TOWARDS SAFE AND EFFICIENT *BT* MOSQUITOCIDES; DETERMINING CRY AND CYT PROTEIN PROFILES AND THE PRODUCTION OF TYPE I B EXOTOXIN BY LOCAL *BT* STRAINS 4 AND 6E

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Bacillus thuriengiensisis a ubiquitous, gram positive and spore forming bacterium which has been used to control certain insect species among the orders Lepidoptera, Diptera and Coleoptera. The genetic diversity of Bacillus thuringiensis (Bt) strains show differences according to the regions where they have been isolated. Hence, each habitat may contain novel Bt strains, which have toxic effect on target spectra of insects. The aim of this study was to identify the crystalline protein content of the local isolates with reference to Bacillus thuringiensis subsp. israelensis (Bti). To detect the protein profiles of crystal proteins obtained from Bti, Bt4 and Bt6e (local isolates), SDS-PAGE analysis was optimized. To identify the presence of cyt genes, the local strains were cultured on blood agar and observed for hemolysis. Both the local strains were subjected to HPLC analysis on a C-18 column for type 1 β exotoxin. The mobile phase was 50 mM KH₂PO₄ (pH 3.0) in double distilled water at a flow rate of 2 mL/min at 25 °C. It was found that Bti exhibited three main bands, 125-135kDa (Cry 4A and Cry 4B), 68kDa (Cry 11Aa) and 29kDa (Cyt 1Aa) and all the three including Bt4 and Bt6e contained Cyt1Aa protein. Bt4 showed a distinct band at 49kDa and Bt6e showed two distinct bands at 112kDa and 45kDa concluding that the Bt strains differed in their content of proteins. Both the local strains showed βhemolytic character, indicating the presence of cyt genes. Type 1 β exotoxin was not produced by Bt6e, but Bt4 produced a significant quantity (114.15 μg/ml). Accordingly Bt6e could be integrated into mosquito control programs, whereas Bt4 is not ideal due to the production of type 1 β exotoxin. This work has identified, that these novel Bt isolates differ in their protein profiles, driving them to be used as potential candidates for the industrial production of mosquito pathogenic Bt isolates.

Keywords: Bacillus thuringiensis, Bti, Bt4, Bt6e, Cry proteins