Comparative genomics and phylogeny unveil lineage diversification of *Citrobacter rodentium* polyvalent bacteriophages

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ABSTRACT

Citrobacter rodentium is a mouse-restricted pathogen that has long been used as an *in vivo* model for two important human intestinal pathogen enteropathogenic *Escherichia coli* (EPEC) and enterohaemorrhagic *E. coli* (EHEC). And yet, in contrast to *E. coli*, little is known about the bacteriophages (phages) – bacterial viruses – that infect *C. rodentium*, reflecting in part a need to isolate and comparatively analyze phages associated with this bacterial species. Here, we isolated two novel virulent phages CrRp3 and CrRp10 that infect *C. rodentium* and conduct *in vitro* and comparative genomic studies with other, related phages. Whole-genome analyses revealed that CrRp3 and CrRp10 phages are members of *Sp6virus* and T4virus genera, respectively. In addition, we show that these phages have pervasively mosaic genome architectures by actively evolving with several horizontal genetic exchange and mutational events. Phylogenetic analyses showed that these phages are more closely related to *E. coli* phages than those infecting *Citrobacter* genus, suggesting that CrRp3 and CrRp10 may have evolved from *E. coli* phages rather than *Citrobacter* spp. phages.

INTRODUCTION

The genus *Citrobacter* belongs to the family of Enterobacteriaceae and comprises eleven different species of facultative anaerobic Gram-negative bacilli, widely distributed in water, soil, food and intestinal tract of human and animals. Previously recognized as colonizers with low virulence or environmental contaminants, they are now known to account for up to 6% of all nosocomially acquired life threatening *Enterobacteriaceae* infections, such as urinary tract, respiratory, wound, bone, bloodstream, and central nervous system infections (1, 2). While the majority of human infections are caused by *Citrobacter freundii* and *Citrobacter koseri*, *Citrobacter youngae*, *Citrobacter braakii*, and *Citrobacter amalonaticus* are also important human pathogens; all of which are increasing difficult to treat due to the rise of multidrug resistance (MDR) (1-3). By contrast, *Citrobacter rodentium* is a naturally occurring mouse-restricted pathogen, which is highly similar to the important human pathogens enteropathogenic *Escherichia coli* (EPEC) and enterohaemorrhagic *E. coli* (EHEC) (4-6). Thus, *C. rodentium* has become an important *in vivo* model of several human intestinal diseases and disorders (7, 8).

We have little knowledge of how potentially important processes bacteriophages (phages) – viruses that prey on bacteria – play in shaping bacterial population phenotypes in perhaps the most important and clinically relevant microbial ecosystem – the human microbiome. Despite knowing that antagonistic interactions between phages and bacteria play a key role in driving and maintaining microbial diversity, eco-evolutionary processes in humans and animals have not received much research attention (9, 10). As both obligate parasites and vectors of horizontal gene transfer, a better understanding of phage strain diversification and how viral diversity might impact bacterial diversity and populations is required. Furthermore, there is renewed interest in the use of phages to eliminate or modulate bacterial population, namely phage therapy, partly due to their specificity for host bacterial species and ability to kill MDR pathogens (11, 12).

Phages are the most abundant biological entities on the planet, with an estimated 10^{31} present in the biosphere (13), suggesting there is an untapped biodiversity of *Citrobacter* phages. And yet, only a limited number of *Citrobacter* phages have been previously described, most of which only at the genome nucleotide level. Furthermore, most previously characterized phages infect *C. freundii*, including members related to *T4virus* (Merlin, Miller, Moon) (14-16), *FelixO1virus* (Michonne, Mordin, Moogle) (17-19), *T1virus* (Stevie) (20), and *T7virus* (phiCFP-1, SH1, SH2, SH3, SH4, SH5) (21, 22) genera. Genome sequences are also publically available for the *T7virus* related phages CR8 and CR44b that specifically infect *C. rodentium*(23). Among all these phages, only the *C. freundii T7virus* related phages have been experimentally studied.

In this study, we isolated and characterized two novel virulent phages, CrRp3 and CrRp10, which could

infect and lyse the mouse-restricted intestinal pathogen *C. rodentium*, as well as several human intestinal pathogenic *E. coli* strains, suggesting these phages could be effective for their control. CrRp10 is, to the best of our knowledge, the first sequenced *C. rodentium Myoviridae* phage. Furthermore, comparative genomic analyses provide evidence that these phages may have diversified from other *E. coli Sp6virus* and *T4virus* phages, respectively.

RESULTS

Biological characteristics of two novel *C. rodentium* phages. We isolated phage strains vB_CroP_CrRp3 (CrRp3) and vB_CroM_CrRp10 (CrRp10) from different wastewater samples from Paris France after they were found to form distinct clear plaques on the *C. rodentium* strain ICC180. Figure 1A is transmission electron microscopy image that revealed each phage has a different Caudovirales morphotype (24). CrRp3 has an icosahedral head and short tail, which implies it belongs to the *Podoviridae* family, while the icosahedral head and long tail of CrRp10 morphologically classifies this phage in the *Myoviridae* family.

