More Evidence of Collusion: a New Prophage-Mediated Viral Defense System Encoded by Mycobacteriophage Sbash

Gabrielle M. Gentile,a Katherine S. Wetzel,a Rebekah M. Dedrick,a Matthew T. Montgomery,a Rebecca A. Garlena,a Deborah Jacobs-Sera,a Graham F. Hatfulla

+Department of Biological Sciences, University of Pittsburgh, Pittsburgh, Pennsylvania, USA

ABSTRACT The arms race between bacteria and their bacteriophages profoundly influences microbial evolution. With an estimated $10^{23}$ phage infections occurring per second, there is strong selection for both bacterial survival and phage coevolution for continued propagation. Many phage resistance systems, including restriction-modification systems, clustered regularly interspaced short palindromic repeat-Cas (CRISPR-Cas) systems, a variety of abortive infection systems, and many others that are not yet mechanistically defined, have been described. Temperate bacteriophages are common and form stable lysogens that are immune to superinfection by the same or closely related phages. However, temperate phages collude with their hosts to confer defense against genomically distinct phages, to the mutual benefit of the bacterial host and the prophage. Prophage-mediated viral systems have been described in *Mycobacterium* phages and *Pseudomonas* phages but are predicted to be widespread throughout the microbial world. Here we describe a new viral defense system in which the mycobacteriophage Sbash prophage colludes with its *Mycobacterium smegmatis* host to confer highly specific defense against infection by the unrelated mycobacteriophage Crossroads. Sbash genes 30 and 31 are lysogenically expressed and are necessary and sufficient to confer defense against Crossroads but do not defend against any of the closely related phages grouped in subcluster L2. The mapping of Crossroads defense escape mutants shows that genes 132 and 141 are involved in recognition by the Sbash defense system and are proposed to activate a loss in membrane potential mediated by Sbash gp30 and gp31.

IMPORTANCE Viral infection is an ongoing challenge to bacterial survival, and there is strong selection for development or acquisition of defense systems that promote survival when bacteria are attacked by bacteriophages. Temperate phages play central roles in these dynamics through lysogenic expression of genes that defend against phage attack, including those unrelated to the prophage. Few prophage-mediated viral defense systems have been characterized, but they are likely widespread both in phage genomes and in the prophages integrated in bacterial chromosomes.

KEYWORDS bacteriophage, *Mycobacterium*, viral defense

The bacteriophage population is vast, dynamic, and old (1). The microbial arms race between bacteriophage predators and bacterial prey has promoted evolution of a phage population that is genetically enormously diverse (1), with strong selection for bacterial acquisition of phage resistance mechanisms to defend against viral attack and the need for phage coevolution (2). The phage response to resistance involves both development of antidefense systems (including antiresistance and anticlustered regularly interspaced short palindromic repeat-Cas [CRISPR-Cas] systems) and alteration of host range so that a new spectrum of bacterial strains can be infected (3–6).

Although a plethora of bacterially encoded phage defense systems have been...
described (7–9), recent evidence suggests that temperate bacteriophages collude extensively with their bacterial hosts to provide defense against viral attack (10, 11); these extend beyond well-known homotypic defenses (i.e., against themselves) such as repressor-mediated superinfection immunity to novel heterotypic defense systems (i.e., against genomically different phages), including some with exquisite specificity (11). Such prophage-mediated heterotypic defense systems may be widespread, but few have been described in detail or are mechanistically understood.

A substantial proportion of bacteriophages are temperate, forming stable lysogens in their bacterial hosts (12, 13). Establishment of lysogeny occurs at relatively high frequencies, especially at high bacterial densities and high multiplicities of infection, and a high proportion of sequenced bacterial genomes carry one or more prophages (14–16). Expression of prophage-carried genes during lysogeny can profoundly influence host physiology, endowing pathogenic properties or additional metabolic pathways (15, 17). However, prophage-encoded viral defense systems are capable of providing survival advantages in multiple environments and may thus be common among temperate phages. These systems have been largely overlooked because of the dearth of well-defined collections of phages that infect common bacterial host strains.

Over 10,000 individual phages have been isolated using *Mycobacterium smegmatis* mc²¹⁵⁵ as a host, and over 1,700 have been completely sequenced (https://phagesdb.org) (18). These span a considerable range of genetic diversity and can be assorted into 34 genome types based on shared nucleotide sequences and gene content (29 clusters [A, B, C,…, Z, AA, AB, AC] and five singletons) (19). There is substantial diversity within the clusters, 13 of which can be divided into subclusters (19). Although over one-half of the different types correspond to temperate phages, prophage-mediated defense systems have been described only for phages grouped in cluster N, which collectively have at least five distinct defense systems (11).

Here we describe a new heterotypic prophage-mediated defense system encoded by the cluster I2 mycobacteriophage Sbash. Two Sbash genes, 30 and 31, are required for defense, and the system targets the cluster L2 phage Crossroads with remarkable specificity; it does not defend against any other mycobacteriophage tested, including other closely related subcluster L2 phages. Isolation and characterization of defense escape mutants showed that Crossroads genes 132 and 141 are required for identification of Crossroads as the target of defense. We also show that distantly related homologues of the Sbash 30/31 defense genes present in *Gordonia* phage CarolAnn also defend against Crossroads when expressed in *M. smegmatis*.

**RESULTS**

*Mycobacteriophage Sbash encodes at least two distinct phage defense systems*. Mycobacteriophage Sbash was isolated in Durban, South Africa, has a 55,832-bp genome, and is grouped in cluster I (20). Cluster I has only five member phages (https://phagesdb.org): three in subcluster I1 and two—Sbash and Che9c—in subcluster I2 (21). Sbash and Che9c are temperate but heteroimmune (Fig. 1A and B), sharing extensive nucleotide similarity across the structural genes in the left side of the genome and at the extreme right genome ends (Fig. 1A). The central regions have 7 to 8 shared genes (e.g., Sbash 33 to 42) and several that are present in only one of the two phages (Sbash 28–34; Che9c 28–34) (Fig. 1A). It was shown previously that genes centrally located in cluster N mycobacteriophage genomes are lysogenically expressed and confer defense against other mycobacteriophages (11). We therefore constructed Che9c and Sbash lysogens and tested them for susceptibility to infection by the use of a panel of genomically diverse phages that infect *M. smegmatis* mc²¹⁵⁵.

