

Inhibition of *Streptococcus pneumoniae* growth by masarimycin.

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Abstract:

Despite renewed interest, development of chemical biology methods to study peptidoglycan metabolism has lagged in comparison to the glycobiology field in general. To address this, a panel of diamides were screened against the Gram-positive bacterium *Streptococcus pneumoniae* to identify inhibitors of bacterial growth. The screen identified the diamide masarimycin as a bacteriostatic inhibitor of *S. pneumoniae* growth with an MIC of 8 μ M. The diamide inhibited detergent-induced autolysis in a concentration-dependent manner, indicating perturbation of peptidoglycan degradation as the mode-of-action. Cell based screening of masarimycin against a panel of autolysin mutants, identified a higher MIC against a Δ lytB strain

lacking an endo-N-acetylglucosaminidase involved in cell division. Subsequent biochemical and phenotypic analyses suggested that the higher MIC was due to an indirect interaction with LytB. Further analysis of changes to the cell surface in masarimycin treated cells identified the overexpression of several moonlighting proteins, including elongation factor Tu which is implicated in regulating cell shape. Checkerboard assays using masarimycin in concert with additional antibiotics identified an antagonistic relationship with the cell-wall targeting antibiotic fosfomycin, which further supports a cell-wall mode-of-action.

Introduction

Antibiotic resistance is a growing global threat. Drug-resistant *Streptococcus pneumoniae* alone is estimated to cause more than 2 million infections with an excess of 1.3 billion USD in medical costs per annum.[1, 2] In light of this, there is need for the development of new therapeutics. Peptidoglycan (PG) is the primary structural heteropolymer conferring strength and cell shape determination in both Gram-negative and Gram-positive organisms (Figure 1). The PG polysaccharide backbone is composed of β -1,4-linked N-acetylmuramic acid (MurNAc) and N-acetylglucosamine (GlcNAc). Attached to the C-3 lactyl moiety of MurNAc is a stem peptide that is involved in cross-linking the adjacent glycan strands to form the three-dimensional structure of the cell wall. The incorporation of new material into the stress bearing layer of the existing cell wall requires the delicate homeostasis of biosynthetic and degradative enzymes to prevent lysis.[3, 4] Disruption of this interplay between degradative and biosynthetic enzymes via chemical inhibition could provide unique insights into their biological role. The degradative enzymes, collectively referred to as autolysins, are a broad class of enzymes that are differentiated based on their bond selectivity. Deciphering physiological activity of autolysins has been a formidable task as functional redundancy complicates attribution of biological activity.[5] Recent biophysical [4, 6] and computational studies [7] of bacterial autolysins have begun to unravel their roles in the release of stress in the cell wall to allow for incorporation of new material. A renaissance in PG metabolism research has started to provide new chemical biology tools to study synthesis [8-11] and the role endopeptidases play in methicillin resistance.[12] While the cell wall, and PG in particular, have provided a wealth of clinically relevant antimicrobial targets [13], our understanding of the complex interplay between degradative and synthetic steps is still developing.

Previously, we had screened a panel of 21 diamides for antibacterial activity against the Gram-positive *Bacillus subtilis*. [14] This screen identified the diamide masarimycin (formerly fgkc) (Figure 1 inset) as a bacteriostatic inhibitor of *B. subtilis* growth that targets the major active N-

acetylglucosaminidase (GlcNAcase) LytG (glycosyl hydrolase family 73 (GH73)) *in vitro*. Here we report on the screening of this panel of diamides against *S. pneumoniae*, identifying masarimycin as a bacteriostatic inhibitor of cell growth. Although an initial examination experiment implicates a related *S. pneumoniae* GlcNAcase (LytB, GH73) in masarimycin's activity, it is not the direct molecular target. A series of subsequent mode-of-action studies in *S. pneumoniae* highlights the challenges involved in target identification.

Materials and Methods

Strains and compounds. *Streptococcus pneumoniae* 6305 and R6 were purchased from ATCC (Mannassas, VA), and *S. pneumoniae* IU1945 (Δ lytB, Δ lytC, Δ dacA, Δ dacB, Δ pmp23, Δ bbp1a) mutants were kindly provided by Dr. Malcolm E. Winkler at the University of Indiana.[15] *S. pneumoniae* TIGR4 and TIGR4 Δ lytB strains were previously reported.[16] *S. pneumoniae* strains were grown in Mueller Hinton (MH) broth (MilliporeSigma, St. Louis, MO) supplemented with 5 % (v/v) sheep blood (Lampire Biological, Pipersville PA) or MH agar plates containing 1.5 % (m/v) Bacto agar and 5 % (v/v) sheep blood at 37°C under anaerobic conditions.

Staphylococcus aureus was grown in MH broth or solid media, *Clostridiodes difficile* was grown in brain heart infusion (BHI) and *Escherichia coli* DH5 α on Luria Bertani (LB) broth or solid media. Diamide inhibitors were synthesized as described previously.[14] Other reagents, unless otherwise specified, were purchased from MilliporeSigma (St. Louis, MO).

