

Biomimetic hydroxyapatite nanocrystals are an active carrier for *Salmonella* bacteriophages

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Purpose: The use of bacteriophages represents a valid alternative to conventional antimicrobial treatments, overcoming the widespread bacterial antibiotic resistance phenomenon. In this work, we evaluated whether biomimetic hydroxyapatite (HA) nanocrystals are able to enhance some properties of bacteriophages. The final goal of this study was to demonstrate that biomimetic HA nanocrystals can be used for bacteriophage delivery in the context of bacterial infections, and contribute – at the same time – to enhance some of the biological properties of the same bacteriophages such as stability, preservation, antimicrobial activity, and so on.

Materials and methods: Phage isolation and characterization were carried out by using Mitomycin C and following double-layer agar technique. The biomimetic HA water suspension was synthesized in order to obtain nanocrystals with plate-like morphology and nanometric dimensions. The interaction of phages with the HA was investigated by dynamic light scattering and Zeta potential analyses. The cytotoxicity and intracellular killing activities of the phage–HA complex were evaluated in human hepatocellular carcinoma HepG2 cells. The bacterial inhibition capacity of the complex was assessed on chicken minced meat samples infected with *Salmonella* Rissen.

Results: Our data highlighted that the biomimetic HA nanocrystal–bacteriophage complex was more stable and more effective than phages alone in all tested experimental conditions.

Conclusion: Our results evidenced the important contribution of biomimetic HA nanocrystals: they act as an excellent carrier for bacteriophage delivery and enhance its biological characteristics. This study confirmed the significant role of the mineral HA when it is complexed with biological entities like bacteriophages, as it has been shown for molecules such as lactoferrin.

Keywords: biomimetic hydroxyapatite nanocrystals, *Salmonella* Rissen, bacteriophages, antibiotic resistance

Introduction

Inappropriate abuse of antibiotics has led to the development of drug-resistant microorganisms.¹ Antibiotic resistance is a constantly evolving phenomenon and represents a serious problem with high death toll and a substantial economic impact worldwide; it complicates patient management and treatment strategy and prolongs hospital stays. In Europe antibiotic-resistant bacteria infect 4 million people every year.^{2,3} Therefore, therapies that can serve as an alternative to antibiotics need to be developed.

A valid and alternative approach to solve the widespread phenomenon of “antibiotic resistance” of different pathogens such as *Salmonella*,⁴ *Staphylococcus aureus*,⁵ *Escherichia coli*,⁶ and *Streptococcus*⁷ – in addition to well-known antimicrobial peptides⁸ – is the use of bacteriophages or phages. Bacteriophages are the most abundant viruses

found in the biosphere;^{9,10} they grow quickly and exponentially, and they are efficacy since the first interaction with bacteria.¹¹ At the same time, phages have some disadvantages such as low stability over time and low resistance (or short half-life) in acidic environments such as that of the stomach.^{12,13}

These limitations could be overcome by stabilizing the phages by combining them with nanoparticles of different materials such as carbon, silica, metal oxide, graphene, and hydroxyapatite (HA). HA, in particular, has been frequently used in several studies to stabilize, protect, and deliver molecules or radionuclides.^{14–22}

HA represents the major components of bone, tooth, and cartilaginous tissues. It possesses several properties such as biocompatibility, biomimetic dimensions, osteoconductivity, and degradability, which make it suitable for several applications^{16,23} when combined with different biological molecules.¹⁹ Indeed, HA nanocrystals have been successfully employed to build bone scaffolds and implant coating materials and vehicles for drug targeting.^{24–29} Furthermore, HA nanocrystals have low toxicity and remarkable stability, as reported in our previous studies.³⁰

In industrialized countries, infections acquired by food and water represent a big concern for public health, and about 50% of foodborne infections in humans are caused by *Salmonella* spp.³¹ Despite control measures and monitoring carried out by the healthcare authorities, cases of *Salmonella* contaminations are still very frequent.³²

Salmonella serovars such as Enteritidis and Typhimurium are accountable for most of the salmonellosis cases.³³ In recent years, infections by other serovars such as Kentucky and Rissen became more frequent; the latter *Salmonella* strain, especially, has been isolated from pork and chicken products and in human and swineherds gastrointestinal tract.^{34,35}

Based on these considerations, we decided to investigate whether the biomimetic HA nanocrystals – which mimic the natural bone mineral – could interact with *Salmonella* bacteriophages and whether the newly developed complex could control *Salmonella* bacterial infection.

Materials and methods

Bacteria

All the *Salmonella* spp. strains used in this study were provided by the Istituto Zooprofilattico Sperimentale Del Mezzogiorno (Portici, Naples, Italy). Among these, *S. Rissen* was selected as reference strain for bacteriophage (phage) isolation.

Phage isolation

Phage (SR ϕ 1) from *S. Rissen* was isolated as described by Capparelli et al.³⁶ Briefly, *S. Rissen* was grown in 5 mL of Luria Bertani broth (LB; Sigma-Aldrich Co., St Louis, MO, USA) and, when the culture reached the exponential growth phase, Mitomycin C (1 μ g/mL) was added. Then, the suspension was incubated at 37°C for 30 minutes. After incubation, it was centrifuged twice at 5,000 rpm for 10 minutes; each time the supernatant was discarded and the pellet resuspended in 5 mL of LB in order to remove any residue of Mitomycin C. Later, the suspension was incubated at 37°C for 4 hours. Then another centrifugation step was carried out but this time, the supernatant – containing bacteriophages – was collected, filtered through a 0.22 μ m membrane, and screened for the presence of phages using the double-layer agar (DLA) method. The last step included a lower nutrient agar layer and an upper soft agar layer (4 mL of 0.7% bacteriological agar [Sigma-Aldrich Co.] mixed with 10⁷ colony forming unit [CFU]/mL of bacterial strain used for phage isolation and previously filtered bacteriophage solution). After overnight incubation at 37°C, the plates were observed for the presence of clear zone (plaque formation) over the surface of the double agar plate. The phages isolated were stored at –20°C in buffer SM.³⁷ However, the pellet was streaked onto *Salmonella* Chromogenic Agar Base (CM1007, Oxoid; Thermo Fisher Scientific, Waltham, MA, USA) to confirm the presence of *Salmonella* spp. bacteria.

The titer of phage, expressed as plaque forming units (PFUs), was evaluated by using the DLA technique as reported by Sambrook et al.³⁸

Phage host range

Host range of phages was evaluated against 14 different *Salmonella* strains by the overlay method.³⁹ Briefly, the test consisted of spotting 100 μ L of SR ϕ 1 on the surface of a double agar layer as reported above.

