Antibiotic-Induced Cell Chaining Triggers Pneumococcal Competence by Reshaping Quorum Sensing to Autocrine-Like Signaling
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Graphical Abstract

Highlights
- Identification of a mechanism by which antibiotics induce competence in *S. pneumoniae*
- Antibiotics targeting penicillin-binding protein 3 promote chain formation
- Cell chains retain, rather than diffuse, the quorum-sensing peptide CSP
- Chaining populations feature a longer competence and transformation time window

In Brief
*Streptococcus pneumoniae* can take up exogenous DNA by activating competence. Aztreonam and clavulanic acid can induce competence by targeting PBP3, leading to cell chaining. Cell chaining reshapes quorum sensing to autocrine-like signaling and increases the time window in which cells can take up DNA, potentially accelerating the spread of antibiotic resistance.

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Antibiotic-Induced Cell Chaining Triggers Pneumococcal Competence by Reshaping Quorum Sensing to Autocrine-Like Signaling

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SUMMARY

Streptococcus pneumoniae can acquire antibiotic resistance by activation of competence and subsequent DNA uptake. Here, we demonstrate that aztreonam (ATM) and clavulanic acid (CLA) promote competence. We show that both compounds induce cell chain formation by targeting the β,δ-carboxypeptidase PBP3. In support of the hypothesis that chain formation alters CSP diffusion kinetics, Indeed, ATM or CLA presence affects competence synchronization by shifting from global to local quorum sensing, as CSP is primarily retained to chained cells, rather than shared in a common pool. Importantly, autocrine-like signaling prolongs the time window in which the population is able to take up DNA. Together, these insights demonstrate the versatility of quorum sensing and highlight the importance of an accurate antibiotic prescription.

INTRODUCTION

Streptococcus pneumoniae (the pneumococcus) is a member of the commensal microbiota of the human nasopharynx. However, it is also considered one of the leading bacterial causes of morbidity and mortality worldwide, being responsible for a wide variety of invasive and non-invasive diseases (Prina et al., 2015; Wahl et al., 2018).

Transformation, defined as the uptake and assimilation of exogenous DNA, is an important mechanism largely responsible for the rapid spread of antimicrobial resistance in the pneumococcus (Croucher et al., 2011). This process is regulated by competence (Figure 1A), a physiological state that involves about 10% of the pneumococcal genome (Aprianto et al., 2018; Claverys et al., 2009). Competence is induced by a classical two-component quorum-sensing system in which the comC-encoded competence-stimulating peptide (CSP) is cleaved and exported by the membrane transporter ComAB to the extracellular space. CSP stimulates autophosphorylation of the membrane-bound histidine-kinase ComD, which subsequently activates the cognate response regulator ComE (Figure 1A) (Martin et al., 2013; Pestova et al., 1996). Upon a certain threshold CSP concentration, a positive-feedback loop overcomes counteracting processes and the competent state is fully activated. One of the genes regulated by ComE, comX, encodes a sigma factor, which activates the genes required for DNA repair, DNA uptake, and transformation. CSP can be retained by producing cells (Prudhomme et al., 2016), but CSP also diffuses and can induce competence in neighboring cells (Christie, 2016; Hävarstein et al., 1995; Moreno-Gámez et al., 2017). Other environmental factors such as pH, oxygen, phosphate, and diffusibility of the growth medium also influence competence development (Chen and Morrison, 1987; Claverys and Havarstein, 2002; Echeneique et al., 2000). Thus, the initiation of competence can be considered as a combination of diffusion sensing and autocrine-like signaling (Doganer et al., 2016; Moreno-Gámez et al., 2017).

The competent state is activated in response to several antibiotics, which thereby allow the bacterium to take up foreign DNA and potentially acquire antimicrobial resistance determinants (Prudhomme et al., 2006; Slager et al., 2014; Stevens et al., 2011). Spread of antibiotic resistance is exacerbated by the fact that, coregulated with competence, S. pneumoniae expresses several bacterial killing factors, thereby using interbacterial predation to acquire foreign DNA (Kjos et al., 2016; Veening and Blokesch, 2017; Wholey et al., 2016).

We have shown previously that antimicrobials targeting DNA replication, such as fluoroquinolones, cause an increase in the copy number of genes proximal to the origin of replication (oriC) due to replication stalling (Slager et al., 2014). As the competence operons comAB and comCDE are located near oriC, these antibiotics induce competence. Aminoglycoside antibiotics such as kanamycin are thought to activate competence by causing the accumulation of misfolded proteins via mistranslation. Since these misfolded proteins are targeted by
Identification of Clinically Relevant Antibiotics that Induce Competence

To monitor competence development, we utilized the ComX-dependent promoter P\textsubscript{ssbB}, driving expression of firefly luciferase (luc). We selected antibiotics on basis of their use for the treatment of several pneumococcal respiratory infections (otitis media, pneumonia, or exacerbations of chronic respiratory diseases), as well as for the treatment of respiratory infections with other bacterial etiologies (Table S1). Cells of encapsulated strain D39V (Slager et al., 2018) were grown in C+Y medium at pH 7.3, a pH non-permissive for natural competence development under our experimental conditions (Moreno-Gámez et al., 2017), and antibiotics were added at concentrations below the minimum inhibitory concentration (MIC) to prevent large growth defects and cell killing. Only when antibiotics induce competence, the ssbB promoter is activated and firefly luciferase is produced. In line with previous reports, four antibiotics belonging to the fluoroquinolone and aminoglycoside classes of antibiotics robustly induced competence (Figure 1B) (Moreno-Gámez et al., 2017; Prudhomme et al., 2006; Slager et al., 2014; Stevens et al., 2011). Antibiotics from the macrolide and linezolid classes were not able to induce competence (Table S1).

