

Genome Organization, Phylogenies, Expression Patterns, and Three-Dimensional Protein Models of Two Acetylcholinesterase Genes from the Red Flour Beetle

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Abstract

Since the report of a paralogous acetylcholinesterase (AChE, EC3.1.1.7) gene in the greenbug (*Schizaphis graminum*) in 2002, two different AChE genes (*Ace1* and *Ace2*) have been identified in each of at least 27 insect species. However, the gene models of *Ace1* and *Ace2*, and their molecular properties have not yet been comprehensively analyzed in any insect species. In this study, we sequenced the full-length cDNAs, computationally predicted the corresponding three-dimensional protein models, and profiled developmental stage and tissue-specific expression patterns of two *Ace* genes from the red flour beetle (*Tribolium castaneum*; *TcAce1* and *TcAce2*), a globally distributed major pest of stored grain products and an emerging model organism. *TcAce1* and *TcAce2* encode 648 and 604 amino acid residues, respectively, and have conserved motifs including a choline-binding site, a catalytic triad, and an acyl pocket. Phylogenetic analysis shows that both *TcAce* genes are grouped into two insect *Ace* clusters and *TcAce1* is completely diverged from *TcAce2*, suggesting that these two genes evolve from their corresponding *Ace* gene lineages in insect species. In addition, *TcAce1* is located on chromosome 5, whereas *TcAce2* is located on chromosome 2. Reverse transcription polymerase chain reaction (PCR) and quantitative real-time PCR analyses indicate that both genes are virtually transcribed in all the developmental stages and predominately expressed in the insect brain. Our computational analyses suggest that the *TcAce1* protein is a robust acetylcholine (ACh) hydrolase and has susceptibility to sulfhydryl agents whereas the *TcAce2* protein is not a catalytically efficient ACh hydrolase.

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Introduction

Acetylcholinesterase (AChE, EC3.1.1.7) is an essential enzyme at the synapses of cholinergic neurons in the central and peripheral nervous systems in all animals. It catalyzes the hydrolysis of the neurotransmitter acetylcholine (ACh), thus terminating neurotransmission. AChE has long been of academic and industrial interest and studied extensively at the biochemical, biophysical, and molecular levels in mammals because this enzyme is a target of palliative Alzheimer drugs, nerve agents, and insecticides [1]. In insects, AChE has also been extensively studied because it serves as the target site for organophosphate and carbamate insecticides, and involves in insecticide resistance known as target-site insensitivity [2–8].

The first insect AChE gene (*Ace*) was sequenced from *Drosophila melanogaster* in 1986 [9]. After the first *Ace* paralogous gene was reported in the greenbug (*Schizaphis graminum*) in 2002 [10], the *D. melanogaster Ace* gene was designated as *Ace* orthologous gene. It is now clear that *D. melanogaster* has only one *Ace* gene as confirmed by its genome sequence [11], whereas most other insect species have two different *Ace* genes (i.e., *Ace1* and *Ace2*) [12]. *Ace1* commonly refers to the *Ace* paralogous (AP-*Ace*) gene and *Ace2* the

Ace orthologous (AO-*Ace*) gene in relation to the *D. melanogaster Ace* [8].

To date, the cDNAs encoding AChEs have been sequenced from at least 43 insect species. Among them, both *Ace1* and *Ace2* have been reported from each of 27 species, including *Bombyx mandarina* (GenBank accession numbers: EU262633 for *BmAce1* and EU262632 for *BmAce2*); *Sitobion avenae* [13]; *Rhopalosiphum padi* [13]; *Anopheles gambiae* [14]; *Liposcelis decolor* (GenBank accession numbers: FJ647186 for *LdeAce1* and FJ647187 for *LdeAce2*); *Orchesella villosa* [15]; *Liposcelis entomophila* (GenBank accession numbers: EU854149 for *LeAce1* and EU854150 for *LeAce2*); *Blattella germanica* [16]; *Bemisia tabaci* [17]; *Culex pipiens quinquefasciatus* (GenBank accession numbers: XM_001847396 for *CqAce1* and XM_001842175 *CqAce2*); *Bombyx mori* [18]; *Acyrtosiphon pisum* (GenBank accession numbers: XM_001948618 for *ApAce1* and XM_001948953 for *ApAce2*); *Nasonia vitripennis* (GenBank accession numbers: XM_001600408 for *NvAce1* and XM_001605518 for *NvAce2*); *Pediculus humanus corporis* [19]; *Cydia pomonella* [20]; *Helicoverpa assulta* [21]; *Aedes albopictus* (GenBank accession numbers: AB218421 for *AaAce1* and AB218420 for *AaAce2*); *Aphis gossypii* [22]; *Culex tritaeniorhynchus* [23]; *Myzus persicae* [24]; *Culex pipiens* [25]; *Plutella xylostella* [26–28]; *Chilo suppressalis* [29]; *Pediculus*

humanus capitis [19]; *Aedes aegypti* [30]; *Liposcelis bostrychophila* (GenBank accession numbers: FJ647185 for *LbAce1* and EF362950 for *LbAce2*) and *Alphitobius diaperinus* [31]. The remaining insect species may also have two *Ace* genes but only one *Ace* gene (*Ace1* or *Ace2*) has been documented.

The existence of two *Ace* genes in insects has attracted much attention to the study of their functions, particularly their roles in insecticide resistance [8,32] and as targets for developing new insecticides [12,33–37]. Beetles (coleopterans) are the most evolutionarily successful metazoans, accounting for 25% of all known animal species, far more than any other taxonomic orders [31]. Despite the diversity and economic importance of coleopterans, *Ace* genes have been reported from only two species: *Leptinotarsa decemlineata* [5,6] and *Alphitobius diaperinus* [31]. Although *T. castaneum* (the red flour beetle) is one of the most notorious stored grain pests in the world and is now regarded as an emerging model organism, its *Ace* genes were only predicted from genomic sequence and detailed information on these genes has been limited.

In this paper, we report two *Ace* genes from *T. castaneum*. Our study of the two genes focuses on the genome organization, three-dimensional (3D) protein models, phylogenies, and expression patterns of the two genes at different developmental stages of the insect, in an effort to better understand the functions of the two genes and obtain insights into better strategies for insect pest control.

Results

ACHE cDNA and deduced amino acid sequences

Based on the predicted sequences of two *T. castaneum Ace* genes in NCBI (XM_968369 and XM_965681), we designed specific

primers (Table 1) to determine the full-length cDNAs of the two genes from the brain of *T. castaneum*. Each of several polymerase chain reaction (PCR) primer pairs was able to generate overlapping fragments for each gene and it was then possible to assemble them into its full-length cDNA of the protein coding region. The two deduced amino acid sequences show significant similarities to AChE1 (AP-AChE) and AChE2 (AO-AChE) proteins of other insects in GenBank according to our BLASTP analysis. Therefore, the two *T. castaneum Ace* genes are named *TcAce1* (AP-*Ace*) and *TcAce2* (AO-*Ace*), and their protein products are named TcAce1 and TcAce2, respectively. The *TcAce1* cDNA contains 2148 base pairs (bp) and has an open reading frame (ORF) of 1944 bp, encoding a protein of 648 amino acid residues, whereas the *TcAce2* cDNA contains 1,890 bp and has an ORF of 1,812 bp, encoding a protein of 604 residues. However, we were not able to obtain the 5'-untranslated region (5'-UTR) of the *TcAce2* cDNA (Fig. 1).

TcAce1 and *TcAce2* belong to typical *Ace1*- and *Ace2*-type genes, respectively, as judged by their sequence similarities with other known insect *Aces* (Fig. 2). The deduced amino acid sequences (TcAce1 and TcAce2) of *TcAce1* and *TcAce2* exhibit six and four *N*-glycosylation sites (N-X-S or N-X-T) [38], respectively (Fig. 1). Predicted isoelectric points (pI) and molecular masses of TcAce1 and TcAce2 are 6.58 and 5.39, and 72.81 and 68.15 kDa, respectively (<http://www.scripps.edu/~cdputnam/protcalc.html>). Both TcAce1 and TcAce2 are predicted to contain a cleavable signal peptide, which suggests that these proteins can be secreted and function in an extracellular environment. Both proteins have a C-terminal Cys residue (C617 in TcAce1 and C600 in TcAce2) that is likely to form an intermolecular disulfide bond. According to the analysis using PredGPI (<http://gpcr.biocomp.unibo.it/>)

Table 1. PCR primers used to amplify cDNA sequences of both *TcAce1* and *TcAce2* genes and to analyze their gene expressions.