Multiplicity of infection (MOI) of phage

MOI is the ratio of virus particles to host cells.⁴⁰ To establish the best MOI, *S. Rissen* strain was grown in LB at 37°C till 10⁴ CFU/mL. Later, 100 μ L of bacterial suspension was treated with 100 μ L of SR ϕ 1 at different ratios, ranging from 10^{–3} to 10³ PFU/CFU, in a 96-well microplate. Positive and negative controls were represented by a mixture containing 100 μ L of *S. Rissen* (10⁴ CFU/mL) plus 100 μ L of SM buffer or 100 μ L of LB broth plus 100 μ L of SM buffer, respectively. After 16–18 hours of incubation at 37°C, the

optical density was measured at OD 600 nm to determine the optimal MOI. This last parameter shows that the lower phage titer is able to kill the majority of the bacteria, and it was used for all the subsequent experiments.

One-step growth curve

The “burst size” and phage life cycle were evaluated by the one-step growth assay. The culture of *S. Rissen* (10^8 CFU/mL), in exponential growth phase, was infected with phage according to the optimal MOI (previously selected). The mixture was incubated at 37°C for 5 minutes to favor phage adsorption, and later, it was serially diluted up to 10^{-4} in 20 mL of LB. The above mixture was incubated at 37°C for 90 minutes, and 100 μ L of sample was taken every 5 minutes and plated, during the whole incubation period. Latent period was the interval between the beginning of the adsorption (not including 5 minutes pre-incubation) and the onset of the first burst (corresponding to the initial rise in phage titer). Burst size was the ratio between the final count of released phage particles and the initial amount of infected bacterial cells.

Electron microscopic analysis

SR ϕ 1 (10^8 PFU/mL) was purified by CsCl density gradient ultracentrifugation and dialyzed against SM buffer overnight at 4°C.⁴¹ Phage particles were negatively stained with 2% of phosphotungstic acid (pH 7.2) for 20 minutes. Later, phages were observed by using a Philips EM 300 electron microscope.

Phage phylogenetic analysis

Phylogenetic analysis was performed using the maximum likelihood method.⁴² Data alignment was carried out with Blosum 65 (gap open penalty = 11; gap extension penalty = 3) and tree was built by using Jukes-Cantor and UPGMA models.

Phage genome sequencing, assembly, and annotation

Genomic DNA was quantified using the Qubit dsDNA BR Assay Kit (Thermo Fisher Scientific); DNA purity was assessed with a Nanodrop (Thermo Fisher Scientific), and DNA size was determined with a 2200 Tape Station Instrument (Agilent Technologies, Santa Clara, CA, USA). Illumina libraries were produced starting from 1 μ g of genomic DNA, which was sheared using the Covaris S220 instrument (Covaris Inc. Woburn, MA, USA) and the TruSeq DNA

Sample Prep Kit (Illumina, San Diego, CA, USA) following the manufacturer’s guidelines. Sequencing was performed on a NextSeq500 instrument with the 150-nt paired-end protocol (Illumina) according to the manufacturer’s guidelines. Illumina reads underwent quality filtering and trimming using Sickle and were quality corrected with Bayes Hammer before being assembled de novo. Genomes were assembled de novo from Illumina reads using SPAdes 2.9.0 with multiple k-mer combinations. The resultant contigs were scaffolded using SSPACE 3.0. Five micrograms of high-molecular-weight genomic DNA (peak >60 kb) was used to prepare ~20 kb-insert single-molecule real-time (SMRT)-bell libraries according to the manufacturer’s guidelines (Pacific Biosciences, Menlo Park, CA, USA). The library templates were sequenced using the SMRT sequencing technology on a PacBio RSII sequencer (Pacific Biosciences) at Macrogen, Inc (Korea). PacBio subreads were extracted using Bash5tools (version 0.8.0), filtrated, and assembled de novo using Falcon-Integrate with the suggested settings for bacterial genome. The assembled genome sequence was polished by Quiver v 0.9.2, and gene annotation was performed using RAST web service (<http://rast.nmpdr.org/>).⁴³

Biomimetic HA nanocrystal synthesis and characterization

Biomimetic HA nanocrystals were produced as described by Nocerino et al.⁴⁴ Briefly, HA nanocrystals were precipitated from an aqueous solution of $(\text{CH}_3\text{COO})_2\text{Ca}$ (75 mM) by slow addition (one drop per second) of an aqueous solution of H_3PO_4 (50 mM), keeping the pH constant at 10 (by the addition of $(\text{NH}_4)\text{OH}$ solution). The synthesis was performed at room temperature. After this last process, the suspension of HA was washed with distilled water in order to remove ammonium ions and favor the interaction between nanocrystals.

Transmission electron microscopy (TEM) investigation was carried out using a 1200 EX microscope, linked to X-ray analysis detectors and a 3010 UHR operating at 300 kV (JEOL Ltd, Tokyo, Japan). Few droplets of the samples (in ultrapure water) were deposited on perforated carbon foils supported on conventional copper microgrids. The surface area was determined using a Sorpty 1750 instrument (Carlo Erba Reagents S.r.l., Milan, Italy) using N_2 absorption at 77 K.⁴⁵

Synthesis of complex HA–SR ϕ 1

The HA–SR ϕ 1 complex was prepared by mixing 10 mL of HA (100 mg/mL) with 10 mL of SR ϕ 1 (10^7 PFU/mL) under stirring wad. The suspension was stable, and no precipitation

phenomena were observed during the synthesis process. Later, aliquots of 1 mL of the mixture were incubated – under shaking – at room temperature for the following time intervals: 30, 90, 180, and 300 minutes and 24 hours. After proper incubation, the sample was centrifuged, and the pellet was suspended in SM buffer. The concentration of the active phage particles was evaluated by the DLA method. After overnight incubation, the optimal incubation time was selected according to the results. Next, other tests were carried out by maintaining constant concentration of the HA but increasing the bacteriophage titer. The results showed that, by increasing the phage titer, the concentration of the active phage particles estimated by the DLA method, after the optimal time of incubation, remained constant at 10^7 PFU/mL (the titer used for the above complex synthesis) (data not shown). Based on these data, we evidenced that the maximum loading capacity of the HA used for this study was 10^7 PFU/mL of bacteriophages.

Complex HA–SR ϕ 1 characterization Zeta potential, dynamic light scattering (DLS) analysis, and pH stability

SR ϕ 1, HA–SR ϕ 1, and HA were analyzed for the measurement of zeta potential and DLS in appropriate disposable folded capillary cells (DTS1070; Malvern Instruments, Malvern, UK) using a Zetasizer Nano ZS (Malvern Instruments). For the DLS analysis, the instrument was, in addition, equipped with a 633 nm He–Ne laser and an avalanche photodiode detector placed at a detection angle of 173° . Each analysis was carried out in triplicate for three independent experiments. The analysis temperature was 25°C and about 1 mL of sample (at pH value of 7) was used for the test. The results were examined, and, for each sample, the zeta potential average and DLS measurement value were determined.

