Marine Viruses Exploit Their Host’s Two-Component Regulatory System in Response to Resource Limitation

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Summary

Phosphorus (P) availability, which often limits productivity in marine ecosystems, shapes the P-acquisition gene content of the marine cyanobacteria Prochlorococcus [1–4] and its viruses (cyanophages) [5, 6]. As in other bacteria, in Prochlorococcus these genes are regulated by the PhoR/PhoB two-component regulatory system that is used to sense and respond to P availability and is typical of signal transduction systems found in diverse organisms [7]. Replication of cyanophage genomes requires a significant amount of P, and therefore these phages could gain a fitness advantage by influencing host P acquisition in P-limited environments. Here we show that the transcription of a phage-encoded high-affinity phosphate-binding protein gene (pstS) and alkaline phosphatase gene (phoA)—both of which have host orthologs—is elevated when the phages are infecting host cells that are P starved, relative to P-replete control cells. We further show that the phage versions of these genes are regulated by the host’s PhoR/PhoB system. This not only extends this fundamental signaling mechanism to viruses but is also the first example of regulation of lytic phage genes by nutrient limitation in the host. As such, it reveals an important new dimension of the intimate coevolution of phage, host, and environment in the world’s oceans.

Results and Discussion

Unicellular cyanobacteria Prochlorococcus and Synechococcus are the dominant photosynthetic organisms in the oceans and they contribute significantly to global primary production [8]. The environmental availability of phosphorus (P), often a limiting nutrient in marine ecosystems, exerts strong selective pressure on Prochlorococcus genomes, which is manifested in the suite of P-acquisition genes they contain [1–4]. This selective pressure is also visible in cyanophages (viruses that infect cyanobacteria); 9 of the 16 sequenced T4-like cyanophages isolated on Prochlorococcus and Synechococcus [9] contain pstS, encoding a periplasmic high-affinity phosphate-binding protein, and 2 contain phoA, an alkaline phosphatase gene. We have suggested previously that these genes of host origin play a role in the acquisition of phosphorus required for phage DNA replication [6]. Consistent with this hypothesis, the frequency of occurrence of pstS gene in phages in the wild was shown to be higher in oceanic sites with lower phosphate [5], as is true of their Prochlorococcus hosts [4, 10].

Both pstS and phoA are upregulated in Prochlorococcus [2] and Synechococcus [11] during P starvation, a signal transduction response regulated by the bacterial phosphate sensing two-component system. We postulated that the expression of the phage versions of these P-acquisition genes (hereafter referred to as “phage pstS” and “phage phoA”) might be regulated by P availability to the hosts—possibly through the PhoR/PhoB two-component regulatory system in the host cells. To address this question, we used a cultured Prochlorococcus strain (NATL2A) and the T4-like cyanophage P-SSM2, which encodes pstS, to measure expression of selected phage and host genes, under P-depleted and P-replete conditions (Figure 1). Transcript levels of the host pstS gene increased within 8 hr in the P-limited host cells (Figure 1B) and rose steadily thereafter, signaling the onset of P starvation, which was manifested as a decrease in the culture growth rate (Figure 1A). Reintroduction of phosphate at 46 hr (arrow, Figure 1B) resulted in a rapid decline of pstS transcripts as the cells were released from P starvation. Similar patterns have been observed in other strains of Prochlorococcus under P starvation [2].

We used this system to examine the effect of host P starvation on infection kinetics and pstS expression level in infecting phage. The cultures were infected 47 hr after resuspension in P-depleted media (shaded interval in Figure 1B) when P starvation was well established in the host cells, as indicated by upregulation of pstS (Figure 1B). Both host and phage pstS transcript levels were then measured over time. Host pstS transcript levels in infected P-starved cells stayed higher than those in the P-replete cultures (Figure S1A available online), as was observed in uninfected hosts (Figure 1B). Transcript levels of phage pstS increased in P-starved hosts relative to the P-replete control (Figures 1C and S1B), whereas transcripts of genes g61 and g20—early and late T4-like phage structural genes [12–14], respectively—did not (Figures 1C and S1B), suggesting differential regulation of pstS and phage structural genes. Addition of phosphate 4 hr after infection (arrow, Figure 1C) resulted in a reduction of both host (similar to Figure 1B) and phage pstS transcripts within 4 hr (Figure 1C), suggesting that both phage and host pstS genes could be regulated by the same mechanism.

