

Article



Genomic and Biological Profile of a Novel Bacteriophage, Vibrio phage Virtus, Which Improves Survival of *Sparus aurata* Larvae Challenged with *Vibrio harveyi*

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Abstract: Due to the emergence of multidrug-resistant bacteria, commonly known as "superbugs", phage therapy for the control of bacterial diseases rose in popularity. In this context, the use of phages for the management of many important bacterial diseases in the aquaculture environment is auspicious. *Vibrio harveyi*, a well-known and serious bacterial pathogen, is responsible for many disease outbreaks in aquaculture, resulting in huge economic and production losses. We isolated and fully characterized a novel bacteriophage, Vibrio phage Virtus, infecting *V. harveyi* strain VH2. Vibrio phage Virtus can infect a wide spectrum of *Vibrio* spp., including strains of *V. harveyi*, *V. owensii*, *V. campbellii*, *V. parahaemolyticus*, and *V. mediterranei*. It has a latent period of 40 min with an unusually high burst size of 3200 PFU/cell. Vibrio phage Virtus has a double-stranded DNA of 82,960 base pairs with 127 predicted open reading frames (ORFs). No virulence, antibiotic resistance, or integrase-encoding genes were detected. In vivo phage therapy trials in gilthead seabream, *Sparus aurata*, larvae demonstrated that Vibrio phage Virtus was able to significantly improve the survival of larvae for five days at a multiplicity of infection (MOI) of 10, which suggests that it can be an excellent candidate for phage therapy.

Keywords: Vibrio harveyi; phage therapy; aquaculture; gilthead seabream

1. Introduction

Vibrio harveyi belongs to the Vibrionaceae family and is an opportunistic, serious pathogen responsible for many disease outbreaks in marine animals worldwide. It is established as the main cause of gastroenteritis and vibriosis in various fish, crustacean, and molluscan species [1,2]. V. harveyi is ubiquitous and usually grows in temperatures above 18 °C, as its optimal temperature is 25 °C [3]. Climate change and the overall rise of the water temperature in the oceans, along with the intensification of aquaculture, favor the increase in vibrios, and hence the vibriosis incidents have increased alarmingly [3–7]. To date, the management of Vibrio infections has relied mostly on antibiotics such as oxytetracycline, flumequine, and ampicillin. [8]. However, the extensive use of such treatments is associated with the development of multidrug-resistant bacteria, affecting not only the management of the diseases in aquaculture but also humans since antimicrobial resistance (AMR) can be transmitted from livestock to humans [9]. Therefore, a new strategy for tackling the problems related to antibiotics in aquaculture is urgently required. Phage therapy, the use of phages as biocontrol agents, is considered a promising alternative [10,11]. The ease with which phages may be isolated, their abundance and host specificity, along with the high cost and effort required for the development of novel antimicrobial agents, have shifted the attention of scientific community to phages. There is an increasing number of studies regarding the application of phage therapy in aquaculture, yielding promising results [12–14]. However, phage therapy requires a thorough understanding

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Copyright: © 2022 by the authors. Licensee MDPI, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons Attribution (CC BY) license (https://creativecommons.org/licenses/by/4.0/). of the bacteriophages being used, determining their genomic and biological characteristics. Here, we isolated and fully characterized a lytic bacteriophage, Vibrio phage Virtus, infecting the pathogenic, antibiotic-resistant *Vibrio harveyi* strain VH2 [15], and tested its efficacy in vitro against its host and in vivo using gilthead seabream, *Sparus aurata*, larvae.

2. Materials and Methods

2.1. Bacterial Strains

Twenty-six strains of *Vibrio harveyi*, *V. alginolyticus*, *V. owensii*, *V. anguillarum*, *V. campbellii*, *V. parahaemolyticus*, *V. campbellii*, and *V. rotiferianus* (Table 1) used in this study were obtained from the bacterial collection of the Laboratory of Aquaculture Microbiology, Institute of Marine Biology, Biotechnology and Aquaculture (IMBBC), Hellenic Center for Marine Research (HCMR) in Heraklion, Crete. The bacterial strains were previously identified either through their NCBI or ENA accession numbers for the type strains, biochemical test (BIOLOG GEN III), and PCR method (sequencing of 16s rRNA and toxR amplifications). Moreover, a strain of *Phaeobacter piscinae*, a kind offer of Prof. Lone Gram (DTU), was also included in the assays. All the bacterial strains were maintained in microbeads (MicroBank) at -80 °C and were grown in lysogeny broth (10 g/L tryptone, 5 g/L yeast extract, 10 g/L NaCl, 1 L deionized water, 0.75 g/L MgSO4, 1.5 g/L KCl, 0.73 g/L CaCl₂) at 25 °C when used.

2.2. Isolation and Purification of Bacteriophages

Water samples were collected from a fish tank in the broodstock section of HCMR in Gouves, Heraklion, Crete. An overnight *Vibrio harveyi* VH2 culture in LB broth (2.5 mL) was added in 25 mL of concentrated LB and used for the enrichment of the water samples followed by incubation at 25 °C with a shaking speed of 80 rpm for 24 h. Subsequently, enrichments were centrifuged at 13,000 rpm for 3 min, and supernatants were filtered through a 0.22 μ m sterile filter (GVS Life Sciences, Sanford, ME, USA). A total of 10 μ L of each sample was spotted on bacterial lawns of the host strain. After an incubation for 24 h, the clearest plaques were collected and serially propagated against the host using the double agar layer method according to Clokie et al. as described in Misol et al. [16,17]. The same procedure was repeated five times. The purified phage was selected for further characterization and was designated Vibrio phage Virtus.

2.3. Transmission Electron Microscopy

A purified, high-titer (>10¹¹ PFU mL⁻¹) aliquot of the novel phage obtained following PEG centrifugation was used for electron microscopy observation. Negatively stained (4% w/v uranyl acetate, pH 7.2) samples were prepared at the Electron Microscopy Laboratory of the University of Crete as described previously by Misol et al. [17] and observed with a JEOL transmission electron microscope at 80 kV. Morphology and size of virions were obtained from digital micrographs using ImageJ (software version 1.53p) [18]. Measurements (n = 15) were obtained for capsid width, tail length, baseplate length, and baseplate width.

2.4. Lysogeny Test

To examine whether Vibrio phage Virtus can follow a lysogenic cycle, we developed phage-resistant host mutants following the method described in Thomas Denes et al. [19]. Briefly, a high titer of Vibrio phage Virtus was added in liquid cultures of host bacteria *V. harveyi* VH2, following a 24 h incubation. Samples were then taken from each culture and plated on LB agar. After incubation for 24 h, more than 20 colonies were isolated from each plate. Colonies that were resistant to phage and retained resistance through subsequent recultures, were selected and stored as phage-resistant mutants. We assume that if the phage was temperate, some of the phage resistant mutants would be lysogenized. Following that, prophage induction was conducted to 11 resistant mutants according to

Jackel et al. [20] with minor modifications. Briefly, aliquots of overnight bacteria cultures at 10⁷ CFU mL⁻¹ were mixed with top molten LB agar (0.75% agar) and poured on bottom LB agar. The Petri dishes were then placed in 10 cm to a UV lamp (6 W, 254 nm) for 5 s. The induction of any possible lysis was assessed the following day by examining plates for plaques.

2.5. Host Range Test

Host range analysis was conducted according to Misol et al. [17]. Fresh cultures of the bacterial strains used in this study (Table 1) inoculated LB broths at a concentration of approximately 10^7 CFU mL⁻¹. They were then mixed with top soft LB agar (0.75% agar) and poured on bottom LB/2 agar, which contained half of the tryptone and yeast content of LB agar. After the solidification of top agar, three spots of 10 µL of Vibrio phage Virtus were used for phage enumeration.

Table 1. Bacterial strains used in the study.

