Identification of Pneumococcal Factors Affecting Pneumococcal Shedding Shows that the \( dlt \) Locus Promotes Inflammation and Transmission

M. Ammar Zafar,\(^a\) Alexandria J. Hammond,\(^b\) Shigeto Hamaguchi,\(^c\) Weisheng Wu,\(^d\) Masamitsu Kono,\(^e\) Lili Zhao,\(^f\) Jeffrey N. Weiser\(^b\)

\(^{a}\)Department of Microbiology and Immunology, Wake Forest School of Medicine, Winston-Salem, North Carolina, USA
\(^{b}\)Department of Microbiology, New York University, New York, New York, USA
\(^{c}\)Division of Infection Control and Prevention, Osaka University Hospital, Osaka, Japan
\(^{d}\)BRCF Bioinformatics Core, University of Michigan, Ann Arbor, Michigan, USA
\(^{e}\)Department of Otolaryngology-Head and Neck Surgery, Wakayama Medical University, Wakayama, Japan
\(^{f}\)Department of Biostatistics, University of Michigan, Ann Arbor, Michigan, USA

**ABSTRACT** Host-to-host transmission is a necessary but poorly understood aspect of microbial pathogenesis. Herein, we screened a genomic library of mutants of the leading respiratory pathogen *Streptococcus pneumoniae* generated by mariner transposon mutagenesis (Tn-Seq) to identify genes contributing to its exit or shedding from the upper respiratory tract (URT), the limiting step in the organism’s transmission in an infant mouse model. Our analysis focused on genes affecting the bacterial surface that directly impact interactions with the host. Among the multiple factors identified was the \( dlt \) locus, which adds \( \alpha \)-alanine onto lipoteichoic acids (LTA) and thereby increases Toll-like receptor 2-mediated inflammation and resistance to antimicrobial peptides. The more robust proinflammatory response in the presence of \( \alpha \)-alanylation promotes secretions that facilitate pneumococcal shedding and allows for transmission. Expression of the \( dlt \) locus is controlled by the CiaRH system, which senses cell wall stress in response to antimicrobial activity, including in response to lysozyme, the most abundant antimicrobial along the URT mucosa. Accordingly, in a \( \text{lysM}^{-/} \) host, there was no longer an effect of the \( dlt \) locus on pneumococcal shedding. Thus, our findings demonstrate how a pathogen senses the URT milieu and then modifies its surface characteristics to take advantage of the host response for transit to another host.

**IMPORTANCE** *Streptococcus pneumoniae* (the pneumococcus) is a common cause of respiratory tract and invasive infection. The overall effectiveness of immunization with the organism’s capsular polysaccharide depends on its ability to block colonization of the upper respiratory tract and thereby prevent host-to-host transmission. Because of the limited coverage of current pneumococcal vaccines, we carried out an unbiased *in vivo* transposon mutagenesis screen to identify pneumococcal factors other than its capsular polysaccharide that affect transmission. One such candidate was expressed by the \( dlt \) locus, previously shown to add \( \alpha \)-alanine onto the pneumococcal lipoteichoic acid present on the bacterial cell surface. This modification protects against host antimicrobials and augments host inflammatory responses. The latter increases secretions and bacterial shedding from the upper respiratory tract to allow for transmission. Thus, this study provides insight into a mechanism employed by the pneumococcus to successfully transit from one host to another.

**KEYWORDS** bacterial transmission, *Streptococcus pneumoniae*, host-pathogen interactions, inflammation, transposon mutagenesis
Successful pathogens require the ability to transit to new hosts. Studying host-to-host transmission within an experimental setting, however, has proven to be problematic because of a lack of tractable, representative animal models and the inherent complexities of the steps involved (exit from the host, transit and survival in the environment, and establishment in a new host). As a result, we have limited biological insight into the contributions of either bacterial or host factors in a pathogen’s dissemination.

Our current understanding of host-to-host transit comes mainly from epidemiological studies that address modes of transmission (1–6). The transmission route utilized by one of the leading bacterial pathogens, *Streptococcus pneumoniae* (the pneumococcus), is via the nasal secretions of colonized individuals and requires close contact (7, 8). Although colonization of the nasopharynx is generally considered asymptomatic (the carrier state), under certain circumstances pneumococci can invade sterile sites within the host, resulting in disease manifestations, including otitis media, pneumonia, sepsis, and meningitis (8, 9). Colonization and transmission are most common among young children, especially in crowded settings, such as day care centers, or in association with episodes of viral upper respiratory tract infection (8, 10, 11).

Each year, approximately a million individuals succumb to infections associated with *S. pneumoniae* (12, 13). For this reason, the World Health Organization has labeled *S. pneumoniae* a priority pathogen, a designation that emphasizes the need for new strategies to combat its spread (14). Due to the challenges of studying pneumococcal transmission, most studies have focused on its disease manifestations (within-host events), where significant advances have broadened our understanding of pneumococcal pathogenesis (7). Recently, we utilized an infant mouse model to study the biology of pneumococcal transmission (the between-host events) (15). Collectively, these studies demonstrate that infection induces a mild acute inflammatory response on the mucosal surfaces of the upper respiratory tract (URT) that promotes nasal secretions and increases the numbers of pneumococci shed from colonized pups (16).

A high level of bacterial shedding is required to overcome the tight population bottleneck observed during intralitter transit from colonized to uncolonized pups (17).

We recently showed that the two major virulence determinants of the pneumococcus also contribute to its transmission among infant mice. Pneumolysin, the organism’s sole toxin that forms pores in host membranes, facilitates transmission by enhancing mucosal inflammation and the frequency of high-shedding events (18). We also observed that certain pneumococcal capsule types are transmitted at higher rates and that this effect is independent of the genetic background (19). This role of capsule type correlates with its effect on shedding and the ability of the organism to escape from entrapment from negatively charged URT mucus.

