Oscillations in continuous culture populations of Streptococcus pneumoniae: population dynamics and the evolution of clonal suicide

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Agents that kill or induce suicide in the organisms that produce them or other individuals of the same genotype are intriguing puzzles for ecologists and evolutionary biologists. When those organisms are pathogenic bacteria, these suicidal toxins have the added appeal as candidates for the development of narrow spectrum antibiotics to kill the pathogens that produce them. We show that when clinical as well as laboratory strains of \textit{Streptococcus pneumoniae} are maintained in continuous culture (chemostats), their densities oscillate by as much as five orders of magnitude with an apparently constant period. This dynamic, which is unanticipated for single clones of bacteria in chemostats, can be attributed to population-wide die-offs and recoveries. Using a combination of mathematical models and experiments with \textit{S. pneumoniae}, we present evidence that these die-offs can be attributed to the autocatalytic production of a toxin that lyses or induces autolysis in members of the clone that produces it. This toxin, which our evidence indicates is a protein, appears to be novel; \textit{S. pneumoniae} genetic constructs knocked out for \textit{lytA} and other genes coding for known candidates for this agent oscillate in chemostat culture. Since this toxin lyses different strains of \textit{S. pneumoniae} as well as other closely related species of \textit{Streptococcus}, we propose that its ecological role is as an allelopathic agent. Using a mathematical model, we explore the conditions under which toxins that kill members of the same clone that produces them can prevent established populations from invasion by different strains of the same or other species. We postulate that the production of the toxin observed here as well as other bacteria-produced toxins that kill members of the same genotype, ‘clonal suicide’, evolved and are maintained to prevent colonization of established populations by different strains of the same and closely related species.

**Keywords:** evolution; chemostats; \textit{Streptococcus pneumoniae}; allelopathy; population dynamics; mathematical models

1. INTRODUCTION

From an ecological and evolutionary perspective, it is not at all surprising that bacteria produce toxic agents that kill other species (Wiener 1996) or genetically different clones of the same species (Riley & Gordon 1999; Czaran \textit{et al.} 2002; Riley & Wertz 2002). More interesting evolutionarily are the toxins produced by bacteria that kill or induce suicide (apoptosis and autolysis) in genetically identical members of their own species. Included among these agents are the murein hydrolases and the choline-binding proteins responsible for autolysis and alloysis in \textit{Streptococcus pneumoniae} (Steinmoen \textit{et al.} 2002, 2003; Moscoso & Claverys 2004; Guiral \textit{et al.} 2005), and the proteins produced by sporulation-committed \textit{Bacillus subtilis} that kill non-committed members of the same genotype (Gonzalez-Pastor \textit{et al.} 2003; Ellermeier \textit{et al.} 2006; Claverys & Havarstein 2007).

Three general classes of hypotheses have been presented for the selection pressures responsible for the evolution and maintenance of mechanisms that are lethal to members of the same genotype. One has been likened to cannibalism, where the killed cells provide nutrients which in times of dearth enable the population at large to survive (Claverys & Havarstein 2007). A variant of this cannibalism hypothesis has been proposed for \textit{B. subtilis}, wherein the nutrients released by killed cells enable the remaining ‘killer’ cells to postpone the terminal commitment to sporulation, an energetically costly process that delays vegetative growth until new resources become available (Gonzalez-Pastor \textit{et al.} 2003). In the second class of hypothesis, gastronomy is not the reason for killing one’s own, but rather the acquisition of genetic information via transformation (a more intellectual goal?). In accord with this hypothesis, killing members of one’s own genotype is coincidental to the lysis of bacteria of the same or related species which may carry genes that provide an advantage to the killer population, perhaps by enabling them to invade new habitats or survive environmental stresses such as antibiotics (Claverys & Havarstein 2007; Claverys \textit{et al.} 2007). The third class of hypothesis of which we are aware considers lysis to be a stress response, which promotes the release of virulence factors (Pinas \textit{et al.} 2008). Presumably, these factors would facilitate the ability of the surviving members of the killer population to colonize susceptible hosts.