Strain ID	Species	Country	Location	Date	Host
DSM 19623	V. harveyi	USA	Massachusetts	-	Talochestria capensis
SA 2.1	V. harveyi	Saudi Arabia	Red Sea	July 2019	Sparus aurata
DSM 2171	V. alginolyticus	Japan	-	-	Trachurus trachurus
Gal 90	V. harveyi	Greece	Central Greece	September 2020	Sparus aurata
Vh No22	V. harveyi	Greece	Ionian Islands	September 2015	Dichentrachus labrax
Kef 62	V. harveyi	Greece	Ionian Islands	July 2020	Dichentrachus labrax
Kef 75	V. harveyi	Greece	Ionian Islands	July 2020	Dichentrachus labrax
Gal 56	V. harveyi	Greece	Central Greece	May 2020	Dichentrachus labrax
Gal 77	V. harveyi	Greece	Central Greece	July 2020	Sparus aurata
Gal 72	V. harveyi	Greece	Central Greece	June 2020	Dichentrachus labrax
Gal 94	V. harveyi	Greece	Central Greece	September 2020	Sparus aurata
L. SUSI	V. parahaemolyticus	Philippines	Philippines	April 2018	Shrimp
V1	V. alginolyticus	Greece	-	May 2018	Sparus aurata
LAR194	V. mediterranei	Greece	Central Greece	May 2020	Artemia nauplii
SM1	V. harveyi	Greece	Central Greece	-	Seriola dumerili
MAN113	V. splendidus	Greece	Saronikos Gulf	September 2019	Seriola dumerili
Varv A4/1.1	V. harveyi	Greece	Central Greece	July 2019	Sparus aurata
VH2	V. harveyi	Greece	Crete	2007	Seriola dumerili
VhP1 Liv	V. harveyi	Greece	Crete	April 2015	Seriola dumerili
VhP1 Spl	V. harveyi	Greece	Eastern Aegean	September 2015	Dichentrachus labrax
DY05	V. owensii	Greece	Eastern Aegean	September 2015	Dichentrachus labrax
SA 6.2	V. owensii	Saudi Arabia	-	July 2019	Oreochromis niloticus
VIB391	V. campbellii	Thailand	-	October 2016	Shrimp
Kef 56	V. rotiferianus	Greece	Ionian Islands	May 2020	Dichentrachus labrax
VhSerFre	V. harveyi	Greece	Crete	August 2015	Seriola dumerili
Phaeobacter S26	Phaeobacter piscinae	Greece	Euboea	July 2013	Artemia nauplii

2.6. Efficiency of Plating (EOP)

Efficiency of plating (EOP) was performed according to Kutter et al. [21], as described in Misol et al. [17]. The phage was serially diluted to $\approx 10^{-1}$, 10^{-2} , 10^{-3} , 10^{-4} , 10^{-5} , 10^{-6} , and 10^{-7} and spotted on the bacterial lawns of the 25 bacterial strains. Three spottings were used in order to assess phage titer after the agar plates were incubated at 25 °C for 24 h. The EOP was calculated as a percentage of the number of plaque-forming units formed on a bacterial strain against the number of plaque-forming units formed on the host *Vibrio harveyi* VH2. EOP more than 10 was categorized as high, EOP between 9.9 and 0.5 was considered medium, while EOP less than 0.5 was considered low.

2.7. Stability of Phage in Different Temperatures, pH Values, and Organic Solvents

The stability of the phage against different temperatures, pH values, organic solvents, and common disinfectants was assessed in order to determine phage versatility in therapy conditions. Phage aliquots of $\approx 10^{11}$ PFU mL⁻¹ were exposed to different temperatures (25, 30, 35, 40, 45, 50, 55, and 65 °C) for 1 h before being rested at 25 °C for 10 min. Following serial dilutions, the aliquots were spotted on host bacterial lawn. Vibrio phage Virtus stored at 4 °C for 24 h was used as a control. Stability studies for acidic and alkaline pH conditions were conducted according to the methods described by Pan et al. [22], with some modifications. Briefly, phages were suspended in LB adjusted with 1 M NaOH or HCl (Thermo Fisher Scientific, NJ, USA) to yield a pH range of 1-10, and incubated at 4 °C for 2 h, and then the phage aliquots were serially diluted and spotted on host bacterial lawn to determine the titer and the survival of the phage. The sensitivity of Vibrio phage Virtus to chloroform was determined by exposing $\approx 10^{11}$ PFU mL⁻¹ of the phage aliquots to 10% chloroform at 4 °C for 1 h, while the stability of the Vibrio phage Virtus against commonly used disinfectants in aquaculture was measured by exposing $\approx 10^{11}$ PFU mL⁻¹ of Vibrio phage Virtus to 0.001% benzalkonium chloride (BKC), 3% hydrogen peroxide (H₂O₂), 1% sodium hypochlorite (NaOCl), 70% ethanol (EtOH), and 1% formaldehyde (CH₂O) at 25 °C for 1 h. Vibrio phage Virtus incubated at 25 °C for 1 h were used as control. Each treatment was serially diluted and spotted on host bacterial lawn. The phage titer was assessed after the agar plates were incubated at 25 °C for 24 h. All assays were conducted in triplicate.

2.8. One-Step Growth

One-step growth of Vibrio phage Virtus was determined according to Clokie et al. as described in Misol et al. with some modifications [16,17]. Briefly, 1 mL of host culture inoculated LB broth until it reached exponential phase ($\approx 10^8$ CFU mL⁻¹) and was then centrifuged at 13,000 rpm for 3 min. The supernatant was then discarded, and the pellet was washed and resuspended in 1 mL of SM buffer (5.8 g/L NaCl, 2 g/L MgSO4, 50 mL 1 M Tris-Cl (pH 7.5) and 2% gelatin, 1 L deionized H₂O). This step was then repeated twice before the pellet was finally resuspended in 1 mL of LB. The fresh host culture was then inoculated with Vibrio phage Virtus at MOI 0.01. Following incubation for 10 min at 25 °C, the infected Vibrio harveyi VH2 culture was transferred to LB with the final volume of 30 mL. At 10 min intervals, 1 mL aliquots were collected from the infected host culture and were centrifuged for 13,000 rpm for 3 min. Subsequently, the supernatants were collected, serially diluted, and spotted on the host bacterial lawn on LB/2 agar plates. The phage titer was assessed after the incubation of agar plates at 25 °C for 24 h. For the assessment of the eclipse period, the same procedure was followed, but instead of centrifuging the samples, chloroform was added. Burst size was calculated as the ratio of the final count of liberated virions at the end of the burst period to the initial count of infected bacterial cells at the beginning of the latent period.

2.9. In Vitro Cell Lysis

The in vitro cell lysis of Vibrio phage Virtus against *Vibrio harveyi* VH2 was carried out by loading 180 μ L of fresh host bacterial culture in each well of sterile 96-well plates. The plates were then read at OD₆₀₀ using a TECAN microplate reader (Infinite PRO 200) at 25 °C with orbital shaking. A total of 20 μ L of Vibrio phage Virtus was then added at MOIs 0.1, 1, 10, and 100 when host culture was at the exponential phase ($\approx 10^7$ CFU mL⁻¹). Phages added to LB without host bacteria served as the control. The growth curves of the cultures were then measured every 10 min for 24 h. All assays were performed in triplicate.

2.10. DNA Extraction and Purification

The DNA extraction of Vibrio phage Virtus was carried out using the phenol-chloroform method according to Higuera et al. [23]. The extracted DNA was visualized for quality on 1% agarose gel electrophoresis at 80 kV for 1 h with a 50 kbp ladder. Milli-Q[®] Reference Water (Merck KGaA, Darmstadt, Germany) was used as a negative control. The extracted DNA of Vibrio phage Virtus was then stored in –20 °C.