To estimate the stability of SR ϕ 1 and HA–SR ϕ 1 over time, the titer of phage – or mixed with HA – was evaluated for 2 successive months, at weekly intervals. During this time, the samples were stored at $+4^\circ\text{C}$.⁴⁶

In addition, the effects of an acid or alkaline pH, on SR ϕ 1 or HA–SR ϕ 1 complex, were evaluated by mimicking different pH conditions (ranging from 2 to 10). Briefly, the phage suspensions were incubated at 37°C for 1 hour in buffer SM at the following pH values: 2, 4, 7, and 10. After incubation, the phage titer was estimated by the DLA method as reported by Jun et al.⁴⁷ Each assay was performed in triplicate.

X-ray analysis

X-ray powder diffraction (XPD) patterns were obtained at room temperature by using a Rigaku RINT2500 rotating

anode laboratory diffractometer (50 kV, 200 mA) equipped with the silicon strip Rigaku D/teX Ultra detector. An asymmetric Johansson Ge (111) crystal was used to select the monochromatic Cu $K\alpha_1$ radiation ($\lambda=1.54056 \text{ \AA}$). The measurements were carried out in transmission geometry by introducing the samples in a Lindemann glass capillary of 0.3 mm diameter. The XPD patterns were first indexed by the program QUALX⁴⁸ by matching the XPD patterns with the ICSD database. They were further analyzed by using a whole-profile Rietveld-based fitting program⁴⁹ to determine the crystalline domain size as follows:

1. The instrumental resolution function (IRF) was evaluated by fitting the XPD pattern of a LaB6 NIST standard recorded under the same experimental conditions as those used for measuring the samples. The IRF data file was provided separately to the program in order to allow subsequent refinement of the diffraction patterns of the samples.
2. The crystal structure of the samples, once determined, was input into the program and refined. The inhomogeneous peak broadening of the diffraction peaks was described by a phenomenological model based on a modified Scherrer formula:

$$\beta_{h,k,l} = \frac{\lambda}{h \cos \theta} = \frac{\lambda}{\cos \theta} \sum_{imp} a_{imp} y_{imp} (\theta_h, \Phi_h)$$

where $\beta_{h,k,l}$ represent the size contribution to the integral width of the (h,k,l) reflection and are real spherical harmonics. After refinement of the a_{imp} coefficients, the program calculates the apparent size of the crystal domains along each reciprocal lattice vector (h,k,l) direction.

Cytotoxicity trials

MTT assay

Human hepatocellular carcinoma HepG2 (HB-8065; American Type Culture Collection [ATCC], Manassas, VA, USA) cells were grown in minimal essential media (MEM) plus 10% FBS, 2 mM glutamine, 1% nonessential amino acids (NEAA), 100 U/mL penicillin, and 100 $\mu\text{g/mL}$ streptomycin (all from Thermo Fisher Scientific). Cells (200 μL at 30,000 cells/well) were placed in a 96-well plate and were treated with SR ϕ 1 (10^7 PFU/mL), HA–SR ϕ 1 (100 mg/mL plus 10^7 PFU/mL), or HA (100 mg/mL) for 24, 48, and 72 hours, respectively.

The positive and negative controls contained 10% DMSO or PBS, respectively. All the conditions tested were performed in triplicate. An aliquot of 20 μL of MTT dissolved in PBS at a concentration of 5 mg/mL was added to each well. After 2 hours, the supernatant was discarded and

150 μL of isopropanol was added. The plate was incubated at 37°C for 30 minutes and then, the optical density was measured at 560 nm.

Lactate dehydrogenase (LDH) assay

An LDH assay was performed using a CytoTox 96 Non-Radio cytotoxicity assay kit (Promega Corporation, Fitchburg, WI, USA) at 24, 48, and 72 hours according to the manufacturer's protocol. The LDH levels were evaluated for all the conditions tested for the MTT assay.

Internalization of fluorescent HA–SR ϕ I

In order to evaluate the internalization of the complex, HA nanoparticles were labeled by fluorescein isothiocyanate (FITC). Briefly, HA (100 mg/mL) was mixed with 20 mL of 3-aminopropyltriethoxysilane (Sigma-Aldrich Co.) in 100 mL of anhydrous ethanol and stirred at 74°C for 3 hours. Later, fluorescein (6 $\mu\text{g}/\text{mL}$; Sigma-Aldrich Co.) was added and continued to react at 74°C for about 20 hours in the dark. The mixture was centrifuged at 5,000 rpm for 1 minute and the pellet was rinsed three times with anhydrous ethanol and deionized water, and later suspended in SM buffer.⁵⁰ Subsequently, SR ϕ I (10^7 PFU/mL) was added to the HA previously labeled. An aliquot of 10 μL of the complex HA–SR ϕ I at different dilutions (1:1,000, 1:100, and 1:10) was added to HepG2 cells (1.0×10^5 per well) for 24 hours. The control cells were treated with 10 μL of SM buffer. After treatment, the medium was discarded, and cells were rinsed twice with PBS and fixed in 4% paraformaldehyde for 10 minutes at room temperature. DAPI was used to counterstain nuclei. Slices were observed using a Zeiss LSM 710 Confocal Laser Scanning Microscope (Carl Zeiss MicroImaging GmbH). Samples were vertically scanned with a $63\times$ or $40\times$ (1.40 NA) Plan-Apochromat oil-immersion objective. Images were obtained with Zeiss ZEN Confocal Software (Carl Zeiss MicroImaging GmbH).

Intracellular killing activity

HepG2 cells were placed in a 24-well plate (0.2×10^6 cells/well), incubated overnight (37°C , 5% CO_2) in MEM supplemented with 10% FBS (Thermo Fisher Scientific). The experiment was carried out in two different ways in order to evaluate different types of antimicrobial activity of the complex according to different infection processes. First, the cells were infected with *S. Rissen* (10^8 CFU/mL) for 1 hour. The medium was discarded, and the cells were washed with PBS (three times) and incubated in MEM plus 10% FBS and gentamicin (12.5 $\mu\text{g}/\text{mL}$)³⁶ at 37°C in 5% CO_2 for 3 hours. Later, the cells were treated with SR ϕ I (10^7 PFU/mL), HA–SR ϕ I (100 mg/mL plus 10^7 PFU/mL),

or HA (100 mg/mL) for 24 hours as reported by McLaughlin et al.⁵¹ and Withanage et al.⁵²

In the second infection process, the cells were infected with *S. Rissen* (10^8 CFU/mL) and simultaneously treated with SR ϕ I (10^7 PFU/mL), HA–SR ϕ I (100 mg/mL plus 10^7 PFU/mL), or HA (100 mg/mL) for 1 hour. The cells were then treated with gentamicin as described above. After 3 hours of incubation at 37°C in 5% CO_2 , the medium was discarded, and the cells were incubated in MEM containing 10% FBS for 24 hours.

