

1 **Pneumococcal attachment to epithelial cells is enhanced by the secreted peptide VP1**
2 **via its control of hyaluronic acid processing**

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9 **ABSTRACT**

10 The Gram-positive bacterium *Streptococcus pneumoniae* (pneumococcus) is an
11 important human pathogen. It can either asymptotically colonize the nasopharynx or spread
12 to other tissues to cause mild to severe diseases. Nasopharyngeal colonization is a
13 prerequisite for all pneumococcal diseases. We describe a molecular pathway utilized by
14 pneumococcus to adhere to host cells and promote colonization. We demonstrate that the
15 secreted peptide VP1 enhances pneumococcal attachment to epithelial cells. Transcriptional
16 studies reveal that VP1 triggers the expression of operons involved in the transport and
17 metabolism of hyaluronic acid (HA), a glycosaminoglycan present in the host extracellular
18 matrix. Genetic experiments in the pneumococcus reveal that HA processing locus (HAL)
19 promotes attachment. Further, overexpression of HAL genes in the $\Delta vp1$ background, reveal
20 that the influence of VP1 on attachment is mediated via its effect on HA. In addition, VP1 also
21 enhances degradation of the HA polymer, in a process that depends on the HAL genes. siRNA
22 experiments to knockdown host HA synthesis support this conclusion. In these knockdown
23 cells, attachment of wild-type pneumococci is decreased, and VP1 and HAL genes no longer
24 contribute to the attachment. Finally, experiments in a murine model of colonization reveal that
25 VP1 and HAL genes are significant contributors to colonization. Our working model, which
26 combines our previous and current work, is that changes in nutrient availability that influence
27 CodY and Rgg144 lead to changes in the levels of VP1. In turn, VP1 controls the expression of
28 a genomic region involved in the transport and metabolism of HA, and these HAL genes
29 promote adherence in an HA-dependent manner. VP1 is encoded by a core gene, which is
30 highly induced *in vivo* and is a major contributor to host adhesion, biofilm development,
31 colonization, and virulence. In conclusion, the VP1 peptide plays a central role in a pathway

32 that connects nutrient availability, population-level signaling, adhesion, biofilm formation,
33 colonization, and virulence.

34 AUTHOR SUMMARY

35 *Streptococcus pneumoniae* (the pneumococcus) is a major human pathogen. This
36 bacterium asymptotically colonizes the human upper respiratory tract from where it can
37 disseminate to other tissues causing mild to severe disease. Colonization is a prerequisite for
38 dissemination and disease, such that the molecules that control colonization are high-value
39 candidates for therapeutic interventions. Pneumococcal colonization is a population-level
40 response, which requires attachment to host cells and biofilm development. VP1 is a signaling
41 peptide, highly induced in the presence of host cells and *in vivo*, promotes biofilm
42 development, and serves as a potent virulence determinant. In this study, we build on the
43 molecular mechanism of VP1 function to reveal novel bacterial and host molecules that
44 enhance adherence and colonization. Our findings suggest that host hyaluronic acid serves as
45 an anchor for pneumococcal cells, and that genes involved in the transport and metabolism of
46 HA promote adherence. These genes are triggered by VP1, which in turn, is controlled by
47 regulators that respond to nutrient status of the host. Finally, our results are strongly supported
48 by studies in a murine model of colonization. We propose that VP1 serves as a marker for
49 colonization and a target for drug design.

50 INTRODUCTION

51 The Gram-positive bacterium *Streptococcus pneumoniae* (also known as the
52 pneumococcus) is an important human pathogen. A recent global study on lower respiratory
53 infections determined that the pneumococcus contributed to morbidity more than all other
54 etiologies combined: it was responsible for an estimated 1.18 million deaths (1,2). This
55 pathogen can either asymptotically colonize the nasopharynx or spread to other tissues to
56 cause mild to severe diseases. It can spread to the middle ear and sinus, leading to otitis
57 media or sinus infections, and the lungs causing pneumonia (3). It can also disseminate into
58 the bloodstream, brain or heart causing sepsis, meningitis, or heart disease, respectively (4,5).
59 It is well established that nasopharyngeal colonization is a prerequisite for all pneumococcal
60 diseases. In this study, we explore the mechanisms of colonization and virulence.

61
62 Pneumococcus secretes many small peptides, which influence colonization, virulence,
63 and adaptation via effects on competence, intra-species competition, and biofilm development
64 (6–18). The **Virulence Peptide 1 (VP1)** is a member of a family of peptides with a conserved N-
65 terminal sequence characterized by a double glycine motif, which directs its export into the
66 extracellular milieu via ABC transporters (7,19). The *vp1* gene is widely distributed across
67 pneumococcal strains, as well as encoded in related streptococcal species. The gene
68 encoding VP1 is a virulence determinant and is highly upregulated in the presence of host cells
69 where it promotes biofilm development (7,20). Pneumococcal biofilms have been documented
70 in the nasopharynx, the middle ear, and the sinus (21–24). This mode of growth enhances
71 pneumococcal dissemination and pathogenesis (25,26). Further, biofilms promote
72 pneumococcal colonization in multiple ways: they facilitate immune evasion, increase
73 resistance to antimicrobials, and provide a platform for DNA acquisition (27–29,29–32). In

74 summary, VP1 is a secreted peptide, induced in the presence of human cells, that promotes
75 biofilm development and virulence.

76

77 Glycosaminoglycans (GAGs) are major components of the extracellular matrix. GAGs
78 are acidic linear polysaccharides with a repeated disaccharide unit. The disaccharides consist
79 of an amino sugar (*N*-acetylglucosamine or *N*-acetylgalactosamine) and a uronic acid
80 (glucuronic acid or iduronic acid) or galactose. These polymers are classified into groups
81 based on the nature of the disaccharides, the mode of glycoside bond, and the sulfation levels
82 (33). The main groups are heparin and heparan sulfates (HS), chondroitin and dermatan
83 sulfate (CS), keratan sulfate, and finally hyaluronic acid (HA) (also known as hyaluronate or
84 hyaluronan) (34). Their molecular weights range from 15 to over 100 kDa, and their synthesis
85 takes place in the Golgi apparatus, with the exception of HA, which is synthesized at the
86 membrane (35). GAGs participate in numerous biological processes of the host and the
87 bacteria. In the host, they influence cell adhesion and growth, cell proliferation and
88 differentiation, and tissue formation (for comprehensive reviews, please see Iozzo and
89 Schaefer, 2015; Pomin and Mulloy, 2018; Taylor and Gallo, 2006). In the context of microbial
90 infections, they serve as sources of nutrients and modulators of the immune response (38–41).
91 In addition, many bacteria utilize GAGs to adhere to host cells (42). This includes the
92 pneumococcus, where treatment of lung cells to decrease HS and CS levels reduces
93 pneumococcal adherence (43). In this manner, GAGs are critical components of the bacterial
94 environment during biofilm formation and host interactions.

