

str-217 (<a +str-217+"#gephebase-summary-title"="" href="https://www.gephebase.org/search-criteria?/and+Gene+Gephebase=">https://www.gephebase.org/search-criteria?/and+Gene Gephebase="str-217"#gephebase-summary-title)	Gephebase Gene	GP00001783	GepheID
Published	Entry Status	Courtier	Main curator

	Trait Category
Physiology (https://www.gephebase.org/search-criteria?/and+Trait+Category=^Physiology^#gephebase-summary-title)	
	Trait
Xenobiotic resistance (insecticide; DEET) (https://www.gephebase.org/search-criteria?/and+Trait=^Xenobiotic resistance (insecticide; DEET)^#gephebase-summary-title)	
	Trait State in Taxon A
sensitive to DEET	
	Trait State in Taxon B
resistant to DEET - Hawaiian strain CB4856	
	Ancestral State
Taxon A	
	Taxonomic Status
Intraspecific (https://www.gephebase.org/search-criteria?/and+Taxonomic+Status=^Intraspecific^#gephebase-summary-title)	

str-217	Generic Gene Name	UniProtKB <i>Caenorhabditis elegans</i>
	Synonyms	GenebankID or UniProtKB <i>Caenorhabditis elegans</i>
CELE_Y102A5C.28; Y102A5C.28	String	
-		
	Sequence Similarities	
-		
	GO - Molecular Function	
GO:0038022 : G protein-coupled olfactory receptor activity (https://www.ebi.ac.uk/QuickGO/term/GO:0038022)		
	GO - Biological Process	
GO:0007186 : G protein-coupled receptor signaling pathway (https://www.ebi.ac.uk/QuickGO/term/GO:0007186)		
GO:0042048 : olfactory behavior (https://www.ebi.ac.uk/QuickGO/term/GO:0042048)		
	GO - Cellular Component	

GO:0005887 : integral component of plasma membrane (https://www.ebi.ac.uk/QuickGO/term/GO:0005887)	Presumptive Null
Yes (https://www.gephebase.org/search-criteria?/and+Presumptive Null=~Yes~#gephebase-summary-title)	Molecular Type
Coding (https://www.gephebase.org/search-criteria?/and+Molecular Type=~Coding~#gephebase-summary-title)	Aberration Type
Deletion (https://www.gephebase.org/search-criteria?/and+Aberration Type=~Deletion~#gephebase-summary-title)	Deletion Size
-	Molecular Details of the Mutation
deletion that leads to a predicted frame shift and early stop codon - the predicted resulting protein has only one transmembrane domain instead of the 7 transmembrane domains - 138-bp deletion in exon 2 and 3 and intervening intron according to https://www.wormbase.org/species/c_elegans/variation/WBVar02076179#02-45-3 and email from Emily Dennis from 9 March 2019	
Candidate Gene (https://www.gephebase.org/search-criteria?/and+Experimental Evidence=~Candidate Gene~#gephebase-summary-title)	Experimental Evidence
A natural variant and engineered mutation in a GPCR promote DEET resistance in <i>C. elegans</i> . (2018) (https://pubmed.ncbi.nlm.nih.gov/30258230)	Main Reference
Dennis EJ; Dobosiewicz M; Jin X; Duvall LB; Hartman PS; Bargmann CI; Vosshall LB	Authors
DEET (N,N-diethyl-meta-toluamide) is a synthetic chemical identified by the US Department of Agriculture in 1946 in a screen for repellents to protect soldiers from mosquito-borne diseases. Since its discovery, DEET has become the world’s most widely used arthropod repellent and is effective against invertebrates separated by millions of years of evolution-including biting flies, honeybees, ticks, and land leeches. In insects, DEET acts on the olfactory system and requires the olfactory receptor co-receptor Orco, but exactly how it works remains controversial. Here we show that the nematode <i>Caenorhabditis elegans</i> is sensitive to DEET and use this genetically tractable animal to study the mechanism of action of this chemical. We found that DEET is not a volatile repellent, but instead interferes selectively with chemotaxis to a variety of attractant and repellent molecules. In a forward genetic screen for DEET-resistant worms, we identified a gene that encodes a single GÅ protein-coupled receptor, str-217, which is expressed in a single pair of chemosensory neurons that are responsive to DEET, called ADL neurons. Mis-expression of str-217 in another chemosensory neuron conferred responses to DEET. Engineered str-217 mutants, and a wild isolate of <i>C. elegans</i> that carries a str-217 deletion, are resistant to DEET. We found that DEET can interfere with behaviour by inducing an increase in average pause length during locomotion, and show that this increase in pausing requires both str-217 and ADL neurons. Finally, we demonstrated that ADL neurons are activated by DEET and that optogenetic activation of ADL neurons increased average pause length. This is consistent with the ‘confusant’ hypothesis, which proposes that DEET is not a simple repellent but that it instead modulates multiple olfactory pathways to scramble behavioural responses. Our results suggest a consistent motif in the effectiveness of DEET across widely divergent taxa: an effect on multiple chemosensory neurons that disrupts the pairing between odorant stimulus and behavioural response.	
	Additional References

RELATED GEPHE

3 (acetyl-CoA carboxylase (ACC), beta-tubulin (ben-1), GLC-1) (https://www.gephebase.org/search-criteria?/or+Taxon ID=~6239~/and+Trait=Xenobiotic resistance/and+groupHaplotypes=true#gephebase-summary-title)	Related Genes
1 (https://www.gephebase.org/search-criteria?/or+Gene Gephebase=~str-217~/and+Taxon ID=~6239~/or+Gene Gephebase=~str-217~/and+Taxon ID=~6239~#gephebase-summary-title)	Related Haplotypes

EXTERNAL LINKS

COMMENTS

https://www.wormbase.org/species/c_elegans/variation/WBVar02076179#02-45-3 - other *C. elegans* strains carry the mutation Gln5*