The cells of both experiments were lysed with Tween 20 (0.05%) and incubated at 37°C in 5% CO_2 for 4 hours, and each lysate was then serially diluted in PBS and plated on XLT4 (Sigma-Aldrich Co.). The plates were incubated at 37°C overnight and evaluated for the presence of bacteria.

Both the above experiments were carried out using mouse macrophage cell line J774A.1 (TIB-67; ATCC) instead of HepG2 cells. J774A.1 cells were grown in DMEM supplemented with 10% FBS, 2 mM glutamine, 1% NEAA, 100 U/mL penicillin, and 100 $\mu\text{g}/\text{mL}$ streptomycin (all from Thermo Fisher Scientific). Cells were maintained in a humidified, 37°C , 5% CO_2 incubator.

Bacterial reduction assay on chicken minced meat

S. Rissen was processed as reported by Sukumaran et al.⁵³ For the test, 250 g of chicken minced meat was divided into five equal parts. Four sections were infected with *S. Rissen* (10^3 CFU/mL) and incubated at room temperature according to ISO 16140-2: 2016;⁵⁴ while the last one was treated only with PBS (negative control). After 30 minutes, three of the four infected parts were treated respectively with SR ϕ I (10^8 PFU/mL), HA–SR ϕ I (100 mg/mL plus 10^8 PFU/mL), or HA (100 mg/mL). All samples were stored at 4°C for 7 days and tested for the presence of the bacteria every 24 hours. In particular, at each time point, 5 g of meat was taken from each sample and homogenized in 45 mL of 0.1% buffered peptone water (AES Laboratoire Groupe, France) at 200 rpm for 1 minute, using the Stomacher 400 Circulator (Seward Ltd, Bohemia, NY, USA). Subsequently, 100 μL of sample was plated on XLT4 agar and incubated at 37°C for 24 hours. The efficacy of each treatment was evaluated by counting the CFU formed on the plate after the incubation time.

Results

Phage isolation and characterization

Phage excision from *S. Rissen* was induced by Mitomycin C procedure.^{36,55} The SR ϕ I host range was tested against 14 *Salmonella* strains, each belonging to a different serovar, and

Table 1 Phage host range against different *Salmonella* strains

Bacteria	Lytic activity
<i>S. Livingstone</i>	+
<i>S. Infantis</i>	+
<i>S. Potsdam</i>	+
<i>S. Thompson</i>	–
<i>S. Mbandaka</i>	+
<i>S. Winston</i>	+
<i>S. Montevideo</i>	–
<i>S. Virchow</i>	+
<i>S. Ohio</i>	+
<i>S. Jerusalem</i>	–
<i>S. Inganda</i>	+
<i>S. Wil</i>	+
<i>S. Typhimurium</i>	+
<i>S. Enteritidis</i>	+

Notes: + indicates lysis after infection with SR ϕ 1; – indicates lack of lysis after infection with SR ϕ 1.

it was active against 11 out of 14 selected strains (Table 1). The MOI test allowed us to select the value of 0.1 as optimal MOI (Figure S1). This value has been used for all the later experiments. The one-step growth analysis highlighted that the latent period of SR ϕ 1 was 60 minutes (Figure S2) and that its burst size was 54 PFU/cell.

TEM analysis of SR ϕ 1 allowed us to classify the phage as a member of the *Podoviridae* family, according to a previous publication.¹⁰ In particular, TEM analysis revealed that the tail length and head diameter were 16 ± 2.0 nm (mean \pm SD) and 57 ± 1.0 nm (mean \pm SD), respectively; and the total length was 73 ± 2.0 nm (mean \pm SD) (Figure 1).

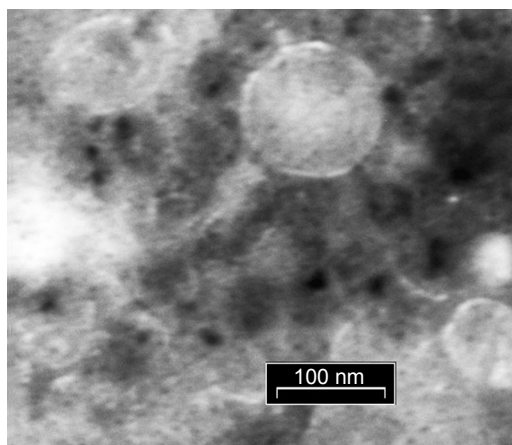


Figure 1 Electron microscopic analysis of phage. SR ϕ 1 observed with a TEM analysis. Scale bar, 100 nm.

Abbreviation: TEM, transmission electron microscopy.

Comparative genomic analysis of phage

DNA sequencing yielded a total of 2,199,543 reads (660 Mb) and an average coverage of 13,200 \times . Phage SR ϕ 1 genome consists of 51,738 bp with a G+C content of 48.4%. The genomic sequence of SR ϕ 1 described in this study has been deposited in the GenBank database (accession number: KY709687). Open reading frames (ORFs) with a length of at least 38 amino acids were selected. A total of 622 ORFs were predicted to be present within the genome. However, only 87 ORFs (13.98%) were predicted to be functional based on gene predictions and annotation of the genome (Figure 2). Concrete gene information such as positions, directions, sizes, and putative functions of SR ϕ 1 coding DNA sequences (CDSs) are shown in Table S1. The genome contains 87 predicted CDSs, of which 70 matched with already identified phage genes. Identified genes included 30 genes affecting bacteriophage physiology, 12 phage structures, 10 DNA replication, and three bacterial lysis. The remaining 32 CDSs encoded hypothetical proteins. Phage SR ϕ 1-predicted genome did not show genes coding for toxins, antibiotic resistance, or *Salmonella* virulence.

To gain an insight into the characteristics of the phage SR ϕ 1 genome, we performed a phylogenetic analysis, comparing its genome to the predicted genomes of other 37 fully sequenced phages. The phylogenetic analysis confirmed that phage ϕ 1 is a *Podoviridae*, specifically showing 19 homologous genes with *PHAGE_Salmon_SPN9CC* (NC017985) (Figure 3).

Biomimetic HA nanocrystal characterization

The biomimetic HA nanocrystals used in this study had a composition very close to that of the human body.⁵⁶

The TEM analysis revealed the length of HA nanocrystals (~30–40 nm) and its plate-like morphology (Figure S3). The high reactivity of HA is ascribed to its amorphous surface¹⁶ and to its high surface area of about 110 m²/g, which is only slightly lower than that of biological nanocrystals (120 m²/g).³⁰

Moreover, the degree of crystallinity and the presence of carbonate ions in the structure of biomimetic HA (data not shown) clearly confirm that the HA selected for this study is – structurally – very similar to that of the bone, not only in the size of the nanocrystals but also in ionic substitution.