The beta-lactam subclass antibiotics, carbapenems and cephalosporins, also did not induce competence at any of the concentrations tested (Table S1). In contrast, the addition of ATM and the combination of amoxicillin and CLA resulted in activation of P\textsubscript{ssbB-luc}. To test whether amoxicillin, CLA, or the combination of amoxicillin-CLA was responsible for competence induction, the compounds were also tested individually. Surprisingly, competence was not induced by the beta-lactam amoxicillin, but by CLA, an inhibitor of beta-lactamases. As the human nasopharynx is often colonized by non-typeable pneumococci, characterized by the absence of a polysaccharide capsule (Sá-Leão et al., 2006), we also tested whether ATM and CLA could induce competence in an unencapsulated derivative strain (strain ADP26). The deletion of the capsule did not affect competence induction by either of the drugs (Figure S1A).

To confirm whether ATM and CLA induce competence in a strain with reduced susceptibility to beta-lactams, we tested a strain (ADP305) with a mutation in PBP2X (PBP2X\textsuperscript{T550S}), which confers a MIC of 0.5 \mu g/mL and 0.64 \mu g/mL to penicillin G and cefotaxime, respectively. As shown in Figure S1B, both antibiotics were still able to induce competence in this strain. Together, this now extends the list of antibiotics capable of inducing competence to the following compounds: HPUra, mitomycin C, hydroxyurea, aminoglycosides, fluoroquinolones, trimethoprim, the beta-lactam ATM, and the inhibitor of beta-lactamases CLA.

ATM and CLA Promote Horizontal Gene Transfer

To examine whether competence induction by ATM and CLA leads to increased horizontal gene transfer (HGT), we co-incubated two pneumococcal strains that are genetically identical except for a unique antibiotic resistance marker (tetracycline and kanamycin, respectively) integrated at different genomic locations. Since the extracellular pH is an important factor for...
ATM and CLA Do Not Induce Competence via HtrA or Altering Gene Dosage

So far, two different molecular mechanisms of competence induction by antibiotics have been described. The first mechanism is via substrate competition of the HtrA protease, which degrades both CSP and misfolded proteins (Cassone et al., 2012; Stevens et al., 2011), and the second via gene dosage alterations leading to higher comAB and comCDE copy numbers (Slager et al., 2014).

We confirmed that strain ADP309, carrying a mutation in htrA that renders the catalytic domain inactive (HtrAS234A), is hyper-competent compared with the wild-type (Figure S2A) (Stevens et al., 2011). However, competence was still induced in this strain by ATM and CLA, as well as by the aminoglycosides gentamycin and tobramycin (Figure S2B).

To test whether ATM and CLA induce competence via altering the gene dosage of the early competence operons, we performed marker frequency analysis. As shown in Figure 2A, a shift in origin-to-terminus ratio was observed after the addition of HPura; however, the presence of ATM or CLA did not lead to an increase of the oriC-ter ratio. HPura analysis from Slager et al. (2014) is shown in red, as a positive control of oriC-ter ratio shift. Abbreviations: RE, rapid exposure; AE, adaptive exposure.

ATM and CLA Target PBP3 and Induce Cell Chaining

It is well known that both ATM and CLA have an impact on cell wall synthesis. Specifically, it has been shown that they can directly interact with PBP3 (Kocaoglu et al., 2015; Severin et al., 1997). To assess whether perturbing cell wall synthesis could lead to activation of competence, we employed clustered regularly interspaced short palindromic repeats (CRISPR) interference (CRISPRi), allowing us to downregulate essential genes involved in cell wall biosynthesis (Liu et al., 2017). Competence development was not influenced by downregulation of either genes involved in peptidoglycan precursor synthesis (murA-F) or genes encoding class B PBPs (transpeptidase only) ppb2b and ppb2x (Figure S3). However, when the genes encoding class A (dual transglycosylase and transpeptidase) PBP1A, or the d,D-carboxypeptidase PBP3 were repressed using CRISPRi,
Competence was strongly induced under otherwise non-permissive conditions (Figure 3A). To corroborate that pbp2b and pbp2x do not upregulate competence, we repeated the same experiment in a permissive pH for natural competence. As expected, pbp1a and pbp3 repression resulted in a stronger induction of competence, while repression of pbp2b and pbp2x did not influence competence development (Figure S2B).

To confirm that ATM and CLA bind PBP1A and/or PBP3, we used fluorescently labeled Bocillin (Bocillin-FL). As shown before (Kocaoglu et al., 2015; Severin et al., 1997), ATM and CLA bind PBP3, with ATM having a higher affinity to PBP3 than CLA (Figure 3B). As we were not able to clearly separate PBP1A and PBP1B, we cannot conclude whether ATM and/or CLA also bind to one of these PBPs. Since pbp3 is not essential (Li et al., 2017), we constructed a deletion mutant. In line with the CRISPRi results, the Δpbp3 strain (strain ADP30) displayed a hyper-competent phenotype (Figure S4A). Importantly, ATM and CLA did not further induce competence in the Δpbp3 strain, indicating that PBP3 is the main target of these compounds (Figure S4A).