Primer name	Sequence (5'-3')	Tm (°C)	Product size (bp)	Location ^a
<i>PCR for cDNA sequences</i>				
<i>TcAce1</i> -F ^b	CGGCCTGCTTACTAGTGATTCTAC	60.66	1714	17-1730
<i>TcAce1</i> -R	ACATCGAGGGTGAGAACTCC	60.50		
<i>TcAce1S</i> -F	CAACGACCGTTGTGCAAATA	60.56	320	N/A-219
<i>TcAce1S</i> -R	CAGGGGATCATCTTCGGAGT	61.39		
<i>TcAce1E</i> -F	CGTTTGGACACCCACTTTCT	60.01	402	1665-N/A
<i>TcAce1E</i> -R	GTCGTGTTGATTTGAATACCTCAC	60.99		
<i>TcAce2</i> -F	AGACCTCATCACGCTGTTTG	55.2	1034	750-1783
<i>TcAce2</i> -R	CTGGGTTATCCCGAAGCTTG	61.86		
<i>TcAce2S</i> -F	GTCGTAGAGGCGTGCCTG	61.44	1014	N/A-895
<i>TcAce2S</i> -R	TCTCCCCGACATGTAAGTCT	59.93		
<i>TcAce2E</i> -F	AACCAGTGACAGACGACGTG	59.78	267	1627-N/A
<i>TcAce2E</i> -R	CGCAACCGATGCGTTAATA	61.86		
<i>Quantitative real-time PCR</i>				
<i>TcAce1</i> (Q)-F	CCGTTGTCGCCAGTCATTG	55.3	121	1069-1189
<i>TcAce1</i> (Q)-R	AGTAGTAGCCTTCTTCTGTGTTAG	55.4		
<i>TcAce2</i> (Q)-F	AGACCTCATCACGCTGTTTG	55.2	179	750-928
<i>TcAce2</i> (Q)-R	CCTCCACCAGGACCTTCC	54.9		
<i>TcRps3</i> -F	CCGTCGTATTCGTGAATTGAC	54.8	130	279-408
<i>TcRps3</i> -R	TCTAAGAGACTCTGCTTGTGC	54.7		

^aProduct location refers to the PCR fragment corresponding to the *Ace* gene nucleotide sequence of *T. castaneum* from NCBI database (*TcAce1*: XM_968369; *TcAce2*: XM_965681). N/A refers to the sequence based on the genomic sequence in Beetlebase (<http://beetlebase.org/>).

^bF and R refer to forward and reverse primers, respectively.

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Figure 1. The cDNA and deduced amino acid sequences of two *Ace* genes from *Tribolium castaneum*. The amino acid sequences were numbered from the start of the mature proteins. The start codon ATG were bold and underlined, and the stop codon TAA at the end of the coding region were bold and marked with asterisks. The putative signal peptides of the deduced amino acid sequences were underlined with red dots. Potential N-linked glycosylation sites were bold and shaded. The sequences were deposited in the GenBank (accession numbers: HQ260968 for *TcAce1* and HQ260969 for *TcAce2*). doi:10.1371/journal.pone.0032288.g001

predgpi/), *TcAce1* appears to contain a GPI-anchor, which is linked to the C-terminal residue after a proteolytic cleavage at the ω site (D619 in *TcAce1*), whereas *TcAce2* doesn't seem to contain a GPI-anchor. Both proteins have relatively high sequence identity (47% for *TcAce1* and 59% for *TcAce2*) to the C-terminal residues 526–543 (*i.e.*, QTCAFVNRFLPKLLSAT) of the recombinant mouse AChE (mAChE) [39,40]. This level of sequence identity is identical or higher than the corresponding sequence identity (47%) between mAChE and *D. melanogaster* AChE (DmAChE). All these suggest that residues 589–605 in *TcAce1* and residues 578–594 in *TcAce2* are likely responsible for the formation of a dimeric four-helix bundle at the C-terminus as seen in mAChE [40] and DmAChE [41].

Chromosomal locations of *TcAce1* and *TcAce2*

The exon-intron organizations of *TcAce1* and *TcAce2* were revealed by comparisons of the full-length cDNAs with their corresponding genomic sequences (<http://beetlebase.org/>). The lengths of *TcAce1* and *TcAce2* genomic DNA sequences are 2,986 bp and 32,243 bp, respectively. Genome structure analysis shows that the two genes are located on different chromosomes of *T. castaneum*; *TcAce1* is located on chromosome 5, whereas *TcAce2* on chromosome 2. *TcAce1* has two exons and one intron, whereas *TcAce2* has six exons and five introns (Fig. 3).

Phylogenetic relationship of *T. castaneum* AChE to other AChEs

The phylogenetic tree of the deduced amino acid sequences of AChEs from the Pacific electric ray (*Torpedo californica*), twospotted spider mite (*Tetranychus urticae*), and all the insect species available in GenBank was generated using the neighbor-joining method. Phylogenetic analysis suggests that there are two major groups (Ace1 and Ace2); *TcAce1* and *TcAce2* belong to the *Ace1*- and *Ace2*-type genes, respectively (Fig. 4). As expected, the cDNA-deduced *TcAce1* has high protein sequence identities to BgAce1 (70%), HaAce1 (64%), AgAce1 (61%), SaAce1 (57%), and SgAce1 (57%). The cDNA-deduced *TcAce2* has high protein sequence identities to LdAce2 (84%), HaAce2 (69%), BgAce2 (68%), AgAce2 (60%), SaAce2 (57%), and DmAce (55%) (Table 2). *TcAce1* and *TcAce2* exhibited 39% and 38% protein sequence identities to *T. californica* AChE, respectively. However, the protein sequence identities of Ace1 and Ace2 in the same insect species are 36% (between *TcAce1* and *TcAce2*), 31% (between SaAce1 and SaAce2), 35% (between AgAce1 and AgAce2), 35% (between BgAce1 and BgAce2), and 32% (between HaAce1 and HaAce2) (Table 2).

It is worth noting that, per the present nomenclature, the reported PhAce1, PhcAce1, AgoAce1, MpAce1 and CtAce1 in the Ace2 group should change to PhAce2, PhcAce2, AgoAce2, MpAce2 and CtAce2, respectively, whereas the reported PhAce2, PhcAce2, AgoAce2, MpAce2 and CtAce2 in the Ace1 group should be named PhAce1, PhcAce1, AgoAce1, MpAce1 and CtAce1, respectively (Fig. 4).

Three-dimensional models

TcAce1 (TcAP-AChE). Eighteen 10-ns molecular dynamics simulations of the substrate-bound *TcAce1* protein homology

model derived from a human butyrylcholinesterase (hBChE) crystal structure (Protein Data Bank ID: 2J4C [42]) resulted in a time-averaged model with a distorted catalytic triad and partial unfold of the omega loop. This result indicates that the homology model is structurally unstable, presumably due to the low sequence identity of the omega loop between *TcAce1* and hBChE that results in gaps in the omega loop. Another homology model was therefore built from a simulation-refined model of the African malaria mosquito (*A. gambiae*) AP-AChE (Protein Data Bank ID: 2AZG) [33] that has a higher sequence identity to *TcAce1* (73%) than the identity between hBChE and *TcAce1* (46%). Twenty-two 10-ns molecular dynamics simulations of this model liganded with its substrate yielded a time-averaged model without distortions in the catalytic triad and the omega loop. In this final model of *TcAce1* (Fig. 5), C354 is exposed to solvent and accessible to covalent bonding at the opening of the active-site gorge [33]; R407 is enclosed by F143, F146, and F406 via cation-pi interactions; Y189, Y396, and Y400 adopt conformations that make the gorge relatively open; ACh adopts the fully extended conformation with its ammonium group forming a cation-pi interaction with the indole ring of W152 and its carbonyl oxygen atom anchored at the oxyanion hole comprising of G186, G187, and A268; E266 forms a hydrogen bond with Y198 at the bottom of the active-site gorge; E393, H507, and S267 form the catalytic triad. The *TcAce1* active site is very similar to those of *A. gambiae* and *S. graminum* AP-AChEs [33,34], and different from the human AChE active site in that Y449 in the human enzyme is replaced by Asp creating void space at the bottom of the *TcAce1* active site (Fig. 6). In *T. californica* AChE, rotation of Y442, which corresponds to Y449 of human AChE, reportedly controls the opening of a 3.4-Å-wide channel that enables rapid clearance of substrate hydrolysis products [43]. In DmAChE, the counterpart of Y449 is also mutated to Asp; hence the crystal structure of DmAChE (Protein Data Bank ID: 1DX4 [41]) has a channel with a diameter of ~5 Å that is formed by G79, W83, W472, L479, and D482 and connects the active-site gorge to solvent [44]. Because of the mutation Tyr of human to Asp of *TcAce1*, the final model of *TcAce1* has a similar channel that is comprised of G148, W152, W499, M506, and D509. The *TcAce1* channel is, however, partially blocked by M151 and P498.

TcAce2 (TcAO-AChE). A homology study identified the crystal structure of DmAChE (PDB ID: 1DX4 [41]) as a template with a sequence identity of 60% and generated a homology model of *TcAce2*. This model has a protruded large loop conformation for residues 145–162, which is due to the omission of the corresponding loop (residues 103–136) in the 1DX4 crystal structure. An initial set of 21 10-ns molecular dynamics simulations of the *TcAce2* homology model without residues 145–162 resulted in a time-average model with the catalytic triad distorted.