CrRp3 and CrRp10 both have the ability to reduce bacterial population growth in liquid culture; the more potent virus appears to be the *Podoviridae* CrRp3 (Fig. 1B). That is, CrRp3 exhibited early lytic activity dynamics at the lowest MOI (0.001) tested that required ~60 min to show signs of reversing bacterial population growth, whereas CrRp10 required ~135 min. Increasing the phage concentration (i.e. higher MOIs of 0.1 and 10) abolished any differences in early lytic activity to initiate bacterial density reduction (Fig. 1B). However, late lytic activity dynamics again revealed differences in lytic potency. CrRp3 was able to nearly eliminate bacterial density by ~2.2 h at each MOI tested, while CrRp10 required at least a further 30 min to show bacterial elimination (Fig. 1C). Furthermore, *C. rodentium* was able to gain resistance to CrRp3 infection within 4 h, whereas for the duration of the study, it did not against CrRp10 infection (Fig. 1C).

Genome structure and general features. Table 1 shows the general genomic features of CrRp3, CrRp10, as well as all other *Citrobacter* phage genomes available on the public database. CrRp3 has a genome size of 44.3 kb and CrRp10 a size of 171.5 kb and displayed average GC contents of 45.1% and 35.5%, respectively. Although these characteristics are in line with their family classification (i.e. *Podoviridae* and *Myoviridae*, respectively), CrRp10 displays the lowest GC% of all sequenced *Citrobacter* phages to date. Furthermore, it appears that all *C. rodentium* phages, including CrRp3 and CrRp10, exhibit GC contents significantly lower than their host *C. rodentium* (54.5% GC content). Likewise, *C. freundii* displays a higher GC content (~51.5%) than phages infecting it, with the exception of SH4 (52.6% GC content) (Table 1).

The gene annotated features of CrRp3 and CrRp10 are listed in Table S1 and Table S2, respectively. CrRp3 has a terminally repetitive dsDNA genome that consists of 54 coding sequences (CDSs) with 35% having putative functions. CrRp10 has a circularly permuted dsDNA genome that consists of 267 CDSs with 50% having putative functions and 10 tRNAs. CrRp10 to the best of our knowledge is the first *Myoviridae* infecting *C. rodentium* to have its genome completely sequenced. Neither phages exhibited gene similarities to known bacterial virulence-associated genes or lysogeny-associated genes. In addition, each genome appears to not encode antibiotic resistance genes, which is consistent with previous findings of these genes being rare in phages (25).

Phylogenetic analysis. Using the Genome-BLAST Distance Phylogeny (GBDP) method (26), we show that CrRp3 is more closely related to *Podoviridae* infecting *Escherichia* rather than those infecting any *Citrobacter* species. That is, *Citrobacter* (Table 1) and *Escherichia* (Table S3) phages have a heterogeneous clustering of closely related *Podoviridae* into three distinct clades, each with several tree branches supported by high bootstrap values (Fig. 2A). CrRp3 clusters with only *Escherichia* phages in clade 1 (C1), whereas phages infect the human pathogen *C. freundii* (phiCFP1, SH1, and SH2) cluster in clade 2 (C2) and (SH3 and SH4) clade 3 (C3). Interestingly, the only previously described *C. rodentium* phages, CR8 and CR44b, have genomes that also cluster C3 despite, as previously mentioned, these phages having little protein sequence homology (Fig S1). Lastly, phage CVT2 branches separately. This could be expected due to being isolated from the gut of termites on an uncharacterized *Citrobacter* species (27). The genome relationships of CrRp10 among other *Citrobacter* and *Escherichia Myoviridae* also show a heterogeneous clustering (Fig. 3A). CrRp10 groups with several *E. coli* phages (Ime09, vB_EcoM_UFV13, slur02, slur07 and slur14) in C1, with all exhibiting similarly sized branches. Other known *Myoviridae*, which infect *C. freundii*, are distantly related by clustering in the three other clades with other related *Escherichia* phages. Interestingly, C3 is composed of almost exclusively of *C. freundii* phages (IME, CF2, Miller, CfP1, and Margaery); with the exception of the *E. coli* phage Lw1.

Comparative genomic analysis with other, related phages. To elucidate the mechanisms of the different host specificities of phylogenetically close phages, comparative genomic analysis was conducted using BLAST genome alignment. The genomic comparison at the amino acid level of CrRp3 and CrRp10 against all *Citrobacter* phages showed that they are unique among other sequenced *Citrobacter*

phages (Fig. 4). CrRp10 is most closely related with phage Moon, with and average amino acid identity of 70% over 50% coverage.

CrRp3 showed a 90.7% gene homology to phage genes from the *Sp6virus* genus in the subfamily Autographivirinae (family *Podoviridae*) reinforcing the taxonomic relationship (Table S4). CrRp3 has the highest genome nucleotide and structure similarity to the *E. coli* phages K1-5 and K1-E, both of which belong to the *Sp6virus* genus (Fig. 2B). Almost half of CrRp3 genes products have as best hit either K1-5 or K1-E (Table S4). These, include the DNA and RNA polymerases DNA ligase and major capsid proteins, among many hypothetical proteins. CrRp3 genes products that differ from K1-5 include the head tail connector protein, endolysin, tailspike protein, lyase, minor structural protein and several proteins with unknown function (Fig. 2B). Interestingly, CrRp3 lyase and minor structural protein are the only genes products with similarity to those in *Citrobacter* phage CR8 (Table S1). By contrast, CrRp10 genome shares significant synteny with gene products of *E. coli* phage Ime09 belonging to the *T4viruses* genus in the within the subfamily *Tevenvirinae* (family *Myoviridae*) (98% nucleotide identity over the complete length) (Table S4, Fig. 3A). There are few genome features unique to CrRp10 compared to Ime09, including its tail fiber gene having a highly diverse C-terminal region (3% dissimilarly over 80% of the length). The *T4virus* genus is known to have extended host ranges, largely due to their tail fiber protein having the unique ability to bind to several outer membrane proteins or lipopolysaccharide (LPS) receptors (28). Another striking feature in the CrRp10 genome is the recombination event to gain a DUTPASE with high sequence similarity to the phage e11/2. This latter phage has been shown to infect the EHEC (29), which suggests that CrRp10 might be a good candidate for development of a new therapeutic agent to inhibit important *E. coli* O157:H7 strains. Other recombination events in CrRp10 have added several putative endonucleases with high similarity to other related Enterobacteriaceae phages (Table S2).