MIC assays. MIC values were determined using the resazurin method.[17] Briefly, cells were initially grown from the freezer on MH agar plates containing 5% defibrinated sheep blood. For all assays second passage cells of *S. pneumoniae* 6305, TIGR4, or R6 were used and grown overnight in MH broth, and standardized to an OD_{600nm} = 0.4. Inhibitors were analyzed *via* serial dilution into PBS media in microtitre plates. For masarimycin, dilutions were initially made in DMSO down to a concentration of 100 μ M, further dilutions were then made into PBS. Plates containing MH broth were inoculated with a 1/20 dilution of the OD_{600nm} = 0.4 cell culture with a final concentration of 1 % DMSO. Cultures were grown statically under anaerobic conditions for 24 h at 37 °C, followed by addition of 30 μ L of a 0.01% (m/v) solution of resazurin. The plates were incubated for 15 min to allow stabilization of color production. MICs were read directly off the plate; MICs were recorded as the lowest concentration that completely inhibited growth. MIC assays with *S. aureus* were performed in MH broth, *C. difficile* in BHI broth, while *E. coli* was performed in LB.

105 *Morphological studies of S. pneumoniae.* Cultures were prepared from second passage *S.*
106 *pneumoniae* 6305, R6 and Δ lytB [15] as previously described for MIC determination. Cells were
107 washed in phosphate-buffered saline and chemically fixed in 20 mM HEPES pH 6.8 containing
108 1% formaldehyde. [18] Samples were fixed overnight at 4 °C to limit *de novo* cell wall
109 biosynthesis during fixation and stained with 0.1% (m/v) methylene blue (solution in 20%
110 ethanol). Samples were gently heated to 60 °C for 15-20 min to bring cells to a common focal
111 plane. Slides were visualized using bright-field microscopy with a Zeiss Primo Star microscope
112 at 1000× magnification. Micrographs were acquired using an Axiocam ERc5s camera and Zen
113 lite software.

114 *Autolysis assays.* Cellular autolysis assays were performed as previously described by Cornett
115 and Shockman.[19] Briefly, *S. pneumoniae* 6305 were grown in MH broth containing 5% (v/v)
116 defibrinated sheep blood under anaerobic conditions. Cells were harvested by centrifugation
117 (8,000 rpm, 5 min) and washed with PBS. Cells were suspended in PBS and autolysis induced
118 with the addition of Triton X-100 to a final concentration of 0.1% (v/v) and turbidity monitored at
119 600 nm over 60 min. Rates were calculated using the linear portion of the autolysis curves with
120 the rate of autolysis in the absence of inhibitor set at 100%.

121 *Chain dispersing assay.* Dispersion of the Δ lytB chain morphology with purified LytB was carried
122 out as previously described using the TIGR4 and associated Δ lytB strains. [16] LytB was added
123 to the cell suspension at a final concentration of 2 μ M. The final concentration of masarimycin in
124 the assays was 40 μ M.

125 *DNA intercalation assays.* To determine if masarimycin intercalates DNA, DNA mobility shift
126 assays were performed as previously described using BamHI-linearized pUC18 plasmid.[20]
127 The known DNA intercalator actinomycin D was used as a control.

128 *Dnase I assays.* Degradation of the pUC18 plasmid DNA was assayed using 150 ng of
129 linearized pUC18 plasmid as described by Huang *et al.*[21] Compound titrations in DMSO were
130 added and reactions were initiated with 0.002 units of Dnase I. The reactions were incubated for
131 15 min at 37°C before being subjected to agarose gel electrophoresis. EDTA was used as a
132 control for Dnase I inhibition.

133 *Lipoteichoic acid detection by Western blot.* Lipoteichoic acid profiles were analyzed as
134 previously described.[22] Briefly, *S. pneumoniae* R6 cells were cultured overnight, harvested
135 (3000 x g), resuspended in 6 M urea, and incubated at 37°C for 5 min to solubilize proteins.
136 Samples were standardized by total protein content and separated by SDS-PAGE (16%) and

transferred to PVDF membrane. The membrane was incubated with a 1:5000 α -phosphocholine monoclonal antibody (SSI Biotech, Santa Cruz, CA). Blots were analyzed by chemiluminescence.

Analysis of cell-wall associated proteins. Changes to cell wall protein profiles were analyzed as previously described.[23] Briefly, second passage *S. pneumoniae* R6 were inoculated 1/100 into MH broth and grown statically for 6 h anaerobically at 37°C. Masarimycin was added to a final concentration of 0.75x MIC, the effect of solvent was controlled by the addition of DMSO to a second flask and the cells grown overnight at 37°C anaerobically. Cells were harvested into PBS containing sucrose (20% w/v) and pelleted at 8,000 rpm for 10 min. The pellets were washed with PBS containing 20% (w/v) sucrose and centrifuged again. The washed and pelleted bacteria were resuspended in 2 mL of 50 mM glycine-NaOH (pH 12.0) containing 20% (w/v) sucrose and incubated for 30 min at room temperature with gentle shaking. The suspension was centrifuged (10,000 rpm, 20 min). The supernatant was collected, and adjusted to pH 7 with 1 M HCl, proteins precipitated with acetone and analyzed by 1D-SDS PAGE and silver staining. All lanes were standardized to total A_{280nm} loaded onto the gel. Bands were excised from the gel and sent for identification by mass spectrometry at the National Resource for Proteomics (University of Arkansas).