To examine the effects of ATM and CLA and the downregulation of pbp1a and pbp3 on cell morphology, we performed microscopy analysis on exponentially growing cells (optical density measured at 595 nm [OD595], 0.1). In contrast to cells with downregulated pbp2b or pbp2x (Berg et al., 2013; Land et al., 2013; Liu et al., 2017; Peters et al., 2014), individual cell size and morphology were only slightly altered by ATM, CLA, or pbp1a and pbp3 perturbation. However, in all cases, pneumococci formed longer chains of unseparated cells (Figure S5A). When cells were grown until stationary phase (OD595, 0.4), chain formation was even more evident (Figure 3D).

Other beta-lactams, such as amoxicillin, ampicillin, piperacillin, or cefotaxime, also have a strong affinity for PBP3, but also for PBP2X (Kocaoglu et al., 2015). The inactivation of PBP2X seems to counteract the effect of PBP3 depletion, because these drugs did not induce chain formation (Figure S5B). These results suggest that cell chaining by ATM and CLA could be responsible for competence induction. To test this hypothesis, we performed a multi-dose checkerboard experiment with eight different concentrations of both ATM and CLA (Figure S5C). Indeed, the effects of ATM and CLA on competence activation are additive, until a certain maximum effect size, likely corresponding to the maximum chaining capacity.

Cell Chaining Is Responsible for ATM- and CLA-Induced Competence

To test whether ATM and CLA induce competence by specific binding to PBP3 or because of cell chaining, we generated a knockout of the gene encoding the major autolysin LytB (strain ADP21). LytB mutants are well known to form chains due to their lack in muralytic activity at cell poles (De Las Rivas et al., 2002; Garcia et al., 1999; Rico-Lastres et al., 2015). In line with the hypothesis that cell chaining induces competence, the ΔlytB mutant showed a hypercompetent phenotype, and readily developed competence even at pH 7.3, at which wild-type cells do not become naturally competent (Figure S4B). Importantly, complementation by ectopic expression of LytB in the ΔlytB (ADP43) restored the normal diplococcus phenotype and restored competence development to wild-type-like (Figures 3D, S5A, and S5D).

Finally, to test whether ATM or CLA induction is lost in the ΔlytB mutant, we have tested the effect of ATM and CLA in the ΔlytB and the complementation strain (Figure S5D). In the absence of isopropyl β-D-1-thiogalactopyranoside (IPTG) (chaining phenotype; Figure S5D), this strain is naturally hypercompetent. Under these conditions, ATM and CLA can only slightly accelerate competence development, relative to the control condition. LytB complementation by the addition of IPTG in ADP43 restores the normal phenotype, and as a result, the strain behaves as DLA3, confirming the role of chain formation in the regulation of competence (Figure S5D).
for the two lowest inoculation densities, due to the OD detection limit of the rapid synchronization of the population, we repeated a previous experiment. To first establish whether CSP produced by our wild-type pneumococci expressing a SsbB-GFP fusion (Aprianto et al., 2016) together with a ΔcomC mutant strain that also contains the SsbB-GFP fusion and constitutively expresses a cytoplasmic RFP (Moreno-Gúmez et al., 2017). We observed that wild-type cells became competent after 80 min as shown by the expression of SsbB-GFP, and ΔcomC cells started to express SsbB-GFP in the same time frame, independent of whether cells touch each other or not (Figure 5A). This validates our assumption that wild-type cells share CSP in a common pool and can trigger competence in neighboring cells, without the necessity of direct cell contact.

Next, we tested whether the results observed at the population level were reproducible in single-cell-level experiments. First, we established the noise level of false-positive particles in flow cytometry using the ΔcomC strain (which cannot become naturally competent), which turned out to be less than 1% of the cells (Figure S6A). Interestingly, we observed a strong correlation between the detection of the first subpopulation of positive single cells via flow cytometry (2.5% and 4.1% of 12,000 cells per histogram in ATM and control conditions, respectively) and the first value of ≥100 RLU in the plate reader, which was considered a positive signal for competence activation (Figure S6B). Similar results were obtained by fluorescence microscopy, ruling out the presence of an early, pre-existing subpopulation of competent cells below the detection limit of our flow cytometer or plate reader (Figure S6C).

For a better understanding of how competence is initiated and spread at the single-cell level, we studied untreated wild-type cells, ATM-treated wild-type cells, and the ΔlytB mutant at four different inoculum sizes, analyzing 36,000 single particles every 10 min by flow cytometry (three replicates of 12,000 particles) using the SsbB-GFP reporter (Figure 5B). The single-cell flow cytometry data showed that competence development is density dependent in all three conditions, rather than time dependent. For instance, in the wild-type, the onset of competence in cultures with an inoculum size of 10^8 was delayed by more than 2 hr relative to inoculums of 10^2 (green areas, Figure 5B). Note that the SsbB-GFP fusion is much more stable than the luciferase reporter used in plate reader assays, and that GFP-based assays, therefore, do not reflect the narrow window of transcriptional activity that is (more) visible in the corresponding luciferase assays.