Lytic spectrum. Next, we tested the bacterial genera, species and strains lysis spectrum by which the virulent phages CrRp3 and CrRp10 along with other representative phages are capable of infecting and lysing bacterial cells (Table 2). While, in addition to their isolation strain of C. rodentium, CrRp3 can infect the *E. coli* strain K-12, while CrRp10 displays a much broader host range infecting K-12 and several pathotype strains of *E. coli*, as well as the *Erwinia carotovora* strain CFBP2141. Although the *E. coli* phage LF82_P10 also exhibits a relatively broad host range (10), it cannot infect *C. rodentium*. Moreover, most *E. coli* strains tested were resistant to CrRp3 and CrRp10, as well as the *Pseudomonas aeruginosa* and *Serratia marcescens* strains.

DISCUSSION

In this study, we report the genomic and phenotypic characterization of two novel virulent phages, CrRp3 and CrRp10, which infect the mouse-restricted pathogen *C. rodentium*. In addition to doubling the available *C. rodentium* phage genomes on the public database, these phages provide new evolutionary relationships with the expanding group of viruses belonging to the *Sp6likevirus* and *T4virus* genera, respectively. CrRp10 is the first reported virulent *Myoviridae* with genome sequence characterization, as the previously isolated and characterized *C. rodentium Myoviridae* phiCr1 genome could not been sequenced (30). CrRp3 and CrRp10 appear to be quite distantly related to the previously sequenced *C. rodentium* phages CR8 and CR44b (both *T7virus*) (23), as well as other phages that infect the human pathogen *C. freundii*.

Comparative genomics revealed that CrRp3 and CrRp10 may have evolved independently from closely related *E. coli* phages presumably because it was advantageous to gain new specificities to infect *C. rodentium*. The evolution of phages is a multifaceted and complex process, strongly influenced by mutational and horizontal acquisition of genetic elements and their subsequent infection of new hosts (10). Interestingly, the tail associated genes and endolysin gene from CrRp3 appear to have evolved the most from the same gene products of *E. coli* phage K1-5 (Fig. 2B). Phage K1-5 has been shown to exhibit two tail fiber genes, one carrying a lyase domain and the second one an endosialidase domain, which allow it to infect both *E. coli* K1 and K5 polysaccharide capsule types (31). Interestingly, CrRp3 lyase is more closely related to that from the *Citrobacter* phages CR8 and CR44b (Table S1), also known to infect *C. rodentium*, which suggests a mosaic genome structure that may be driven by recombination of modules from varied species. This is consistent with other phages of the genus *Sp6virus*, which exhibit a high genetic identity and structure (and highly specific RNA polymerase) (32), with the modest differences observed related to gene products for presumably adaptation to host constraints.

C. rodentium and EPEC and EHEC are A/E pathogens derived from a common ancestral origin (6). However, the genome of *C. rodentium* exhibits several features associated with a bacterium that has recently passed through an evolutionary bottleneck, including several large-scale genomic rearrangements and functional gene loss in core genomic regions (5, 6, 33, 34). This has led researchers to postulate that *C. rodentium* may have emerged alongside the development of laboratory mice model of human *E. coli* infection (6, 34). Several studies have demonstrated the reciprocal selection of phages on bacterial populations and bacteria on phage populations. There is also increasing evidence that this process can maintain phage diversity, influence phage virulence, and increase phage evolvability

(10). The genomic characteristics of CrRp3 and CrRp10 strengthen the hypothesis that their host *C. rodentium* has recently evolved from *E. coli*. Our work does not exclude the potential for isolation of other *C. rodentium* phages more closely related to phages infecting other *Citrobacter* species, for example, *C. freundii* phages Merlin and Moon.

Phages CrRp3 and CrRp10 exhibit polyvalence, infecting strain across several genera within the gram-negative *Enterobacteriaceae*, including *Citrobacter*, *Escherichia*, and *Erwinia*. By contrast, the previously characterized *C. rodentium* phage phiCR1, which target lipopolysaccharide (LPS) as a receptor, was shown to be highly species-specific infecting only *C. rodentium* and not closely related *Enterobacteriaceae*, including *C. freundii* and *E. coli* (30). Although, phiCR1 host range supports the general notion that phages are largely species-specific (35, 36), several other phages infecting various *Enterobacteriaceae* (35, 37, 38) have also been demonstrated to be polyvalent, as well as some among phages of staphylococci (39). Polyvalent phages infecting strains across genera raises the question as to their so called "optimal" host and whether a phage's host should be inferred by genomic relationships rather than bacterial strain plaquing *in vitro* (40). This may also be a relevant question to the identification of the bacterial hosts for phages identified from virome metagenomes.

