CRISPR-based herd immunity limits phage epidemics in bacterial populations

Pavel Payne[1,2*], **Lukas Geyrhofer**[3], **Nicholas H. Barton**[2], **Jonathan P. Bollback**[1,2*]

***For correspondence:**
Pavel.Payne@liverpool.ac.uk (PP);
J.P.Bollback@liverpool.ac.uk (JPB)

[1]Institute of Science and Technology Austria, Am Campus 1, Klosterneuburg, Austria; [2]Institute of Integrative Biology, University of Liverpool, Liverpool, UK; [3]Technion – Israel Institute of Technology, Haifa, Israel

Abstract Pathogens are a driving force in evolution, giving rise to a diversity of host immune defenses. In order for a pathogen to spread in a population a sufficient number of its members must be susceptible to infection, as resistant individuals can prevent the spread of a pathogen among susceptible hosts in a process known as herd immunity. While herd immunity has been extensively studied in vertebrate populations, little is known about its role, if any, in the dynamics between bacteria and their phage pathogens. Here we explore the dynamics of T7 phage epidemics in structured and unstructured *Escherichia coli* populations consisting of differing mixtures of susceptible and resistant individuals harboring CRISPR immunity to the phage. Using both experiments and mathematical modelling we describe the conditions under which herd immunity arises in bacterial populations. Notably, the effects of herd immunity depend strongly on the presence of spatial structure in the population, the bacterial growth rate, and phage replication rate. The results of our model can apply to other host–pathogen systems to determine the herd immunity threshold from the relative speed of an epidemic wave in partially resistant populations. In addition, our findings suggest that herd immunity plays an important role in bacterial communities, as seen in other host–pathogen systems, allowing for stable coexistence of bacteria and their phages and the maintenance of polymorphism in bacterial immunity.

Introduction

"Herd immunity" has been used in a variety of ways by different authors (see ***Fine et al.*** (***2011***)). Here, we define it as a phenomenon where a fraction of resistant individuals in a population reduces the probability of transmission of a pathogen among the susceptible individuals. Furthermore, if the fraction of resistant individuals in a population is sufficiently large the spread of a pathogen is suppressed. Experimental research into the phenomenon has focused mostly on mammals ***Jeltsch et al.*** (***1997***); ***Mariner et al.*** (***2012***), birds ***Boven et al.*** (***2008***); ***Meister et al.*** (***2008***), and invertebrates ***Konrad et al.*** (***2012***); ***Wang et al.*** (***2013***). In human populations the principles of herd immunity were employed to limit epidemics of pathogens through vaccination programs ***Fine et al.*** (***2011***), which in the case of smallpox lead to its eradication between 1959 and 1977 ***Fenner*** (***1993***).

Alongside advances in vaccination programs, the formalization of a general theory of herd immunity was developed. The theory is based on a central parameter, R_0, which describes the fitness of the pathogen, as measured by the number of subsequent cases that arise from one infected individual in a population (for a historical review of R_0 see ***Heesterbeek*** (***2002***)). Thus, R_0 indicates the epidemic spreading potential in a population. Given R_0 the herd immunity threshold

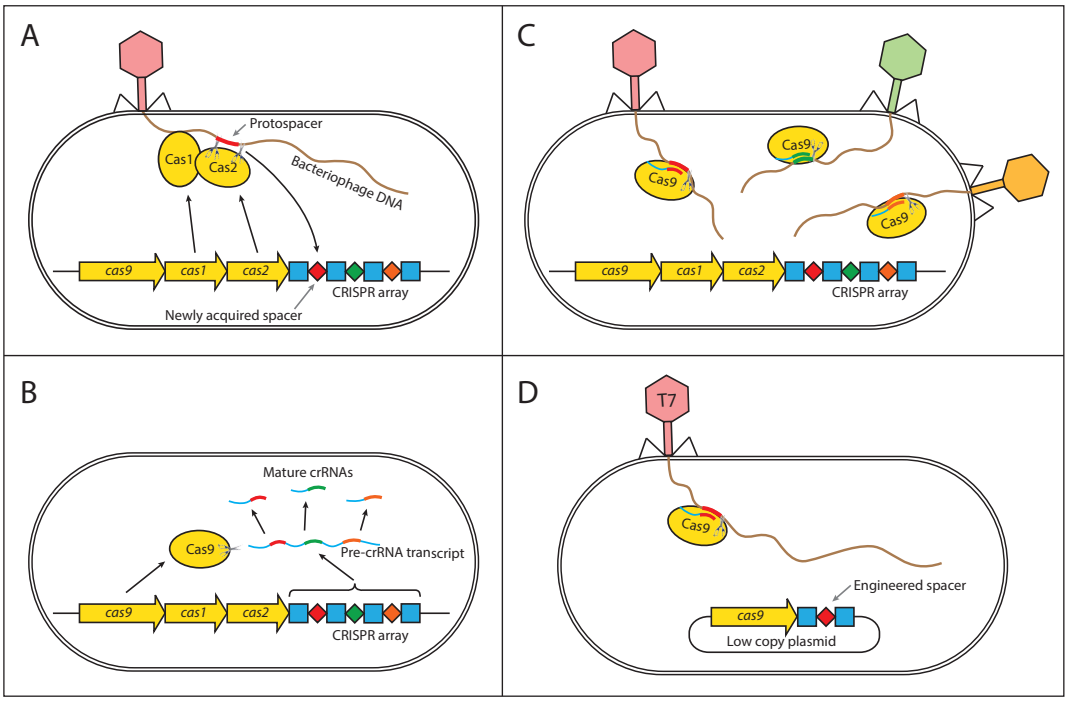


Figure 1. Mechanism of CRISPR/Cas type II immunity. The CRISPR/Cas system provides immunity to phages and its main features can be described by three distinct stages. (A) Acquisition. When a cell gets infected by a phage, a protospacer on the invading phage DNA (indicated as a red bar) is recognized by Cas1 and Cas2. The protospacer is cleaved out and ligated to the leader end (proximal to the Cas genes) of the CRISPR array as a newly acquired spacer (red diamond). (B) Processing. The CRISPR array is transcribed as a Pre-crRNA and processed by Cas9 (assisted by RNaseIII and trans–activating RNA, not shown) into mature crRNAs. (C) Interference. Mature crRNAs associate with Cas9 proteins to form interference complexes which are guided by sequence complementarity between the crRNAs and protospacers to cleave invading DNA of phages whose protospacers have been previously incorporated into the CRISPR array. (D) A truncated version of the CRISPR system on a low copy plasmid, which was used in this study lacks *cas1* and *cas2* genes and was engineered to target a protospacer on the T7 phage chromosome to provide *Escherichia coli* cells with immunity to the phage. The susceptible strain contains the same plasmid except the spacer does not target the T7 phage chromosome.

is defined as,

$$H = \frac{R_0 - 1}{R_0}\,, \tag{1}$$

which determines the required minimum fraction of resistant individuals needed to halt the spread of an epidemic and is effected by the specific details of transmission and the contact rate among individuals ***Grassly and Fraser*** (***2008***). Many theoretical studies have addressed the influence of some of these details, in particular maternal immunity ***Anderson and May*** (***1992***), age at vaccination ***Anderson and May*** (***1982***); ***Nokes and Anderson*** (***1988***), age related or seasonal differences in contact rates ***Schenzle*** (***1984***); ***Anderson and May*** (***1985***); ***Yorke et al.*** (***1979***), social structure ***Fox et al.*** (***1971***), geographic heterogeneity ***Anderson and May*** (***1984***); ***Lloyd and May*** (***1996***); ***Real and Biek*** (***2007***), and the underlying contact network of individuals ***Ferrari et al.*** (***2006***).

Interestingly, little work has focused on the potential role of herd immunity in microbial systems which contain a number of immune defense systems and have an abundance of phage pathogens. These defenses vary in their potential to provide herd immunity as they target various stages of the phage life cycle, from adsorption to replication and lysis. Early defense mechanisms include the prevention of phage adsorption by blocking of phage receptors ***Nordström and Forsgren*** (***1974***), production of an extracellular matrix ***Hammad*** (***1998***); ***Sutherland et al.*** (***2004***), or the excretion of competitive inhibitors ***Destoumieux-Garzón et al.*** (***2005***). Alongside these bacteria have evolved innate immune systems that target phage genomes for destruction. These include host restriction-

modification systems (RMS) ***Blumenthal and Cheng*** (***2002***), argonaute-based RNAi-like systems ***Swarts et al.*** (***2014***), and bacteriophage-exclusion (BREX) systems ***Goldfarb et al.*** (***2015***). In addition to innate systems, bacteria have evolved an adaptive immune system called CRISPR-Cas (clustered regularly interspaced short palindromic repeat) ***Sorek et al.*** (***2013***). In order for any of these immune systems to provide herd immunity, they must prevent further spread of the pathogen, i.e., provide a 'sink' for the infectious particles reducing the average number of successful additional infections below one. Unlike the early defense mechanisms that may simply prevent an infection but not the further reproduction of infectious particles, the RMS, BREX, argonaute-based RNAi-like, and the CRISPR-Cas systems degrade foreign phage DNA after it is injected into the cell, and thus continue to remove phage particles from the environment, which increases their potential to provide herd immunity. Among these the CRISPR-Cas system is unique in that it is adaptive allowing cells to acquire immunity upon infection (see Fig 1A, B, and C), which can lead to polymorphism in immunity and give rise to herd immunity.

In addition to immune system-specific factors, the reproductive rate of phage depends strongly on the physiology of the host bacterium ***Hadas et al.*** (***1997***), and the underlying effective contact network which may vary greatly in bacterial populations depending on the details of their habitat. Thus, herd immunity will be influenced by the physiological state of the bacteria and the mobility of the phage in the environment through passive diffusion and movement of infected individuals. Taken together these details call into question the applicability of the traditional models of herd immunity from vertebrates to phage-bacterial systems. Thus, experimental investigation and further development of extended models that take into account the specifics of microbial systems are required.

To investigate under which conditions herd immunity may arise in bacterial populations, we constructed an experimental system consisting of T7 phage and bacterial strains susceptible and resistant to it. Our experimental system can be characterized by the following features. First, we used two strains of *Escherichia coli*, one with an engineered CRISPR-based immunity to the T7 phage, and the other lacking it (Fig 1D). Second, we examined the dynamics of the phage spread in different environments – spatially structured and without structure. Furthermore, we developed and analyzed a spatially explicit model of our experimental system to determine the biologically relevant parameters necessary for bacterial populations to exhibit herd immunity.

Results

Properties of resistant individuals

We engineered a resistant *E. coli* strain by introducing the CRISPR-Cas Type II system from *Streptococcus pyogenes* with a spacer targeting the T7 phage genome (see Material and Methods). We further characterized the ability of the system to confer resistance to the phage. We find a significant level of resistance as measured by the probability of cell burst when exposed to T7 (Fig 2A). However, resistance is not fully penetrant as approximately 1 in 1000 resistant cells succumb to infection. In addition, we observe that as phage load increases (multiplicity of infection, MOI) the probability that a cell bursts increases (Fig 2A). In order to determine the herd immunity threshold in our experimental system, we constructed the resistant strain such that upon infection the cell growth is halted, yet the cell still adsorbs and degrades phages (Fig 2B,C). This feature is important as it prevents the action of frequency dependent selection which in naturally growing populations will favor the resistant strain until its frequency reaches the herd immunity threshold. Thus, in our system if the frequency of the resistant strain is below the herd immunity threshold, the resistant cells remain below the threshold and are unable to stop the epidemic and the whole population collapses. In contrast, if the frequency of resistant individuals in the population is above the herd immunity threshold, the resistant individuals provide complete herd immunity and the population survives. These properties allow us to quantify the expanding epidemic in both liquid media and on bacterial lawns (without and with spatial structure, respectively) using high throughput techniques.

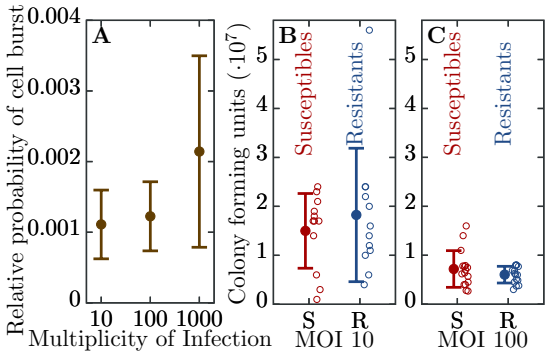


Figure 2. Efficiency of bacterial resistance. (A) The probability that a resistant cell bursts, relative to a susceptible cell, at three different initial multiplicities of infection (MOI). The probability that a resistant cell bursts at MOI 1000 is significantly higher than at MOI 10 (p = 0.019, $t_{4,0.05}$ = 3.031) or at MOI 100 (p = 0.022, $t_{5,0.05}$ = 2.674). The error bars show the standard deviations from the mean. Note that this measure is not a widely used 'efficiency of plating' but it determines the probability of burst of single resistant cells (see Materials and Methods for details). (B) The number of colony forming units (CFUs) post phage challenge (see Materials and Methods). The mean number of CFUs after the bacterial cultures were exposed to the phage is not significantly different between susceptible and resistant strains at MOI 10 (p = 0.239, $t_{22,0.05}$ = 0.721) and (C) at MOI 100 (p = 0.27, $t_{30,0.05}$ = 1.124), indicating that the resistant cells' growth is halted after the cells are infected by a phage. The error bars show the standard deviations from the mean. There were no detectable CFUs in either susceptible or resistant cell cultures at MOI 1000. It should be noted that the indicated MOI values do not correspond to the average number of phages that adsorb to cells in the experiments. For MOI 10 we estimated the mean number of phages per cell as 0.229 and for MOI 100 as 0.988 (see Materials and Methods for details). It was impossible to determine the mean for MOI 1000 as there were no detectable CFUs under such conditions.

