivalent amino acid from GABA<sub>A</sub>Cl<sub>s</sub> results in a drastic increase in the antagonist potency of fluralaner and a dramatic elimination of the IVM activation of currents and the IVM potentiation of the glutamate responses.

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## Materials and Methods

**Chemicals.** Fluralaner (99%), A1209 (99%), and A341 (99%) were synthesized according to a previously reported method (Mita et al., 2005, 2009, 2010). GABA and IVM ( $B_{1a} \geq 90\%$ ,  $B_{1b} \leq 5\%$ ) were purchased from Sigma-Aldrich (St. Louis, MO, USA). Other general chemicals including sodium hydrogen L-glutamate were purchased from Wako Pure Chemical Industries, Ltd. (Osaka, Japan), unless otherwise noted.

**Wild-type and mutant *GluCl* cDNAs.** Two LGIC splice variants that show robust agonist responses were used in this study. cDNAs encoding GluCl (variant A) (accession No. AB177546) and GABA<sub>Cl</sub> (RDL variant ac) (accession No. AB177547, AB824728, AB824729) subunits from the housefly (*Musca domestica*; WHO/SRS strain) were subcloned into the plasmid vectors pcDNA3 and pBluescript KS(-), respectively (Eguchi et al., 2006; Ozoe et al., 2013). The introduction of mutations into the cDNAs was performed using a QuikChange Site-Directed Mutagenesis Kit (Agilent Technologies, Santa Clara, CA, USA) and verified by DNA sequencing.

**Two-electrode voltage clamp electrophysiology.** The lobes of the ovary were surgically removed from female African clawed frogs (*Xenopus laevis*) anesthetized by immersion in a 0.1% (w/v) Tricaine solution. Follicle cells were treated with collagenase (2 mg/ml; Sigma-Aldrich) in a calcium-free standard oocyte solution ( $Ca^{2+}$ -free SOS) (100 mM NaCl, 2 mM KCl, 1 mM  $MgCl_2$ , 5 mM HEPES, pH 7.6) for 1–2 h at 20 °C. After the treatment, the oocytes were washed with  $Ca^{2+}$ -free SOS supplemented with 2.5 mM sodium pyruvate, gentamycin (50 µg/ml; Thermo Fisher Scientific, Waltham, MA, USA), penicillin (100 U/ml; Thermo Fisher Scientific), and streptomycin (100 µg/ml; Thermo Fisher Scientific) and incubated for 1–2 days at 16 °C.

The *Musca GluCl* and *Rdl* cDNAs containing a T7 promoter site upstream of the

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coding region were amplified by PCR. The PCR products were purified using an illustra GFX PCR DNA and Gel Band Preparation Kit (GE Healthcare Bio-Sciences, Pittsburgh, PA, USA). After sequence verification, the amplified cDNA templates (100 ng) were *in vitro* transcribed into capped poly(A) cRNAs using a mMESAGE mMACHINE® T7 Ultra Kit (Thermo Fisher Scientific). The quality and quantity of the prepared cRNAs were evaluated by agarose gel electrophoresis and absorption spectroscopy, respectively. Purified cRNA (5 ng) was injected into each oocyte using a Nanoliter 2000 injector (World Precision Instruments, Sarasota, FL, USA). The injected oocytes were incubated for 2–4 days at 16 °C.

The oocytes expressing *Musca* GluCl<sub>s</sub> or GABA<sub>A</sub>Cl<sub>s</sub> were placed in a chamber perfused with SOS (100 mM NaCl, 2 mM KCl, 1.8 mM CaCl<sub>2</sub>, 1 mM MgCl<sub>2</sub>, 5 mM HEPES, pH 7.6). Glass microelectrodes were filled with 2 M KCl to yield a resistance of 0.5-1.6 MΩ. Electrophysiological recordings were performed using an Oocyte Clamp OC-726C amplifier (Warner Instruments, Hamden, CT, USA) at a holding potential of -80 mV at 20 °C. The data were digitized using a Lab-Trax-4/16 converter (World Precision Instruments) and analyzed using the Data-Trax2 software (World Precision Instruments). IVM, fluralaner and fluralaner analogs dissolved in dimethyl sulfoxide (DMSO) were diluted with SOS to produce solutions containing the indicated concentrations for each compound and <0.01% DMSO. To analyze the antagonism of the GluCl<sub>s</sub> or GABA<sub>A</sub>Cl<sub>s</sub> by fluralaner and its analogs, glutamate or GABA dissolved at the EC<sub>50</sub> of each channel (Table 1) in SOS was applied to the oocytes for 3 s at 30–60-s intervals with the perfusion of fluralaner until maximum inhibition was achieved. Oocytes were perfused with the solution for 3 min to analyze channel activation by IVM. The ability of IVM to potentiate and antagonize glutamate responses was analyzed in a manner



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similar to the evaluation of fluralaner antagonism but using glutamate at its EC<sub>5</sub> and EC<sub>90</sub>, respectively. The EC<sub>5</sub> and EC<sub>90</sub> are the concentrations at which the potentiation and antagonism of glutamate responses are observed, respectively (Fuse et al., 2016). All experiments were replicated using at least six oocytes from at least two frogs. The data are presented as the means ± SEM. The EC<sub>50</sub> and IC<sub>50</sub> values were obtained from dose-response relationships by the four-parameter logistic regression using OriginPro 8J SR4(ver.8.0951) (LightStone, Tokyo, Japan). Unpaired *t*-tests were performed to evaluate statistical significance; *p*-values for L315 mutants are reported with a Bonferroni correction for multiple tests.

**Homology modeling and docking simulation.** The amino acid sequences of the *Musca* GluCl-A subunit and the *Caenorhabditis elegans* GluCl- $\alpha$  subunit were aligned using ClustalW2. A *Musca* GluCl homology model was constructed using MOE software (version 2014.04; Chemical Computing Group, Montreal, Canada). The X-ray crystal structure of the *C. elegans* GluCl- $\alpha$  channel (PDB code: 3RHW) was used as a template.

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## Results

**Responses of wild-type and mutant GluCl<sub>s</sub> to glutamate.** The amino acid at position 36' (index number starting from a conserved TM2 cationic residue numbered 0') in TM3 is particularly important in determining the sensitivity of IVM in LGICs (Lynagh and Lynch, 2010). To examine the effects of intersubunit amino acids on the potencies of fluralaner and IVM against GluCl<sub>s</sub>, we first substituted four GluCl amino acids (Ile253, Met257, Leu315, and Thr316) near G36' (Gly312 in the *Musca* GluCl subunit) with the positionally equivalent amino acids of the *Musca* GABA<sub>A</sub>Cl (RDL) subunit (Fig. 2), generating four single mutants (I253A, M257L, L315F, and T316V) and one double mutant (M257L/T316V). Because the amino acid at position 253 of the *Musca* GluCl subunit was identical to the amino acid of the *Musca* GABA<sub>A</sub>Cl RDL subunit at the equivalent position, Ile253 was substituted with Ala. All wild-type and mutant GluCl<sub>s</sub> expressed in *Xenopus* oocytes responded to glutamate to elicit currents (Fig. 3A). The L315F mutant was  $\approx 36$ -fold less sensitive to glutamate compared with the wild-type channel (Table 1). Although this mutation may allosterically affect glutamate binding to the orthosteric site, the dose-response curve indicated that the channel functioned normally to induce currents in response to glutamate.

**Fluralaner inhibition of glutamate-induced currents in mutant GluCl<sub>s</sub>.** Fluralaner inhibited glutamate-induced currents in wild-type *Musca* GluCl<sub>s</sub> expressed in *Xenopus* oocytes (Fig. 3B,D), with an IC<sub>50</sub> of 146 nM (Table 1). However, wild-type *Musca* GluCl<sub>s</sub> were  $\approx 24$ -fold less sensitive to fluralaner than were wild-type *Musca* GABA<sub>A</sub>Cl<sub>s</sub>, as previously reported (Ozoe et al., 2010). Fluralaner was  $\approx 2$ -fold less potent toward the I253A mutant than in the wild-type GluCl, whereas it was  $\approx 4$ -fold and  $\approx 2$ -fold more potent in the M257L and M257L/T316V mutants, respectively, than in the wild-

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type GluCl (Fig. 3D; Table 1). The potency of fluralaner in the T316V mutant did not differ significantly from that in the wild type. Notably, the L315F mutant was strongly inhibited by fluralaner, with an  $IC_{50}$  of 1.06 nM, indicating that this mutant is  $\approx 138$ -fold more sensitive to fluralaner inhibition than the wild type (Fig. 3C,D; Table 1). The  $IC_{50}$  of fluralaner in the L315F mutant was even  $\approx 6$ -fold smaller than its  $IC_{50}$  in the inhibition of GABA-induced currents in *Musca* GABACls (Fig. 3D; Table 1).