2.11. Genomic Analysis

The genome of Vibrio phage Virtus was sequenced using the DNBSEQ[™] sequencer using paired-end technology (PE100) at BGI, Hong Kong. The workflow for library preparation for the platform included fragmentation, size selection, end repair and A-tailing, bubble adaptor ligation, PCR amplification, denaturation, splint circularization, enzymatic digestion and purification, and DNB making. Raw reads were filtered if more than 25% matched the adapter sequence, if more than 50% bases had quality values lower than 20 and if there were more than 3% N in the read. Filtering was completed using the SOAPnuke software. The raw reads were quality inspected and were assembled by Unicycler v0.4.8 in PATRIC [24]. QUAST v4.6.3[25] and BBMap v38.88 [26]were used to map the reads back to the assembled genome, while PhageTerm was used to predict phage termini [27] through the Galaxy server [28]. RASTk, Glimmer, and GeneMark were used for gene prediction. Sixpack, a naive gene caller, was used as validation to annotate genes that may have been missed by Glimmer. Moreover, potential protein-coding genes were manually checked to ensure the presence of a phage start codon (ATG/GTG or TTG), and a Shine-Dalgarno feature was added to all features that had a detectable match. Proteins of Vibrio phage Virtus were manually annotated using (i) NCBI Basic Local Alignment Search Tool (BLAST) [29] adjusted at non-redundant (nr) protein database, (ii) Gene Ontology [30], (iii) InterPro [31], and (iv) TMHMM 2.0 [32,33]. Predicted proteins of Vibrio phage Virtus were also manually annotated with NCBI Conserved Domain Database (NCBI CDD) [34]. All ORF predictions and annotations were manually inspected. Integrase, virulence, and antibiotic-resistance-encoding genes in Vibrio phage Virtus were for searched using the INTEGRALL Database webserver [35], Virulence Factor Database (VFDB) [36], and VirulenceFinder and ResFinder webservers [37]. The host Vibrio harveyi VH2 genome was analyzed for prophage-like sequences using Phage Search Tool Enhanced Release (PHASTER) [38]. A computational analysis using Bacphlip [39] was conducted in order to assess phage lifestyle on the basis of phage proteome. For protein structural homologies, only probabilities above 90% were accepted for manual protein function assignment of the Vibrio phage Virtus predicted ORFs. All hits were in existing databases with expected E-value below 10-3. The genome of Vibrio phage Virtus with annotated predicted ORFs was then visualized in a circular representation with Geneious software (Geneious v9.1, Biomatters, Auckland, Australia) and CGview.

2.12. Genome Alignment and Phylogenetic Analysis of Vibrio phage Virtus

The whole proteome of Vibrio phage Virtus was searched for similarity with other phages using the NCBI BLASTP nr protein database. The phage genomes with significant similarities were then downloaded and aligned with Vibrio phage Virtus using the progressiveMauve: Multiple Genome Alignment [40] for analysis of the genomic synteny. Pairwise alignment with of Vibrio phage Virtus with vB_VcaS_HC was conducted using Geneious Alignment with a cost matrix of 65% similarity (5.0/–4.0) on the basis of the Needleman and Wunsch (1970) and Smith and Waterman (1981) algorithms [41,42]. ViPTree was used to investigate the taxonomic position of Virtus and its host [43]. MEGA X was used to analyze the phylogeny and molecular evolution of the novel phage in comparison with other *Vibrio* phages [44]. Eighteen large terminase subunits of described Vibrio phages were downloaded from the NCBI database and were aligned with the large terminase subunit of Vibrio phage Virtus using MUSCLE algorithm [45]. Gaps in the

amino acid sequence alignments were trimmed. A maximum likelihood phylogenetic tree was constructed using the TN93 model [46] with bootstrap test = 1000. The tree was visualized using the Interactive Tree of Life web server [47].

2.13. In Vivo Phage Therapy Trial in Gilthead Seabream Larvae

Gilthead seabream (*Sparus aurata*) larvae were selected as a model to assess the therapeutic potential of Vibrio phage Virtus. Gilthead seabream eggs at the same developmental stage were obtained from HCMR hatchery, washed three times with sterile sea water, and placed individually in a 96-well microplate (1 egg/well) containing 200 µL sterile sea water. After one day of incubation, the quality of eggs was evaluated according to Panini et al. [48]. The challenge test started when eggs were hatched.

Bacteria used in the challenge test were grown in LB overnight and diluted 1:100 in fresh LB. After a 2 h incubation at 25 °C, cells were centrifuged and washed twice with buffer A (saline 0.9%, MgCl₂ 10 mM). The bacterial suspensions were adjusted to $\approx 10^7$ CFU mL⁻¹ with buffer A. No treatment occurred in the first group of larvae. The second group was treated with Vibrio phage Virtus alone (without addition of bacteria) at an approximate concentration of 10⁸ PFU ml⁻¹ and served as a negative phage control. The third group was treated with 10⁶ CFU ml⁻¹ of a *Phaeobacter piscinae* S26 strain, which has probiotic properties and served as a control to assess the effect of the addition of the same quantity of non-pathogenic bacteria on the viability of the larvae. The fourth group was treated with 10⁶ CFU mL⁻¹ Vibrio harveyi VH2. The fifth and sixth groups were treated with 10⁶ CFU mL⁻¹ Vibrio harveyi VH2 and Vibrio phage Virtus at 10 MOI. A second dose of Vibrio phage Virtus was administered the following day in the sixth group, at the same MOI. Phage suspensions were treated with 10% (w/v) PEG overnight at 4 °C to remove possible endotoxins in the phage lysate and were diluted in SM buffer (NaCl 100 mM, MgSO₄7H₂O 8 mM, Tris-Cl 1 M; pH 7.5). The phage titer was also determined prior to the experiment with double agar assay. Phage suspensions were added to the corresponding treatments two hours after infection. In addition, all controls were treated the same way, but instead of phage lysate, saline 0.9% was added to each well. The survival of gilthead seabream larvae was monitored daily for the following five days. A Kaplan-Meier survival curve was then constructed using GraphPad Prism version 9.0.0 for Windows (GraphPad Software, San Diego, CA, USA).

2.14. Statistical Analysis

One-way ANOVA was performed for the thermal and pH stability, and effects of organic solvents assays along with Dunnett's multiple comparison test [49]. Tukey's HSD post hoc test [50] was used as a multiple comparison tool after ANOVA was performed. Kaplan–Meier survival analysis [51] was performed for the in vivo phage therapy trial in gilthead seabream larvae. All statistical analyses were carried out using GraphPad Prism version 9.0.0 for Windows, GraphPad Software, San Diego, CA, USA).

2.15. Data Availability

The genome sequence of phage Vibrio phage Virtus is available in GenBank under accession number OK381870. The associated BioProject and BioSample accession numbers are PRJNA764828 and SAMN21529761, respectively.

3. Results

3.1. Isolation and Morphology of Vibrio phage Virtus

Vibrio phage Virtus was isolated from fish tank water collected from the broodstock section of the aquaculture facilities of the Institute of Marine Biology, Biotechnology and Aquaculture of the Hellenic Centre for Marine Research in Heraklion, Greece, against *Vibrio harveyi* VH2 [15]. Throughout the propagation steps, Vibrio phage Virtus showed a consistent plaque morphology producing pinhole-type plaques with a diameter of 0.42 ±

0.05 mm (n = 40). Transmission electron microscopy (TEM) showed that Vibrio phage Virtus has a long non-contractile, conspicuously striated tail and an icosahedral capsid (Figure 1), morphologically consistent with the *Siphoviridae* family. The phage capsid was 70 ± 05 nm in width, and the tail was 220 ± 10 nm long and 12 ± 2 nm wide. Finally, the baseplate had a width of 20 ± 02 nm and a length of 13 ± 01 nm.



Figure 1. Transmission electron microscopy picture of Vibrio phage Virtus showing a typical morphology of siphoviruses.