The purpose of this study was the high-throughput identification of the complete array of pneumococcal factors that affect its transmission using the infant mouse model. We used the technique of mariner transposon mutagenesis (Tn-Seq) to screen a genomic transposon library to identify loci that negatively affect pneumococcal shedding, the limiting step in its transmission (16, 17, 20). Many of the genes identified impact the bacterial surface or host environment. In this report, we focused on one such factor, the *dlt* locus, which modifies teichoic acids through D-alanylation (21, 22).

*RESULTS*

**Identification of pneumococcal genes that affect transmission of the pneumococcus.** To identify genes that potentially impact pneumococcal transmission, we used mariner transposon mutagenesis (Tn-Seq) to generate a library of ~16,000 random mutants in a streptomycin-resistant derivative of strain TIGR4 (T4S) (Fig. 1A). In total, 28 pools, each containing ~500 random mutants, were screened in infant mice (pups).
Each pool (4,000 to 8,000 CFU) was inoculated intranasally (i.n.) into two 4-day-old infant mice. Samples obtained from each of the pups were kept separate and considered biological replicates for each pool. Shedding was used as a proxy for transmission because of the tight population bottleneck between hosts and its close correlation with...

**FIG 1** (A) Schematic representation of the mariner transposon mutagenesis screen (Tn-Seq) carried out in vivo in infant mice (pups). Pups at day 4 of life were infected with 4,000 to 8,000 CFU of a random mutant pool (~500 mutants), and the bacteria shed daily from nasal secretions were collected, quantified, and pooled from days 5 to 9. DNA was isolated from the pooled samples (output I). At 9 days of age, URT lavage fluids in PBS were obtained to quantify the colonization density and DNA isolated (output II). (B) The genetic organization of the genes within the dlt locus of *S. pneumoniae* with the TIGR4 designation and their names are listed. Adjusted P values for shedding for each gene are listed, and the solid red lines represent the gene deletions constructed for the current study. (C and D) Survival assay using *S. pneumoniae* strain TIGR4 Strep-T (T4S), the dltB- and dlt- negative mutants, and the corrected strain (the dltB+ strain) in the presence of various concentrations of nisin or bacitracin. Survival was quantified by incubating serially diluted bacteria in tryptic soy broth with the indicated concentration of the antimicrobial peptide for 3 h at 37°C in 5% CO₂. Each strain was independently tested at least three times, with statistically significant differences being calculated using the Kruskal-Wallis analysis of variance with Dunn’s posttest. *, P ≤ 0.05; **, P ≤ 0.01; ***, P ≤ 0.001.

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the rate of transmission (15, 17–19). Shedding was quantified daily over 5 days (days 5 to 9), colonies were pooled, and DNA was isolated (output I). At day 9 of life, nasal lavage fluid specimens were collected from the URT and placed in phosphate-buffered saline (PBS) to determine the colonization density and DNA isolated (output II). Isolated DNA from the outputs was used as a template for PCR amplification and addition of unique sequence tags to distinguish the different outputs. These samples were processed as described in detail in the Materials and Methods section. Input pools were processed in a similar manner.

Data analysis was carried out by using TnseqDiff, which utilizes a two-step approach in determining the conditional essentiality of a gene (the output compared to the input) (23). To identify factors that affect shedding, genes revealed through the TnseqDiff approach were further filtered to find candidates defective in shedding that did not impact colonization by using an arbitrary cutoff of an adjusted \( P \) value of \(<0.05\) for shedding and an adjusted \( P \) value of \(>0.05\) for colonization. Using these parameters, there were \(>300\) candidates that negatively affected pneumococcal shedding (see Table S1 in the supplemental material). No pneumococcal genes that positively impacted shedding were identified. There were an average of \(>10\) unique transposon insertions per open reading frame.

As the pneumococcal surface is dynamic and interfaces with the host, further analyses focused on candidates that modify its surface or the surrounding host environment. This targeted approach revealed a list of 20 candidate genes (Table 1) whose functions included (i) the biosynthesis of capsular polysaccharide, which was previously implicated in shedding (19), (ii) adherence (pilus attachment, choline binding proteins), and (iii) interactions with glycans in host glycoconjugates and mucus (deglycosylation and mucin binding).

Identification and analysis of the dlt locus. A candidate of particular interest was the gene \(dltB\) (Table 1), which forms part of the \(dlt\) locus (\(dltA, dltB, dltC,\) and \(dltD\) \([dltA-D]\)). The Dlt pathway modifies the anionic glycopolymer teichoic acid (TA), a prominent surface feature of Gram-positive bacteria, by the addition of \(\alpha\)-alanine onto its ribitol or glycerol (in \(Staphylococcus aureus\) backbone (21, 22, 24)). In \(Bacillus subtilis\), inactivation of the Dlt pathway prevents \(\alpha\)-alanylation of both lipoteichoic acid (LTA) and wall teichoic acid (WTA) (25). \(\alpha\)-Alanylation has the effect of decreasing the negative charge of TAs, which could alter interactions with host factors and inflammatory responses. We were intrigued by the potential role of the Dlt pathway in altering the host inflammatory response, as our previous report demonstrated that expression of the pneumococcal cytolysin, pneumolysin, causes an inflammatory response that enhances shedding and transmission (18).

\(DltB\), identified in our shedding screen, is proposed to function as a chaperone in the secretion of the \(\alpha\)-alanine carrier protein (DltC) (26). To confirm that the \(dlt\) locus impacts pneumococcal shedding, we constructed a mutant with an unmarked, in-frame deletion of \(dltB\) (the \(dltB\)-negative \([\Delta dltB]\) mutant) and also a mutant with complete in-frame knockout of the entire \(dlt\) locus (the \(dltA-D\)-negative \([\Delta dltA-D]\) mutant) (Fig. 1B).