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Electronic supplementary material is available at \texttt{http://dx.doi.org/10.1098/rspb.2008.1415} or via \texttt{http://journals.royalsociety.org}.
In this paper, we present evidence for a new mechanism by which bacteria kill members of their same genotype that we discovered serendipitously. While performing experiments to estimate the rate constant of recombination (Levin 1981) in chemostat populations of *S. pneumoniae* (pneumococcus) strain R6, we observed oscillations in density of up to five orders of magnitude with an apparently constant period. These oscillations were not restricted to this domesticated laboratory strain but it was also obtained for the three clinical isolates of pneumococcus strain R6, we observed oscillations in (Levin 1981) in chemostat populations of *S. pneumoniae* (pneumococcus) strain R6, we observed oscillations in the concentration of an agent (a toxin) that is released by the bacteria via an autocatalytic process and either lyses bacteria via an autocatalytic process and either lyses or induces their autolysis. Our experiments indicate that the toxin responsible for these oscillations is a protein that can also lyse or induce lysis in closely but not distantly related species of *Streptococcus*. Since we can rule out the six known suspects anticipated for this lysis, we conjecture that this oscillation-driving toxin is novel, i.e. has not been previously identified or characterized.

We present and, with the aid of a mathematical model, explore the properties of a fourth hypothesis for the evolution and maintenance of the observed, and other, toxins that kill members of the same clone: allelopathy to prevent the invasion of established populations by competing strains of the same and different species. We briefly discuss why resistance to the suicidal agent observed here has not evolved and the potential for using these toxins for the treatment of bacterial infections.

### 2. MATERIAL AND METHODS

#### (a) Bacteria

The strains and species of bacteria used in this investigation, their genetic markers and their source are listed in table 1.

#### (b) Culture media

**(i) Liquid**

THY (per litre): 30 g Todd–Hewitt broth (Difco) plus 5 g yeast extract (Difco).

**(ii) Solid (agar) for sampling**

TSA: tryptic soy agar (Fisher) with 5 per cent sheep blood (Difco) and 16 g agar per litre.

MHA: Mueller–Hinton base (Difco) with 16 g agar per litre.

#### (c) Chemostat culture and sampling

The chemostats used in these experiments are ‘home-made’ (see www.eclf.net for the design and photographs). Chemostat populations were established from 80°C freezer stocks of *S. pneumoniae* via a two-step process: 200 µl aliquots of frozen cells were thawed in 2 ml of THY and grown to an OD₆₀₀ ≈ 0.3 (corresponding to a density of approximately 2e⁸

### Table 1. List of strains and *streptococcus* species used in this investigation. (The source of each strain and the relevant genotypic features for this study are provided. Genetic marker employed (when relevant) is also provided.)

| strain/species ID | relevant genotypic features | genetic marker | source |
|------------------|----------------------------|----------------|--------|
| R6 | non-encapsulated derivative of D39 | chromamphenicol resistant | laboratory isolate |
| LytA | Δ-LytA in the R6 background | kanamycin resistant | A. Tomasz |
| Δ-ComC originally FP5 | Δ-ComC in the RX1 background | | Iannelli et al. (2005) |
| Δ-ComD originally FP48 | Δ-ComD in the RX1 background | | McGee |
| PMEN-6 | PMEN-6 multiply antibiotic resistant | | |
| clinical isolate, serotype 23F | | |
| PMEN-18 | PMEN-18 multiply antibiotic resistant | | |
| clinical isolate, serotype 14 | | |
| PMEN-20 | PMEN-20 multiply antibiotic resistant | | |
| clinical isolate, serotype 6B | | |
| TIGR4 | encapsulated strain | streptomycin resistant | laboratory strain |
| TIGR4 Δ-spxB | Δ-spxB in the TIGR4 background | kanamycin resistant | Regev-Yochay et al. (2007) |
| 6A | encapsulated strain of serotype 6A | | |
| Δ-blpR | Δ-blpR in the 6A background | erythromycin resistant | Dawid et al. (2007) |
| RH-1 | R6 derivative, but *bgl*:spc | | |
| RH-3 | RH1 but Δ-ComE | | |
| RH-17 | RH1 but Δ-Cbdp | | |
| *Streptococcus mitis* | | | |
| *Streptococcus oralis* | | | |
| *Streptococcus sanguinis* | | | |
| *Streptococcus salivarius* | | | |
| *Streptococcus gordonii* | | | |
| *Streptococcus pyogenes* | | | |
| *Streptococcus australis* | | | |

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cells ml$^{-1}$) at which time they were diluted 10-fold in THY broth and regrown to OD$_{600}$ ≈ 0.3. Aliquots of 100 µl of these regrown cultures were inoculated into 20 ml of THY broth in the chemostat vessels for an initial density of approximately $1 \times 10^6$ cells ml$^{-1}$. The chemostat vessels (tubes) were incubated at 35°C in a water bath. A peristaltic pump maintained a constant flow of fresh media at a rate of 2 ml per hour, which is the same rate at which the waste, excess resources and live or dead cells were removed from the vessel. The chemostat cultures were agitated and aerated by a vacuum-induced bubbling of filter-sterilized air. The samples were taken by removing approximately 500 µl directly from the chemostat vessel. Viable cell densities (CFU) were estimated by serially diluting the samples in 0.85 per cent saline and plating on tryptic soy agar containing 5 per cent sheep blood. The experiments with low densities of *S. pneumoniae* were performed in chemostats with a reservoir containing THY and 0.85 per cent saline in a ratio of 1 : 28.

