

Discovery of phosphonic acid natural products by mining the genomes of 10,000 actinomycetes

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Although natural products have been a particularly rich source of human medicines, activity-based screening results in a very high rate of rediscovery of known molecules. Based on the large number of natural product biosynthetic genes in microbial genomes, many have proposed “genome mining” as an alternative approach for discovery efforts; however, this idea has yet to be performed experimentally on a large scale. Here, we demonstrate the feasibility of large-scale, high-throughput genome mining by screening a collection of over 10,000 actinomycetes for the genetic potential to make phosphonic acids, a class of natural products with diverse and useful bioactivities. Genome sequencing identified a diverse collection of phosphonate biosynthetic gene clusters within 278 strains. These clusters were classified into 64 distinct groups, of which 55 are likely to direct the synthesis of unknown compounds. Characterization of strains within five of these groups resulted in the discovery of a new archetypical pathway for phosphonate biosynthesis, the first (to our knowledge) dedicated pathway for H-phosphinates, and 11 previously undescribed phosphonic acid natural products. Among these compounds are argolaphos, a broad-spectrum antibacterial phosphonopeptide composed of amino-methylphosphonate in peptide linkage to a rare amino acid *N*⁵-hydroxyarginine; valinophos, an *N*-acetyl L-Val ester of 2,3-dihydroxypropylphosphonate; and phosphonocystoximate, an unusual thiohydroximate-containing molecule representing a new chemotype of sulfur-containing phosphonate natural products. Analysis of the genome sequences from the remaining strains suggests that the majority of the phosphonate biosynthetic repertoire of Actinobacteria has been captured at the gene level. This dereplicated strain collection now provides a reservoir of numerous, as yet undiscovered, phosphonate natural products.

natural products | genome mining | phosphonic acid | antibiotic

The use of genomic data to enable discovery of novel biological processes, often referred to as genome mining, has the potential to revolutionize numerous areas of modern biology, including the field of natural product discovery. These biologically produced small molecules have been the source of, or inspiration for, nearly two-thirds of all human medicines (1), yet research in this area has dwindled in recent years due to, among other reasons, high costs and increasing rates of rediscovery (2–4). Although a number of solutions to this dilemma have been proposed, many within the natural product biosynthesis field have suggested that the solution lies in genomics (5, 6). Thus, by focusing research efforts on strains that encode genes for the biosynthesis of uncharacterized natural products, one can dereplicate, streamline, and accelerate the discovery process. Indeed, genome mining has led to the discovery of several novel natural products, but most efforts have been limited to individual strains or small collections (5–9). Mining of metagenomic libraries has also produced a number of new compounds (10–12), but it has not been

generally adopted. If we hope to revitalize the use of natural products in the pharmaceutical industry, genome mining must be shown to be a high-throughput discovery process complementary or superior to existing methods (13, 14). Here, we demonstrate the feasibility of this approach in a campaign to identify the full genetic repertoire of phosphonic acid natural products encoded by a collection of over 10,000 actinomycetes.

Phosphonic acid natural products possess several traits that make them ideal candidates for large-scale genome mining. First and foremost, phosphonates have great pharmaceutical potential, with a commercialization rate of 15% [three of 20 isolated compounds (15)]. This percentage is much higher than the 0.1% average estimated for natural products as a whole (16). The potent bioactivity of phosphonates derives from their chemical mimicry of essential metabolites, including phosphate esters and anhydrides, as well as carboxylate reaction intermediates (15). Given the ubiquitous presence of these chemical moieties in biology, phosphonates are unrivaled in the range of targets they can potentially affect. Consistent with this idea, phosphonate natural products with herbicidal, insecticidal, antibacterial, antiparasitic, antiviral, and antihypertensive activities are known (15, 17). Notable examples include fosfomycin (Monurol), clinically prescribed for

Significance

The discovery of natural products, an important source of human medicines, is critical for the development of new therapeutics against health threats, including cancer and multidrug-resistant pathogens. Yet, in recent years, industrial development of pharmaceuticals from natural products has been stymied due to a variety of reasons, including the repeated discovery of previously known compounds. Here, we demonstrate large-scale genomics as one potential solution to this problem by mining a collection of 10,000 actinomycetes for novel phosphonic acids, an important class of natural products with antimicrobial, antiviral, antimalarial, and herbicidal activities. The framework described here provides a foundation for rapid, large-scale discovery of other classes of natural products and their use as lead compounds in the pharmaceutical industry.