Preparation and characterization of the complex (HA–SR ϕ 1)

The interaction of SR ϕ 1 with HA, carried out as reported in Patent IT102017000050733,⁵⁷ was evaluated at different



Figure 2 Functional genome map of phage SR ϕ 1.

Notes: Hypothetical functions of encoded proteins were determined by comparison of amino acid sequences to the non-redundant databank using BLASTP. Annotation was verified using PHAST, a web server designed to rapidly and accurately identify, annotate, and graphically display prophage sequences within bacterial genomes or plasmids. The + and – stranded ORFs were colored in yellow.

Abbreviations: BLASTP, Basic Local Alignment Search Tool Protein; ORFs, open reading frames; PHAST, PHAge Search Tool.

incubation time intervals. The results showed that SR ϕ 1 was interacted with HA already after 30 minutes. Furthermore, the graph showed that the amount of complexes SR ϕ 1–HA increased over time (Figure 4A). Based on this result, we considered overnight incubation to be the optimal time interval for the complex synthesis.

The interaction between HA and SR ϕ 1 was studied also through the zeta potential analysis, in which SR ϕ 1 was negatively charged (-11.28 ± 1.16) while HA was positively charged (2.9 ± 0.9 mV). Therefore, the SR ϕ 1 and HA complex showed a positive zeta potential of 0.9 ± 1.60 mV (Figure 4B).

In addition, the stability of the SR ϕ 1 and the HA–SR ϕ 1 complex was assessed after the complex synthesis for about 2 months at 7 days intervals. Stability for long periods is essential for using phages in several biocontrol applications.⁵⁸ The titer of the phage SR ϕ 1 alone started to decrease already after 1 week; instead, when the phage was complexed with HA, its titer was stable for up to 6 weeks without any variation at the different time points analyzed (Figure 4C).

The analysis of the HA–SR ϕ 1 complex stability at different pH values (ranging from 2 to 10) assessed the strong stability of the complex, compared to that of SR ϕ 1 alone

(Figure 4D). Further characterization of the complex was carried out by DLS analysis. During this test, no rapid aggregation phenomenon was observed. As shown in Figure 4E, the diameter of HA was 300 nm while only diameter of SR ϕ 1 was smaller (about 200 nm) than the HA. When HA and SR ϕ 1 were complexed, the estimated hydrodynamic diameter was about 400 nm, thereby confirming the interaction between these two elements as reported by Wang et al.⁵⁹

XPD analysis showed that HA nanocrystals⁶⁰ have a hexagonal crystalline structure (structure parameters are reported in Table S2). XPD patterns were further studied by using a whole-profile Rietveld-based fitting program⁴⁹ as shown in Figure S4. The Rietveld analysis allowed us to determine cell parameters and crystalline domain size along the [002] and [110] crystallographic directions (summarized in Table S3). This analysis also showed the structural variation of the HA crystalline domain size with or without SR ϕ 1 (Figure 5, black and red dots, respectively). When the HA percentage was increased, the samples without SR ϕ 1 did not show any change in the crystalline domain size along [002] and [110] directions. Instead, when the HA amount was increased, in the case of the complex (HA–SR ϕ 1), the crystalline domain size decreased (Figure 5).

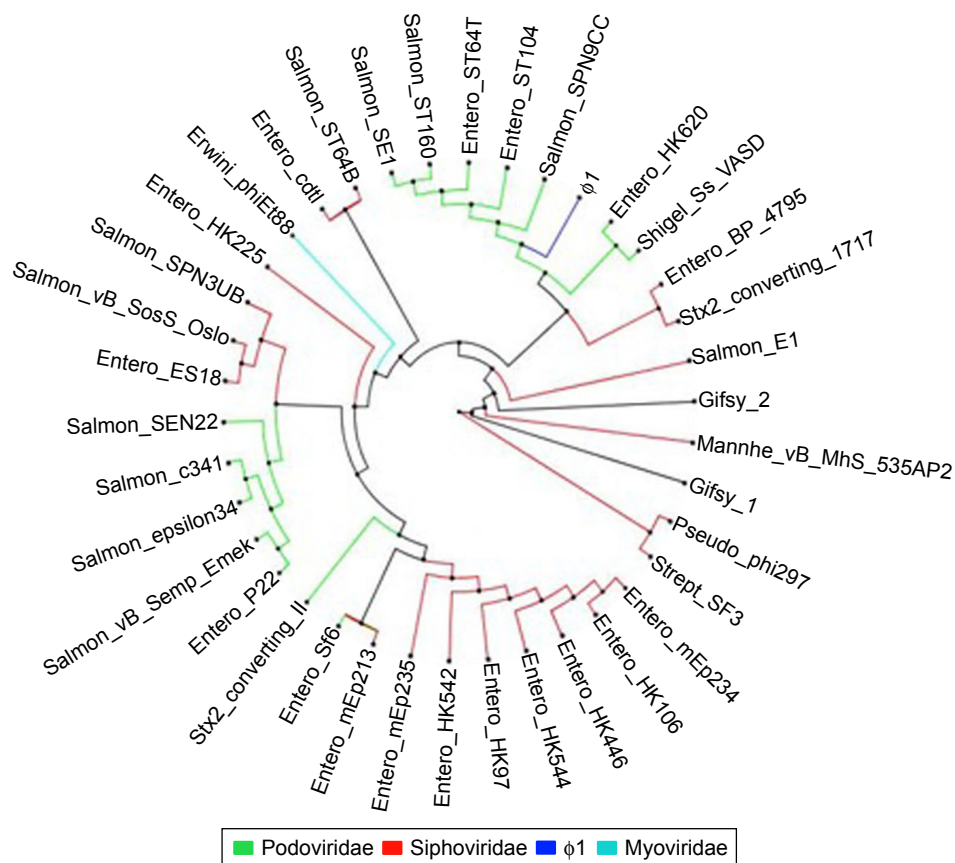


Figure 3 Assignment of SR $\phi 1$ to the *Podoviridae* family.

Notes: The phylogenetic tree shows a strong DNA identity between SR $\phi 1$ and five members of the *Podoviridae* (three *Salmonella* and two *Enterococcus* phages). The tree is based on the alignment of 39 phage genomes. The bar indicates branch length scale.

Cytotoxicity trials

To evaluate the cytotoxic effect of SR $\phi 1$, HA, and the complex HA–SR $\phi 1$, a cell viability test was carried out by evaluating the reduction of MTT to formazan in human hepatocellular carcinoma HepG2 cells. The results showed that none of the treatments affected the cell vitality up to 72 hours (Figure S5A).

