



International Journal of ChemTech Research CODEN (USA): IJCRGG, ISSN: 0974-4290, ISSN(Online):2455-9555 Vol.9, No.12 pp 114-124, 2016

Cloning, sequencing and expression of the xylanase gene from *Bacillus pumilus* GH in *Escherichia coli*

Maha T. H. Emam¹*, Karima A. Mohamed¹, Fatma M. I. Badawy² and S. A. Ibrahim²

¹Genetics & Cytology Dept., National Research Centre, Dokki, Cairo, Egypt. ²Genetics Dept., Faculty of Agriculture, Ain Shams University, Egypt

Abstract : The thermostable endo-1,4-beta-xylanase gene of *Bacillus pumilus* GH strain was isolated from chromosomal DNA using specific primers designed from *Bacillus pumilus* xylanase gene given in gene bank database then cloned into pET29a (+) vector and transformed into *E. coli* DH5a. The positive clone was selected, sequenced and submitted to gene bank with the accession number KT757524.1. The open reading frame of the xylanase gene was 687 bp encoding a protein of 228 amino acids with a molecular mass of 23 kDa. The sequence of *Bacillus pumilus* GH xylanase gene showed 99 % similarity with other xylanase genes of different *Bacillus pumilus* strains, differ only in two nucleotide bases at positions 579 and 600. The recombinant plasmid was subcloned into the expression host *E.coli* BL21 (DE3) and successfully expressed. The total activity of xylanase was 9 U/ml, 52% (4.7U/ml) of the activity was extracellular and 48 % (4.3 U/ml) intracellular.

Key words: *Bacillus pumilus*; *E. coli*; xylanase; gene cloning and expression; sequence analysis.

Maha T. H. Emam et al /International Journal of ChemTech Research, 2016, 9(12): 114-124.
