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Cyclic AMP Directly Activates NasP, an N-Acyl Amino Acid Antibiotic Biosynthetic Enzyme Cloned from an Uncultured β-Proteobacterium

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The cyclic AMP (cAMP)-dependent biosynthesis of N-acetylphenylalanine antibiotics by NasP, an environmental DNA-derived N-acyl amino acid synthase, is controlled by an NasP-associated cyclic nucleotide-binding domain and is independent of the global cAMP signal transducer, cAMP receptor protein. A 16S rRNA gene sequence found on the same environmental DNA cosmid as NasP is most closely related to 16S sequences from β-proteobacteria.

High-throughput sequencing of DNA extracted directly from environmental samples indicates that uncultured bacteria are likely to be a very rewarding source of interesting new proteins (26). While it is easy to infer the existence of a large number of novel proteins from the data generated by environmental DNA sequencing projects, it has been much more challenging to functionally access these new proteins (12, 13). Screening environmental DNA clones using simple assays that respond to a wide array of stimuli should increase the likelihood of functionally cloning new proteins from environmental DNA and, in turn, identifying new biological phenomena from the genomes of uncultured bacteria. Large-scale screening of environmental DNA libraries for antibacterially active cosmids clones has identified a number of new N-acyl amino acid synthases that confer the production of long-chain N-acyl amino acid antibiotics on Escherichia coli (4–7). The identification of multiple unique N-acyl amino acid-producing clones in environmental DNA libraries suggests that these compounds could play an important, if undefined, role for soil microbes and that their biosynthesis might therefore be highly regulated. Here we report the discovery and characterization of an environmental DNA-derived N-acyl amino acid synthase (NasP) that is directly regulated by cyclic AMP (cAMP), defining a new role for this widely studied universal second messenger.

The first insight into how N-acyl amino acid biosynthesis is regulated in some uncultured bacteria comes from NasP, an environmental DNA-derived N-acyl amino acid synthase that confers the production of long-chain N-acylphenylalanines on Escherichia coli (Fig. 1). While all previously described N-acyl amino acid synthases are predicted to contain one domain, a conserved-domain search with NasP indicated the presence of two domains, an N-terminal N-acyl amino acid synthase domain and a C-terminal cyclic nucleotide-binding domain that is related to both eukaryotic and prokaryotic cAMP-binding domains (14). Signal transduction pathways present in both eukaryotes and prokaryotes employ cAMP as a second messenger. In prokaryotes, cAMP is biosynthesized from ATP by an adenylate cyclase and its presence is then detected by cAMP receptor proteins, a small family of transcription factors found in the gamma subdivision of the proteobacteria (2). Although cAMP receptor proteins are thought to be the main cAMP signal transducer in prokaryotes, non-cAMP receptor protein-associated cAMP-binding domains have been found in the genomes of many sequenced bacteria (10, 15, 17). If these domains prove to be functional, it would indicate that cAMP directly regulates a broad range of enzymatic activities in bacteria. The functional cloning of NasP from environmental DNA provides an opportunity to confirm a new role for cAMP-binding domains in bacteria and to investigate the mechanism by which N-acyl amino acid biosynthesis is regulated in some uncultured bacteria.

The environmental DNA cosmid clone CSL142, from which NasP was cloned, was originally identified in a high-throughput screen for antibacterially active environmental DNA clones (3). One- and two-dimensional nuclear magnetic resonance analysis of the antibacterially active material purified from ethyl acetate extracts of this clone indicated that the active material was a mixture of long-chain N-acylphenylalanines (see the supplemental material). The major clone-specific compounds present in the extracts were analyzed by high-resolution mass spectrometry and found to contain both fully saturated and monounsaturated fatty acid side chains ranging from 14 to 16 carbons in length (Fig. 1). Sequencing of transposed cosmids that no longer conferred antibacterial activity on E. coli indicated that a single open reading frame (NasP) with an N-terminal N-acyl amino acid synthase domain and a C-terminal cyclic nucleotide-binding domain was responsible for the production of these compounds. Sequences that show significant sequence identity (>20%) to NasP in a BLAST search are all hypothetical proteins of unknown function (NP_924045 and NP_924043 from Gloeobacter violaceus PCC 7421 and ZP_01079393 from Synechococcus sp. strain RS9917).