Moreover, the amount of the extracellular enzyme LDH, a cell death indicator,⁶¹ was assessed under the same experimental conditions described above. Again, based on the amount of LHD produced, we concluded that none of the treatments produced cytotoxic effects (Figure S5B).

Internalization of fluorescent complex

As *Salmonella* spp. is an intracellular pathogen, the capacity of the complex to penetrate HepG2 eukaryotic cells was evaluated by laser scanning confocal microscopy analysis. Figure 6 evidences the cell uptake of the complex. In particular, the complex HA–SR $\phi 1$ (HA colored in green) was internalized into the cytoplasm of the HepG2 cells after 24 hours of treatment. Nuclei were counterstained with

DAPI dye (blue). No fluorescent signal was detected in the control cells.

Intracellular killing activity

Once the ability of the complex to penetrate host cells was determined, the intracellular killing activity was gauged in two different experimental conditions: the HA–SR $\phi 1$ complex was administered at 1 hour postinfection or together with *Salmonella* bacteria. The results showed that 1 hour postinfection, only HA–SR $\phi 1$ showed lytic activity against bacteria, reducing the bacterial load significantly (Figure 7A) while the effect of phages alone was minimal. In the second experiment, in which all the treatments were carried out simultaneously with the bacteria infection, both SR $\phi 1$ and HA–SR $\phi 1$ showed comparable lytic activity (Figure 7B). The data reported in Figure 7A evidenced that 1 hour after the infection with *S. Rissen*, the presence of biomimetic HA favored the uptake of phages into the HepG2 cells, allowing phages to kill also the intracellular bacteria.

Furthermore, Figure 7B showed that the biomimetic HA nanocrystals did not compromise the lytic activity of the

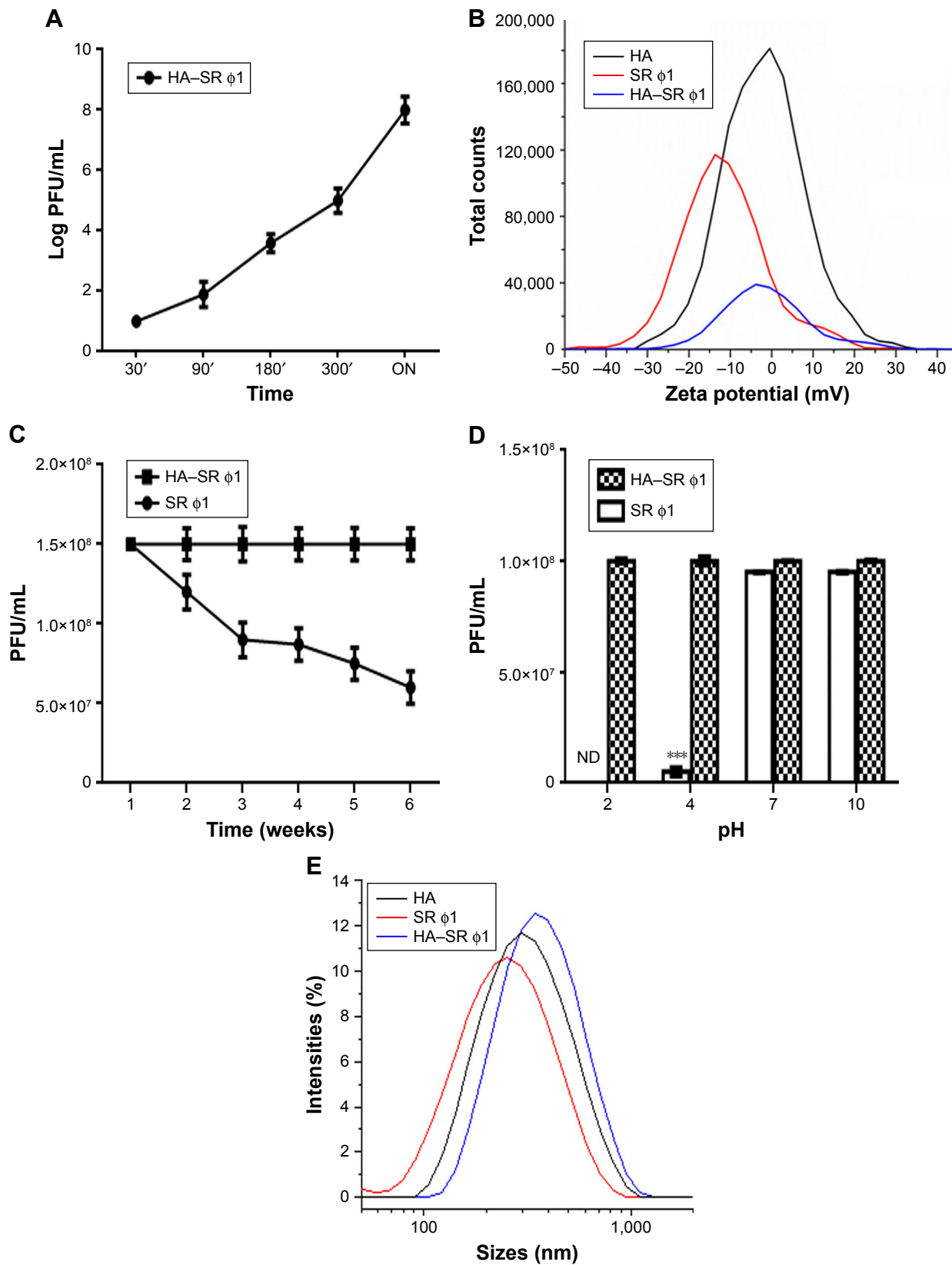


Figure 4 (A) The absorption of SR ϕ 1 on HA after different times (30, 90, 180, 300 minutes and overnight) of incubation. Each value is the mean \pm SD of three independent experiments with three replicates each. (B) Zeta potential analysis of SR ϕ 1, HA-SR ϕ 1, and HA. (C) The half-life of the SR ϕ 1 and the complex HA-SR ϕ 1. Each value is the mean \pm SD of three independent experiments with three replicates each. (D) Effects of pH on the stability of SR ϕ 1 and HA-SR ϕ 1. *** $P < 0.001$. Each value is the mean \pm SD of three independent experiments with three replicates each. Statistical analysis was performed with Student's *t*-test. (E) DLS analysis of the SR ϕ 1, HA-SR ϕ 1, and HA. **Abbreviations:** DLS, dynamic light scattering; HA, hydroxyapatite; ND, not detectable; PFU, plaque forming unit.