Previous studies showed that \(\alpha\)-alanylation of the TA of \(S. pneumoniae\) results in increased resistance to cationic antimicrobial peptides (AMP), in an effect thought to be related to the altered surface charge of \(\alpha\)-alanylation (27). To validate the phenotype of our constructs, we assessed resistance to the cationic AMP nisin, which has previously been shown to be affected by \(dlt\) (27), and confirmed that interruption of the pneumococcal \(dlt\) locus decreased survival in the presence of nisin (Fig. 1C). The \(dlt\) mutants were also more sensitive to the cyclic antimicrobial peptide bacitracin. This suggests a more general effect of Dlt-mediated cell surface modification, since bacitracin acts by a different mechanism: the disruption of peptidoglycan biosynthesis (28) (Fig. 1D). In the presence of both of these antimicrobials, the survival defect was specific to the \(dlt\) locus, as correction of the genotype (producing the \(dltB\)-positive \([dltB^+]\) mutant) restored the wild-type (WT) phenotype.
TABLE 1  Surface-exposed or surface-acting factors in pneumococcal sheddinga

| TIGR4 designation | P value for shedding | Gene name and description | Function (reference) |
|-------------------|----------------------|---------------------------|----------------------|
| SP_0112           | 0.019949             | artP; amino acid ABC transporter | Arginine ABC transporter (64) |
| SP_0148           | 0.000324             | gshT; ABC transporter       | Predicted glutathione ABC transporter (65) |
| SP_0198           | 0.030898             | Hypothetical protein        | Lipoprotein, putative d-stereospecific aminopeptidase (44, 45) |
| SP_0314           | 0.013614             | hysA; halorutonate lyase    | Surface enzyme that cleaves halorutonate, a component of the host extracellular matrix (66, 67) |
| SP_0352           | 0.000093             | cps4G; glycosyltransferase  | Biosynthesis of capsular polysaccharide (68) |
| SP_0368           | 0.000004             | eng; endo-alpha,N-acetylgalactosaminidase | Secreted O-glycosidase that modifies host O-linked glycans (69) |
| SP_0391           | 0.001690             | cbpF; choline binding protein F | Modulates the autolytic function of LytC (70) |
| SP_0467           | 0.036190             | srtC; sortase C             | Involved in covalent attachment of the pilus to the peptidoglycan cell wall (71, 72) |
| SP_0614           | 0.016540             | estA; esterase              | Enhances neuraminidase activity by removing acetylation from sialic acid (73) |
| SP_0667           | 0.000046             | cbpl; choline binding protein L | Contributes to resistance against phagocytosis and is involved in adhesion (74) |
| SP_0771           | 0.021157             | slrA (pplA); peptidyl-prolyl cis-trans-isomerase, cyclophilin type | Catalyzes isomerization of proline containing tetrapeptides (75) |
| SP_0899           | 0.037124             | Conserved hypothetical protein | Surface-exposed lipoprotein of unknown function (76) |
| SP_0965           | 0.023346             | lytB; endo-beta-N-acetylgalactosaminidase | Nonautolytic peptidoglycan hydrolyase, acts as a glucosaminidase and is involved in cell division cycle (77) |
| SP_1000           | 0.016753             | etrx2; thioredoxin family protein | Surface-exposed lipoprotein that provides oxidative stress resistance (78) |
| SP_1004           | 0.000050             | phtE; histidine triad protein | Surface-exposed protein that promotes adhesion to host cell surfaces (79) |
| SP_1154           | 0.014494             | zmpA; zinc metalloproteinase | Cleaves human immunoglobulin A (IgA1) (80, 81) |
| SP_1492           | 0.035625             | mucBP; cell wall surface anchor family protein | Binds mucin, has a potential role in pneumococcal adherence (82, 83) |
| SP_1573           | 0.000314             | lytC; lysozyme              | Surface-bound lysozyme that acts as a pneumococcal cell wall hydrolase (84) |
| SP_1796           | 0.045895             | fusA (suxS); carbohydrate ABC transporter, substrate binding protein | Required for utilization of fructose oligosaccharide, uses inulin as a carbon source (85–87) |
| SP_1872           | 0.034338             | piuA; iron compound ABC transporter, iron compound-binding protein | Lipoprotein component of iron ABC transport Piu system (88, 89) |
| SP_1963           | 0.036813             | Putative hemolysin          | CBS domain-containing protein with homology to putative cytolysin proteins (64) |
| SP_1964           | 0.033659             | endA; DNA entry nuclease    | Surface endonuclease that degrades the DNA scaffold of neutrophil extracellular traps (NETs) (90) |
| SP_2084           | 0.041972             | pstS; phosphate ABC transporter, phosphate-binding protein | Subunit of the phosphate ABC transporter, implicated in penicillin resistance (91, 92) |
| SP_2099           | 0.003095             | ppplB; bifunctional penicillin-binding protein | Synthesis of peptidoglycan (transglycosylation and transpeptidation) (93) |
| SP_2175           | 0.014521             | dltB; membrane protein involved in D-alanine export | Addition of D-alanine to teichoic acids (27) |

aStreptococcus pneumoniae genes identified by mariner transposon mutagenesis (Tn-Seq) screening to be involved in pneumococcal shedding and predicted to be either surface-exposed or surface-acting factors.

Contribution of the dlt locus to pneumococcal shedding and transmission.

Because of day-to-day variation, shedding was measured daily (days 1 to 5 postinoculation) and values were pooled for comparison (15). Both the dltB- and dlt-negative mutants showed reduced median shedding compared to either the corrected mutant (the dltB+ mutant) or the WT type 4 strain (Fig. 2A). These results included inpairlier comparisons of strains from pups from multiple litters to control for host and environmental effects on shedding. The contribution of the dlt locus to shedding was not due to differences in colonization density, as all the strains colonized equally well at the conclusion of the shedding experiment at age 9 days (Fig. 2B).

