

Nucleotide sequence of a cyclodextrin glucosyltransferase gene, *cgtA*, from *Bacillus licheniformis*

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Cyclodextrin glucosyltransferases (CGTases), EC 2.4.1.19, convert starch into cyclic glucosyl oligosaccharides (cyclodextrins) having a hydrophilic surface and a hydrophobic core (1). Thus, cyclodextrins bind to and solubilize hydrophobic materials. Recently, a potential therapeutic benefit for cyclodextrins was demonstrated (2). Here we report the nucleotide sequence of a unique gene encoding a CGTase cloned from a strain of *Bacillus licheniformis* (3). The *B. licheniformis* gene, called *cgtA*, was cloned using procedures similar to those described (4) except the *B. licheniformis* library was prepared in a pUC19 derivative. *E. coli* transformants expressing a full length *cgtA* gene were initially identified on the basis of starch clearing ability as described (4). The *cgtA* clones were confirmed by measuring the conversion of 2% maltodextrin solution to cyclodextrin following growth in liquid culture. Cyclodextrins were identified and quantitated by HPLC using a cyclodextrin assay column purchased from Advanced Separation Technologies, Whippany, NJ; α and β cyclodextrins were the

principal products obtained. The *cgtA* gene, sequenced as described (5), is contained within a 2516 base pair Sau3A to SphI fragment encoding an open reading frame of 718 amino acids. The translated sequence exhibits 58% and 66% amino acid similarity, respectively, to the *B. macerans* CGTase (4) and either *Bacillus* sp. 1011 (6) or 38-2 (7). This similarity extends throughout the entire open reading frame except for the amino terminal leader sequences.

REFERENCES

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-209 GATCATAACTCGAGGGCCATGCCTTTAAACCCAAAGCGGTTTAGGACAACCCCGGTAACCTCCCAGGAGATACCGGGTGTGGCGGTGC
-119 GAAGGAGGAAGTCTTTAAGCCACAGAATGTCCTCTCGTACAGTTGTAAGCGCTGTTATTCGAGAGCCAGGTTAGTTGTTTTGTTGAAA
-29 ACTACTACATTCATGAAGGGTGGATTACC ATGTTTCAAATGGCCAAACCGCTTCTCCTCAGTACCACGCTAACGTTTCAGCCTGCTTGCC
61 GGCAGTGCAATGCCTTTCCGCTGCTTCCCGGATTTATGCCGATGCCGATACAGCTGTCAACCAACAGCAGAATTCAGTACGGATGTC
151 ATCTATCAAGTTTTACGGACCGATTTCTGGATGGTAACCCATCCAACAACCTACCGGGGCTGCCTTTGATGGCAGCTGACGCAACCTG
241 AAGCTGTACTGCGGTGGGACTGGCAGGGATGGTCAACAAAATCAATGACAACTATTCAGTGACTGGGTGTCACGGCCCTCTGGATC
331 TCCCAGCCTGTCGAAAAATTTTCCGCTACCATCAACTACAGCGGTGTAACCAATACAGCCTATCACGGCTATTGGGCACGGGATTTCAAG
421 AAGACCAATCCGTAATTCGGAACCATGACCGATTTTCAGAACTCTGGTAACCAACCGCCATGCGAAAGGCATCAAAAATCATATTGATTTTC
511 GCGCCAAACCATACGTCCTCCGTCATGAAACCGATACCTCTTCGCGGAGAACCGGCAAACTGTACGATAACGGCAACCTGGTTGGCGGG
601 TACACCAATGATACCAACGGATATTTCCACCACAATGGCGGCTCCGATTTCTCCACTTTGAGAAATGGCATTACAAAAACCTTACGAT
691 CTGGCCGATCTGAATCACAATAACAGCAGCATCGATACATATTTCAAAGACGCCATCAAGCTGTGGCTTGATATGGCGCTGGACGGCATT
781 CGTGTGATGCGGTCAAGCACATGCCTCAGGGCTGGCAGAAAGTGGATGTCTATCTATGACACAAAGCCGGTATTACCTTCGGC
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1141 GCCTTACAGCTACCTCACGGCGTACCTGCCATCTATTATGGTACCAGCAGTATCTGACTGGGAACGGCGACCCGATACACCGGGGC
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1681 ACCGTCTACTTCGGTACAACAGCCGTTACGGGCTCAGCCATTACTTTCATGGGAAGACTCAGATCAAAGTCAACATTCACCAGTAGCA
1771 GCGGGTACTATGCAGTGAAGTAGCTGCCAATGGTGTGAACAGCAACCGCTATAACGATTCACCATCCTTAGCGCGATCAGGTATCG
1861 GTGCGGTTGCTCATCAATAATGCCAACTGCGCTGGCGGAGAAATATCTACCTGACAGGCAACGTGCCGAACTCGGTAACCTGACCCACA
1951 GGTGAGCTTCCATGGACCGGCTTTCAATCAGTGCATCCAGCCTACCGACTTGGTATTATGACGTAAGTGTTCGCGCGGGAACAG
2041 CTGGAATCAAGTCTTCAAGAAAACGGCGCTACCATACGTGGGAAGTGGATCCAATCACACCTTTACAACACCGCAGTGGTACT
2131 GCCACAGTAACGATAAACTGGCAATACCTAGCCATAATCCATTAATGAAGCCGGAGCAGCGCTTATACAGCTACACTCCGGGTTTTGA
2221 CGTTTAATCTGTTTCTGGTTTACGGTAAGTTCGAATGGACTCAATGTGGTCCGGCAAGGATTCGGTTCGAGCAGCAGCATGC

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