A set of 108 phages were serially diluted and plated onto lawns of *M. smegmatis* mc²¹⁵⁵ and lysogens of Sbash and Che9c and the efficiencies of plating (EOP) determined (Fig. 1B). Most of the phages showed little or no difference in EOP, but substantial reductions were observed for several phages on each of the lysogens. These included Crossroads, Giles, and Typha on the Sbash lysogen and phages Euphoria, Llama, Wee, Charlie, Bunnies, Jebeks, and Phayonce on the Che9c lysogen (Fig. 1B); the
FIG 1 Viral defense systems in mycobacteriophages Sbash and Che9c. (A) The genome maps of phages Che9c and Sbash are shown with shared nucleotide sequence similarity represented as spectrum-colored shading between the genomes, with violet representing the most similar and red the least similar above a BLASTN E value threshold of 10^-4. Genes are shown as colored boxes above and below genome rulers. An expanded view of the centers of the Che9c and Sbash-Mediated Phage Defense.

B

| Phage (cluster) | Che9c | Sbash |
|-----------------|-------|-------|
| Bethlehem (A1)  | 1/1/1 | 1/1/1 |
| BlekhKnuckles (A1) | 1/1/1 | 1/1/1 |
| Bbx1 (A1)       | 1/1/1 | 1/1/1 |
| Euphoria (A1)   | 1/1/1 | 1/1/1 |
| MrGordo (A1)    | 1/1/1 | 1/1/1 |
| El28Brutus (A11) | 1/1/1 | 1/1/1 |
| Che12 (A2)      | 1/1/1 | 1/1/1 |
| D29 (A2)        | 1/1/1 | 1/1/1 |
| Trie (A2)       | 1/1/1 | 1/1/1 |
| Penny1 (A3)     | 1/1/1 | 1/1/1 |
| Peaches (A4)    | 1/1/1 | 1/1/1 |
| Skippet (A4)    | 1/1/1 | 1/1/1 |
| Gladiator (A6)  | 1/1/1 | 1/1/1 |
| Alma (A9)       | 1/1/1 | 1/1/1 |
| Fang (B1)       | 1/1/1 | 1/1/1 |
| Orion (B1)      | 1/1/1 | 1/1/1 |
| PG1 (B1)        | 1/1/1 | 1/1/1 |
| Rosebush (B2)   | 1/1/1 | 1/1/1 |
| Chandler (B3)   | 1/1/1 | 1/1/1 |
| Bx21 (C1)       | 1/1/1 | 1/1/1 |
| SirHarley (D1)  | 1/1/1 | 1/1/1 |
| Hawkeye (D2)    | 1/1/1 | 1/1/1 |
| Eureka (E)      | 1/1/1 | 1/1/1 |
| Lilac (E)       | 1/1/1 | 1/1/1 |
| Phaux (E)       | 1/1/1 | 1/1/1 |
| Pumpkin (E)     | 1/1/1 | 1/1/1 |
| Toto (E)        | 1/1/1 | 1/1/1 |
| Oomer (F1)      | 1/1/1 | 1/1/1 |
| Brocalys (F1)   | 1/1/1 | 1/1/1 |
| Buzz2Lysyeay (F1) | 1/1/1 | 1/1/1 |
| Che8 (F1)       | 1/1/1 | 1/1/1 |
| Daeneray (F1)   | 1/1/1 | 1/1/1 |
| Dante (F1)      | 1/1/1 | 1/1/1 |
| DeadP (F1)      | 1/1/1 | 1/1/1 |
| D Lane (F1)     | 1/1/1 | 1/1/1 |
| Firehouse51 (F1)| 1/1/1 | 1/1/1 |

| Phage (cluster) | Che9c | Sbash |
|-----------------|-------|-------|
| Fruitloop (F1)  | 1/1/1 | 1/1/1 |
| Gumbie (F1)     | 1/1/1 | 1/1/1 |
| Ishubes (F1)    | 1/1/1 | 1/1/1 |
| Liana (F1)      | 1/1/1 | 1/1/1 |
| Piply (F1)      | 1/1/1 | 1/1/1 |
| Ramsey (F1)     | 1/1/1 | 1/1/1 |
| SG4 (F1)        | 1/1/1 | 1/1/1 |
| SweetyN (F1)    | 1/1/1 | 1/1/1 |
| SweetyO (F1)    | 1/1/1 | 1/1/1 |
| Wee (F1)        | 1/1/1 | 1/1/1 |
| XFactor (F1)    | 1/1/1 | 1/1/1 |
| Avan (F2)       | 1/1/1 | 1/1/1 |
| Zapper (F2)     | 1/1/1 | 1/1/1 |
| Squirty (F3)    | 1/1/1 | 1/1/1 |
| TChen (F4)      | 1/1/1 | 1/1/1 |
| BPs (G1)        | 1/1/1 | 1/1/1 |
| Halo (G1)       | 1/1/1 | 1/1/1 |
| Hope (G1)       | 1/1/1 | 1/1/1 |
| Liefie (G1)     | 1/1/1 | 1/1/1 |
| Sneeze (G1)     | 1/1/1 | 1/1/1 |
| Damien (H1)     | 1/1/1 | 1/1/1 |
| Konstantine (H1)| 1/1/1 | 1/1/1 |
| Predor (H1)     | 1/1/1 | 1/1/1 |
| Babioki (I1)    | 1/1/1 | 1/1/1 |
| Brujia (I1)     | 1/1/1 | 1/1/1 |
| Island3 (I1)    | 1/1/1 | 1/1/1 |
| Che9c (I2)      | 1/1/1 | 1/1/1 |
| Sbash (I2)      | 1/1/1 | 1/1/1 |
| Omega (J)       | 1/1/1 | 1/1/1 |
| Acesphaia (K1)  | 1/1/1 | 1/1/1 |
| Zeej (K2)       | 1/1/1 | 1/1/1 |
| ShedlockHoimes (K3) | 1/1/1 | 1/1/1 |
| Cheetobro (K4)  | 1/1/1 | 1/1/1 |
| FlorenBharron (K4) | 1/1/1 | 1/1/1 |
| Taquito (K4)    | 1/1/1 | 1/1/1 |
| Wintermute (K4) | 1/1/1 | 1/1/1 |

C

Typha
Crossroads
Sbash
Giles
Pipsqueaks

mc²155 mc²155p(Sbash) mc²155pGG07

(Continued on next page)
Sbash and Che9c lysogens showed immunity to themselves but not to each other, and the cluster I phages Brujita and Island3 showed reduced plating on Che9c, probably reflecting homoimmunity. In general, the Sbash and Che9c profiles are nonoverlapping, suggesting the presence of distinct defense systems in the two prophages (Fig. 1B). To determine whether the Sbash defense against Crossroads, Giles, and Typha is repressor mediated, we tested whether these phages can infect a recombinant strain (mc2155pGG07) expressing the Sbash repressor (gp43; Fig. 1C). All three phages infect mc2155pGG07 efficiently, and the Sbash prophage-mediated defense is thus distinct from repressor-mediated superinfection immunity. Here we further characterized the Sbash defenses; Che9c defenses are to be described elsewhere.