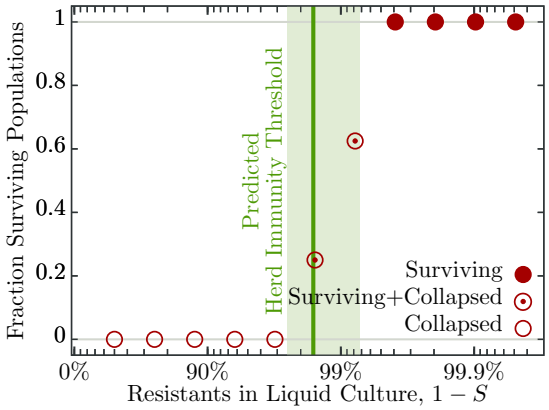


Figure 3. Fraction of surviving populations at $18\,h$ post phage infection. Bacterial populations consisting of various fractions of resistant to susceptible individuals infected with ≈ 50 phages, corresponding to a multiplicity of infection (MOI) of $\approx 10^{-4}$, to resemble an epidemic initiated by the burst size from one infected individual (see Table 2 for burst size estimates). Each population phage challenge is replicated 16 times. The solid dark green line shows the model prediction, Eqn. (4), for the herd immunity threshold (H), given latent period (λ), bacterial growth rate (α), and phage burst size (β). Shaded area indicates ± 1 standard deviation.

Specifically, it allows us to control for the complex dynamics of the system arising from frequency dependent selection and simultaneous changes in the physiological states of the cells (growth rates depending on the nutrient concentrations) and phage (burst size, latent period depending on the cells' physiology).

Herd immunity in populations without spatial structure

To understand the influence of spatial population structure, or lack thereof, we first measured the probability of population survival (i.e., whether the cultures are cleared or not) in well mixed liquid environments (no spatial structure) consisting of differing proportions of resistant to susceptible

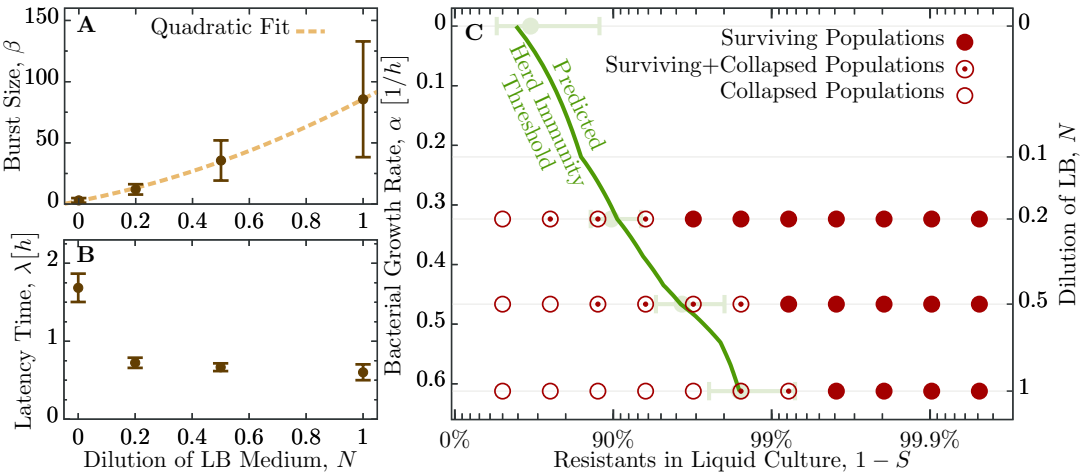


Figure 4. Herd immunity threshold in liquid culture as a function of bacterial growth. (A) Phage burst size (β) change as a function of nutrient concentrations. Brown dashed line shows a numerical quadratic fit to the observed values of β. (B) Latent period (λ) increase across the range of nutrient concentrations. Values for β and λ are given in Table 2. (C) Population survival analysis upon phage challenge as a function of the fraction of resistant cells and the intrinsic growth rate (nutrient availability, N). The empty circles, circled dots, and filled circles represent the outcomes of 18 replicates done in 3 independent batches, as indicated in the legend. The green line shows the model predicted herd immunity threshold given by Eqn. (5), using a quadratic fit for β/λ and inverting the Monod kinetics of bacterial growth (see Fig 6B) numerically. Light green error-bars are estimates from Eqn. (5) using actual experimental results for growth parameters.

individuals and T7 phage. When the percentage of resistant individuals is in excess of 99.6% all 16 replicate populations survive a phage epidemic (i.e., show no detectable difference in growth profiles to the phage free controls; Fig 3). Populations with 99.2% and 98.4% resistant individuals show intermediate probabilities of survival – 10 out of 16 replicate populations and 4 out of 16 replicate populations survive, respectively (Fig 3). The likely explanation as to why some populations survive and others collapse is due to the stochastic nature of phage adsorption after inoculation: If the population composition is close to the herd immunity threshold a stochastic excess of phage particles adsorbing to susceptible cells may trigger an epidemic, whereas if chance increases the number of phages adsorbing to resistant individuals, the epidemic is suppressed. However, when populations have fewer than 96.9% resistant individuals all 16 replicate populations fail to survive and collapse under the epidemic (Fig 3).

As mentioned in the introduction, phage and bacterial physiology may affect the herd immunity threshold. To test this we altered bacterial growth by reducing the concentration of nutrients in the medium (Fig 6) which concurrently alters the T7 phage's latent period and burst size (Fig 4A,B and Table 2). Indeed, we observe as bacterial growth rates decline the fraction of resistant individuals necessary for population survival decreases (Fig 4C). When the populations are grown in a 50% diluted growth medium, the fraction of resistant individuals required for a 100% probability of survival is 99.2%; when the fraction of resistant individuals is 75% or less populations go extinct. In a 20% growth medium the fraction of resistant individuals required for survival decreases to 96.9%, while the fraction when all replicates collapse to 50%.

From the experimental observations of the herd immunity threshold values we infer the phage R_0 using Eqn. 1. In an undiluted growth medium the phage R_0 falls between 32 and 256 and decreases to between 4 and 128 in 50% and between 2 and 32 in 20% nutrient medium. These data indicate that bacterial populations can exhibit herd immunity in homogeneous liquid environments. However, bacteria typically live in spatially structured environments such as surfaces, biofilms or micro-colonies, therefore we extended our experiments to consider the potential impact of spatially structured populations.

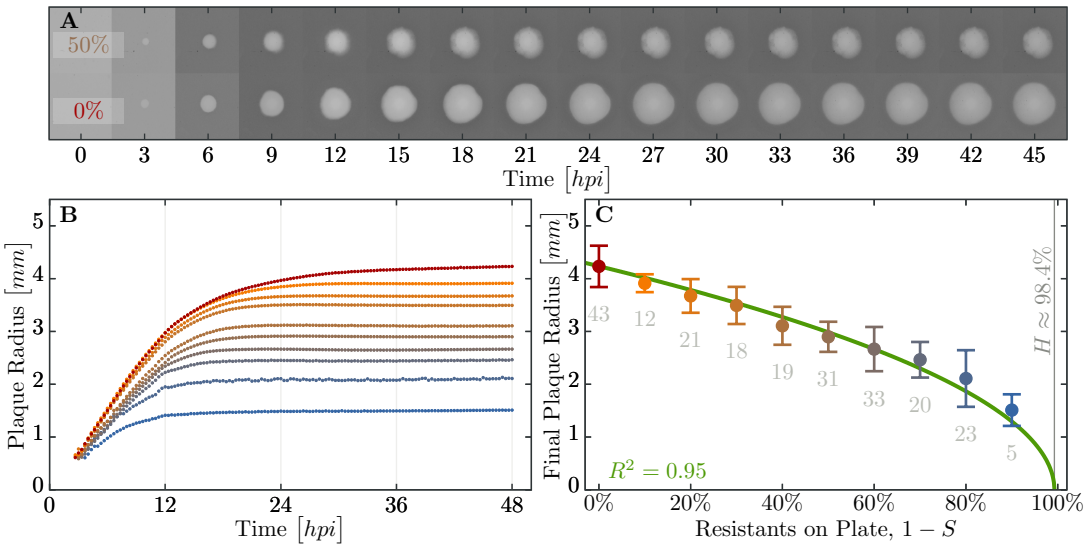


Figure 5. Properties of expanding phage epidemics on bacterial lawns. (A) Example of plaque morphology and size change over 48 hours for populations with 50% resistant cells (top) and a control with 100% susceptible cells (bottom). (B) Mean plaque size area through time. Colors indicate the different fraction of resistant individuals (color coding as in panel C). Note the distinct two phases of plaque growth – initially, phage grow fast with exponentially growing bacteria but slow once the nutrients are depleted (≈ 10 hours). The plaque radius is reduced, relative to 100% susceptible population, even when only a small fraction of resistant individuals are in the population. (C) Final plaque radius at 48 *hpi*. Green line shows the prediction from the model for the plaque radius r. Grey numbers indicate the number of plaques measured. Error bars indicate the standard deviations.

Herd immunity in spatially structured populations

In order to discern the role, if any, spatial structure plays in herd immunity we conducted a set of experiments in spatially structured bacterial lawns on agar plates. Spatially structured bacterial populations provide a more fine grained measure of herd immunity, compared to the population survival assays done in liquid culture. On bacterial lawns, phages spread radially from a single infectious phage particle and the radius of plaque growth on different proportions of resistant to susceptible individuals can be easily quantified. In addition, these data allow for estimating the speed of the epidemic wave front in these different regimes using real-time imaging (Fig 5A).

We observe a decline in the number of plaque forming units (data not shown) and a significant decrease in final plaque sizes as the proportion of resistant individuals in the populations increases (Fig 5B,C). A reduction in the final plaque size compared to a fully susceptible population was statistically significant with as few as 10% resistant individuals in a population ($p = 0.004$, $t_{53,0.05} = 2.744$). In order to determine the effect of resistant individuals during the earlier phase of bacterial growth (until the bacterial density on the agar plate reaches saturation; Fig 6A), we analyze the velocities of plaque growth between 0 and 24 hours post inoculation (*hpi*). We find that the speed is significantly reduced after $11\,hpi$ when the population consists of as few as 10% of resistant individuals ($p = 0.0317$, $t_{32,0.05} = 1.923$). As the fraction of resistant individuals further increases, the speed declines significantly at earlier and earlier time points: $6\,hpi$ with 20% ($p = 0.0392$, $t_{62,0.05} = 1.79$), and $5.67\,hpi$ with 30% ($p = 0.0286$, $t_{53,0.05} = 1.943$). In fact, when the fraction of resistant individuals exceeds 40%, the reduction in the speed of the spread is statistically significant immediately after the plaques are visually detectable (Fig 7). It should be noted that all populations with such low percentages of resistant individuals in liquid environment collapsed, indicating that spatial structure significantly facilitates herd immunity.

The results presented in this and the previous section would allow us to use Eq. (1) to infer a value for R_0 from the observed threshold between surviving and collapsing bacterial populations, Figs 3 and 4. We also observe that herd immunity is strongly influenced by spatial organization of

the population, Fig 5. How the exact value of H (and subsequently the "classical" epidemiological parameter R_0) is affected by various factors such as bacterial growth rate, phage burst size and latent period is, however, difficult to resolve experimentally. Similarly, quantification of the effect of spatial structure is hardly achievable solely by experimental investigation. In order to disentangle the roles of all the factors mentioned above, we proceed with development and analysis of a mathematical model of the experimental system.

Modelling bacterial herd immunity

We developed a model of phage growth that takes several physiological processes into account: bacterial growth during the experiment, bacterial mortality due to phage infection, and phage mortality due to bacterial immunity. Furthermore, we use the previously reported observation that phage burst size β and latent period λ depend strongly on the bacterial growth rate α (see Table 1).