In the loop of residues 145–162, there are two histidine residues, four arginine residues, two lysine residues, one aspartate residue, and one glutamate residue. At the physiological pH of 7.4, this loop has a net charge of +4. To avoid a possible effect of the highly-charged loop on the catalytic triad distortion, a second set of 21 10-ns simulations of the *TcAce2* homology model possessing residues 145–162 was performed. The triad was still distorted in an average conformation of all trajectories saved at 1.0-ps intervals

DmAce	-----MATSCRQSRVLPMSLPLPLT-----TPLPLVLLG-----	29
AgAce2	-----MASAYYHQASVAGVGN-----VLVLLG-----	22
TcAce2	-----MGSNLVVVVVVVVVAVSASVRAVSWP-----SEETT-----	34
LdAce2	-----MG-----QLSTLCLFVTVASVCGVSWP-----SDETT-----	28
HaAce2	-----MPVRTTIFEMISNTKIVFTKLLCCFVSGAVARSWANH-----HDTTSTTQTTP-----	50
BgAce2	-----MATQCLLLAGCICTAMAIAGSKPHFNSSASSTP-----HSTIGGGSP-----	46
SaAce2	-----MSVDVYTSAVTLLCCSAVLRPSSGGADGGGGGGGGGAGGGGGGGGG-----	53
TcAce1	-----MTG-AWAAC-----LLVILLPSPICPSHRGRHHPPEHAETHMSRDPDPHRS-----EERFRADAPDK-----	60
BgAce1	-----MDVGDVG-GLSVSGSDTDLPLRDIHREHYIKDHHVSHHSGGGGDPYSMSRSDPSVGRGPEMDMRRDPVDFRRHSIDMDR-----	83
HaAce1	-----MRVVLAALTAALARALAGPHEHRAKHHAEPHPFPAPAPPQYRGHGEAVRYNPELDTILRPLEDEHETSSKRAKFEAEATSSKRAKYD-----	89
SaAce1	-----MDQWLLWFGVLAFTVGLSLRHARHQVSGTPTVEEILEPQLIEDTDHVFQRASDMFAQPEYTEKRNLNHRRSSEFSNGQ-----	82
SgAce1	-----MDQWLLWFGVLAFTVGLSLRHARHQVSGTPTVEEILEPQLIEDTDHVFQRASDMFAQPEYTEKRNLNHRRSSEFSNGQ-----	82
AgAce1	-----MEIRGLMLGRRLRGRVPLGLLGVTAALLLPPFVALVGRHIELNNGAAGSHQLSAAAGVGLASQAGSSVMSVPAAGASSSSSSLLSSAEDDVARITLSDADAFPTPYGHGESVRTIDAEELGT-----	136
TcaAce	-----MNLVTSLSGLVLLHLVLCQADHS-----	25
DmAce	-----SLHLSCVGVDIRLVVQTSSGPRGR-SVTYQGREVHYTGYPAKPPVEDLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	143
AgAce2	-----ATVCPAYAILDRLVVQTSSGPIRGR-SVTYQGREVHYTGYPAKPPVEDLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	136
TcAce2	-----TRPPQARDFHSDDLVEVETSSGLARGK-AKTVLGREVHYTGYPAKPPVEIQLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	148
LdAce2	-----TKPSQKDFHTDPLVETSSGLVARGY-SKTVLGREVHYTGYPAKPPVEIQLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	142
HaAce2	-----TTSVPYKFNHNDPLIVETKSGLVKGY-AKTVMGREVHYTGYPAKPPVEIQLRFKRPVPAEPWHG-----VLEATAMPNSIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	164
BgAce2	-----NTNPKGLDYHDDLVEVETSSGLVARGY-ARVVLKGEVHYTGYPAKPPVEIQLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	160
SaAce2	-----GGGSAVDVDTDESPPVYVSSGMYGQY-TKIANKRVRYTGYPAKPPVEIQLRFKRPVPAEPWHG-----VLDATGLSATIQERYEYFPFGSGEEIWPNTNVSEDIYINWVPAKAR-----	167
TcAce1	-----REFTRDSEDDPLVIQTKKGRVIRGSLTAATGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----VMNTTSQPNSIQIDTVFGDFPGATMWNPNVPLNEDIYINWVPAKAR-----	173
BgAce1	-----YGHGGDPLDLAHESRMSDESGHGMQDDPLVIQTKKGRVIRGSLTAATGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----EYNATKMPNSIQIDTVFGDFPGALWNPNTPLSEDIYINWVPAKAR-----	216
HaAce1	-----ERFYSNHERADEEPMLEDEGKPEDEPLVTRKGRVIRGSLTAATGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----EILATTLPHISIQIDTVFGDFPGATMWNPNVPLNEDIYINWVPAKAR-----	226
SaAce1	-----DNDFESSGETYSAYKSDPLVHTNKGRIRGITAASSGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----ILNCTPPNNTIQIDTVFGDFPGATMWNPNVPLNEDIYINWVPAKAR-----	207
SgAce1	-----DTFASSGETYSAYKSDPLVHTNKGRIRGITAASSGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----ILNCTPPNNTIQIDTVFGDFPGATMWNPNVPLNEDIYINWVPAKAR-----	207
AgAce1	-----LEHVHSGATPRRRLTRRESNDSANDPLVNTDKGRIRGITAASSGKVDWLGIPYAKQKPLGRLFRHPRPAEKWEG-----VLTNTTPNNTIQIDTVFGDFPGATMWNPNVPLNEDIYINWVPAKAR-----	266
TcaAce	-----ELLVNTKSGVMGTRVPLVSS-HISAFGLPFAEPPVGMFRFRPEPKPWSG-----VWNASTYNNIQIDTVFGDFPGSGEEMWPNRSEDIYINWVPAKAR-----	126
DmAce	-----LRHGRGANGGHEHPNGKADTDHLLHNGNQNTNGLPILWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	279
AgAce2	-----LRHGRGLNFG-SNDYFQDDDFQR-----QHSSKGLLMLWYGGGFMGTSATLDLYNAEILAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	266
TcAce2	-----TRHHEKGLPQ-----DRPKVPLVWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	262
LdAce2	-----TRHHEKGLPQ-----DRPKVPLVWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	256
HaAce2	-----VRHHDQKPLA-----ERPKVPLVWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	278
BgAce2	-----LRHSG-----ERPKVPLVWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	269
SaAce2	-----TRHHSNNAHH-----AKIPVPLVWYGGGFMGTSATLDLYNADIMAAVGNVIVASQYRVGAFGLFLHAPMPSFAEAEAPNGVLDQALAIKWLKDNHAFGGNPEWMTLFGESAGS-----	279
TcAce1	-----PT-----SAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TPDVPNGAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	270
BgAce1	-----PK-----NAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TPDVPNGAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	313
HaAce1	-----PK-----NAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TPDVPNGAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	317
SaAce1	-----PQ-----NAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TEDVPGNAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	304
SgAce1	-----PQ-----NAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TEDVPGNAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	304
AgAce1	-----PK-----NAAVMWVFGGGFYSGATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----TPEAPNGAGLFDQMLAQWRVNDIAAFGGNPNVTLFGESAGA-----	363
TcaAce	-----PK-----STTVMWVYGGGFMGTSATLDYDHPKTLVSEENVIVSMQYKVASLGLFLYFD-----SQEAPNGVLDQMLAQWRVNDIAAFGGNPNVTLFGESAGS-----	224
DmAce	-----SSVNAQLMSPVTRGLVYKRGMMQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	413
AgAce2	-----SSVSLHLLSPVTRGLVYKRGMMQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	400
TcAce2	-----GSVSLHLLSPVTKLARGLILQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	396
LdAce2	-----GSISIHLLSPVTKLARGLILQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	390
HaAce2	-----GSISIHLLSPVTKLARGLILQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	412
BgAce2	-----GSISIHLLSPVTKLARGLILQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	413
SaAce2	-----GSVSHLLSPVTRGMVRRGIIQSGTMNAPWSHMTSEKAVEIGKALINDCN-----NASMLKTNPAHMS-----RISVDAKTSVQQNNSYS-----GILSPSAPTIDGAFPLADPMTLMKATDLKDYDILMGNVDEGTYFLLYD-----	403
TcAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----ERHLSAVDIDKDKKDDIDLNVNNGWTL-----GICESPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	401
BgAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----SRSDIRAVDIDKDKKDDIDLNVNNGWTL-----GICESPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	444
HaAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----SRDMDGPMIELRKKSPDELNVNNGWTL-----GICESPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	448
SaAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----DRNATHKTVLRLKVNSSAMVEKEDWH-----AICFPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	435
SgAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----DRNTIHKTVLRLKVNSSAMVEKEDWH-----AICFPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	435
AgAce1	-----VSVSLHLLSPLSRNLSQAIMSGSSTAPWALISREESILRGLRLAEAVGPH-----EPLSKSDAVELRGGKDPVHVNNGWTL-----GICESPFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	494
TcaAce	-----ASVGMHLLSPGSRDLFRALILQSGSNPCWASVSAEGRRAVELGRNLSL-----NSD-EELIHLRKKPQELLDVENVLPDIFRFRFPVVDGAFIDESPTRALANKFKTKTILMGSNTEEGYFIIYY-----	356
DmAce	-----FIDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----GNPGYQNGQIGRAVDHFFPTNEAYQALAEARGASVHYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	546
AgAce2	-----FIDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----GNPGYQNGQIGRAVDHFFPTNEAYQALAEARGASVHYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	534
TcAce2	-----FIDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----GNPGYQNGQIGRAVDHFFPTNEAYQALAEARGASVHYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	530
LdAce2	-----FIDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----GNPGYQNGQIGRAVDHFFPTNEAYQALAEARGASVHYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	524
HaAce2	-----FIDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----GNPGYQNGQIGRAVDHFFPTNEAYQALAEARGASVHYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	546
BgAce2	-----FMDYFDKDDATALPRDKYLEIMNIFGKATQAREAIIFQYTSWE-----LADGYNQKQIMADVVDYFFPTNFAEQFAEHTKYVYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	537
SaAce2	-----FVDFDRTSATLPEKFRVIVNIFKDRQLERDAIYQYSGWEKKEVDKYSNQGKSDVVADYFFPTNFAEQFAEHTKYVYHYFTHRTSTSLWGEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	549
TcAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	535
BgAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	578
HaAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	582
SaAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	569
SgAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	569
AgAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	628
TcaAce	-----APG-PSKDSSEKISREDFMSGKLVSHANDLGLDAVTLQYTDWMD-----DNNGIKNRGLDDIVGDHNVPLMHFVNYKFKPNGTYLFFPHRASNLVPEWGMVGHGDEIYFFGQPLNNSLQYRVPVRELGRK-----	489
DmAce	-----MLSAVIEFAKTGNPAQD-----GEEWPNFSKEDPVYVIFSTDD-----KIEKLARGPLAARFNFNDLPLKVR-----SWAG-----TCDGSGSASIPRLQLLIGLAALYICAAALRKRKF-----	649
AgAce2	-----MVLVSSEFAKTGNPALE-----GEHWPLYTRENPIYIFNAEGEDDLRGEYGRGPMATSFNFNDLPLKLR-----AWSVLPKDCPKLDDHITSIASTARAPTVALLIALSLAVARLVAA-----	645
TcAce2	-----IMQAFARFAATGKPVTD-----DYNWPLYTKDQYQYIFNAD-----KNGIG-----KGRPRATAFNFDLPLKLR-----DNGK-----FN-----RCSLHH-----	604
LdAce2	-----IMQAFARFAATGKPVTD-----DYNWPLYTKDQYQYIFNAD-----KNGIG-----KGRPRATAFNFDLPLKLR-----DNGSEAPCVNTYLSKTRSSSLELLPPSTSLVLIWIMTLNLAAL-----	629
HaAce2	-----IMQSFTRFALTKGPKHP-----DEKWPYLSRTPHYTYTADG-----TSGPAGPRGRPRASAFNFDLPLKLR-----ELEHMPDCGAVT-----GPYSSAVGATLPIVLLTTLATVALL-----	647
BgAce2	-----IMEIYSRFALTKGPKIAN-----EADWYTYTKDKPKYIYLNSD-----TYGAT-----KGRPTLN-----FNFDLPLKLANHP-----DPDYCASGVAETANLGNVNSATSCLLGPKHLLYFVTLVARLLLSL-----	648
SaAce2	-----IMEAFTRFSLTKGTPVSD-----DIDWPLYNESNPYIYHWNAA-----ELHGYGCPRAAEFNFDLPLKLANHP-----QEDYDPSMPTNENCTSSATINPRISFTIIFVPLPAHGLF-----	664
TcAce1	-----LTELFRKEENVYVNRQELRAVLELNPYVNAISRAQIVFEYTDWLN-----PDDPKRNDALDKMVDYHFTNVEFAHRYAETGNVYMYFKHRSNPNPWSWTGVMHGEIYNYVFGELNPKAKNYQPEIELSR-----	648
BgAce1	-----MBRYWANFAKTGNPMSSEDTWTATYVPHVHTAYGREYLTLDVN-----STETGRGPRKQLFNFKKYLPLQLI-----AVTSNLNQPP-----ESCPDTSGAEMNKESLILLIYLVKMLIILLWGRGWV-----	692
HaAce1	-----LMRYWANFARSGNPSINPDSGDKIHPVHTASGREYLSLAVN-----SSSIGHGLRVKELFNFKKYLPLQLI-----ATTSKILLELFLKLN-----	664
SaAce1	-----LMRYWTFNAKTGNPSKTEGSSVTPKPVHTAYGKEFLTLDTN-----NTSIGHGLRVKELFNFKKYLPLQLI-----ATTSKILLELFLKLN-----NCTTISGGTKNMIKLSLWTVMTTAVLV-----	676
SgAce1	-----LMRYWTFNAKTGNPSKTEGSSVTPKPVHTAYGKEFLTLDTN-----NTSIGHGLRVKELFNFKKYLPLQLI-----ATTSKILLELFLKLN-----NCTTISGGTKNMIKLSLWTVMTTAVLV-----	676
AgAce1	-----LMRYWTFNAKTGNPNPN-----TASSEFVPHHTAHRHYLEGLN-----TSFVGRGPRKQLFNFKKYLPLQLI-----AATSNLPGAP-----PSEPCSSAFYRPLDILVLSLTLATVIRFLK-----	737
TcaAce	-----LMHYWTFNAKTGNPNP-----HSQEKWPLFTTKQKFDLNTL-----PMKVHQLRVKQLFNFKKYLPLQLI-----NATETIDEAERQWTEFHRRSSYMMHKNQDHYSRHESCAEL-----	596