**Identification of Sbash genes conferring viral defense.** To determine which Sbash genes are lysogenically expressed, RNA was isolated from a Sbash lysogen and analyzed by transcriptome sequencing (RNAseq) and the sequence reads were mapped to the Sbash genome (Fig. 2A; see also Fig. S1 in the supplemental material). Expression of five groups of genes was observed near the center of the viral genome, positioned near the ends of the integrated prophage (Fig. 2A). These included the repressor and integrase genes (genes 40 to 43) adjacent to attL, two operons of leftward-transcribed genes (genes 37 to 39 and 32 to 34), and two rightward-transcribed operons (genes 30, 31, and 35 to 36; Fig. 2A). Most of these genes are of unknown function, with the exceptions being gene 35, which encodes a homologue of the Esx-A component EspB (22), gene 39, which codes for a predicted membrane-localized protein, and gene 42, which codes for an InrE-like protein (Fig. 2A).

To determine which of these genes are responsible for viral defense, we used Bacteriophage Recombineering of Electroporated DNA (BRED) mutagenesis (23) to construct deletion mutants of genes 30, 31, 34, 35, 37, 38, and 39 (Fig. 2B). All of these mutants grow lytically and form stable lysogens in *M. smegmatis*. Lysogenic strains of the mutants were recovered and tested for phage susceptibility (Fig. 2C). Five of the mutant derivatives (mutants Δ34, Δ35, Δ37, Δ38, Δ39) had susceptibility profiles similar to that of wild-type Sbash for all the tested phages (Fig. 2C), but both the Δ30 and Δ31 mutant lysogens lost defense against Crossroads while maintaining defense against Giles (Fig. 2C). Sbash 30 and 31 thus are required for defense against Crossroads but not for defense against Giles. All of the deletion mutants tested retained defense against Giles, and Sbash genes 32, 33, 36, 41, and 42 remain plausible candidates for Giles defense genes.

To test whether Sbash genes 30 and 31 are sufficient for Crossroads defense, we constructed a series of recombinant plasmids containing subsets of Sbash genes (Fig. 3A). A plasmid carrying Sbash 30 and 31 together with the upstream expression signals (pGG05) conferred a level of Crossroads defense similar to that of a Sbash lysogen, and both gene 30 and gene 31 were required; plasmids pGG10 and pGG24 carrying gene 30 and gene 31 alone, respectively, did not confer defense (Fig. 3B). Thus, Sbash genes 30 and 31 are necessary and sufficient for defense against Crossroads infection; they do not defend against Typha (data not shown). None of the plasmids conferred defense against Giles, perhaps either because noncontiguous genes are required for defense or because the defense genes are not efficiently expressed in the recombinant strains. Although the genetic basis for defense against Giles infection

**FIG 1 Legend (Continued)**

Sbash genomes with putative gene functions indicated is shown at the bottom. Gene numbers are shown within the gene boxes, and the assigned Pham numbers are shown above the boxes, with the number of Pham members indicated in parentheses. (B) Plating efficiencies of mycobacteriophages on lysogens of phages Che9c and Sbash. A total of 108 mycobacteriophages were serially diluted and plated onto lawns of *M. smegmatis* mc²155, mc²155(Che9c), and mc²155(Sbash). The plating efficiencies on each of the lysogens are shown relative to plating on the nonlysogen. The cluster assignment is shown in parentheses following each phage name, and phages are sorted according to their cluster designation; Giles and Crossroads are highlighted in red type. Plating efficiencies between 1 and 10⁻², between 10⁻² and 10⁻¹, and at 10⁻¹ or below are highlighted in green, yellow, and red, respectively. (C) Sbash-mediated defense against Typha, Crossroads and Giles is not repressor mediated. Ten-fold serial dilutions of phages Typha, Giles, Crossroads, Sbash, and Pipsqueaks were plated onto lawns of mc²155, mc²155(Sbash), and mc²155pGG07, which carries the Sbash repressor (see Fig. 3). Like strain mc²155(Sbash), strain mc²155pGG07 confers immunity to Sbash but does not inhibit infection by Typha, Crossroads or Giles. Phage Pipsqueaks is a control that is not affected by Sbash defenses. The pGH1000 vector control is shown in Fig. 3.
remains unclear, there is clearly a second viral defense system distinct from that corresponding to Sbash 30 and 31.

**Sbash prophage-mediated defense specificity for phage Crossroads.** In the initial screening for defense patterns (Fig. 1B), none of the other cluster L phages tested (Rumpelstiltskin [L2]; LeBron [L1]; Whirlwind [L3]; DyoEdafos [L4]) were subject to Sbash defense.
Crossroads and Rumpelstiltskin are closely related to each other, but Rumpelstiltskin has a deletion of ~5 kbp relative to Crossroads near its right end (Fig. 4A and B). We tested seven additional subcluster L2 phages (Kahlid, LilDestine, Wilder, Gardann, MkaliMitinis3, Faith1, and BigCheese) for infection of a Sbash lysogen to further define the defense specificity (Fig. 4C). Somewhat surprisingly, none of these other subcluster L2 phages had reduced EOP on the Sbash lysogen (Fig. 4C), and the Sbash prophage targeted Crossroads with exquisite specificity.

Crossroads genes 132 and 141 are involved in Sbash-mediated defense targeting. To understand how Crossroads is targeted for defense, we isolated a series of
defense escape mutants (DEMts) that overcome Sbash gp30/gp31-mediated defense (Fig. 5A). A total of 24 mutants were isolated that escape defense by Sbash or by Sbash genes 30 and 31 (i.e., using plasmid pGG05), and the phage DNA was purified and sequenced (Table 1). Sequencing of the parental phage stock from which the lysates were prepared for mutant isolation revealed a 5,004-bp deletion (Δ38089–43093) that removed all or parts of genes 49 to 58, which are evidently dispensable for lytic growth and are not involved in Sbash-targeting (Table 1); all of the mutants carried this deletion. The parental phage with the deletion was shown to be fully susceptible to Sbash defense. Relative to this parent phage, all 24 mutants contained single mutations that included small deletions or insertions and single nucleotide changes (Table 1) (Fig. 5B and C). Two pairs of mutants with the same mutations were isolated from the same lysates on the Sbash lysogen and the pGG05 recombinant (pgg344/pgg400 and pgg354/pgg408) and are presumably siblings (Table 1).
Characterization of Crossroads defense escape mutants. (A) Plating of Crossroads defense escape mutants (DEMs) on host strains. Crossroads mutants were isolated as escapees on lawns of either a Sbash lysogen or a pGG05 recombinant strain, purified, and shown to escape Sbash prophage defense. Ten-fold serial dilutions of DEMs (phgg409, phgg411, phgg399, phgg407) were plated on bacterial lawns as indicated.