The main processes in our model system can be defined by the following set of reactions,

$$B_\mathrm{s} + Y\,N \xrightarrow{\alpha} 2\,B_\mathrm{s}\,, \tag{2a}$$

$$B_\mathrm{r} + Y\,N \xrightarrow{\alpha} 2\,B_\mathrm{r}\,, \tag{2b}$$

$$B_\mathrm{s} + P \xrightarrow{A} \left(B_\mathrm{s} P\right) \xrightarrow{1/\lambda} \beta\,P\,, \tag{2c}$$

$$B_\mathrm{r} + P \xrightarrow{A} \left(B_\mathrm{r} P\right) \begin{cases} \xrightarrow{\text{fast}} B_\mathrm{r}\,, \\ \overset{\text{slow}}{\dashrightarrow} \beta\,P\,. \end{cases} \tag{2d}$$

Susceptible (B_s) and resistant (B_r) cells grow at a rate α (no significant difference in growth rate between strains, p = 0.066, $t_{70,0.05}$ = 1.867), (2a) and (2b), by using an amount Y of the nutrients N. Phage infection first involves adsorption to host cells, (2c) and (2d), with the adsorption term A specified below. Infected susceptible bacteria produce on average β phage with a rate inversely proportional to the average latency λ. In contrast, resistant bacteria either survive by restricting phage DNA via their CRISPR-Cas immune system or – less likely – succumb to the phage infection. However, when the MOI is large even resistant cells become susceptible to lysis resulting in the release of phage progeny (see Fig 2) ***Westra et al.*** (***2015***); ***Chabas et al.*** (***2016***).

In our system, bacteria eventually deplete the available nutrients, $N(t > T_\mathrm{depl}) = 0$, resulting in the cessation of growth. This decline in bacterial growth affects phage growth – latency increases and burst size decreases, such that phage reproduction declines dramatically (see Table 2). We define the critical time point at which cells transition from exponential growth to stationary phase as,

$$T_\mathrm{depl} \approx \frac{1}{\alpha} \log\left(\frac{B_\infty}{B_0}\right)\,. \tag{3}$$

Here, B_0 and B_∞ are the initial and final bacterial densities, respectively. In the initial exponential growth phase, our estimates from experimental data for growth parameters are $\alpha = 0.63\,h^{-1}$, $\beta = 85.6\,\mathrm{phages/cell}$ and $\lambda = 0.60\,h$, for bacteria and phages, respectively (Table 1 and Table 2). After time T_depl, bacterial growth rate is set to zero ($\alpha = 0$) and phage growth is reduced to $\beta_\mathrm{depl} = 3.0\,\mathrm{phages/cell}$ and $\lambda_\mathrm{depl} = 1.69\,h$. Such a two state model – constant growth rate while nutrients are present and no growth after depletion – describes the observed population trajectories in experiments quite well (see Fig 6).

Modelling herd immunity in populations without structure

An important parameter for estimating herd immunity is the fraction S of susceptible bacteria in the population. As a first estimate, a phage infection spreads in well mixed bacterial cultures if $\beta S > 1$ for a continuous chain of infections: the product of burst size β of phage particles and the probability S of infecting a susceptible host has to be larger than one. As a first approximation, one could identify R_0 with the burst size β, which is compatible with the observed herd immunity thresholds when inverting Eq. (1).

However, the growing bacterial population could outgrow the phage population, if the former reproduces faster, which introduces deviations from the simple relation between R_0 and H as shown in (1). We capture this dynamical effect in a correction to the previous estimate as $\beta S > 1 + \lambda\alpha$ (see Materials and Methods): more phages have to be produced for the chain of infections to persist in growing populations. The correction $\frac{\lambda}{1/\alpha}$ is the ratio of generation times of phages over bacteria – usually, such a correction is very small for non-microbial hosts and can be neglected. Ultimately, herd immunity is achieved if the threshold defined by $H = 1 - S_c$ is exceeded, with S_c the critical value in the inequality above. Rearranging, we obtain an expression for the herd immunity threshold

$$H = \frac{\beta - 1 - \lambda\alpha}{\beta} \,. \tag{4}$$

This estimate of H coincides to a very good extent with the population compositions of susceptible and resistant bacteria where we observe the transition from surviving and collapsed populations in experiments (see Fig 3). Simulations presented in the Appendix (section Simulation of recovery rate) show a range in the bacterial population composition with non-monotonic trajectories for B_s and B_r (see Fig 9B), which is comparable to the range in composition we find both outcomes, i.e., some surviving and some collapsing populations in experiments. For such parameter choices, stochastic effects could then decide the observed fates of bacteria.

As presented above, the herd immunity threshold changes when the bacterial cultures grow in a diluted growth medium. In a set of independent experiments we measured bacterial growth rate α, phage burst size β and phage latent period λ under such conditions (see Fig 6B and Table 2). From these data we estimated the dependence of the phage burst size on the bacterial growth rate, $\beta(\alpha)$, using a numerical quadratic fit (Fig 4A). Similarly, we estimated the dependence of the phage latent period on the bacterial growth rate, $\lambda(\alpha)$ (Fig 4B). Using these estimates we calculated the expected growth rate–dependent herd immunity threshold

$$H(\alpha) = \frac{\beta(\alpha) - 1 - \lambda(\alpha)\alpha}{\beta(\alpha)} \,, \tag{5}$$

which gives a very good prediction of the shift in the herd immunity threshold to lower values for slower growing populations (green line in Fig 4C). This verification of our model shows that it correctly captures the dependence of the herd immunity threshold on bacterial and phage growth parameters.

Modelling herd immunity in spatially structured populations

The dynamics of phage spread differ between growth in unstructured (e.g., liquid) and structured (e.g., plates) populations. In order to quantify the effect of spatial structure in a population, we extend our model to include a spatial dimension. In structured populations growth is a radial expansion of phages defined by the plaque radius r and the expansion speed v, for which several authors have previously derived predictions ***Kaplan et al.*** (***1981***); ***Yin and McCaskill*** (***1992***); ***You and Yin*** (***1999***); ***Fort and Méndez*** (***2002a***); ***Ortega-Cejas et al.*** (***2004***); ***Abedon and Culler*** (***2007***); ***Mitarai et al.*** (***2016***).

We assume phage movement can be captured by a diffusion process characterized with a diffusion constant D, which we estimate in independent experiments as $D = 1.17\,(\pm 0.26)\cdot 10^{-2}\ mm^2/h$ (see Materials and Methods, Fig 8). However, we assume that only phages disperse and bacteria are immobile as the rate of bacterial diffusion does not influence the expanding plaque on timescales relevant in the experiment. Adsorption of phages on bacteria is modeled with a adsorption constant $\delta^\star$. The spreading infection will sweep across the bacterial lawn with the following speed

$$v = 2\sqrt{D\delta^\star}\sqrt{\beta S - 1 - \lambda\alpha} \,, \tag{6}$$

which is computed in more details in the Materials and Methods. This expression (6) indicates that the population composition crucially influences the spreading speed at much lower fractions of

resistant bacteria than the herd immunity threshold (4), where phage expansion stops completely. Consequently, the resulting plaque radius r decays with increasing fractions of resistants and reaches zero at H. A prediction for r can be obtained by integrating (6) over time. Using the resulting expression we estimated the adsorption constant $\delta^\star$ from the growth experiments as it is difficult in practice to measure on plates. The green line in Fig 5B is the best fit, yielding the value $\delta^\star = 4.89(\pm 0.19) \cdot 10^{-2}\,\mathrm{bacteria/phage}\,h$ for the adsorption constant.

Our results for spatially structured populations allows us to speculate on a general epidemiological question: If an infection is not stopped by herd immunity in a partially resistant population, by how much is its spread slowed down? By generalizing (6) we can derive a relative expansion speed, compared to a fully susceptible population,

$$v_{\mathrm{rel}} = \sqrt{1 - \frac{1-S}{H}}\,. \tag{7}$$

This expression, (7), is devoid of any (explicit) environmental conditions, which are not already contained in the herd immunity threshold H itself. Thus, it could apply to any pathogen-host system. Ultimately, this relative speed approaches zero with a universal exponent of $1/2$, when the fraction of resistant individuals $1-S$ approaches the herd immunity threshold H. However, a few caveats exist when using (7), as several conditions have to be fulfilled: Obviously, a pathogen is expected not to spread in a population exhibiting complete herd immunity – the relative speed should only hold for populations below the herd immunity threshold. Moreover, if dispersal cannot be described by diffusion, but rather dominated by large jumps ***Hallatschek and Fisher*** (***2014***), the diffusion approach we used for traveling waves is not applicable, and thus also renders (7) inadequate.

Discussion

The spread of a pathogen may be halted or slowed by resistant individuals in a population and thus provide protection to susceptible individuals. This process, known as herd immunity, has been extensively studied in a wide diversity of higher organisms ***Jeltsch et al.*** (***1997***); ***Mariner et al.*** (***2012***); ***Boven et al.*** (***2008***); ***Meister et al.*** (***2008***); ***Konrad et al.*** (***2012***); ***Wang et al.*** (***2013***). However, the role of this process has largely been ignored in microbial communities. To delve into this we set out to determine under what conditions, if any, herd immunity might arise during a phage epidemic in bacterial populations as it could have profound implications for the ecology of bacterial communities.

We show that herd immunity can occur in phage-bacterial communities and that it strongly depends on bacterial growth rates and spatial population structure. Average growth rates of bacteria in the wild have been estimated as substantially slower than in the laboratory (generation time is $\approx$ 6.5 fold longer ***Gibson et al.*** (***2016***)), a condition that we have shown to facilitate herd immunity. Furthermore, bacterial populations in the wild are also highly structured, as bacteria readily form micro-colonies or biofilms ***Hall-Stoodley et al.*** (***2004***) and grow in spatially heterogeneous environments such as soil or the vertebrate gut ***Fierer and Jackson*** (***2006***), a second condition we have shown to facilitate herd immunity. These suggest that herd immunity may be fairly prevalent in low nutrient communities such as soil and oligotrophic marine environments.

In an evolutionary context, herd immunity may also impact the efficacy of selection as the selective advantage of a resistance allele will diminish as the frequency of the resistant allele in a population approaches the herd immunity threshold, H. This has two important implications. First, while complete selective sweeps result in the reduction of genetic diversity at linked loci, herd immunity may lead to only partial selective sweeps in which some diversity is maintained. Second, herd immunity has a potential to generate and maintain polymorphism at immunity loci, as has been shown for genes coding for the major histocompatibility complex (MHC) ***Wills and Green*** (***1995***). Polymorphism in CRISPR spacer contents have been demonstrated in various bacterial ***Tyson and Banfield*** (***2008***); ***Sun et al.*** (***2016***); ***Kuno et al.*** (***2014***) and Archaeal ***Held et al.*** (***2010***) populations and

communities ***Pride et al.*** (***2011***); ***Zhang et al.*** (***2013***); ***Andersson and Banfield*** (***2008***). While these studies primarily explain polymorphisms in CRISPR spacer content as a result of rapid simultaneous independent acquisition of new spacers, we suggest that observed polymorphisms may result from frequency-dependent selection on resistance loci arising from herd immunity. It has also been suggested that herd immunity might favor stable coexistence between hosts and their pathogens ***Hamer*** (***1906***); ***Fine*** (***1993***); ***Childs et al.*** (***2014***).

We also developed a mathematical model and show how the herd immunity threshold H (Eqn. (4)) depends on the phage burst size β and latent period λ, and on the bacterial growth rate α. This dependence arises as phages have to outgrow the growing bacterial population, as host and pathogen have similar generation times in our microbial setting. In addition to these parameters, we also describe how the speed v (Eqn. (6)) of a phage epidemic in spatially structured populations depends on phage diffusion constant D, phage adsorption rate $\delta^{\star}$, and the fraction of resistant and susceptible individuals in the population. All of which are likely to vary in natural populations. We also derived the relative speed of spread for partially resistant populations, as measured relative to a fully susceptible population, and show that it can be parametrized solely with the herd immunity threshold H (Eqn. (7)). This relative speed of the spread of an epidemic should be applicable to any spatially structured host population where the spread of the pathogen can be approximated by diffusion. Both our experiments and the modelling show that even when the fraction of resistant individuals in the population is below the herd immunity threshold the expansion of an epidemic can be substantially slowed, relative to a fully susceptible population.

In conclusion, we have presented an experimental model system and the connected theory that can be usefully applied to both microbial and non-microbial systems. Our theoretical framework can be useful for identifying critical parameters, such as H (and to some extent R_0), from the relative speed of an epidemic expansion in partially resistant populations so long as the process of pathogen spread can be approximated by diffusion. This approximation has been shown to be useful in such notable cases as rabies in English foxes ***Murray et al.*** (***1986***), potato late blight ***Scherm*** (***1996***), foot and mouth disease in feral pigs ***Pech and McIlroy*** (***1990***), and malaria in humans ***Gaudart et al.*** (***2010***).