Figure 2. Alignment of deduced AChE protein sequences encoded by TcaAce (CAA27169, *Torpedo californica* Ace); TcAce1 (HQ260968, *Tribolium castaneum* Ace1, this paper); TcAce2 (HQ260969, *T. castaneum* Ace2, this paper); SaAce1 (AY819704, *Sitobion avenae* Ace1); SaAce2 (AY707319, *S. avenae* Ace2); DmAce (X05893, *Drosophila melanogaster* Ace); AgAce1 (XM_321792, *Anopheles gambiae* Ace1); AgAce2 (BN000067, *A. gambiae* Ace2); BgAce1 (DQ288249, *Blattella germanica* Ace1); BgAce2 (DQ288847, *B. germanica* Ace2); HaAce1 (DQ001323, *Helicoverpa assulta* Ace1); HaAce2 (AY817736, *H. assulta* Ace2); SgAce1 (AF321574, *Schizaphis graminum* Ace1) and LdAce2 (L41180, *Leptinotarsa decemlineata* Ace2). Numbering of the amino acid sequences was from the N-terminus of mature proteins. Identical amino acids were indicated by asterisks and conservative substitutions by dots. The catalytic triad residues were marked with arrowhead. The number 1, 2, 3 on the boxed amino acids indicated the residues forming intramolecular disulfide bonds. The positions of aromatic residues lining the active site gorge in *T. californica* AChE were marked with block arrows. The cholinesterase signature sequence was underlined.

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during the last 1-ns period of all 21 simulations or in an average of each cluster of the trajectories generated by a cluster analysis. As to the loop conformation of residues 145–162, the cluster analysis showed that 24% of the trajectories had residues 145–162 folded in contact with surface residues such as R21, D148, D501, and D203.

A third set of 22 10-ns simulations of TcAce2 with residues 145–162 adopting the folded conformation was then carried out. The initial conformation of the third set of simulations was obtained from averaging all the trajectories of TcAce2 with the folded loop conformation of residues 145–162 followed by manual adjustment of the side-chain torsions to restore the hydrogen bond network of the catalytic triad. A cluster analysis of all the trajectories saved at 1.0-ps intervals during the last 1.0-ns period of all 22 simulations showed that residues 145–162 remain the folded conformation. However, the average of all the trajectories had a distorted catalytic triad. Visual inspection of all 22 simulations found that conformations of the first of the 22 simulations have a catalytic triad engaging in a hydrogen-bond network. The final model of TcAce2 was then obtained from averaging all trajectories saved at 1.0-ps intervals during the last 1.0 ns period of the first simulation.

Although the homology model of TcAce2 was based on the DmAChE crystal structure, the active site of the simulation-refined TcAce2 model (Fig. 7) is very different from that of the DmAChE crystal structure. It is also different from those of human AChE and insect AP-AChEs (*i.e.*, AChE1s). In the refined model of TcAce2 (Fig. 7), Y114, Y345, Y395, and W342 form an aromatic cluster that completely block the entrance of the active site; E388, H502, and S259 form a catalytic triad; ACh has a cation- π interaction with W126, but it does not adopt the fully extended conformation, nor is its carbonyl oxygen atom located in the oxyanion hole. In contrast to the TcAce1 model with residues 146–154 and 493–509 that partially shield ACh from interacting with solvent, the TcAce2 model has its corresponding residues (120–128 and 488–504) adopt conformations that leave ACh to be exposed to solvent (Fig. 8).

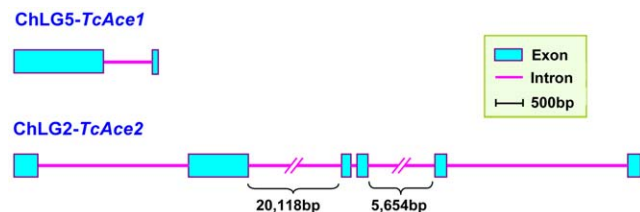


Figure 3. Schematic diagram of the organization of two Ace genes from *Tribolium castaneum*. The full lengths of the two genomic sequences were 2,986 bp for TcAce1 and 32,243 bp for TcAce2. Genome structure showed that two different Ace genes in *T. castaneum* located on different chromosomes. TcAce1 located on chromosome 5 and TcAce2 on chromosome 2. TcAce1 has two exons and one intron, whereas TcAce2 has six exons and five introns.

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TcAce gene expression profiles

The transcript levels of TcAce1 and TcAce2 were evaluated by reverse transcription PCR (RT-PCR) and quantitative real-time PCR (qPCR) in tissues of *T. castaneum* at different developmental stages (Fig. 9). Both TcAce genes were transcribed in all the stages examined, including 1-day (d) and 3-d eggs; 5-d and 20-d larvae; 1-d, 3-d and 6-d pupae; and 2-d and 14-d adults. The lowest expression levels of these genes were found in eggs, particularly for TcAce2 whose expression level was undetectable by RT-PCR in 1-d eggs (Fig. 9A and 9B). The expression patterns of TcAce1 and TcAce2 were very similar. In addition, the TcAce1 and TcAce2 genes also exhibited similar tissue-specific expression patterns (Fig. 9C and 9D). As expected, these genes were predominately expressed in the brain, although their expressions were also detected in the gut and carcass after the brain and ventral nerve cord were removed.