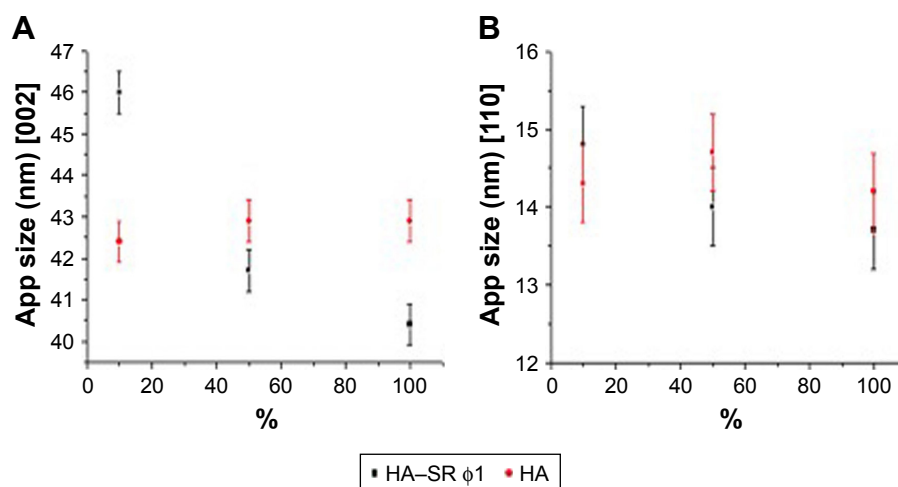


Figure 5 XPD analysis.

Note: Apparent size along the: (A) [002] and (B) [110] crystallographic directions vs the HA percentage for the complex (HA-SR ϕ 1) (black dots) and for HA alone (red dots).
Abbreviations: App, apparent; HA, hydroxyapatite; XPD, X-ray powder diffraction.

phages, when the complex was applied concurrently with the bacteria. The trend observed in HepG2 cells was also observed for murine macrophage cell line J774A.1 (data not shown).

Bacterial reduction assay on chicken minced meat

The efficacy of the complex HA-SR ϕ 1 to reduce *Salmonella* contamination was investigated in chicken minced meat which is considered one of the meat categories intended to be cooked before consumption (together with mechanically separated meat and meat preparations) with the highest level of noncompliance, as reported by European Food Safety Authority (EFSA).⁶² *S. Rissen* colonies, in the sample treated with SR ϕ 1, were reduced by 0.3 log CFU/g, while the sample treated with only HA was contaminated very similar to that of the positive control (5.5 log CFU/g). Instead, in the case of treatments with HA-SR ϕ 1, the bacterial load of *S. Rissen* was reduced by 3 log CFU/g (Figure 8).

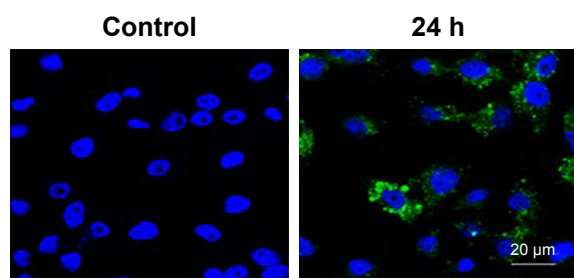


Figure 6 Confocal microscopy analysis.

Notes: Control cells or cells treated with the complex at 24 hours. Scale bars, 20 μ m.

Discussion

Different material nanoparticles such as carbon, silica, metal oxide, graphene, and HA have been used in several studies to stabilize, protect, deliver, or enhance some biological properties of molecules.^{14–16,30,44}

The characteristics of HA used in this study (such as composition, structure, size, and morphology) and the results obtained allow us to confirm that this mineral, due to its features, was able to chemically interact not only with molecules but also with biological structures like bacteriophages. Its low degree of crystallinity and the presence of carbonate ions in the crystal structure are other important characteristics that make HA extremely reactive in biological systems, and particularly suitable to interact and transport the bacteriophages, as reported in this study.

The bacteriophage SR ϕ 1 – isolated in this study – showed an efficient bacteriolytic activity against *S. Livingstone*, *S. Infantis*, *S. Potsdam*, *S. Thompson*, *S. Mbandaka*, *S. Winston*, *S. Montevideo*, *S. Virchow*, *S. Ohio*, *S. Jerusalem*, *S. Inganda*, *S. Wil*, *S. Typhimurium*, and *S. Enteritidis* as shown in Table 1. These results evidenced that it could be used as an antimicrobial agent against the strains of *Salmonella* spp. selected for this study. Further investigations are needed to evaluate whether the phage SR ϕ 1 could show antimicrobial activity against other *Salmonella* strains and serovars.

Based on these results, we evaluated whether the use of HA could improve some properties of the bacteriophage. First, we used the mineral HA, which did not affect the cell viability and was not toxic for human cells (Figure S5).

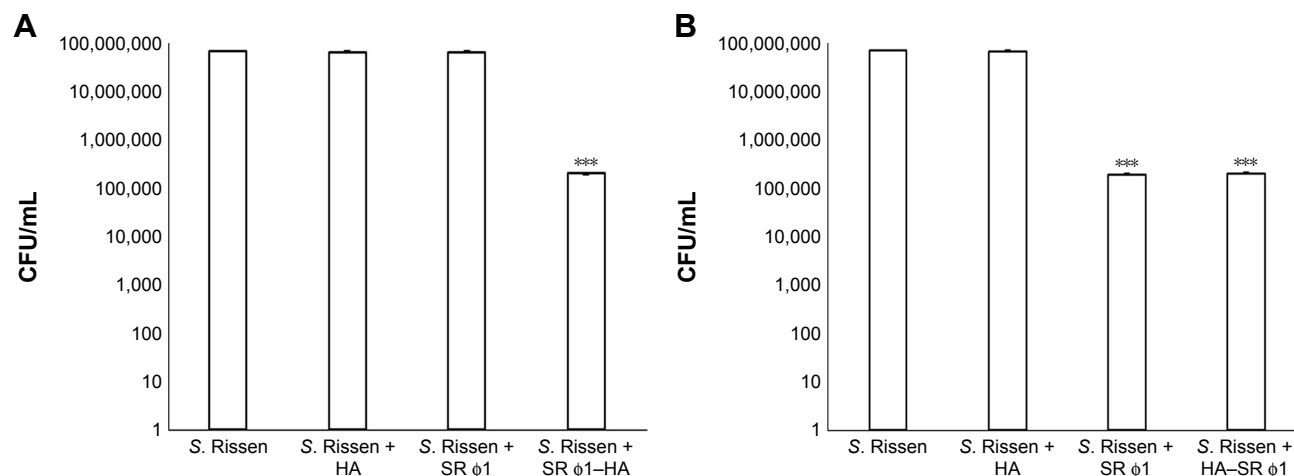


Figure 7 Intracellular killing activity.