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acute cystitis; FR-900098 and fosmidomycin, antimicrobials undergoing clinical trials for malaria; and phosphinothricin, the active component in several commercial herbicides (Liberty, Basta, and Rely). Second, the methodology needed for gene-based discovery of phosphonate biosynthetic loci has been rigorously established (17–23). This method relies on the fact that all but two characterized phosphonate biosynthetic pathways begin with the enzyme phosphoenolpyruvate (PEP) mutase (encoded by *pepM*) (14). Thus, amplification of an internal fragment of *pepM* with degenerate PCR primers is sufficient to identify strains or plasmid clones that encode phosphonate biosynthetic pathways. Third, gene-based surveys have proven that phosphonate biosynthesis is relatively common in microorganisms, with the largest reservoir of unexplored pathways observed within Actinobacteria (23). Finally, the carbon-phosphorus bond endows phosphonates with unique chemical properties that enable their chemispecific detection from complex mixtures of metabolites using MS and NMR spectroscopy, even when the structure of the molecule in question is unknown (24, 25). These methods facilitate the isolation and characterization of phosphonate natural products by enabling their direct detection during production screening and purification.

Results and Discussion

Identification and Genome Sequencing of Actinobacteria That Contain Phosphonate Biosynthetic Pathways. A collection of over 10,000 actinomycetes, composed of *ca.* 2,500 isolated in our laboratory (23) and nearly 7,500 from the Agricultural Research Service (ARS) Culture Collection (NRRL) at the National Center for Agricultural Utilization Research, were screened for the presence of *pepM*. The NRRL collection houses the largest public repository of Actinobacteria, encompassing species from 117 different genera, important industrial strains, and the retired strain collections of several prominent streptomycete researchers (e.g., S. A. Waksman, D. Gottlieb, E. B. Shirling). To enable rapid genetic screening, genomic DNA was isolated from all strains and arrayed into a 96-well format. Blind PCR screening of this library identified 403 putative phosphonate producers. Subsequent draft genome sequencing of all candidate strains confirmed the presence of *pepM* in 278 strains, including 244 from the NRRL collection; 30 from our laboratory collection; and four additional strains specifically added to the study because of their known ability to produce phosphonothrixin (26), FR-900098 (27), and fosmidomycin (28) (*SI Appendix, Table S1*). The majority (87%) of the *pepM*⁺ strains are *Streptomyces*, with the remainder scattered among 24 genera, a distribution that mirrors the composition of the strain collection. A single copy of *pepM* was found in all but five strains, each of which encodes two distinct *pepM* homologs. Significantly, all known phosphonate producers deposited in

the NRRL collection, including producers of fosfomycin, dehydrophos, and phosalacine, were rediscovered in our blind screening, validating the sensitivity and thoroughness of our *pepM*-based approach.

Dereplication of Phosphonate Biosynthetic Pathways and Production Screening. Although recent years have seen great improvements in subtractive MS- and NMR-based screening methods for new natural products (29, 30), in our hands, ³¹P NMR spectroscopy has proven more reliable than MS-based methods for detection of phosphonates of unknown structure. However, NMR screening of 10,000 strains is impractical. Therefore, the genomic data were used to identify, classify, and dereplicate phosphonate biosynthetic pathways within the candidate strains before characterization of the natural products they produce. For this process, we used two established methods for linking genetic data to phosphonate natural product diversity (23). Initially, we examined the relatedness of the *PepM* proteins encoded by the 278 strains. This analysis revealed 168 unique sequences that distributed into 78 groups at 80% sequence identity (Fig. 1A). Because natural product biosynthetic genes are almost always clustered together in a single chromosomal locus, we also examined the similarity of the genes surrounding *pepM*. This parallel approach uncovered 64 discrete gene neighborhoods (Fig. 1B). Known phosphonate producers map to nine of these groups, most of which also include additional strains. Thus, our genome mining effort revealed 55 new potential producers of hydroxynitrilaphos (cyanohydroxymethylphosphonate), 8 of plumbemycin, 7 of fosfazinomycin, 2 of dehydrophos, and 1 of fosfomycin. We also identified the putative biosynthetic gene cluster for phosphonothrixin in the genome sequence of the only known producer, *Saccharothrix* ST-888 (26). Our observation of phosphonothrixin in the spent media of four of the 13 strains that carry this gene cluster (*SI Appendix, Fig. S2 and Table S2*) provides strong evidence that these genes direct the synthesis of this herbicidal compound (31). After eliminating strains linked to known products, we were left with a dereplicated collection of 192 strains grouped into 69 *PepM* clades and 55 *pepM* gene cluster families. By both measures of diversity, greater than 85% of the pathways reported here direct synthesis of new phosphonate natural products.