Discussion

Since the first insect AChE orthologous gene (*i.e.*, the one later named Ace2 or AO-Ace) and the first insect AChE paralogous gene (*i.e.*, the one later named Ace1 or AP-Ace) were reported in *D. melanogaster* in 1986 [9] and in *S. graminum* in 2002 [10], respectively, cDNAs encoding both Ace1 and Ace2 have been sequenced from each of at least 27 insect species. However, the gene models and the genomic organizations of Ace1 and Ace2 have not been well established in insects. In this study, we confirmed the two Ace gene (TcAce1 and TcAce2) models in *T. castaneum* by sequencing the coding regions of their cDNAs followed by comparative analyses of their cDNA and genomic sequences. The TcAce1 and TcAce2 genes are significantly different not only in the length of their genomic DNA (TcAce1 with 2,986 bp and TcAce2 with 32,243 bp) but also in the intron/exon organizations. Specifically, TcAce1 possesses only one intron whereas TcAce2 has five introns. Furthermore, TcAce1 is located on chromosome 5 (ChLG5), whereas TcAce2 is on chromosome 2 (ChLG2; Fig. 3). Apparently, the intron/exon organizations and the chromosomal locations of these genes in *T. castaneum* are different from their counterparts in other insect species [8,14,18,23,45].

Despite the significant differences in genomic structures and the chromosomal locations of TcAce1 and TcAce2, the deduced protein sequences of the two genes exhibit all the common features of an AChE sequence wise. These features include (1) a conserved active-site triad, including S267 in TcAce1 and S259 in TcAce2 (S200 in *Torpedo*), E393 in TcAce1 and E388 in TcAce2 (E327 in *Torpedo*), and H507 in TcAce1 and H502 in TcAce2 (H440 in *Torpedo*); (2) a choline binding site, W152 in TcAce1 and W126 in TcAce2 (W84 in *Torpedo*); (3) three pairs of cysteines putatively forming intramolecular disulfide bonding (C135~C162, C321~C334, and C469~C591 in TcAce1; and C109~C136, C313~C328, and C464~C580 in TcAce2); (4) a cysteine forming intermolecular disulfide bonding (C617 in TcAce1 and C600 in TcAce2); (5) 10 conserved aromatic amino acid residues out of 14

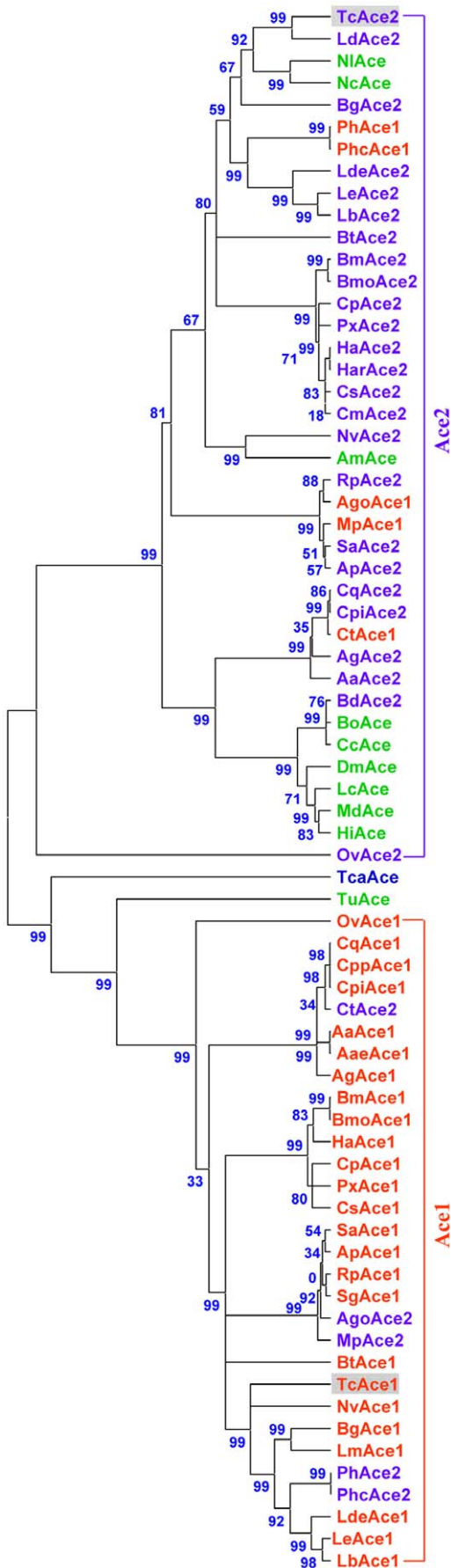


Figure 4. Rooted phylogenetic tree of deduced *Ace* amino acid sequences from the Pacific electric ray (*Torpedo californica*), two-spotted spider mite (*Tetranychus urticae*) and 43 insect species constructed by the neighbor-joining method. The name is made up of a species abbreviation (first letter of the genus followed by the first one or two letters of the specific name). Sequences used: TcaAce (CAA27169, *T. californica* Ace); TcAce1 (HQ260968, *Tribolium castaneum* Ace1, this paper); TcAce2 (HQ260969, *T. castaneum* Ace2, this paper); BmAce1 (EU262633, *Bombyx mandarina* Ace1); BmAce2 (EU262632, *B. mandarina* Ace2); SaAce1 (AY819704, *Sitobion avenae* Ace1); SaAce2 (AY707319, *S. avenae* Ace2); RpAce1 (AY667435, *Rhopalosiphum padi* Ace1); RpAce2 (AY707318, *R. padi* Ace2); DmAce (X05893, *Drosophila melanogaster* Ace); AgAce1 (XM_321792, *Anopheles gambiae* Ace1); AgAce2 (BN000067, *A. gambiae* Ace2); LdeAce1 (FJ647186, *Liposcelis decolor* Ace1); LdeAce2 (FJ647187, *L. decolor* Ace2); OvAce1 (FJ228227, *Orchesella villosa* Ace1); OvAce2 (FJ228228, *O. villosa* Ace2); LeAce1 (EU854149, *Liposcelis entomophila* Ace1); LeAce2 (EU854150, *L. entomophila* Ace2); BgAce1 (DQ288249, *Blattella germanica* Ace1); BgAce2 (DQ288847, *B. germanica* Ace2); BtAce1 (EF675188, *Bemisia tabaci* Ace1); BtAce2 (EF675190, *B. tabaci* Ace2); CqAce1 (XM_001847396, *Culex quinquefasciatus* Ace1); CqAce2 (XM_001842175, *C. quinquefasciatus* Ace2); BmoAce1 (NP_001037380 *Bombyx mori* Ace1); BmoAce2 (NP_001108113 *B. mori* Ace2); ApAce1 (XM_001948618, *Acyrtosiphon pisum* Ace1); ApAce2 (XM_001948953, *A. pisum* Ace2); NvAce1 (XM_001600408, *Nasonia vitripennis* Ace1); NvAce2 (XM_001605518, *N. vitripennis* Ace2); PhAce1 (AB266605, *Pediculus humanus corporis* Ace1); PhAce2 (AB266606, *P. humanus corporis* Ace2); CpAce1 (DQ267977, *Cydia pomonella* Ace1); CpAce2 (DQ267976, *C. pomonella* Ace2); HaAce1 (DQ001323, *Helicoverpa assulta* Ace1); HaAce2 (AY817736, *H. assulta* Ace2); AaAce1 (AB218421, *Aedes albopictus* Ace1); AaAce2 (AB218420, *A. albopictus* Ace2); AgoAce1 (AF502081, *Aphis gossypii* Ace1); AgoAce2 (AF502082, *A. gossypii* Ace2); CtAce1 (AB122151, *Culex tritaeniorhynchus* Ace1); CtAce2 (AB122152, *C. tritaeniorhynchus* Ace2); SgAce1 (AF321574, *Schizaphis graminum* Ace1); MpAce1 (AF287291, *Myzus persicae* Ace1); MpAce2 (AY147797, *M. persicae* Ace2); CpiAce1 (AJ489456, *Culex pipiens* Ace1); CpiAce2 (AM159193, *C. pipiens* Ace2); MdAce (AY134873, *Musca domestica* Ace); PxAce1 (AY970293, *Plutella xylostella* Ace1); PxAce2 (AY061975, *P. xylostella* Ace2); CsAce1 (EF453724, *Chilo suppressalis* Ace1); CsAce2 (EF470245, *C. suppressalis* Ace2); CmAce2 (FN538987, *Cnaphalocrocis medinalis* Ace2); PhcAce1 (AB266614, *Pediculus humanus capitis* Ace1); PhcAce2 (AB266615, *P. humanus capitis* Ace2); TuAce (AY188448, *T. urticae* Ace); AaeAce1 (EF209048, *Aedes aegypti* Ace1); LmAce1 (EU231603, *Locusta migratoria manilensis* Ace1); LbAce1 (FJ647185, *Liposcelis bostrychophila* Ace1); LbAce2 (EF362950, *L. bostrychophila* Ace2); NIAce (FM866396, *Nilaparvata lugens* Ace); CcAce (EU130781, *Ceratitis capitata* Ace); AmAce (AB181702, *Apis mellifera* Ace); BdAce2 (AY155500, *Bactrocera dorsalis* Ace2); HiAce (AY466160, *Haematobia irritans* Ace); CppAce1 (AY762905, *Culex pipiens pallens* Ace1); HarAce2 (AF369793, *Helicoverpa armigera* Ace2); BoAce (AF452052, *Bactrocera oleae* Ace); LcAce (U88631, *Lucilia cuprina* Ace); NcAce (AF145235, *Nephotettix cincticeps* Ace); LdAce2 (L41180, *Leptinotarsa decemlineata* Ace2).

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aromatic residues lining the catalytic gorge of AChE; and (6) the conserved sequence FGESAG, flanking S267 in TcAce1 and S259 in TcAce2 (Fig. 2).