3.2. Host Range and Efficiency of Plating (EOP) of Vibrio phage Virtus against Multiple Antibiotic Resistant Strains

Vibrio phage Virtus was able to infect 13 out of 25 strains tested (Table 2). It infected 8 of the 16 strains of *V. harveyi*; the single strains of *V. parahaemolyticus*, *V. campbellii*, and *V. mediterranei*; and one out of two strains of *V. owensii*. The strains of *V. alginolyticus*, *V. rotiferianus*, and *V. splendidus* tested were not susceptible to the phage. EOP of Vibrio phage Virtus was high for four strains of *V. harveyi* (SA 1.2, VhP1 Spl, VH2, Kef 75), and moderate for five strains of other *Vibrio* spp. (VhSerFrE, Vh28, L. SUSI, SA 6.2, VIB391). All strains used in the assay are from the HCMR collection and have been identified to species level through sequencing.

Efficiency of Plating of Vibrio Phage Virtus										
		EOP								
SPECIES/51 KAIN	10 ⁰	10-1	10-2	10-3	10-4	10-5	10-6	10-7	(%)	
	Vibrio harveyi									
SA 2.1	++	++	++	+	+	+	+	-	High	
Varv A4/1.1	-	-	-	-	-	-	-	-	NF	
DSM 19623	++	+	+	+	+	-	-	-	Medium	
Vh No22	-	-	-	-	-	-	-	-	NF	
VhP1 Spl	+++	++	++	++	++	++	+	+	High	
VH2	++++	+++	+++	+++	+++	+++	+++	+++	High	
VhSerFre	++++	+++	+++	-	-	-	-	-	Medium	

Table 2. Host range and efficiency of plating of Vibrio phage Virtus against selected Vibrio spp.

-	-	-	-	-	-	-	-	NF
+++	+++	++	++	++	++	+	+	High
+	-	-	-	-	-	-	-	Low
+	-	-	-	-	-	-	-	Low
-	-	-	-	-	-	-	-	NF
-	-	-	-	-	-	-	-	NF
-	-	-	-	-	-	-	-	NF
-	-	-	-	-	-	-	-	NF
-	-	-	-	-	-	-	-	NF
		Vibri	o parah	aemolyt	icus			
++++	+++	-	-	-	-	-	-	Medium
		Vił	orio algi	nolyticı	IS			
-	-	-	-	-	-	-	-	NF
-	-	-	-	-	-	-	-	NF
		Vil	brio med	literran	ei			
+++	+	+	-	-	-	-	-	Low
		V_{i}	ibrio spl	lendidus	3			
_	-	-	-	-	-	-	-	NF
			Vibrio o	wensii				
++++	+++	+++	++	+	-	-	-	Medium
-	-	-	-	-	-	-	-	NF
		V	'ibrio ca	mpbellii				
++++	+++	+++	++	+	-	-	-	Medium
		Vi	brio rot	iferianu	S			
+	-	-	-	-	-	-	-	Low
	- +++ + - - - - - - - - - - - - - - - -	 ++++ +++ + - + - ++++ +++ ++++ +++ ++++ +++ ++++ ++++ +++ ++++	$\begin{array}{cccccccccccccccccccccccccccccccccccc$		- - - - ++++ +++ +++ ++ + - - - -	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	$\begin{array}{cccccccccccccccccccccccccccccccccccc$	- -

EOP: efficiency of plating; **NF:** no plaque formation; ++++: single large clearing zone +++: substantial turbidity throughout clearing zone; ++: ≥20 small plaques; +: <20 small plaques; **high:** EOP > 10.0%; **medium:** 0.5% < EOP < 9.9%; **low:** EOP < 0.5%.

3.3. Thermal and pH Stability of Vibrio phage Virtus and Exposure to Organic Solvents and Common Disinfectants

Exposure to different temperatures showed that Vibrio phage Virtus was stable between 4 and 55 °C (Figure 2a). No statistically significant difference (F(4, 10) = 0.1923, p = 0.9369) of its titer was observed at the temperatures assessed, while a complete inactivation was observed from 65 °C and above. The optimum pH of Vibrio phage Virtus was 6 (Figure 2b). Complete inactivation was observed at low pH values, while statistically significant reduction of the titer was observed at pH 3, 4, 5, 8, 9, and 10 compared to the control (F(9, 20) = 150.5, p < 0.001). A one-way ANOVA was performed to compare the effect of 6 different organic solvent solutions and common disinfectants to phage titer (F(6, 14) = 46.08, p < 0.001) (Figure 3). Hydrogen peroxide and chloroform did not affect Vibrio phage Virtus titer (p = 0.6609, p = 0.2975). However, there was a significant reduction when the phage was exposed to 70% ethanol (p < 0.001). Complete inactivation was observed in BKC, NaClO, and CH₂O.







Figure 2. (**A**) Effect of different temperatures on the stability of Vibrio phage Virtus. Incubation at 4 °C was used as control. (**B**) Effect of pH in the stability of Vibrio phage Virtus. Incubation with pH



= 7 was used as control. Phage titer was measured against *V. harveyi* VH2. Error bars were shown for the mean of n = 3. Statistical significance indicated by **** at p < 0.0001.

Figure 3. Effect of different organic solvents to the stability of Vibrio phage Virtus. Incubation with LB was used as control. Phage titer was measured against *V. harveyi* VH2. Error bars were shown for the mean of n = 3. Statistical significance indicated by **** at p < 0.0001 compared to the control.

3.4. One-Step Growth of Vibrio phage Virtus

One-step growth assay (Figure 4) showed that Vibrio phage Virtus has a latent phase of 40 min and an eclipse phase of 20 min. The rise phase was estimated between 40 and 110 min. The plateau phage was reached at 110 min. In this assay, the burst size of Vibrio phage Virtus was 3200 PFU per cell.



Figure 4. One-step growth of Vibrio phage Virtus measured against *V. harveyi* VH2 at multiplicity of infection (MOI) 0.01. Error bars were shown for the mean of n = 3.

3.5. In Vitro Cell Lysis

In vitro lysis assay with *Vibrio harveyi* VH2 showed that Vibrio phage Virtus was able to lyse the host bacterial population from MOI 0.1 to 100 after 24 h of incubation (Figure 5). The growth of the bacteria treated with the Vibrio phage Virtus was inhibited at 7, 5, 3.5, and 2 h post infection for MOIs 0.1, 1, 10, and 100, respectively, and a significant reduction of their titer compared to the untreated control was maintained until the end of the experiment. The titer of *V. harveyi* VH2 was reduced by 40–50% at MOIs 0.1, 1, and 100 compared to the control group over a 24 h period.



Figure 5. In vitro lysis of Vibrio phage Virtus against *V. harveyi* VH2 at MOIs 0.1, 1, 10, and 100 for 24 h. Bacterial growth indicated by the absorbance (OD₆₀₀) read. Error bars were shown for the mean of n = 3.

3.6. Whole Genome Sequencing and Assembly

Genome sequencing of Vibrio phage Virtus resulted in 6,207,226 clean reads with an average read length of 100 bp and 100% correct base calls. The GC content was 47.42%. Genome assembly resulted in a single contig with a minimum genome coverage of 5×. Genome length was 82,960 bp with coverage depth of 7912.21 ×. According to PhageTerm analysis, the Vibrio phage Virtus genome did not have any termini and was found to be terminally redundant and circularly permuted.