95

96 HA consists of disaccharides of *N*-acetyl-D-glucosamine (GlcNAc) and D-glucuronic
97 acid (GlcUA). This GAG is present on the apical surface of human bronchial epithelial cells and
98 in the airway mucosa (44,45). It is also abundant in the synovial fluid, skin, umbilical cord, and

99 vitreous body and is the only non-sulfated glycosaminoglycan in the lung (46–48). This
100 unbranched polysaccharide provides mechanical support, activates immunity, and modulates
101 cell proliferation, migration, and intracellular signaling within the host (49–51). Many
102 streptococcal genomes encode multiple operons for HA transport and metabolism. In the
103 pneumococcus, it has been shown that HA processing and internalization is carried out by a
104 unique set of enzymes including a hyaluronidase, a transporter system, and a hydrolase
105 (52,53). The current model suggests that HA is digested into disaccharides by a hyaluronidase
106 (*hysA* or *hyla*), predicted to be covalently linked to the cell wall via an LPxTG motif. The
107 disaccharides are imported into the bacteria and phosphorylated via a phosphoenolpyruvate-
108 dependent phosphotransferase system (PTS) system. The PTS system is composed of
109 general and substrate-specific components. Enzyme I (EI) and the histidine-containing
110 phosphocarrier protein (HPr) are cytoplasmic subunits, shared by multiple PTS systems. In
111 contrast, Enzyme II (EII) provides specificity to the PTS via substrate recognition. EII has four
112 enzymes: EIIA and EIIB are cytosolic, and EIIC and EIID are the transmembrane components
113 and form a pore. During internalization via the PTS, the HA disaccharides are phosphorylated.
114 These disaccharides are processed by an unsaturated glucuronyl hydrolase (*ugl*) into the
115 monosaccharide components. Finally, the GlcUA can be further processed into glyceraldehyde
116 3-phosphate and pyruvate by a set of four genes (the isomerase *kduL*, the reductase *kduD*,
117 the kinase *kdgK*, and the aldolase *kdgA*). Operons for the hydrolase, the PTS-EII system, and
118 the GlcUA metabolism enzymes are all adjacent in the genome, and under the control of
119 RegR, a negative regulator located immediately downstream (54). Note that the unsaturated
120 glucuronyl hydrolase not only cleaves HA into GlcNAc and GlcUA, it can also cleave
121 chondroitin sulfate into their component monosaccharides (53,55). The HA pathway has been
122 shown to influence pneumococcal biology. Specifically, *hysA* and the genes in the PTS-EII-

123 encoding operon (EIIB, EIIC, EIID, and *ugl*), are required for pneumococcal growth on human
124 HA, and contribute to pneumococcal biofilm formation and virulence (54,56,57).

125

126 In this study, we show that VP1 promotes pneumococcal adherence via activation of HA
127 transport and metabolism. Our experiments in a murine model of pneumococcal carriage
128 demonstrate that both VP1 and HA processing promote colonization. We propose a model
129 where the nutritional status of the host induces expression of the pneumococcal secreted
130 peptide VP1, via CodY and Rgg144 (7,18,58). Signaling by VP1 triggers HA metabolism in the
131 bacteria, which in turn, regulates attachment to host cells and colonization in the upper
132 airways.

133 RESULTS

134 ***vp1* enhances pneumococcal attachment to epithelial cells**

135 We have previously shown that *vp1* plays a role in biofilm development (7). Thus, we
136 investigated whether *vp1* influences attachment to epithelial cells, an early step in biofilm
137 formation. For these studies, we utilized strain PN4595-T23, a representative of the drug-
138 resistant and pandemic PMEN1 pneumococcal lineage (59–62). To show that the results
139 extend beyond one lineage, we also employed TIGR4, a clinical isolate frequently used as a
140 model strain (63). For host cells, we utilized the human lung epithelial cells line A549. These
141 cells have been extensively used to study pneumococcal behavior (20,64,65). Early works
142 demonstrated that A549 cells display features of type II alveolar pneumocytes: they are
143 capable of surfactant production (66–68), they produce cell surface-associated MUC1 and
144 secrete MUC5AC mucin in air-liquid cultures (69,70), and they produce HA. Further, our
145 experiments on adhesion are extended to NIH-3T3 fibroblasts as they express HA and
146 chinchilla middle ear epithelial cells (CMEE) as these primary cells were used in the original
147 characterization of VP1 function (7,71).

148

149 We established that the wild-type strain displays strong attachment to A549 cells after
150 one hour of exposure (Fig S1A). Thus, we used this time point to compare adhesion between
151 wild-type, *vp1* deletion mutant ($\Delta vp1$) and *vp1* complement strains ($\Delta vp1:vp1$) by using spot
152 plating assays and enumerating the total bound bacteria on TSA plates. The $\Delta vp1$ displays a
153 reduction in attachment compared to the wild-type cells. Furthermore, the wild-type phenotype
154 is rescued in a *vp1* complement strain ($\Delta vp1:vp1$) (Fig 1A). These data suggest that VP1
155 promotes attachment to the lung epithelia. This role for the *vp1* gene product is not specific to
156 A549 cells, as *vp1* also promotes adherence to chinchilla middle ear epithelial cells (CMEE)

157 and NIH-3T3 fibroblasts (Figs S1B and S1C). Similarly, these results extend beyond the
158 PMEN1 lineage, deletion of the operon encoding *vp1* (*vpo*) in a TIGR4 background also
159 displays decreased attachment relative to the wild-type strain and this phenotype is restored to
160 wild-type levels in a *vpo* complement strain (Fig S1D). Adherent bacteria were visualized by
161 fluorescence microscopy of A549 cells inoculated with PMEN1 cells (Fig 1B). The confocal
162 orthogonal view shows localization at the surface of A549 cells (**Fig S1E**). Finally, the
163 difference in attachment is not the result of variation in growth rate between strains, as growth
164 in DMEM media is the same for wild-type, $\Delta vp1$, and $\Delta vp1:vp1$ strains, as previously reported
165 (7). We conclude that *vp1* promotes pneumococcal attachment to mammalian epithelial cells.

166

167 **The *vp1* product positively regulates expression of HA transport and metabolism genes.**

168 The VP1 pro-peptide possesses a double glycine motif common to secreted peptides
169 that bind surface-bound histidine kinase receptors from pneumococcal two-component signal
170 transduction system (72). Further, synthetic VP1 binds to the pneumococcal surface (7). Thus,
171 we hypothesized that VP1 signals to producing and neighboring cells and induces expression
172 of genes that participate in the attachment.

173

174 To test this hypothesis, we compared the genome-wide expression of the wild-type
175 PN4595-T23 strain and the isogenic $\Delta vp1$ mutant using the pangenome pneumococcal array
176 (13) (Fig 2A) Strains were grown in chemically defined media (CDM) supplemented with
177 glucose (CDM-Glu), as *vp1* expression is induced in this media relative to rich media (7). In
178 support of the hypothesis, twenty-eight genes displayed a 3-fold difference in expression
179 between the wild-type strain and the $\Delta vp1$ strain (Fig 2A and S1 Table). Many of these genes

180 are organized into three genomic regions (Fig 2B). Microarray results were validated by qRT-
181 PCR for a subset of genes (S2 Table). Taken together, we observed a significant number of
182 genes with transcription levels that respond to *vp1*.