(B) Organization at the extreme right end of the Crossroads genome showing genes 121 to 145. The genome is displayed as described for Fig. 1.

(C) Mapping of Crossroads DEM mutations. Defense escape mutants of Crossroads were purified and sequenced, and the locations of the mutations were mapped to gene 132 and 141 and the 141 to 142 intergenic region.

(D) DEM mapping in the Crossroads 141-to-142 intergenic region. The DNA sequence is shown for the beginning of the leftward-transcribed gene 141 and its amino acid sequence, together with the sequences immediately upstream. The ATG translation start codon for gene 141 is underlined, and the ribosome binding site (RBS) is indicated. Four DEM mutations are

(Continued on next page)
Six of the DEM mutants mapped in Crossroads gene 132, corresponding to four different mutations (Fig. 5C), two of which were single base insertions introducing frameshift mutations; one was a 9-bp duplication (dup) resulting in a three-residue insertion, and one introduced a single amino acid substitution (R185S) (Table 1) (Fig. 5C). The two frameshift mutations likely led to loss of function and the inability to be recognized by the Sbash gp30/31 system. Crossroads gene 132 was well conserved, and closely related homologues were present in 41 of the 44 cluster L phages. However, these (including Crossroads) contained a second, more distantly related homologue (e.g., Crossroads 140; Fig. 5B), and Crossroads gp140 shared 32% amino acid identity with Crossroads gp132 (Fig. 5B). It is likely that neither is needed for lytic growth, as one cluster L phage (Bromden) has lost a large genome segment containing these genes. However, there are few bioinformatic clues as to their specific functions. Interestingly, several subcluster L2 phages had identical (e.g., Wilder, LilDestine, and Kahlid) or nearly identical (e.g., Faith1 and Gardann; one amino acid difference) homologues of Crossroads 132 but were not subject to Sbash defense (Fig. 4B and C).

To examine the role of Crossroads 132, we used BRED engineering to construct a Crossroads deletion mutant (Crossroads Δ132). The Δ132 mutant grew well and formed plaques that were similar in size to those seen with wild-type Crossroads, demonstrating that 132 is not required for lytic growth (Fig. 5E). The Δ132 mutant plated with an EOP value of 1 on both the Sbash lysogen and a recombinant strain carrying pGG05, showing that 132 was required for Sbash 30/31 targeting (Fig. 5E). Expression of Crossroads 132 from a recombinant strain carrying plasmid pKSW06 (see Table S2 in the supplemental material) at least partially restored Sbash-mediated defense of the Cross-

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**TABLE 1** Defense escape mutants of phage Crossroads

| Mutant name | Defender | Length (bp) | Mutation | Gene (substitution) |
|-------------|----------|-------------|----------|---------------------|
| Crossroads-wt | 76,129 | | | |
| Crossroads-par | 71,124 | Δ38089–43093 | Δ49–58 | |
| phgg344 | pGG05 | 71,124 | G68548T | gp132 R185S |
| phgg348 | pGG05 | 71,025 | 74303Δ99 bp | 141–142 intergenic |
| phgg350 | pGG05 | 71,124 | A73608T | gp141 V214E |
| phgg354 | pGG05 | 71,123 | 68435 9 bp (dup) | gp132 222 Δ HDK |
| phgg356 | pGG05 | 71,125 | 68575 1 bp | gp132 175f/s |
| phgg359 | pGG05 | 71,118 | Δ73760–73765 (Δ6 bp) | gp141 ΔE161, ΔT162 |
| phgg364 | pGG05 | 71,124 | Δ73760–73765 (Δ6 bp) | gp141 ΔE161, ΔT162 |
| phgg398 | Sbash | 71,125 | 68395 1 bp | gp132 236f/s |
| phgg399 | Sbash | 71,124 | G68231T | gp141 A338D |
| phgg400 | Sbash | 71,124 | G68548T | gp132 R185S |
| phgg403 | Sbash | 71,124 | G73236T | gp141 A338D |
| phgg407 | Sbash | 71,124 | G73275T | gp141 A325E |
| phgg408 | Sbash | 71,133 | 68435 9 bp (dup) | gp132 222 Δ HDK |
| phgg409 | Sbash | 71,124 | C73681A | gp141 V190F |
| phgg411 | Sbash | 71,124 | G73317T | gp141 A311E |
| phgg413 | Sbash | 71,133 | 74335 9 bp (dup) | 141–142 intergenic |
| phgg417 | Sbash | 71,124 | T74303C | 141–142 intergenic |
| phgg419 | Sbash | 71,150 | 74312 26 bp (dup) | 141–142 intergenic |
| 1_1A | Sbash | 71,124 | G73437A | gp141 S271F |
| 3_4A | Sbash | 71,118 | Δ73760–73765 (Δ6 bp) | gp141 ΔE161, ΔT162 |
| 3_3A | Sbash | 71,124 | A73009G | gp141 *413Q |
| 1_4A | Sbash | 71,118 | Δ73760–73765 (Δ6 bp) | gp141 ΔE161, ΔT162 |
| 1_3A | Sbash | 71,118 | Δ73760–73765 (Δ6 bp) | gp141 ΔE161, ΔT162 |
| 1_2A | Sbash | 71,124 | G73101T | gp141 A383E |

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**FIG 5** Legend (Continued)

shown: a single base substitution at coordinate 74,303, 26-bp and 9-bp duplications, and a 99-bp deletion. (E) Crossroads 132 or 129 to 141 is required for targeting by Sbash 30/31. Serial dilutions of a Crossroads mutant in which gene 132 is deleted (C’roadsΔ132) or genes 139 through 141 are deleted (C’roadsΔ139–141) together with Crossroads (C’roads), Sbash, and Pipsqueaks were plated onto lawns of M. smegmatis mc2155, a Sbash lysogen strain, and a pGG05 recombinant strain.
roads Δ132 mutant and DEMs phgg356 and phgg408 (Fig. 6A), confirming that the escape phenotype resulted from loss of gp132 function.

Fourteen of the mutants mapped in Crossroads gene 141, whose specific function is unknown but is related to AAA ATPases (Walker A box residues at 191 to 198; Fig. 5B). The mutants correspond to nine different mutations, one of which is a 6-bp deletion removing codons 161 and 162 and was isolated independently five times (Table 1) (Fig. 5B). Oddly, one mutation is within the termination codon and a single nucleotide change introduces a glutamine codon, enabling readthrough and inclusion of 28 additional C-terminal amino acids. Seven mutations introduced single amino acid substitutions, primarily within the C-terminal half of the predicted gp141 protein (Fig. 5B). The remaining four mutants mapped to the 370-bp 141-to-142 intergenic region (Fig. 5D) and included a 99-bp deletion, duplications of 9 bp or 26 bp, and a single nucleotide substitution at coordinate 74303 (Fig. 3B). All of these changes were upstream of the 141 start codon (comparative analysis and coding potential strongly support the use of this start codon), and these mutations presumably influence 141 expression rather than gp141 activity per se.