phoH encodes an ATP binding protein with unknown function [15] and is considered a phosphate (pho) regulon gene because it is upregulated by P starvation in E. coli [16, 17]. Its expression is not upregulated during P starvation in marine cyanobacteria [2], however, suggesting that it may not play the same role as in E. coli [18]. Nonetheless, because of its prevalence in T4-like cyanophages [9] and association with the pho regulon in E. coli, we examined its expression in our experiments. We found that the expression of phoH in the phage (Figures 1C and S1B) and host (Figure S1A) was not affected by P starvation, and therefore its role in both host and phage remains a mystery.

That the level of P starvation of the host cell selectively influences the degree of upregulation of phage pstS suggests coevolution of regulatory systems between host and phage—probably involving the PhoR/PhoB two-component regulatory system widely used by bacteria including...
cyanobacteria during P starvation [19, 20]. The obvious way to test this hypothesis would be to measure phage \(pstS\) expression in a mutant host without the PhoR/PhoB system. Because there is no genetic system for \textit{Prochlorococcus}, we used a \textit{phoR} knockout mutant of its close relative, \textit{Synechococcus WH8102} [11], and the T4-like cyanophage S-SM1 [9], which contains \(pstS\) (and also \textit{phoA}).

We first established that upon the onset of P starvation in host cells, \(pstS\) was not upregulated in the \textit{phoR} mutant, although it was in the WT cells—simply confirming the results of Tetu et al. [11] (data not shown). Once \(pstS\) was upregulated in the P-starved WT host, we infected P-starved and control cultures of both the mutant and WT strains with phage and measured phage production and the expression levels of a suite of phage genes over the course of infection (Figure 2). Phage production in P-starved cells was reduced relative to the P-replete controls for both the WT and \textit{phoR} mutant (Figures 2A and 2B), confirming that the mutant was indeed P starved and agreeing with our results for \textit{Prochlorococcus} phages (Figure S1F). The \textit{phoR} mutant (doubling time 1.39 ± 0.04 days) grew slower than the WT (1.39 ± 0.04 days) in P-replete conditions, which could explain why phage production was lower in the \textit{phoR} mutant than in the WT host (Figures 2A and 2B). Most importantly, phage \(pstS\) and \textit{phoA} transcripts increased relative to controls in P-starved WT host cells (Figure 2C), as in our \textit{Prochlorococcus} phage/host system (Figure 1C), but they did not in the mutant lacking the PhoR/PhoB regulatory system (Figure 2D). The significance of the expression patterns of genes \(g172\) and \textit{hsp20} in this experiment will become clear below.

The evidence presented thus far is compelling that the transcription of the phage \(pstS\) (and \textit{phoA}) is regulated by the host PhoR/PhoB two-component system. What is the mechanism? Host genes regulated by the PhoR/PhoB system (\textit{pho regulon genes}) have well-conserved \textit{pho} box sequences in their promoters, which bind the transcriptional activator PhoB [16]. \textit{pho} box sequences, which contain conserved
tandem 8 bp sequences 70.8% identical to the predicted host *Synechococcus* WH8102 *pho* box [22]. We purified recombinant *Synechococcus* WH8102 PhoB protein (see Experimental Procedures) and PCR amplified a ~200 bp DNA fragment upstream of *g172* (Figure 3A) containing this putative *pho* box and used it in binding assays. A gel mobility shift assay showed that the recombinant *Synechococcus* WH8102 PhoB protein binds to this DNA fragment (Figure 3C) and that this binding reaction is sequence specific (Figure S2B), suggesting that this putative *pho* box is functional. Phage gene *g172*, which is not found in any host genomes, is between the putative *pho* box and the phage *pstS* gene (Figure 3A). The putative *pho* box sequence in front of *g172* could be formed or gained after *g172* and *pstS* got inserted in the phage genome. Or, this putative *pho* box could be gained together with *pstS* from the host genome and *g172* was inserted between them during gene recombination events, which could destroy the *pho* box. In both scenarios, there must be strong selective pressure to maintain this putative *pho* box in front of *pstS* in the phage, even though it need not be in that location in the host genome. Thus, cyanophages not only carry *pstS* and *phoA* genes, but the expression of these genes is tied into the host PhoR/PhoB phosphate-sensing two-component regulatory system, which regulates the P-starvation response of the host. Although we can’t completely exclude the possibility of indirect regulation of these phage genes by the PhoR/PhoB system, the results of our binding assay strongly favors direct regulation.