As observed in plate reader experiments, the presence of ATM (Figure 5B, red) and the deletion of lytB (Figure 5B, orange) both led to earlier competence development from all inoculation densities, compared to the control condition (Figure 5B, green); however, the synchronization of competent cells in the presence of ATM or absence of lytB was reduced. This was especially obvious at lower inoculation densities, where cells had more time to form chains. Interestingly, the loss of synchronization in the presence of ATM is largely reversed by the exogenous addition of 100 nM synthetic CSP, at the moment the first competent cells were detected (Figure S7A), confirming that there is a large portion of live cells that did not sense enough CSP to develop competence in the absence of exogenous CSP. Indeed, in the presence of ATM or a lytB deletion, synchronization of ≥60% of the population takes nearly twice as long as in control conditions.
Overall, these results show that the initiation of competence is density dependent, with CSP acting as a quorum-sensing agent. However, this sensing can be disrupted or complicated by several factors, such as the presence of long chains retaining CSP, acidification of the medium by fermentation, or other phenomena that affect the diffusion of CSP into the common pool. Furthermore, once competence has initiated at lower cell densities, contact-dependent triggering of competence may play a role (Prudhomme et al., 2016) as exhibited by reduced propagation kinetics (Figure 5B).

Cell Chaining Reduces the Shared CSP Pool
To elucidate whether production and export of CSP are affected in chaining cells, we employed the HiBiT tag detection system (Aggarwal et al., 2018; Wang et al., 2018). The HiBiT tag was placed under the control of the comCDE promoter, either with (strain ADP308) or without (strain ADP312) the leader peptide sequence of comC. As an additional control, we deleted comAB from strain ADP308 (strain ADP311). If the HiBiT peptide carries the leader sequence, it is recognized, cleaved, and secreted by ComAB. Then, extracellularly, it reacts with a HiBiT-dependent luciferase variant (LgBiT), added to the medium, resulting in bioluminescence (Figure 6A, left). In the absence of the comC leader sequence, HiBiT accumulates in the cytoplasm and no luminescence is generated (Figure 6A, right). The extracellular bioluminescence produced by this reporter was similar in the wild-type (ADP308) and the ∆lytB mutant (ADP310) (Figure 6B). We used strains ADP311 (∆comAB) and ADP312 (no comC leader) to confirm that luminescence resulted from active export of the HiBiT tag and was not caused by cell lysis. In both strains, HiBiT cannot be exported and therefore accumulates in the cytoplasm. Indeed, although we detected some lysis after 120 min, the bioluminescence observed is significantly less compared to the strains that export the peptide (Figure 6B). Combined, these results strongly suggest that comC transcription and ComAB activity is not affected by cell chaining and CSP is exported at similar rates in chains of cells.

As the amount of CSP released is similar in wild-type and ∆lytB cultures, we hypothesized that the chain-induced phenotype retains CSP and decreases the amount of CSP released to the shared pool, reducing the synchronization of the population (Figure 6C). Thus, chain formation would reshape global quorum-sensing signaling, where all cells communicate and synchronize competence in a short lapse of time, into local quorum-sensing signaling, where chains retain and sense most of their own produced CSP. To test this hypothesis, we analyzed the ability of wild-type D39V and the ∆lytB strain to induce competence in a coincubated ∆comC strain that harbors the SsbB-GFP fusion. The ∆comC strain is only able to become competent if there is free CSP in the medium, but cannot produce its own CSP. As shown in Figure 6D, competence in the ∆comC strain was detected roughly 40 min earlier when mixed with wild-type cells than with the ∆lytB mutant. This seems in contrast with the fact that the ∆lytB strain is hypercompetent and therefore should release CSP into the medium earlier than the wild-type (in individual populations, the ∆lytB mutant became competent 60 min earlier than the wild-type; Figure 5B). Furthermore, the fraction of activated

conditions, as measured from the first positive time point (Figure S7B). The addition of exogenous CSP, in the presence of ATM nearly compensates for this loss in synchronization.

To test whether addition of CSP, eliminates the differences observed in Figure 5B between wild-type and the ∆lytB strain, we added three different concentrations of CSP, (1, 10, and 100 nM) 60 min into the experiment. This time point is well before the onset of natural competence, so the ComD receptor is not produced at high levels or saturated yet. Indeed, for all three CSP concentrations, competence profiles of wild-type and ∆lytB cells are nearly identical (Figure S7C).
Figure 6. Cells in Chains Have Similar CSP Production Levels but Retain More CSP, Leading to an Extended Transformation Period

(A) Graphical representation of the HiBiT experiment. Left, ComC (called CSP once outside the cell) and HiBiT are regulated by the comCDE promoter, and both precursors have a leader peptide signal, which is recognized, cleaved, and exported by ComAB. Once outside the cell, HiBiT interacts with the soluble protein LgBiT and yields bioluminescence (Wang et al., 2018). Right, HiBiT lacks the ComC leader peptide and accumulates in the cytoplasm, since it cannot be recognized and exported by ComAB.