Closely related homologues of Crossroads 141 were present in 37 of the 44 sequenced cluster L phages, but their absence from seven of the phages, including the subcluster L2 phage BigCheese (Fig. 4A and B), indicated that it is not required for lytic growth. Interestingly, none of the 141 DEM mutants have large deletions or frameshift mutations within the open reading frame, and none of the single amino acid substitutions are in the Walker A motif or other known catalytic residues (Fig. 5C). Six of the substitutions were in residues that are highly conserved among the mycobacteriophage homologues, four of which are conserved alanines (A311E, A325E, A338D, and A383E). Attempts to knock out 141 from Crossroads were unsuccessful, suggesting that although it is not required for lytic growth, eliminating it resulted in a nonviable phenotype. Thus, the 141 DEM phenotypes likely resulted from reductions in expression or activity rather than from complete loss of function. Phage BigCheese, which lacks 141, has also lost genes 139 and 140 (Fig. 4B), suggesting that loss of one or both of these may rescue the nonviable phenotype of a 141 null mutant. We thus attempted to knock out 140 and 141 together, as well as 139 through 141. We were unable to recover a knockout of 140 and 141, but a knockout of 139 to 141 (mimicking the BigCheese genome arrangement; Fig. 4B) was readily obtained. Therefore, viable inactivation of 141 requires loss of 139 and possibly of both 139 and 140. We then tested the ability of the Crossroads Δ139–141 mutant to escape Sbash 30-to-31-mediated defense. The Crossroads Δ139–141 mutant grew to a high titer, formed plaques that were similar in size to those seen with wild-type Crossroads, and plated with an EOP value of 1 on both a Sbash lysogen and a pGG05 recombinant strain (Fig. 5E). Taken together, these observations are consistent with the interpretation that escape from Sbash can be achieved by mutational reduction of gp141 activity or by altering its expression by mutation in the 141-to-142 intergenic region. Escape from defense can also occur by loss of gp141 but requires accompanying loss of gp139 and gp140.

Attempts at complementation of the 141 DEM defect are complicated by the toxicity of Crossroads 141 when expressed in M. smegmatis, even in the absence of Sbash (Fig. 6B and C). However, by using very small amounts (5 ng/ml) of anhydrotetracycline (Atc) and extended incubation times (Fig. 6B), we could show that 141 expression (from plasmid pGG38, which contains an Atc-inducible 141 gene) in a Sbash lysogen substantially restored the reduced EOP of DEMs phgg348, phgg364, and phgg399 (Fig. 6B), consistent with the interpretation that reduced expression or activity of Crossroads gp141 is directly responsible for the escape phenotype.

Only some DEM mutants of Crossroads 141 are toxic in M. smegmatis. Expression of Crossroads gp132 in M. smegmatis was well tolerated, using induction of expression with 100 ng/ml Atc (Fig. 6C). Furthermore, expression of gp132 in a Sbash lysogen was also well tolerated (Fig. 6C). The Sbash 30/31 defense thus seems distinct from those abortive infection (abi) mechanisms that involve toxin-antitoxin-mediated
FIG 6 Expression of Crossroads gp132 and gp141. (A) Strains were constructed carrying extrachromosomal expression vector pCCK38 or plasmid pKSW06, in which Crossroads 132 expression can be induced by addition of ATc, in either M. smegmatis mc2155 (top row) or a Sbash lysogen (bottom row). Lawns were prepared on solid media with (Induced) or without (Uninduced) ATc, and 10-fold serial dilutions (from left to right) of phages were spotted. Phages tested were Crossroads, Crossroads Δ 132, DEM mutants phgg356 and phgg408, Sbash, and Pipsqueaks. Expression of gp132 in mc2155(Sbash) results in an approximately 100-fold reduction in EOP of strain Δ 132 and the DEMs relative to the uninduced control. It is unclear why some of these phages have reduced plating efficiencies on M. smegmatis (Sbash) pCCK38 relative to M. smegmatis mc2155pCCK38. (B) Strains containing integration-proficient vector pCCK39 or plasmid pGG38 in which Crossroads 141 can be induced with ATc were plated as described in the panel A legend, except the ATc concentration was reduced to 5 ng/ml and the plates were incubated for 3 days (all others were incubated for 1.5 days). Phages Crossroads, phgg348, phgg364, phgg399, Sbash, and Pipsqueaks were spotted as described in the panel A legend. Expression of gene 141 substantially reduced the plating efficiencies of DEM mutants in a Sbash lysogen. (C) Liquid cultures of strains containing the plasmids indicated on the left were grown to an OD600 of 0.5, serially diluted 10-fold, and spotted onto solid media with (Induced) or without (Uninduced) ATc. The genes/mutants in each plasmid are noted at the right. Plasmids pGG37 and pKSW06 are integrating and extrachromosomal plasmids, respectively, containing Crossroads 132, and neither displayed toxicity in either a Sbash lysogen or a nonlysogen.
cell death (24, 25), although Crossroads gp132 alone could be insufficient to activate growth inhibition. Expression of Crossroads gp141 in nonlysogenic M. smegmatis was highly toxic, and growth was strongly inhibited (Fig. 6C), even in the absence of the Sbash prophage. Interestingly, in testing five different DEM mutants with mutations within 141, two (Δ161 and Δ162; V190F) exhibited substantial toxicity—albeit milder than that seen with wild-type 141—and three (A338D, A383E, and S271F) showed little or no toxicity at all. We also tested expression of the Crossroads 141 homologues in phages Wilder (140) and LilDestine (136), both of which were found to be toxic when expressed (Fig. 6C). Taken together, these observations suggest that the toxicity of Crossroads gp141 is unrelated to its role in targeting Crossroads for defense by the Sbash system.

**RNAseq profiles of Crossroads and DEM mutants.** To examine the impact of the DEM mutations mapping upstream of 141 on gene expression, we analyzed the RNAseq profiles of wild-type Crossroads and DEMs phgg348, phgg413, phgg417 and phgg419 (Table 1) (Fig. 7A), at 30 min and 150 min after infection, which we refer to here as “early” and “late” time points, respectively. In wild-type Crossroads, many of the leftward-transcribed genes at the right end of the genome (genes 122 to 144) were expressed early, together with gene 36 encoding a Lsr2-like protein (Fig. 7A). Rightward early transcription starts just upstream of gene 41 and transcribes genes 41 to 48 and also occurs at a location near gene 58 to transcribe genes 59 to 118. At the late time point, early transcripts were still detected, but rightward transcription proceeded from upstream of gene 59 all the way through to the right genome end; the virion structure and assembly genes, especially genes 6 to 16, were highly expressed also (Fig. 7A). It is unusual that transcripts were present for both strands of genes 122 to 144 at the late time point (Fig. 7A).