Production screening of all 244 *pepM*⁺ strains from the NRRL collection by ³¹P NMR spectroscopy suggested that at least 45 produced phosphonates. Three of these strains, *Streptomyces monomycin* NRRL B-24309, *Streptomyces* sp. NRRL S-474, and *Streptomyces* sp. NRRL S-481, were of particular interest because they elicited positive responses in a phosphonate-specific bioassay (18) and because the genomic information illustrated that their biosynthetic gene clusters were novel.

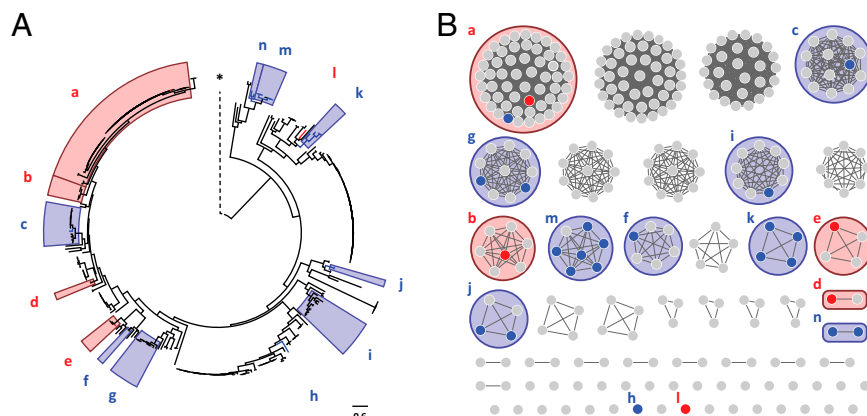


Fig. 1. Diversity of phosphonate biosynthesis in Actinobacteria. (A) Maximum-likelihood phylogeny of actinomycete *PepMs* using methylisocitrate lyase as the outgroup (*). Complete labeling of strain names is provided in *SI Appendix, Fig. S1*. (B) Network diagram of phosphonate gene clusters. In both diagrams, highlighting indicates clades and strains containing pathways for previously described phosphonates (blue) and those phosphonates newly isolated in this study (red). Groups encoding known phosphonate natural products are as follows: a, hydroxynitrilaphos; b, phosphonocystoximate; c, phosphonothrixin; d, valinophos; e, argolaphos; f, dehydrophos; g, fosfazinomycin; h, fosmidomycin; i, plumbemycin and phosacetamycin; j, FR-900098; k, phosphinothricin peptides; l, DMPT; m, fosfomycin; n, 2-hydroxyethylphosphonate polyglycans. Phosphonate gene clusters are available (www.igb.illinois.edu/labs/metcalf/gcf/Phosphonates.html).

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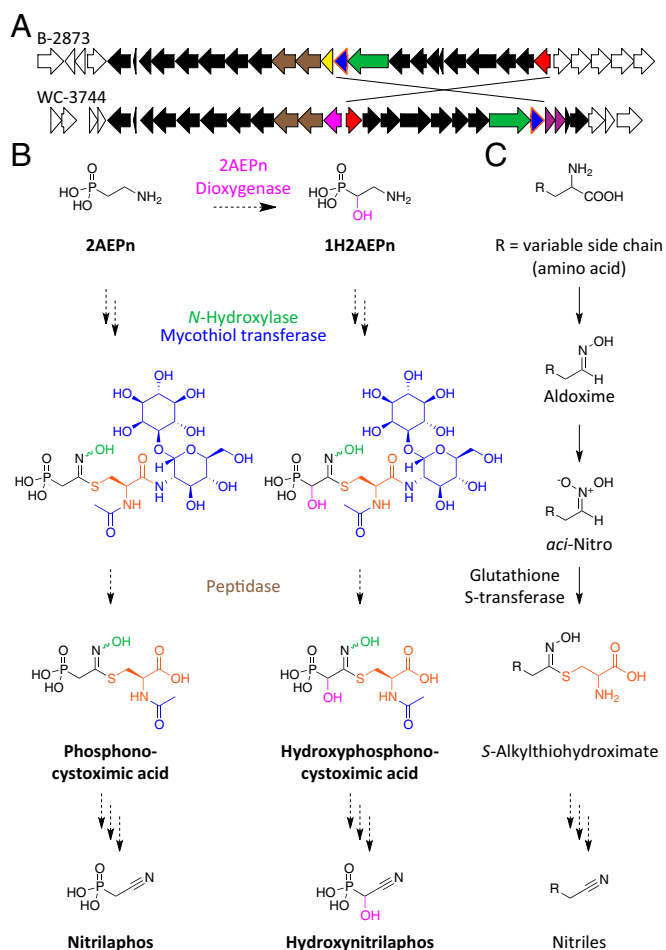