3.7. Genomic Features of Vibrio phage Virtus

The genome size of Vibrio phage Virtus is 82,960 bp. The genome arrangement was dense, as suggested by the 1.53 genes per kbp. A total of 127 ORFs were identified with Rapid Annotation using Subsystem Technology (RASTk) server, Glimmer.hmm 2.0, and GeneMark. Comparison of the predicted ORFs showed that all ORFs called by Glimmer.hmm 2.0 and GeneMark were also called by RASTk. Manual inspection of each predicted ORF and gap between ORFs, as well as subsequent alignment in the NCBI nr database, validated that the 127 predicted ORFs were present in the Vibrio phage Virtus genome. No tRNA was found in the genome. A total of 119 ORFs used a start codon of ATG, 6 ORFs used GTG, and 2 used TTG. A search of the NCBI nr database showed that 121 ORFs (95.27%) had significant hits (expected value $\leq 10^{-3}$) with an average similarity of 85.62%. A total of 109 ORFs (85.8%) were determined to have best hits with Vibrio phage vB_VcaS_HC MK559459.1, which infects V. campbellii, while 12 ORFs (9.44%) had the best hits with another six similar Vibrio phages: Vibrio phage 1 (JF713456.1), Vibrio phage Ares1 (MG720309.1), Vibrio phage Thalassa YP (MG649967), Vibrio phage vB_ValS_PJ32 (MT735629.1), Vibrio phage vB_VhaS-a (KX198615.1), and Vibrio phage vB_VpaS_VP-RY-9 (MW411580.1). In addition, protein structural homolog search for the predicted ORFs showed 9 hits in the Gene Ontology database, 7 hits with InterPro, and 24 hits with the NCBI CDD. Overall, 43 (33.8%) ORFs were annotated on the basis of amino acid sequence and protein structural homologies. No homologs of integrase, virulence, or antibiotic-resistance-encoding genes were found in Vibrio phage Virtus. Computational analysis based on phage proteome revealed that there is 92.5% probability that Vibrio phage Virtus follows a lytic lifestyle.

3.8. Genomic Arrangement and Functional Annotations of Vibrio phage Virtus

Generally, the genome of Vibrio phage Virtus did not have any modular arrangement (Figure 6). However, some genes encoding for head and tail proteins (ORF 112, ORF 114, ORF 115, ORF 117, ORF 120) were arranged in subclusters as well as some genes encoding for DNA replication and nucleotide metabolism proteins (ORF 6, ORF 8, ORF 10, ORF 11, ORF 13, ORF 16). Genes that were functionally annotated are shown in Table 2.



Figure 6. Visual representation of the Vibrio phage Virtus genome in which the genome GC content is shown by the inner black line and the GC skew by the inner purple/green line. The predicted ORFs are shown as arrows. The color of the ORFs refers to annotated biochemical function: phage assembly proteins (brown); DNA-replication-, repair-, and recombination-associated proteins (purple); auxiliary metabolic proteins (light green); hypothetical (dark green).

3.8.1. Phage Structural Proteins

Proteins required for phage assembly, including major tail protein (ORF 117), major capsid protein (ORF 112), tail length tape measure protein (ORF 120), tail-completion protein (ORF 116), head completion adaptor (ORF 114), neck protein (ORF 115), portal protein (ORF 33), and minor head protein (ORF34). In addition, the large terminase subunit involved for DNA packaging for tailed phages was identified at ORF 22.

3.8.2. DNA Replication, Repair, and Recombination

Proteins for DNA replication, recombination, and repair were also identified: RecA (ORF 13), HNH endonuclease (ORF 30), DNA polymerases (ORF 32, 39), DNA helicases (ORF 6, 10, 89), DNA primase (ORF 8), and other regulatory elements (ORF 11, 16, 36).

3.8.3. Miscellaneous Proteins

Several transmembrane proteins were detected (ORF 2, 37), including a possible K+dependent Na⁺/Ca⁺ exchanger at ORF 111 (Table 3). Additionally, auxiliary metabolic genes were detected; rubredoxin-type fold protein (ORF 15), a transporter (ORF 31), and a gene coding the pyruvate phosphate dikinase (PPDK), whose product plays a key role in the Embden–Meyerhof–Parnas (EMP) glycolytic pathway (ORF 27).

Table 3. Summary table of Vibrio phage Virtus ORFs that were annotated with relevant information on the basis of significant amino acid sequences and protein structural homologies (E-value $\leq 10^{-3}$).

	Predicted Functions	Start	End	Length	Direction	InterPro	NCBI CDD Best Hit	E-Value
ORF1	Hypothetical protein	44	478	434	Forward		pfam18925 DUF5675	7.64 × 10 ⁻³⁰
ORF2	TMhelix containing protein	480	1019	539	Forward		-	
ORF3	AAA family ATPase	1143	2366	1223	Forward		cl38936 P-loop_NTPase super family	2.77 × 10 ⁻³⁸
ORF4	Winged helix-turn-helix DNA-binding domain protein	2753	4246	1493	Forward		cl41463 PspC_subgroup_2 super family	6.13 × 10 ⁻³
ORF5	Hypothetical protein	4246	4839	593	Forward			
ORF6	Replicative DNA helicase	4824	6272	1448	Forward		cl38936 P-loop_NTPase super family	1.16 × 10 ⁻²⁰
ORF7	Hypothetical protein	6344	6928	584	Forward			
ORF8	DNA primase	6921	7907	986	Forward		cl40740 DnaG super family	1.07 × 10 ⁻⁹
ORF9	Hypothetical protein	7921	8841	920	Forward		-	
ORF10	Putative DNA helicase	8889	10316	1427	Forward		cl34083 SSL2 super family	$1.43\times10^{\scriptscriptstyle -32}$
ORF11	Putative DNA-binding domain protein	10415	10867	452	Forward		cl02600 HTH_MerR-SF super family	1.22 × 10 ⁻³
ORF12	Hypothetical protein	10874	11257	383	Forward			
ORF13	RecA protein	11268	12344	1076	Forward	IPR013765	cl38936 P-loop_NTPase super family	1.94 × 10 ⁻⁵⁰
ORF14	Hypothetical protein	12325	12738	413	Forward		No hit	
ORF15	Rupredoxin-type fold protein	12728	13723	995	Forward		cl37788 NOB1_Zn_bind super family	8.11 × 10 ⁻³
ORF16	Ribonuclease-H-like domain protein	13723	14277	554	Forward		No hit	
ORF17	Hypothetical protein	14386	15360	974	Forward		No hit	
ORF18	Hypothetical protein	15427	15696	269	Forward		No hit	
ORF19	Hypothetical protein	15721	16464	743	Forward		No hit	
ORF20	Hypothetical protein	16542	17273	731	Forward		No hit	
ORF21	Coil containing protein	17288	18097	809	Forward		No hit	
ORF22	Terminase large subunit	18084	20057	1973	Forward		No hit	
ORF23	Hypothetical protein	20076	20318	242	Forward		No hit	
ORF24	Hypothetical protein	20434	20697	263	Forward		No hit	
ORF25	Hypothetical protein	20739	21005	266	Forward		No hit	
ORF26	Coil-containing protein	20995	21417	422	Forward		No hit	
ORF27	Pyruvate, phosphate dikinase	21445	23520	2075	Forward	IPR010121	cl35801 PRK09279 super family	1.21×10^{-180}
ORF28	Hypothetical protein	23525	24013	488	Forward		No hit	
ORF29	Putative protein-tyrosine phosphatase	24006	24551	545	Forward		cl28904 PTP_DSP_cys super family	8.26 × 10 ⁻¹⁷
ORF30	Hypothetical protein	24577	25620	1043	Forward		No hit	
ORF31	Transporter	25631	26473	842	Forward		No hit	