Proteomic analysis of SDS-PAGE gel bands. Each SDS-PAGE gel band was subjected to in-gel trypsin digestion as follows. Gel segments were destained in 50% methanol, (Fisher), 50 mM ammonium bicarbonate, (Sigma-Aldrich), followed by reduction in 10 mM Tris[2-carboxyethyl]phosphine (Pierce) and alkylation in 50 mM iodoacetamide. (Sigma-Aldrich). Gel slices were then dehydrated in acetonitrile, (Fisher), followed by addition of 100 ng porcine sequencing grade modified trypsin (Promega) in 50 mM ammonium bicarbonate (Sigma-Aldrich) and incubation at 37°C for 12-16 hours. Peptide products were then acidified in 0.1% formic acid. (Pierce). Tryptic peptides were separated by reverse phase Xselect CSH C18 2.5 μ m resin (Waters) on an in-line 150 x 0.075 mm column using a nanoAcquity UPLC system (Waters). Peptides were eluted using a 60 min gradient from 98:2 to 65:35 buffer A:B ratio. [Buffer A = 0.1% formic acid, 0.5% acetonitrile; buffer B = 0.1% formic acid, 99.9% acetonitrile.] Eluted peptides were ionized by electrospray (2.4 kV) followed by MS/MS analysis using higher-energy collisional dissociation (HCD) on an Orbitrap Fusion Tribrid mass spectrometer (Thermo) in top-speed data-dependent mode. MS data were acquired using the FTMS analyzer in profile mode at a resolution of 240,000 over a range of 375 to 1500 m/z. Following HCD activation, MS/MS data were acquired using the ion trap analyzer in centroid mode and normal mass range with precursor mass-

dependent normalized collision energy between 28.0 and 31.0. Proteins were identified by database search using Mascot (Matrix Science) with a parent ion tolerance of 3 ppm and a fragment ion tolerance of 0.5 Da. Scaffold (Proteome Software) was used to verify MS/MS based peptide and protein identifications. Peptide identifications were accepted if they could be established with less than 1.0% false discovery by the Scaffold Local FDR algorithm. Protein identifications were accepted if they could be established with less than 1.0% false discovery and contained at least 2 identified peptides. Protein probabilities were assigned by the Protein Prophet algorithm.[24]

Antagonism assay and fractional inhibitory concentration index (FIC_{Index}) determination.

Fractional inhibitory concentration index (FIC_{Index}) was conducted to determine the interaction between masarimycin and a range of antibiotics with defined mode-of-action in a 96 well-plate microdilution broth assay. A checkerboard assay was performed with each masarimycin pair as previously described. [25] Plates were inoculated with 5 μ L of a $OD_{600nm} = 0.2$ culture of *S. pneumoniae* R6 and growth monitored as previously described for the MIC assays. FIC_{Index} was determined using the formulae:

(Eq 1) $FIC = X / MIC_x$, where X is the lowest inhibitory concentration of the drug in the presence of the co-drug, and MIC_x is the lowest inhibitory concentration of the drug in the absence of the co-drug.

(Eq 2) $FIC_{Index} = FIC_{masarimycin} + FIC_{antibiotic}$

Drug interactions were rated as synergistic ($FIC_{Index} \leq 0.5$), additive ($0.5 < FIC_{Index} \leq 1.0$), indifferent ($1.0 < FIC_{Index} \leq 4.0$), and antagonistic ($FIC_{Index} > 4.0$), based on published standards.[26]

NADP/NADP(H) ratio assays. Measurement of NADP/NADPH intracellular ratios was determined via the Amplite colorimetric NADP/NADPH ratio assay kit (Kit #15274; AAT Bioquest Inc, Sunnyvale CA) following manufacturer protocols. *S. pneumoniae* R6 second passage cells grown on MH with 5% defibrinated sheep blood under anaerobic conditions were used to inoculate 5 mL MH broth cultures ($OD_{600nm} = 0.5$) containing either masarimycin (1x, 3x MIC) or vehicle control. Cultures were grown under anaerobic conditions for 1.5 h and centrifuged (8,000 rpm, 10 min). Cell pellets were resuspended in PBS and lysed via sonication. Lysed cells were then used in assays following manufacturer's instructions. Samples were analyzed in a 96 well plate format using a Molecular Devices SpectraMax190 with detection at 570 nm. Data was analyzed using GraphPad Prism.

Results and Discussion

We screened a previously reported [14] panel of 21 diamides against *S. pneumoniae* using the resazurin microtiter assay (Figure S1).[17] Of the 21 compounds screened, masarimycin (formerly fgkc) was identified as a single digit micromolar bacteriostatic inhibitor (Figure S2) of *S. pneumoniae* growth with an MIC of 8 μ M against all three strains of *S. pneumoniae* that we tested – 6305, R6, and TIGR4 (Figure S3A). Three strains were chosen for screening to account for the known genomic plasticity among *S. pneumoniae* isolates which can manifest as varying antibiotic sensitivity between strains. [27-29] TIGR4 was included as it is capable of causing invasive disease.[30] These results for masarimycin are comparable to those obtained against *B. subtilis* (MIC = 3.8 μ M, bacteriostatic).[14] To further investigate masarimycin spectrum, the compound was screened against the Gram-positive organisms *Clostridiodes difficile*, *Staphylococcus aureus* and Gram-negative *Escherichia coli* which possess at least one GH73. In all cases, no antimicrobial activity was observed up to 150 μ M (Figure S3B). Extrapolating from the activity of masarimycin against *B. subtilis*, where it inhibits LytG, a GH73 enzyme, we hypothesized that the target of masarimycin in *S. pneumoniae* was also a GH73 family glycosidase. *S. pneumoniae* possesses one glycosidase classified as a member of the GH73 family (www.cazy.org) – LytB, a cell division associated endo- β -GlcNAcase belonging to cluster 4 of GH73.[16, 31-33] In contrast, *B. subtilis* LytG is an exo-acting GlcNAcase active in vegetative growth and belongs to GH73 cluster 2.[34]