Notes: (A) First experiment. (B) Second experiment. Positive control is represented by *Salmonella* Rissen infected cells. *** $P < 0.001$. Each value is the mean \pm SD of three independent experiments with three replicates each. Statistical analysis was performed with Student's *t*-test.

Abbreviation: CFU, colony forming unit.

The complex HA–SR ϕ 1 showed enhanced lytic activity against *S. Rissen* than SR ϕ 1 alone, thus indicating that the interaction with HA increased the antibacterial activity of the bacteriophage. Furthermore, the mineral contributed to stabilize the phage activity over time and at different pH values, making the complex suitable for long-term treatment and effective for controlling infection in harsh niches, showing acid pH (such as stomach, urinary tract, etc), or in different physiological environments of healthy tissues (including blood).

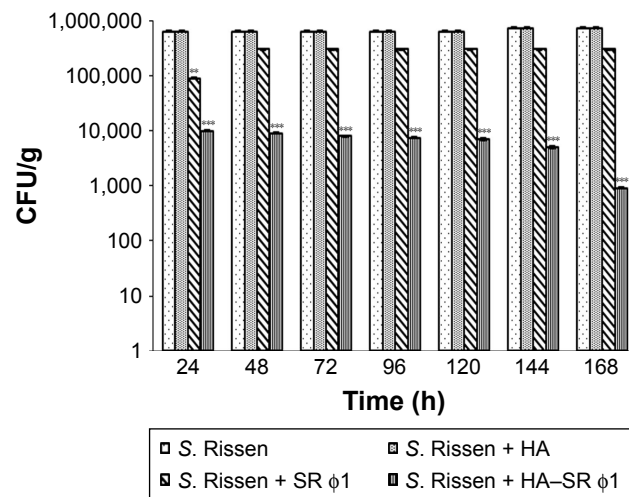


Figure 8 Bacterial reduction assay on chicken meat.

Notes: The samples of the meat were infected with *Salmonella* Rissen (10^3 CFU/mL) and were treated with SR ϕ 1 (10^7 PFU/mL), HA–SR ϕ 1 (100 mg/mL plus 10^7 PFU/mL), and HA (100 mg/mL). Positive control was represented by *S. Rissen* infected meat. ** $P < 0.01$, *** $P < 0.001$. Each value is the mean \pm SD of three independent experiments with three replicates each. Statistical analysis was performed with Student's *t*-test.

Abbreviations: CFU, colony forming unit; HA, hydroxyapatite; PFU, plaque forming unit.

The interaction of HA with phage is a good strategy to overcome the problem related to the reduction of bacteriophage viability in acidic conditions.¹²

The zeta potential analysis and DLS measurements demonstrated the interaction of the phage with HA. In fact, Figure 4B showed the behavior of the zeta potential of SR ϕ 1 and HA alone and when they were complexed. In the latter case, neutralization of the negative surface charges of SR ϕ 1 was observable. Because the charges of two elements are reversed, neutralization of the surface charges could be principally due to electrostatic interactions. Furthermore, the results of the DLS analysis allow us to hypothesize that when the two elements are complexed, the HA acts as a scaffold for the phages.

These results are also supported by Wang et al.⁵⁹ Indeed, they demonstrated the ability of the bacteriophages to bind calcium ions, or non-stoichiometric HAs with positive zeta potential.⁵⁹ The HA used in this study, just like the ones mentioned above, has a positive zeta potential, a non-stoichiometric Ca/P ratio; all these characteristics induce the interaction of the HA with proteins and carboxyl groups of the phage capsid by the development of an electrostatic bond.

Some studies have shown the capacity of HA to permeate the cell membrane through the energy-dependent process of clathrin-mediated endocytosis⁶³ or phagocytosis⁶⁴ but have, also, highlighted that the permeabilization process is influenced by the equilibrium of multiple features such as dimension, charge, shape, and surface area.^{24,64–66}

It has been reported that the bacteriophages are also able to bypass the cell membrane.⁶⁷

Based on these evidences, we tested the capacity of the complex, and in particular the contribution of the mineral HA to penetrate into the eukaryotic cells. The results demonstrated that, in the same infection condition, the presence of the mineral increases the number of phages that are able to penetrate into cells more efficiently and consequently kill intracellular bacteria.³⁶ This mineral could be a good candidate in the use of phage therapy against obligate or facultative intracellular bacteria like *Mycobacterium* spp., *Chlamydia* spp., *Rickettsia* spp. or *Salmonella* spp., *Listeria* spp., and *Brucella* spp.⁶⁸

In addition, another experiment was carried out on HepG2 cells or J774A.1 cells not infected with bacteria. The results confirmed the ability of the HA to enter into eukaryotic cells and the capacity of bacteriophage to bypass the cell membrane (both reported in literature) and, at the same time, highlighted the role of HA in increasing the number of phages that pass through the cell membrane (data not shown) although there was no infection.

The last test, carried out to evaluate the efficacy of the complex to control bacterial infection in a food matrix such as chicken minced meat and for long time, showed the ability of the complex to reduce bacterial load also in the case of food contamination.

Conclusion

The main drawback in using phages are their lack of stability over time and their low activity against intracellular infections due to their low efficiency in penetrating eukaryotic cells. In addition, phages do not tolerate the low pH present in the stomach and when used in food processing their activity can be compromised. In this study, these problems were successfully addressed by complexing phages with HA (nontoxic mineral for humans). This complex is stable, allows phages to enter eukaryotic cells more efficiently than phages alone, and, at the same time, the complexed phages were stable at very low pH. These results evidence the important contribution of HA making it a promising approach to overcome problems which could emerge when biological entities are used.

One of the most important problems in phage therapy is the application of phages as biocontrol agents against contamination in food.⁶⁹ This drawback has been addressed, and it has been resolved by carrying out a test on infected meat. The results showed how HA enhances the lytic activity of phage to control bacterial meat infection. Strikingly, the complex is able to control *Salmonella* infection in food.

Furthermore, the approach proposed here (use of biomimetic HA nanocrystals as a carrier for bacteriophages) can

be extended to different fields of interest such as biomedical, agricultural, and other commercial applications.

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Author contributions

RC, FCa, and NR have made major contributions to the conception of the study; AF, FI, MP, FCo, TS, CG, SP, RV, BDV, and ML contributed to the acquisition, analysis, or interpretation of the data; AF, FI, MP, CG, and BDV performed the experiments; RC wrote the manuscript. All authors contributed to data analysis, drafting and revising the article, gave final approval of the version to be published, and agree to be accountable for all aspects of the work.

Disclosure

AF, FI, NR, ML, FCo, and RC are the inventors of the Patent IT102017000050733 presented to the Italian Ministry of Economic Development on May 10, 2017. All the other authors report no conflicts of interest in this work.

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