Ethanolmine and glucosamine residues are characteristic of a C-terminal glycolipid anchor in most G2 AChE [46,47]. Our analysis by using PredGPI predictor suggests that DmAChE and TcAce1 contain a GPI-anchor at the C-terminal. Although TcAce2 has higher sequence identity level with DmAChE than TcAce1, TcAce2 does not appear to contain a GPI-anchor at the C-terminal. Because the C-terminal Cys residue in AChE is reportedly for the intermolecular disulfide linkage [48], C617 in TcAce1 and C600 in TcAce2 are likely involved in the intermolecular disulfide linkage, although the corresponding Cys residue is missing in DmAChE [46]. We also analyzed the hydrophilic and hydrophobic of non-homologous amino acid residues of TcAce1 and TcAce2 and compared homologous

Table 2. Percent identities of amino acid residues among the AChEs of *Tribolium castaneum*, *Torpedo californica* and other seven insect species.

Name	Tca	TcAce1	SaAce1	AgAce1	BgAce1	SgAce1	HaAce1	DmAce	TcAce2	SaAce2	AgAce2	BgAce2	LdAce2	HaAce2
Tca	—	39	38	42	40	38	39	35	38	37	37	38	37	37
TcAce1		—	57	61	70	57	64	31	36	31	34	36	34	34
SaAce1			—	51	59	96	55	32	36	31	33	34	33	34
AgAce1				—	57	51	59	34	37	33	35	39	35	36
BgAce1					—	59	62	33	36	33	33	35	36	34
SgAce1						—	55	32	36	30	33	34	34	33
HaAce1							—	31	35	30	32	35	34	32
DmAce								—	55	48	62	51	53	51
TcAce2									—	57	60	68	84	69
SaAce2										—	50	53	55	52
AgAce2											—	55	57	56
BgAce2												—	63	60
LdAce2													—	66
HaAce2														—

NOTE: TcaAce, *Torpedo californica*; TcAce, *Tribolium castaneum*; SaAce, *Sitobion avenae*; DmAce, *Drosophila melanogaster*; AgAce, *Anopheles gambiae*; BgAce, *Blattella germanica*; HaAce, *Helicoverpa assulta*; SgAce, *Schizaphis graminum*; LdAce, *Leptinotarsa decemlineata*.
doi:10.1371/journal.pone.0032288.t002

sequences of TcAce1 and TcAce2 with mAChE and DmAChE at the C-terminal for possible formation of the four-helix bundle. Our analysis suggests that C-terminal sequences of both TcAce1 and TcAce2 may form the dimeric four-helix bundle.

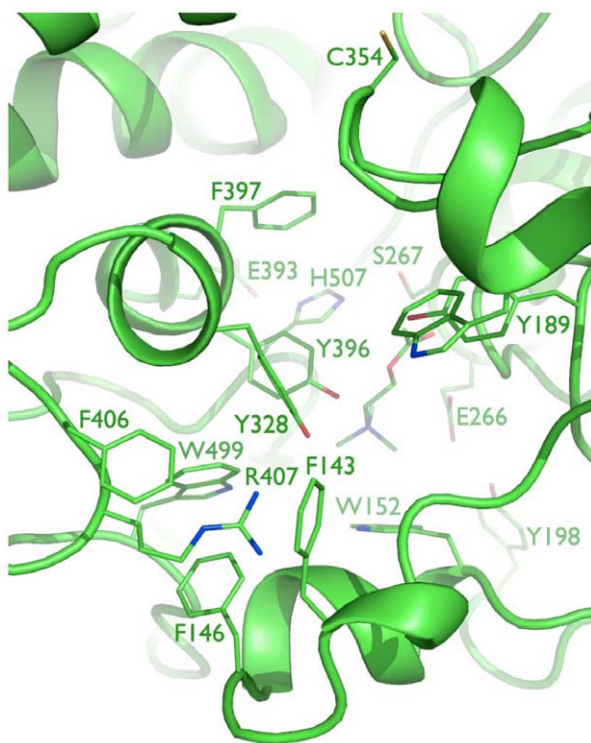


Figure 5. Close-up view of the active site of TcAce1 with a perspective from the free cysteine at the opening of the active-site gorge down to ACh and the catalytic triad at the bottom of the gorge.

doi:10.1371/journal.pone.0032288.g005

According to comparisons of TcAce1 and TcAce2 with Ace proteins from other insect species, TcAce1 showed high sequence identities to the AChE1 (AP-AChE) proteins from *Sitobion avenae* (57%), *A. gambiae* (61%), *Blattella germanica* (70%), *S. graminum* (57%), and *Helicoverpa assulta* (64%; Table 2). Similarly, TcAce2 also showed high amino acid identities to the AChE2 (AO-AChE) proteins of *S. avenae* (57%), *D. melanogaster* (55%), *A. gambiae* (60%), *B. germanica* (68%), *Leptinotarsa decemlineata* (84%), and *H. assulta* (69%). Based on our analysis of deduced amino acid sequences from the two genes among respective insect species, the sequence identity levels within the paralogous (*Ace1*) or orthologous (*Ace2*) genes range from 48 to 96%, whereas the sequence identity levels between the *Ace1* and *Ace2* genes of the same insect species range only from 31 to 36% in all insect species examined (Table 2). These results support the hypothesis that the two *Ace* genes were originated from an old duplication before the diversification of insect species [32].

Furthermore, a phylogenetic tree, which was generated from the highly conserved regions of all insect and *T. urticae* AChE

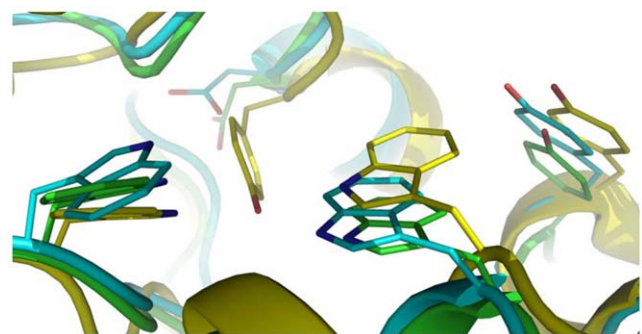


Figure 6. Comparison of the bottom of the active-site gorge in TcAce1 to those in human AChE and *Anopheles gambiae* AP-AChE. Tyr449 in human AChE (yellow) is mutated to Asp509 in TcAce1 (green) and Asp441 in *A. gambiae* AChE (cyan).

doi:10.1371/journal.pone.0032288.g006

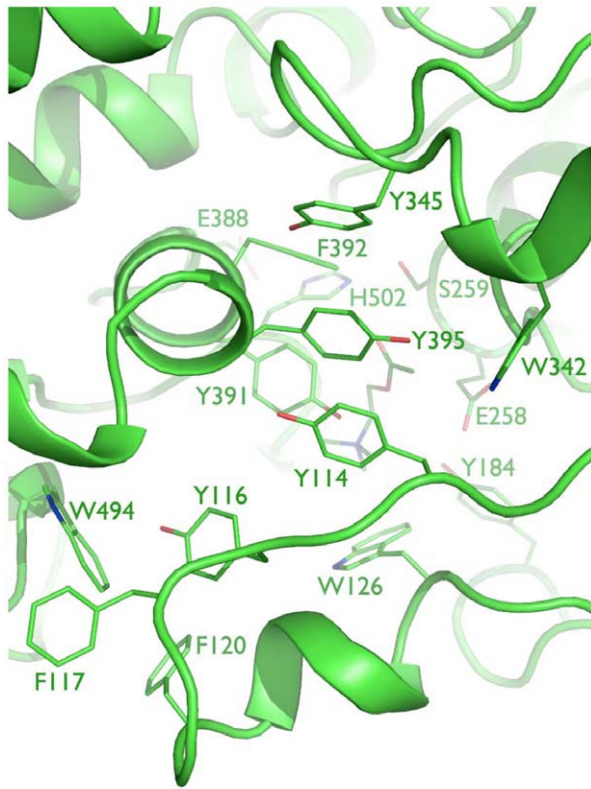


Figure 7. Close-up view of the active site of TcAce2 with a perspective looking down to acetylcholine and the catalytic triad at the bottom of the gorge.
doi:10.1371/journal.pone.0032288.g007

amino acid sequences available in GenBank and the corresponding one in *T. californica* using the neighbor-joining method, revealed two insect AChE clusters. *TcAce1* was grouped into the insect *Ace1* cluster and *TcAce2* into the insect *Ace2* cluster. The significant divergence between *TcAce1* and *TcAce2* suggests that these genes were evolved from their corresponding *Ace* gene lineages in insect species [49]. These results also suggest, for the first time, that the divergence of *Ace1* and *Ace2* might occur prior to insect speciation and that the *Ace1* gene might be lost in *D. melanogaster* and other species in Cyclorapha suborder of Diptera during the evolutionary process [14,25,32]. Thus, it is likely that both the *Ace* genes in insects may have different functions because these genes have evolved during the evolutionary histories of these insect species.