183

184 Our analysis shows that two neighboring operons display higher expression in the wild-
185 type strain relative to the $\Delta vp1$; these operons have been previously implicated in the
186 processing of HA (53). Together, these two operons encode eleven genes that displayed an
187 average 4.28-fold decrease in expression in the $\Delta vp1$ relative to the wild-type strain (Figs 2B
188 and 3A). One operon is on the positive strand and encodes seven genes. In this study, for
189 simplicity, we refer to this as the PTS-HAL operon as it encodes the four EIIABCD enzymes
190 that form the PTS importer (Fig 3A). This importer is required for phosphorylation and
191 internalization of HA (EII genes are: SPN23F_02950, and SPN23F_02970 to SPN23F_02990).
192 The operon also contains the *ugt* gene that encodes an unsaturated glucuronyl hydrolase
193 (SPN23F_02960), the preprotein translocase YajC that is a subunit of the bacterial holo-
194 translocon (SPN23F_03000) (73), and finally a predicted heparinase (SPN23F_3010). Our
195 analysis with Phyre2 (74), indicates that the heparinase is a glycosidase that belongs to the
196 superfamily of heparinases type II/III; member of this family bind and degrade heparin and
197 heparin sulfate (75). It is noteworthy, that 60 base-pairs downstream of the end of *hep*, is the
198 negative regulator RegR (SPN23F_3020) (76). Analyses of the region upstream of RegR with
199 the promoter identification programs BDPG and BPRM suggests the presence of an active
200 promoter with the -35 within the *hep* sequence (77,78). The second operon regulated by VP1
201 is immediately upstream of the PTS-HAL operon on the negative strand, it encodes four
202 enzymes predicted to metabolize glucuronic acid into glyceraldehyde-3 phosphate and

203 pyruvate (SPN23F_02910 to SPN23F_02940). These transcriptional findings suggest that *vp1*
204 triggers the expression of genes implicated in the transport and metabolism of HA.

205

206 **The *vp1* product promotes degradation of the HA polymer via its control of the HA**
207 **processing and transport operon.**

208 Once we established that VP1 stimulates expression of multiple genes involved in HA
209 metabolism, we tested whether VP1 could influence the processing of the HA polymer. To this
210 end, we measured the hyaluronidase activity in pellets and supernatants of the PN4595-T23
211 strain carrying deletions of *vp1*, PTS-HAL, and various genes involved in HA processing (Fig
212 3B). In this assay, bacterial pellets or supernatants were mixed with the HA polymer,
213 degradation of HA was measured as a change in optical density at OD₄₀₀ and reported as
214 hyaluronidase concentration (76). As controls, we employed the $\Delta hysA$, as this gene encodes
215 for a hyaluronidase and its deletion should result in reduced hyaluronidase relative to the wild-
216 type strain (53,79). As predicted, this strain displayed 2.6-fold lower hyaluronidase levels ($p <$
217 0.0001 relative to the wild-type strain). We also employed a $\Delta regR$ mutant, as this gene
218 encodes for a repressor of *hysA* and its deletion should lead to increased hyaluronidase
219 activity (54). The $\Delta regR$ strain exhibited elevated hyaluronidase activity ($p = 0.0006$). The $\Delta vp1$
220 strain displayed 1.7-fold lower hyaluronidase compared to the wild-type strain ($p < 0.0001$).
221 This defect was rescued in the $\Delta vp1:vp1$ ($p < 0.0001$ relative to $\Delta vp1$ strain). The ΔPTS -HAL
222 mutant and the $\Delta vp1$ - ΔPTS -HAL double mutant showed a 1.4- and 1.5-fold reduction in
223 hyaluronidase relative to the wild-type strain ($p < 0.0005$). This reduction was similar to the
224 $\Delta vp1$ strain compared to the wild-type strain. The majority of the hyaluronidase activity was
225 captured within the pellets, suggesting that the majority of the degradation enzymes are

226 associated with the cell and not secreted into the supernatant. These data suggest that *vp1*
227 and the PTS-HAL processing and transport operon promote degradation of HA.

228

229 To determine whether the ability to break down HA is conserved beyond the PMEN1
230 strain PN4595-T23, we performed the same set of experiments in a TIGR4 background. The
231 locus in TIGR4 resembles that of the PMEN1 strain, with the addition of a predicted
232 transposon at the end of the operon (see graphic Fig S2A). The results were similar in that the
233 genes encoded by the *vp1* operon (*vpo*) and HA processing and transport operon (PTS-HAL)
234 contributed to the degradation of the HA polymer. In contrast to PN4595-T23, the majority of
235 the TIGR4 activity was localized to the supernatant (Fig S3C), suggesting differences in the
236 subcellular localization of the hyaluronidase activity between strains.

237

238 Finally, to investigate whether VP1 exerts its influence via genes encoded in the HA
239 processing and transport operon, we tested whether overexpression of the PTS-HAL operon in
240 a $\Delta vp1$ background could rescue the wild-type phenotype in both TIGR4 and PN4595-T23
241 strains. To this end, using the $\Delta vp1$ backgrounds, we replaced the native promoter of PTS-
242 HAL with a constitutive promoter (from *amiA* gene) (strain $\Delta vp1$ -OE PTS-HAL)(80). These
243 strains displayed HA degradation comparable to the isogenic wild-type strains ($p = 0.21$) (Fig
244 3B and S2C). These data suggest that *vp1* exerts its effect on HA metabolism via control of the
245 gene products encoded in the PTS-HAL operon.

246

247 **Pneumococcus attaches to host HA in a process controlled by VP1 and mediated by**
248 **molecules involved in HA processing.**

249 We have demonstrated that the *vp1* product enhances attachment of pneumococci to
250 lung epithelial cells (Fig 1), and promotes transcription and activity of genes in the HA
251 processing and transport operon associated with HA metabolism (Figs 2 and 3). Thus, we
252 hypothesized that *vp1* influences attachment via its effect on the PTS-HAL operon. To test this
253 hypothesis, we performed attachment assays in a strain with a deletion in the PTS-HAL operon
254 (Δ PTS-HAL); it displayed decreased attachment relative to the wild-type strain (Fig 4A).
255 Moreover, the attachment defect observed in the Δ *vp1* mutant was rescued to wild-type levels
256 in the Δ *vp1*-OE PTS-HAL strain where the genes in the PTS-HAL operon are expressed from a
257 constitutive promoter. These data strongly suggest that the products of *vp1* and PTS-HAL
258 promote adhesion and that the effect of *vp1* is exerted via its influence on PTS-HAL. Further,
259 attachment levels for the double mutant Δ *vp1 Δ PTS-HAL resembled that of the Δ PTS-HAL
260 strain consistent with these gene products acting within the same pathway (Figs S3A and
261 S3B). The Δ *regR* strain, with increased levels of expression for genes encoded in the PTS-
262 HAL operon, resembled wild-type levels. The Δ *hysA* strain displayed similar attachment to the
263 wild-type strain, suggesting that genes in PTS-HAL are not promoting attachment via indirect
264 effects on HysA (Fig S3A). The attachment of the wild-type and mutant strains were also
265 visualized by confocal microscopy, and representative images are displayed in Fig 4B.
266 Together, these data demonstrate that the *vp1* promotes attachment via its control of the
267 genes in the PTS-HAL operon, responsible for processing and transport of HA.*

268

269 HA production in mammalian cells is controlled by at least three different, membrane-
270 anchored hyaluronan synthases: *HAS1*, *HAS2*, and *HAS3* (35). Previous transcriptomic
271 studies have reported that A549 cells exclusively express *HAS2* and *HAS3*, while *HAS1*
272 remains undetectable (81). Our qPCR analysis corroborates these findings (Fig S3C). To

273 evaluate whether the HA produced by the A549 cells is implicated in pneumococcal
274 attachment, we employed siRNA to knockdown the expression of *HAS2* and *HAS3*
275 (independently and together) on A549 cells, and 48 h later, assessed the attachment strength
276 of the wild-type strain. As a control for treatment, we used a scrambled construct for siRNA
277 experiments, as well as *HAS1* because it has the low expression levels in A549 (Fig S3C and
278 as described by Chow et al., 2010). At the 48h time point, the siRNA treatment reduced levels
279 of *HAS2* and *HAS3* to below 20% when compared to the scrambled control (Fig S3D). Our
280 attachment assays revealed a significant reduction in pneumococcal attachment to host cells
281 where *HAS2* or *HAS3* were knockdown, with maximum reduction in the double knockdown
282 cells (Fig 4C). These data suggest that pneumococcus binding is enhanced in the presence of
283 host HA.