Materials and Methods

Experimental methods

Engineering resistance

Oligonucleotides AAACTTCGGGAAGCACTTGTGGAAG and AAAACTTCCACAAGTGCTTCCCGAA were ordered from Sigma-Aldrich, annealed and ligated into pCas9 plasmid (pCas9 was a gift from Luciano Marraffini, Addgene plasmid #42876) carrying a *Streptococcus pyogenes* truncated CRISPR type II system. For the detailed protocol see ***Jiang et al.*** (***2013***). The oligonucleotides were chosen so that the CRISPR system targets an overlap of phage T7 genes 4A and 4B. Therefore, the CRISPR system allows the gene 0.7, coding for a protein which inhibits the RNA polymerase of the cell, to be expressed before the T7 DNA gets cleaved ***García and Molineux*** (***1995***). The subsequent growth of the cells is halted and phage replication is inhibited. The plasmid was electroporated into *Escherichia coli* K12 MG1655 (F- lambda- *ilvG*- *rfb*-50 *rph*-1). The T7 wildtype phage was used in all experiments.

Efficiency of the CRISPR-Cas system

Efficiency of the engineered CRISPR-Cas system was tested using the following protocol: Overnight culture was diluted 1 in 10, cells were infected with the T7 phage and incubated for $15\,\mathrm{min}$ in $30°C$. Cells were spun down for 2 min in room temperature at $21130g$. Supernatant was discarded and the cell pellet was resuspended in 950 μl of 1X Tris-HCl buffer containing 0.4% ($\approx 227\mu M$) ascorbic acid pre-warmed to $43°C$ and incubated in this temperature for 3 min to deactivate free phage particles ***Murata and Kitagawa*** (***1973***). Cultures were serially diluted and plated using standard plaque assay protocol on a bacterial lawn of susceptible cells to detect bursting infected cells. The supernatant was tested for free phage particles, which were not detected.

Determining the mean number of phages per cell

The cultures that were plated using standard plaque assays in the "Efficiency of the CRISPR-Cas system" experiment were also plated on LB agar plates containing 25 $\mu g/ml$ chloramphenicol to determine the number of surviving CFUs. The numbers of bursting and surviving susceptible cells were subsequently used to determine the actual mean number of adsorbed phages per cell. The fraction of susceptible cells surviving the phage challenge experiment was assumed to correspond to the Poisson probability that a cell does not encounter any phage, which was than used to determine the mean of the Poisson distribution, which corresponds to the mean number of phages per cell.

Herd immunity in a liquid culture

Herd immunity in a liquid culture was tested in 200 μl of LB broth supplemented with 25 $\mu g/ml$ chloramphenicol in Nunclon flat bottom 96 well plate in a Bio-Tek Synergy H1 Plate reader. Bacterial cultures were diluted 1 in 1000 and mixed in the following ratios of resistant to susceptible cells: 50:50, 75:25, 87.5:12.5, 93.75:6.25, 96.88:3.13, 98.44:1.56, 99.22:0.78, 99.61:0.39, 99.8:0.2, 99.9:0.1, 99.95:0.05, 100:0 %. T7 phage was added at a multiplicity of infection (MOI) of $\approx 10^{-4}$ (≈ 50 plaque forming units (pfu) per culture) to resemble an epidemic initiated by the burst size from one infected cell and the cultures were monitored at an optical density 600 nm for 18 hours post inoculation (hpi).

Time-lapse imaging of plaque growth

Soft LB agar (0.7%) containing 25 $\mu g/ml$ chloramphenicol was melted and poured into glass test tubes heated to $43°C$ in a heating block. After the temperature equilibrated, 0.9 ml of a bacterial culture consisting of resistant and susceptible cells (ratios 10% – 100% of susceptible cells, 10% increments) were diluted 1 in 10 and added to the tubes. Then, 100 μl of bacteriophage stock, diluted such that there would be ≈10 plaques per plate, was added to the solution. Tubes were vortexed thoroughly and poured as an overlay on LB agar plates containing 25 $\mu g/ml$ chloramphenicol. The plates were placed on scanners (Epson Perfection V600 Photo Scanner) and scanned every 20 minutes in "Wide Transparency mode" for 48 hours. A total of 3 scanners were employed with a total of 12 plates, plus a no phage control plate and 100% resistant control outside the scanners (see Fig 10). Time-lapse images were used to calculate the increase of individual plaque areas using image analysis software PerkinElmer Volocity v6.3.

Bacterial growth on soft agar

Growth rate of susceptible bacteria in soft LB agar (0.7%) was measured by sampling from a petri dish with a soft agar overlay with bacteria prepared in the same way as the plaque assays except an absence of the phage. Sampling was performed in spatially randomized quadruplicates at the beginning of the experiment and subsequently after 2, 4, 6, 8, 10, 12, 14, 16, 24, 32, 40, and 48 hours using sterile glass Pasteur pipettes (Fisherbrand art.no.: FB50251). Samples were blown out from the Pasteur pipette using an Accu-jet pro pipettor into 1 ml of M9 buffer pre-warmed to $43°C$, vortexed for 15 seconds and incubated for 10 minutes in $43°C$ with two more vortexing steps after 5 and 10 minutes of incubation. Samples were serially diluted and plated on LB agar plates containing 25 $\mu g/ml$ chloramphenicol. How bacterial densities change over time, measured as CFU/ml, is shown in Fig 6A.

Bacterial growth rates in liquid culture

Nutrient-dependent growth rate of susceptible bacteria was measured in Nunclon flat bottom 96 well plate in Bio-Tek Synergy H1 Plate reader in $30°C$. Overnight LB cultures were diluted 1:200 in media consisting of LB broth mixed with 1X M9 salts in ratios 10:90, 20:80, 30:70, 40:60, 50:50, 60:40, 70:30, 80:20, 90:10 and 100:0. Final volume was 200 μl. Optical density at 600 nm was measured every 10 min. Natural logarithm of the optical density values was calculated to determine the growth rate using a maximal slope of a linear regression of a sliding window spanning 90 min.

	Estimate	*Units*
$\alpha_{\max}$	$0.720\ (\pm 0.011)$	$[h^{-1}]$
K_c	$0.257\ (\pm 0.012)$	Dilution N of LB $[0 \dots 1]$

Table 1. Estimated parameters for bacterial growth using Monod kinetics. Undiluted LB medium ($N = 1$) is assumed to have $15\,mg/ml$ nutrients ($10\,mg/ml$ Tryptone, $5\,mg/ml$ yeast extract). The full dataset is shown in Fig 6.

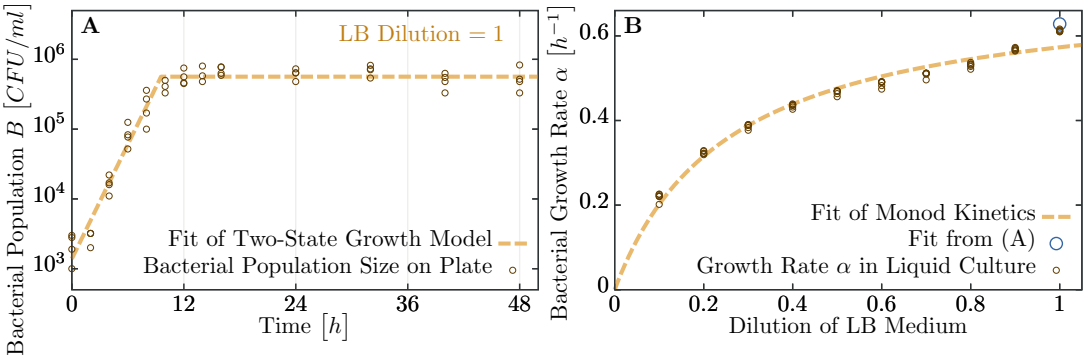


Figure 6. Measuring bacterial growth without phage. (A) Trajectory of population size on agar plates over time. For modeling, we assume two states of growth (green curve): first, the bacterial population grows exponential until the time T_{depl}, when nutrients are depleted. From this time on, growth rate is assumed to be zero and the population saturates at a maximal size B^{final}. Experimental observations fit this proposed growth curve to a very good extent. After all, half of all nutrients are used up in the last generation – with generation times of less than one hour, this the switch between growth and no-growth should be fast. (B) Growth rates of bacteria in diluted medium follow closely Monod's empirical law, given by expression (8). Fit parameters are found to be $\alpha_{\max} \approx 0.720\,h^{-1}$ and $K_c \approx 0.257$ (with the latter in dimensionless units as dilution of LB medium), see also Table 1.

Medium	Dilution N	Latent period $\lambda\ [min]$	Burst size β	Burst size/hour $\beta/\lambda\ [h^{-1}]$
LB 0	0.0	$101.1\ (\pm 10.9)$	$3.0\ (\pm 1.9)$	$1.8\ (\pm 1.1)$
LB 20	0.2	$43.4\ (\pm 3.9)$	$12.0\ (\pm 4.2)$	$16.6\ (\pm 6.0)$
LB 50	0.5	$40.0\ (\pm 3.0)$	$35.6\ (\pm 16.4)$	$53.4\ (\pm 24.9)$
LB 100	1.0	$36.1\ (\pm 6.1)$	$85.6\ (\pm 47.3)$	$142.1\ (\pm 82.1)$

Table 2. Estimated parameters for phage growth. See also Fig 4A,B.

The resulting growth rates for various nutrient concentrations fit well with Monod's growth kinetics,

$$\alpha = \alpha_{\max} \frac{N}{K_c + N}\ . \tag{8}$$

Results for the two fitting parameters, $\alpha_{\max}$ and K_c, are listed in Table 1. The whole dataset, including the fit, is displayed in Fig 6B.

Test for a difference in growth rates of resistant and susceptible bacteria was done in LB broth in the same manner as nutrient-dependent growth rate measurements. Two-sample t-test was performed on acquired growth rate data.

All growth media used in growth rate measurements were supplemented with 25 $\mu g/ml$ chloramphenicol.

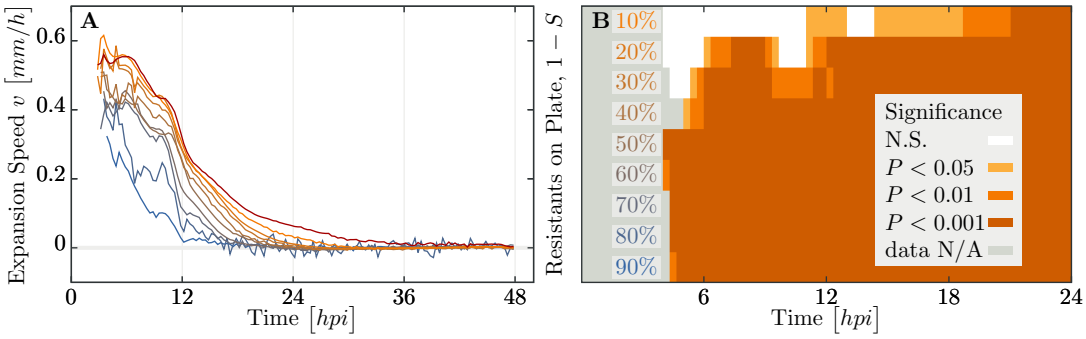


Figure 7. Speed of phage epidemic expansion on bacterial lawns. (A) Speed of expanding phage epidemics for all population compositions is initially high, before it drops once nutrients are depleted at around $10\,hpi$ (hours post infection). (B) Plaque speed significance. Comparing velocities of plaque spread with the 100% susceptible control. Linear regression of a sliding window spanning 4 hours of the radius sizes was calculated for all individual plaques and all compositions of the populations between t_0 and t_{24}. Slopes of the linear regressions for all compositions of the populations were compared using a two-sided heteroscedastic t-test against the 100% susceptible dataset.

Phage burst sizes

Phage burst sizes in bacteria growing at different growth rates were measured by one-step phage growth experiments. The burst sizes were calculated as the ratio of average number of plaque before burst to average number of plaques after burst. Consecutive sampling before and after burst were used for the calculation if they were not significantly different from each other (two sided t-test, $p > 0.05$).

Phage latent periods

Phage latent periods were determined as the time interval between the first and the last significantly different consecutive sampling between those used for phage burst size calculations.

Speed of phage expansion

The speed of the phage expansion was measured as difference in radii of plaques over time. Statistical tests allowed to infer that the reduction of expansion speed is significant already for small deviations from the 100% susceptible control experiment, as described and shown in Fig. 7.

Phage diffusion in soft agar

Soft M9 salts soft agar (0.5%) was supplemented with SYBR safe staining (final conc. 1%) and poured into glass cuvettes (VWR type 6040-OG) to fill $\sim 2\,cm$ of the cuvette height. After soft agar solidification, the same stained soft agar was supplemented with T7 phage particles to a final concentration $10^{11}\,\mathrm{pfu/ml}$ and poured on top of the agar without phages. The cuvettes were monitored in $30°C$ every hour for 40 hours at the SYBR safe emission spectrum peak wave length $524\,nm$ illuminated with the SYBR safe excitation spectrum peak wave length $509\,nm$. The diffusion constant was estimated as the best fit parameter for the spread of fluorescent phages through the soft agar over time.