(B) CSP is exported at a similar rate in wild-type and ΔlytB mutant cells. Bioluminescence (relative luminescence units [RLU]) can be correlated with CSP export. In both the wild-type (ADP308) and the ΔlytB mutant (ADP310), the export rates are similar until the saturation point (1 × 10^7 RLU). Cells were grown in C+Y at competence-permissive pH 7.6. At pH 7.6, cells become naturally competent but with a delay relative to pH 7.9, facilitating the visualization of the inducing effect of chaining. However, competence development occurs later than the RLU saturation point. Neither the comAB mutant (ADP311) nor the HiBiT version without the leader peptide (ADP312) showed any signal during the first 120 min (values below the threshold line of 100 RLU). After that, potentially due to cell lysis, the signal increased but was negligible compared to the exported version of the peptide. Two replicates are shown for each time point and condition.

(C) Graphical representation of the experimental setup. Coincubation (1:1 proportion) of wild-type D39V (black, left panel) or ΔlytB (black, right panel) with ΔcomC mutant (pink) cells. D39V releases more CSP (green dots) into the common pool than the ΔlytB, and more ΔcomC cells become competent (green halo).

(legend continued on next page)
ΔcomC cells incubated with wild-type cells was nearly twice as high as for cells coincubated with ΔlytB (Figure 6D). The initial presence of chains was prevented by bead-beating, and there was no significant difference in either growth rate or survival rate between the ΔlytB and wild-type. Therefore, these results support the conclusion that wild-type D39V releases more CSP into the common pool than the ΔlytB mutant, leading to earlier competence activation in the ΔcomC strain.

Finally, we studied the effect of CSP concentration on the synchronicity of competence development throughout the population. To this end, we added different concentrations of exogenous synthetic CSP1, either at the beginning of the experiment or 90 min after, just before the onset of competence (Figure S7D). Interestingly, the dynamics of competence propagation are similar for different CSP concentrations, with a concentration-dependent delay in the onset of competence. When CSP1 was added after 90 min (roughly three doubling times), this delay in offset was not visible. The dynamics of propagation were similar, with more than 60% of the population becoming competent 20 min after the addition of CSP1. Together, these data show that chained pneumococci have distinctly different kinetics of competence activation and signal propagation from unchained, untreated wild-type diplococci, and do not contribute as much to the extracellular pool of CSP.

Natural Competence in Chained Bacteria Extends the Transformation Window
To investigate the biological relevance of the chain-induced phenotype, we performed transformation experiments, adding external DNA every 20 min in the D39V and ΔlytB strains. As shown in Figure 6E, the chaining phenotype increases the window where bacteria can take up and integrate exogenous DNA, from 100 min (in D39V) to 140 min (in ΔlytB).

DISCUSSION
Many clinically used antibiotics are able to induce competence (Figure 1), which can subsequently lead to the acquisition of antibiotic resistance. Two molecular mechanisms underlying antibiotic-induced competence have been described: altered gene dosage by DNA-targeting antibiotics (Slager et al., 2014), and reduced CSP degradation by HtrA under mistranslation conditions (Stevens et al., 2011). The principal contribution of this work is the identification of a third mechanism, by which certain cell-wall-targeting antibiotics can induce competence. Specifically, the antibiotic ATM, which is used to treat respiratory infections caused by Gram-negative bacteria, and CLA, which is frequently co-administered with the broad-spectrum antibiotic amoxicillin, induce competence (Figure 1B).

Both ATM and CLA target the non-essential PBP3 of *S. pneumoniae* (Figure 3B) (Kocaoglu et al., 2015; Severin et al., 1997), and we show that this causes cell chaining (Figures 3D and S5A). Using CRISPR-mediated deletion of *pbp3* and deletion of the major autolysin *LytB*, we confirmed that competence is upregulated when pneumococci form chains instead of having the normal diplococcal appearance (Figures 3A and S5A).

It is interesting to note that our observations reconcile observations made across different laboratories concerning the dynamics of pneumococcal competence. For instance, at a single-cell level, we confirmed that competence occurs first in a small subpopulation, and then spreads to the whole population, as suggested before (Prudhomme et al., 2016). However, the way in which competence is propagated still remains a cause of debate; some evidence indicates that CSP is, to some extent, retained by producer cells and competence propagates by cell-cell contact (Figure 7; Prudhomme et al., 2016). However, other data showed that CSP is released into a common, shared pool and sensed by the whole population in a typical quorum-sensing manner, which does not require direct contact between cells (Figure 5A; Moreno-Gámez et al., 2017). Here, we show that despite the appearance of a small initial subpopulation of competent cells, in normal conditions (diplococcal phenotype), competence is rapidly spread and synchronized (Figures 4 and 5B). Under cell-chaining conditions or when the medium acidifies after several hours of cultivation, the dynamics of competence propagation seems to depend more on short-range communication between cells (Figure 5B). However, the similar dynamics of population-wide competence development in the presence of various concentrations of exogenous CSP1, even in the presence of ATM, supports the existence and importance of a quorum-sensing mechanism, in addition to a contact-dependent mechanism of competence propagation (Figure S7). The presence of chains could decrease the local or global diffusivity of the CSP in the medium, enhancing local quorum-sensing signaling.