284

285 If products of the *vp1* gene and the PTS-HAL operon exert their influence on attachment
286 via HA, a dramatic decrease in host HA levels should reduce the effect of these pneumococcal
287 molecules on host attachment. Consistently, the $\Delta vp1$ strain displayed similar levels of
288 attachment on host cells independent of the levels of *HAS2* *HAS3*. Further, the Δ PTS-HAL
289 strain also displayed the same levels of attachment in host cells regardless of the levels of
290 *HAS2* and *HAS3* (Fig 4C). Moreover, attachment of the Δ PTS-HAL strain resembled levels of a
291 wild-type strain on cells with reduced *HAS2* and *HAS3*, that is, substantially lower than
292 attachment to wild type cells. Finally, a strain overexpressing the PTS-HAL operon in the $\Delta vp1$
293 background attached better in host cells that express HA. Thus, pneumococcus binding is
294 enhanced by host HA, in a process that depends on the products of *vp1* and PTS-HAL.

295

296 The influence of HA in promoting pneumococcal binding is further corroborated by an
297 experiment with exogenous HA. In one experiment, the addition of exogenous HA diminishes
298 binding to A549, consistent with bacteria binding to detached HA (Fig S3E). In a second
299 experiment, we coated 6-well plates with HA from *S. equis* (Fig S4A). A spot assay revealed
300 that pneumococcal attachment was substantially higher on a surface coated with HA, relative
301 to those without HA (Fig S4B). As observed on cells, the levels of attachment for $\Delta vp1$ or
302 ΔPTS -HAL strains were lower than that of the wild-type strain in HA-coated surfaces, and
303 overexpression of the PTS-HAL operon in the $\Delta vp1$ restored attachment to wild-type levels
304 (Fig 4D). Moreover, these genetic changes had no significant effect on binding on surfaces
305 that were not treated with HA. All together, these data demonstrate that the products of genes
306 involved in the HA processing and transport promote the binding of pneumococcus to host HA
307 in a process regulated by VP1.

308

309 **VP1 and the HA processing and transport operon promote pneumococcal colonization**

310 To establish the role of VP1 and of the PTS-HAL operon in pneumococcal survival in
311 the nasopharynx, we employed the murine model of colonization. Cohorts of five mice were
312 inoculated with single strains, using an inoculation dose of 20 μ l, where the wild-type strain
313 colonizes the nasopharynx, but not the lungs (83,84). The bacterial load was measured two
314 hours and seven days post-inoculation. Enumeration of pneumococci in the nasopharyngeal
315 lavages two hours post-infection did not show any difference in bacterial load among the
316 strains, indicating the accuracy of infection and the consistency in obtaining nasopharyngeal
317 content (Fig 5A). Throughout the course of infection, the number of recovered wild-type
318 pneumococci remained constant (Figs 5A and 5b).

319

320 Samples recovered seven days post-inoculation revealed significant differences in
321 colony counts across the pneumococcal strains (Fig 5B). Our previous work, in the chinchilla
322 model of pneumococcal disease, established that *vp1* is a virulence determinant based on the
323 reduced mortality and decreased dissemination associated with the $\Delta vp1$ strain relative to the
324 wild-type strain (7). Akin to the chinchilla model, the $\Delta vp1$ strain displayed a dramatic decrease
325 in the murine model, where four out of five animals cleared the infection ($p < 0.0001$).
326 Complementation of *vp1* ($\Delta vp1:vp1$) partially rescued the wild-type phenotype. The difference
327 between wild-type and complement ($\Delta vp1:vp1$) is likely due to changes in the dose driven by
328 variation of the promoter. These data suggest that VP1 contributes to infection in multiple
329 tissue types, as well as in carriage and disease.

330

331 By the seventh day post-inoculation, we also detected a significant reduction in colony
332 counts in the ΔPTS -HAL and the double $\Delta vp1$ - ΔPTS -HAL mutants compared to the wild-type
333 strain ($p < 0.0001$). Moreover, overexpression of the PTS-HAL operon in the $\Delta vp1$ background
334 significantly increased the recovered pneumococci compare to $\Delta vp1$ ($p < 0.001$). Yet,
335 overexpression of PTS-HAL in the wild-type background had no effect ($p = 0.542$). The
336 observation that the PTS-HAL operon can compensate for the absence of *vp1* is consistent
337 with the model where VP1 acts upstream of HA processing (Figs 5 and 6). These results
338 strongly suggest that VP1 and HA transport and metabolism play a critical role in
339 pneumococcal colonization of the upper airways.

340 DISCUSSION

341 The binding of pneumococci to epithelial cells of the host mucosal surfaces is a crucial
342 step during colonization and precedes dissemination to other tissues (3,85,86). In this study,
343 we reveal a pneumococcal pathway that promotes adherence via host HA. We previously
344 documented that the signaling peptide *vp1* is dramatically induced during host infection,
345 enhances biofilm formation, and is a potent virulence determinant in a chinchilla model of
346 middle ear infection (7). Here, we build on this work, providing mechanistic details that reveal
347 links across signaling, adherence, and colonization. Specifically, we demonstrate that VP1
348 controls genes involved in the transport and metabolism of the host glycosaminoglycan, HA.
349 Our *in vitro* studies demonstrate that genes in the PTS-HAL operon enhance attachment of the
350 pneumococcus to epithelial cells. Our *in vivo* studies on a murine model of colonization
351 suggest that VP1 and the operon for HA transport and metabolism play critical roles in
352 nasopharyngeal colonization.

353

354 VP1 is part of a regulatory network involved in nutritional sensing. Previous work
355 demonstrated that the master nutritional regulator CodY negatively regulates the level of *vp1*
356 transcripts, and positively regulated by the transcriptional regulator Rgg144 (18,87). The
357 activity of Rgg144 (nomenclature based on gene name in D39, SPD_0144) is regulated by a
358 cognate peptide SHP144 (short hydrophobic peptide 144), which is secreted from cells and
359 internalized into producing and neighboring cells where it binds to Rgg144 (7,18). Rgg144 is a
360 negative regulator of the capsule: it directly binds a predicted promoter upstream of the
361 capsular genes, inhibits the expression of capsular genes, and leads to a reduction in the size
362 of the type 2 capsule (18). The bacterium capsule hinders cell adhesion by concealing
363 bacterial receptors (88). In addition, our work on murine models shows that the Rgg144-

364 SHP144 system serves as a colonization factor in the nasopharynx and contributes to
365 pathogenesis in pneumonia (18). These data are consistent with a model where the Rgg144-
366 SHP144 system plays an essential role in colonization by decreasing the capsule and inducing
367 *vp1*, and that together these events promote adhesion to epithelial cells in the nasopharynx
368 and the lungs. Of note, adhesion experiments in this study were performed in CDM
369 supplemented with glucose. In this condition, levels of *rgg144* are low, and levels of *vp1* are
370 high (7,18). Thus, while Rgg144-SHP144 likely plays a key role in multiple aspects of
371 colonization in our animal model, it is unlikely to be a major contributor *in vitro* experiments
372 presented in this study. We propose that VP1 is a critical component of a complex cellular
373 response to changes in nutritional status.