ORF32	DNA polymerase I	26473 28830	2357	Forward	IPR002298	cl34031 PolA super family	8.32 × 10 ⁻⁷¹
ORF33	Portal proten	28832 30697	1865	Forward	IPR006944	cl19194 Phage_portal super family	2.47 × 10 ⁻³⁹
ORF34	Minor head protein	30701 31789	1088	Forward		cl10072 Phage_Mu_F super family	1.69 × 10-3
ORF35	HNH endonuclease	31789 32304	515	Forward		pfam13392 HNH_3	5.16×10^{-10}
ORF36	DNA methylotransferase	32304 32954	650	Forward	IPR007757	cl01947 MT-A70 super family	2.11 × 10 ⁻²⁸
ORF37	TMhelix containing protein	33138 33554	416	Forward		No hit	
ORF38	Hypothetical protein	33566 34021	455	Forward		No hit	
ORF39	Putative DNA polymerase I	34149 34544	395	Forward		cl02626 DNA_pol_A super family	1.35 × 10-5
ORF40	Hypothetical protein	34547 34732	185	Forward		No hit	
ORF41	Hypothetical protein	34755 35123	368	Forward		No hit	
ORF42	Hypothetical protein	35134 35394	260	Forward		No hit	
ORF43	Hypothetical protein	35401 35889	488	Forward		No hit	
ORF44	Hypothetical protein	35892 36176	284	Forward		No hit	
ORF45	Hypothetical protein	36182 36481	299	Forward		No hit	
ORF46	Hypothetical protein	36566 36874	308	Forward		No hit	
ORF47	Hypothetical protein	36926 38263	1337	Forward		No hit	
ORF48	Hypothetical protein	38309 38902	593	Forward		No hit	
ORF49	SEC-C motif protein	38928 39290	362	Forward		pfam02810 SEC-C	2.55×10^{-6}
ORF50	Hypothetical protein	39299 39469	170	Forward		No hit	
	putative zinc- or iron-						
ORF51	chelating-domain-containing	39466 39927	461	Forward		No hit	
	protein						
ORF52	Hypothetical protein	39911 40432	521	Forward		No hit	
ORF53	Hypothetical protein	40432 40674	242	Forward		No hit	
ORF54	Hypothetical protein	40674 40853	179	Forward		No hit	
ORF55	Hypothetical protein	41325 41765	440	Forward		No hit	
ORF56	Hypothetical protein	41753 41956	203	Forward		No hit	
ORF57	Hypothetical protein	41953 42135	182	Forward		No hit	
ORF58	Hypothetical protein	42132 42845	713	Forward		No hit	
ORF59	Coil containing protein	42826 43638	812	Forward		No hit	
ORF60	Hypothetical protein	43644 43952	308	Forward		No hit	
ORF61	Hypothetical protein	44081 44587	506	Forward		No hit	
ORF62	DUF550-domain-containing protein	44651 45433	782	Forward		cl04522 DUF550 super family	5.51 × 10 ⁻⁷
ORF63	Hypothetical protein	45417 45575	158	Forward		No hit	
ORF64	Hypothetical protein	45683 45877	194	Forward		No hit	
ORF65	Hypothetical protein	45892 46221	329	Forward		No hit	
ORF66	Hypothetical protein	46274 46669	395	Forward		No hit	
ORF67	Hypothetical protein	46746 47126	380	Forward		No hit	
ORF68	Hypothetical protein	47178 47639	461	Forward		No hit	
ORF69	Hypothetical protein	47636 47962	326	Forward		No hit	
ORF70	Hypothetical protein	47946 48461	515	Forward		No hit	
ORF71	Hypothetical protein	48448 48855	407	Forward		No hit	
ORF72	Hypothetical protein	48861 49031	170	Forward		No hit	
ORF73	Hypothetical protein	49427 49699	272	Forward		No hit	
ORF74	Hypothetical protein	49714 49920	206	Forward		No hit	
ORF75	Hypothetical protein	50017 50349	332	Forward		No hit	
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ORF76	Hypothetical protein	50408 50743	335	Forward		No hit	
ORF77	Hypothetical protein	50800 51246	446	Forward		No hit	
ORF78	Hypothetical protein	51236 51598	362	Forward		No hit	
ORF79	Phage protein (ACLAME 851)	51644 52147	503	Forward		No hit	
ORF80	Hypothetical protein	52281 52481	200	Forward		No hit	
ORF81	VHS1018	52523 52774	251	Forward		No hit	
ORF82	Coil-containing protein	52822 53127	305	Forward		No hit	
ORF83	Hypothetical protein	53131 53367	236	Forward		No hit	
ORF84	Hypothetical protein	53380 53580	200	Forward		No hit	
ORF85	Hypothetical protein	53580 53723	143	Forward		No hit	
ORF86	Hypothetical protein	53847 54092	245	Forward		No hit	
ORF87	Hypothetical protein	54108 54341	233	Forward		No hit	
ORF88	Hypothetical protein	54468 54668	200	Forward		No hit	
	JI I I I I I					cl28899 DEAD-	
ORF89	DNA Helicase	54715 54993	278	Forward		like helicase N super	8.48 × 10 ⁻³
011107	21	01110 01170		1 of Ward		family	0.10 10
ORF90	Hypothetical protein	54990 55247	257	Forward		No hit	
ORF91	Hypothetical protein	55244 56659	1415	Forward		No hit	
ORF92	Hypothetical protein	56753 57649	896	Forward		No hit	
ORF93	Hypothetical protein	57724 58683	959	Forward		No hit	
ORF94	Hypothetical protein	58814 59128	314	Forward		No hit	
ORF95	Coil-containing protein	59106 59690	584	Forward		No hit	
ORF96	Hypothetical protein	59766 60128	362	Forward		No hit	
ORF07	Hypothetical protein	60179 60571	302	Forward		No hit	
ODE00	Hypothetical protein	60568 61002	124	Forward		No hit	
OPE00	Hypothetical protein	61070 61422	252	Forward		No hit	
ORF 99	Hypothetical protein	61079 61432	353 404	Forward		No hit	
ORF100 ODE101	Hypothetical protein	61008 62602	494	Forward		No hit	
ORF101	Hypothetical protein	61996 62695	093 201	Forward		No liit	
ORF102	Hypothetical protein	62697 62976	201	Forward		No lut	
ORF103	Hypothetical protein	63124 63414	290	Forward		No hit	
OKF104	Hypothetical protein	63428 63730	302	Forward		No nit	
OKF105	Hypothetical protein	63742 64266	524 979	Forward		No nit	
ORF106	Hypothetical protein	64266 64538	272	Forward		No hit	
ORF107	Coil-containing protein	64607 65395	788	Forward		No hit	
ORF108	Hypothetical protein	65388 67358	1970	Forward		No hit	
ORF109	Hypothetical protein	67424 67774	350	Forward		No hit	
ORF110	XkdF	67903 68514	611	Forward		cl24270 Peptidase_S78_2 super family	2.76 × 10 ⁻¹⁹
ORF111	Hypothetical protein	68517 69686	1169	Forward		cl36772 2A1904 super family	2.50 × 10 ⁻⁵
ORF112	Major capsid protein	69767 70717	950	Forward	IPR024455	cl27082 Phage_capsid super family	4.87×10^{-5}
ORF113	Coil-containing protein	70795 71064	269	Forward		No hit	
ORF114	Head completion adaptor	71080 71724	644	Forward		No hit	
ORF115	Neck protein	71721 72185	464	Forward		No hit	
ORF116	Tail-completion protein	72182 72664	482	Forward		No hit	
ORF117	Major tail protein	72718 73512	794	Forward	IPR016893	No hit	
ORF118	Hypothetical protein	73601 74035	434	Forward	10 _ 00 / 0	No hit	
ORF119	VHS1060 protein	74122 74235	113	Forward		No hit	
ORF120	Tail length tape measure protein	74241 78374	4133	Forward		pfam10145 PhageMin_Tail	1.22 × 10 ⁻²²

ORF121	Hypothetical protein	78371 78757	386	Forward	No hit
ORF122	Hypothetical protein	78767 79756	989	Forward	No hit
ORF123	Hypothetical protein	79766 80659	893	Forward	No hit
ORF124	Hypothetical protein	80662 81960	1298	Forward	No hit
ORF125	Hypothetical protein	81963 82487	524	Forward	No hit
ORF126	Hypothetical protein	82488 82790	302	Forward	No hit
ORF127	Hypothetical protein	82787 82960	173	Forward	No hit