Next, we determined whether the attenuated shedding seen in dlt mutants leads to reduced transmission. Using a 1:1 ratio of pups colonized by inoculation (index pups) to uninoculated littermates (contact pups), no transmission events were detected with either the dltB- or dlt-negative strains that showed diminished shedding (Fig. 2C). Importantly, transmission to levels previously reported for the WT strain was restored by correction of the deletion in dltB (producing the dltB+ strain) (15, 18). Again, all
constructs colonized pups at a high density at the conclusion of the transmission experiment at age 14 days (Fig. 2D). Thus, we validated the initial observations from the Tn-Seq screen and confirmed that the dlt locus does not impact colonization but is needed for sufficient pneumococcal shedding to allow for host-to-host transmission.

Because expression of the pneumolysin toxin (Ply) also increases shedding and transmission (18), we examined whether the dlt locus had an additive effect with Ply. The dlt- andply-negative double mutant showed low median shedding that was indistinguishable from that of the dlt-negative single mutant (Fig. 2A). The major effect of the loss of ply, however, was fewer high-shedding events (>300 CFU) rather than altered median shedding (18). The proportion of high-shedding events was significantly lower for the dlt- and ply-negative double mutant than for the dlt-negative single

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**Table:**

| Strain | Total Contacts | Colonized Contacts | Transmission (%) | p-value |
|--------|----------------|--------------------|-----------------|---------|
| WT*    | 14             | 4                  | 29              |         |
| dltB-  | 16             | 0                  | -               | 0.0242  |
| dlt-   | 15             | 0                  | -               | 0.0282  |
| dltB+  | 12             | 4                  | 33              |         |

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**Fig 2** The dlt locus affects pneumococcal shedding and transmission without impacting colonization. (A) Pups were challenged i.n. at 4 days of age with the indicated construct, and bacteria shed daily from nasal secretions were collected and quantified from days 5 to 9. Median values are indicated, and each symbol represents the number of CFU measured from a single pup. The dotted horizontal line represents the 300-CFU threshold described in Results. The dashed vertical line separates the intralitter comparisons of two pneumococcal strains. Data are for 6 to 10 pups per group. (B) Colonization density of each pneumococcal construct in cultures of URT lavage fluids obtained from pups at 9 days of age. Median values are shown. (C) Summary of the rate of transmission of dlt locus mutants and the corrected strain (the dltB+/H11001 mutant) from colonized index pups at the age of 4 days to naive contact pups in the same litter. The transmission rate was determined by the number of contact pups (at a 1:1 ratio to index mice) colonized by S. pneumoniae at the age of 14 days. Data for the WT are historic data (15). dlt locus mutants were compared to the dltB+/H11001 group by using Fisher’s exact test. (D) Colonization density of each pneumococcal dlt construct tested for transmission (index mice) in cultures of URT lavage fluids obtained from pups at 14 day of age. *, P ≤ 0.05; **, P ≤ 0.01; ****, P ≤ 0.0001, n.s, not significant.
mutant, suggesting that the toxin and α-alanylation may act through different pathways to induce shedding.

**Role of α-alanylation in URT inflammation.** *In vitro* studies have shown that LTA, once it is released after bacteriolysis, triggers an inflammatory response via recognition by the pathogen recognition receptor Toll-like-receptor 2 (TLR2) (29–31) and that α-alanylation of pneumococcal LTA augments TLR2-mediated inflammation (31, 32). Therefore, we postulated that disruption of the dlt locus might impact shedding by dampening the proinflammatory responses of the URT during colonization. As predicted, at age 7 days (2 days postchallenge), the numbers of neutrophils (CD45+Ly6G+ and CD11b+ events) detected by flow cytometry of the nasal lavage fluid specimens from infant mice infected with the dltB-negative mutant were reduced compared to the numbers for infant mice infected with the WT strain (Fig. 3A). Additionally, there was decreased URT transcription of the chemokine CXCL-2, which stimulates neutrophil recruitment, and the proinflammatory cytokine interleukin-1β (IL-1β) in the lavage fluid specimens obtained from infant mice colonized with the dltB-negative mutant (Fig. 3B). The observed differences in the inflammatory response were not due to differences in colonization density, as both strains colonized to a high density at this earlier time point (Fig. 3C). Thus, our *in vivo* data correlated with the findings of the *in vitro* studies linking α-alanylation to increased inflammation and provided further evidence correlating inflammation and shedding.

**The effect of α-alanylation of LTA on shedding requires signaling through TLR2.** Next, we examined if the increased inflammation observed with strains expressing α-alanylated TAs was due to increased TLR2-mediated signaling, as suggested by the *in vitro* studies.

We previously reported that tlr2−/− pups show decreased URT inflammation and reduced shedding (18). The difference in median shedding between mice infected with the dlt-negative mutant and mice infected with its parent strain, as noted in WT mice, was no longer observed in tlr2−/− infants, and both strains colonized the knockout mice at equivalent densities (Fig. 4A and B). Moreover, in tlr2−/− mice infected with the dlt-negative mutant or its parent strain, there was no difference in neutrophil numbers or the transcript levels of chemokines/cytokines upregulated by the presence of dlt (Fig. 4C and D). Together, our *in vivo* data suggest that α-alanylation of LTA increases inflammation in a TLR2-dependent manner and that this increased host response drives pneumococcal shedding.