**IVM actions on mutant GluCls.** IVM alone activated slow, irreversible currents in wild-type *Musca* GluCls expressed in *Xenopus* oocytes (Fig. 4A), with an  $EC_{50}$  of 18.8 nM (Table 1). The wild-type *Musca* GluCls were  $\approx 66$ - to  $\approx 184$ -fold more sensitive to IVM than were wild-type *Musca* GABACls (Table 1). The potency of IVM was  $\approx 5$ -fold higher in the I253A mutant than in the wild type, whereas the potencies of the M257L and T316V mutants were not significantly different from that of the wild-type GluCl (Fig. 4C; Table 1). The difference in the maximal current amplitude between the wild type and mutants could be ascribed to differences in the expression levels of channels in oocytes (Fig. 4C). Surprisingly, the L315F mutant, which showed an enhanced sensitivity to fluralaner, lacked sensitivity to IVM in terms of activation (Fig. 4B,C). Next, we examined IVM potentiation and antagonism of glutamate-induced currents in the L315F mutant because IVM exerts a triple action on GluCls and GABACls depending on the conditions (Fuse et al., 2016). In L315F GluCls, IVM did not potentiate the currents induced by a low concentration of glutamate ( $EC_5$ , 50  $\mu$ M) (Fig. 5A,B) but inhibited those induced by a high concentration of glutamate ( $EC_{90}$ , 3 mM), with an  $IC_{50}$  of  $5.48 \pm 1.20$  nM (SEM,  $n = 6$ ), which is not significantly different from the  $IC_{50}$  ( $4.92 \pm 2.23$  nM) for the wild type (Fig. 5C,D).

**Effects of fluralaner analogs on L315F mutant GluCls.** As fluralaner showed

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marked antagonism of the L315F GluCl, we examined whether a similar potency enhancement could be observed for fluralaner analogs (Fig. 6; Fig. 7). The isoxazolines A1209 and A341 (Fig. 1) are fluralaner analogs that show >500-fold and >100-fold higher antagonism of *Musca* GABA<sub>A</sub> receptors, respectively, compared with *Musca* GluCl receptors (Fig. 6A,B,D; Fig. 7A,B,D; Table 1). The L315F mutation resulted in >6000- and >100-fold enhancement of the potency of A1209 and A341 in GluCl receptors (Fig. 6C,D; Fig. 7C,D; Table 1), which were, respectively, much greater than and comparable to the enhancement observed for fluralaner. The L315F GluCl was outstandingly sensitive to A1209. Although the isoxazoline A341 exhibited little antagonism in wild-type GluCl receptors, the antagonist potency of this compound in L315F GluCl receptors was comparable to that in wild-type GABA<sub>A</sub> receptors (Fig. 7).

**Effects of aromatic amino acid substitution at position 315 on the potency of fluralaner.** As the L315F substitution enhanced the potency of fluralaner against *Musca* GluCl receptors, we examined the effects of other amino acid substitutions on the potency of fluralaner (Fig. 8). We injected cRNAs encoding five mutants (L315Y, L315W, L315H, L315A, and L315M) into oocytes. The aromatic amino acids included Tyr, Trp, and His, which has the aromatic heterocycle imidazole in the side chain. Ala and Met were chosen as hydrophobic aliphatic amino acids. Although the oocytes injected with the cRNAs for L315W and L315A failed to respond to glutamate, the other oocytes did respond to glutamate (Fig. 8A). Fluralaner inhibition of glutamate-induced currents was  $\approx 24$ -fold and  $\approx 8$ -fold more potent in the L315Y and L315H mutants, respectively, than in the wild type (Fig. 8B,C,E; Table 1). In contrast, the potency of fluralaner did not differ significantly between the L315M mutant and the wild-type channel (Fig. 8E; Table 1). These findings indicate that aromatic amino acids at position 315 are effective in

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enhancing the antagonist potency of fluralaner in *Musca* GluCl<sub>s</sub> but that an aliphatic amino acid is not.

**Effects of aromatic amino acid substitution at position 315 on the potency of IVM.** We examined the effects of the L315Y, L315H, and L315M substitutions on IVM-induced currents in *Musca* GluCl<sub>s</sub> (Fig. 9). The former two substitutions abolished the IVM-induced activation of *Musca* GluCl<sub>s</sub> (Fig. 9A,B,D). In contrast, IVM activated currents in the L315M mutant, although the potency was  $\approx 2$ -fold lower than in the wild-type channel (Fig. 7C,D; Table 1). These findings indicate that aromatic amino acids, but not an aliphatic amino acid, at position 315 eliminate the IVM-induced activation of GluCl<sub>s</sub>.

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## Discussion

The transmembrane subunit interface (TSI) of pentameric LGICs plays critical roles in the actions of insecticides and other drugs (Forman and Miller, 2016; Nakao et al., 2013). The TSI in mammalian GABACls has been extensively studied as a binding site for general anesthetics such as propofol, etomidate, and barbiturates (Forman and Miller, 2016). Interestingly, both convulsive and anesthetic barbiturates modulate GABACls by binding to this region (Jayakar et al., 2015). IVM activates currents and potentiates and antagonizes the agonist-induced currents of LGICs including GABACls and GluCl<sub>s</sub>, with the latter being more sensitive than the former, by possibly binding in the TSI (Estrada-Mondragon and Lynch, 2015; Fuse et al., 2016; Hibbs and Gouaux, 2011). A single TM3 amino acid in the TSI, Gly at position 36' (G36') (Fig. 2), which is conserved in IVM-sensitive LGICs, has been shown to be critical for these actions of IVM (Fuse et al., 2016; Lynagh and Lynch, 2010). The substitution of G36' with bulkier amino acids results in a reduction or the loss of sensitivity to IVM. The importance of G36' in insecticide actions was indicated by a report that a G36'D mutation was identified in the abamectin-resistant strain of two-spotted spider mites (*Tetranychus urticae*) (Kwon et al., 2010). A G36'E substitution disrupted *T. urticae* GluCl activation by abamectin and milbemycin (Mermans et al., 2017). Furthermore, G36' mutations were reported to diminish or eliminate the ability of a meta-diamide insecticide to inhibit GABA-induced currents in the *Drosophila* GABACls (Nakao et al., 2013). Interestingly, the analogous mutation of a positionally equivalent Gly was identified in the nicotinic acetylcholine receptor  $\alpha 7$  subunit of spinosad-resistant western flower thrips (*Frankliniella occidentalis*) and the nicotinic acetylcholine receptor  $\alpha 6$  subunit of spinosad-resistant tomato leafminers (*Tuta absoluta*) (Puinean et al., 2013; Silva et al., 2016). Together, these reports indicate that a

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conserved Gly in TM3 plays a key role and that the TSI seems to form the sites of action for a variety of ligands.

The ectoparasiticide fluralaner was shown to exhibit selective antagonism of GABACls over GluCls (Ozoe et al., 2010). To examine whether any amino acid substitution enhances the low potency of fluralaner in GluCls, we focused on replacing amino acids around G36' in the TSI of *Musca* GluCls with the positionally equivalent amino acids of *Musca* GABACls, which were sensitive to fluralaner (Table 1). We have shown that the substitution of an amino acid, Leu315, located one  $\alpha$ -helical turn below G36', with aromatic amino acids (but not with an aliphatic amino acid) dramatically enhances the potency of fluralaner (Fig. 8). These findings may explain the high potency of fluralaner in GABACls, which possess Phe at the equivalent position. Our docking simulation of the *S* enantiomer of fluralaner, which is the active component (Ozoe et al., 2010), to a *Musca* GluCl homology model revealed that the aromatic ring of fluralaner lies near Leu315 (Fig. 10). It remains to be clarified whether the enhanced potency depends on a  $\pi$ - $\pi$  stacking interaction (Zhao et al., 2015) between the aromatic ring of substituted amino acids at position 315 and the phenyl group of fluralaner.