Moreover, virulent phages similar to CrRp3 and CrRp10 are being re-investigated as potential antimicrobial agents to both combat bacterial diseases and the dissemination of MDR bacteria (11). Because mice are resistant to EPEC and EHEC infections, *C. rodentium* is widely used as an *in vivo* model system for several important human gastrointestinal diseases (41, 42). However, to the best of our knowledge, phages that infect *C. rodentium* have not yet been explored for antibacterial potential in preclinical animal models. The newly characterized C. rodentium phages, in particular the polyvalent *Myoviridae* phage CrRp10 resilient against resistance development, may lead studies into innovative antimicrobial agents for food safety, veterinary and clinical use.

MATERIAL AND METHODS

Bacterial strains, phage isolation, and culture conditions. The *C. rodentium* strain ICC180 (43) was used isolate phages from waste water. *C. rodentium* and other gram-negatives were grown at 37°C in Luria-Bertani (LB) medium on an orbital shaker. LB mixed with 1.5% agar provided a solid medium on which bacteria were cultured. Citrobacter phages were collected from Paris France municipal wastewater and isolated on lawns of early log phase seeded agar of *C. rodentium*. Select plaque lysates were serial passaged five times by spiking liquid ICC180 cultures grown to an optical density (OD) OD_{600} 0.25 with phages titer at a multiplicity of infection (MOI) of 0.1. When required, phage strains were further purified by cesium chloride density gradient (1.3, 1.5 and 1.6 g/ml) using ultracentrifugation at 140,000 g for 3 h. Single bands were dialyzed in cold H_2O once and cold Tris buffer twice.

Phage characterization. Phage inhibition of bacterial population growth was measured in changes in culture optical densitometry. Microtiter plate wells were filled with 100 µl of 2x LB concentrate spiked with 2 $x10^6$ CFU of *C. rodentium* strain ICC180. Phages were diluted in PBS and added to wells at different MOIs and PBS was added to achieve a total assay volume of 200 µL. A Promega GloMax plate reader was used to measure OD_{600} nm at 15-min intervals for 18 h, while being incubated at 37°C and orbital shaken for 30 secs prior to each read. Phage host ranges were determined by spotting 4 µL of 10^7 PFU phages onto air-dried lawns of mid-log growing test bacteria strains on agar and grown overnight (Table 2).

Aliquots of 10 µl of cesium chloride purified phages dialyzed against Tris buffer were applied to carbon-coated copper grids, negatively stained with 2% uranyl acetate for 30 s and observed under a transmission electron microscope Tecnai Biotwin 120 FEI- 1 (FEI Company, USA) operating at 120kV.

Genome sequencing and bioinformatics analyses. Phage DNA was extracted from sterile DNase and RNase pretreated cesium chloride purified phages by a phenol-chloroform extraction as previously described (44). DNA samples were sequenced using an Illumina MiSeq (Illumina Inc., San Diego, CA) with 2x250 bp read length. For phage sequence analysis, the quality of Illumina reads was visualized by FastQC v0.10.1 (http://www.bioinformatics.babraham.ac.uk/projects/fastqc/) . Quality controlled trimmed reads were assembled to a single contig using CLC Assembler (Galaxy Version 4.4.2). Protein-coding genes in the assembled contigs were predicted using Prodigal (45), and tRNAs were predicted using tRNAscan-SE (46). Additional annotation of genes was done by comparing against the NCBI NR, COG (47), and TIGRfam (48) databases. In addition, genomes were also manually annotated using HHPRED server (49). Genomic comparisons among related viral genomes and reference genomes were performed using tBLASTx or BLASTN (50).

Phylogenetic trees. The phylogenies of phages CrRp3 and CrRp10 were constructed using VICTOR (51). All pairwise comparisons of the nucleotide sequences were conducted using the Genome-BLAST Distance Phylogeny (GBDP) method (26) under settings recommended for prokaryotic viruses (51). GBDP approach was used for phylogenetic inference from all publically available *Podoviridae Citrobacter* phages, including phages CR8 and CR44b that infect *C. rodentium*, and 37 of the closest related *E. coli*

phages (Table S3). All available *E. coli* and *Citrobacter* phage genomes were downloaded from https://www.ncbi.nlm.nih.gov/genome/browse/. The resulting intergenomic distances (100 replicates each) were used to infer a balanced minimum evolution tree with branch support via FASTME including SPR post processing (52). The trees were rooted at the outgroup (*Synechococcus* phages**)** and visualized with FigTree V1.4.3 (http://tree.bio.ed.ac.uk).

Nucleotide sequence accession numbers. The complete genome sequence and annotations of phages vB_CroP_CrRp3 and vB_CroM_CrRp10 have been deposited GenBank under accession numbers MG775042 and MG775043, respectively.

Acknowledgements. We thank Elena Resen, Luisa De Sordi, and Marta Mansos Lourenço of Institut Pasteur for their assistance. CMM was supported by the European Molecular Biology Organization (ALTF 1562-2015) and Marie Curie Actions program from the European Commission (LTFCOFUND2013, GA-2013-609409), while DRR was supported by an European Respiratory Society Fellowship (RESPIRE2–2015–8416). The funders had no role in study design, data analyses, or manuscript preparation.