**Bioassay for toxin**

To test for the presence and relative concentration of a toxin, samples taken at different times were centrifuged for 2 min at 13 000g, and then the supernatant was stored at $\sim 20°C$. No CFUs were observed when 100 µl samples of the centrifuged supernatants were plated on tryptic soy agar with 5 per cent sheep blood, indicating that the supernatants were cell free. Between 1.5 and 2.0 µl of these supernatants were spotted onto soft-agar THY lawns of *S. pneumoniae* on a base of Mueller–Hinton (hard) agar. These lawns were prepared by growing *S. pneumoniae* R6 or other strains of this or other species of streptococci in THY to an OD$_{600}$ ≈ 0.3 nm and adding 100 µl to 4 ml of soft agar at approximately 55°C (approx. $5 \times 10^8$ cells ml$^{-1}$). The plates containing the supernatant-spotted lawns were scored after 24 hours of incubation at 35°C in a CO$_2$ incubator. To test whether the zones of inhibition on the agar were due to replicating phage, the material from zones of inhibition was added to fresh soft agar with *S. pneumoniae* and transferred to new lawns. No plaques were observed.

To ascertain whether the toxin was hydrogen peroxide (H$_2$O$_2$), we added approximately 5000 units of catalase (Worthington Biochemical) to 3 ml of soft agar used for the lawns, or incubated the supernatants from the chemostats for 30 min with 5000 units catalase before spotting. To determine whether the toxin had a protein nature, we incubated the supernatant at 55, 65 and 75°C for 30 min before assaying its activity. Also, prior to assaying for activity, the supernatant was treated for 30 min at 37°C with proteinase K at a final concentration of 1.5 mg ml$^{-1}$. Initial screening to determine the size range of the putative protein toxin was performed using membrane centrifugal filters (MicroconTM) with cut-offs of 3, 10, 30 and 50 kDa following the manufacturer's protocols.

**Numerical analysis (simulations)**

Numerical solutions to the differential equations of our models (simulations) were programmed in R (a free software environment for statistical computing) and/or BERKELEY MADONNA. The codes for these simulations are available on [www.ecclf.net](http://www.ecclf.net).

**RESULTS**

We open this paper with the serendipitous observation that motivated this investigation, profound oscillations in the density of *S. pneumoniae* R6 in a chemostat (figure 1a).

Similar observations were obtained with all three clinical isolates of *S. pneumoniae* that we examined (figure 2a). These dramatic oscillations in colony-forming units (CFU) are also apparent from changes in the optical density (OD$_{620}$ nm) in these cultures. If these bacteria were killed but not lysed, the OD of the chemostat flow rate employed (0.1 per hour) and the culture would remain turbid for more than 24 hours. Instead, we observed a complete loss of turbidity over roughly 10 hours, which we interpret as evidence that the decline in the CFU estimates of density in these chemostats can be attributed to the bacteria lysing.

By spotting the supernatants of centrifuged samples on soft-agar lawns of *S. pneumoniae* R6 (see §2), we assayed for the presence of an exogenous agent released by the bacteria, a ‘toxin’, that could account for the declines in the densities. The results of these assays were positive: zones of inhibition were observed (figure 1b). Based on the clarity of the zone, the concentration of this toxin appears to reach an apex shortly after the bacteria reach their maximum density (figure 1b).

In an effort to better understand the population dynamic mechanisms responsible for the observed oscillations in density and the conditions under which these
oscillations would occur, we developed a simple model of these population dynamics. In this model, $R$ is the concentration of a limiting resource (μg ml$^{-1}$), $B$ is the density of the bacteria (cells per ml) and $T$ is the concentration of the toxin (arbitrary units per ml). We assume a Monod (1949) model for resource concentration-dependent growth of the bacteria, $\Psi(R) = vR/(R+K)$, where $v$ is the maximum growth rate of the bacteria (per hour) and $K$ is the concentration of the resource at which the bacteria are growing at half their maximum rate (μg ml$^{-1}$). The limiting resource from a reservoir where it is maintained at a concentration of $C$ (μg ml$^{-1}$) enters a vessel of unit volume (1 ml) at a constant rate $w$ (per hour), which is the same rate at which excess resources, bacteria and toxin are removed. The bacteria take up the resource at a rate proportional to their density, growth rate and a conversion efficiency parameter, $e$ (μg) (Stewart & Levin 1973). In addition to being washed out with the flow, the toxin decays at a rate $d$ (per hour).