Using a reported simulation-refined model of *A. gambiae* AP-AChE [33] and a crystal structure of DmAChE [41] with high sequence identities to TcAce1 (73%) and TcAce2 (60%), respectively, and the same multiple molecular dynamics simulation method to model and refine TcAce1 and TcAce2, we obtained models of TcAce1 and TcAce2 both which are in complex with acetylcholine. The TcAce1 model has an active site that is almost identical to those of *A. gambiae* and *S. graminum* AP-AChEs, and it has C354 at the opening of the active-site gorge just like the insect-specific C286 of *A. gambiae* and C289 of *S. graminum* that are susceptible to sulfhydryl agents [33,34]. The TcAce2 model has an active site with an entrance comprised of G122, W126, W494, M501, and D504. This entrance corresponds to the small opening at the bottom of the active-site gorge

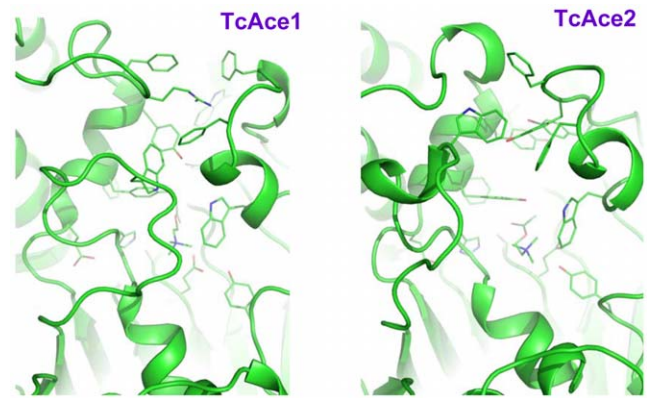


Figure 8. Comparison of loop conformations of residues 146-154 and 493-509 in TcAce1 with the corresponding ones of residues 120-128 and 488-504 in TcAce2.
doi:10.1371/journal.pone.0032288.g008

of DmAChE [44] or *T. californica* AChE when Y442 moves away from W84 [43]. In the TcAce2 model, the region that corresponds to the entrance of TcAce1 is completely blocked by Y114, Y345, Y395, and W342. Of the four aromatic residues, Y395 and W342 correspond to Y334 and W279 of *T. californica* AChE, respectively, and belong to the 14 conserved aromatic residues that line the active-site gorge of *T. californica* AChE [50]. In other words, the entrance of the active-site gorge of TcAce2 appears to be reversed relative to that of TcAce1. Unlike ACh in the TcAce1 model, ACh does not adopt the fully extended conformation and its carbonyl oxygen atom is not placed in the oxyanion hole in the TcAce2 model.

In addition, R576 in the TcAce2 model is close to E388 (the separation between the side-chain N atom of R576 and the side-chain O atom of E388 is 3.9 Å), a composite residue of the catalytic triad, in contrast to the corresponding arginine residue that is away from the catalytic glutamate residue in TcAce1, *A. gambiae* AP-AChE and human AChE (Fig. 10). Analysis of all the trajectories of the third set of simulations of TcAce2 showed that 95% of the trajectories has a hydrogen bond between R576 and E388, accompanied by hydrogen bonds between H502 and E505 and between S259 and E258, leading to disruption of the catalytic triad. These computational observations suggest that TcAce1 is a robust ACh hydrolase and susceptible to sulfhydryl agents and that TcAce2 is not a catalytically efficient ACh hydrolase, although further study is needed to comprehensively elucidate physiological functions of *Ace1* and *Ace2* genes. In view of these computational results, it is logical to investigate whether TcAce2 functions more as a cholinesterase-like adhesion molecule (CLAM) [51,52] than TcAce1. However, our sequence analysis using ClustalW 2.0.12 shows that TcAce1 has a slightly higher sequence homology to *D. melanogaster* gliotactin, which is one of the three *D. melanogaster* CLAMs, than TcAce2 (Table 3); both TcAce1 and TcAce2 have dipole moments that are comparable to those of other AChEs (Table 4). The orientations of the dipole moments of TcAce1 and TcAce2 are almost the same. The two dipole moments are approximately along the beta strand that corresponds to Strand 5 of the *T. californica* AChE crystal structure [53] and nearly identical to those of other AChEs but orthogonal to that of *Galactomyces geotrichum* lipase [52,54]. These sequence and dipole moment analyses do not support the hypothesis that TcAce2 functions as a CLAM.

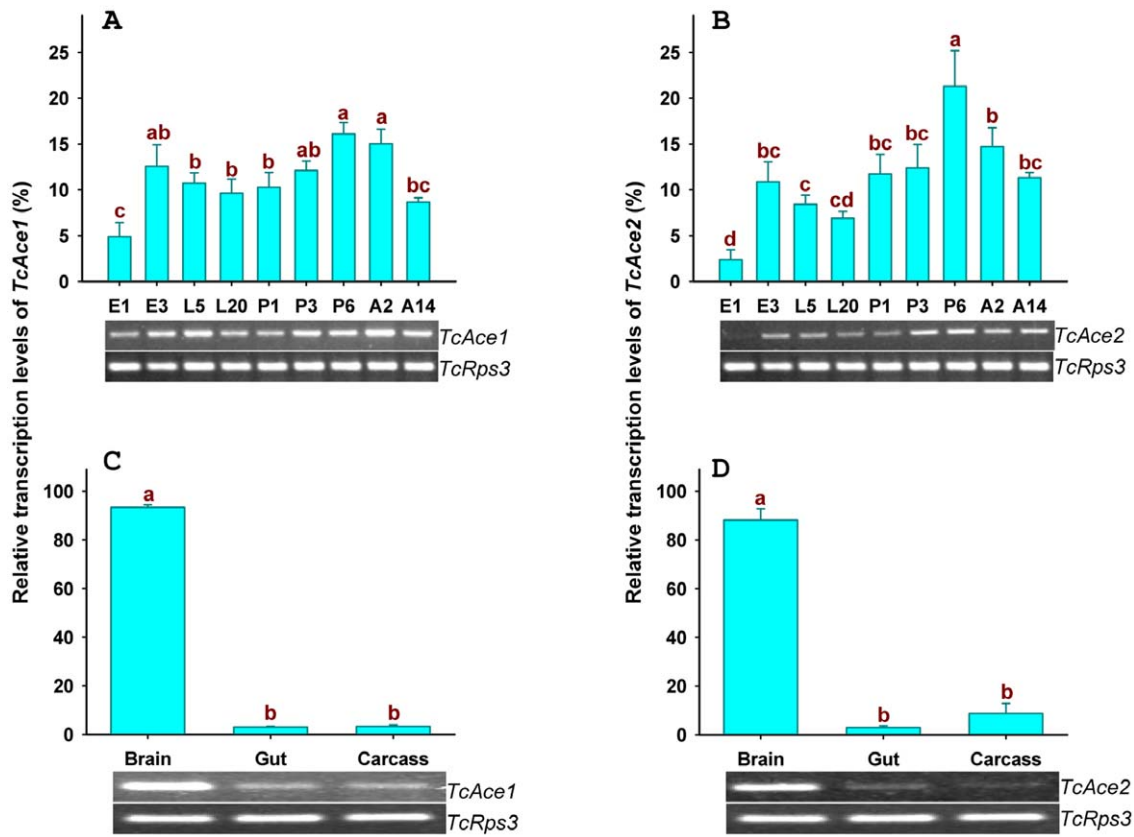


Figure 9. The two *TcAce* genes mRNA expression levels of developmental stages and different tissues were determined by RT-PCR as shown by gel pictures at the bottom of each panel and quantitative real-time PCR as shown by histograms. Real-time PCR data were normalized to *TcRps3* gene expression. E1, 1-d eggs; E3, 3-d eggs; L5, 5-d larvae; L20, 20-day larvae; P1, 1-d pupae; P3, 3-d pupae; P6, 6-d pupae; A2, 2-d adults; A14, 14-d adults. Gut (including midgut and hindgut), Carcass (not including head, gut and nerve system). Standard error bars were based on three replicates. One-Way ANOVA-Fisher's LSD was used in statistical analysis of quantitative real-time PCR data. doi:10.1371/journal.pone.0032288.g009

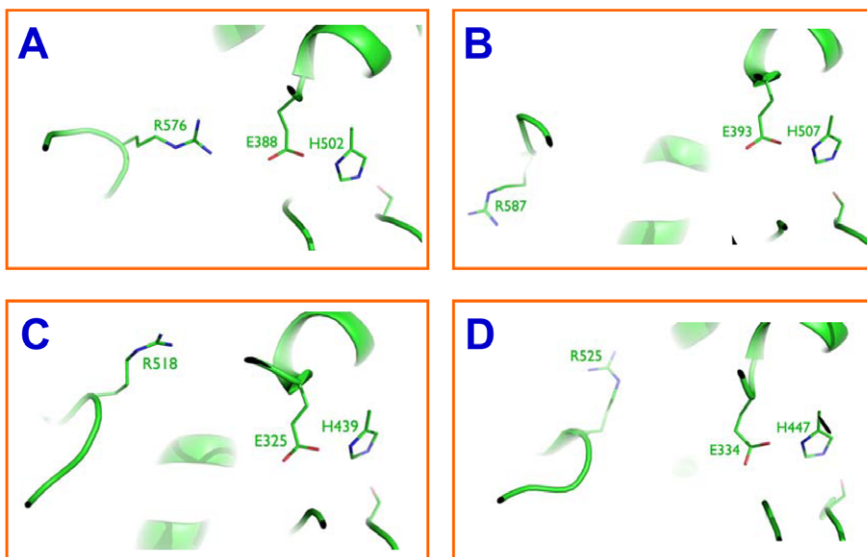


Figure 10. Separation of Arg from the catalytic triad in TcAce2 (A), TcAce1 (B), *Anopheles gambiae* AP-AChE (C), and human AChE (D). doi:10.1371/journal.pone.0032288.g010

Table 3. Percentages of amino acid sequence identity of TcAce1 and TcAce2 to cholinesterase-like lipase and adhesion proteins.