Pneumococcal competence is a population-sensing process that, to a certain extent, is influenced by stochastic parameters, such as basal ComAB and ComCDE expression, replication state, and many more indirect factors. Therefore, single cells produce and sense CSP at different rates, and differences in local CSP concentration will occur. These differences, along with heterogeneity in cells’ CSP-sensing potential, will lead to slight timing differences of competence activation on a single-cell level, thereby leading to the formation of initial subpopulations of competent cells that then activate the rest of the population. Also, competent cells produce cell-wall hydrolases and might reduce growth and kill non-competent siblings.
Interestingly, several factors, such as pH or antibiotics, can modify the rates at which single cells produce and/or sense CSP (Moreno-Gámez et al., 2017; Prudhomme et al., 2016). Our results suggest that chain formation by the presence of ATM or CLA modifies the balance between CSP production and sensing, increasing the self-sensing of CSP between cells within the same chain. Thus, single cells that produce more CSP than average are more likely to share this CSP with cells of the same chain (autocrine-like signaling), reducing the shared pool of CSP (Figure 7, right) (Bareia et al., 2018). We propose to keep using the term quorum sensing (QS) to describe competence activation and signal propagation, as it is clear in the field, as nicely stated by Paul Williams “that the size of the ‘quorum’ is not fixed but depends on the relative rates of production and loss of the signal molecule, which will, in turn, vary depending on the local environmental conditions” (Williams, 2007). In addition, Williams also pointed out that QS can also be considered in the context of “diffusion or compartment sensing,” where the signal molecule supplies information with respect to the local environment and spatial distribution of the cells rather than, or as well as, “global cell population density” (Williams, 2007). This beautifully sums up the observations made here for competence development in S. pneumoniae.

Amoxicillin/CLA (Augmentin) has been available for over 20 years and continues to be one of the most widely used antibiotics, especially in the treatment of respiratory tract infections. However, CLA is a beta-lactamase inhibitor that is useless for the specific treatment of pneumococcal infections, as there have been no reports of S. pneumoniae producing beta-lactamases. Our study suggests that in such cases CLA can best be omitted for antibiotic therapy as it would drive pneumococcal evolution and potentiate antibiotic resistance development by upregulating competence.

Additionally, it has been described that the presence of pneumococcal chains enhances adhesion and colonization (Rodriguez et al., 2012), facilitating the persistence in the nasopharynx in pneumococcal (or polymicrobial) biofilms. This chained phenotype could result in a prolonged time window, during which cells are able to take up exogenous DNA (Figure 6E), explain the rapid adaptation and evolution in response to antibiotic-induced stress in pneumococcal strains colonizing the nasopharynx (Croucher et al., 2011). Thus, it will be interesting to see how competence is synchronized and propagated in more realistic environments, closely resembling the polymicrobial environment that is present in the human nasopharynx. Continued molecular epidemiology studies will be crucial to determine the role and long-term effects of antibiotic therapy.
and vaccination on pneumococcal prevalence and antibiotic resistance.

STAR METHODS

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SUPPLEMENTAL INFORMATION

Supplemental Information includes seven figures and six tables and can be found with this article online at https://doi.org/10.1016/j.celrep.2018.11.007.

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AUTHOR CONTRIBUTIONS

Conceptualization, A.D. and J.-W.V.; Methodology, A.D. and J.S.; Investigation, A.D. and J.S.; Writing – Original Draft, A.D. and J.-W.V.

DECLARATION OF INTERESTS

The authors declare no competing interests.

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**STAR METHODS**

**KEY RESOURCES TABLE**

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Bacterial and Virus Strains** | This paper | N/A |
| Bacterial strains are listed in Table S6 | This paper | N/A |
| **Chemicals, Peptides, and Recombinant Proteins** | | |
| ATM | Sigma-Aldrich | A6848; CAS: 78110-38-0 |
| Potassium clavulanate | Sigma-Aldrich | 33454; CAS: 61177-45-5 |
| Nano-Glo® HiBiT Extracellular Detection System | Promega | N2420 |
| BOCILLIN-FL Penicillin | Thermofisher | B13233 |
| D-Luciferine | Synchem | bc219; CAS: 115144-35-9 |

**CONTACT FOR REAGENT AND RESOURCE SHARING**

Further information and requests for reagents may be directed to, and will be fulfilled by the Lead Contact Jan-Willem Veening (Jan-Willem.Veening@unil.ch).

**EXPERIMENTAL MODEL AND SUBJECT DETAILS**

**Bacterial strains**

All pneumococcal strains used in this study are derivatives of the clinical isolate *S. pneumoniae* D39V (Avery et al., 1944; Slager et al., 2018) unless specified otherwise. See Table S6 for a list of the strains used and the Supplemental information for details on the construction of the strains.

*S. pneumoniae* was grown in C+Y medium at 37°C. C+Y was adapted from Adams and Roe (Adams and Roe 1945) and contained the following compounds: adenosine (68.2 μM), uridine (74.6 μM), L-asparagine (302 μM), L-cysteine (84.6 μM), L-glutamine (137 μM), L-tryptophan (26.8 μM), casein hydrolysate (4.56 g L⁻¹), BSA (729 mg L⁻¹), biotin (2.24 μM), nicotinic acid (4.44 μM), pyridoxine (3.10 μM), calcium pantothenate (4.59 μM), thiamin (1.73 μM), riboflavin (0.678 μM), choline (43.7 μM), CaCl₂ (103 μM), K₂HPO₄ (44.5 mM), MgCl₂ (2.24 mM), FeSO₄ (1.64 μM), CuSO₄ (1.82 μM), ZnSO₄ (1.58 μM), MnCl₂ (1.29 μM), glucose (10.1 mM), sodium pyruvate (2.48 mM), saccharose (861 μM), sodium acetate (22.2 mM) and yeast extract (2.28 g L⁻¹).