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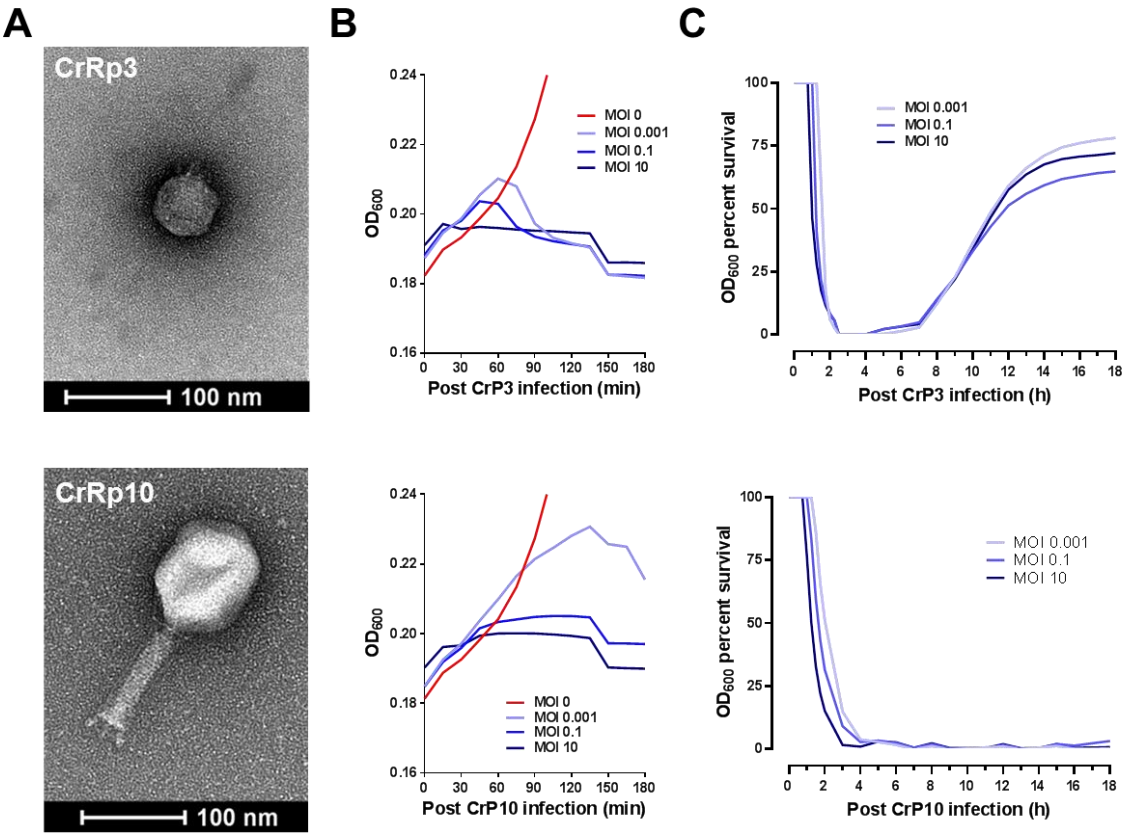

Figure 1. Morphological and Biological characterization of CrRp3 and CrRp10. A) Electron micrographs of *Citrobacter rodentium* phages negatively stained uranyl acetate. **B)** Early phage lysis dynamics and **C)** percent survival of *C. rodentium* cell populations at different initial multiplicity of infection (MOI) of CrRp3 (top) and CrRp10 (bottom) compared to growth in uninfected cultures (n = 3).

Table 1. Citrobacter bacteriophages and their genome features.

Phage	Host	Source	Phage family*	Size (kb)	GC%	Accession no.	Ref.
CrRp3	*C. rodentium*	municipal wastewater	P	44.3	45.1	MG775042	This study
CR44b	*C. rodentium*	sewage effluent	P	39.2	50.5	NC_023576	(23)
CR8	*C. rodentium*	sewage effluent	P	39.7	49.7	NC_023548	(23)
CVT22	*Citrobacter sp.*	termite gut	P	47.6	41.6	NC_027988	(27)
phiCFP-1	*C. freundii*	seawater	P	38.6	50.3	NC_028880	N/A
SH1	*C. freundii*	"	P	39.4	51	NC_031066	N/A
SH2	*C. freundii*	"	P	39.2	50.7	NC_031092	N/A
SH3	*C. freundii*	"	P	39.4	50.6	NC_031123	N/A
SH4	*C. freundii*	"	P	39.3	52.6	NC_031018	N/A
CrRp10	*C. rodentium*	municipal wastewater	M	171.5	35.5	MG775043	This study
IME-CF2	*C. freundii*	hospital wastewater	M	177.7	43.2	NC_029013	N/A
Margaery	*C. freundii*	"	M	178.2	44.9	NC_028755	N/A
Merlin	*C. freundii*	"	M	172.7	38.8	NC_028857	(14)
Michonne	*C. freundii*	"	M	90.0	38.8	NC_028247	(17)
Miller	*C. freundii*	"	M	178.2	43.1	NC_025414	(15)
Moogle	*C. freundii*	"	M	88.0	39	NC_027293	(18)
Moon	*C. freundii*	"	M	170.3	38.9	NC_027331	(16)
CfP1	*C. freundii*	sewage effluent	M	180.2	43.1	NC_031057	N/A
Stevie	*C. freundii*	soil	S	49.8	42.8	NC_027350	(20)

