F322 Cloning and nucleotide sequence of a *Peanibacillus* sp. KCTC 8848P xylanase gene and its expression in *Saccharomyces cerevisiae*.

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The xylanase gene(xynA) from Paenibacillus sp. KCTC 8848P has been cloned in Escherichia coli and its complete nucleotide sequence was determined. The structural gene of xynA revealed 636 bp encoding a mature protein of 184 amino acids and a signal peptide of 28 amino acids. The amino acid sequence of the xynA gene showed high homology to the xylanase of Aeromonas caviae with 83% identity and the xylanases of other Bacillus spp with 79% identity, and belonging to family G xylanases. Crude xylanase from E. coli transformant was endoxylanase to degrade xylan to xylose, xylobiose and xylotriose. The xynA gene containing its own signal sequence was introduced into yeast expression vector. The Saccharomyces cerevisiae transformants harboring the recombinant plasmids secreted xylanase into culture medium.

F323 Molecular cloning and nucleotide sequence of an endo- β -1,4-glucanase gene from *Paenibacillus* sp. KCTC 8848P and its expression in Saccharomyces cerevisiae

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An endo- β -1,4-glucanase(Egl) gene from Paenibacillus sp. KCTC 8848P was cloned in $Escherichia\ coli$, and its complete nucleotide sequence was determined. The Egl gene had open reading frame(ORF) of 1191bp encoding 397 amino acids with calculated molecular weight of 44539 Da. The proposed signal sequence consisted of 32 amino acid residues. The deduced amino acid sequence revealed 94% identity to endo- β -1,4-glucanase of $Bacillus\ polymyxa$, and it belongs to the cellulase family A. To examine whether the signal sequence of Egl can act functionally in a yeast, the Egl gene containing its own signal sequence was introduced between ADCI promoter and CYCI transcription terminator of the yeast expression vector. The $Saccharomyces\ cerevisiae$ transformants harboring such recombinant plasmids secreted endo- β -1,4-glucanase into culture medium.