ACHe	GcNeutralLipase	DmNeurologin	DmNeurotactin	DmGliotactin
<i>Torpedo californica</i> AChE	23	19	17	25
TcAChE1	19	21	17	27
TcAChE2	20	20	18	25

The NCBI session numbers of GcNeutralLipase, DmNeurologin, DmNeurotactin, and DmGliotactin are P79066, AAF52450, CAA37831, and AAC41579, respectively.
doi:10.1371/journal.pone.0032288.t003

Materials and Methods

Insect culture

The Georgia-1 (GA-1) strain of *T. castaneum* was reared on whole-wheat flour containing 5% (by weight) of brewers' yeast at 30°C and 65% relative humidity under standard conditions in the laboratory of Kansas State University (Manhattan, Kansas, United States of America) based on the method of Halisack and Beeman [55].

Total RNA isolation and reverse transcription

Total RNA was isolated from *T. castaneum* samples using TRIzol reagent following the recommended procedure by Invitrogen (Carlsbad, California, United States of America). The RNA was treated with DNase I (Fermentas, Glen Burnie, Maryland, United States of America) according to the manufacturer's instruction and the first-strand cDNA template was synthesized from 3.0 µg of total RNA by using First Strand cDNA Synthesis Kit (Fermentas) with oligo (dT)₁₈ as the primer.

Subcloning and sequencing of cDNA

To obtain the cDNAs corresponding to the entire protein coding regions of *TcAce1* and *TcAce2*, we designed specific primers based on *TcAce* gene predictions and their genomic organization (Table 1). The PCR products of each reaction were subjected to electrophoresis on 1% agarose gel containing ethidium bromide. The PCR bands were excised and purified using QIAEX II Agarose Gel Extraction Kit (Qiagen, Valencia, California, United States of America). The purified fragment was subcloned into a pGEM-T Easy Vector (Invitrogen) according to the manufacturer's instruction. The ligation DNA mixtures were used to transform bacterial cells by using Z-Competent *E. coli* Transfor-

mation Kit and Buffer Set™ (Zymo Research Corporation, Irvine, California, United States of America). Plasmids were isolated from the bacterial cells and used for DNA sequencing (KSU DNA Sequencing and Genotyping Facility, Manhattan, Kansas, United States of America). Signal P software was used to predict signal peptide [56].

Phylogenetic analysis of AChEs

ClustalW software (www.ebi.ac.uk/clustalw/) [57], was used to perform multiple sequence alignments prior to phylogenetic analysis. Phylogenetic analysis was done using MEGA 4.0 [58] for construction a neighbor-joining tree to examine the evolutionary relationships among *T. californica*, *T. urticae* and 43 insect species. To evaluate the branch strength of the phylogenetic tree, a bootstrap analysis of 1000 replications was performed.

AChE model prediction

Model preparation. The starting conformations of TcAce1 and TcAce2 used in the multiple molecular dynamics simulations were generated by the SWISSMODEL homology program using a computer model (Protein Data Bank ID: 2AZG [33]) and a crystal structure (Protein Data Bank ID: 1DX4 [41]) as templates, respectively. ACh was manually docked into the active site of TcAce1 or TcAce2 according to the bound ACh conformation in the crystal structure of *T. californica* AChE (Protein Data Bank ID: 2ACE [59]). All His, Glu, Asp, Arg, and Lys residues of the ACh-bound TcAce1 or TcAce2 were treated as HIP, GLU, ASP, ARG, and LYS, respectively. The topology and coordinate files were generated by the PREP, LINK, EDIT, and PARM modules of the AMBER 5.0 program [60]. The complex was refined by energy minimization using the SANDER module of the AMBER 5.0 program with a dielectric constant of 1.0 and 500 cycles of

Table 4. The electrostatic characteristics of TcAce1, TcAce2, and other proteins.

PDB ID (Species)	Total atoms	Total residues	Net charge of N terminal deletion (e)	Net charge of C terminal deletion (e)	Net charge of deletion between N and C termini (e)	Net charge of the structure or model (e)	Dipole moment (Debye)	Dipole moment per atom (Debye)
TcAce1 (<i>Tribolium castaneum</i> , AP)	4201	529	-5	0	0	-3	1490	0.35
TcAce2 (<i>Tribolium castaneum</i> , AO)	4353	545	-1	+1	0	-19	1163	0.27
1QO9 (<i>Drosophila melanogaster</i> , AO)	4273	540	0	-2	+1	-18	669	0.16
2AZG (<i>Anopheles gambiae</i> , AP)	4243	536	0	-1	0	-8	1718	0.40
2HCP (<i>Schizaphis graminum</i> , AP)	4302	540	-7	+2	0	-8	1312	0.30
2ACE (<i>Torpedo californica</i>)	4143	527	-2	-1	-1	-8	1819	0.44
1J06 (<i>Mus musculus</i>)	4177	535	+2	-2	0	-9	867	0.21
2X8B (<i>Homo sapiens</i>)	4179	536	0	-2	0	-10	1384	0.33
1THG (<i>Galactomyces geotrichum</i> , lipase)	4287	543	+1	0	0	-15	814	0.19

doi:10.1371/journal.pone.0032288.t004

steepest-descent minimization followed by 10,000 cycles of conjugate-gradient minimization. The energy-minimized ACh complex with TcAce1 or TcAce2 was solvated with 5,897 or 6,744 TIP3P water molecules [61], leading to a system of 20,703 or 23,244 atoms, respectively. The water molecules were obtained from solvating the complex using a pre-equilibrated box of 216,000 TIP3P molecules, whose hydrogen atom charge was set to 0.4170, where any water molecule was removed if it had an oxygen atom closer than 2.2 Å to any solute atom or a hydrogen atom closer than 2.0 Å to any solute atom, or if it was located further than 10.0 Å along the x-, y-, or z-axis from any solute atom.

Multiple molecular dynamics simulations. The solvated protein complex was energy-minimized for 100 cycles of steepest-descent minimization followed by 100 cycles of conjugate-gradient minimization to remove close van der Waals contacts in the system, then heated from 0 to 300 K at a rate of 10 K/ps under constant temperature and volume, and finally simulated independently with a unique seed number for initial velocities at 300 K under constant temperature and pressure using the PMEMD module of the AMBER 8.0 program [62] with the AMBER force field (ff99SB) [63,64]. All simulations used (1) a dielectric constant of 1.0, (2) the Berendsen coupling algorithm [65], (3) a periodic boundary condition at a constant temperature of 300 K and a constant pressure of 1 atm with isotropic molecule-based scaling, (4) the Particle Mesh Ewald method to calculate long-range electrostatic interactions [66], (5) a time step of 1.0 fs, (6) the SHAKE-bond-length constraints applied to all the bonds involving the H atom, (7) saving the image closest to the middle of the “primary box” to the restart and trajectory files, (8) formatted restart file, and (9) default values of all other inputs of the PMEMD module. All simulations were performed on a cluster of Apple Mac Pros with 80 Intel Xeon cores (3.0 GHz) and a cluster of Apple Xserves with 590 G5 processors (2.2/2.3 GHz).

Simulation analysis. Average structures were obtained by using the CARNAL module of AMBER 5.0. Cluster analyses were performed by using the PTRAJ module [67] of AMBER 10.

Dipole moment calculations. All dipole moment calculations were performed using the Protein Dipole Moments Server (<http://bioinfo.weizmann.ac.il/dipol/indexj.html>) [68]. Protein structures with minimal deletions were used and ligands and structural water molecules were removed before the dipole moment calculations.

Analysis of expression of *TcAce1* and *TcAce2* by RT-PCR and qPCR

The expression patterns of both *TcAce1* and *TcAce2* genes were analyzed at various developmental stages including embryos (1 day and 3 days eggs), early larvae (5 days larvae), late larvae (20 days larvae), early pupae (1 day pupae), middle pupae (3 days larvae), late pupae (6 days pupae), early adults (2 days adults) and two weeks old adults. To analyze tissue specific expression, we

collected samples from the following dissected late pupa tissues (pooled from thirty late pupae): brain, gut (midgut and hind gut) and carcass (the whole body excluding brain, ganglia and gut). For all the samples, 3.0 µg of total RNA were treated with DNase I (Fermentas) to remove any genomic DNA contaminations, and then used as templates for the first strand cDNA synthesis. The cDNAs prepared from total RNA were used as templates for amplification and detection of specific *TcAce* sequences. The gene-specific primers were designed by using the Beacon Designer 2.0 software (Premier Biosoft International, Palo Alto, California, United States of America) and are shown in Table 1. For reverse transcription PCR (RT-PCR), cDNA fragments of each *TcAce* were amplified using the PCR conditions as follows: 94°C for 1.5 min followed by 30 cycles (26 cycles for *TcRps3* gene) of 94°C 30 s, 55°C 30 s and 72°C 45 s. A final extension at 72°C for 5 min was added at the end of the PCR. The relative mRNA expression of each *TcAce* was assessed by qRT-PCR using SYBR-Green in the Bio-Rad iCycler iQ™ multi-color real-time PCR detection system (Bio-Rad Laboratories, Hercules, CA, USA) based on the method of Giulietti et al. [69]. All the experiments were performed in triplicate and normalized to the mRNA level of ribosomal protein S3 (*Rps3*) as a reference gene for each sample [70]. The relative mRNA expression levels were calculated according to the $2^{-\Delta\Delta C_t}$ method [71].

Statistical analysis

The data from the qPCR analysis were subjected to ANOVA followed by Fisher’s least significant difference (LSD) multiple comparisons to separate the means among the treatments by using ProStat software (Poly Software International, Pearl River, New York, United States of America).

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Author Contributions

Conceived and designed the experiments: YL YPP XG YP KYZ. Performed the experiments: YL JY XZ YPP KYZ. Analyzed the data: YL YPP YP JY XZ KYZ. Contributed reagents/materials/analysis tools: YP KYZ. Wrote the paper: YL YPP KYZ. Contributed with revisions: YL YPP XG YP JY XZ KYZ.

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