In contrast to the enhancement of the potency of fluralaner, we found that the same aromatic substitution abolished the direct IVM activation of GluCls and the IVM potentiation of glutamate-induced currents, while the antagonism remained unchanged (Fig. 5,9). The L315M mutant, which has a non-aromatic amino acid at position 315, retained the ability to be activated by IVM. This finding is consistent with the fact that the homomeric AVM-sensitive *C. elegans* GluCl- $\alpha$  channel has Met at an equivalent position, whereas the AVM-insensitive  $\beta$  channel has Gln at this position (Cully et al., 1994) (Supplementary Fig. 1). It will be interesting to investigate whether this Gln is

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responsible for the insensitivity to AVMs. The amino acid Thr316 of the *Musca* GluCl subunit (Fig. 2), adjacent to Leu315, is positionally equivalent to the amino acid that was reported to form a hydrogen bond with IVM in an X-ray crystal study of the *C. elegans* GluCl- $\alpha$  channel (Hibbs and Gouaux, 2011). The T316V substitution did not change the potency of IVM in the present study, suggesting that the hydrogen bonding does not contribute substantially to IVM binding in *Musca* GluCls. Instead, our data suggest that Leu315 is located adjacent to bound IVM. How the substitution of the amino acid Leu315 impairs the IVM-induced activation of GluCls and the IVM-induced potentiation of glutamate responses remains to be examined.

Finally, we generated the *Musca* RDL subunit with an inverse mutation (the substitution of Phe with Leu at an equivalent position (Fig. 2A)) to evaluate whether this mutation results in low sensitivity of GABACls to fluralaner. However, because this mutation led to a spontaneously open channel, we were unable to determine the potency of fluralaner in this mutant.

In conclusion, we have shown that Leu315 located in the TSI of *Musca* GluCls and the positionally equivalent amino acid of GABACls play key roles in determining the selectivity of fluralaner and IVM toward these channels. This finding may be applicable to other GluCls and GABACls, given that the TM3 sequence is highly conserved among insect and mite species (Supplementary Fig. 1). As predicted, a very recent publication has indicated that activation by IVM was strongly reduced and that activation by okaramine B, an insecticidal indole alkaloid, was completely abolished in the silkworm (*Bombyx mori*) GluCl containing an L319F mutation, which is equivalent to the L315F mutation in the *Musca* GluCl (Furutani et al., 2017). More importantly, we have shown in the present study that the L315F mutation has opposite impacts on the selectivity of



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fluralaner and IVM for GluCl<sub>s</sub>. This implies that even if IVM-insensitive arthropod pests with an equivalent mutation emerge, these arthropod pests would be sensitive to fluralaner. These findings should prove useful for understanding the mode of action of these parasiticides and further development of pest control agents.

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### **Authorship Contributions**

*Participated in research design:* Nakata, Asahi, Nakahira, F. Ozoe, and Y. Ozoe.

*Conducted experiments:* Nakata, Fuse, Yamato, and F. Ozoe.

*Performed data analysis:* Nakata, Fuse, and Yamato.

*Wrote or contributed to the writing of the manuscript:* Nakata, Fuse, and Y. Ozoe.

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#### Footnotes

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## Figure legends

**Fig. 1.** Chemical structures of fluralaner, its analogs, and ivermectin B<sub>1a</sub>.

**Fig. 2.** Location of the amino acid substitution in *Musca* GluCl<sub>s</sub>. A, Amino acid alignment of the TM1 and TM3 of *Musca* GluCl and GABA<sub>A</sub>Cl (RDL) subunits. Substituted amino acids are highlighted in red. B, Top view of the channel domain with the side chains of substituted amino acids indicated by highlighting. C, Closeup of the side chains of substituted amino acids in the transmembrane subunit interface (TSI).

**Fig. 3.** Inhibition of glutamate-induced currents by fluralaner in wild-type and mutant *Musca* GluCl<sub>s</sub>. A, Dose-response curves of glutamate-induced currents. Data points indicate the means  $\pm$  SEM (n = 6). Normalized relative to maximal current amplitudes. B, Current trace of glutamate (EC<sub>50</sub>)-induced currents during fluralaner perfusion in the wild-type channel. Note that the slight current recovery in the last glutamate application is within the range encompassed by variation. C, Current trace of glutamate (EC<sub>50</sub>)-induced currents during fluralaner perfusion in the L315F mutant. D, Dose-response curves of fluralaner inhibition of agonist-induced current in wild-type and mutant GluCl<sub>s</sub> compared with that in the wild-type GABA<sub>A</sub>Cl. Normalized relative to responses induced by the EC<sub>50</sub>s of agonists. Data points indicate the means  $\pm$  SEM (n = 6).

**Fig. 4.** Responses of wild-type and mutant *Musca* GluCl<sub>s</sub> to IVM. A, Current trace showing the IVM activation of wild-type GluCl<sub>s</sub>. B, Current trace when IVM was applied to the L315F mutant. C, Dose-response curves of IVM-induced currents in GluCl<sub>s</sub>.

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Normalized relative to responses induced by the EC<sub>50</sub>s of glutamate. Data points indicate the means  $\pm$  SEM (n = 6).

**Fig. 5.** Responses of wild-type and mutant *Musca* GluCl<sub>s</sub> to IVM. A, Current trace showing the absence of the IVM potentiation of glutamate (EC<sub>5</sub>) responses in L315F GluCl<sub>s</sub>. B, Dose-response curves for evaluating the IVM potentiation of glutamate responses in wild-type and L315F GluCl<sub>s</sub>. Compared to responses induced by glutamate (EC<sub>5</sub>) alone. Data points indicate the means  $\pm$  SEM (n = 6). C, Current trace showing the IVM inhibition of glutamate (EC<sub>90</sub>)-induced currents in the L315F mutant. D, Dose-response curves of the IVM antagonism of wild-type and L315F GluCl<sub>s</sub>. Normalized relative to responses induced by the EC<sub>90</sub>s of glutamate. Data points indicate the means  $\pm$  SEM (n = 6).

**Fig. 6.** Inhibition of glutamate-induced currents by a fluralaner analog, A1209, in wild-type and mutant *Musca* GluCl<sub>s</sub>. A, Current trace showing the inhibition of GABA (EC<sub>50</sub>)-induced currents in wild-type GABA<sub>A</sub>Cl<sub>s</sub>. B, Current trace showing the inhibition of glutamate (EC<sub>50</sub>)-induced currents in wild-type GluCl<sub>s</sub>. C, Current trace showing the inhibition of glutamate (EC<sub>50</sub>)-induced currents in L315F GluCl<sub>s</sub>. D, Dose-response curves of the A1209 inhibition of glutamate- and GABA-induced currents. Normalized relative to responses induced by the EC<sub>50</sub>s of agonists. Data points indicate the means  $\pm$  SEM (n = 6).

**Fig. 7.** Inhibition of glutamate-induced currents by a fluralaner analog, A341, in wild-type and mutant *Musca* GluCl<sub>s</sub>. A, Current trace showing the inhibition of GABA (EC<sub>50</sub>)-

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induced currents in wild-type GABA<sub>A</sub> receptors. B, Current trace showing the inhibition of glutamate (EC<sub>50</sub>)-induced currents in wild-type GluCl<sub>s</sub>. C, Current trace showing the inhibition of glutamate (EC<sub>50</sub>)-induced currents in L315F GluCl<sub>s</sub>. D, Dose-response curves of the A341 inhibition of glutamate- and GABA-induced currents. Normalized relative to responses induced by the EC<sub>50</sub>s of agonists. Data points represent the means ± SEM (n = 6).

