

The Molecular Characterization of the *est31* gene, in the Colombian strain of *Culex quinquefasciatus*

The molecular basis of organophosphate resistance in the Colombian strain of *C. quinquefasciatus* (COL), is due to the two amplified esterases genes *est 31* and *est1*. These two esterases are situated in the same amplicon, but more than 10kb apart. This study was carried out using the PCR analysis of the genomic DNA and cDNA of COL strain using five primer pairs, designed from the *est* sequence.

The *est* genomic sequence, was 89% similar to the *est* genomic sequence. The sequence data also revealed, that ATG start codon, to the down stream 2227 nucleotides of *est* and *est* are highly homologous.

The partial sequence of *est* cDNA has 1406 nucleotides and 463 inferred amino acids. Five introns and six exons were identified within the region. The full-length *est 21* cDNA has a 1623 bp open reading frame and 540 inferred amino acids. A few alteration of sizes and greater variability in sequence were observed in the introns (81.4%) compared to exons. However there were ten amino acid changes and 71 silent nucleotide differences compared to the *est* sequence. The two esterases (*Est* & *Est 0*) have a different mobility in native PAGE gels. This could be due to different in the charge of *est* protein. They are probably alleles of the same gene, which have been amplified separately.

The selective advantage for *est* over *est* / *est* amplicon could be due to their genomic organization of the genes.