374

375 Our studies in the murine model of colonization reveal a dramatic defect for the *vp1*
376 deletion mutant, where four out of five mice cleared the infection (Fig 5). The strain with a
377 deletion of the HA transport and metabolism operon also displayed a decrease in bacterial
378 load, but not of the same magnitude as the $\Delta vp1$. These data suggest that: (1) VP1 mediates
379 attachment via control of the HA transport and metabolism operon, and (2) VP1 plays
380 additional roles in colonization, which are mediated by yet unknown factors.

381

382 Many pneumococcal surface molecules serve as adhesins, mediating attachment of the
383 bacteria to host cells (40,89,90). A subset of adhesins binds to extracellular matrix
384 components. Pneumococcal virulence protein A and B (PavA and PavB) (91–93) and enolase
385 (94,95) bind to fibronectin and plasminogen. Similarly, the plasmin- and fibronectin-binding
386 proteins A and B (PfbA and PfbB) bind to the extracellular matrix and are required for
387 adherence to lung and laryngeal epithelial cells (96,97). The choline-binding protein A (CbpA

388 or PspC) binds vitronectin (98), and PsrP binds to keratin (99). Pneumococcus also encodes
389 surface-exposed glycosidases, which have been shown to contribute to adhesion not only via
390 modification of the host surface that reveal receptors but also via direct binding (100). The
391 glycosidase NanA (64) and the β -galactosidase BgaA (101), both process GAGs. NanA
392 releases terminal sialic acid residues and BgaA releases terminal galactose residues from
393 glycoconjugates. However, these enzymes promote attachment to host cells in a manner
394 independent of their enzymatic activity. This observation is particularly evident for BgaA, where
395 binding is mediated via a carbohydrate-binding module (102). Thus, the pneumococcal can
396 employ glycosidases in host attachment. In this study, deletion of the surface exposed HysA
397 strongly influenced HA processing (Fig 3), but not attachment (Fig 4). Thus, we deduce that
398 the glycosidase HysA does not mediate the VP1-regulated defect in adhesion.

399

400 The data presented in this study strongly suggest that the transport and processing of
401 host HA plays an essential role in adhesion. However, we have not yet established the
402 molecular mechanism responsible for this outcome. Levels of regulators, carbohydrates, and
403 transporters are interconnected in the bacterial cells via highly complex networks. Regulators
404 control gene expression, including levels of transport systems and biosynthesis enzymes.
405 Carbohydrates can function as substrates, and can positively or negatively influence the
406 activity of regulators. Transporters not only control the import of carbohydrates, but the extent
407 and position of phosphorylated residues on the cytoplasmic Hpr in the PTS system also
408 controls CcpA-dependent carbon catabolite repression (103). CcpA influences the levels of
409 almost one-fifth of pneumococcal genes, including ABC transporters and regulators (103,104).
410 Thus, many molecular mechanisms may explain the connections between transporters and
411 adhesions.

412

413 The HA transport and metabolism locus manipulated in this study encodes all the PTS-
414 EII components (EIIA-EIIB, EIIC, and EIID) that specifically mediate HA import. The locus also
415 encodes two additional genes for a putative heparinase and YajC, a component of the
416 machinery associated with the secretion of proteins with N-terminal signal sequences (73),
417 Data presented in Fig 4, where host HA levels are decreased using SiRNA knockdowns,
418 strongly suggest that the PTS and VP1 mediate attachment, via a yet uncharacterized effector,
419 in an HA-dependent manner.

420

421 Multiple lines of evidence suggest that GAGs facilitate attachment of multiple bacterial
422 species to host cells. Several groups have explored and advanced the utilization of HA to
423 facilitate attachment of bacteria to abiotic surfaces (49,105,106). A recent report showed that
424 multidrug-resistant *Staphylococcus aureus* and *Pseudomonas aeruginosa* form more robust
425 biofilms on surfaces coated with collagen and HA than on uncoated control surface (42). In
426 addition, levels of GAGs on A549 cells were positively associated with the attachment of
427 several pathogens, including *Staphylococcus aureus*, *Streptococcus pyogenes*, and the
428 pneumococcus (43). Similarly, attachment of multiple bacterial species (both Gram-positive
429 and Gram-negative) was positively associated with GAG levels on corneal epithelial cells HCE-
430 2, well known for expressing a variety of GAGs on their surface (40). In yet another example,
431 HA is the primary attachment site of *M. tuberculosis* in A549 cells (107). While HA has not
432 been directly implicated in the attachment of pneumococcus, previous studies demonstrate
433 that the pneumococcus can utilize HA as a carbon source. In particular, *hysA* and genes in the
434 PTS-EII-encoding operon (EIIB, EIIC, EIID, and *ugtI*), are required for pneumococcal growth in
435 media where HA is the only carbohydrate (56). Furthermore, HA supports *in vitro*

436 pneumococcal biofilm formation (57). These studies suggest that multiple bacteria may utilize
437 host HA for attachment or nutrients and that control of each or both of these processes
438 influences colonization and virulence.

439

440 In this study, we reveal a novel pathway involved in the attachment of the
441 pneumococcus to host epithelial cells. Our working model is that Rgg144 and CodY monitor
442 nutrient availability and, both of these transcription factors influence levels of the secreted
443 peptide *vp1*. VP1, in turn, controls a genomic region involved in the transport and metabolism
444 of HA, and these genes promote adherence in an HA-dependent manner. VP1 is encoded by a
445 core gene, is highly expressed during infection, and is a significant contributor to both
446 colonization and virulence. Thus, this molecule is a high-value candidate for the development
447 of anti-pneumococcal therapies.

448 **MATERIALS AND METHODS**

449 **Bacterial strains and culture conditions.** Wild-type *S. pneumoniae* strain PN4595-T23
450 (GenBank ABX001), was selected as a representative of the PMEN1 lineage (59–62). The
451 clinical isolate TIGR4 (The Institute for Genomic Research, Aaberge et al., 1995), was
452 graciously provided by Dr. Jason Rosch. The construction of the *vp1* deletion mutant ($\Delta vp1$)
453 and the *vp1* complemented strain ($\Delta vp1:\Delta vp1$) in the PN4595-T23 background was reported
454 previously (7). Strain PN4595-T23 was modified to also generate the PTS-HAL operon
455 deletion (Δ PTS-HAL), the *hysA* deletion ($\Delta hysA$), and the *regR* deletion ($\Delta regR$). We generated
456 a double mutant *vp1*-PTS ($\Delta vp1-\Delta$ PTS-HAL) and a PTS-HAL overexpressor in the $\Delta vp1$ and
457 wild-type backgrounds ($\Delta vp1$ -OE PTS-HAL and OE PTS-HAL, respectively) (S5 Table). A
458 similar set of strains was constructed in the TIGR4 background (S5 Table). In this set, we
459 generated a deletion mutant strain of the operon encoding *vp1* (*vpo*) and the *vpo*
460 complemented strain ($\Delta vpo::\Delta vpo$). For growth on solid media, strains were streaked onto
461 Trypticase Soy Agar II plates (TSA) containing 5% sheep blood (BD BBL, New Jersey, USA).
462 For growth in liquid culture, colonies from a frozen stock were grown overnight on TSA plates
463 and inoculated into Columbia broth (Remel Microbiology Products, Thermo Fisher Scientific,
464 USA). The media was supplemented with antibiotics as needed. Cultures were incubated at
465 37°C and 5% CO₂ without shaking. Experiments in chemically defined medium (CDM) were
466 performed utilizing a published recipe (104), and glucose was used at a final concentration of
467 55mM. Growth in CDM was initiated by growing a pre-culture in Columbia broth for 2-3 hours
468 and then diluted to an absorbance of 0.1 at 600 nm, this culture was then grown in CDM-Glu.