**Fig. 8.** Inhibition of glutamate-induced currents by fluralaner in *Musca* L315 mutant GluCl<sub>s</sub>. A, Dose-response curves of glutamate-induced currents in wild-type and mutant GluCl<sub>s</sub>. Normalized relative to maximal current amplitudes. Data points indicate the means ± SEM (n = 6). B, Current trace of glutamate (EC<sub>50</sub>)-induced currents during fluralaner perfusion in the L315Y mutant. C, Current trace of glutamate (EC<sub>50</sub>)-induced currents during fluralaner perfusion in the L315H mutant. D, Current trace of glutamate (EC<sub>50</sub>)-induced current during fluralaner perfusion in the L315M mutant. E, Dose-response curves of fluralaner inhibition of glutamate-induced currents in wild-type and mutant GluCl<sub>s</sub>. Normalized relative to responses induced by the EC<sub>50</sub>s of glutamate. Data points indicate the means ± SEM (n = 6).

**Fig. 9.** Responses of *Musca* L315 mutant GluCl<sub>s</sub> to IVM. A, Current trace showing the absence of the IVM activation in L315Y GluCl<sub>s</sub>. B, Current trace when IVM was applied to L315H GluCl<sub>s</sub>. C, Current trace showing the IVM activation of L315M GluCl<sub>s</sub>. D, Dose-response curves showing the presence and absence of IVM-induced currents in mutants. Normalized relative to responses induced by the EC<sub>50</sub>s of glutamate. Data points indicate the means ± SEM (n = 6).

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**Fig. 10.** Docking of the *S* enantiomer of fluralaner into the transmembrane subunit interface (TSI) of a wild-type *Musca* GluCl homology model. The  $\alpha$ -helical transmembrane segments (TM1-TM4) of two adjacent subunits are shown in different colors. The docked fluralaner molecule lies near Leu315, suggesting a  $\pi$ - $\pi$  stacking interaction between the phenyl group of fluralaner and the aromatic ring of an amino acid substituted at position 315 of *Musca* GluCl. The CPK coloring is used for the fluralaner stick model.

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TABLE 1

Potencies of glutamate (Glu), fluralaner (Flu), Flu analogs, and IVM in wild-type and mutant forms of *Musca* GluCl<sub>s</sub> and GABA<sub>A</sub>Cl<sub>s</sub> expressed in the *Xenopus* oocytes

Channel	Type	Glu	Flu	A1209	A341	IVM
		EC <sub>50</sub> ( $\mu$ M)	IC <sub>50</sub> (nM)	IC <sub>50</sub> (nM)	IC <sub>50</sub> (nM)	EC <sub>50</sub> (nM)
GluCl	Wild type	8.93 $\pm$ 0.21	146 $\pm$ 14	>10000	>10000	18.8 $\pm$ 5.0 <sup>a</sup>
GluCl	I253A	4.47 $\pm$ 0.32**	308 $\pm$ 21**	NT	NT	4.01 $\pm$ 1.10*
GluCl	M257L	3.32 $\pm$ 0.62**	39.2 $\pm$ 13.5**	NT	NT	6.91 $\pm$ 2.72
GluCl	L315F	323 $\pm$ 57*	1.06 $\pm$ 0.25**	1.63 $\pm$ 0.49	86.9 $\pm$ 10.2	>3000
GluCl	L315Y	81.7 $\pm$ 3.9**	6.07 $\pm$ 1.27**	NT	NT	>3000
GluCl	L315W	NR	NA	NA	NT	NA
GluCl	L315H	63.3 $\pm$ 9.7*	18.0 $\pm$ 1.6**	NA	NT	>3000
GluCl	L315M	12.1 $\pm$ 2.2	77.2 $\pm$ 30.7	NT	NT	33.5 $\pm$ 1.5*
GluCl	L315A	NR	NA	NT	NT	NA
GluCl	T316V	17.7 $\pm$ 1.4**	136 $\pm$ 34	NT	NT	20.2 $\pm$ 2.3
GluCl	M257L/T316V	9.68 $\pm$ 2.09	65.4 $\pm$ 10.7**	NT	NT	NT
GABA <sub>A</sub> Cl	WT	6.97 $\pm$ 1.21	6.05 $\pm$ 1.47	18.8 $\pm$ 6.0	53.2 $\pm$ 9.3	1250 $\pm$ 400 <sup>b</sup>

NR, No response. NA, Not applicable. NT, Not tested. <sup>a</sup>6.79  $\pm$  1.48 nM according to Fuse et al., 2016. <sup>b</sup>Fuse et al., 2016. The data are the means of six experiments  $\pm$  SEM. Unpaired *t*-tests were performed to evaluate statistical significance; *p*-values for L315 mutants are reported with a Bonferroni correction for multiple tests. \**p*<0.05, \*\**p*<0.01 (relative to wild-type GluCl).

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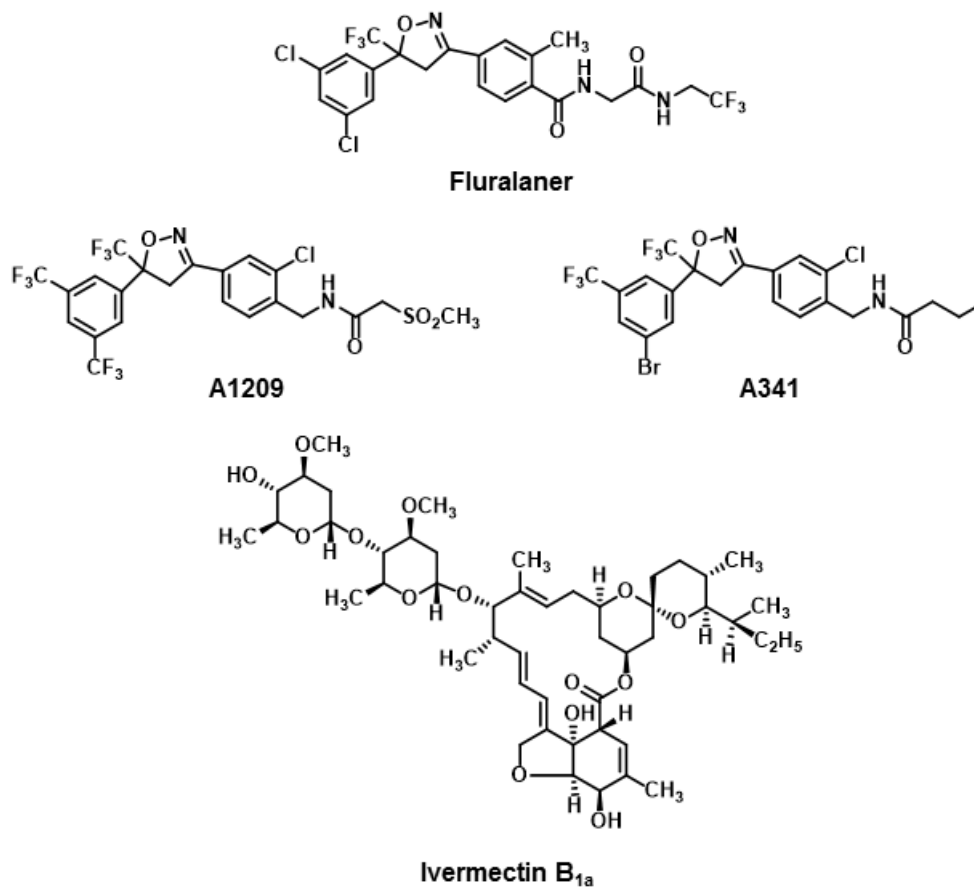