Fig. 3. Comparisons between phosphonocystoximic acid and hydroxynitrilaphos biosynthesis. (A) Putative gene clusters. The majority of genes are shared between the two pathways, with some notable differences. Genes for α -ketoglutarate-dependent dioxxygenase (magenta) and PnPy decarboxylase (purple) are absent from the phosphonocystoximate gene cluster, which instead contains a gene for pyridoxamine-phosphate oxidase (yellow). Half of the shared genes encoding the two pathways are inverted relative to each other. Other labeled genes putatively encode for PepM (red), mycothiol transferase (blue, outlined in red), *N*-hydroxylase (green), and peptidases (brown). (B) Proposed biosynthetic pathways for phosphonocystoximic acids and cyanomethylphosphonic acids. (C) Proposed biosynthetic pathway for glucosinolates, adapted from Halkier and Gershenzon (41). Several intermediates in glucosinolate biosynthesis and metabolism share similarities to the compounds from these pathways. Glutathione is the source of Cys (orange) in glucosinolate biosynthesis, whereas mycothiol (orange and blue) may contribute *N*-acetylcysteine in phosphonocystoximic acid biosynthesis.

strains are biosynthetic intermediates or the products of degradation remains to be determined.

Valinophos, a Phosphonopeptide Produced by a New Archetypical Pathway. *Streptomyces durhamensis* NRRL B-3309 was the next strain selected for further analysis because of the unusual content of its *pepM* gene neighborhood. Two novel phosphonate natural products, 2-hydroxy-3-phosphonopropanoic acid and (*R*)-2,3-dihydroxypropylphosphonic acid (DHPPA), were purified from spent media (Fig. 4A). These compounds are likely early intermediates in the formation of a fourth new phosphonate that we name valinophos, an *N*-acetyl-L-Val ester of DHPPA that was also produced by *S. durhamensis*. Only DHPPA was purified in sufficient

quantity to assess bioactivity, which slightly inhibited growth of *M. smegmatis* (SI Appendix, Fig. S5).

We propose a biosynthetic pathway for valinophos that is supported by the content of the *pepM* gene cluster (Fig. 4B) and the predicted biochemistry of the encoded proteins. Like the phosphonocystoximate cluster, a known example of an enzyme that could be coupled with PEP mutase (VlpA) to drive the thermodynamically unfavorable rearrangement of PEP to PnPy is absent. Instead, VlpB, a putative short-chain dehydrogenase, may reduce PnPy to 2-hydroxy-3-phosphonopropanoic acid (phosphonolactate) in a reaction analogous to the highly favorable reduction of pyruvate to lactate by NADH-dependent lactate dehydrogenases (equilibrium constant, $K_{eq} = 1 \times 10^5$) (42). Analogous to the proposed formation of 1-oxo-2-phosphorylethylphosphonic acid by the phosphoglycerate kinase DhpB in dehydrophos biosynthesis (37), phosphorylation of phosphonolactate by VlpD would yield 2-hydroxy-3-oxo-3-phosphoxypropylphosphonic acid. Reduction by VlpC, homologous to *N*-acetyl γ -glutamyl phosphate (AGP) reductase, results in 2-hydroxy-3-oxopropylphosphonic acid that could be converted to DHPPA via another reduction by VlpE (alcohol dehydrogenase). Although ATP-grasp ligases typically adjoin substrates by amide linkages, these enzymes have also been shown to form ester bonds (43). Here, ATP-grasp ligases VlpF and VlpG are proposed to install L-Val onto DHPPA sequentially and then to acetylate the terminal amine to yield valinophos. Interestingly, *vlpBCDE* appears as a conserved cassette within the *pepM* gene clusters for phosphonothrixin. This observation suggests the early steps for valinophos are the archetype of a new branch of phosphonate biosynthetic pathways, from which phosphonothrixin also originates.