Given the potential connection to LytB and cell wall metabolism, we examined whether masarimycin inhibited autolytic activity in *S. pneumoniae* (Figure 2A). Exposure of *S. pneumoniae* to low concentration of the non-ionic detergent (Triton X-100) induces autolysis.[19] This autolytic activity was inhibited in a concentration-dependent manner by masarimycin. The elevated concentration for near complete inhibition of autolysis in the whole cell assay is likely due to the broad dysregulation of autolysins induced by Triton X-100. Given the observed inhibition of autolytic activity, the MIC of masarimycin was determined for a series of *S. pneumoniae* mutant strains lacking the autolysins *lytA*, *lytB*, *pmp23*, *dacA*, or *dacB*, as well as the bifunctional *pbp1a*. It has been demonstrated that deletion of any gene that affects PG biosynthesis, stability, or regulation can make the bacterium more susceptible to compounds that target the cell wall. [27] While none of the mutants were more sensitive to masarimycin, higher MICs were observed with *lytB* (GH73 GlcNAcase), *pmp23* (muramidase), and *dacA* (*pbp3*, D,D-carboxypeptidase) mutants. This is counter to what was observed in a screen of autolysin mutants in *B. subtilis*.[14] The near 4-fold decrease in sensitivity to masarimycin in the

Δ lytB mutant suggests that changes to the cell wall imparted by the lack of LytB reduces sensitivity to masarimycin.

To further explore these results, morphological changes induced by sub-MIC (0.7x) concentrations of masarimycin, and antibiotics with known modes-of-action were investigated (Figure S4). Treatment of *S. pneumoniae* with the cell wall-acting antibiotics bacitracin, vancomycin, as well as the protein synthesis inhibitor kanamycin presented a phenotype of clumping cells. Sub-MIC treatment with masarimycin showed a change similar to these antibiotics. The clumping phenotype observed with kanamycin has been associated with antibiotics that target intracellular protein synthesis.[35] In light of this, the clumping phenotype could not be directly attributed to a cell wall mode-of-action. Comparison of the masarimycin-induced phenotype with the reported phenotypes of a Δ lytB[15], Δ pmp23[36], or Δ dacA[37] mutants, for which higher MICs with masarimycin were observed, did not correlate. Additionally, the masarimycin-induced phenotype did not correspond to phenotypes of other *S. pneumoniae* autolysin mutants. [38-40] The masarimycin phenotype more closely resembles a morphology in which the autolysin is still produced but is catalytically inactive, such as that reported for E61Q and D68N mutants of pmp23. [36]

To further probe alterations to the cell wall suggested by the autolysis and genetic screen assays, lipoteichoic acid (LTA) disruption was monitored by Western blot in the presence of sub-MIC masarimycin. LTA has been shown to regulate autolysin activity in several species and LytB possesses a choline binding domain, a component of *S. pneumoniae* LTA.[41-43] Additionally, it has been suggested that LytB function is altered when cell wall choline content is depleted.[44] Results indicated that changes to LTA and choline incorporation in the cell wall was not a contributor to the observed masarimycin-induced autolysis, genetic screen, and morphology phenotypes (Figure S5A).

Next, changes to cell-wall-associated protein profiles were examined using high-pH extraction[23] of the *S. pneumoniae* cell surface (Figure S5B). Upon treatment with 0.75x MIC masarimycin the appearance of several overexpressed proteins was observed. Proteomic analysis of SDS-PAGE gel bands identified several cell surface and moonlighting proteins that are present only in the masarimycin treated sample (Table 1, Supplementary dataset 1). Of note is the upregulation of elongation factor Tu, a known moonlighting protein that has been implicated in regulating cell shape by modulating the formation of MreB filaments in *B. subtilis* and *E. coli*.[45] The overexpression of proteins involved in complex carbohydrate catabolism (BgaA, MalX) have been previously shown to be regulated by the two-component system

CiaR/H, which is also associated with sensing cell-wall stress in *S. pneumoniae*. [46] These changes to proteins on the cell surface suggest that masarimycin is interfering with cell wall remodeling.

Table 1. Overexpressed proteins observed in SDS-PAGE gel (Figure S5B) of *S. pneumoniae* R6 cell-surface protein extracts when exposed to 0.75x MIC masarimycin.