Fig. 1



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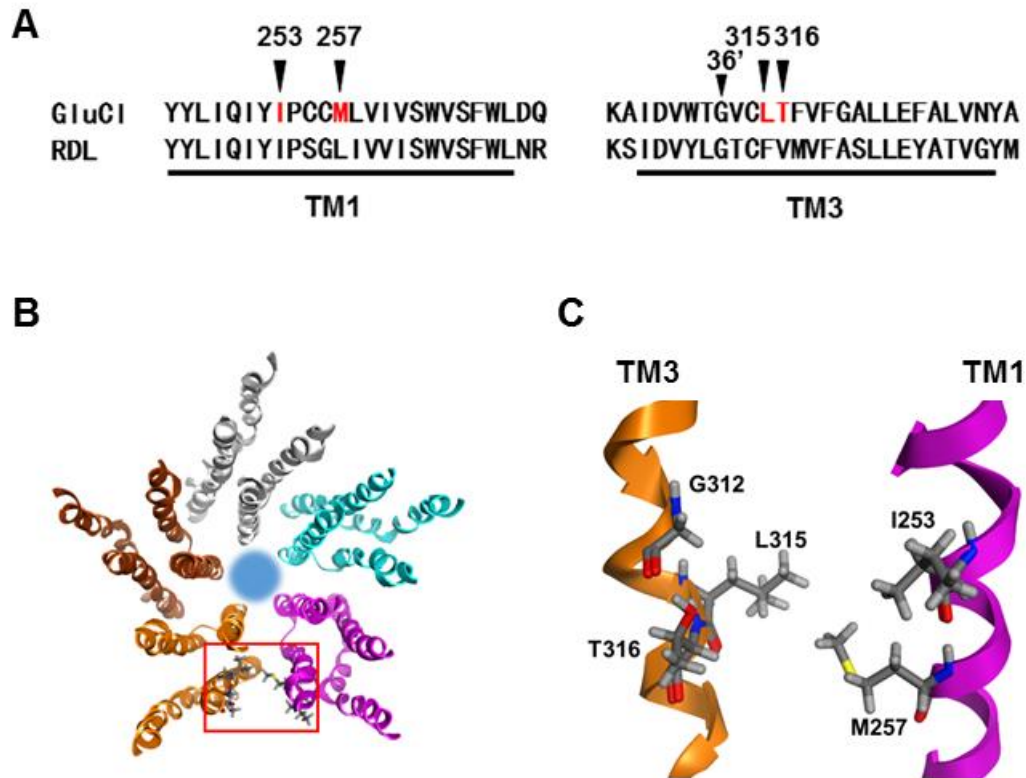


Fig. 2

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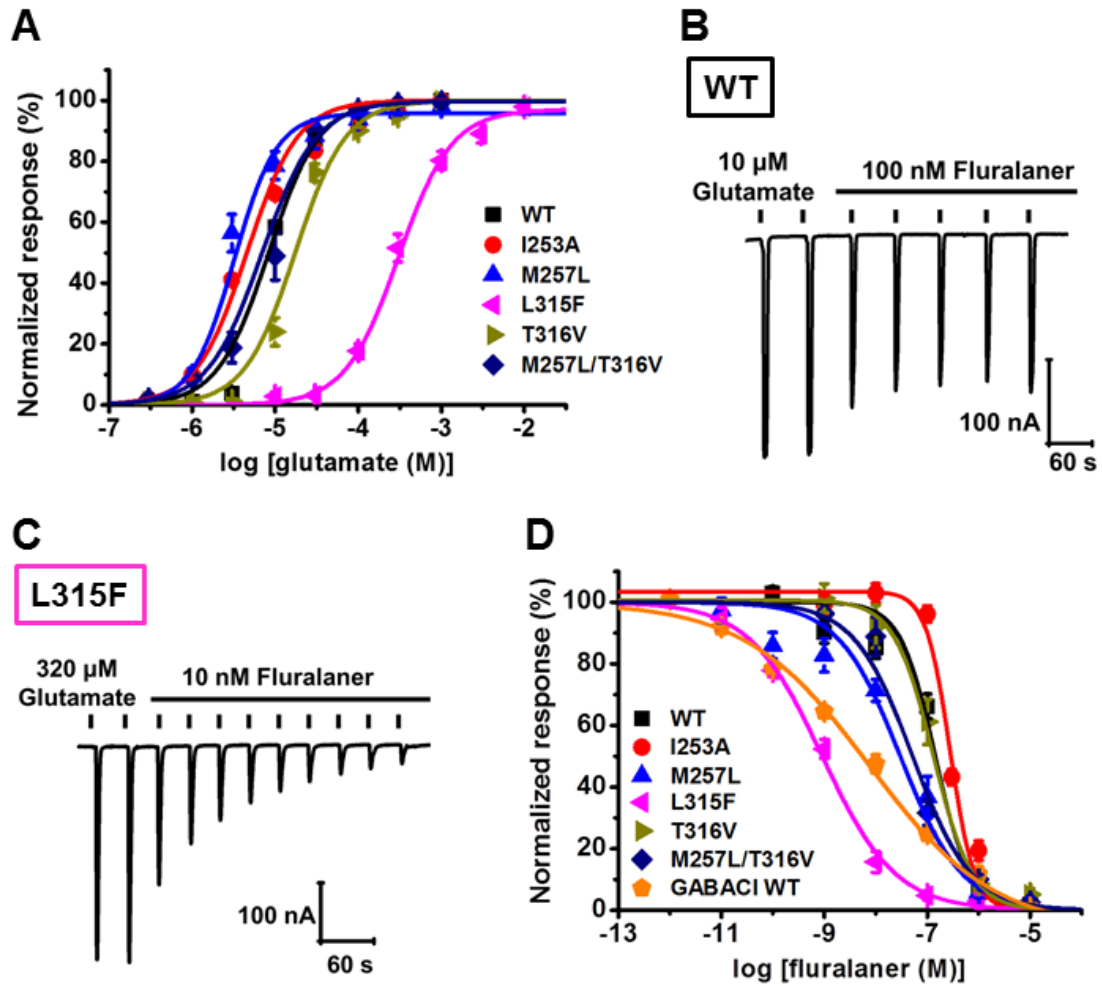


Fig. 3

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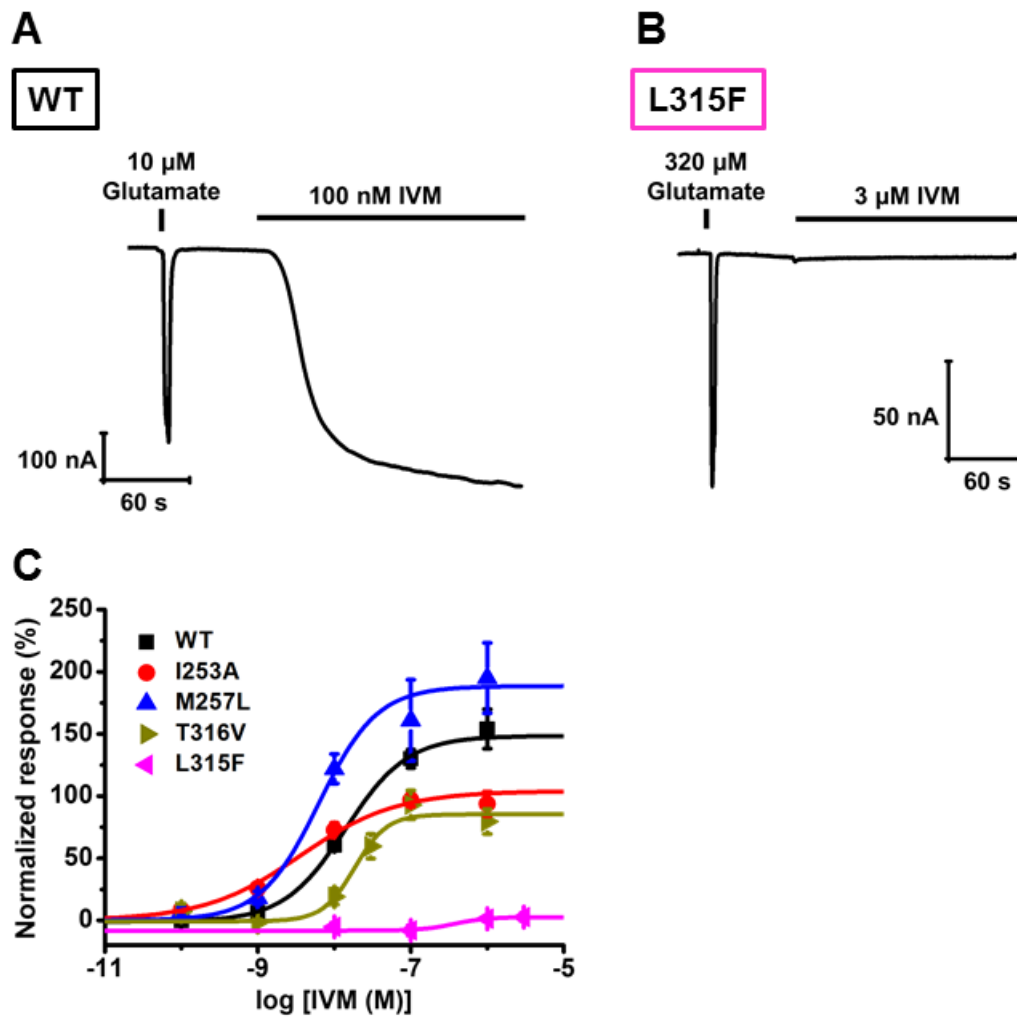