First we computed the luminosity L_i of fluorescence (a gray-scale value defined as $L = 0.2126R + 0.7152G + 0.0722B$ from the RGB image) as average over the width of the cuvette for pixel row i, and corrected the profiles of luminosity L_i by subtracting the background value. This background value was estimated as a linear fit at the end of the profile without phages, where only the gray value of the agar was measured. Moreover, luminosity saturates at values above ~ 0.4 where it does not have a simple linear dependence on fluorescence: diffusion would lead to a decrease of the signal behind the inflection point of the profile and increase after the inflection point, but images only show increasing profiles – the bulk density does not decay. Thus, any estimate should only take the part of the profile that is below the threshold value of 0.4 into account (see Fig 8).

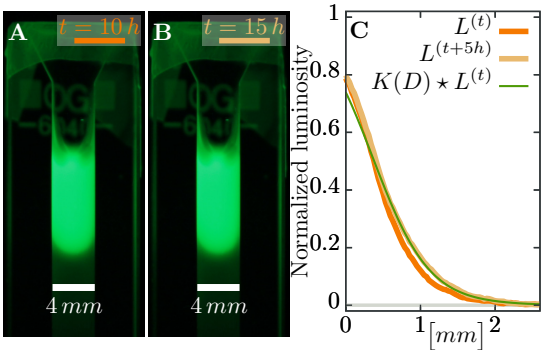


Figure 8. Estimating diffusion constant of phages. (A), (B) Phage are slowly expanding on agar which can be observed via their fluorescence. Pictures are taken $5\,h$ apart. (C) The diffusion constant D can be estimated as best-fit parameter in a *heat kernel* $K(D)$: $K(D)$ propagates the fluorescence profile $L^{(t)}$ at time t forward (via a convolution to "smear" out the signal) to the profile $L^{(t+\Delta t)}$ at the next measured time point. The difference between the expected change and the actual profile is quantified as total squared deviation, see Eqn. (9), which we minimize to obtain D. Consequently, we can estimate the diffusion constant as $D \approx 1.17 \cdot 10^{-2}\, mm^2/h$. The green line uses this estimated parameter D and shows the change between the profile at $t = 10\,h$ (orange line) and the profile at $t = 15\,h$ (light brown line), assuming diffusive spread of phages. See Materials and Methods for more information.

The diffusion constant D itself was estimated as the minimal value of the total squared deviation of the convoluted profile $L^{(t)}$ (at time t) with a heat kernel $K(D)$ compared to the profile $L^{(t+1)}$ at time $t+1$,

$$D = \left\langle \min_D \sum_i \left(\left(\sum_j \frac{e^{-(i-j)^2/4D}}{\sqrt{4\pi D}} L_j^{(t)} \right) - L_i^{(t+1)} \right)^2 \right\rangle . \tag{9}$$

Such a convolution with the heat kernel $K_{ij}(D) = (4\pi D)^{-1/2} \exp\left(-(i-j)^2/4D\right)$ assumes that the only change in the profile is due to diffusion for a time span of length 1 with i and j indices of pixels. Thus, expression (9) estimates the diffusion constant in units of $\mathrm{pixel}^2/\mathrm{frame}$, where frame is the time difference between two images. Several estimates are averaged over different snapshots in the whole experiment that spans $40\,h$ in intervals of $1\,h$ each.

The final estimate in appropriate units is

$$D \approx 1.17\ (\pm 0.26) \cdot 10^{-2}\ \ mm^2/h\ , \tag{10}$$

which is in agreement with previous measures of phage diffusion ***Stent and Wollman*** (***1952***); ***Bayer and DeBlois*** (***1974***); ***Briandet et al.*** (***2008***).

Modelling

Phage growth

In the main text we stated that relevant processes for phages growing on bacteria are given by the set of reactions (2). In the following, we will analyze an extended version of our model, which takes all these processes into account. We try to justify our approximations and explain the reasoning behind leaving parts of the full model out. While reactions for single bacteria or phages are inherently stochastic in nature, we assume that the involved numbers are large enough such that the dynamics can be described with deterministic differential equations for the populations. Furthermore, reaction rates are identified with the inverse of the average time scale of the process.

Thus, the full model is given by the coupled differential equations,

$$\begin{aligned}
\partial_t B_\mathrm{s} &= \alpha B_\mathrm{s} - A[B_\mathrm{s}, P|B_j] \,, && \text{(11a)}\\
\partial_t B_\mathrm{r} &= \alpha B_\mathrm{r} - A[B_\mathrm{r}, P|B_j] + \rho I_\mathrm{r} \,, && \text{(11b)}\\
\partial_t I_\mathrm{s} &= A[B_\mathrm{s}, P|B_j] - (1/\lambda) I_\mathrm{s} \,, && \text{(11c)}\\
\partial_t I_\mathrm{r} &= A[B_\mathrm{r}, P|B_j] - (1/\lambda) I_\mathrm{r} - \rho I_\mathrm{r} \,, && \text{(11d)}\\
\partial_t P &= (\beta/\lambda)\left(I_\mathrm{s} + I_\mathrm{r}\right) - \sum_{i\in\{\mathrm{s,r}\}} A[B_i, P|B_j] - \sum_{i\in\{\mathrm{s,r}\}} A[I_i, P|B_j] \,, && \text{(11e)}\\
\partial_t N &= -\alpha/Y\left(B_\mathrm{s} + B_\mathrm{r}\right) \,. && \text{(11f)}
\end{aligned}$$

Both bacterial populations B_i grow with rate α and decay via adsorption of phages $A[B_i, P|B_j]$, an expression that is specified below. Infected populations I_i gain numbers by adsorption and decrease via bursting. Resistant bacteria also can recover from their infected state with a recovery rate ρ. Phages grow by bursting cells, and lose numbers by adsorption to the various bacterial populations. Moreover, explicit dynamics for nutrients is considered, which are drained by each grown cell inversely proportional to the yield Y, the conversion factor between nutrient concentration and cell numbers. Essentially, this last equation acts as a timer, when we switch from abundant resources to the depleted state: all growth parameters change significantly upon nutrient depletion. Nevertheless, despite the possible deviations, we assume depletion time is given by the simple estimate (3) and only treat the two possible states of abundant and depleted nutrients.

Adsorption of phages, given by the term $A\left[B_i, P|B_j\right]$, can be influenced by the whole distribution of populations within the culture. In liquid medium, a common assumption is that this term is proportional to the concentrations of both the phages and cells ***Weitz*** (***2016***),

$$A\left[B_i, P|B_j\right] = \delta B_i P \,, \qquad (12)$$

with an adsorption constant δ. This expression assumes constant mixing of the population and relatively short contact times between phages and bacteria. In general, this system of equations is akin to Lotka-Volterra dynamics, which has been analyzed in great detail, eg. ***Hofbauer and Sigmund*** (***1998***); ***Nowak*** (***2006***).

For our ensuing analysis, we neglect the population of infected resistant bacteria I_r. Upon examining (11d) we find that most cells to leave their infected state by reducing phage DNA via CRISPR/Cas instead of bursting if $\rho \gg 1/\lambda$. If furthermore $\rho \gg \delta P$, which is true at least in the initial stages of the experiment, essentially all infected resistant bacteria immediately recover from a phage infection. Consequently, with both conditions, the resistant infected bacteria tend to vanish, $I_\mathrm{r} \to 0$, and their dynamics can be neglected. Only in the Appendix (section Simulation of recovery rate) we release this assumption to explicitly cover the full dynamics of (11) in simulations to estimate values for ρ.

Exponentially growing bacteria lead to double exponential phage growth

For convenience, we transform the populations to the total bacterial density $B = B_\mathrm{s} + B_\mathrm{r}$ and introduce the fraction of susceptible cells $S = B_\mathrm{s}/B$. The crucial assumption for the remainder of this section is that phages burst immediately after infection, $\lambda = 0$, such that we can ignore all infected populations. While not a very biological condition, it allows to analyze the model in more detail. Using these simplifications, we obtain

$$\begin{aligned}
\partial_t B &= (\alpha - \delta S P) B \,, && \text{(13a)}\\
\partial_t S &= -S(1-S)\delta P \,, && \text{(13b)}\\
\partial_t P &= (\beta S - 1)\delta B P \,. && \text{(13c)}
\end{aligned}$$

If we assume that in initial stages of phage growth the number of phages is small, ie. $\delta P \ll \alpha \sim \mathcal{O}(1\,h^{-1})$, the dynamics of bacteria and the fraction of susceptibles simplify to $\partial_t B = \alpha B$ and $\partial_t S = 0$.

Note that this term δP also occurs in the linear phage dynamics, where it cannot be neglected. In this instance, we need to view δB as a coefficient, which is likely much larger initially. This set of simplified equations can be solved in closed form,

$$S(t) = S_0 \,, \tag{14a}$$

$$B(t) = B_0 \exp(\alpha t) \,, \tag{14b}$$

$$P(t) = P_0 \exp\big((S_0\beta - 1)\delta B_0(\exp(\alpha t) - 1)/\alpha\big) \,. \tag{14c}$$

The structure of phage dynamics is particularly important here – it exhibits a double-exponential dependence on time t, which is a very fast, almost explosive, growth. Such double-exponential growth leads to very large population sizes within a short amount of time (but after an extended initial delay). This general behavior of the solution is independent of the actual growth rate of phages, which only has to be positive. Thus, inspecting the exponent in (14c) yields the condition

$$\beta S_0 > 1 \tag{15}$$

for phage growth to be positive. Incidentally, relation (15) is the naive estimate for the number of successful additional infections arising from a single burst. The double exponential time-dependence is central for our arguing that the dynamics can be described by threshold phenomena, given by conditions like (15): Usually, phages are negligible in the dynamics until they grow fast enough to large enough size, such that it is too late for the bacterial population to deal with the overwhelming phage population.

An important question in the context of these solutions is whether nutrients run out before this double-exponential growth of phages occurs. Hence, we compute the time T_δ defined as when phages reach a population of $P(T_\delta) = 1/\delta$ assuming phages grow as (14c) until then. After T_δ the assumptions that allowed to obtain (14c) are not valid anymore. Inverting (14c) for time leads to

$$T_\delta = \frac{1}{\alpha} \log\left(1 + \frac{\alpha \log(1/\delta P_0)}{(\beta S_0 - 1)\delta B_0}\right) \,. \tag{16}$$

Subsequently, we can compare this estimate T_δ to the depletion time $T_{\text{depl}} = (1/\alpha)\log(B_\infty/B_0)$. When rearranging the inequality $T_{\text{depl}} > T_\delta$ in terms of the (initial) fraction of susceptibles S_0, we obtain

$$\beta S_0 > 1 + \frac{\alpha \log(1/\delta P_0)}{\delta(B_\infty - B_0)} \,. \tag{17}$$

This expression (17) is a condition for phages to reach "large" population sizes before nutrients are depleted by bacteria. The final population density B_∞ usually fulfills $\delta B_\infty \gg 1$, such that the correction given by the second term of (17) can be considered small. Thus, if phages grow ($\beta S_0 > 1$), they also grow very fast with a double-exponential time-dependence and reach a considerably large population size before bacteria stop multiplying (for almost all parameter values).

Extending analysis to finite burst times

The analysis above only treated the case $\lambda \to 0$. However, we reported that the latency time λ increases significantly when bacterial growth rate α declines, see Table 2. Considering finite latency times entails dealing with an infected bacterial population I. (However, we identify $I \equiv I_s$ and set $I_r = 0$.)

To this end, note that we can rearrange (11a) to $\big(1 + \lambda\partial_t\big) I = \lambda\delta SBP$ using the adsorption model in (12). Hence, we can use the differential operator $(1+\lambda\partial_t)$ and apply it directly to (11e) to reduce the dependence on I in this equation at the cost of introducing higher order derivatives. In particular, we obtain

$$\lambda\partial_t^2 P + \big(1 + \lambda\delta B\big)\partial_t P + \delta B(\beta S - 1 - \lambda\alpha)P = 0 \,, \tag{18}$$

where we also inserted $\partial_t B \approx \alpha B$ in the last term, as we aim again for a solution at initial times where $\delta P \ll \alpha$. The effects of the limit $\lambda \to 0$ are directly observable – no terms are undefined in

this limit. In particular, we find that equation (18) and $\lambda = 0$ lead directly to the dynamics of phages we just analyzed above, obtaining solution (14c).