Dedicated Pathway for H-Phosphinate Biosynthesis. *Nonomureae* sp. NRRL B-24552 was the final strain chosen for study due to the similarity of its *pepM* gene cluster to the gene cluster involved in the synthesis of phosphinothricin tripeptide (PTT), the only known phosphinate natural product. However, unlike the well-characterized PTT biosynthetic gene clusters, *Nonomureae* sp. NRRL B-24552 lacks the genes needed for tripeptide biosynthesis (*phsABC*; nonribosomal peptide synthetases) and formation of the C-P-C moiety of phosphinothricin (*phpK*; P-methylase) (19), suggesting that this organism may instead produce an H-phosphinate (H-P-C bond motif) (Fig. 5A). Consistent with this idea, we purified 2-phosphinomethylmalic acid and desmethylphosphinothricin (DMPT) from extracts of this strain (Fig. 5B). DMPT

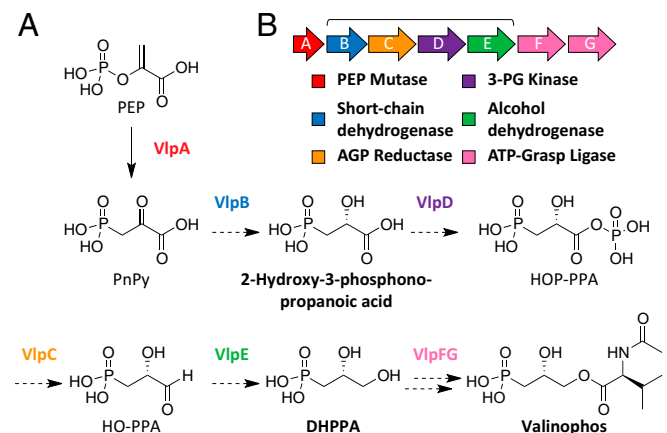


Fig. 4. *S. durhamensis* B-3309. (A) Isolated compounds and proposed pathway for valinophos biosynthesis. (B) Putative Vlp gene cluster, with the genetic cassette conserved in phosphonothrixin biosynthesis indicated by brackets. HO-PPA, 2-hydroxy-3-oxopropylphosphonic acid; HOP-PPA, 2-hydroxy-3-oxo-3-phosphoxypropylphosphonic acid; 3-PG, 3-phosphoglycerate.

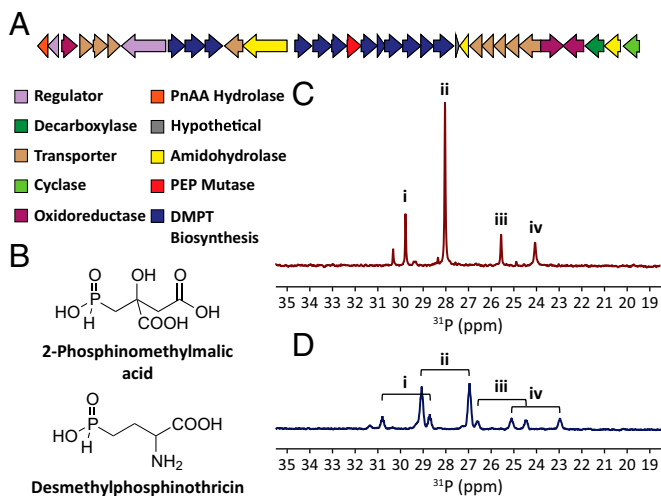


Fig. 5. *Nonomuraea* B-24552. (A) Putative H-phosphinate biosynthetic genes. Core genes encoding for predicted enzymes in DMPT biosynthesis are shown in dark blue (*pepM* is shown in red). (B) Isolated compounds. (C) ^1H -decoupled ^{31}P NMR analysis of spent medium revealing several compounds with chemical shifts characteristic of H-phosphinates. (D) ^1H -coupled ^{31}P NMR spectra of concentrated extract. The four largest signals (i–iv) exhibit distinctive splitting patterns and coupling constants consistent with H-phosphinates (i, $J_{\text{H-P}} = 505$ Hz; ii, $J_{\text{H-P}} = 512$ Hz; iii, $J_{\text{H-P}} = 518$ Hz; iv, $J_{\text{H-P}} = 519$ Hz).

is not a known natural product; it has been observed only as a byproduct in *Streptomyces hygroscopicus* strains with mutations in the PTT pathway (44). Based on the content of the *Nonomuraea* sp. NRRL B-24552 *pepM* gene cluster and the results of ^{31}P NMR analyses of spent media (Fig. 5C), we believe it is likely DMPT is further modified to produce additional H-phosphinates in this organism.