3.9. Genomic Synteny of Vibrio phage Virtus with Other Similar Phages

Pairwise alignment between Vibrio phage Virtus and vB VcaS HC showed that they have a genetic identity of 94.2% (Figure 7). The areas coding the proteins required for the phage structural assembly were generally conserved; however, there were significant nucleotide disagreements in genes who relate to DNA replication and nucleotide metabolism, i.e., ORF 10, ORF 11, ORF 15, and in areas coding miscellaneous proteins. A gene coding a homing endonuclease (ORF 35), two genes coding hypothetical proteins (ORF 86, ORF 87), and a non-coding area were present in the Vibrio Virtus genome but not in vB_VcaS_HC. The Vibrio phage Virtus had the highest degree of genomic synteny with vB_VcaS_HC (Figure 8) sharing eight collinear blocks. The longest shared collinear block had a sequence length almost 20,000 bp. Furthermore, the common collinear blocks had similar genomic arrangements and shared high DNA sequence similarities. Alignment with another three similar vibrio phages: Vibrio phage 1, Vibrio phage Ares1, and Vibrio alginolyticus phage vB_ValS_PJ32, also showed eight shared collinear blocks of similar length with high genomic synteny and sequence similarities. On the contrary, both Vibrio phage Virtus and vB_VpaS_VP-RY-9 shared six collinear blocks, but with very low sequence similarities.



Figure 7. Pairwise alignment of Vibrio phage Virtus with vB_VcaS_HC. From the top, the first bar represents mean pairwise identity over all nucleotide pairs (green: 100% identity, brown: at least 30% and under 100% identity, red: below 30% identity). Predicted ORFs are shown by arrows. The color of the ORFs refers to annotated biochemical function; phage assembly proteins (orange), DNA-replication-, repair-, and recombination-associated proteins (purple); auxiliary metabolic proteins (blue); hypothetical (dark green).



Vibrio phage v8 VprS VP-RY-9. MW411580

Figure 8. Whole genome alignment with progressive MAUVE of Vibrio phage Virtus with similar phages. From the top is Vibrio phage Virtus, vB_VcaS_HC, Vibrio phage 1, Vibrio phage Ares1, vB_ValS_PJ32, and vB_VpaS_VP-RY-9. The colored collinear blocks indicate homologous regions between genome sequences, while the height of the similarity profile in the collinear blocks indicate average level of conservation in the regions of the genome sequence. Inverted blocks indicate homologous regions that align in the complement orientation.

3.10. Phylogenetic Analysis

Wide genome proteomic tree analysis confirmed that Vibrio phage Virtus belongs to the *Siphoviridae* taxonomic family (Figure 9). In addition, Vibrio phage Virtus was predicted to infect hosts from the Gammaproteobacteria class, which includes the Vibrionaceae family.

Phylogeny using large terminase subunits of vibrio phages (Figure 10) showed that Vibrio phage Virtus has a recent common ancestor with vB_VcaS_HC. Moreover, Vibrio phage Virtus has a high bootstrap support (100%) with Vibrio phage 1 and Vibrio phage Ares1, indicating that they share a common evolutionary history. In addition, the branch length is proportional to the amount of evolutionary divergence, and hence Vibrio phage Virtus and vB_VcaS_HC phages share a similar number of amino acid substitutions in their large terminase subunit since diverging from their common ancestor.



Figure 9. Determination of taxa and host group for Vibrio phage Virtus according to the proteomic tree produced by VIPTree. Vibrio phage Virtus was determined to belong to the Siphoviridae family and to infect Gammaproteobacteria group (red star and line). Vibrio phage Virtus (asterisk) proteome was compared with 4892 dsDNA phage proteomes. The branch length scale was calculated as log values. The inner and outer ring indicate the taxonomic virus family and host group, respectively.



Figure 10. Phylogenetic tree of Vibrio phage Virtus with other Vibrio phages. The large terminase subunits of similar phages were downloaded from the NCBI database and aligned using MUSCLE,

and a maximum likelihood (bootstrap = 1000) phylogenetic tree was constructed using MEGA X. The tree was visualized using the Interactive Tree of Life (ITOL). The bootstrap support value is denoted in each branch.

3.11. In Vivo Phage Therapy in Gilthead Seabream Larvae

In vivo phage therapy trials with gilthead seabream larvae were conducted to assess the efficacy of Vibrio phage Virtus in controlling *Vibrio harveyi* VH2 (Figure 11). VH2 was found to be very pathogenic, significantly reducing the survival of larvae to just 6% compared to the control group in which 92% of larvae survived during the 5-day trial (X^2 (1, 192) = 148.6, p < 0.001). Survival of gilthead seabream larvae was significantly increased when treated with Vibrio phage Virtus at a MOI of 10 compared to the group treated with V. *harveyi* VH2 (X^2 (1, 190) = 33.4, p < 0.001). Moreover, no significant reduction was observed between the single dose and the two doses of treatment (data not shown). The phage control group (no bacteria added) also had no significant difference compared to the control (X^2 (1, 191) = 0.07865, p = 0.7791), indicating the safety of the phage suspension and possibly the absence of endotoxins (Figure 10).



Figure 11. Survival of gilthead seabream larvae infected with *V. harveyi* VH2 in an experimental phage therapy trial during a period of 5 days. Gilthead seabream larvae that were infected with VH2 were inoculated with Vibrio phage Virtus with different multiplicities of infection (MOI) at 2 h post-infection. *Phaeobacter piscinae* S26 were used to evaluate the effect of non-pathogenic bacteria at the same concentration to fish larvae.

4. Discussion

Vibrio harveyi outbreaks are increasing, as climate change becomes more imminent, threatening a broad range of marine organisms such as abalones, shrimps, corals, and various fishes [52–55] and leading to severe economic and production losses in aquaculture worldwide [56,57]. The biggest problem associated with these outbreaks is that many strains are highly resistant to antibiotic treatments [58–60]. Because of this, an increasing number of studies aiming to control vibriosis have been conducted by employing phages as therapeutic agents. To our knowledge, 21 bacteriophages have been previously isolated against *V. harveyi*, including 16 siphoviruses [61–66], 4 myoviruses [17,62,67,68], and 1 podovirus [69]. Two bacteriophages, VHML and Siphophage 1 VHS1, were found to be temperate [66,68], while the rest are considered to be lytic. Here, we isolated a novel lytic

bacteriophage, Vibrio phage Virtus, against *V. harveyi* VH2, and tested its efficacy as a potential candidate for therapy.

Whole sequence homolog search and pairwise alignment revealed that a Vibrio phage, vB_VcaS_HC, which infects Vibrio campbelii, shared a high similarity with Vibrio phage Virtus. Both methods yielded 94.2% genetic identity between the phages. The threshold to distinguish two different species is 95%, and thus Vibrio phage Virtus probably belongs to a novel species of the Siphoviridae family [70]. Both phages shared similar genomic arrangements with nucleotide similarities according to genomic synteny analysis. Interestingly, Vibrio phage Virtus was isolated in Heraklion, Greece, while vB_VcaS_HC was isolated in Qingdao, China. This suggests that this particular phage has a rather wide geographical distribution. Phage geographical distribution depends on the abundance and metabolic state of the host, since phage survival depends on the presence of susceptible hosts [71]. Thus, a phage with wide geographical distribution indicates that either its host is ubiquitous or that the phage has a broad host range. Moreover, specific phage traits such as latent period and burst size may also influence phage dispersal and what geographical patterns it follows [72]. High burst size and long latent period improve the probability of a successful dispersal and are indicatives of a cosmopolitan phage. In our case, Vibrio phage Virtus was capable of infecting hosts from different species, unlike most phages, which are usually species-specific [73]. On the contrary, vB_VcaS_HC had a very narrow host range. However, no direct comparison can be made, since the host range is related to the bacterial strains used, which were different in the two studies. It is suggested that a broad host range is an important evolutionary trait for phages [74], although often with a decreased virulence as a cost, which reflects the antagonistic pleiotropy [75]. However, phages with a broad spectrum of hosts are desirable for therapy, especially for pathogens that are abundant and diverse such as the vibrios [76,77].