In principle, (18) is a hyperbolic reaction-diffusion-equation, which is known to occur upon transformation (or approximation) of time-delayed differential equations ***Fort and Méndez*** (***2002b***). For initial times we can use the solutions $B(t) = B_0 \exp(\alpha t)$ and $S(t) = S_0$. To proceed, we introduce the auxiliary variable

$$z(t) = -\delta B_0 \exp(\alpha t)/\alpha \ , \tag{19}$$

and assume $P(z)$ as a function of this new variable z. We need to transform the differential operators of time derivatives, and obtain $\partial_t = \frac{\partial z(t)}{\partial t}\partial_z = \alpha z \partial_z$ and $\partial_t^2 = (\alpha z \partial_z)(\alpha z \partial_z) = \alpha^2(z\partial_z + z^2\partial_z^2)$. Inserting these expressions in (18) and multiplying the whole equation with $(\alpha^2 \lambda z)^{-1}$ yields the dynamics for phages,

$$0 = z\partial_z^2 P(z) + (b - z)\partial_z P(z) - aP(z) \ , \tag{20}$$

where the two extant constants are $a = 1 - (\beta S_0 - 1)/(\lambda\alpha)$ and $b = 1 + 1/(\lambda\alpha)$. Equation (20) is called "Kummers equation" with confluent hypergeometric functions ${}_1F_1$ as solutions (***Abramowitz and Stegun, 1964***, pg. 504),

$$P(z) = A\ {}_1F_1\left(a, b; z\right) + B\, z^{1-b}\, {}_1F_1\left(a - b + 1, 2 - b; z\right) \ . \tag{21}$$

The two integration constants A and B can be determined via the initial conditions $P(t = 0) = P_0$ and $(\partial_t P)(t = 0) = -\delta B_0 P_0$. Using these conditions, the shape of the solution is again similar to before with $\lambda = 0$ (double exponential time-dependence), although $\lambda > 0$ introduces some skew. The most important aspect of this solution (21) is to compute the parameter combination where it switches from a decreasing to increasing function over time. A careful analysis reveals that at the parameter value $a = 0$ the behavior of the solution changes. Consequently, we find the condition for growing phage populations,

$$\beta S_0 > 1 + \lambda\alpha \ , \tag{22}$$

which is a non-trivial extension including finite latency times λ.

Note, however, that this relation (22) does not indicate a correction to the general epidemiological parameter R_0, which can be identified with β in our model. Rather, it shows that a growing bacterial population requires the phage population to grow even faster for a continuous chain of infections in an epidemic. The term $\lambda\alpha$ denotes the ratio of generation times of pathogen over host, which in most cases is small and negligible compared to 1. For bacteria and phages, however, which have similar generation times, such a correction is needed to describe the effects of growing host population sizes. In contrast, many other epidemiological models assume the host population size constant and only pathogens are increasing (or decreasing) in number.

While our result (22) suggest that it also should hold in the limit $\alpha \to 0$, it might not necessarily be so. This specific limit is actually quite important for the time when nutrients are depleted in the experiments. However, at several instances in the calculations above we implied a positive $\alpha > 0$. The most important of these is the transformation to $z(t) = -\delta B(t)/\alpha$, which actually exhibits two problems: dividing by α should not be allowed and $B(t)$ is essentially constant and cannot serve as a variable in a differential equation. We also neglected the second term in $\partial_t B = (\alpha - \delta S P)B$ throughout our calculation. For $\alpha = 0$ this second term is dominant in bacterial dynamics and would generate non-linear phage dynamics if inserted for $\partial_t B$ right before stating (18). However, we expect that albeit the process will run *very* slow, and might not be measurable in experiments, the simple condition $\beta S_0 > 1$ could indicate phage expansion and bacterial decay.

Growth of phages on plated bacterial lawn

Spatial modelling of phage expansion has produced several predictions for how plaque radius r and expansion speed v are influenced by experimentally adjustable parameters ***Kaplan et al.*** (***1981***); ***Yin and McCaskill*** (***1992***); ***You and Yin*** (***1999***); ***Fort and Méndez*** (***2002a***); ***Ortega-Cejas et al.*** (***2004***);

Abedon and Culler (***2007***); ***Mitarai et al.*** (***2016***). Here, we try to use a minimal model to estimate these two observables, based on the considerations of previous sections.

One of the main complications arises from the fact that all densities in (11) have a spatial dimension in addition to their time dependence, $B_i = B_i(\vec{x}, t)$, $i \in \{\mathrm{s}, \mathrm{r}\}$. As explained in the main text we only consider phage diffusion, heterogeneities in all other densities are generated only by phage growth. The additional spatial dimension imposes a particular contact network between phages and bacteria, which are not entirely random encounters anymore: One can expect that the size of the bacterial neighborhood $\hat{B}$ phages are able to explore is only slightly determined by the actual density B, and can be assumed constant for most of the experiment, $\hat{B}(B) \approx const$. Consequently, the adsorption term can be written in the following way,

$$A\left[B_i, P|B_j\right] = \delta^\star \frac{B_i}{\sum_j B_j} P \,, \tag{23}$$

which only depends on the *relative* frequencies of bacterial strains. The adsorption constant $\delta^\star$ is both the rate of adsorption and inter-host transit time as determined by the diffusion constant D. Thus, one can expect the formal dependence $\delta^\star = \delta^\star\left(D, \hat{B}(B)\right)$. For our particular experimental setup, however, $\delta^\star$ will be treated as a constant. This adsorption term (23) leads to the dynamics of phages

$$\partial_t P = D\nabla^2 P + G[P, S] \,, \tag{24}$$

where we collected all contributions to phage growth in $G[P, S]$ and added the spatial diffusion term $D\nabla^2 P$. For simplicity, we consider only expansion in a single dimension ($\nabla^2 \equiv \partial_x^2$), which has been found to coincide well with the dynamics of plaque growth ***Yin and McCaskill*** (***1992***). The growth term for phages is then defined as,

$$G\left[P, S\right] = \delta^\star\left(S\beta - 1 - \lambda\alpha\right) P \,, \tag{25}$$

where we also consider the correction $\lambda\alpha$ obtained from the analysis in liquid culture. Reaction-diffusion equations similar to (24) have been first analyzed almost 80 years ago ***Fisher*** (***1937***); ***Kolmogorov et al.*** (***1937***) and since then treated extensively, e.g. ***Murray*** (***2002***); ***van Saarloos*** (***2003***). They admit a traveling wave solution – here, this corresponds to phages sweeping over an uninfected bacterial lawn. In general, the asymptotic expansion speed for the traveling wave solutions is given by the following expression,

$$\begin{aligned} v &= 2\sqrt{D(\partial_P G)\left[0, S\right]} \\ &= 2\sqrt{D\delta^\star}\sqrt{S\beta - 1 - \lambda\alpha} \,. \end{aligned} \tag{26}$$

Only the linearized growth rate of phages at very low densities is relevant for the expansion speed, $\partial_P G\left[P = 0, S\right]$. Thus, the fraction of susceptible individuals S should be unchanged from its initial value S_0. It should be noted, that only for $S_0\beta > 1 + \lambda\alpha$ does Eqn. (26) remain valid, otherwise we have $v = 0$. Such a scenario is relevant when nutrients are depleted and phage growth parameters changes to β_{depl} and λ_{depl}.

The expression for the expansion speed also shows the need for the spatial adsorption model in (23), in contrast to the liquid case (12). If adsorption would directly depend on the bacterial density B, the additional linear dependence on B in (25) would lead to an exponentially increasing speed during the experiment. This is in clear contradiction to experimental observations.

The density of phages behind the expanding front is large and as previously noted at large MOIs the CRISPR-Cas system fails to provide effective immunity (see section Materials and Methods and appendix Infection load and efficiency of the CRISPR/Cas system). However, in comparison to an un-structured environment (e.g., liquid) the structured environment effectively limits transit of phage from within a plaque to the expanding front: The combined effect of growth and diffusion usually generates a much faster expansion of phages during plaque formation, than diffusion alone.

Only when nutrients are depleted, can pure diffusion processes explain the slow decrease in speed observed in experiments (see Fig 7A). Our model assumes a sharp drop to $v = 0$ at T_{depl} for small S.

In order to derive an expression for the plaque radius r, we integrate the expansion speed (6) over time, $r(t) = \int_0^t dt' v(t')$. Employing the simplification that only two values of phage growth are necessary to describe the dynamics – before T_{depl} phages grow normally with β and λ, after T_{depl} phage growth changes to β_{depl} and λ_{depl} – we can evaluate the integral for the radius directly, arriving at,

$$r(t) = \begin{cases} 2t\sqrt{D\delta^\star}\sqrt{S\beta - 1 - \lambda\alpha}\,, & 0 < t < T_{\text{depl}}\,, \\ 2\sqrt{D\delta^\star}\left(T_{\text{depl}}\sqrt{S\beta - 1 - \lambda\alpha} + \left(t - T_{\text{depl}}\right)\sqrt{S\beta_{\text{depl}} - 1}\right), & T_{\text{depl}} < t\,. \end{cases} \tag{27}$$

Using this expression we estimated the adsorption constant $\delta^\star$ from the growth experiments as it difficult to measure in practice. This estimate is done for radii exactly at the time of nutrient depletion T_{depl}, and excluding the control experiment with only susceptible cells.

Predictions of our model show a discrepancy from experimental results on plates. We independently estimated $\beta_{\text{depl}} = 3.0$, which results in $H_{\text{depl}} = \left(\beta_{\text{depl}} - 1\right)/\beta_{\text{depl}} \approx 0.67$. Thus, all experiments with $S > 0.33$ should exhibit expanding plaques after nutrients are depleted. In the experimental setup plaques stop expanding in all mixtures of resistant to susceptible cells ($S \leq 0.9$), which would correspond to $\beta_{\text{depl}} < 1.1$. This value is, however, still within experimental accuracy of our estimates of β_{depl}.

Acknowledgments

We are grateful to Remy Chait for his help and assistance with establishing our experimental setups and to Tobias Bergmiller for valuable insights into some specific experimental details. We thank Luciano Marraffini for donating us the pCas9 plasmid used in this study. Finally, we want to express our gratitude to Seth Barribeau, Andrea Betancourt, Călin Guet, Mato Lagator, Tiago Paixão and Maroš Pleška for valuable discussions on the manuscript.

Author Contributions

Conceived and designed the experiments: PP JPB. Performed the experiments: PP. Analyzed the data: PP LG. Conceived and designed the model: PP LG JPB NHB. Encoded and analyzed the model: LG. Wrote the manuscript: PP LG NHB JPB.

References

Abedon ST, Culler RR. Optimizing bacteriophage plaque fecundity. Journal of theoretical biology. 2007; 249(3):582–592. http://dx.doi.org/10.1016/j.jtbi.2007.08.006.

Abramowitz M, Stegun IA. Handbook of mathematical functions: with formulas, graphs, and mathematical tables, vol. 55. Courier Corporation; 1964.

Anderson RM, May RM. Directly transmitted infections diseases: control by vaccination. Science. 1982 Feb; 215(4536):1053–1060. http://science.sciencemag.org/content/215/4536/1053, doi: 10.1126/science.7063839.

Anderson RM, May RM. Spatial, temporal, and genetic heterogeneity in host populations and the design of immunization programmes. IMA journal of mathematics applied in medicine and biology. 1984; 1(3):233–266.

Anderson RM, May RM. Age-related changes in the rate of disease transmission: implications for the design of vaccination programmes. The Journal of Hygiene. 1985 Jun; 94(3):365–436. http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2129492/.

Anderson RM, May RM. Infectious Diseases of Humans: Dynamics and Control. OUP Oxford; 1992.

Andersson AF, Banfield JF. Virus Population Dynamics and Acquired Virus Resistance in Natural Microbial Communities. Science. 2008 May; 320(5879):1047–1050. http://science.sciencemag.org/content/320/5879/1047, doi: 10.1126/science.1157358.

649 **Bayer M**, DeBlois R. Diffusion Constant and Dimension of Bacteriophage φX174 as Determined by Self-Beat
650 Laser Light Spectroscopy and Electron Microscopy. Journal of virology. 1974; 14(4):975–980.

651 **Blumenthal RM**, Cheng X. Restriction-modification systems. Modern microbial genetics. 2002; p. 177–225.

652 **Boven Mv**, Bouma A, Fabri THF, Katsma E, Hartog L, Koch G. Herd immunity to Newcastle disease virus in
653 poultry by vaccination. Avian Pathology. 2008 Feb; 37(1):1–5. http://www.tandfonline.com/doi/abs/10.1080/
654 03079450701772391, doi: 10.1080/03079450701772391.

655 **Briandet R**, Lacroix-Gueu P, Renault M, Lecart S, Meylheuc T, Bidnenko E, Steenkeste K, Bellon-Fontaine MN,
656 Fontaine-Aupart MP. Fluorescence correlation spectroscopy to study diffusion and reaction of bacteriophages
657 inside biofilms. Applied and environmental microbiology. 2008; 74(7):2135–2143.

658 **Chabas H**, van Houte S, Høyland-Kroghsbo NM, Buckling A, Westra ER; **The Royal Society**. Immigration of
659 susceptible hosts triggers the evolution of alternative parasite defence strategies. Proc R Soc B. 2016;
660 283(1837):20160721.

661 **Childs LM**, England WE, Young MJ, Weitz JS, Whitaker RJ. CRISPR-induced distributed immunity in microbial
662 populations. PloS one. 2014; 9(7):e101710.

663 **Destoumieux-Garzón D**, Duquesne S, Peduzzi J, Goulard C, Desmadril M, Letellier L, Rebuffat S, Boulanger P.
664 The iron–siderophore transporter FhuA is the receptor for the antimicrobial peptide microcin J25: role of
665 the microcin Val11–Pro16 β-hairpin region in the recognition mechanism. Biochemical Journal. 2005 Aug;
666 389(3):869–876. http://www.biochemj.org/content/389/3/869, doi: 10.1042/BJ20042107.