Regulation of phage *pstS* and *phoA* genes by P availability to the host also suggests that these phage genes play a role in the acquisition of phosphorus for phage DNA replication during infection. For *Prochlorococcus* (Figure S1F) and *Synechococcus* (Figures 2A and 2B) infecting phages with the *pstS* gene, we found that phage production was reduced by P starvation (Figures S1F, 2A, and 2B), but the timing of the infective cycle was not changed (Figures S1C–S1F). In a related cyanophage/host system, in which the phage does not carry the *pstS* gene, the lytic cycle is lengthened during infection of P-starved host cells [23], consistent with our hypothesis that phage *pstS* expression may augment the P supply to the host cell to facilitate an expedient lytic cycle. Phage-encoded *phoA* may facilitate access to organic P pools, which again would give phage a selective advantage in phosphate-limited environments. Cyanophage S-SM1 gene *g172* may play a role in this process as indicated by the fact that its expression is also induced by P starvation, although its function is still unknown. Because of the lack of genetic tools, we couldn’t knock out these cyanophage genes to see how they affect the lytic cycle, although the selective advantage for cyanophages to have *pstS* gene is supported by the fact that it is enriched in phage genome fragments at oceanic sites with lower phosphate concentrations [5].

Two-component regulatory systems have been found in the three kingdoms of life [7] enabling cells to better acclimate to changing environmental conditions. Although nutrient limitation has been shown to affect the lysis-versus-lysogenization decision of coliphage λ through the host ppGpp level [24, 25], to our knowledge this is the first example in which a lytic virus exploits a host two-component system to be responsive to “environmental conditions” within the host cell, which in turn is responding to nutrient limitation in the external milieu. This extends the selection pressures on the host cell to its infecting viruses, making ever more intimate the coevolution of viruses, hosts, and their environment.
Experimental Procedures

Strains and Growth Conditions
Axenic *Prochlorococcus* NATL2A was grown in 0.2 μm filtered Sargasso seawater-based Pro99 medium [26] amended with 10 mM HEPES (pH 7.5) and 12 mM sodium bicarbonate. Axenic *Synechococcus* WH8102 was grown in SN medium [27] made with seawater from Woods Hole, MA. Kanamycin (25 μg ml⁻¹) was used to maintain the phoR mutant [11]. Cultures were maintained at 21°C under constant cool white light (30 μmol quanta m⁻² s⁻¹).

Prior to infection, the NATL2A culture was harvested by centrifugation (15,000 x g for 10 min), washed twice in either PO₄-replete (Pro99 with PO₄) or -depleted (Pro99 without PO₄) medium, and resuspended in the same medium. WH8102 and the *phoR* mutant cultures were transferred 1:25 to PO₄-replete or -depleted SN medium. Cyanophage P-SSM2 lysate was concentrated by centrifugation (5,000 x g) with Amicon Ultra-15 30K Centrifugal Filter Units (Millipore), washed twice in filtered Sargasso seawater, and resuspended in the same medium. Cyanophage S-SSM1 lysate was concentrated the same way and resuspended in filtered seawater from Woods Hole. Infection was carried on with a multiplicity of infection of 3. Total cell concentration was determined by flow cytometry (Influx, Cytopeia-BD), and phage concentration was determined by the most probable number assay [29].

Quantification of Phage and Host Genomic DNA during Infection
Phage and host genomic DNA was quantified with a quantitative PCR method described previously [29]. In brief, infected *Prochlorococcus* cells were filtered through polycarbonate filters (0.2 μm pore-size) to separate extracellular phage (filtrate), from host cells containing both host and intracellular phage (genomic DNA). The latter were recovered from the filters. The qPCR primers used to measure DNA in the filter and filtrate fractions are listed in Table S1.