*Podoviridae (P), Myoviridae (M), Syphoviridae (S)

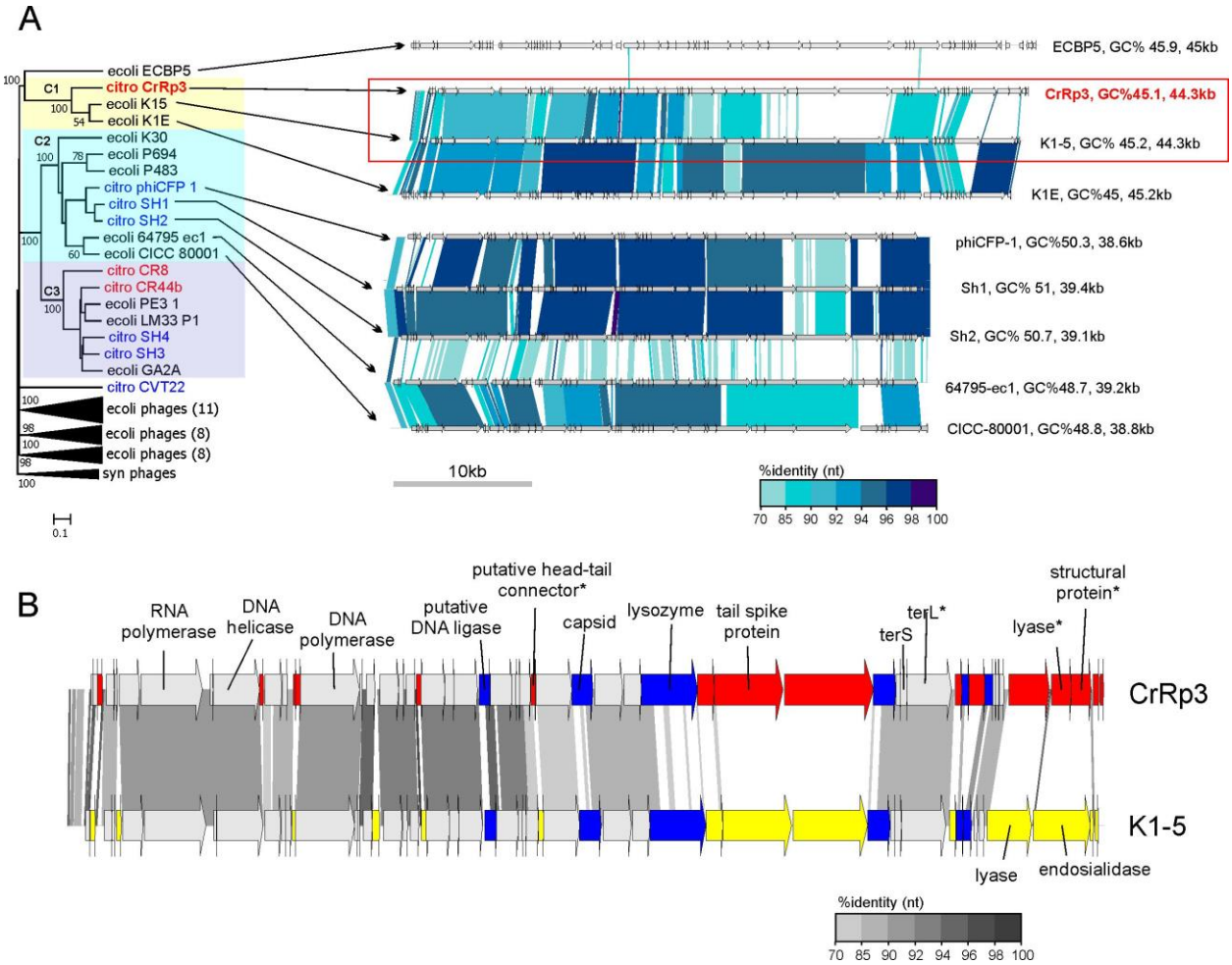

Figure 2. Genome structure and phylogeny of *Citrobacter rodentium* phage CrRp3. A) Genomic relationship of CrRp3 with other *Citrobacter* and *Escherichia* phages at nucleotide level. The tree shows bootstrap values (percentages of 100 replicates) below the branches and was rooted using *Synechococcus* phages as outgroup. Phages reported to infect *C. rodentium* are labeled in red and those that infect *C. freundii* and phage CVT22 are labeled in blue. **B)** Gene functional comparison of CrRp3 and *E. coli* phage K1-5. Genes are colored according to the relationship between CrRp3 and K1-5, with red labels being exclusive to CrRp3, yellow labels being exclusive to K1-5, while blue labels are homologous but highly variable. Gene products marked with (*) are those with some similarity to other *Citrobacter* phages (see Fig. 4).