The toxin kills the bacteria at a rate equal to the product of its concentration, the bacterial density and a constant $x$. We assume that the toxin is produced at a rate proportional to the product of its concentration, the density of bacteria and a constant $y$. Our biological justification for including the concentration of the toxin in its rate of production is the observation that other known secreted products in S. pneumoniae, such as the quorum-sensing peptide responsible for the induction of competence in S. pneumoniae, the competence-stimulating peptide (CSP) and the bacteriocin-like protein (BLP), involve a positive feedback (autocatalysis) between the concentration of these agents and their production (Havarstein et al. 1995, 1996; Pestova et al. 1996; de Saizieu et al. 2000; Berge et al. 2002).

With these definitions and assumptions, the rates of change in the density of bacteria and concentration of the resource and toxin are given by

$$\frac{dR}{dt} = w(C-R) - \Psi(R)Be,$$

(3.1)

$$\frac{dB}{dt} = \Psi(R)B - xBT - wB,$$

(3.2)

$$\frac{dT}{dt} = yBT - dT - wT.$$

(3.3)

The density of bacteria that are killed by the toxin, $D$, is calculated from a fourth differential equation,

$$\frac{dD}{dt} = xBT - wD.$$

(3.4)

To illustrate the properties of this model, we use numerical solutions to these differential equations. The growth rate parameter $v$ used in these simulations is in the range estimated for S. pneumoniae R6 in the THY broth. The rate of flow, $w$, corresponded to that used in our experiments. The value of the Monod constant, $K$, is similar to that estimated for Escherichia coli in a glucose-limited minimal medium. For the maximum density (as determined by $C$ and $e$), the toxin killing and production constants ($x$ and $y$) and the toxin decay rate ($d$), we employed values that would produce the period and amplitude of the oscillations (figure 3a) similar to that observed in our experiments (figure 1a).

For an analytical understanding of the properties of this model and a dynamic summary of its properties, see the electronic supplementary material. The central point of this analysis is straightforward and interesting. Most importantly, this analysis shows that the resource concentration, $C$, and conversion efficiency, $e$, are not only the primary determinants of maximum population density but also control the dynamical behaviour of this system. To examine the transition between the different dynamic behaviours, we let $a = v/d + wyC$, $b = w$ (note that necessarily $b > 1$) and $f = (d + w)/w$ (with $k = K/C < 1$, typically $k \sim 10^{-2} - 10^{-5}$ holding for the different dynamics considered below). In accord with our analysis if and only if $b > a$ (with the parameters used in figure 3, this corresponds to $C > 50$), there exists a 'three-species' equilibrium with $R$, $B$ and $T$ all present. There is a very sharp transition in the dynamical behaviour when $a = 1$ and for $a < 1$ (i.e. $C > 500$), the oscillations generated decay very slowly (of the order of 10$^5$ hours, with parameters and initial values of the variables employed in our simulations; figure 3a). For $b > a > 1$, conditions that would be obtained when $C$ is lower (300 μg ml$^{-1}$), the oscillations are strongly damped (figure 3b).

To test the validity of the model's prediction that the oscillations would be damped if the cell density is lower, we inoculated S. pneumoniae R6 into a chemostat with the

Figure 2. (a) Clinical isolates show oscillatory dynamics that qualitatively resemble those obtained with the laboratory strains (black triangles, PMEN-18; grey squares, PMEN-6; grey circles, PMEN-20). The source of these and other strains used in this study are listed in table 1. (b) Cross activity of supernatants among different strains. Spot assays performed on lawns (backgrounds) of the laboratory strain R6, and three clinical isolates (PMEN-18, PMEN-6 and PMEN-20). The supernatant from single-clone chemostats of S. pneumoniae: (i) R6 (1), (ii) PMEN-6 (4), (iii) PMEN-20 (2) and (iv) PMEN-18 (3), and tested on lawns of each cell line.