We can control competence development by changing the pH in the medium. The underlying mechanism is not fully understood, but it is believed that it is related to the production and export of CSP (Moreno-Gámez et al., 2017). For this reason, we always grow a preculture in C+Y at pH 6.8, because at this pH, even the hypercompetent strains such as ΔlytB or Δpbp3 mutants, are not able to accumulate enough CSP to induce competence before cells reach stationary phase.

**METHOD DETAILS**

**Luminescence assays of competence induction**

To monitor competence development, strains either contain a transcriptional fusion of the firefly *luc* and the *gfp* gene with the late competence gene *ssbB* or a full translational *ssbB-gfp* fusion. Cells were pre-cultured in C+Y (pH 6.8) at 37°C to an OD₅95nm of 0.4. Right before inoculation, cells were collected by centrifugation (8000 rpm for 3 minutes) and resuspended in fresh C+Y at pH 7.3, which is non-permissive for natural spontaneous competence under these experimental conditions. All experiments were
started with an inoculation density of OD$_{595nm}$ 0.004, unless indicated. Luciferase assays were performed in 96-wells plates with a Tecan Infinite 200 PRO illuminometer at 37°C as described before (Slager et al., 2014). Luciferin was added at a concentration of 0.45 mg/mL to monitor competence by means of luciferase activity. Optical density (OD$_{595nm}$) and luminescence (relative luminescence units [RLU]) were measured every 10 minutes. For the CRISPRi experiments, cells were grown as above, and diluted 100x in the presence of a range of IPTG indicated for each condition, depending on whether the targeted gene is essential or not. Despite the fine-tuning regulation of CRISPRi, there is some leakiness that could slightly affect the growth rates and time of natural competence development. For this reason, in these experiments, we do not compare the effect between strains but we compare the control with the addition of IPTG in every strain.

**Detection of the PBPs using Bocillin-FL**

Samples were prepared as described before (Kocaoglu et al., 2015) with slight modifications. Briefly, 4 mL of cells were grown in C+Y pH 6.8 until OD 0.15 and harvested by centrifugation (16,000 × g for 2 min at 4°C). Cell pellets were washed in 1 mL PBS, pH 7.4. Cells were pelleted and resuspended in 50 μL PBS with or without the indicated concentration of ATM or CLA. After 30 min of incubation at room temperature, cells were pelleted, washed in 1 mL PBS, and resuspended in 50 μL PBS containing 5 μg/ml Bocillin-FL. After 10 min of incubation at room temperature, cells were washed again in 1 mL PBS. Next, cells were sonicated on ice (power 30%, three cycles of 10 s interval with a 10 s cooling time on ice (Sonoplus, Bandelin)). Then samples were centrifuged at max speed for 15 min at 4°C and pellets were resuspended in 100 μL cold PBS. The protein concentration was adjusted to 2 mg/ml as determined by Bradford by diluting with PBS. 5x SDS-PAGE loading buffer was added to each sample and heated 10 minutes at 95°C. Proteins were separated by gel electrophoresis (10% acrylamide) for 2.5 h at 180 V, 400 mA, and 60 W. The gel was scanned using a Typhoon gel scanner (Amersham Biosciences, Pittsburgh, PA) with a 526-nm short-pass filter at a 25-μm resolution.

**Intraspecies HGT**

We calculated the *in vitro* HGT efficiency using two genetically identical pneumococcal strains, differing only with the integration of two antibiotic resistance markers at two different locations of the genome. Strains DLA3 and MK134 (tetracycline and kanamycin resistant, respectively), (Slager et al., 2014) were grown to OD$_{595nm}$ 0.4 in C+Y pH 6.8 at 37°C (non-permissive conditions for natural competence activation). Then, a mixed 100-fold dilution of both strains were grown in C+Y pH 7.3 (non-permissive conditions) and pH 7.5 (permissive conditions) to OD$_{595nm}$, to promote the transfer of genes. When cells reached OD$_{595nm}$ 0.4 again (approximately 3 hours), serial dilutions of cultures were plated in Columbia agar + 5% sheep blood with 250 μg/ml of kanamycin plus 1 μg/ml tetracycline for the recovery of the number of recombinants, and without antibiotics to obtain the total viable counts, respectively). Plates were incubated for 16h at 37°C with 5% CO2.

**Interspecies DNA transfer**

*S. pneumoniae* strain D39V was grown to OD$_{595nm}$ 0.4 in C+Y pH 6.8 at 37°C, and *E. coli* carrying the plasmid pLA18 (integrates the tetracycline resistant marker tetM, via double crossover at the non-essential bgaA gene in *S. pneumoniae*, and contains a high copy Gram-negative origin of replication; Slager et al., 2014) was grown overnight with shaking, in LB supplemented with 100 μg/ml of ampicillin (resistant marker also contained in the plasmid, outside the double integration region). Both strains were diluted to OD$_{595nm}$ 0.004 and co-incubated with or without 28 μg/ml of ATM in C+Y pH 7.3. After 3h, serial dilutions were plated either with 1 μg/ml of tetracycline (to recover transformants) or 50 μg/ml of ATM (to recover only the total viable pneumococci). Transformation efficiency was calculated by dividing the number of transformants by the total number of viable count. Three independent replicates of each condition were performed.