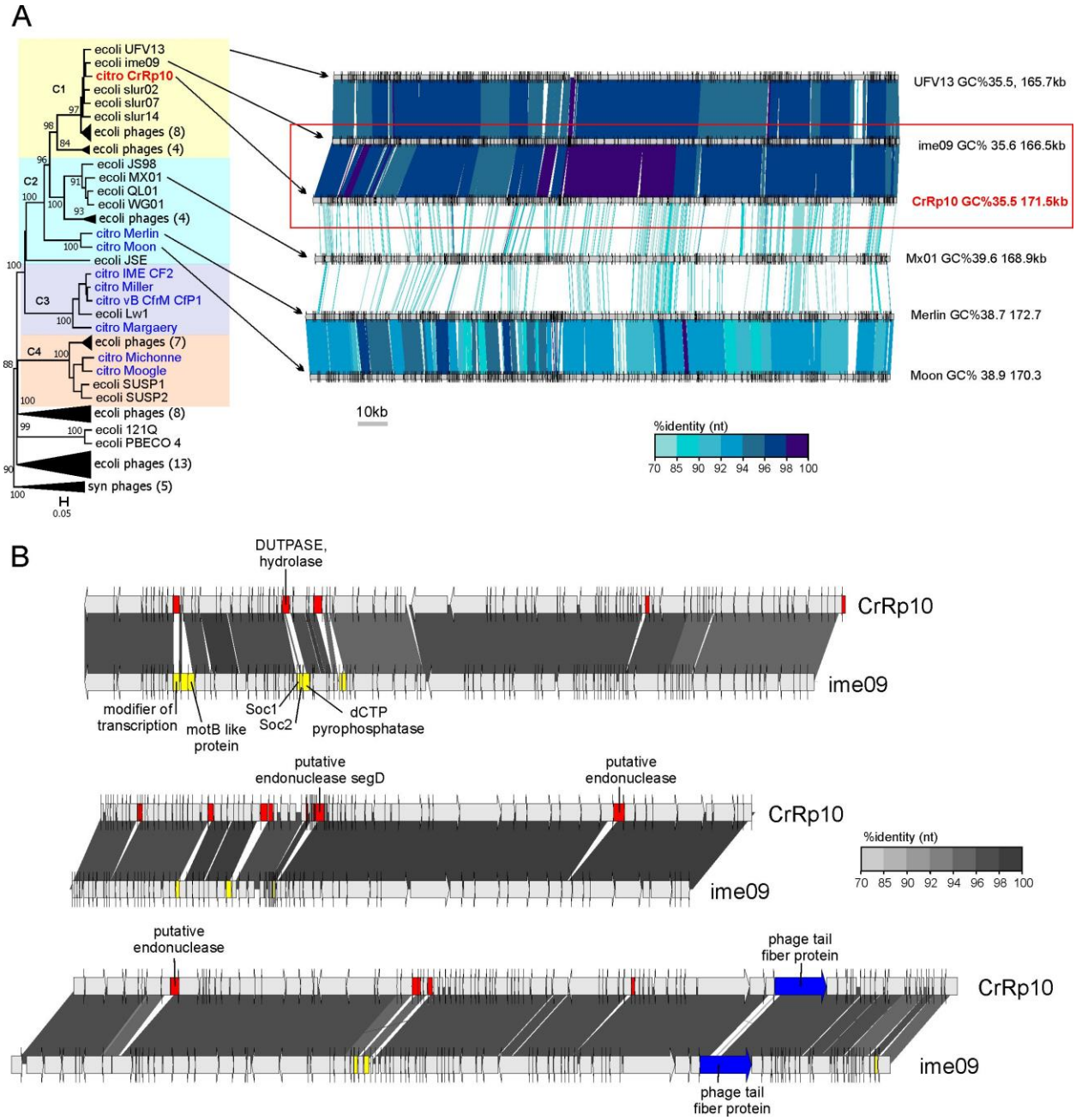

Figure 3. Genome structure and phylogeny of *Citrobacter rodentium* phage CrRp10. A) Genomic relationship of CrRp10 with other *Citrobacter* and *Escherichia* phages at nucleotide level. The tree shows bootstrap values (percentages of 100 replicates) below the branches and was rooted using *Synechococcus* phages as outgroup. Phages reported to infect *C. rodentium* are labeled in red and those that infect *C. freundii* are labeled in blue. **B)** Gene functional comparison of CrRp10 and *E. coli* phage ime09. Genes are colored according to the relationship between CrRp10 and ime09, with red labels being exclusive to CrRp10, yellow labels being exclusive to ime09, while blue labels are homologous but highly variable.