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THY broth diluted 28-fold. The results of this experiment are presented in figure 4. After the population recovered from an initial decline in density, which is attributable to an extended lag period in this nutrient-poor medium, the maximum density achieved is approximately two orders of magnitude lower than the maximum densities we observed in chemostats run with the full-strength THY broth (figure 1a). As predicted by the model, the oscillations in the diluted medium are considerably damped (figure 4) relative to that in the undiluted THY broth (figure 1a). While we cannot rule out the possibility that the physiological effects of low nutrient concentrations, rather than the reduction in cell density, are responsible for the dampening of the oscillations, we admit to some satisfaction with the consistency between our theoretical predictions and experimental results. After all, hypotheses can only be rejected.

To date, our efforts to characterize this toxin have been through exclusion experiments using strains of *S. pneumoniae* R6, TIGR4 and 6A, which do not produce known candidates for this toxin, and a limited biochemical characterization of the toxic supernatant.

One candidate for this toxin is H2O2, which is released by *S. pneumoniae* at concentrations that are lethal to the cells of this and other species of bacteria (Regev-Yochay et al. 2006, 2007). To test whether H2O2 is responsible for the oscillations, we added approximately 5000 units of catalase (Worthington Biochemical) to the 3 ml of soft agar used to grow the lawns of target cells, or incubated the supernatants from the chemostats for 30 min with 5000 units catalase before spotting. With both the treatments, the catalase would hydrolyse any H2O2. The results of this test indicated that H2O2 is not the toxin; lysis was observed on *S. pneumoniae* lawns containing catalase and with catalase-treated chemostat supernatants (figure 1c). Additional evidence that the toxin is not H2O2 comes from the observation of density oscillations in chemostats inoculated with *S. pneumoniae* with a deletion in the gene encoding the synthesis of pyruvate oxidase (TIGR4ΔspxB) rendering this strain unable to produce hydrogen peroxide (Regev-Yochay et al. 2006, 2007; see figure S2 in the electronic supplementary material).

Our initial characterization of supernatants taken at the peak of toxin production suggests this agent is likely to be a protein. These supernatants do not generate zones of inhibition on lawns of *S. pneumoniae* R6 following incubation under denaturing conditions at relatively high temperatures or after protease treatment (figure 1c). Size fractionation using Microcon filters and the spot assay indicates that the relative size of the toxin is between 30 and 50 kDa (300 and 500 amino acids).

On first consideration, it may seem that the protein responsible for these oscillations is the same as that responsible for autolysis in batch cultures of *S. pneumoniae* (Sanchez-Puelles et al. 1986; Ronda et al. 1987; Havarstein et al. 2006). This does not appear to be the case. Mutants defective for the major pneumococcus lysis, *N*-acetylmuromoyl-l-alanine amidase (*lytA*), do not display autolysis in batch culture (figure 5a), yet they oscillate in density when introduced into a chemostat (figure 5b).

It is somewhat gratifying that the products of *lytA* and hydrogen peroxide can be ruled out as the toxins, because there is no evidence for their being produced autocatalytically as our model predicts they would have to be. On the other side, our experiments have also ruled out ‘suspects’ that directly or indirectly induce lysis in pneumococcus and for which there is evidence for autocatalytic production: CSP and BLP (Claverys & Havarstein 2002; Peterson et al. 2004; Dawid et al. 2007; Lux et al. 2007). In chemostat culture, mutants that are defective for the production of these secreted peptides (plus the CSP receptor, *comC*, *comD* and *blpR*), oscillate in a manner.

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similar to wild-type strain R6 in chemostat culture (see figures S3 and S4 in the electronic supplementary material). In addition, a mutant defective in the competence response regulator (Δ-ComE), which is unable to induce competence or activate downstream genes regulated by competence induction, also shows oscillatory dynamics in a chemostat similar to the parental strains from which it was derived (see figure S5 in the electronic supplementary material).

A particularly appealing candidate for the toxin responsible for the density oscillations is the murine hydrolase coded for by the gene CbpD. As Johnsborg et al. (2008) recently demonstrated using a β-galactosidase release assay, murine hydrolase lyses pneumococcus in liquid culture. This enzyme is approximately the molecular weight we estimated for the toxin in the supernatants of our chemostats, and, as we show below, the host range of bacteria sensitive to the S. pneumoniaeR6 murine hydrolase resembles that of the supernatant from the oscillating S. pneumoniaeR6 chemostats (Johnsborg et al. 2008). But unfortunately, strains of pneumococcus R6 knocked out for the production of murine hydrolase protein, ΔCbpD, oscillate to an extent no different from the ΔCbpD+ strain from whence it was derived (see figure S5 in the electronic supplementary material), and the supernatant of these ΔCbpD chemostats produced zones of inhibition on R6 lawns similar to those we observed with CbpD+ strains (data not shown).