**Lysozyme treatment leads to upregulation of the dlt locus.** Lysozyme, an antimicrobial enzyme that functions by degrading the bacterial peptidoglycan backbone, is one of the most abundant antimicrobial proteins present on the mucosal surface of the URT (33). To evade lysis by the high concentration of lysozyme in its URT niche, *S. pneumoniae*, like other mucosal pathogens, must modify its cell wall (34). The

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**FIG 3** The absence of α-alanylation (dltB negative) leads to reduced URT inflammation. (A) Number of neutrophils determined by flow cytometry (CD45+, CD11b+, and Ly6G+ events) in URT lavage fluids obtained from WT (T4S)-inoculated or isogenic dltB-negative construct-inoculated pups at age 7 days. Values are ±SEM (n = 8 to 11). (B) Gene expression in pups colonized with the dltB-negative mutant relative to that in WT strain-inoculated pups at age 7 days, as measured by qRT-PCR, for the chemokine/cytokine shown. Values are ±SEM (n > 10). (C) Colonization density in URT lavage fluids obtained at age 7 days, with the median values being shown. *P ≤ 0.05; n.s, not significant.
The pneumococcal dlt locus has been reported to confer resistance against lysozyme in diverse bacterial species, such as *Bacillus subtilis*, *Clostridium difficile*, and *Enterococcus faecalis* (26, 35–37). As the Dlt pathway modifies TA, we hypothesized that it might also contribute to the resistance of *S. pneumoniae* against lysozyme-mediated killing, as observed for bacitracin, another antimicrobial that targets peptidoglycan (Fig. 1D). In the presence of physiologic concentrations of lysozyme in the URT (33), the WT strain was resistant to its lytic effects, but survival was reduced when the dlt locus was disrupted (Fig. 5A). Restoration of the intact dlt locus (producing the dltB− strain) led to WT levels of survival.

It has been suggested that in response to AMPs the CiaRH two-component regulatory system, present in different streptococcal species, upregulates expression of the dlt locus (38). Therefore, we examined whether the cell wall stress caused by lysozyme would also lead to Dlt pathway upregulation. As shown in Fig. 5B, we observed a 2-fold induction of the dlt locus, as evidenced by an increase in dltA transcript levels, that was dependent on CiaRH, since for a ciaRH-negative mutant treated with lysozyme, there was no upregulation of dltA expression. Our finding suggests that CiaRH-mediated sensing of cell wall perturbation by lysozyme upregulates the Dlt pathway. We then determined whether the contribution of the dlt locus required lysozyme by the use of lysM−/− mice, which lack the enzyme in URT secretions. In the absence of the cell wall stress from lysozyme, when our study predicts that expression of the dlt locus would not be upregulated, shedding was lower than that in WT mice and there was no longer an effect of the dlt locus (Fig. 5C). Deletion of the dlt locus or the lack of lysozyme had no effect on the colonization density (Fig. 5D).

**DISCUSSION**

The long-term goal of this project is to provide a thorough understanding of the biology of pneumococcal transmission. Widespread immunization of children with the pneumococcal conjugate vaccine (PCV) since 2000 has led to a dramatic reduction in
invasive pneumococcal disease, including for unimmunized groups, such as adults and the elderly (39, 40). This indirect protection, or herd immunity, which accounts for most of the overall efficacy of PCV, has been attributed to reduced carriage and transmission within the community (41, 42). PCV, however, is based on capsular polysaccharide and targets only a limited number of the >95 known serotypes (43). A more comprehensive understanding of pneumococcal surface factors contributing to its transmission, therefore, could yield novel targets that could broaden and make more effective prevention strategies through immunization.

In the current study, we used a genomic approach to identify pneumococcal genes affecting the exit of the pneumococcus from the colonized host (i.e., shedding) through nasal secretions using an infant mouse model. The function and the role of a large proportion of pneumococcal genes in transmission remain to be determined. Shedding was used as a proxy for transmission because of the tight population bottleneck during transmission (17). It was necessary to focus on shedding, an earlier event that is the limiting step in host-to-host transit (15, 17–19). We validated this approach by confirming that pneumococci carrying one of the hits from the Tn-Seq screen, the dlt locus, that showed reduced shedding were also attenuated in transmission among infant mice. Besides confirming the effects of the dlt locus, to date we have validated the effects of two other hits listed in Table 1, SP_1963 (annotated as a putative hemolysin) and SP_1964 (endonuclease; endA). Pneumococci carrying both were confirmed to have a defect in shedding, but they did not impact colonization density. Prior studies using in vitro or in vivo or screens of random pneumococcal transposon mutants have been used to identify essential genes or within-host factors (virulence factors), whereas our in vivo screen is the first to examine factors involved in between-host events (44–47).

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**FIG 5** α-Alanylation of pneumococcal lipoteichoic acids provides protection against host lysozyme. (A) Lysozyme survival assay using S. pneumoniae strain TIGR4 (T4S), the dltB- and dlt-negative mutants, and the corrected strain (the dltB+ strain). Survival was quantified by incubating serially diluted bacteria in tryptic soy broth with 1.0 mg/ml of chicken egg lysozyme for 3 h at 37°C in 5% CO2. Each strain was independently tested at least three times. (B) Gene expression of dltA for pneumococcal strains treated with or without lysozyme for 60 min in tryptic soy broth at 37°C, as measured by qRT-PCR. (C) Congenic lysM−/− pups were colonized at age 4 days with the WT strain (T4S) or with its isogenic dlt-negative construct using intralitter comparisons. Daily shedding in nasal secretions was quantified from ages 5 to 9 days, with median values being indicated and with each symbol representing the number of CFU observed from a single pup on a single day. Data are for 4 to 5 pups per group. (D) Colonization density in URT lavage fluids obtained at age 9 days, with the median value being shown. * P ≤ 0.05; ** P ≤ 0.01; n.s, not significant.
There are several important caveats to the approach used in our genomic screen. First, we identified over 300 genes that impact pneumococcal shedding, suggesting that ~15% of all nonessential TIGR4 genes significantly affect its shedding (output I) and not its colonization (output II). One reason that our screen identified such a high number of genes affecting shedding could be the less stringent cutoff ($P < 0.05$) used because shedding is an inherently variable readout. Additionally, our screen was carried out with a single highly annotated pneumococcal strain (TIGR4), although pneumococcal isolates show marked differences in genome content and transmissibility (19, 48–50). Moreover, mice are not a natural host for the pneumococcus, even though the infant mouse model recapitulates many of the key features of human contagion (need for close contact, increased rate during viral coinfection and among infants). It is also likely that random transposon insertions might only partially abrogate the function of a gene or lead to polar effects, and TnseqDiff accounts only for inserts that occur within coding regions. Finally, trans-acting factors, such as the effect of ply on inflammation, could be obscured when screening large pools of mutants. This could explain why the effect of the loss of dlt appeared to be greater for the constructs (the dltB- and dlt-negative constructs) tested individually than for the transposon mutant pools.