469

470 **Strain construction and transformation.** To generate the Δ PTS-HAL mutant strain we used
471 site-directed homologous recombination to replace genes SPN23F_02950 to SPN23F_0310

472 with a kanamycin-resistance gene (*kan*). Likewise, the PTS-HAL operon in the TIGR4 strain
473 (SP_0321 to SP_0327) was replaced with *kan*. The *kan* region was originally amplified from an
474 *rpsL* cassette (80). Gene IDs for PN4595-T23 and TIGR4 are listed in S3 Table. The same
475 strategy was used for the single gene replacements, of *regR* (SPN23F_03020) and *hysA*
476 (SPN23F_02890). To decrease the likelihood of polar effects in all our constructs, the strong
477 artificial transcriptional terminator B1002 was inserted downstream of *kan* (109). Briefly, the
478 transforming DNA was prepared by ligation of the kanamycin gene and promoter, with DNA
479 (one to two kilobases) from the flanking regions upstream and downstream of the PTS-HAL
480 operon. Flanking regions were amplified from the parental strains using either Q5 polymerase
481 or OneTag polymerase (New England Biolabs). The OE PTS-HAL was generated in the $\Delta vp1$
482 and wild-type backgrounds by replacing the native PTS operon promoter with an *amiA*
483 promoter and *kan* gene. Assembly of these transforming fragments was achieved by either
484 sticky-end ligation of restriction enzyme-cut PCR products or by Gibson Assembly using
485 NEBuilder HiFi DNA Assembly Cloning Kit. The resulting constructs were transformed into
486 PN4595-T23 or TIGR4. Primers used to generate the constructs are listed in S4 Table.
487 PN4595-T23 and TIGR4 strains were transformed with approximately 1 μg of DNA. Liquid
488 cultures were supplemented with 125 $\mu\text{g ml}^{-1}$ of CSP1 (EMRLSKFFRDFILQRKK, for TIGR4) or
489 CSP2 (EMRISRIILDFLFLRKK, for PN4595-T23), at an absorbance of 0.05 at 600 nm
490 (GenScript, NJ, USA), and incubated at 37°C. After 2 hours, the treated cultures were plated
491 on Columbia agar containing the appropriate concentration of antibiotic for selection,
492 spectinomycin, 100 $\mu\text{g ml}^{-1}$, kanamycin 150 $\mu\text{g ml}^{-1}$). Resistant colonies were cultured in
493 Columbia broth. Sequences were confirmed using DNA sequencing of the PCR amplicons
494 (Genewiz, Inc., USA). All strains generated in this study are listed in S5 Table.

495

496 **RNA purification, reverse transcription (RT) and qPCR.** Bacterial samples were collected
497 on RNALater (ThermoFisher), and pellets were lysed with 1x lysis mix containing 2 mg ml⁻¹ of
498 proteinase K, 10 mg ml⁻¹ of lysozyme, and 20 µg ml⁻¹ mutanolysin in TE buffer (10 mM Tris·Cl,
499 1 mM EDTA, pH 8.0) for 20 min. Total RNA was isolated using the RNeasy (Zymo Research)
500 following manufacturer instructions. Contaminant DNA was removed by incubating total RNA
501 samples with DNase (2U/µl) at 37°C for at least 15 min and then checked by amplification of
502 *gapdh* (no visible band should be observable in RNA only samples). RT reaction was
503 performed using 1 µg of total RNA using SuperscriptVILO kit for 1 h. Five ng of total cDNA was
504 subjected to real-time PCR using PowerUp SYBR Green Master Mix in the ABI 7300 Real-
505 Time PCR system (Applied Biosystems) according to the manufacturer's instructions. All qRT-
506 PCR amplification was normalized to pneumococcal *gapdh* and expressed as fold change with
507 respect to the wild-type strain. qPCR primers are listed in S4 Table. Primers were obtained
508 from IDT (Integrated DNA Technologies).

509

510 **Spot assays and CFU counts.** Bacteria were inoculated from overnight plates into 20 ml of
511 Columbia broth. Liquid cultures were grown statically and monitored at an optical density of
512 600 nm using Nanodrop 2000c spectrophotometer (Thermo Scientific). Bacterial counts were
513 assessed by streaking 10-fold dilutions on TSA or TSA-antibiotic containing plates. Spot
514 assays were performed with 5 µl of ten-fold dilutions.

515

516 **Microarray Analysis:** We utilized the Pneumococcal Supragenome Hybridization Array
517 (SpSGH) to compare gene expression between the wild-type PN4595-T23 strain and the $\Delta vp1$
518 as described previously (13). Pneumococcal cultures were collected and normalized to an
519 absorbance of 0.3 at 600 nm. RNA extraction, cDNA preparation, and cDNA labeling were

520 performed as described previously (110). Cyber T was used for data analysis (111). Microarray
521 experiments were performed in triplicate. Genes with at least a 3-fold difference between
522 strains and a Bayesian p -value < 0.05 were selected for further analysis. Microarray data for
523 selected genes was confirmed by qPCR.

524

525 **Preparation of HA-coated surfaces.** Hyaluronic acid from *Streptococcus equi* (Sigma-
526 Aldrich) was bound to tissue culture 24-wells plates following a previously described method
527 with modifications (49). Briefly, twenty-four well plates (VWR) were treated with 0.5 ml of
528 concentrated sulfuric acid for 10 min at 60°C, washed extensively with distilled water, then
529 incubated for 2 h at 37°C with 0.5 ml 5 mg ml⁻¹ of hyaluronic acid. Finally, the plates were
530 rinsed with distilled water and allowed to stand for 1h at room temperature. Plates were not
531 dried during the entire procedure. Control plates were treated solely with either sulfuric acid or
532 HA alone. Hyaluronic acid binding was assessed with 0.5 ml of 1% of Alcian Blue 8GX (Sigma-
533 Aldrich) for 10 min at room temperature, then washed extensively with distilled water.

534

535 **Hyaluronidase activity measurement.** Hyaluronidase activity was measured as described
536 previously (76,112). Briefly, pneumococcal cultures were grown in Columbia broth to an
537 absorbance of 0.7 at 600 nm. Then the bacterial pellet and supernatant were obtained by
538 centrifugation. Prior to the reaction, bacterial pellets were washed thoroughly with PBS at room
539 temperature then diluted in 250 μ l of assay buffer while supernatant samples were diluted 1:1
540 in assay buffer. Hyaluronic acid isolated from *Streptococcus equi* (Sigma Aldrich) was utilized
541 at a final concentration of 10 μ g ml⁻¹ in assay buffer (150 mM NaCl, 200 mM sodium acetate,
542 pH 6) for 15 min at 37°C. Stop buffer (2% NaOH, 2.5% cetrimide) was added to stop the
543 reaction, and the absorbance at 400 nm was determined. Commercial hyaluronidase from

544 bovine testes (Sigma-Aldrich) was used as positive control. Assay results were calculated
545 using the standard curve from the positive controls and by comparison to that of a blank
546 comprised of assay buffer solely or without bacterial material.

