Expression of a Fusion Protein with the Cry1Ac Protein and the AaHIT Protein in **Bacillus thuringiensis** Cry-B Strain

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Expression of a fusion protein between B. thuringiensis crystal protein, the Cry1Ac1 and the AaHIT (Androctonus australis Hector insect toxin; Buthid scorpion) in B. thuringiensis Cry B strain was examined. The cry1Ac1 gene was cloned in B. thuringiensis-E. coli shuttle vector, pHT3101, under the control of the native cry1Ac1 gene promoter (pProAc) and a gene encoding AaHIT was inserted in XhoI site in the middle of the cry1Ac1 gene (pProAc-ScoR). B. thuringiensis Cry B strain carrying pProAc-ScoR (ProAc-ScoR/CB) produced an inclusion body of irregular shape and the expressed fusion protein is approximately 60 kDa in size. Sporulated cells and spore-crystal mixtures of ProAc-ScoR/CB had insecticidal activity against Plutella xvlostella larvae, showing LT₅₀ of ProAc-ScoR/CB lower than that of ProAc/CB. These results suggest that the fusion protein including a B. thuringiensis crystal protein and the AaHIT may be functionally expressed in B. thuringiensis. Further study is focusing on elucidation of the synergism of the two insecticidal proteins against insect larvae.