Fig. 4

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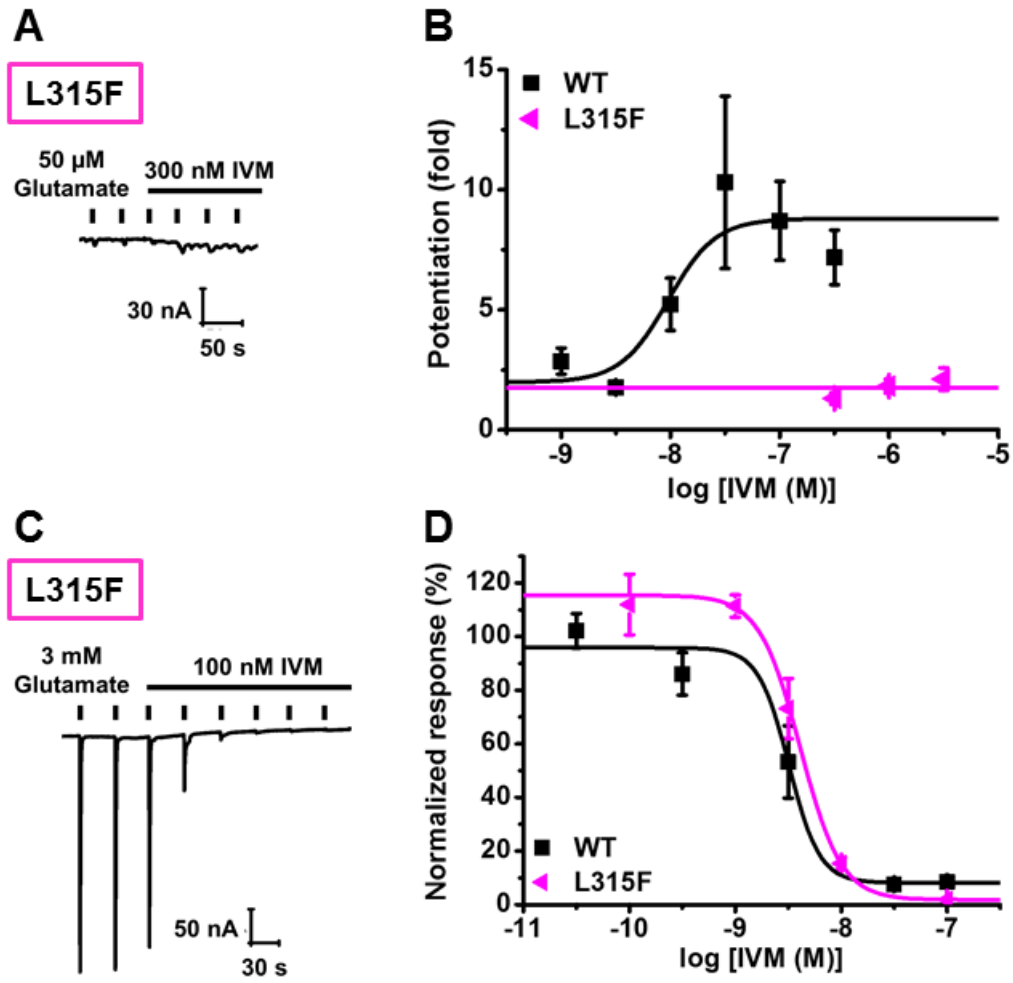


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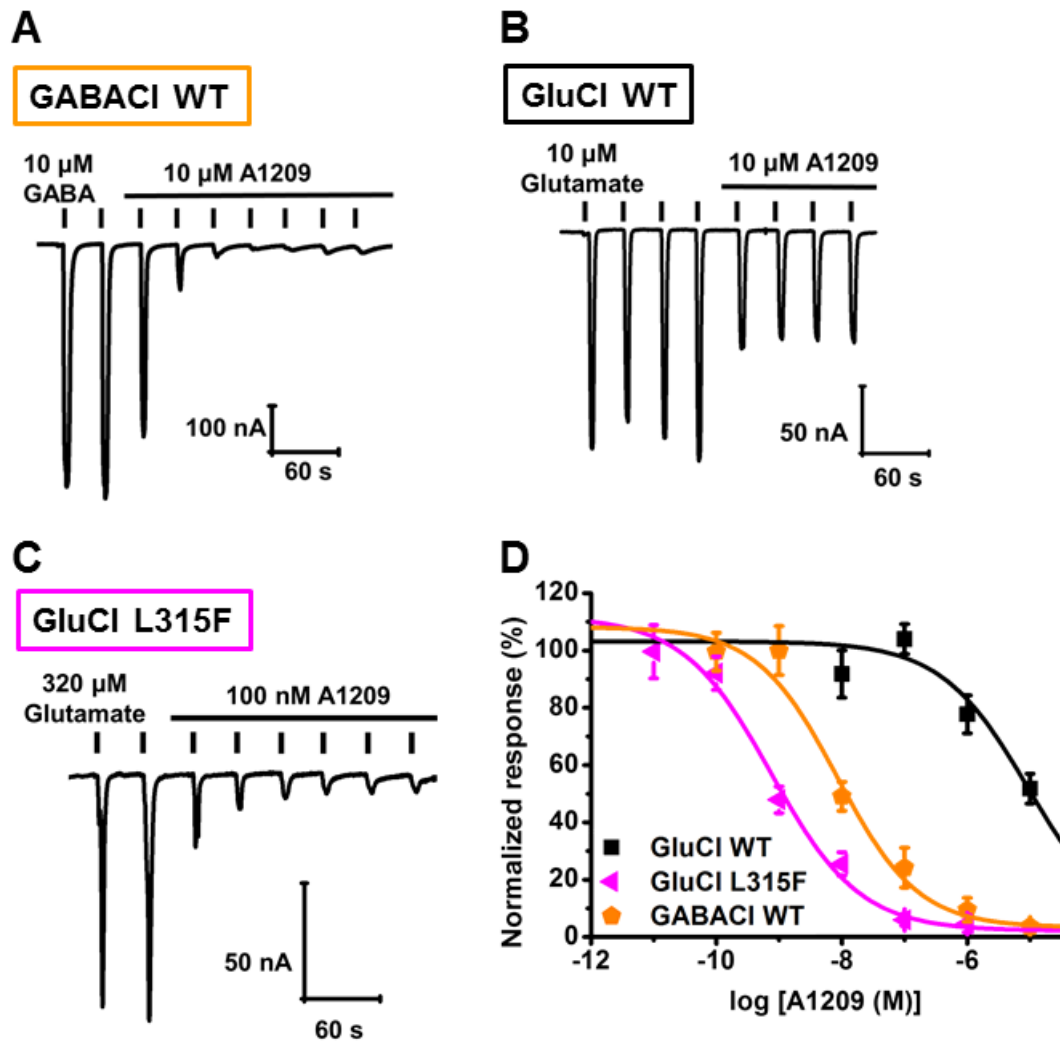