667 **Fenner F**. Smallpox: emergence, global spread, and eradication. History and Philosophy of the Life Sciences.
668 1993; 15(3):397–420.

669 **Ferrari MJ**, Bansal S, Meyers LA, Bjornstad ON. Network frailty and the geometry of herd immunity. Proceedings
670 of the Royal Society B: Biological Sciences. 2006 Nov; 273(1602):2743–2748. http://rspb.royalsocietypublishing.
671 org/cgi/doi/10.1098/rspb.2006.3636, doi: 10.1098/rspb.2006.3636.

672 **Fierer N**, Jackson RB. The diversity and biogeography of soil bacterial communities. Proceedings of the National
673 Academy of Sciences of the United States of America. 2006 Jan; 103(3):626–631. http://www.pnas.org/content/
674 103/3/626, doi: 10.1073/pnas.0507535103.

675 **Fine PE**. Herd immunity: history, theory, practice. Epidemiologic Reviews. 1993; 15(2):265–302.

676 **Fine P**, Eames K, Heymann DL. "Herd Immunity": A Rough Guide. Clinical Infectious Diseases. 2011 Apr;
677 52(7):911–916. http://cid.oxfordjournals.org/content/52/7/911, doi: 10.1093/cid/cir007.

678 **Fisher RA**. The wave of advance of advantageous genes. Annals of Human Genetics. 1937; 7(4):355–369.

679 **Fort J**, Méndez V. Time-delayed spread of viruses in growing plaques. Physical review letters. 2002; 89(17):178101.
680 http://dx.doi.org/10.1103/PhysRevLett.89.178101.

681 **Fort J**, Méndez V. Wavefronts in time-delayed reaction-diffusion systems. Theory and comparison to experiment.
682 Reports on Progress in Physics. 2002; 65(6):895.

683 **Fox JP**, Elveback L, Scott W, Gatewood L, Ackerman E. Herd immunity: basic concept and relevance to public
684 health immunization practices. American Journal of Epidemiology. 1971 Sep; 94(3):179–189.

685 **García LR**, Molineux IJ. Rate of translocation of bacteriophage T7 DNA across the membranes of Escherichia coli.
686 Journal of Bacteriology. 1995 Jul; 177(14):4066–4076. http://www.ncbi.nlm.nih.gov/pmc/articles/PMC177138/.

687 **Gaudart J**, Ghassani M, Mintsa J, Rachdi M, Waku J, Demongeot J. Demography and Diffusion in Epidemics:
688 Malaria and Black Death Spread. Acta Biotheoretica. 2010; 58(2):277–305. http://dx.doi.org/10.1007/
689 s10441-010-9103-z, doi: 10.1007/s10441-010-9103-z.

690 **Gibson**, Wilson, Feil, Eyre-Walker. personal communication; 2016, unpublished results.

691 **Goldfarb T**, Sberro H, Weinstock E, Cohen O, Doron S, Charpak-Amikam Y, Afik S, Ofir G, Sorek R. BREX is a novel
692 phage resistance system widespread in microbial genomes. The EMBO journal. 2015 Jan; 34(2):169–183. doi:
693 10.15252/embj.201489455.

694 **Grassly NC**, Fraser C. Mathematical models of infectious disease transmission. Nature Reviews Microbi-
695 ology. 2008 Jun; 6(6):477–487. http://www.nature.com/nrmicro/journal/v6/n6/full/nrmicro1845.html, doi:
696 10.1038/nrmicro1845.

Hadas H, Einav M, Fishov I, Zaritsky A. Bacteriophage T4 development depends on the physiology of its host Escherichia coli. Microbiology (Reading, England). 1997 Jan; 143 (Pt 1):179–185.

Hall-Stoodley L, Costerton JW, Stoodley P. Bacterial biofilms: from the Natural environment to infectious diseases. Nature Reviews Microbiology. 2004 Feb; 2(2):95–108. http://www.nature.com/nrmicro/journal/v2/n2/abs/nrmicro821.html, doi: 10.1038/nrmicro821.

Hallatschek O, Fisher DS. Acceleration of evolutionary spread by long-range dispersal. Proceedings of the National Academy of Sciences. 2014; 111(46):E4911–E4919.

Hamer WH. Epidemic Disease in England: The Evidence of Variability and of Persistency of Type. Bedford Press; 1906.

Hammad Amm. Evaluation of alginate-encapsulated Azotobacter chroococcum as a phage-resistant and an effective inoculum. Journal of Basic Microbiology. 1998 Mar; 38(1):9–16. http://onlinelibrary.wiley.com/doi/10.1002/(SICI)1521-4028(199803)38:1<9::AID-JOBM9>3.0.CO;2-4/abstract, doi: 10.1002/(SICI)1521-4028(199803)38:1<9::AID-JOBM9>3.0.CO;2-4.

Heesterbeek JaP. A Brief History of R 0 and a Recipe for its Calculation. Acta Biotheoretica. 2002 Sep; 50(3):189–204. http://link.springer.com/article/10.1023/A%3A1016599411804, doi: 10.1023/A:1016599411804.

Held NL, Herrera A, Cadillo-Quiroz H, Whitaker RJ. CRISPR Associated Diversity within a Population of Sulfolobus islandicus. PLoS ONE. 2010 Sep; 5(9). http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2946923/, doi: 10.1371/journal.pone.0012988.

Hofbauer J, Sigmund K. Evolutionary games and population dynamics. Cambridge University Press; 1998.

Jeltsch F, Müller MS, Grimm V, Wissel C, Brandl R. Pattern formation triggered by rare events: lessons from the spread of rabies. Proceedings of the Royal Society of London B: Biological Sciences. 1997 Apr; 264(1381):495–503. http://rspb.royalsocietypublishing.org/content/264/1381/495, doi: 10.1098/rspb.1997.0071.

Jiang W, Bikard D, Cox D, Zhang F, Marraffini LA. RNA-guided editing of bacterial genomes using CRISPR-Cas systems. Nature Biotechnology. 2013; http://www.nature.com/nbt/journal/vaop/ncurrent/full/nbt.2508.html, doi: 10.1038/nbt.2508.

Kaplan DA, Naumovski L, Rothschild B, Collier RJ. Appendix: a model of plaque formation. Gene. 1981; 13(3):221–225. http://dx.doi.org/10.1016/0378-1119(81)90027-5.

Kolmogorov AN, Petrovsky I, Piscounoff N. Study of the diffusion equation with growth of the quantity of matter and its application to a biology problem. Bull Univ Moscow, Ser Int A. 1937; 1(1).

Konrad M, Vyleta ML, Theis FJ, Stock M, Tragust S, Klatt M, Drescher V, Marr C, Ugelvig LV, Cremer S. Social Transfer of Pathogenic Fungus Promotes Active Immunisation in Ant Colonies. PLOS Biol. 2012 Apr; 10(4):e1001300. http://journals.plos.org/plosbiology/article?id=10.1371/journal.pbio.1001300, doi: 10.1371/journal.pbio.1001300.

Kuno S, Sako Y, Yoshida T. Diversification of CRISPR within coexisting genotypes in a natural population of the bloom-forming cyanobacterium Microcystis aeruginosa. Microbiology. 2014; 160(5):903–916. http://mic.microbiologyresearch.org/content/journal/micro/10.1099/mic.0.073494-0, doi: 10.1099/mic.0.073494-0.

Lloyd AL, May RM. Spatial Heterogeneity in Epidemic Models. Journal of Theoretical Biology. 1996 Mar; 179(1):1–11. http://www.sciencedirect.com/science/article/pii/S0022519396900429, doi: 10.1006/jtbi.1996.0042.

Mariner JC, House JA, Mebus CA, Sollod AE, Chibeu D, Jones BA, Roeder PL, Admassu B, Klooster GGMv'. Rinderpest Eradication: Appropriate Technology and Social Innovations. Science. 2012 Sep; 337(6100):1309–1312. http://science.sciencemag.org/content/337/6100/1309, doi: 10.1126/science.1223805.

Meister T, Lussy H, Bakonyi T, Šikutová S, Rudolf I, Vogl W, Winkler H, Frey H, Hubálek Z, Nowotny N, Weissenböck H. Serological evidence of continuing high Usutu virus (Flaviviridae) activity and establishment of herd immunity in wild birds in Austria. Veterinary Microbiology. 2008 Mar; 127(3–4):237–248. http://www.sciencedirect.com/science/article/pii/S0378113507004154, doi: 10.1016/j.vetmic.2007.08.023.

Mitarai N, Brown S, Sneppen K. Population dynamics of phage and bacteria in spatially structured habitats using Phage λ and Escherichia coli. Journal of bacteriology. 2016; 198(12):1783–1793. http://dx.doi.org/10.1128/JB.00965-15.

745 **Murata A**, Kitagawa K. Mechanism of Inactivation of Bacteriophage J1 by Ascorbic Acid. Agricultural and
746 Biological Chemistry. 1973; 37(5):1145–1151. doi: 10.1271/bbb1961.37.1145.

747 **Murray JD**, Stanley EA, Brown DL. On the Spatial Spread of Rabies among Foxes. Proceedings of the Royal
748 Society of London B: Biological Sciences. 1986; 229(1255):111–150. http://rspb.royalsocietypublishing.org/
749 content/229/1255/111, doi: 10.1098/rspb.1986.0078.

750 **Murray JD**, Mathematical biology I: an introduction, Vol. 17 of interdisciplinary applied mathematics. Springer,
751 New York, NY, USA,; 2002.

752 **Nokes DJ**, Anderson RM. Measles, mumps, and rubella vaccine: what coverage to block transmission? The
753 Lancet. 1988; 332(8624):1374.

754 **Nordström K**, Forsgren A. Effect of Protein A on Adsorption of Bacteriophages to Staphylococcus aureus.
755 Journal of Virology. 1974 Aug; 14(2):198–202. http://jvi.asm.org/content/14/2/198.

756 **Nowak MA**. Evolutionary dynamics. Harvard University Press; 2006.

757 **Ortega-Cejas V**, Fort J, Méndez V, Campos D. Approximate solution to the speed of spreading viruses. Physical
758 Review E. 2004; 69(3):031909. http://dx.doi.org/10.1103/PhysRevE.69.031909.

759 **Pech RP**, McIlroy JC. A Model of the Velocity of Advance of Foot and Mouth Disease in Feral Pigs. Journal of
760 Applied Ecology. 1990; 27(2):635–650. http://www.jstor.org/stable/2404308.

761 **Pride DT**, Sun CL, Salzman J, Rao N, Loomer P, Armitage GC, Banfield JF, Relman DA. Analysis of strepto-
762 coccal CRISPRs from human saliva reveals substantial sequence diversity within and between subjects
763 over time. Genome Research. 2011 Jan; 21(1):126–136. http://genome.cshlp.org/content/21/1/126, doi:
764 10.1101/gr.111732.110.

765 **Real LA**, Biek R. Spatial dynamics and genetics of infectious diseases on heterogeneous landscapes. Journal of
766 The Royal Society Interface. 2007 Oct; 4(16):935–948. http://rsif.royalsocietypublishing.org/content/4/16/935,
767 doi: 10.1098/rsif.2007.1041.

768 **van Saarloos W**. Front propagation into unstable states. Physics reports. 2003; 386(2):29–222.

769 **Schenzle D**. An Age-Structured Model of Pre- and Post-Vaccination Measles Transmission. Mathematical
770 Medicine and Biology. 1984 Jan; 1(2):169–191. http://imammb.oxfordjournals.org/content/1/2/169, doi:
771 10.1093/imammb/1.2.169.

772 **Scherm H**. On the velocity of epidemic waves in model plant disease epidemics. Ecological Mod-
773 elling. 1996; 87(1–3):217 – 222. http://www.sciencedirect.com/science/article/pii/0304380095000305, doi:
774 http://dx.doi.org/10.1016/0304-3800(95)00030-5.

775 **Sorek R**, Lawrence CM, Wiedenheft B. CRISPR-Mediated Adaptive Immune Systems in Bacteria
776 and Archaea. Annual Review of Biochemistry. 2013; 82(1):237–266. http://dx.doi.org/10.1146/
777 annurev-biochem-072911-172315, doi: 10.1146/annurev-biochem-072911-172315.

778 **Stent GS**, Wollman EL. On the two-step nature of bacteriophage adsorption. Biochimica et biophysica acta.
779 1952; 8:260–269.

780 **Sun CL**, Thomas BC, Barrangou R, Banfield JF. Metagenomic reconstructions of bacterial CRISPR loci constrain
781 population histories. The ISME journal. 2016; 10(4):858–870. http://www.nature.com/ismej/journal/v10/n4/
782 abs/ismej2015162a.html.

783 **Sutherland IW**, Hughes KA, Skillman LC, Tait K. The interaction of phage and biofilms. FEMS microbiology
784 letters. 2004; 232(1):1–6.