547

548 **Mammalian cells and media growth conditions.** Epithelial lung human carcinoma A549 and
549 fibroblast NIH-3T3 cells were propagated and maintained at 37°C and supplemented of 5%
550 carbon dioxide at pH 7 in DMEM media contained 1% L-glutamine, 5% fetal bovine serum
551 (FBS) without antibiotics. Chinchilla middle ear CMEE cells were maintained as previously
552 reported (7,71).

553

554 **siRNA knockdown.** All transfection reagents were provided by Sigma-Aldrich. Subconfluent
555 A549 cultures (20% to 30%) were seed in 24-wells plates and transfected with 3 pmol of a
556 mixture containing pre-designed siRNA. Each mixture consisted of multiple siRNA targeting
557 specifically either *HAS1*, *HAS2*, and *HAS3*. Universal control siRNA was used as a negative
558 control. siRNA was transfected in a final volume of 100 µl of OptiMEM containing 3 µl of
559 MISSION siRNA transfection reagent and no antibiotics. Twenty-four hours later the medium
560 was replaced by DMEM supplemented with fetal bovine serum (FBS). Forty-eight hours after
561 transfection, cells were used in adherence assays. To verify the reduction in *HAS* expression,
562 A549 cells were lysed using the RNeasy kit and relative *HAS1*, *HAS2*, *HAS3*, and *ACTINB*
563 expression was verified by qPCR, as described above.

564

565 **Pneumococcal host cell adherence assay.** A549, NIH-3T3 or CMEE were seed in 24-wells
566 plates or on Lab-Tek II Chamber Slides (Thermo Fisher) at a density of 1×10^5 , and

567 maintained without antibiotic until confluence. Before infection, the cell monolayer medium is
568 replaced with fresh medium and then infected with pneumococci resuspended in DMEM
569 supplemented with 1% FBS using a multiplicity of infection (MOI) of 50 bacteria per cell.
570 Infected cells were incubated at 37°C in 5% CO₂ for 1 h. Then the media of post-infected cells
571 was aspirated and washed thoroughly three times with PBS to remove unbound bacteria. The
572 total number of adherent bacteria was released with saponin (1% w/v) for 15 minutes at 37 °C
573 and plated on TSA plates followed by colony formation and enumeration. The effect of
574 exogenous HA on attachment interference was performed through the addition of HA from
575 *Streptococcus equi* at concentrations ranging between 0.01 to 1 mg/ml to the A549
576 monolayers before the addition of the bacteria. Each experiment was repeated at least four
577 times, and results were expressed as mean ± S.E.M.

578

579 **Confocal microscopy.** A549 cells were seeded on Nunc Lab-Tek 4-chamber (VWR) and
580 inoculated with pneumococcal strains with a MOI of 50 as described above. After 1 hour of
581 infection, cells were washed 3-times with PBS and fixed with 4% PFA, washed with PBS and
582 stained with BODIPY FL Vancomycin (Thermo Fisher Scientific), Alexa Fluor 568-Phalloidin
583 (Thermo Fisher Scientific) following manufacturer's instructions. Finally, cells were covered
584 with Vectashield mounting media with DAPI (Vector Laboratories). Confocal microscopy was
585 performed on the stage of Carl Zeiss LSM-880 META FCS, using 488 nm laser line for the
586 fluorescence tag and 561 nm laser line for the fluorescence Alexa tag. Images were analyzed
587 using ZEN Black software (Carl Zeiss Microscopy, Thornwood, NY) and assembled in Fiji
588 (ImageJ. 2.0.0rc48). Laser intensity and gain were kept the same for all stacks and images.

589

590 ***In vivo* studies:** Colonization ability of pneumococcal strains were tested essentially as
591 previously described (83,84). Nine-week-old CD1 mice were raised in the pre-clinical facility of
592 the Leicester University. Mice were anesthetized lightly with 2.5% isoflurane over oxygen (1.5
593 to 2 liter/min), and they were inoculated intranasally with 20 μ l PBS, pH 7.0, containing
594 approximately 5×10^5 CFU/mouse. When inoculating, mice were kept horizontally to prevent the
595 dissemination of inoculum into the lower respiratory tract. A group of mice (n = 5) for each
596 cohort were killed by cervical dislocation at 2 hours and 7 days after infection. After removing
597 the mandible, nasopharyngeal cavity was accessed using a 18G needle through the soft
598 palate, and the content of nasopharynx was collected by injecting 500 μ l PBS, pH 7.0. This
599 process was repeated once more using the same lavage fluid in order to optimize the recovery
600 of bacteria. Undiluted and serially diluted lavage fluid was plated on blood agar base plates
601 and CFU/nasopharynx was calculated. The results were analyzed by two-way ANOVA
602 followed by Tukey's multiple comparisons test.

603

604 **Ethics statement:** Mouse experiments were done under the UK Home Office approved
605 project (permit no: P7B01C07A) and personal (permit no. I66A9D84D) licenses. The protocols
606 used were approved by both the UK Home Office and the University of Leicester ethics
607 committee. We used anesthetic during the procedure to minimize the suffering. Moreover, the
608 animals were kept in individually ventilated cages, had access to water and feed *ad libido*, and
609 their health were checked regularly in accordance with the legal and institutional requirements.

610

611 **Statistical analysis.** Statistical significance was assessed using Student's two-tailed *t*-test for
612 independent samples, one-way ANOVA and Bonferroni's multiple comparisons test for

613 independent samples, and two-way ANOVA and Tukey's multiple comparisons test for animal
614 data. p -values less than 0.05 were considered statistically significant.

615 **Acknowledgments**

616 We are very grateful to Dr. Aaron Mitchell and Dr. Frederick Lanni for their insightful
617 suggestions and guidance during this project. We also thank Dr. Samantha King for productive
618 discussions, particularly in regards to hyaluronic acid. We thank Dr. Haibing Teng from the
619 Molecular Biosensor and Imaging Facility (MBIC) at Carnegie Mellon University for her
620 assistance with confocal imaging. We thank Dr. Phil G. Campbell for providing A549 cells. This
621 work was supported by NIH grant R01 AI139077-01A1 to N.L.H., the Eberly Family Career
622 Development Professorship of Biological Science, an award from the Samuel and Emma
623 Winters Foundation, as well as support from Carnegie Mellon University. The authors declare
624 that there are no conflicts of interest.

625

626 **Author Contributions**

627 RAC contributed to the conception, design, experimentation, analysis and interpretation,
628 and manuscript preparation; EE contributed with design and construction of bacterial clones,
629 as well as data interpretation; OG contributed with *in vivo* experiments and data analysis, HY
630 contributed with *in vivo* experiments, analysis and interpretation, and manuscript preparation;
631 NLH contributed design, analysis and interpretation, and manuscript preparation.