Gel Band	Protein ID	Biological Function
1	1. BgaA β -galactosidase 2. Iga immunoglobulin A1 protease	1. Plays role in growth and adhesion[47] 2. Covalently linked to cell surface by sortase A. Involved in host immune evasion. [48]
2	Spr0440, endo- β -GlcNAcase	Surface protein, role in commencement of neuroinvasion[49]
3	1. Elongation factor Tu 2. PykF Pyruvate kinase	1. Moonlighting function in regulation of cell shape[50] 2. Moonlighting protein [51, 52]
5	MalX maltooligosaccharide transporter	Complex carbohydrate catabolism regulated by CiaR/H system which is involved in sensing cell wall stress.[46, 53]
6	GapA glyceraldehyde 3-phosphate dehydrogenase	Moonlighting protein[52]

To further investigate the higher MIC observed with the Δ lytB mutant, inhibition of LytB activity was investigated in an established chain dispersing assay using the TIGR4 Δ lytB strain in the presence of exogenously added LytB.[16] When the Δ lytB mutant was treated with exogenous LytB, dispersion of the chains was observed (Figure 3). When a 5-fold MIC (40 μ M) concentration of masarimycin was added, LytB catalyzed chain dispersion was not inhibited. In vitro analysis with Remazol Brilliant Blue labeled PG [16, 54] and purified LytB confirmed these results. This lack of inhibition of the biochemical activity of purified LytB further suggests that the higher MIC of masarimycin observed for the Δ lytB mutant may be due to changes in cell wall structure, stability, or metabolism. These results further highlight the difficulty in target identification of small molecule inhibitors. The reduced MIC values observed in the genetic screen against the Δ lytB strain is likely due to a more complex interplay between the deletion

of LytB and the actual target(s) of masarimycin. Consistent with this hypothesis, treatment of ΔlytB with 0.7x MIC masarimycin (Figure 4) resulted in the conversion of the ΔlytB chaining phenotype to the clumping phenotype observed in Figure S4. Co-administration of sub-MIC concentrations of cefoxitin, a DacA (PBP3) selective β -lactam, with masarimycin resulted in the reduction of the clumping phenotype in the ΔlytB mutant but not wildtype cells. This further suggests that masarimycin's mode-of-action is impacted by alterations to the cell wall caused by either genetic deletion or chemical inhibition of autolysins. LytB is implicated in PG remodeling during cell division while DacA is associated with division rather than remodeling. [15] These observations suggest that masarimycin is influencing cell wall biosynthesis, turnover, stability, and/or regulation in these mutant backgrounds. Given their roles in cell wall remodeling during division (LytB) or directly involved in division or division complex (DacA, Pmp23) and the overexpression of elongation factor Tu suggests that masarimycin is either directly or indirectly impacting the cell wall during division.

Taken collectively, the reduction in autolysis, phenotypic changes and overexpression of surface associated proteins observed with masarimycin, along with the reduced sensitivity against multiple mutant strains, suggest that these results could be the result of meta-phenotypes - a phenotype that results from the alteration of more than one pathway.[55] For instance, the clumping meta-phenotype observed in *S. pneumoniae* in the presence of masarimycin could be generated via a direct mechanism (e.g. inhibition of an enzyme associated with cell wall metabolism) or an indirect one (e.g. alterations in autolysin expression levels, changes in metabolic flux through cell wall associated pathways).

To further interrogate the potential mode-of-action, masarimycin was screened in checkerboard assays with antibiotics with well-defined mechanisms (Table 2). Using pre-established guidelines [26] for interpreting $\text{FIC}_{\text{index}}$, two antibiotics, levofloxacin (DNA gyrase) and fosfomycin (MurA, first committed step of PG biosynthesis) demonstrated mild antagonism ($\text{FIC}_{\text{index}}$ 4.5). Antagonistic relationships can be used to map genetic networks and reveal novel connections between pathways.[25] Antagonism with fosfomycin suggests a functional connection to the target of masarimycin. To further probe the antagonism with fosfomycin, we looked to see if masarimycin was indirectly impacting fosfomycin's target MurA. The subsequent step, catalyzed by MurB reduces the product of MurA, UDP-GlcNAc-enolpyruvate, to UDP-MurNAc utilizing NADPH. [56] Given the impact that changes to the cell wall have on masarimycin activity, we wondered if the observed antagonism with fosfomycin might be due to altered redox potential in the cell, a consequence of a destabilized cell envelope. This could

impact PG metabolite flux through MurA/B by reducing the levels of NADPH in the cell. Alterations in redox potential and oxidative stress can negatively influence fosfomycin sensitivity.[57] It has been previously demonstrated in *S. aureus* that alterations in metabolic flux of precursors and cofactors can contribute to fosfomycin resistance.[58] Changes in NADP/NADPH levels were measured in a colorimetric assay in the presence/absence of masarimycin. The ratio of NADP/NADPH did not change when up to 3x MIC masarimycin was present (Figure S6) suggesting the fosfomycin antagonism is not due to an altered redox potential impacting flux through MurB.

Antagonism with levofloxacin provides a counterpoint to the fosfomycin results. Levofloxacin introduces double-stranded DNA breaks upon inhibition of DNA gyrase, and induces the SOS response system.[59] To further evaluate this antagonistic relationship, masarimycin was evaluated for its ability to interact with DNA. Masarimycin was investigated for its ability to inhibit nuclease activity and intercalate DNA in established assays (Figure S7). [20, 21] Masarimycin showed no inhibition of nuclease activity or ability to intercalate DNA, suggesting the antagonism observed is likely not due to direct interaction with DNA or nucleases. Further the additive interaction with rifampicin (RNA synthesis) and the protein synthesis inhibitors tetracycline and kanamycin in Table 2 suggests a pure summation effect of the antibiotics with masarimycin. This suggests the antagonism with levofloxacin is not a product of downstream inhibition of transcription and translation. It should be noted that there is a connection between quinolone bactericidal activity and the expression of stress-induced proteins.[60] Cell wall targeting antibiotics like β -lactams can activate the SOS response system.[61] The antagonism observed with levofloxacin can be explained by the induction of the SOS system by both compounds.[62]

Table 2. Synergy and antagonism screen with masarimycin.