RNA Extraction and Transcript Analysis
Samples were collected by centrifugation at 15,000 x g for 15 min at 4°C, and cell pellet was flash frozen in liquid nitrogen and stored at −80°C. Total RNA was then extracted with the Ambion mirVana RNA isolation kit and DNA was removed with the Turbo DNA-free kit (Ambion). Total RNA was reverse-transcribed with the iScript cDNA Synthesis Kit (Bio-Rad). cDNA copies were quantified with a Quantitect SYBR Green PCR Kit (QIAGEN) with M Primers (Table S1) on a LightCycler 480 Real-Time PCR System (Roche Diagnostics). The qPCR primers were designed to amplify only the gene of interest. The specificity of phage (or host) pstS gene primers was confirmed by showing that they don’t amplify the host (or phage) pstS gene (data not shown). The amplification reaction consisted of an initial activation step of 15 min at 95°C, followed 50 cycles of denaturation (95°C, 15 s), annealing (56°C, 30 s), and extension (72°C, 30 s), followed by 5 min at 72°C. Relative transcript abundance was determined by the ΔΔCT method [30]. The host *mbp* gene was used as an internal control for our gene expression study, as shown by the fact that transcription of this gene is stable in various conditions, including phage infection [29] (Figure S3).

Protein Expression and Purification
Primers 8102phoBNcoI (5’-AAAAAACATGCTATGCGGCCTGCGTTGCCG-3’) and 8102phoBHindIII (5’-TTTTTGAATTCGGACGGCGGTTGC-3’) were used to amplify the *Synechococcus* WH8102 phoB gene. PCR was carried out with 0.02 U/μl KOD HiFi DNA polymerase (Novagen) in 120 mM Tris-HCl (pH 8.0), 10 mM KCl, 6 mM (NH₄)₂SO₄, 0.1% Triton X-100, 0.001% BSA, 1 mM MgCl₂, 0.2 mM dNTPs, and 0.4 μM of each primer. PCR cycling conditions consisted of a hot start at 94°C for 5 min, followed by 25 cycles (98°C for 1 s, 50°C for 2 s, and 74°C for 20 s), followed by incubation at 74°C for 7 min. This gene was then cloned into pET30a plasmid (Novagen) with a His-tag on the 5’ end and transformed into E. coli BL21 (DE3) competent cells. A single colony was grown at 37°C overnight in LB medium containing 50 μg/ml Kanamycin, diluted 1:100 with the same medium, and grown at 37°C until OD₆₀₀ = 0.5. Protein expression was induced by adding IPTG to a final concentration of 0.1 mM and cells were grown at 18°C for 24 hr. Cells were harvested by centrifugation at 6,000 x g for 20 min. The cell pellet was suspended with binding buffer (20 mM sodium phosphate, 500 mM NaCl, 20 mM imidazole [pH 7.4]) and disrupted by sonication. The crude lysate was centrifuged at 10,000 x g for 20 min and the supernatant was loaded on a HisTrap FF crude column (GE Healthcare). Protein purification was performed according to the manufacturer’s instructions. Protein was eluted from the column with elution buffer (20 mM sodium phosphate, 500 mM NaCl, 500 mM imidazole [pH 7.4]). Imidazole was removed from the protein solution by centrifugation (5,000 x g, 15 min) with Amicon Ultra-15 10K Centrifugal Filter Units (Millipore), washed twice with 20 mM NaH₂PO₄ (pH 7.4), and resuspended in the same buffer. Although PhoB needs to be phosphorylated by PhoR to become active, the DNA binding ability of purified recombinant PhoB from *E. coli* is comparable to that of phosphorylated PhoB [20].