Vibrio phage Virtus was found to have an unusually large burst size. To date, only a few phages have been reported to have such large burst sizes in all of dsDNA phages [22,78,79]. The eclipse period was estimated to be longer than the average of most phages, which is 5–15 min [80], and in combination with the long latent period, could possibly lead to high virion productions due to multiple reproduction cycles [81]. However, other factors may also affect the burst size, including the host metabolic activity, ambient environment, and the protein synthesis machinery of the host bacteria [22,82,83], and hence the molecular mechanism associated with the large burst size needs further investigation.

Horizontal gene transfer (HGT) occurs regularly between phages and bacteria populations either by generalized or specialized transductions [69,84,85]. Vibrio phages have occasionally been associated with inducing virulence in their hosts [68,86], and hence a comprehensive profiling of their genomic traits is required before proceeding to therapeutic application. Only 4 of the 21 phages against V. harveyi that were isolated in previous studies have been sequenced and characterized genomically [17,63,86,87]. The genome sizes of these four phages vary between 48 and 286 kbp, including a jumbo bacteriophage Vb_VhaM_pir03. In the Vibrio phage Virtus genome, no integrase, virulence, or antibioticresistance-encoding genes were detected. Moreover, no prophage induction occurred when host mutants with a phage-resistant phenotype were exposed to UV radiation, further supporting the lytic lifestyle of the novel phage. The Vibrio phage Virtus genome is absent of any termini, is circularly permuted, and is terminally redundant, which suggests a headful packaging mechanism [88,89]. An auxiliary metabolic gene coding the pyruvate phosphate dikinase (PPDK) whose product plays a key role in the Embden–Meyerhof– Parnas (EMP) glycolytic pathway was present [90]. PPDK is not commonly found in phages, yet it has been reported before in some vibrio siphoviruses [10,62,84]. Phages that contain auxiliary metabolic genes have mechanisms to manipulate host metabolism into their own benefit [91]. For example, the lytic bacteriophage KVP40 genome includes ORFs that encode proteins that facilitate precursor transport and synthesis of NAD⁺ in the pyridine nucleotide salvage pathway [92]. Moreover, studies have shown that marine viruses genomes, isolated in nutrient-limited environments, were rich in auxiliary metabolic genes compared to the ones isolated in nutrient-rich environments [93], indicating a strong association between phage auxiliary metabolic genes and host resource uptake. Genes encoding phosphorus uptake regulation such as PhoH have been found in Vibrio phages [94], and it has been suggested that they are being used in order to force the host to increase phosphorus acquisition in order to be used during phage DNA replication. Taking this into consideration, it is likely that the phage PPDK gene is co-expressed during infection, increasing host energy uptake, which is ultimately directed to the production of more virions. The presence of auxiliary metabolic genes in the Vibrio phage Virtus genome can also be linked to a possible widespread distribution, since they can lead to a higher burst size and can thereby expand dispersal [72,95].

As shown in stability assays, Vibrio phage Virtus can withstand a wide range of temperatures and pH values, which is very practical for phage therapy. In addition, we showed that Vibrio phage Virtus can be completely inactivated with various organic solvents if this is required to reduce the risk of unwanted dissemination to the environment. In vitro assay showed that Vibrio phage Virtus was able to efficiently reduce the host bacterial populations at different MOIs. The fact that it was able to lyse the bacteria in low MOIs offers a practical advantage for the application in the aquaculture settings, since the required phage quantity is relatively low. However, after 15 h, the host bacterial population started to rise again, suggesting the emergence of resistance, possibly due to the intense selective pressure [96]. Phage resistance is a concerning issue in phage therapy [97] since bacteria populations have various protection mechanisms against phages [98]. The combination of different phages, phage cocktails, has been suggested as a workaround, a practice that has yielded promising results [99–101].

Several studies have shown the successful in vivo application of phages to treat vibriosis in various animal models [17,61,102]]. Wang et al. has shown that vB_VhaS-tm managed to improve survival of abalone by 70% in seven days, while Misol et al. showed that vB_VhaM_pir03 improved Artemia nauplii survival by 15–20% in 48 h [17,102]. Moreover, the survival of giant tiger prawn (Penaeus monodon) was immensely higher when treated with phages compared to antibiotic treatment and the control, as shown by Vinod et al. [61]. Here, in vivo phage therapy trials in gilthead seabream larvae showed that a single dose of Vibrio phage Virtus significantly improved the survival of the larvae by 35% compared to the untreated population. As Levin and Bull [103] suggested, phages decrease bacterial load enough to be eliminated by the fish immune system. In this case, we suggest, that Vibrio phage Virtus decreases the bacterial population in levels that are no longer pathogenic by reducing the colonization of Vibrio harveyi in the larvae skin [104]. Moreover, we showed that a second dose of Vibrio phage Virtus made no difference in the survival of the larvae. It is possible that the phage and the bacteria population had already reached an equilibrium, known as the carrier state. In this state, bacterial populations are heterogenous, as they contain subpopulations in which phages are stably maintained within the host rather than committing to lysis or subpopulations that are resistant to phages and maintain the sensitive population [105]. The molecular mechanism behind the resistance of carrier state bacterial subpopulations is very intriguing and worthy of further investigation. Preliminary experiments for the characterization of resistant mutants (not included here) of host developed in this study revealed a fitness cost to the bacteria (data not shown), which suggests that the defense mechanism is more likely related to cell surface modifications [106,107]. However, this is a mere speculation, since other defense mechanisms have been reported before, such as the acquisition of spacers matching phage genomic material [105]. The emergence of resistance could limit the therapeutic potential of Vibrio phage Virtus, and therefore its synergistic effect with other Vibrio phages is being pursued. However, no fully characterized phage infecting V. harveyi VH2 was available at the time of the study, and thus a comparative or a synergistic treatment with another phage was not possible. Carrier state often results in less pathogenic bacteria populations, as shown in previous studies [105,108]. Furthermore, no significant mortalities were detected in the group treated with only the phage suspensions, indicating the safety of Vibrio phage Virtus to the fish larvae. Although gilthead seabream larvae have been used as an in vivo model to study the therapeutic efficacy of the Vibrio phage Virtus, it should be noted that given the importance of this fish species for the Mediterranean aquaculture [109] and the high prevalence of vibriosis caused by *V. harveyi* [7], the practical usability of Virtus but also similar phages in commercial aquaculture is evident. Of course, there are several issues that remain to be resolved before phage therapy for aquaculture becomes a common practice including regulatory, mass production of phages and resistance development by the bacteria.

In conclusion, we present a comprehensive genomic and biological characterization of Vibrio phage Virtus as a potential and suitable candidate for the biocontrol of *Vibrio harveyi* infections. High virion production and broad host range are the main biological characteristics of Vibrio phage Virtus. As for its genomic profile, Vibrio phage Virtus lacks genes associated with virulence, antibiotic resistance, and transduction potential. On the contrary, its genome contains genes with multiple diverse functions, i.e., PPDK gene, that may contribute to the efficacy of Vibrio phage Virtus. An in vitro assay showed that Vibrio phage Virtus was able to control the host population even at very low MOIs, which favors its practical use in applied therapy. Ultimately, the survival of gilthead seabream larvae challenged with *V. harveyi* was significantly increased when treated with Vibrio phage Virtus, further supporting its effectiveness.

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