Taking into account the dynamic nature of the pneumococcal surface and its interactions with the host, we focused on factors that attenuate shedding, thereby reducing transmission, by modifying its surface. This led us to further characterize the dlt locus, since pneumococcal LTA triggers a strong TLR2-dependent response that is partially dependent upon its $\alpha$-alanination via the Dlt pathway (29–32). Our findings confirmed that mutants disrupting the Dlt pathway showed reduced shedding, a defect that correlated with reduced levels of acute inflammation, with sensing by TLR2 mediating the inflammatory signal. URT inflammation increases the production of mucins, including the major secretory mucin Muc5ac, and the volume of nasal secretions that drive pneumococcal shedding (16). The mechanism of how $\alpha$-alaninated LTA contributes to recognition through host pattern recognition receptors remains to be determined. In this regard, there remains controversy as to the contribution of TLR2-mediated sensing of LTA. Work by Kang et al. showed that LTA can bind to the TLR2 extracellular domain but not cause heterodimerization (51). This work, however, used only the soluble domains, and the authors suggest that their findings do not exclude the possibility that LTA may induce the dimerization of full-length TLR2 with TLR1 or TLR6 under membrane-attached conditions. Furthermore, Gisch et al. showed that with lipid-free LTA (through hydrogen peroxide treatment or the use of a mutant defective in lipidation of the prelipoproteins [a Δlgt mutant]), IL-8 production in vitro was abolished, suggesting that the lipid moiety is important for signaling (52). This report also suggested that lipopeptide-free LTA still had an immunostimulatory effect in human mononuclear cells, supporting the suggestion that the lipid moiety in LTA may still be important for sensing, possibly via TLR2 recognition, an observation consistent with our findings. On the other hand, in Staphylococcus aureus, $\alpha$-alanine modification of LTA, which is structurally distinct from that of S. pneumoniae, did not similarly affect inflammatory responses (53). It is also possible that the dlt locus affects the release of lipoproteins in response to antimicrobial activity, with subsequent sensing via TLR2.

Our previous study demonstrated the importance of TLR2-mediated sensing and the pneumolysin toxin in inflammation, shedding, and transmission but did not establish a link between these pathways. The dlt-dependent effects observed in the current study are epistatic to TLR2 sensing but independent of pneumolysin (Fig. 2A). These results suggest that there is more than a single type of inflammatory response to pneumococcal colonization that affects bacterial shedding and transmission. Furthermore, ply was not identified in the screen because transposon mutations in this gene affected both shedding and colonization. Since we previously showed that mutants with a nonpolar ply in-frame clean deletion have a shedding defect that does not impact colonization, it is possible that the transposon insertions in ply have polar effects. Consistent with our previous findings that link shedding levels with the transmission
rate, we observed that reduced shedding in the absence of a functional dlt locus results in a complete loss of transmission.

Another effect of Dlt-mediated ω-alanylation of LTA observed in our study was increased resistance to the antimicrobial actions of lysozyme. This result adds to the list of pneumococcal enzymes, including pgdA and adr, that modify the pneumococcus cell surface to evade lysis by this highly abundant host factor found in URT secretions (34, 54–56). Our findings obtained with bacitracin, which hinders peptidoglycan biosynthesis, and lysozyme, which cleaves the peptidoglycan backbone, show that the reduced surface charge by ω-alanylation acts in a manner that aids in protecting the pneumococcus against both cationic AMPs and other agents that perturb the integrity of the cell surface. A recent report on group B streptococcus suggests that ω-alanylation increases the cell wall density by changing the conformation of the LTA in a manner that can limit penetration by AMPs and, thus, increase resistance and survival (57). Furthermore, during treatment with physiological levels of lysozyme, there is a CiaRH-dependent upregulation in expression of the Dlt pathway, suggesting a programmed response to cell wall stress. It appears that when targeted by antimicrobial peptides/proteins, such as lysozyme, pneumococci increase ω-alanylation of LTA, which enhances TLR2-mediated inflammation. This increased URT inflammation in turn facilitates increased shedding to allow for exit of the organism to establish itself in a new, more hospitable host. Thus, our study demonstrates that the pneumococcus is a highly adaptable pathogen capable of modifying its cell surface in response to the host environment to promote both its survival and its dissemination.

MATERIALS AND METHODS

Ethics statement. This study was conducted according to the guidelines outlined by National Science Foundation animal welfare requirements and the Public Health Service Policy on Humane Care and Use of Laboratory Animals (58). The New York University Medical Center IACUC oversees the welfare, well-being, and proper care and use of all vertebrate animals.

Growth conditions and strain construction. Pneumococcal strains were grown statically in tryptic soy (TS) broth (Becton, Dickinson (BD)) at 37°C. Upon reaching the desired optical density at 620 nm (OD620), cells were washed and diluted in sterile phosphate-buffered saline (PBS) for inoculation. For quantitative culture, serial dilutions were plated on TS agar-streptomycin (200 μg/ml) supplemented with either 5% sheep blood or catalase (6,300U/plate; Worthington Biochemical Corporation) and incubated overnight at 37°C with 5% CO2. A streptomycin-resistant derivative of the type 4 strain TIGR4 (58) was transformed with a PCR product using primers 5-amino-acid coding sequences of the dltB gene. The transformants were selected on TS agar-kanamycin (125 μg/ml) supplemented with either 5% sheep blood or catalase (6,300 U/plate; Worthington Biochemical Corporation) and incubated overnight at 37°C with 5% CO2. A streptomycin-resistant derivative of the type 4 strain TIGR4 (T4S), P2406, was used throughout the study (15).