Summary. Our successful campaign for phosphonates validates the utility of large-scale genome mining for discovery of new natural products, fulfilling a promise of genomics-enabled drug discovery made 20 y earlier. Including the eight additional molecules we recently reported elsewhere (24, 38, 40, 45) (SI Appendix, Fig. S4), this program has yielded 19 new phosphonate natural products to date. In addition to increasing the number of actinomycete phosphonate gene cluster groups by over eightfold, our efforts doubled the total diversity of phosphonate pathways identified to date. Moreover, rarefaction and extrapolation analyses suggest that the majority of the remaining phosphonate biosynthetic repertoire of Actinobacteria can be found within the dereplicated strain collection (Fig. 6). Accordingly, using our previously published approach (46), we predict that Actinobacteria have a total capacity for ca. 125 distinct pathways for phosphonic acid natural products. The genome mining of 10,000 strains reported here has already identified 78 discrete biosynthetic gene clusters, with multiple producing organisms available for most groups. Lastly, we note that the actinobacterial DNA library and hundreds of genome sequences reported in this study will inform and enable rapid, large-scale genome mining for other natural product classes. We suggest that the future of natural products research and antibiotic development is bright, with many of the best discoveries yet to come.

Materials and Methods

The full description of experimental techniques is provided in SI Appendix.

Construction of the Actinomycete DNA Library. Strains from the US Department of Agriculture ARS Actinobacteria collection were revived from lyophilized vials or slants by inoculation into 150-mm \times 18-mm test tubes containing 5 mL of 172 medium. Cultures were grown at 30 $^\circ\text{C}$ on an angled New Brunswick Scientific roller drum (75 rpm) for 6 d and then arrayed by

aliquoting (2 mL) into 96-well deep blocks. Supernatants were removed after centrifugation, and the remaining cell pellets were frozen at -80 $^\circ\text{C}$ for 24 h. Cell pellets were then allowed to thaw completely at room temperature, and they were immediately refrozen and stored at -20 $^\circ\text{C}$.

DNA was isolated using a 96-well UltraClean-HTP Microbial DNA Kit (MoBio) according to the manufacturer's instructions but with the following modifications. Cell pellets were thawed, resuspended with 300 μL of MicroBead solution containing lysozyme (1 mg/mL) and DNase-free RNase A (100 $\mu\text{g/mL}$), and incubated at 30 $^\circ\text{C}$ for 1 h with gentle rocking. Treated cells were then transferred into the 96-well bead plate. After addition of solution MD1, 96-well blocks were firmly sealed with the provided silicon covers, inverted several times to mix, pulsed in a tabletop centrifuge to draw all liquid to the bottom, and incubated in a 70 $^\circ\text{C}$ water bath for 20 min. After cooling to room temperature, blocks were attached onto a Restech bead mill and cells were lysed by running at setting 20 for 5 min. Blocks were removed, reoriented so that the side closest to the machine body was now furthest away, and shaken again at setting 20 for an additional 5 min. Before DNA elution, spin plates were centrifuged twice to remove residual ethanol and air-dried for 20 min at room temperature. DNA was eluted from spin plates by adding 120 μL of preheated sterile water (70 $^\circ\text{C}$) to wells and waiting 5 min before centrifugation into microplates. DNA stocks were kept at -80 $^\circ\text{C}$ for long-term storage. The completed library contains genomic DNA isolated from 7,488 actinomycetes.