Antibiotic	MIC _{app} antibiotic ^a (μM)	FIC antibiotic	MIC _{app} masarimycin ^b (μM)	FIC masarimycin	FIC _{index}
Ampicillin	0.010	1	2	0.25	1.25
Bacitracin	1.66	0.5	15.6	2	2.5
Cefoxitin	0.69	0.4	4	0.5	0.9
Cefuroxime	0.012	1.59	2	0.25	1.75
Fosfomycin	85.87	0.5	32	4	4.5
Kanamycin	43	0.5	1.66	0.21	0.71
Levofloxacin	1.08	0.5	32	4	4.5
Optochin	7.96	0.5	8	1	1.5
Rifampicin	0.0023	0.124	3.33	0.416	0.54
Tetracycline	0.048	0.75	2	0.25	1
Vancomycin	0.003	1	4	0.5	1.5

^a – MIC for antibiotics alone. ampicillin: 0.010 μM; bacitracin: 3.325 μM; cefoxitin: 1.72 μM; cefuroxime: 0.008 μM; fosfomycin: 171.5 μM; kanamycin: 86 μM; levofloxacin: 2.16 μM; optochin: 15.85 μM; rifampicin: 0.018 μM; tetracycline: 0.065 μM; vancomycin: 0.003 μM.

^b-MIC masarimycin 8 μM

Conclusion

Taking this data holistically, we posit that masarimycin activity is impacted by alterations to the cell wall caused by deletion of certain autolysins (LytB, DacA, Pmp23) in *S. pneumoniae*. These three autolysins are associated with remodeling PG during division (LytB)[31], or associated with division (DacA)[39] or with the Z-ring (Pmp23)[36] components of the cell division complex. Other PG hydrolases [63-65] might also be affected by masarimycin treatment, giving rise to the complex phenotypic results. The data presented here demonstrates that the morphological phenotypes of genetic knockouts of cell wall acting enzymes can be distinct from chemical inactivation and may more closely resemble the phenotype of catalytically inactive mutants. This distinction has previously been observed in *Mycobacterium tuberculosis* shikimate biosynthesis.[66] These data illustrate that the use of genetic and phenotypic screens for target identification may not always lead directly to the molecular target. The complexity involved in deciphering the underlying mechanisms associated with these meta-phenotypes obfuscates target identification. Despite these challenges, the elucidation of the molecular target of masarimycin is on-going. Based on the data provided here, masarimycin may provide a unique molecular scaffold for the development of anti-*S. pneumoniae* therapeutics and can play a role in furthering our understanding of PG metabolism.

The bacterial cell wall and PG biosynthesis has provided a wealth of clinically relevant antibiotic targets. While our understanding of PG biosynthetic and cross-linking steps is well established, our knowledge of the role autolytic enzymes play in the growth and maintenance of the cell wall has remained more elusive. Traditional genetic approaches to studying the biological role of autolysins are complicated by functional redundancy of these enzymes, where other autolysins can compensate for a loss in activity. The results presented here illustrate the complexity of PG metabolism and the difficulty in identifying the molecular target of small molecule inhibitors. Recent reports [4, 6, 7] have begun to elucidate the role of autolysins in relieving stress in the cell wall to allow for incorporation of new material into the stress bearing layer. The results with the diamide masarimycin demonstrate that sensitivity is impacted by alterations in the cell wall caused by the deletion of specific autolysins associated with cell division and separation. Upregulation of elongation factor Tu, a moonlighting protein known to regulate cell shape, indicates that masarimycin is impacting cell wall metabolism. Collectively our data further demonstrate that morphological, genetic, and whole cell assays (autolysis) reveal meta-phenotypes that result from the complex interaction of one or more cellular processes that appear connected to cell wall metabolism. The genetic deletion of one or more autolysins disrupts the equilibrium stoichiometry of the cell wall machinery that likely results in changes to expression levels and activity to both autolytic and biosynthetic enzymes. With interest in the development of chemical biology approaches to study PG metabolism[67-71] receiving renewed attention, the diamide inhibitor masarimycin provides a potential small molecule complement to both traditional genetic and current chemical biology approaches to studying cell wall metabolism.

Conflict of Interest

The authors A.B. and C.W.R hold patents on specific applications of masarimycin.

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Figures and Tables

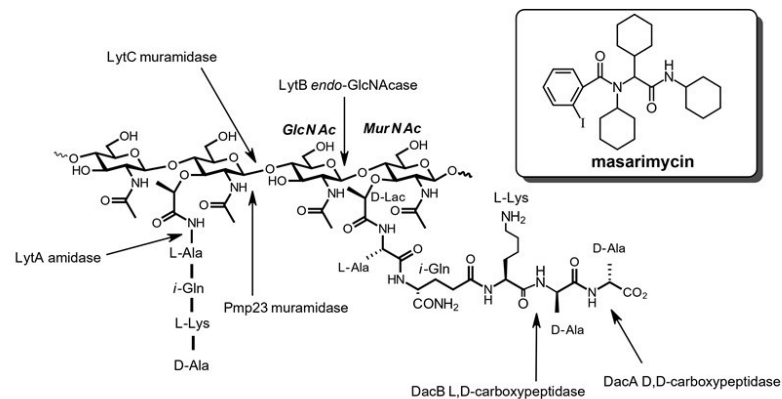
Figure 1. Structure of PG showing the cleavage sites of several of the characterized autolysins in *S. pneumoniae*. Inset: structure of the antimicrobial diamide masarimycin.