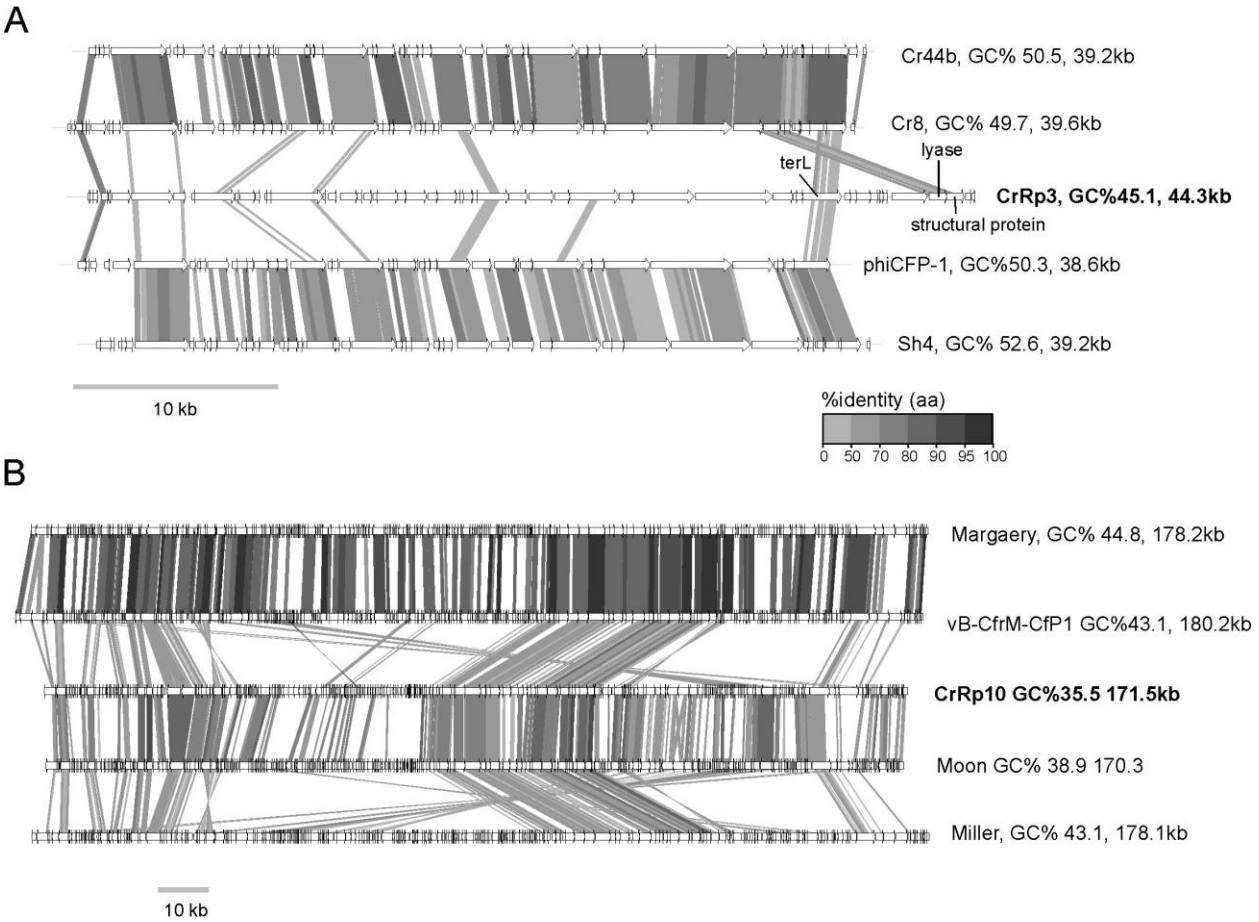

Figure 4. Whole genome alignment of *C. rodentium* phages at the amino acid level. Virulent *Podoviridae* **A)** and *Myoviridae* **B)**.

Table 2. Bacteriophage host ranges.

			Bacteriophage Strain[b]							
Species	**Strain**	**Type[a]**	**CrRp3**	**CrRp10**	**LF82_P10**	**LM33_P1**	**536_P7**	**AL505_P2**	**55989_P2**	**PAK_P1**
Citrobacter rodentium	ICC180		●	●	○	○	○	○	○	○
Erwinia carotovora	CFBL2141		○	●	●					○
Escherichia coli	LF31	AIEC	○	◗	●	○	○	○	◗	○
	LF50	AIEC	○	◗	○	○	●	○	○	○
	LF73	AIEC	○	●	○	○	○	○	○	○
	LF82	AIEC	○	○	●	○	○	◗	○	○
	LF110	AIEC	○	○	●	○	○	○	◗	○
	LF06075	AIEC	○	●	●	○	●	○	●	○
	LF07081	AIEC	○	○	○	○	○	○	○	○
	NRG	AIEC	○	●	●	○	○	○	○	○
	55989	EAEC	○	○	○	○	○	●	●	○
	E22	EPEC	○	○	○	○	○	○	○	○
	LM33	ExPEC	○	○	○	●	○	○	○	○
	Sp15	ExPEC	○	○	○	○	○	○	○	○
	536	UPEC	○	○	○	○	○	○	○	○
	AL505	UPEC	○	○	○	○	○	○	○	○
	MG1655	K-12	○	○	○		●		●	○
	BW25113	K-12	◗	◗	◗					○
	CR63	K-12	○	○	●					○
	OP_{50}	B	○	○	○					○
	BE	B	○	○	◗			●	●	○
	Nissle 1917	probiotic	○	●	●	○	◗	○	○	○
	M1/5		○	○	●					○
	SE15		○	●	○		●			○
	LM40		○	○	○	○	○	○	○	○
	H04		○	○	○	○	○	○	○	○
	BDX03		○	○	○	○	○	○	○	○
	LM02		○	○	○	○	○	○	○	○
	LM08		○	○	○	●	○	○	○	○
Pseudomonas aeruginosa	PAK		○	○	○	○	○	○	○	●
Rouxiella chamberiensis	nov		○	○	○					○
Serratia marcescens	Db11		○	○	○					○
	SM365		○	○	○					○

[a] *Adherent invasive E. coli (AIEC), enteroaggregative E. coli (EAEC), enteropathogenic E. coli (EPEC), extraintestinal pathogenic E. coli (ExPEC), and uropathogenic E. coli (UPEC)*

[b] Bacteriophages applied at 10^7 plaque forming units; ● clear plaque, ◗ partial clearing, ○ resistant, (blank) not tested