Genetic Screen for *pepM*⁺ Strains. The genomic DNA library was screened by PCR using four pairs of degenerate primers designed to amplify a 406-bp conserved fragment within bacterial *pepM* genes (forward: pepMF, CGCCGGCGTCTGCNTNGARGAYAA; reverse: pepMR, GCGCGCATCATGTGRTTNGCVYA; pepMX, GCGCGCATCATGTGGTTNGCCCADAT; pepMW, GCGCGCATCATGTGGTTNGCRTADAT; pepMZ, GCGCGCATCATGTGTTNCCCADAT). Promega Go Taq was used in 25- μL reactions containing each primer (5 μM), 0.5 μL of DMSO, and 0.5 μL of template DNA. Thermocyclers were operated under the following program: (i) initial denaturation for 3 min at 98 $^\circ\text{C}$; (ii) 30 sec at 98 $^\circ\text{C}$; (iii) 30 sec at 58 $^\circ\text{C}$; (iv) 30 sec at 72 $^\circ\text{C}$; (v) 3 min at 72 $^\circ\text{C}$; and (vi) hold at 12 $^\circ\text{C}$. Steps ii–iv were repeated for a total of 35 cycles. PCR amplification of the *pepM* fragment using the genomic DNA of *Streptomyces viridochromogenes* DSM 40736 (PTT producer), *Streptomyces luridus* NRRL 15101, and *Streptomyces rubellomurinus* American Type Culture Collection (ATCC) 31215 was routinely performed to verify the integrity of reaction master mixes. PCR products were separated by electrophoresis through 1% agarose gels with positive controls on each side. The

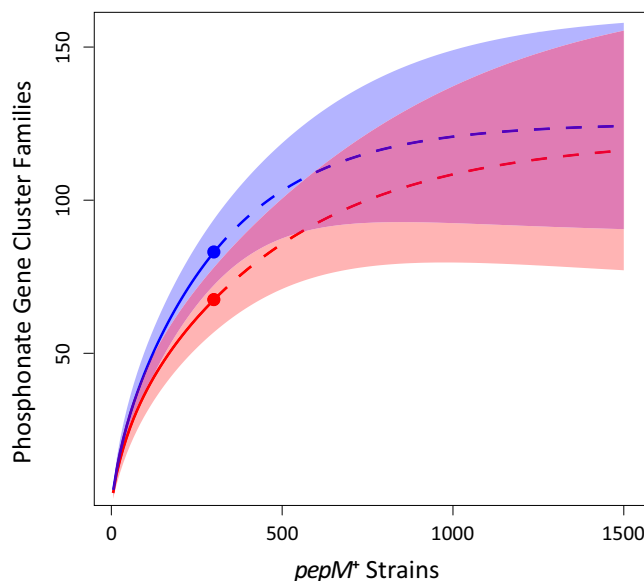


Fig. 6. Extrapolation of unique phosphonate natural product gene cluster families. Calculations were based on amino acid (blue) or gene cluster family (red) data. The current extent of sampling is shown with filled circles. Extrapolation is based on discovery of *pepM*⁺ strains, equivalent to sampling of an additional 40,000 actinomycetes in the same manner as the 10,000 screened for this work. Shaded areas show the 95% confidence intervals for the analyses.

presence of an ~400-bp band matching the size of the positive controls indicated a positive result.

Genome Sequencing and Assembly. *Saccharothrix* ST-888, *Streptomyces lavendulae* Fujisawa 8006, *S. rubellomurinus* subsp. *indigoferus* ATCC 31304, *S. rubellomurinus* ATCC 31215, and all PCR-positive strains from the NRRL collection were revived from freezer stocks on 172 agar plates and regrown as described above. Genomic DNA was isolated using an UltraClean-HTP Microbial DNA kit as described above, except mycelia aggregates were first homogenized in microfuge tubes using sterile pestles. Paired-end, multiplexed sequencing libraries were constructed using Nextera DNA Sample Preparation Kits (Illumina). Samples were sequenced in multiple batches at either the University of Wisconsin Biotechnology Center (Illumina Genome Analyzer IIX) or the University of Illinois at Urbana-Champaign Roy Carver Biotechnology Center (Illumina Hi-Seq 2000 or MiSeq platform). Sequencing reads produced from the Genome Analyzer IIX or the Hi-Seq 2000 with

version 2 chemistry were assembled as previously described (47). ORF prediction was performed with Prodigal version 2.50 (48). The genomes sequenced using the Illumina Hi-Seq 2000 with version 3 chemistry were assembled using IDBA-UD version 1.0.9 (49), and ORF prediction was performed using Prodigal version 2.60. All sequenced strains used in this study are listed in *SI Appendix, Table S1*. DNA sequences have been deposited in GenBank under BioProject PRJNA238534.

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