

Sequence of the *Bacillus thuringiensis* phosphatidylinositol specific phospholipase CDennis J.Henner, Maria Yang, Ellson Chen, Renate Hellmiss, Henry Rodriguez and Martin G.Low¹Genentech Inc., 460 Point San Bruno Boulevard, South San Francisco, CA 94080 and ¹Department of Physiology and Cellular Biophysics, Columbia University, New York, NY 10032, USA
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A number of eukaryotic proteins have been shown to be anchored to membranes by covalent linkage to a phosphatidylinositol molecule(1). Bacterial phospholipases have been instrumental in the identification of proteins which are anchored by this mechanism. To increase the availability of this enzyme, and to possibly understand its mechanism of action, we have isolated the gene for the phosphatidylinositol specific phospholipase C of *B. thuringiensis*. Oligonucleotide probes based on the amino acid sequence (2) of the purified enzyme were used to isolate clones in pBR322. The sequence was determined by the dideoxy sequencing method(3). The amino acids corresponding to the determined amino acid sequence are underlined. When either *E. coli* or *B. subtilis* was transformed with this gene on a plasmid, enzymatic activity could be detected in amounts at least 500-fold higher than nontransformed cultures. No obvious sequence similarities were found in the Dayhoff protein sequence data base, nor were there similarities to the recently reported sequence of a mammalian phosphoinositide-specific phospholipase C (4).

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1 AAGTCCTTAAATTCTAATTAGTATGTAATGATTTCACATTTCAGTTGATGAAATTTCAACTTCTATGCAATTTCATATTGATGAAATGTTATTATTTTATG
101 AACTAGCTTATATTATTCATAAACTTGTGATGTAACAGAAAAGTGAGGGATAATAATGAGCAATAAGAAGTTAATTGAAATTATTATCATATGTA
1 M S N K K L I L K L F I C S
201 GTACAATATTATACATTTGATTTGCTTACATGATAAGAGAGTAGTTGCAGCTAGCCTCTGTTAATGAGCTTGAAATTGGTCAAAATGGATGCCAAC
15 T I F I T F V F A L H D K R V V A A S S V N E L E N W S K W M Q P
301 TATACTGTATAATATCCCGTTAGCACGAATTCAATTCCAGGAACACAGATAAGTGGGACGTTCAAGTGCAAAATCCGATAAGCAAGTGTGGGAATG
48 I P D N I P L A R I S I P G T H D S G T F K L Q N P I K Q V W G M
401 ACAGCAAGAATATGATTTCGTATCAAATGGACCATGGAGCTCGCATTTTGATATAAGAGGACGTTAACAGATGATAATACGATAGTTCTCATCGT
81 T Q E Y D F R Y Q M D H G A R I F D I R G R L T D D N T I V L H H G
501 GGCCATTATATCTTACGTAACTCGATGAATTCTAAATGAAAGCAGAACATTAAAAAGATAACCCGAGTGAACAAATTATTATGCTTTAAAAAA
115 P L Y L Y V T L H E F I N E A K Q F L K D N P S E T I I M S L K K
601 AGAGTATGAGGATATGAAAGGGCAGAGGTCAATTAGTAGTACGTTGAAAAAAATTATTGTTGATCCTATCTTTAAACAGAAAGGAATA
148 E Y E D M K G A E G S F S S T F E K N Y F V D P I F L K T E G N I
701 AAACCTGGAGATGCTCGTGGGGAAATTGACTAAAGGATAGTGGTAGTAATGAACTGGAGGATAATAATTATTATGGCAGATAATGAGA
181 K L G D A R G K I V L L K R Y S G S N E S G G Y N N F Y W P D N E T
801 CGTTTACCAACACTGAAACCAAAATGTAATGTAACAGTCAAGATAAAATTAAGTGAATTATGAGAAACTAAATCTATTAAGATAACGATGGA
215 F T T T T V N Q N V N V T V Q D K Y K V N Y D E K V K S I K D T M D
901 TGAAACGATGAACAAATACCGAGGATTTAAATCATCTATATAATTAAACAGCTTGTCTCTGGTGTACAGCATGGAATAGTCATATTACTACGGCT
248 E T M N N S E D L N H L Y I N F T S L S S G G T A W N S P Y Y Y A
1001 TCCTATATAAAATCTGAAATTGCAACAGATATAAAACAAAGAATCTACAAAGACTAGCTGGCTGGTAATCAAGACTACATAAAATGAAAGTGGCACCAT
281 S Y I N P E I A N D I K Q K N P T R V G W V I Q D Y I N E K W S P L
1101 TATTGTATCAAGAAACTATAAGAGCAATAAGCTTAAAGAATAAGATGATGAGAGTAAGAGCATAAAATATGCTCTTTCTTTCC
315 L Y Q E V I R A N K S L I K E O
1201 ATAACCAACAATAGAAATTGACTTTATAATAGAAAAAA

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