The dltB knockout strain was constructed in a two-step process using the Janus cassette (59). In the first step, the Janus cassette was amplified from genomic DNA, obtained using a MasterPure DNA purification kit (Epicentre/Lucigen), from strain P2408 (60), with flanking regions (~1 kb) upstream and downstream of the dltB gene added via isothermal assembly. Strain P2406 was then transformed with the PCR product, and the transformants were selected on TS agar-kanamycin (125 μg/ml) plates. A PCR-confirmed mutant, P2552 (dltB::Janus KanR Streps), was selected to make an in-frame deletion. The mutant was transformed with a PCR product using primers F1-dltB/R7-dltB and F8-dltB/R6-dltB and genomic DNA obtained from P2406 and selected on TS agar-streptomycin (200 μg/ml) to generate strain P2555 (ΔdltB Streps' Kan'). The in-frame knockout strain (the ΔdltB strain) has a scar containing the first and the last 5-amino-acid coding sequences of the dltB gene.

The corrected mutant, P2556 (the dltB’ mutant), was constructed by transforming strain P2552 with a PCR product generated using primers F1-dltB and R6-dltB to amplify genomic DNA from strain P2406, and the transformants were selected on TS agar-streptomycin (200 μg/ml). The dlt locus knockout mutant was constructed as described above, with the primers used being listed in Table S2 in the supplemental material. Neither the dltB- nor dlt-negative mutant constructs showed a growth defect in nutrient medium.

The ciaRH knockout strain P2597 (ciaRH::erm) was constructed by transforming genomic DNA obtained from an isolate with a ciaRH::erm cassette into P2406 (15). The resulting transformants were selected on TS agar plates supplemented with erythromycin (2 μg/ml) and 5% blood. A second back-transformation was carried out in P2406 with genomic DNA obtained from one of the transformants, and isolates were selected on plates as described above to generate strain P2597 (ciaRH::erm Ermm Streps').

Shedding and colonization in infant mice. WT, thr2-Δ and lysM-Δ CS78BL/6J mice were obtained from The Jackson Laboratory (Bar Harbor, ME) and bred and maintained in our conventional animal facility at the NYU Langone Medical Center. The pups were housed with their dam (mother) for the duration of the experiment and gained weight like uninfected animals. At day 4 of life, the pups were given an intranasal (i.n.) inoculation, without anesthesia, containing ~2,000 CFU of S. pneumoniae suspended in 3 μl of PBS, as described previously (15, 16). Shedding was quantified by gently tapping the nares (20 taps/pup) on a TS agar plate supplemented with streptomycin.
(20 μg/ml) or kanamycin (125 μg/ml) to prevent the growth of contaminants, followed by spreading of the secretions over the agar surface with a sterile cotton-tipped swab. Because of the day-to-day variability in the numbers of shed bacteria, values were obtained daily for 5 days and pooled. To control for environment effects, shedding of different strains was compared for pups within the same litter. To measure colonization density, we euthanized the pups at the ages indicated below by CO₂ asphyxiation, followed by cardiac puncture. The upper respiratory tract (URT) was lavaged with 200 μl of sterile PBS from a needle inserted into the trachea, and fluid was collected from the nares. The limit of detection in lavage fluids was 33 CFU/ml.

Transmission in infant mice. The pneumococcal mono-infection transmission model was described in previous studies (15). Briefly, half of the pups in a litter were randomly selected and, at day 4 of life, infected with the S. pneumoniae strain indicated above. These index mice were then returned to the dam and the other uninfected pups (contact mice) and cohoused for 10 days postinfection. To detect bacterial transmission from the index pups to the contact pups, all pups were euthanized at the age of 14 days, and nasal lavage fluid specimens were cultured.

Transposon library preparation and in vivo selection. Library construction using the mariner transposon was carried out as previously described (20) with slight modifications. Briefly, 28 independent libraries were generated, with each library containing ~500 mutants with transposon insertion mutations in nonessential genes. On a given day, a transposon pool was thawed and grown at 37°C, and two 4-day-old pups (biological replicates) were inoculated i.n. with 4,000 to 8,000 CFU of transposon mutants, such that each pup received an ~10-fold excess of the random mutants (input). Pilot experiments with various inocula confirmed an infectious dose of <10 CFU and the lack of a population bottleneck (the systematic loss of clones) when pups were colonized as described above (data not shown).

After infection, pneumococcal shedding over 5 days was carried out as described above. After shed pneumococci were quantified, colonies were removed with a swab, transferred to 2 ml of TS broth, and incubated at 37°C for an hour, before adding glycerol at a final concentration 20% for storage at −80°C until further processing. Sheding for the two biological replicates was kept separate and was carried out over 5 days. Afterwards, shed bacteria collected over all 5 days (>3,000 total CFU) were pooled and pelleted by centrifugation at 4,300 × g for 10 min. Genomic DNA was isolated as described above, and samples were sequenced as described previously (20) (output II). A cleanup step was added to the ligation mixture to remove excess adaptor using AMPure XP beads (Beckman Coulter), before using PCR to amplify the transposon insertion site.

At day 9 of life, pups infected with S. pneumoniae transposon library mutants were sacrificed and nasal lavage specimens were collected in a 200-μl volume as described above. Ten microliters was used to determine the density, and the remainder of the lavae fluid was plated on TS agar-streptomycin (200 μg/ml) supplemented with catalase and incubated overnight at 37°C. On the next day, the colonies were collected by swabbing and transferred to 5 ml sterile PBS (output II). Genomic DNA was isolated, and samples were prepared as described above.

