Molecular Cloning of Acetylcholinesterases from the Oriental Tobacco Budworm, *Helicoverpa assulta* (Guenée)

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Acetylcholinesterase (AChE) plays an important role in neurotransmission at insect cholinergic synapses. Since an orthologous AChE was identified from Drosophila, it has been suggested that only a single AChE existed in all insect genomes. However recently paralogous AChE has been isolated from mosquitoes and aphids. We fully cloned and isolated two AChE genes, acel and ace2 from the oriental tobacco budworm, Helicoverpa assulta. acel and ace2 encode 664- and 647-amino acid residues, respectively, for the first time in Lepidopterans. Both ace genes have the conserved catalytic triads (acel: 314S, 440E and 554H; ace2: 275S, 404E and 518H), and contain the six cysteines which form the three intramolecular disulfide bonds. The signal peptide of acel and ace2 has 17- and 32-amino acid residues from the start methionine, indicating that both ace1 and ace2 are secreted proteins. In addition, 11 and 7 aromatic residues lining the active site gorge in ace1 and ace2, respectively. Phylogenetic analysis showed that insect AChEs are divided into two clusters, of which ace1 is the paralogous gene whereas ace2 is the othologous gene of Drosophila AChE.