Transcription of gene 141 initiated in the 141-to-142 intergenic region (Fig. 7B), and the deletion mutant (phgg348) and both insertion mutants (phgg413 and phgg419) (Table 1) (Fig. 5C) showed strongly reduced transcription of gene 141 (Fig. 7B), particularly in mutants phgg348 and phgg413, consistent with mutational interruption of the promoter (Fig. 7B); there was also an apparent modest reduction in expression of genes 125 to 138 (Fig. 7). Transcription of 141 was not substantially reduced in the phgg417 point mutant (Fig. 7B), and it is plausible that the DEM phenotype resulted from inefficient translation of gp141. However, the mutation was 55 bp upstream of the predicted translation start site (Fig. 5C) and thus beyond the normal location of a ribosome binding site (RBS).

**Mechanism of Sbash 30-to-31-mediated defense against Crossroads.** To explore how Sbash 30/31 defends against Crossroads infection, we first used RNAseq to determine if Sbash 30 to 31 inhibits an early stage of infection or after DNA injection occurs and gene expression begins. When Crossroads infected a strain expressing Sbash genes 30 and 31 (Fig. 8A), the early expression pattern was similar to that seen with Crossroads transcription following infection of wild-type M. smegmatis (Fig. 7A). Thus, there was no indication of an early block to adsorption, DNA injection, or early transcription (Fig. 8A). However, at the late time point, there was a notable reduction in late gene expression, reflected in the number of RNAseq reads mapping to the virion structure and assembly genes (1 to 32) relative to transcripts present at that time but expressed early (e.g., genes 59 to 118). In infection of wild-type cells, the ratio of Crossroads late transcripts to early transcripts at the late time point was 3.88 (Fig. 7A), but the late transcript/early transcript ratio in the presence of Sbash 30/31 at the late time point was only 0.61 (Fig. 8A). DEM mutants phgg348 and phgg417 had late transcript/early transcript ratios similar to those seen with the wild type at late infection of wild-type M. smegmatis (4.82 and 3.02, respectively), but in a Sbash 30/31 host they at least partially recovered (late transcript/early transcript ratios of 1.75 and 1.71, respectively) from the defect seen in wild-type Crossroads (Fig. 8A).

Sbash genes 30 and 31 share features with the genes encoding the RexAB system of phage lambda (Fig. 8B); Sbash gp31 and RexB both have four predicted transmem-
RNAseq analysis of Crossroads and DEM mutants. (A) RNA was isolated at early (30 min) and late (150 min) time points after infection of M. smegmatis with wild-type Crossroads (green) or DEMs phgg348 (aqua) and phgg417 (red); reads mapping to forward and reverse strands are shown as indicated. Arrows at the bottom indicate the early and late transcribed regions, with gene numbers denoting the positions. (B) Expanded view of RNAseq reads mapping to the reverse strands at the right end of the genome. DEM mutants phgg419 (blue) and phgg413 (magenta) are shown in addition to those shown in panel A and are similarly colored. Arrows indicate leftward transcription of the two genes, 132 and 141, in which DEM mutants mapped. Reads are shown for the reverse strand only.
YSIS OF Sbash 30-TO-31-MEDIATED DEFENSE AGAINST CROSROADS. (A) RNAseq analysis of CROSROADS AND DEM MUTANT INFECTION OF M. smegmatis mc2155pGG05. RNA was isolated at early (30 min) and late (150 min) time points after infection of M. smegmatis mc2155pGG05 with wild-type CROSROADS (green) or DEMs phgg417 (red) and phgg348 (aqua); reads mapping to forward and reverse strands are shown as indicated. Scales were adjusted to reflect approximately similar levels of early gene expression. (B) COMPARISON OF Sbash AND LAMBDA GENOMES. SEGMENTS OF THE TWO PHAGE GENOMES WERE ALIGNED TO SHOW SIMILARITIES BETWEEN Sbash DEFENSE GENES 30 AND 31 AND LAMBDA EXCLUSION GENES rexA AND rexB. THE POSITIONS OF PROMOTERS EXPRESSING Sbash 30/31 AND LAMBDA cI/rexA ARE SHOWN, AND THE LOCATIONS OF A DUF4747 CONSERVED DOMAIN IN Sbash gp30 AND LAMBDA RexA ARE ALSO INDICATED. THE FUNCTION OF Sbash gp29 (CONTINUED ON NEXT PAGE)
brane domains (Fig. 8C), and Sbash gp30 and lambda RexA both contain a DUF4747 motif (Fig. 8B). RexAB excludes infection of T4rlII mutants and is thought to do so by RexA-mediated activation of the ion channel potential of RexB (26, 27). The Sbash system may act similarly, such that when activated, Sbash gp31 mediates membrane depolarization, therefore lowering intracellular ATP concentrations and causing interruption of macromolecular synthesis and abortion of phage growth (Fig. 8D). The RNAseq profiles are consistent with this model, and we similarly observed a reduction in phage DNA amplification. In Crossroads infection of *M. smegmatis*, ~90% of DNA sequencing reads mapped to the Crossroads genome, whereas in infection of a Sbash 30-to-31-expressing strain, fewer than 5% of reads mapped to Crossroads. The interruption of both phage DNA replication and late transcription is consistent with a reduction in intracellular ATP concentration.

The model proposes that the membrane-located Sbash gp31 acts as an ion channel when stimulated by Sbash gp30, which in turn is activated by Crossroads infection. The predicted gp31 topology suggests there are N- and C-terminal segments that are cytoplasmically located together with a region between the second and third membrane-spanning domains (Fig. 8C; see also Fig. S2A). Database searches identified four Sbash gp31 homologues and showed that although few residues are highly conserved, there are eight invariant positions in the putative central cytoplasmic region (Fig. S2A). We constructed two separate mutants (R70A and D73A) with substitutions in this region (Fig. 8C); both were defective in defense (Fig. S2B). This suggests that this is a functionally important part of gp31, and we propose that it is involved in Sbash gp30 interactions.