Fig. 6

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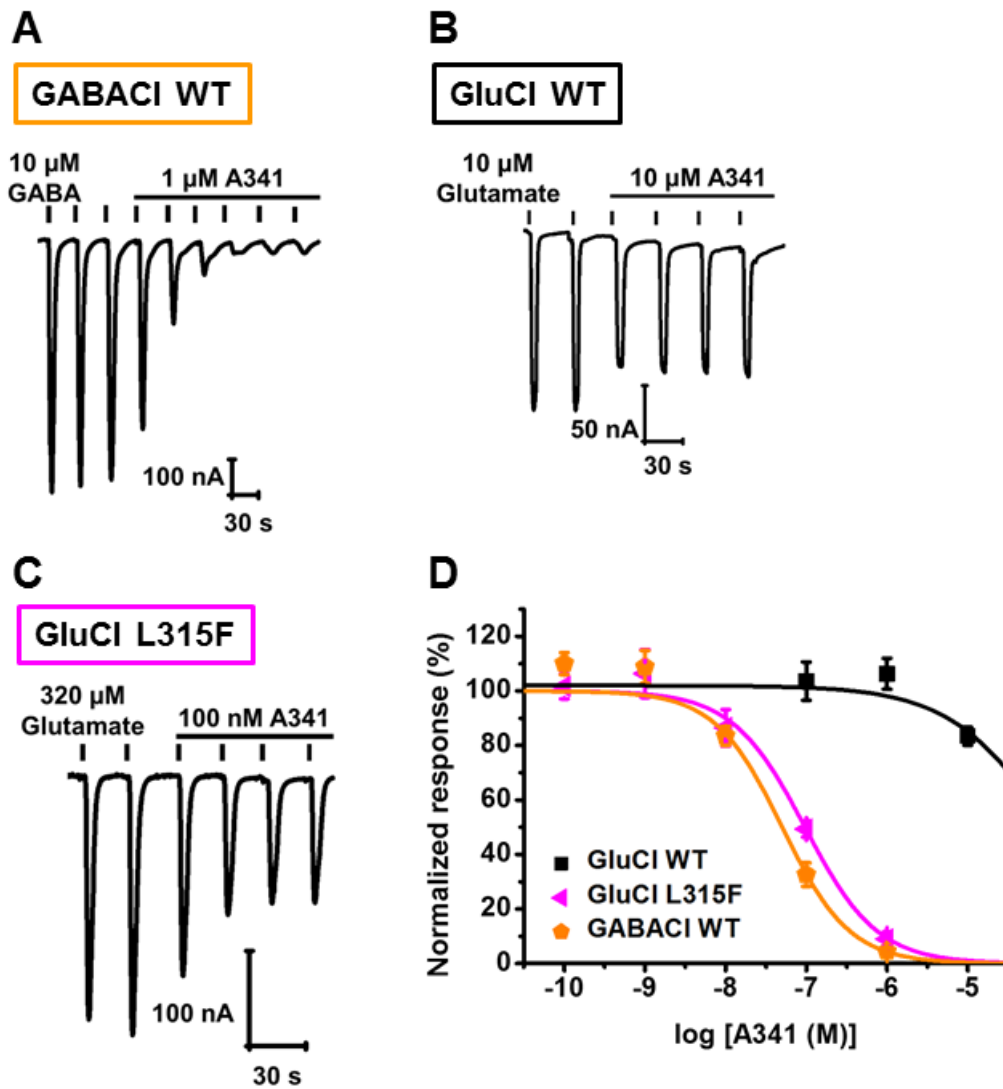


Fig. 7

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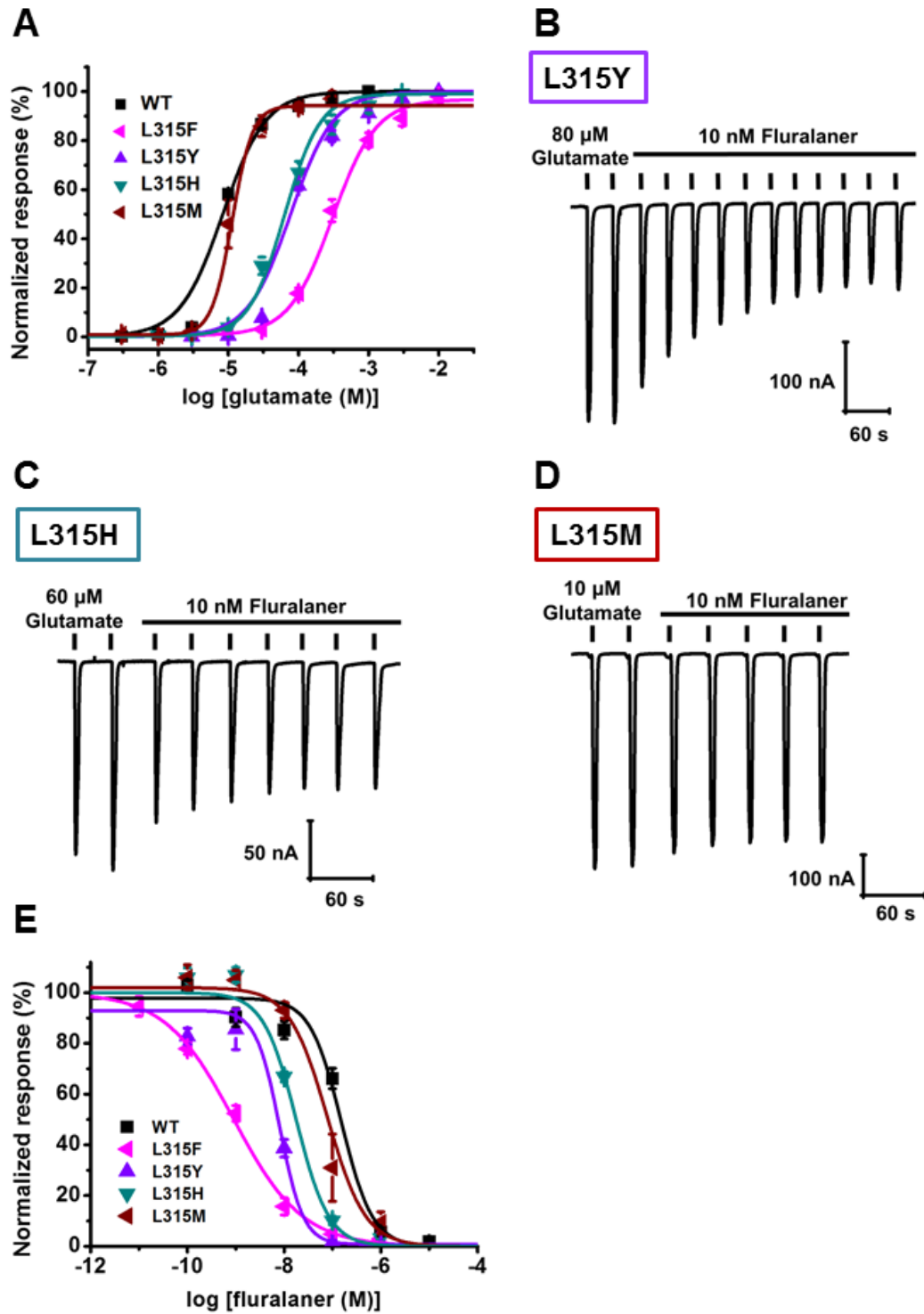


Fig. 8

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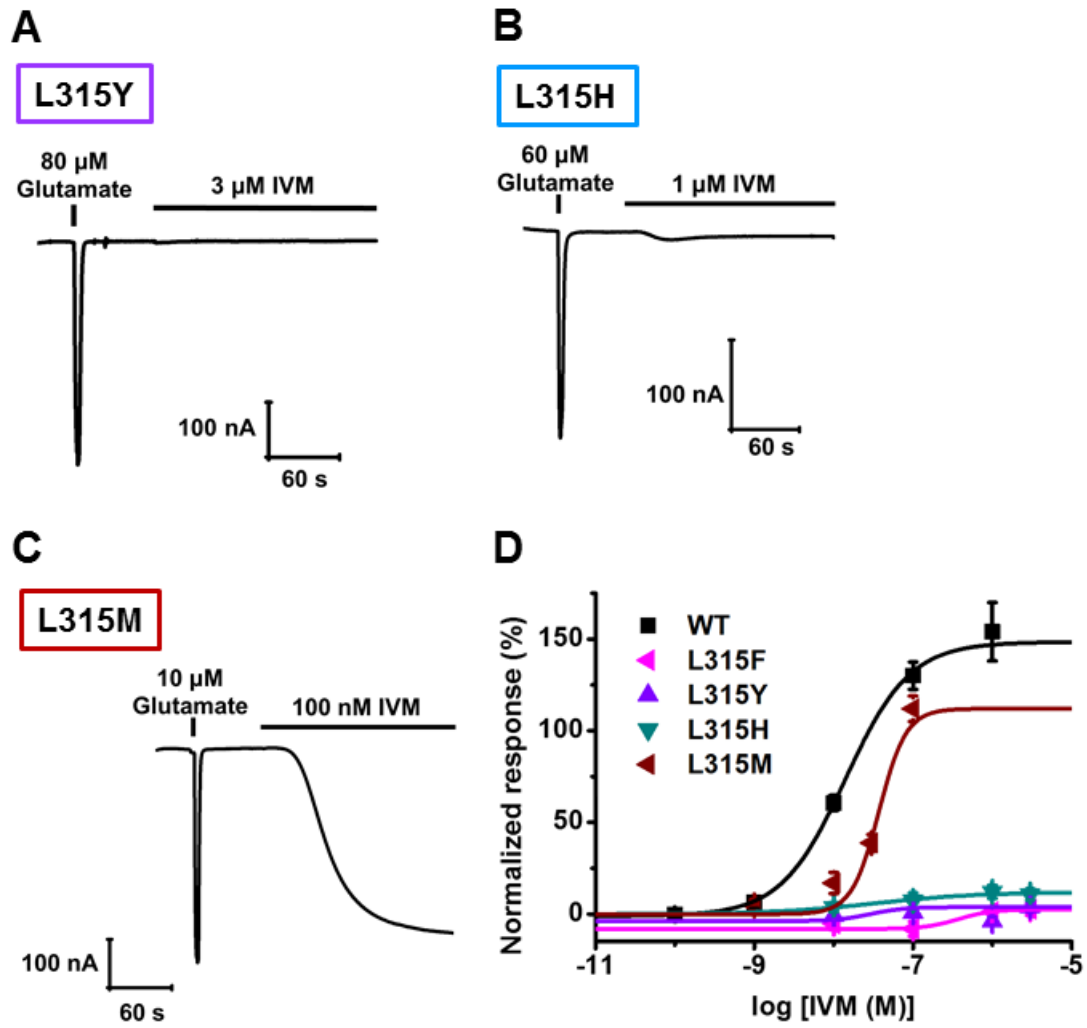