Sequencing and bioinformatics analysis. Transposon junction DNA fragments were subjected to single-end 50 rapid-run sequencing on an Illumina HiSeq 2500 instrument (Illumina, San Diego, CA). The resulting reads were filtered, mapped, and normalized as described previously (61). Briefly, raw sequencing reads were searched for the transposon sequence ACAGGTTG, and the reads without this sequence were discarded. The FASTX-Toolkit (http://hannonlab.cshl.edu/fastx_toolkit/) was used to debarcode the reads based on the barcode sequences at the 5’ ends, and the reads that became shorter than 12 bp after barcode trimming were discarded. The debarcoded reads were then processed using an IN-Seq pipeline as published previously (61). Specifically, reads were aligned to the S. pneumoniae TIGR4 reference genome using the Bowtie program (62), and insertion sites were called for each sample based on the alignment counts. Only uniquely mapped reads with no more than one mismatch were included. Insertion sites within 80% of the gene length from the transcription start sites were treated as candidates that could affect gene functions. Finally, TnseqDiff (23) was used to compare the insertion sites between output and input samples to identify genes potentially essential under different conditions. Comparison was carried out for the following groups: input versus shedding and input versus nasal lavage.

Antimicrobial sensitivity assay. Stocks of nisin (catalog number M5764-1G; Sigma) were prepared in 0.05% acetic acid (10 mg/ml), stocks of bacitracin (catalog number B5150-250KU; Sigma) were prepared in 1 M HCl (50 mg/ml), and stocks of hen egg white lysozyme (Roche) were prepared in distilled water (20 mg/ml). To determine susceptibility to these antimicrobial peptides, strains were grown in TS broth to mid-log phase and diluted 200-fold in a microtiter plate containing a 2-fold concentration gradient of either nisin or bacitracin, with the final volume being 200 μl. The plate was incubated at 37°C in 5% CO₂ for 3 h, after which serial dilutions were plated on TS agar plates and incubated at 37°C in 5% CO₂ overnight. Percent survival was calculated by counting the number of CFU at various concentrations of the antimicrobial peptide versus that for the vehicle control. Each experiment was repeated at least 4 times.

To test sensitivity to lysozyme, strains were grown as mentioned above and diluted 200-fold in a microtiter plate containing either 500 or 1,000 μg/ml lysozyme, with the final volume being 200 μl. Incubation, serial dilutions, and calculation of percent survival were described above.

Flow cytometry. Neutrophils present in the nasopharynx were quantitated as previously described (18). Briefly, nasal lavage samples from individual pups were pelleted by centrifugation at 500 × g for 5 min. Samples were resuspended in 50 μl ACK lysis buffer (Thermo Fisher Scientific) and incubated for 5 min at room temperature. Afterwards, 200 μl of PBS was added to the samples, and cells were pelleted as described above and resuspended in PBS containing 1% bovine serum albumin (BSA). Samples were stained with a LIVE/DEAD Fixable Aqua dead cell stain kit (Invitrogen, Thermo Fisher Scientific). Next,
samples were blocked with a 1:200 dilution of a rat anti-mouse CD16/32 (clone 93; BioLegend). Cells were stained for 30 min at 4°C with fluorophore-conjugated antibodies (diluted 1:150) against the following surface markers: CD11b-V450 (BD), Ly6G-peridinin chlorophyll protein (PerCP)-Cy5.5 (BD), and CD45-allophycocyanin (APC)-Cy7 (BD). Samples were run on a BD LSR II flow cytometer and analyzed with BD FACS Diva software.

qRT-PCR. The upper respiratory tract of the pups was lavaged using RLT lysis buffer (Qiagen) to obtain RNA from the epithelium. Total RNA and cDNA generation was carried out as previously described (60). Quantitative reverse transcription-PCRs (qRT-PCRs) were performed using Power SYBR green master mix (Applied Biosystems), ~10 ng cDNA, and 0.5 mM primers per reaction mixture. Samples were run in duplicate, and each experiment run was repeated. Samples were run on a CFX384 Touch real-time PCR detection system (Bio-Rad). Primers directed toward the GAPDH (glyceraldehyde-3-phosphate dehydrogenase) gene were used as an internal control. RNA expression was quantified using the ΔΔ

d fold method. The primers used in this study were previously described (60, 63).

To determine the ΔΔCT transcript level upon treatment with lysozyme, the bacterial strains were grown in TS broth at 37°C until they reached mid-log phase (OD600 = 0.5), back-diluted 4-fold in fresh TS broth containing either 1 mg/ml lysozyme or water (mock treatment), and grown statically again at 37°C for 1 h. Afterwards, the cultures were spun down at 4°C at 4,300 x g for 10 min. Samples were treated with RNAprotect Bacteria Reagent (Qiagen), and RNA was isolated using an AllPrep bacterial DNA/RNA/protein kit (Qiagen) following the manufacturer’s protocol. cDNA and qRT-PCR were carried out as described above using a primer set against dltA and a primer set against DNA gyrase (gyr) as an internal control.

Statistical analysis. All statistical analyses were performed using GraphPad Prism (version 7.0) software (GraphPad Software, Inc., San Diego, CA). Unless otherwise specified, differences were determined using the Mann-Whitney U test (comparing two groups) or the Kruskal-Wallis test with Dunn’s postanalysis (comparing multiple groups).

SUPPLEMENTAL MATERIAL
Supplemental material for this article may be found at https://doi.org/10.1128/mBio.01032-19.

TABLE S1, XLSX file, 0.04 MB.
TABLE S2, DOCX file, 0.01 MB.

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M.A.Z. and J.N.W. designed the study. M.A.Z., S.H., A.J.H., and M.K. carried out the experiments. M.A.Z., W.W., and L.Z. did the data analysis. M.A.Z. and J.N.W. wrote the manuscript.

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