Figure 2 Screening of the diamide masarimycin against *Streptococcus pneumoniae*. **(A)** The diamide masarimycin inhibits detergent-induced autolysis in a concentration dependent manner. Percent residual activity was calculated using autolysis in the absence of inhibitor set as 100%. Data shown is the average of experiments performed in biological and technical triplicate. Error bars denote standard deviation. **(B)** Activity of masarimycin against *S. pneumoniae* R6 autolysin and cell wall biosynthesis mutants (13) to identify changes to masarimycin sensitivity. Assays were run in biological triplicate and yielded the same MIC values.

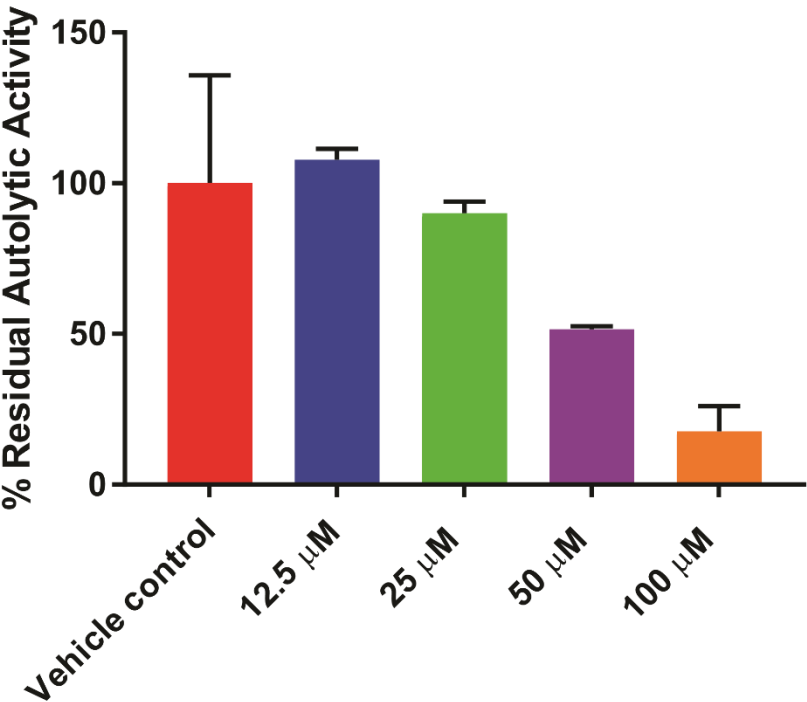
Figure 3. Chain dispersing assay with *S. pneumoniae* TIGR4 Δ lytB strain and purified recombinant LytB (rLytB, 2 μ M). In the presence of 40 μ M masarimycin dispersion of the Δ lytB chain phenotype is not inhibited. Images were taken at 1000x magnification.

Figure 4. Morphological analysis of *S. pneumoniae* Δ lytB mutant-[15] in the presence of sub-MIC masarimycin, the β -lactam cefoxitin (DacA/PBP3 selective) or in combination. Cells were fixed in 1% formaldehyde, stained with methylene blue, and visualized using bright field microscopy under oil immersion.

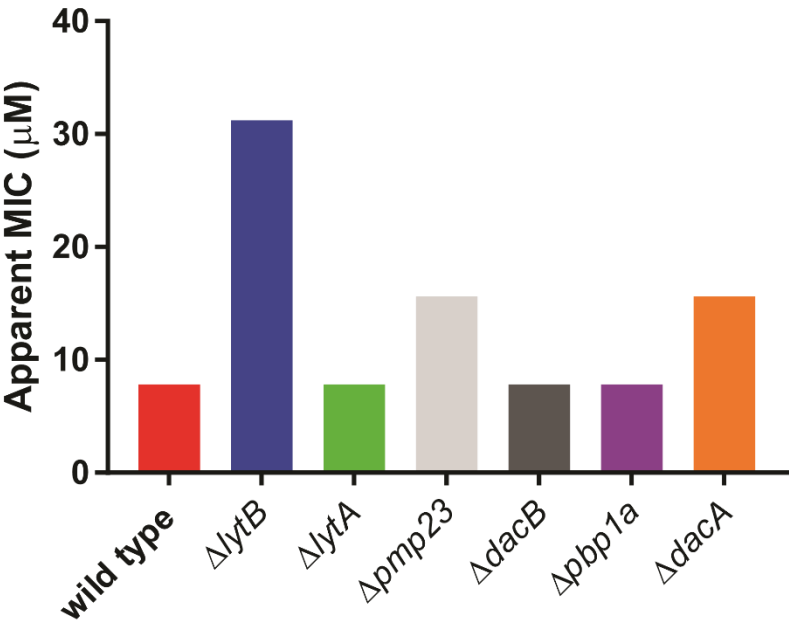
624 Figure 1.