The *Gordonia* phage CarolAnn gp44/gp43 defense system also targets Crossroads. There are no homologues of Sbash genes 30 and 31 elsewhere in the extant collection of mycobacteriophage genomes, but there is a pair of related genes in *Gordonia* phage CarolAnn (see accompanying paper [33]) (Fig. 9A). CarolAnn gp44 and gp43 share 50% and 42% amino acid identity with Sbash gp30 and gp31, respectively, and confer defense against infection of several heterotypic *Gordonia* phages, including Kita and Nymphadora. However, these Gordonia phages are not related to Crossroads and do not have homologues of Crossroads 132 and 141. Instead, Kita and its relatives are targeted for defense through Kita gene 53 and its homologues, which are of unknown function, and are not present in Crossroads. To examine the functional relationship between the two defense systems, plasmid pMM16 expressing CarolAnn 43 and 44 was transformed into *M. smegmatis* and tested for Crossroads infection (Fig. 9B). Somewhat surprisingly, CarolAnn 43/44 conferred defense not only against Crossroads but also against the Crossroads DEMs—including those in gene 132, in gene 141, or in the 141-to-142 intergenic region efficiently escaping the defense system (Fig. 9B). We have also shown that the Sbash genes 30 and 31 confer defense against *Gordonia* phage Kita when expressed in *Gordonia terrae* (see accompanying manuscript [33]). Sbash gp30/gp31 and the CarolAnn gp43/gp44 systems thus act similarly in *M. smegmatis* and *G. terrae*, even though they recognize different and unrelated target phages.

**DISCUSSION**

We have described here a new viral defense system encoded in mycobacteriophage Sbash, which is expressed lysogenically and defends against phage Crossroads with...
CarolAnn genes 43 and 44 confer defense against Crossroads. (A) Genome comparison of mycobacteriophage Sbash and Gordonia phage CarolAnn. Shading between the genomes reflects nucleotide sequence similarities, although there are only two short regions of weak similarity at the extremities of Sbash 30 and CarolAnn 43. Sbash gp30/CarolAnn gp44 and Sbash gp31/CarolAnn gp43 homologous pairs share 50% and 43% amino acid identity, respectively. Sbash 30 and 31 are transcribed rightward and are displaced from the repressor gene (gene 43), whereas CarolAnn 43 and 44 are coexpressed with the repressor (gene 45). (B) Serial dilutions of Sbash, Crossroads, Pipsqueaks, and 12 Crossroads DEM mutants (labeled phggXXX) were plated onto lawns of \textit{M. smegmatis} mc\textsuperscript{2}155pMH94 (vector) and \textit{M. smegmatis} mc\textsuperscript{2}155pMM16, which carries CarolAnn genes 43 and 44.
remarkable specificity. Viral defense plays a central role in bacterial survival in most environments, and Sbash colludes with its \textit{M. smegmatis} bacterial host to mutually benefit the phage and the host. There is at least one other Sbash defense system responsible for defense against phage Giles, and the other subcluster I2 phage, Che9c, codes for at least one additional defense system. It seems probable that such prophage-mediated defense systems are widespread across the bacterial landscape of diversity, and the variety of systems suggests that there is a vast reservoir of such defenses remaining to be discovered. These likely dwarf the number of bacterially encoded systems reported to date (7).

We propose a mechanism of Sbash 30/31 defense reflecting the proposed mechanism for lambda RexAB exclusion of T4rII; Sbash gp30 is activated by Crossroads infection, leading to stimulation of the ion channel potential of gp31, membrane depolarization, and loss of intracellular ATP (Fig. 8D). As a consequence, lytic growth is inhibited and the infection aborts. Consistent with this is the observation that early gene expression of Crossroads proceeds in the presence of Sbash 30/31, indicating that there is not a block to DNA injection or initiation of the transcriptional program (Fig. 7). Reduced levels of transcription and DNA replication occur at later time points, consistent with an overall inhibition of macromolecular synthesis (Fig. 8A). Whereas the lambda RexAB system is thought to be activated by binding of DNA, this seems unlikely for Sbash 30/31, as Crossroads gp132 and gp141 are both required for targeting of Crossroads, and neither is essential for lytic growth. It is plausible that gp132 and gp141 act to directly interact with Sbash gp30 to stimulate gp31 ion channel activity (Fig. 8D).

It is striking that the homologous defense system from \textit{Gordonia} phage CarolAnn confers defense against Crossroads in \textit{M. smegmatis} and that, reciprocally, the mycobacterial Sbash system defends against phage Kita (and its relatives) in \textit{Gordonia} (see accompanying paper [33]). In each instance, defense escape mutants isolated in the cognate system also escape the noncognate system. However, the targeted phages (Crossroads, Kita) are unrelated, and the escape mutants map in entirely different genes. This emphasizes the complexity of these defense systems and how they can respond to different types of attacking phages. It is also noteworthy that Kita gp53 expression leads to growth inhibition in the presence of CarolAnn 43/44, consistent with an abortive infection system as described above (see accompanying paper [33]). Further studies are required to determine the mechanism of activation in these different systems and whether Crossroads and Kita activate via similar mechanisms notwithstanding their sequence differences.

The Sbash prophage targets Crossroads with remarkable specificity, and none of the other closely related subcluster L2 phages are similarly defended against (Fig. 4B). It is striking that phages Wilder, LilDestine, and Crossroads have identical copies of Crossroads 132 and that the 141 homologues differ by five or fewer substitutions. One plausible explanation is that these differences reflect the evolutionary dynamics differing between the prophage-mediated defense systems and the infecting phages, with selection of defense and counterdefense systems, respectively. Thus, it is possible that Crossroads is the outlier and that—unlike the other subcluster L2 phages—it has not yet acquired a mechanism to escape the Sbash system, even though mutants that do so can readily be isolated.

Finally, we note that although phages Sbash and CarolAnn are the only two phages in the collection of 2,800 sequenced actinobacteriophages that encode these defense systems, there are several sequenced isolates of \textit{Mycobacterium fortuitum}, \textit{Mycobacterium immunogenum}, and \textit{Rhodococcus} sp. that have homologues of Sbash gp30 and that also encode a small membrane protein immediately downstream; most are likely encoded by prophages (see Fig. S2 in the supplemental material). This suggests there may be a much larger family of defense systems that show organization and operation similar to those of lambda \textit{rexAB} and whose members have yet to be fully explored.
MATERIALS AND METHODS

Strains and plasmids. *M. smegmatis* mc²155 (29) was used for phage growth and analyses as described previously (4). Phage lysates typically had titers greater than 10⁸ PFU ml⁻¹ and were used for plaque assays and DNA extraction. Lysogens were isolated by spotting 10 µl of high-titer lysate onto a lawn of *M. smegmatis* mc²155, incubating at 37°C for 18 to 24 h, and streaking for isolated colonies from the infected area. Individual colonies were propagated and tested for phage release and immunity to superinfection.