785 **Swarts DC**, Jore MM, Westra ER, Zhu Y, Janssen JH, Snijders AP, Wang Y, Patel DJ, Berenguer J, Brouns SJJ, van der
786 Oost J. DNA-guided DNA interference by a prokaryotic Argonaute. Nature. 2014 Mar; 507(7491):258–261.
787 http://www.nature.com/nature/journal/v507/n7491/full/nature12971.html, doi: 10.1038/nature12971.

788 **Tyson GW**, Banfield JF. Rapidly evolving CRISPRs implicated in acquired resistance of microorganisms to viruses.
789 Environmental Microbiology. 2008 Jan; 10(1):200–207. http://onlinelibrary.wiley.com/doi/10.1111/j.1462-2920.
790 2007.01444.x/abstract, doi: 10.1111/j.1462-2920.2007.01444.x.

791 **Wang Y**, Yang P, Cui F, Kang L. Altered Immunity in Crowded Locust Reduced Fungal (Metarhizium anisopliae)
792 Pathogenesis. PLOS Pathog. 2013 Jan; 9(1):e1003102. http://journals.plos.org/plospathogens/article?id=10.
793 1371/journal.ppat.1003102, doi: 10.1371/journal.ppat.1003102.

Weitz JS. Quantitative Viral Ecology: Dynamics of Viruses and Their Microbial Hosts. Princeton University Press; 2016.

Westra ER, van Houte S, Oyesiku-Blakemore S, Makin B, Broniewski JM, Best A, Bondy-Denomy J, Davidson A, Boots M, Buckling A. Parasite exposure drives selective evolution of constitutive versus inducible defense. Current Biology. 2015; 25(8):1043–1049.

Wills C, Green DR. A Genetic Herd-Immunity Model for the Maintenance of MHC Polymorphism. Immunological reviews. 1995; 143(1):263–292. http://onlinelibrary.wiley.com/doi/10.1111/j.1600-065X.1995.tb00679.x/full.

Yin J, McCaskill J. Replication of viruses in a growing plaque: a reaction-diffusion model. Biophysical journal. 1992; 61(6):1540. http://dx.doi.org/10.1016%2FS0006-3495(92)81958-6.

Yorke JA, Nathanson N, Pianigiani G, Martin J. Seasonality and the requirements for perpetuation and eradication of viruses in populations. American Journal of Epidemiology. 1979 Feb; 109(2):103–123.

You L, Yin J. Amplification and spread of viruses in a growing plaque. Journal of theoretical biology. 1999; 200(4):365–373. http://dx.doi.org/10.1006/jtbi.1999.1001.

Zhang Q, Rho M, Tang H, Doak TG, Ye Y. CRISPR-Cas systems target a diverse collection of invasive mobile genetic elements in human microbiomes. Genome Biology. 2013; 14:R40. http://dx.doi.org/10.1186/gb-2013-14-4-r40, doi: 10.1186/gb-2013-14-4-r40.

Appendix: Additional theoretical considerations

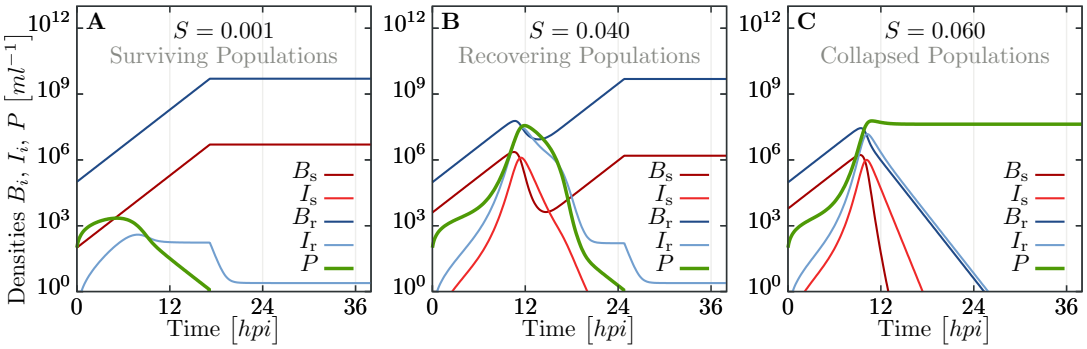
Simulation of recovery rate

Throughout the main text we assumed that resistant bacteria are completely immune to phage infection as their CRISPR/Cas system immediately kills adsorbed phages. However, experimental observation suggest that for fractions close to what we predicted as herd immunity threshold, *all* bacteria eventually die. Thus, in the following section we use numerical simulations to investigate the full set of equations (11), with a particular focus on the question why the whole bacterial population goes extinct. As it turns out, this requires using finite values for the recovery rate ρ (instead of the $\rho \to \infty$ approximation employed previously).

A major difficulty in analyzing the full model (11) is finding appropriate parameter values. In particular, we need values for the adsorption constant δ, the recovery rate ρ and the yield coefficient Y. Undiluted LB medium is known to support a population of $5 \cdot 10^9$ cells$/ml$. Thus one can easily estimate Y as the inverse of this number, when nutrients are measured in units of dilutions, which we already used throughout this publication (undiluted medium corresponds to $N = 1$). Parameter scans in simulations reveal that the actual value of the adsorption constant δ does usually not influence the actual outcome (collapsed or surviving bacterial population), it only adjusts time scales. However, deviations in time scales are insignificant, even when δ is changed by orders of magnitude, $\delta \sim \mathcal{O}\left(10^{-6} \dots 10^{-8}\right)$. They are roughly an hour or less, which is small compared to the expected duration of the experiment that lasts a few hours. For definiteness, we use the value of $\delta = 10^{-7}\ h^{-1}$ for our simulations. That the value of the adsorption constant has only a minor impact on phage growth on bacterial cultures, is also in line with previous findings ***Mitarai et al.*** (***2016***).

The most elusive parameter is the recovery rate ρ. A first indication of the value of ρ can be drawn from our experiments on bursting resistant cells, summarized in Fig 2. As the probability for bursting resistant cells is 3 orders of magnitude smaller than for susceptible bacteria, we can use $1/\lambda \sim \mathcal{O}(1)$ to estimate $\rho \sim \mathcal{O}\left(10^3\right)$. However, our results also indicate that recovery via the CRISPR/Cas system heavily depends on MOI, implying that ρ depends on the actual densities of phages and bacteria. Nevertheless, as experimental determination of recovery is complicated, even more so determining a functional dependence on dynamically changing densities B and P, we assume that ρ is constant.

We ran parameter sweeps in simulations and compared the outcome – collapsed or surviving bacterial populations – to the observed experimental results (see Fig 3). The best agreement of simulations and experiments was reached with $\rho \sim \mathcal{O}\left(1\right)$. Lower values of ρ do not allow the



Appendix 0 Figure 9. Simulated trajectories for all populations in liquid culture for the extended model, including infected and recovering bacteria. Trajectories are obtained by numerically integrating equations (11), using parameters listed in Table 3 and additionally $N = 1$, $Y = 2 \cdot 10^{-10}$ cells^{-1}, $\delta = 10^{-7}$ h^{-1} and $\rho = 1.5$ h^{-1}. (A) For population compositions with a large majority of resistant cells ($S = 10^{-3}$), phages get wiped out fast. (B) For intermediate S (close to parameters where we observe both, collapsed and surviving, populations, see Fig 3), the populations exhibit a complex, non-monotonic trajectory. After fast initial growth of phages, bacterial populations decay but ultimately can recover. (C) If the fraction of susceptibles is too large ($S = 0.06$), the whole bacterial population is infected and succumbs to the overwhelming phage infection. See supporting text for more detailed information.

resistant population to recover from phage infection, while for larger values of ρ, phages are drained from the culture very fast. Such a small value of ρ is most likely related to the recovery at very large MOI, when the densities involved in the dynamics are large, which dominate the overall observed dynamics. At this time phages repeatedly infect the same bacteria and their CRISPR/Cas immune system cannot deal with such an infection load (or only too slow). Thus, we can argue that our final choice $\rho = 1.5$ h^{-1} is the recovery rate when the CRISPR/Cas system is heavily stressed, which is comparable to the actual burst rate $1/\lambda$ for phages.

In Fig 9 we show three exemplary sets of trajectories for bacteria and phage. For a tiny fraction of susceptibles, $S = 10^{-3}$, which is well below the herd immunity threshold (see Fig 3), phages do not thrive on the limited number of favorable hosts and decay fast after a slight increase initially. For intermediate fractions of susceptibles, $S = 0.04$, we observe more complex, non-monotonic trajectories of bacterial populations. For such values of S we also observe mixed outcomes in experiments, see Fig 3. When S is increased further ($S = 0.06$), enough susceptible bacteria exists to produce enough phages and ultimately the whole bacterial population goes extinct.

The purpose of the extended model in this section was to justify the fact that phages can wipe out the whole bacterial population, which was not possible in the simplified model used in the main text. There, the resistant bacterial population was basically unaffected by phages and just acted as

Parameter		Value	Units	Comment
Bacterial growth rate	a	0.63	$1/h$	Table 1, Fig 6
Yield	Y	$2 \cdot 10^{-1}$	$1/\text{cell}$	Measured in dilution of LB
Recovery rate	ρ	1.5	$1/h$	See this appendix
Adsorption constant	δ	10^{-7}	$1/(h\,\text{cell})$	See this appendix
Diffusion constant	D	$1.17 \cdot 10^{-2}$	mm^2/h	See Methods
Burst size	β	85.6	phages/cell	Table 2, Fig 4
Latency time	λ	0.60	h	Table 2, Fig 4
Initial bacterial population	B_0	10^5	cells	
Initial phage population	P_0	10	phages	

Appendix 0 Table 3. Parameters used in simulations shown in Fig 9.

"sink" for phages. However, also in this extended model, we see a very similar behavior in terms of the threshold phenomena reported earlier in the manuscript.

Infection load and efficiency of the CRISPR/Cas system

In the section Modelling we showed that positive phage growth leads eventually to a very fast increase in the phage population, that occurs before nutrients are depleted (for almost all realistic parameters). This behavior of the dynamics was also observed in the extended simulation model presented in the last section. Moreover, as a condition we used that the phage population reaches a size $P \sim 1/\delta$, which is after all arbitrary – it only determines if we can employ useful simplifications and approximations to model equations. However, simulation results presented in the last section indicate that the bacterial population starts to decay soon after such a threshold $P \sim 1/\delta$ is exceeded.

In order to proceed, we investigate the system at time T_δ further. We assume that the phage population is large enough that it will not be degraded by the CRISPR/Cas immune system. The threat to immediate phage extinction is low at this point. The actual equations are hard to solve directly, hence we revert to simple balance equations, ignoring the dynamical component. Specifically, we compare the number of (present and eventually produced) phages to the number of infections needed to wipe out the whole population. To incorporate the effects of the bacterial immune system in resistant bacteria, we assume that they need $M > 1$ infections before they burst and produce only $\kappa\beta$ phages, which reduces the burst size by a (yet unspecified) factor $0 < \kappa < 1$. $\kappa = 1$ implies that resistant cells produce the same number of phages as susceptible cells, while $\kappa = 0$ indicates only cell death. Combining these considerations yields

$$\underbrace{1/\delta}_{\text{phages present}} + \underbrace{\beta S_0 B(T_\delta)}_{\text{phage production } B_\mathrm{s}} + \underbrace{\kappa\beta(1-S_0)B(T_\delta)}_{\text{phage production } B_\mathrm{r}} > \underbrace{S_0 B(T_\delta)}_{\text{infections } B_\mathrm{s}} + \underbrace{M(1-S_0)B(T_\delta)}_{\text{infections } B_\mathrm{r}}, \tag{28}$$

where the left side indicates the total number of phages, while the right side indicates the number of necessary infections to kill all bacteria. The number of bacteria $B(T_\delta)$ can be estimated by inserting the time T_δ from (17) into the exponential growth (14b). Subsequently, we can rearrange (28), obtaining a bound on M:

$$M < \frac{1/\delta B(T_\delta) + S_0(\beta - 1)}{1 - S_0} + \kappa\beta \,. \tag{29}$$

The first term $1/\delta B(T_\delta)$ indicates the ratio of phages to bacteria at time T_δ, and can be considered small for non-extremal parameters compared to the other terms. This fact justifies our assumption that the actual value of δ is not crucial. This number M might allow some insight into the effectiveness of the CRISPR/Cas immune system. For a fraction of susceptibles $S = 0.03$, which corresponds to the minimal value where we observe only collapsed bacterial populations in undiluted LB medium (see Figs 3 and 4), we would obtain the relation $M \lesssim 3 + 86\kappa$. Thus, each resistant bacterial cell could degrade up to $\mathcal{O}\left(10^1 \ldots 10^2\right)$ phages before their CRISPR/Cas system cannot cope with the infection load anymore.

Appendix 0 Figure 10. Image of the scanner system. Photograph of the scanner system used for time-lapse imaging of phage spread in spatially structured bacterial populations. Three scanners (Epson Perfection V600 Photo Scanner) simultaneously scanned 12 plates in total every 20 minutes in $30°C$ for 48 hours per experiment.

Appendix: Additional information on experimental setup

Our experimental setup for the scanner system is shown in Fig. 10.