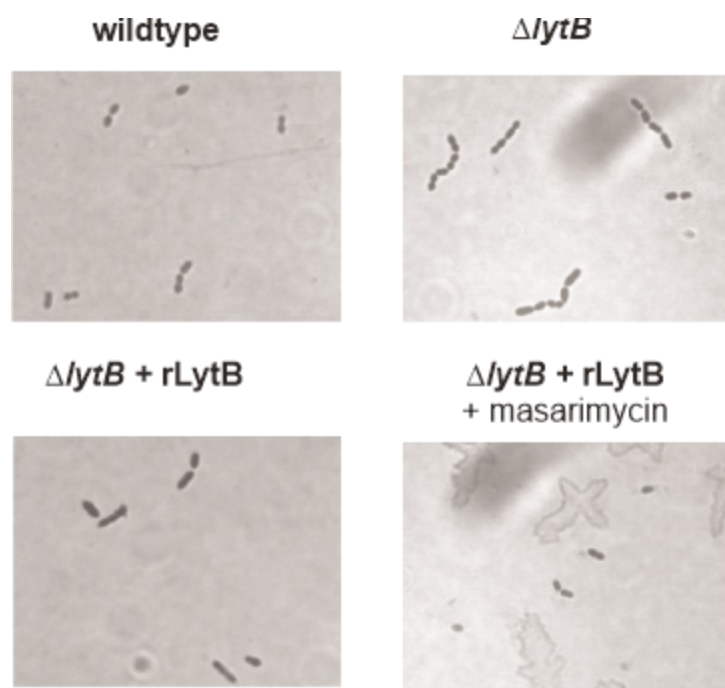
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643 Figure 3.



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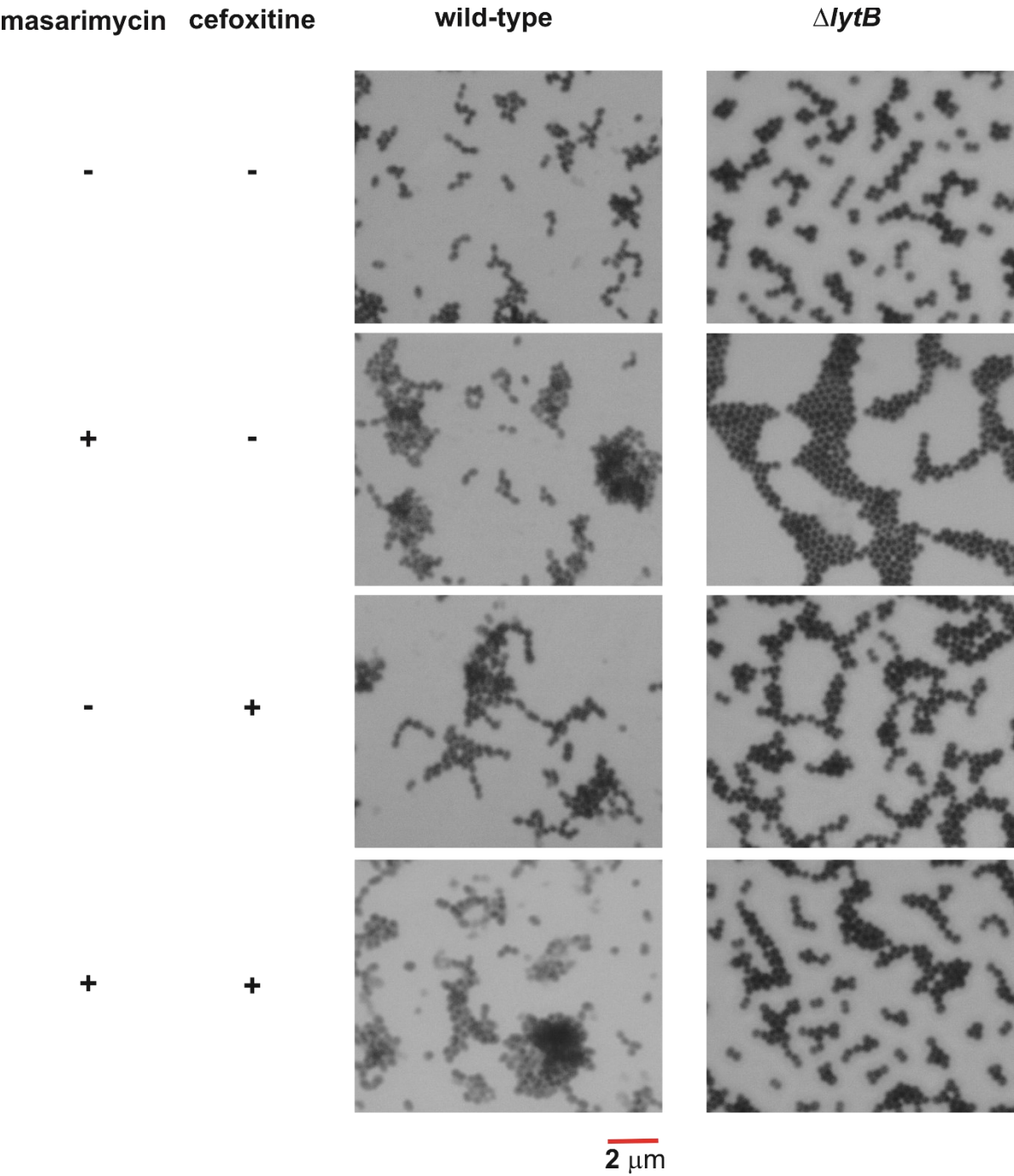
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658 Figure 4.



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