Primers and plasmids used in this study are shown in Tables S1 and S2 in the supplemental material, respectively. Plasmid vectors pJV39 and pGH1000 are hygromycin (Hyg)-resistant, integrated vectors containing the integration cassette of mycobacteriophage L5 and Giles, respectively (30, 31). Cloning of genes into vectors pJV39 and pGH1000 was done using Gibson assembly (New England Biolabs, Inc.) with PCR-amplified substrates inserted at the vector HindIII restriction site. Plasmid vector pCCX39 is a streptomycin (STR)-resistant integrated vector containing the integration cassette of mycobacteriophage L5 (32). The pCCX39 backbone is a Tet-ON plasmid (34) containing an inducible promoter; derivatives of pCCX39 were constructed to include 20 bp upstream of the gene(s) of interest to include the native RBS. Cloning of genes into pCCX39 was done using Gibson assembly (New England Biolabs, Inc.) with PCR-amplified substrates inserted at the vector HindIII restriction site. Gene expression of pCCK39 derivatives described above. All phage-derived DNA fragments to be cloned were amplified from phage Sbsh or phage Crossroads genomic DNA using oligonucleotide primers as shown in Table S1. Plasmid DNAs were extracted from *Escherichia coli* and transformed into electrocompetent *M. smegmatis* mc²155 cells. All constructed plasmids were verified by DNA sequencing. Site-directed mutants were constructed using a Q5 site-directed mutagenesis kit (New England Biolabs, Inc.) to construct plasmid pGG24, in which gene 30 is deleted from plasmid pGG05.

**Phage growth and analyses.** High-titer phage lysates were prepared by diluting a lysate 10-fold into phage buffer with 0.1 M CaCl₂ and plating 10 µl of the 10⁻¹ dilution with 250 µl of mc²155 and 3 ml of liquid Middlebrook 7H9 medium. All plates were incubated at 37°C for 18 to 24 h. Lysates were collected, filtered through a 0.22 µm-pore-size filter, and then stored at 4°C. The phage genome sequences have been reported previously or are available in GenBank. A Phamerator database (Actinobacteriophage_Draft, August 2018) was used for comparative genomic analyses (35). Phage genomic DNA was extracted using either a Wizard DNA kit (Promega) or phenol-chloroform as described previously (36) and was completely sequenced using Illumina MiSeq 150-base single-end runs and assembled as described previously (11).

**Isolation and characterization of defense escape mutants (DEMs).** Defense escape mutants (DEMs) were identified by plating dilutions of independently prepared lysates onto lawns of *M. smegmatis* mc²155 lysogenic strains or onto lawns of a strain carrying plasmid pGG05 (Table S2). Individual plaques of defense escape mutants were recovered from independently prepared lysates, purified, and amplified. Phage DNAs were isolated and sequenced.

**Construction of phage mutants.** Phage mutants were constructed using Bacteriophage Recombining of Electroporated DNA (BRED) as previously described (37). An approximately 500-bp gBlock DNA fragment (Table S3; Integrated DNA Technologies, Inc.) was synthesized and PCR amplified using ~20-bp flanking primers and Q5 high-fidelity DNA polymerase (New England BioLabs, Inc.). This DNA substrate was coelectroporated with phage genomic DNA (150 to 300 ng) into electrocompetent recombining cells (*M. smegmatis* mc²155pJV53). Following recovery for 1 h with shaking at 37°C in 7H9 medium–10% ADC (0.5% Bovine Serum Albumin, 0.2% Dextrose, 0.85% NaCl)–1 mM CaCl₂, the cells were plated with 7H9 top agar and wild-type *M. smegmatis* mc²155 cells. Primary plaque picks were screened by PCR to identify those containing the mutant allele, and mixed plaques were picked, diluted, and replated. Secondary plaques were picked and screened by PCR, followed by amplification.

**RNAseq analysis.** For RNAseq analysis, RNA was isolated from *M. smegmatis* lysogens in late logarithmic growth or from phage-infected cells (multiplicity of infection of 3) at 30 or 150 min after infection. Following DNA removal and rRNA depletion performed using a Turbo DNA-free kit (Ambion) and a Ribo-Zero kit (Illumina), respectively, a TruSeq stranded RNAseq kit (Illumina) was used to prepare libraries. These were evaluated using a BioAnalyzer and run on an Illumina MiSeq system. The fastq reads were analyzed as described previously (38). The Integrative Genomics Viewer (39) was used to visualize and present the RNAseq coverage. RNAseq data sets, with additional method details, are deposited in the Gene Expression Omnibus (GEO) with accession number GSE121829. The average read length was 140 bases. To determine the ratios of late transcripts to early transcripts at the late time point, the number of reads mapping to late genes (genes 1 to 32 and 96 to 113, and genes on the strand opposite that harboring genes 125 to 138 and 140 to 145) was divided by the number of reads mapping to early genes (genes 36 to 37, 41 to 48, and 59 to 118). The ratios were calculated within each sample; thus, the variation of number of reads between each sample (likely due to differences in infection efficiency) did not factor into the calculation.

**Toxicity analysis.** Plasmids were constructed as outlined above, and inducible clones were transformed into strains of interest by electroporation. Transformed cultures were grown to saturation in the presence of STR and were diluted and grown with STR to an optical density at 600 nm (OD₆₀₀) of 0.5. Cultures were then serially diluted 10-fold, and the next to 10⁻⁷ dilutions were spotted onto 7H11-STR
agar plates with (induced) and without (Uninduced) 100 ng/ml ATc. Plates were incubated for 3 days at 37°C and imaged.

**Defense complementation analysis.** M. smegmatis mc²155 and mc²155(Sbash) strains containing pKS006 or pGG38 or empty vector were grown to saturation in the presence of selection, and lawns were poured on 7H10-Hyg or 7H11-STR plates with (induced) and without (Uninduced) ATc; 100 ng/ml ATc was used for the strains containing pKS006 and relevant controls, while 5 ng/ml ATc was used for the strains containing pGG38 and relevant controls. Phage lysates were serially diluted 10-fold and spotted onto lawns, and plates were incubated at 37°C for 1.5 days, except for the strains containing pGG38, which were incubated for 4 days, and then imaged.

**Data availability.** The RNAseq data sets, with additional method details, have been deposited in the Gene Expression Omnibus (GEO) with accession number GSE121829. All phage genome sequences are available at https://phagesdb.org.

**SUPPLEMENTAL MATERIAL**

Supplemental material for this article may be found at https://doi.org/10.1128/mBio.00196-19.

**FIG S1**, PDF file, 0.2 MB.

**FIG S2**, PDF file, 0.4 MB.

**TABLE S1**, PDF file, 0.02 MB.

**TABLE S2**, PDF file, 0.03 MB.

**TABLE S3**, PDF file, 0.03 MB.

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