Fig. 9



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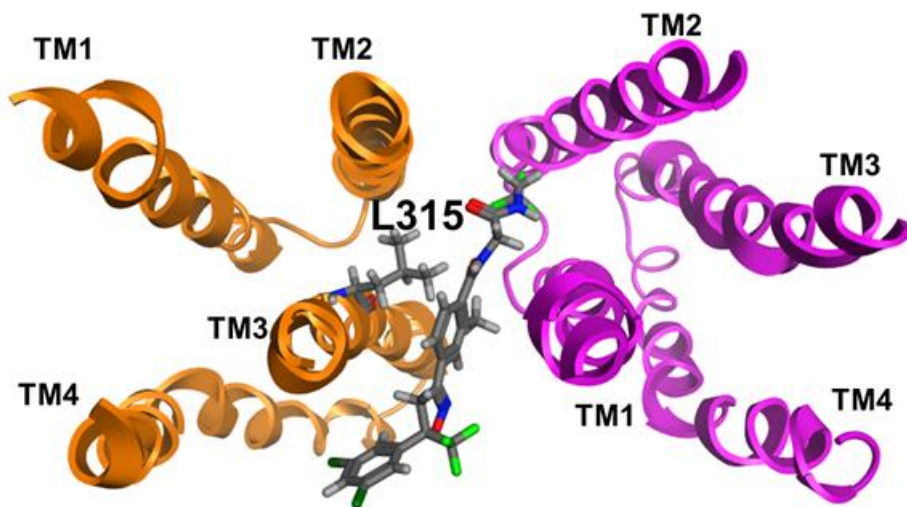


Fig. 10

## Supplementary data

**Title:** A Single Amino Acid Substitution in the Third Transmembrane Region Has Opposite Impacts on the Selectivity of the Parasiticides Fluralaner and Ivermectin for Ligand-Gated Chloride Channels

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**Journal:** Molecular Pharmacology

**GluCl**

<i>Musca domestica</i>		KAIDVWTGVC	TFVFGALLEFALVNYA
<i>Apis mellifera</i>		KAIDVWTGVC	TFVFGALLEFALVNYA
<i>Tetranychus urticae</i>	1	KAVDIWTGCC	IFVFGALIEFAIVNYV
	2	KAIDVWTGVC	AFVFGALLEFALVNYA
	3	KAIDVWTGVC	FFVFGALLEFALVNYA
	4	KAIDVWTESS	TFVFGALLEFALVNYA
	5	KAVDVWTECC	TFVFGALLEFALVNYV
<i>Caenorhabditis elegans</i>	$\alpha$	KAIDVWIGACMTF	IFCALLEFALVNH
<i>Caenorhabditis elegans</i>	$\beta$	KVVDVWLGACQT	TFVFGALLEYAFVSYQ

**RDL**

<i>Musca domestica</i>		KSIDVYLGTC	VMVFASLLEYATVGYM
<i>Periplaneta americana</i>		KSIDVYLGTC	AMVFASLLEYATVGYM
<i>Lucilia cuprina</i>		KSIDVYLGAC	VMVFASLLEYATVGYM
<i>Rhipicephalus microplus</i>		KSIDVYLGTC	VMVFTALLEYAAVGYL
<i>Tetranychus urticae</i>	1	KSIDVFLGTC	VMVFAALLEYAAVGYI
	2	KSIDVFLGTC	VMVFASLLEYATVGYL
	3	KSIDVFLGTC	VMVFASLLEYATVGYL

**Supplementary Fig. 1.** Alignment of the TM3 amino acid sequences of insect, mite, tick, and nematode GluCl and GABACl (RDL) subunits. The key amino acids influencing the fluralaner and IVM selectivity of GluCl and GABACl are indicated by white letters on a black background. Amino acids that differ from that of *Musca domestica* GluCl and GABACl subunits are highlighted in red.

GluCl: *Apis mellifera* (Hymenoptera) (NM\_001077809); *Tetranychus urticae* (Acari) (tetur02g04080, tetur08g04990, tetur10g03090, tetur22g02450, tetur36g00090); *Caenorhabditis elegans*  $\alpha$ -subunit (U14524); *C. elegans*  $\beta$ -subunit (U14525). The TM3 amino acid sequence of the following insects and a tick is identical to that of *M. domestica* (Diptera) GluCl (AB177546): *P. americana* (Blattodea) (LC171470), *Locusta migratoria* (Orthoptera) (DQ643254), *Acyrtosiphon pisum* (Hemiptera) (XM\_016800865), *Laodelphax striatellus* (Hemiptera) (JF430868), *Tribolium castaneum* (Coleoptera) (NM\_001114303), *Anopheles gambiae* (Diptera) (KC902518), *Cochliomyia hominivorax* (Diptera) (KF214912), *Drosophila melanogaster* (Diptera) (U58776), *Lucilia cuprina* (Diptera) (AF081674), *Bombyx mori* (Lepidoptera) (KC342243), *Plutella xylostella* (Lepidoptera) (GQ221939), *Nasonia vitripennis* (Hymenoptera) (FJ851099), and *Rhipicephalus sanguineus* (Acari) (GQ215234, GQ215235).

RDL: *P. americana* (Blattodea) (LC171474); *L. cuprina* (Diptera) (AF024647);

*Rhipicephalus microplus* (Acari) (KF881797); *T. urticae* (Acari) (tetur12g03620, tetur36g00580, tetur36g00590). The TM3 amino acid sequence of the following insects is identical to that of *M. domestica* RDL (AB177547): *L. migratoria* (Orthoptera) (FJ436416), *A. pisum* (Hemiptera) (XM\_001947090), *Cyrtorhinus lividipennis* (Hemiptera) (KJ174466), *L. striatellus* (Hemiptera) (BAF31884), *Nilaparvata lugens* (Hemiptera) (KC841916), *Sogatella furcifera* (Hemiptera) (AB617631), *T. castaneum* (Coleoptera) (NM\_001114337, EF545117), *A. gambiae* (Diptera) (KX431144), *Aedes aegypti* (Diptera) (U28803), *D. melanogaster* (Diptera) (M69057), *B. mori* (Lepidoptera) (AB847423, NM\_001099824), *Heliothis virescens* (Lepidoptera) (AF006189), *P. xylostella* (Lepidoptera) (FJ665610), *Spodoptera litura* (Lepidoptera) (LC171477), *A. mellifera* (Hymenoptera) (KJ485710, KU201321), *N. vitripennis* (Hymenoptera) (XP\_016845785), and *Ctenocephalides felis* (Siphonaptera) (KF881791).