Using the spot assay we tested the cross-sensitivity of R6 and the three clinical isolates to their respective secreted toxins. For each supernatant, zones of inhibition were observed on lawns of all four strains (figure 2b). To further explore the host range of the R6 toxin, we tested for lysis on lawns of the other Streptococcus species. The results of these assays suggest that closely, but not distantly, related species of Streptococcus are susceptible to the S. pneumoniaeR6 toxin (see figure S7 in the electronic supplementary material).

The observations that the activity of the toxin driving these oscillations does not appear to be strain specific and lyases other, closely related, species of Streptococcus raise the possibility (hypothesis) that the ecological role of these toxins is allelopathy. In this interpretation, these toxins either prevent the invasion of established populations by competing strains and species sensitive to this toxin or facilitate invasion into established populations of toxin-sensitive strain species. At this time, we have not tested this allelopathy hypothesis experimentally. However, using an extension of the model equations (3.1)–(3.3), we have explored the a priori plausibility of this hypothesis by ascertaining the conditions under which self-killing toxins would prevent invasion of competitors into established populations of toxin-producing bacteria. In this extended version of our model, there is a second population of bacteria, B2, that is susceptible to the toxin produced by the first, B1, but does not produce that toxin. The system of equations describing this situation is

\[
\frac{dR}{dt} = \omega(C-R) - \Psi(R)w(V_1B_1 + V_2B_2),
\]

\[
\frac{dB_1}{dt} = \Psi(R)V_1B_1 - x_1B_1T - wB_1,
\]

\[
\frac{dB_2}{dt} = \Psi(R)V_2B_2 - x_2B_2T - wB_2,
\]

\[
\frac{dy}{dt} = yB_1T - d_TT - wT,
\]

where \(\Psi(R) = \epsilon R(R + K)\); \(V_1\) and \(V_2\) are the maximum growth rates of these bacteria; and \(x_1\) and \(x_2\) are the killing rates of the producing and invading bacteria, respectively. For convenience, we assume that the efficiency of invasion of resource into bacteria biomass, \(\epsilon\), and the Monod constant, \(k\), are identical for \(B_1\) and \(B_2\). As in the previous model, \(y\) is the rate constant of production of the toxin and \(d_T\) is the inverse of the half-life of the toxin.

As indicated in the analysis of the properties of this model in the electronic supplementary material, there are conditions under which the production of a toxin that is lethal to the bacteria that produce it can prevent invasion by a toxin-sensitive population, even when that invader has a higher growth rate than the established population \(V_2 > V_1\). This, however, requires that the ratio of the rates of toxin-mediated killing of the invading and established strain, \(x_2/x_1\), is larger than \(V_2/V_1\) and that the density of \(B_1\) is sufficiently high to produce adequate concentrations of the toxin.

As can be seen in the simulations presented in figure 6a, in the absence of the toxin, the higher fitness clone, \(B_2\), will invade and replace the established population \(B_1\). If, however, the average density of \(B_1\) is high and \(x_2/x_1 > V_2/V_1\), the invasion of \(B_2\) can be prevented. This is obtained when the density of \(B_1\) oscillates (figure 6b) and when the resource concentration is too low for this established population to oscillate (figure 6c). However, if the density
of \( B_1 \) and thus the concentration of toxin in the environment are too low, the clone with the greater growth rate will invade (figure 6d), albeit at a rate lower than in the absence of toxin production (figure 6a). It should be noted, however, that the conditions for \( B_2 \) to invade an established population of \( B_1 \) are to some extent dependent on the initial density of \( B_1 \) and thus the concentration of the toxin, \( T \), in the environment. If the density of \( B_1 \) is initially low, \( B_2 \) can invade under conditions where it would not invade a toxin-producing \( B_1 \) population with an initially higher density (simulation results not shown). Our numerical simulation results indicate with reasonable parameter conditions, that even if \( B_2 \) has a higher growth rate fitness than \( B_1 \) and produces a toxin that kills \( B_1 \), when rare \( B_2 \) cannot colonize a community dominated by this high-density, toxin-producing established strain (results not shown).

4. DISCUSSION

When we set out to use chemostat cultures to estimate the rate of transformation-mediated chromosomal gene recombination in \( S. pneumoniae \) (now a tale for another report), we did not anticipate the profound oscillations in density reported here; this study is founded on a serendipitous observation. From a population dynamic perspective, these oscillations can be accounted for by the autocatalytic production of an agent, a toxin, which either directly lyse or induces autolysis in other cells of the same clone. The results of our experiments are consistent with the predictions of this model. The concentration of this toxin appears to reach its maximum shortly after the density of the oscillating population reaches its apex and these oscillations are damped when the density of the culture is low.