The role of the NasP-associated cAMP-binding domain in
the regulation of N-acylphenylalanine biosynthesis was examined using NasP expressed as a glutathione S-transferase fusion protein (NasP:GST) in adenylate cyclase (cya)- and cAMP receptor protein (crp)-deficient strains of E. coli (Fig. 2). The GST fusion construct of an N-acyl amino acid synthase that does not contain a cAMP-binding domain, FeeM:GST, was used as the positive control in these studies (5). Both constructs confer the production of N-acyl amino acids on E. coli strains lacking the cAMP receptor protein; however, in adenylate cyclase mutants where no cAMP is present, only the control construct continues to confer the production of N-acyl amino acids on E. coli (Fig. 2). The addition of cAMP (1 mM) to cya crp double mutants transformed with NasP:GST restored the ability of this construct to confer the production of N-acylphenylalanines on the E. coli host (Fig. 2). The biosynthesis of N-acyl amino acids is therefore cAMP dependent yet independent of the cAMP receptor protein. A direct interaction between NasP and cAMP was confirmed by affinity chromatography using cAMP-presenting affinity resin (Fig. 3). Removal of the proposed cyclic nucleotide-binding domain from this construct abrogated binding to the cAMP affinity resin but did not prevent affinity purification using the GST affinity resin (Fig. 3).

In cAMP receptor protein-associated cAMP-binding domains, cAMP binds in an anti conformation with the base oriented away from the sugar phosphate, while in many eukaryotic cAMP-binding domains, cAMP binds predominantly in a syn conformation with the base oriented toward the sugar phosphate (21, 23). The cAMP analog 8-Br-cAMP, which adopts a syn conformation, can therefore be used to preferentially activate eukaryotic protein kinase A (PKA)-like cAMP-binding domains over prokaryotic cAMP receptor protein-like cAMP-binding domains (18, 20). Interestingly, the addition of 8-Br-cAMP to cultures of cya crp double mutants transformed with NasP restored the ability of these cultures to produce N-acyl amino acids. Subtle differences in phosphate binding loops are thought to play a major role in altering the conformation of cAMP bound to a cAMP-binding domain. In prokaryotic cAMP receptor proteins, the variable residue in a conserved 3-amino-acid sequence (R-X-A) found in the phosphate-binding loop is a serine which hydrogen bonds to cAMP bound in the anti conformation. In eukaryotic PKA sequences this variable residue is most frequently an amino acid that cannot fulfill this hydrogen bond (i.e., A, N, V, or Q) (10, 21). The cAMP-binding domain from NasP contains a PKA-like R-A-A phosphate-binding loop sequence motif which would suggest that it binds cAMP in the syn conformation and would explain the activation of N-acyl amino acid biosynthesis by 8-Br-cAMP.

Although NasP appears to bind cAMP in the same conformation as many eukaryotic cAMP-binding domains, a full-length 16S rRNA gene found on the same cosmid clone as NasP indicates that it is, in fact, derived from a bacterium.
16S rRNA gene sequences from cultured bacteria are those from environmental samples (16S rRNA gene associated with NasP is most closely related to 16S rRNA gene sequences from cultured bacteria) but no longer binds the cAMP affinity resin. As is often seen with GST fusion proteins, a 60-kDa protein, likely GroEL, coelutes with GST fusion constructs (24).

**Nucleotide sequence accession numbers.** Nucleotide sequences for NasP (accession number DQ224236) and the CSl142 16S rRNA gene (accession number DQ224237) have been deposited with GenBank.

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