**Microarray experiments**

Pneumococcal transcriptome profiles in the presence or absence of antibiotics were tested under conditions that do not support natural competence development to avoid differences in gene expression due to the activation of the competence pathway. We used strain *S. pneumoniae* ADP62 (D39V non-competent variant, comC::ery), grown in two biological replicates in C+Y (pH 7.6). Two kind of experiments were performed to detect rapid and adaptive exposures to the antibiotics. For the fast response, cells were collected during mid-exponential growth phase (OD 0.15) and incubated 15 minutes with or without 2 μg/ml of CLA or 28 μg/ml of ATM. For the adaptive response, cells at OD 0.15 were diluted 100X with or without the same concentration of antibiotics and grown again until OD 0.15. Results were compared using DNA microarray analysis, as previously described. (Shafeeq et al., 2015). For the identification of differentially expressed genes a Bayesian p < 0.001 and a fold change cut-off ≥ 2 was applied. Microarray data are available at Gene Expression Omnibus (GEO) with accession number GSE111562.

**oriC-ter ratio determination by qPCR**

Cells were grown as described above in the presence of antibiotics. In the real-time qPCR experiments, samples were prepared as previously detailed (Slager et al., 2014). Amplification was performed on a QiQ Real-Time PCR Detection System (Bio-Rad). Amplification efficiencies and analysis were performed as before (Slager et al., 2014).
Chain formation detection
To detect morphological changes, we incubated the different strains in C+Y acid medium (pH 6.8) until OD_{595nm} 0.1 and OD_{595nm} 0.4. Antibiotics or IPTG were added when indicated. 1 μl of cells at the indicated optical density was spotted onto a PBS agarose pad on microscope slides, and phase contrast images were acquired with a Leica DMi8 microscope. Microscopy images conversions were done using Fiji and analysis of the length of the chains was done using MicrobeJ (Ducret et al., 2016). Plotting was performed using the BactMAP/spotprocessR package (R. Van Raaphorst, personal communication; https://github.com/veeninglab/spotprocessR).

Fluorescence microscopy
To detect the morphological changes after incubation with antibiotics, 1 μl of cell suspension was spotted onto a PBS agarose pad on microscope slides. Phase contrast images were acquired with a Leica DMi8 microscope with a DFC9000 GT camera and a 100x/1.42 NA phase/c lens. Images were analyzed with ImageJ. For fluorescence microscopy of strains containing SsbB-GFP fusions, cells were spotted onto agarose slides as detailed above, and visualization was performed using a SpectraX light engine (Lumencor) using the following filters for GFP: Quad mirror (Chroma #89000), excitation at 470/24 nm, emission at 515/40 nm. For mKate2 (RFP): Chroma #69008 with excitation at 575/35 nm and emission at 600-670.

Time-lapses videos were recorded by taking images every 10 minutes. The polyacrylamide gel used as semi-solid growth surface was prepared with C+Y (pH 7.9) and 10% acrylamide.

Flow cytometry
ADP245 (P_{ssbB-ssbB-gfp}, bgA::P_{ssbB-luc}) or ADP249 cells (P_{ssbB-ssbB-gfp}) cells were pre-cultured in C+Y (pH 6.8) at 37°C to an OD_{595nm} of 0.1, washed and diluted as explained before in C+Y (pH 7.9). Cells were thoroughly vortexed to avoid possible chains. Experiments were started with an inoculation density of OD_{595nm} 0.0001, with or without 28 μg/ml of ATM. Optical density (OD_{595nm}) was measured every 10 minutes in 96-wells plates with a Tecan Infinite 200 PRO luminometer at 37°C. Right after every measurement, a sample was taken and measured on a Novocyte Flow Cytometer (ACEA Biosciences). The pneumococci were gated to exclude debris. Twelve thousand bacteria were analyzed for FITC fluorescence using a 488 nm laser (GFP expression) with a flow rate of 9 μl/min. Cells pretreated with CSP1 and cells untreated were used to establish the cutoff value for FITC positive (competence activation). Results were analyzed by Novoexpress software (ACEA Biosciences).

Nano-Glo HiBiT Extracellular Detection System
Cells were pre-cultured in C+Y (pH 6.8) at 37°C to an OD_{595nm} of 0.1, washed and diluted as explained before in C+Y (pH 7.6). Experiments were started with an inoculation density of OD_{595nm} 0.001. Optical density (OD_{595nm}) was measured every 10 minutes in 96-wells plates with a Tecan Infinite 200 PRO luminometer at 37°C. Every 20 minutes, 50 μl of the Nano-Glo Extracellular Detection System reagent was added as specified in the manufacturer’s instructions. Additionally, media and PBS samples were used as controls. Bioluminescence was measured every minute during the 10 minutes after reagent addition.

QUANTIFICATION AND STATISTICAL ANALYSIS
Data analysis was performed using GraphPad Prism and Microsoft Excel. A one-tailed Student’s t-test was used to determine differences on chain formation (Figures 3D and S5A), on transformation efficiency (Figure S1C), and on microarray data analysis (Table S5).

Data shown in plots are represented as mean of at least three replicates ± SEM, as stated in the figure legends. Exact number of replicates for each experiment are enclosed in their respective figure legends.

DATA AND SOFTWARE AVAILABILITY
The authors declare that the data supporting the findings of the study are available in this article and its Supporting Information files, or from the corresponding authors upon request.