We have yet to characterize the toxin responsible for these oscillations or determine the physiological and molecular mechanisms by which it acts. Our results, however, indicate that this toxin is likely to be a protein of between 30 and 50 kDa. By exclusion, our experiments suggest that this toxin is novel in the sense of not being described or characterized earlier. Using strains of \( S. pneumoniae \) deleted for the genes coding for candidates for this toxin, we have excluded the following potential toxins or genetic systems involved in its production:

(i) \( N\)-acetylmuramoyl-L-alanine amidase, \( lytA \), which is responsible for autolysis in batch culture,
(ii) the competence-inducing peptide, \( comC \),
(iii) the BLP system \( blpR \),
(iv) murein hydrolase which lyses pneumococcus in liquid culture, \( CdpB \), and
(v) the cognate response regulator for the competence system, \( comE \).

The densities of strains that do not produce these peptides and proteins oscillate in a manner similar to that observed in the parental strains from which they were constructed. Moreover, using catalase and a strain that does not produce hydrogen peroxide, we have also

![Figure 6. Simulation results: invasion of a clone with higher intrinsic fitness. Changes in the densities of the bacterial populations and concentrations of the toxin (black dashed line) and resource (black dotted line). The parameter values for this simulation are: \( V_1 = 1.0; V_2 = 1.5; k = 0.25; e = 1 \times 10^{-7}; y = 4 \times 10^{-10}; \omega = 0.1; x_1 = 5 \times 10^{-6}; x_2 = 9 \times 10^{-6} \) and \( d = 0.1 \). In these simulations, a single cell of \( B_2 \) (grey solid line) is introduced into the populations of \( B_1 \) (black solid line) at an initial density of \( 10^8 \). In \( b \)–\( d \), the initial concentration of the toxin is 50 units per ml. \( a \) Invasion in the absence of toxin production (\( y = 0 \)). \( b \) Failure to invade a high-density (\( C = 1000 \)) oscillating toxin-producing population. \( c \) Failure to invade a lower density (\( C = 300 \)) non-oscillating toxin-producing population. \( d \) Invasion in the absence of the maintenance of toxin production because of very low resource concentration (\( C = 50 \)). In these simulations, when the density of the invading \( B_2 \) is less than 0.5, the simulation \( B_2 \) was set equal to zero.
excluded this oxidizing agent as being responsible for these density oscillations.

Although density oscillations were observed for all the strains we examined, there appears to be strain differences in the densities reached in these chemostats, with the same media and flow rates, and in the amplitude and, possibly, the period of these oscillations. For two reasons, we have not explored the variation in these dynamics systematically. First, strain variation in these dynamics is secondary to the main focus of this investigation, which is the population dynamic processes responsible for these oscillations and the ecological/evolutionary reason bacteria produce the self-killing toxins generating these oscillations. Second, the frequent sampling of viable cell density (CFU data) required for a quantitatively accurate characterization of oscillations with periods of 30 hours or so is an onerous task that we cannot justify doing given the motivation for this study.

Probably, for many readers and certainly for the authors, the most intriguing questions raised by this study are the ecological role and thereby the selection pressure responsible for the evolution of this oscillation-driving toxin. Despite the fact that this phenomenon occurs with natural isolates as well as laboratory strains of *S. pneumoniae*, it is conceivable that density oscillations of the amplitude observed in figure 1a are an artefact of laboratory culture. In their natural habitat within the human nasopharynx, *S. pneumoniae* may never reach the densities where these oscillations occur. As predicted by our model (figure 3b) and demonstrated experimentally (figure 4), oscillations in density of the amplitude observed in figure 1 are not anticipated in environments that can only support low-density populations. However, whether these oscillations are an artefact of culture conditions or not still remains necessary to account for why, evolutionarily, *S. pneumoniae* produce this and the other toxins that kill genetically identical bacteria.

If the killing of genetically identical members of the same population were the sole function of the toxin, the capacity to produce it or respond to its action would not be favoured by natural selection. If, however, the agent responsible for this killing provides a survival or growth advantage to the producing clone in its natural habitat, its production could be favoured by selection even if some or even many cells of the producing clone are killed. In §1, we briefly described three hypotheses that have been proposed to account for how the production of these suicidal agents could provide an advantage to the toxin-producing population:

(i) Cannibalism: to acquire resources (nutrients and carbon sources) from the killed cells that would increase the likelihood of survival of the whole clone in times of dearth (Sanchez-Puelles *et al.* 1986; Ronda *et al.* 1987; Lewis 2000; Claverys & Havarstein 2007) or postpone entry into a non-replicating state, sporulation (Gonzalez-Pastor *et al.* 2003).