Electrophoretic Mobility Shift Assay
Primers SSM1F167bp (5’-ATGGAATGTCGACAGTTAATTTAC-3’) and SSM1R160_6FAM (5’-GGAGGACATTCAAGGTC-3’) were used to amplify a 200 bp fragment upstream of phase S-SM1 g172. The reverse primer SSM1R160_6FAM was 5’ labeled with the fluorophore 6FAM to enable in-gel detection with a fluorescence scanner. The same primers without labeling were used to amplify the nonlabeled specific competitor DNA fragment. Nonspecific competitor DNA fragment was amplified with primers SSM1F557bp (5’-TAAACGGATATGAGATTGACAACG-3’) and SSM1F187bp_RC (5’-TTGGATAAATTTGTCAGGAGTCACT-3’). PCR products were purified with QiAquick PCR Purification Kit (QIAGEN). For binding assays without competitor, 3 nM labeled DNA fragment was incubated on ice for 30 min with different amounts of PhoB protein in 10 mM Tris-HCl (pH 7.5), 50 mM KCl, 5 mM MgCl₂, 2.5% glycerol, 5 μg/ml poly-dIdC, 2 μg/ml BSA, and 1 mM DTT in a 20 μl reaction. For assay with competitors, 3 nM labeled DNA fragment and 1 μM PhoB protein were incubated with different amounts of nonlabeled specific competitor or nonspecific competitor. The free DNA and protein-bound complexes were separated on 5% native polyacrylamide gel with 1× TBE buffer (89 mM Tris, 2 mM EDTA, 89 mM Boric acid [pH 8.3]).

Supplemental Information
Supplemental Information includes three figures and one table and can be found with this article online at doi:10.1016/j.cub.2011.11.055.

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References
Supplemental Information

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Inventory of Supplemental Information

Figure S1. Transcript levels and infection kinetics of Prochlorococcus NATL2A and cyanophage P-SSM2 during the P-starvation experiment shown in Figure 1C

Figure S2. Competition assays to verify the binding of PhoB to phage pstS promoter as shown in Figure 3C

Figure S3. rnpB transcript levels of Prochlorococcus NATL2A during infection by cyanophage P-SSM2 as mentioned in Experimental Procedures.

Table S1. Quantitative PCR primers described in Experimental Procedures
**Figure S1.** Transcript levels and infection kinetics of *Prochlorococcus* NATL2A and cyanophage P-SSM2 during the P-starvation experiment shown in Figure 1C. The host was infected with cyanophage P-SSM2 at 47 h after the cells were resuspended in P-replete (control) or P-depleted (-P) media. Host (A) and phage (B) transcript levels were normalized to that of the host *rnpB* gene (see Experimental Procedures). Please note that the y axis of this figure is different from that of Figures 1B and 1C. In the latter we show the transcript levels in cells during P starvation relative to P-replete controls. <d.l. indicates phage transcript levels below detection limit at time 0 h. (C) Host gDNA degradation as a function of time after the onset of infection, measured as the disappearance of the host *rnpB* gene. (D) Host cell numbers over the same interval. (E, F) Phage DNA replication and release measured as the number of intra- and extra-cellular phage *pstS* gene copies (each representing a phage genome). When not visible they are smaller than the data points. Dotted line indicates the beginning of phage burst, and is reproduced on Figure 1C. Error bars indicate the s.d. from two biological replicates.
Figure S2. Competition assays to verify that the binding of PhoB to \textit{pstS} promoter is sequence specific. (A) DNA sequence between cyanophage S-SM1 genes \textit{hsp20} and \textit{g172}. The numbers are the genomic positions. The red TAA is the stop codon of \textit{hsp20} and the green ATG is the start codon of \textit{g172}. The putative \textit{pho} box sequence is shown by blue letters. (B) DNA fragment (with black underline in A) containing the putative \textit{pho} box was labeled by a fluorophore. 3 nM labeled DNA fragment and 1 \( \mu \)M PhoB protein were incubated with different amounts of non-labeled specific competitor or nonspecific competitor. The same DNA fragment without labeling was used as specific competitor (lanes 3-6) and a ~200 bp DNA fragment in phage S-SM1 \textit{hsp20} gene (with orange underline in A) was used as non-specific competitor (lanes 7-10). The first lane shows the pattern of free DNA (no PhoB protein was added in the reaction). Free DNA (F) and the shifted protein-DNA complex (S) were separated in native polyacrylamide gels. The protein-DNA complex almost disappeared in lane 6 and remained unchanged in lane 10, confirming that the binding of PhoB to phage \textit{pstS} promoter is sequence specific.
Figure S3. rnpB transcript levels of *Prochlorococcus* NATL2A during infection by cyanophage P-SSM2. The host rnpB transcript copy number was quantified by qPCR and normalized to the total RNA (ng) used in the reverse transcription reaction. Error bars indicate the s.d. from two biological replicates.
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