(ii) The acquisition of adaptive genes through transformation of DNA by lysing cells of other clones and species that bear these genes (Lewis 2000; Steinmoen *et al.* 2002, 2003; Claverys & Havarstein 2007).

(iii) The release of other agents, such as virulence factors that would enable the population to colonize a new habitat or, for parasitic bacteria, an uncolonized host (Pinas *et al.* 2008).

Here, we propose a fourth hypothesis for the production of these clone suicidal toxins: allelopathy to prevent invasion of established populations. We find this hypothesis appealing not only because we presented it, but also because the toxin responsible for the observed oscillations kills other clones of the same species as well as closely related and thereby potentially competing species. Our theoretical analyses suggest that there are relatively broad conditions under which the production of toxins that kill bacteria of the same genotype that produce them can prevent invasion of established populations by toxin-sensitive populations that would otherwise invade because of a growth rate advantage. The necessary conditions for this to occur are that the invader is more sensitive to killing by the toxin than the resident and the concentration of the toxin in the habitat is sufficiently great for toxin-mediated killing to override the growth rate advantage of the invading population. A testable prediction of this hypothesis is that strains would be less sensitive to killing by the toxins they produce than they are to the toxins produced by other strains and species.

In this paper, we have not formally explored the other face of this allelopathy hypothesis, i.e. to facilitate the invasion of established populations of bacteria. For two reasons, we would anticipate that conditions for the invasion of a clone producing a clonal suicide toxin in mass (liquid) culture would be highly restrictive (Chao & Levin 1981; Levin 1988). (i) Unlike bacteriocins, there is no evidence for immunity to this toxin by the producing clone. (ii) All of the strains and species of *Streptococcus* we have examined that are sensitive to this toxin also produce that toxin or a variant of it to which all the other strains and species are sensitive. To be sure, in physically structured habitats, where bacteria are maintained as colonies rather than planktonic cells, there are broad conditions under which bacteriocin production can facilitate invasion of established sensitive populations (Wiener 1996; Kerr *et al.* 2002). Whether this would also be the case for clonal suicide toxins, and how the production of these toxins would affect the population dynamics of bacteria in physically structured habitats, remains to be seen and are intriguing problems for another time.

The four hypotheses for the evolution and maintenance of self-killing toxins considered here are not mutually exclusive and, to our knowledge, none of them has been adequately tested, much less unambiguously supported experimentally. To be sure, there is good evidence that the rate of transformation-mediated recombination is greater for *S. pneumoniae* that produce a lysis-inducing murein hydrolase protein than for otherwise isogenic cells that do not (Johnsborg *et al.* 2008). On the other hand, we know of no evidence that this higher rate of recombination would provide a sufficient advantage to the murein hydrolase-producing population to overcome likely fitness costs associated with synthesizing this protein and its killing isogenic cells.

Particularly intriguing from an evolutionary perspective is why pneumococcus remains susceptible to the toxins it
appealing if indeed viable strains of pneumococcus could the toxin responsible for the lysis reported here be pneumococcus drugs with novel targets will be needed. 2003), this may not be true in the future. Anti-S. pneumoniae matically colonizes many humans, consequently prevents effective chemotherapy (Yu et al., 2003), this may not be true in the future. Anti-pneumococcus drugs with novel targets will be needed. Could the toxin responsible for the lysis reported here be developed as an effective and safe drug to treat pneumococcus infections? This would be particularly appealing if indeed viable strains of pneumococcus resistant to this toxin cannot be generated.

We wish to thank, F. Iannelli for providing us with S. pneumoniae strains FP5 and FP48, Marc Lipstch for the Δ-spaB construct, Leslie McGee for the PMEN strains and ΔlatA mutant, and Suzanne Dawid for the Δ-blpR construct. We wish to acknowledge useful comments and suggestions by Sergei Pilyugin and Rustom Antia. Finally, we want to express our gratitude to Tony Romeo for his help and advice in our attempts to characterize this toxin and look forward to his joining us in continuing this quest. This research was supported by a grant from the US National Institutes of Health, AI40662 (B.R.L.). We wish to thank the reviewers of this report for their flattering as well as helpful comments and suggestions.

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