632 **Figure Captions**

633 **Fig 1. *vp1* enhances pneumococcal attachment to epithelial cells.**

634 (A) A549 lung epithelial cells were exposed to the wild-type PMEN1 strain PN4595-T23, and
635 isogenic *vp1* mutant and *vp1* complemented strains ($\Delta vp1$ and the $\Delta vp1:vp1$, respectively) for
636 1h at 37 °C. After washing the unbound bacteria, the remaining cell-bound bacteria were
637 recovered and quantified on TSA plates. Visualization by a spot assay performed with 5 μ l of
638 ten-fold dilutions (top panel). Quantification of the total number of bacteria bound to epithelial
639 cells based on counts from TSA plates (bottom panel). (B) Representative image of wild-type
640 and the $\Delta vp1$ strains bound to A549 cells. Bacterial cells were stained with BODYPI-FL
641 vancomycin (green), actin was visualized with TRITC-phalloidin (red), and bacterial and host
642 DNA were visualized with DAPI (blue). Data on A represents the mean \pm S.E.M of at least six
643 independent experiments. Statistical significance was calculated using unpaired one-way
644 ANOVA analysis with Bonferroni correction, and multiple comparison between samples were
645 performed. * $p < 0.0001$, ns = not significant with $p = 0.080$.

646

647 **Fig 2. The *vp1* product positively regulates the expression of an operon critical to** 648 **processing of hyaluronic acid.**

649 (A) Microarray transcriptome analysis comparing the expression profiles of the wild-type
650 PMEN1 strain PN4595-T23 and *vp1* derivative strain ($\Delta vp1$) on CDM supplemented with
651 glucose. Data was analyzed for statistical significance using CyberT as per Student's T-test.
652 The dashed blue lines represent 4-fold up- and down-regulation, respectively. The red circle
653 highlights genes associated with hyaluronic acid metabolism. (B) Selected genomic regions
654 with 3-fold difference in expression between the wild-type and the $\Delta vp1$ strains. Targets were

655 sorted by their genomic position on the genome. Operons are indicated by brackets. Color
656 code illustrates expression ratio between the $\Delta vp1$ strain relative to wild-type strain. Gene IDs
657 correspond to those in PMEN1 reference strain ATCC700669 (GenBank FM211187), and the
658 predicted functions based on Pfam. Full set presented in S1 Table.

659

660 **Fig 3. The *vp1* product promotes degradation of the hyaluronan polymer via its controls**
661 **of the operon encoding the PTS-EII system.**

662 (A) Schematic of the genomic region implicated in the metabolism of hyaluronic acid. Arrows
663 correspond to predicted open reading frames. Black and white denote predicted operons. Grey
664 corresponds to genes involved in HA metabolism, but not highly controlled by *vp1*. The IDs
665 refer to gene names in the PMEN1 reference strain ATCC700669. (B and C). Measurements
666 of breakdown of hyaluronan and isogenic deletion mutants for *vp1* ($\Delta vp1$), the hyaluronic acid
667 locus PTS operon (Δ PTS-HAL), *vp1* and the PTS operon ($\Delta vp1$ - Δ PTS-HAL), *hysA* (Δhys), and
668 *regR* ($\Delta regR$), and for a *vp1* complemented strains ($\Delta vp1:vp1$), and a PTS overexpressor in
669 the $\Delta vp1$ background ($\Delta vp1$ -OE PTS-HAL). Both the pellets (B) and the supernatants (C) of
670 bacteria grown in liquid cultures were collected and assessed for hyaluronidase activity. Data
671 on B and C represents the mean \pm S.E.M of at least six independent experiments. Statistical
672 significance was calculated using unpaired one-way ANOVA analysis with Bonferroni
673 correction, and multiple comparison between samples were performed. * $p < 0.05$, ** $p <$
674 0.001 , ns = not significant relative to the wild-type strain. Note that on B, the complemented
675 $\Delta vp1:vp1$ rescues the hyaluronidase deficiency of the $\Delta vp1$ mutant ($p < 0.0001$) and the
676 overexpressor $\Delta vp1$ -OE PTS-HAL rescues the hyaluronidase activity of the mutants $\Delta vp1$ ($p <$
677 0.0001). Further, there is no significant difference between $\Delta vp1$ and Δ PTS-HAL, between
678 $\Delta vp1$ and $\Delta vp1$ - Δ PTS-HAL.

679

680 **Fig 4. Pneumococcus attaches to host HA in a process controlled by VP1 and mediated**
681 **by molecules involved in HA processing.**

682 (A) The wild-type PN4595-T23, $\Delta vp1$, ΔPTS -HAL, and $\Delta vp1$ -OE PTS-HAL strains were tested
683 for attachment to epithelial A549 cells. Cell bound bacteria were enumerated by plating on
684 TSA plates. (B) Representative image of strains attached to epithelial A549 cells. Bacterial
685 cells were stained with BODYPI-FL vancomycin (green), actin was visualized with TRITC-
686 phalloidin (red), and DNA was visualized with DAPI (blue). (C) Adherence measurements on
687 A549 cells. *HAS1*-, *HAS2*-, and *HAS3*-knockdown, and the double-knockdown of *HAS2* and
688 *HAS3* (*HAS2-3*) cells were exposed to wild-type and its derivative $\Delta vp1$, ΔPTS -HAL, and $\Delta vp1$ -
689 OE PTS-HAL strains. siRNA treatment was administered 48 h before addition of bacteria, and
690 adherent bacteria were enumerated by plating on TSA plates. Bound bacteria were
691 enumerated by plating cells on TSA plates. (D) The wild-type PN4595-T23 and its derivative
692 $\Delta vp1$, ΔPTS -HAL, and $\Delta vp1$ -OE PTS-HAL strains were assessed for attachment to hyaluronic
693 acid-coated surfaces in FBS-supplemented DMEM media. The total cell-bound bacteria were
694 recovered and quantified by growth on TSA plates. Data on A, C, and D represents the mean \pm
695 S.E.M of at least four independent experiments. Statistical significance was calculated using
696 unpaired one-way ANOVA (A and D) or two-ways ANOVA (C) analysis with Bonferroni
697 correction, and multiple comparison between samples were performed. * $p < 0.05$, **** $p <$
698 0.0001, ns = not significant. Note that the overexpressor $\Delta vp1$ -OE PTS-HAL rescues the
699 attachment deficiency of the mutant $\Delta vp1$ ($p < 0.0001$) in D.

700

701 **Fig 5. HA processing region is required for pneumococcal colonization.**

702 CD1 mice were inoculated intranasally with 20 ml PBS containing approximately 5×10^5
703 CFU/mouse. At 2 hours (day 0, A), and 7-days (B) post inoculation, nasopharyngeal contents
704 were collected and pneumococci were enumerated. Each column represents the mean of data
705 from five mice. Error bars show the standard error of the mean. Significant differences in
706 bacterial counts are seen comparing to the wild-type strain using two-way ANOVA and Tukey's
707 multiple comparisons test. * $p < 0.0005$, ns = not significant. Note that at day 7, the
708 complemented $\Delta vp1:vp1$ rescues the colonization deficiency of the $\Delta vp1$ mutant ($p = 0.003$).
709 Similarly, the overexpressor OE PTS-HAL rescues the colonization deficiency of the mutants
710 $\Delta vp1$ ($p < 0.0001$). The ΔPTS -HAL ($p = 0.0001$) and the double-mutant $\Delta vp1$ - ΔPTS -HAL ($p =$
711 0.0001) both are significantly different from the wild-type strain and the colonization deficiency
712 in the $\Delta vp1$ strain is greater to the ΔPTS strain ($p = 0.0176$).

713

714 **Fig 6. Working model for VP1 regulation and its role in cell-cell communication in**
715 ***Streptococcus pneumoniae*.**

716 The VP1 is processed and secreted into the extracellular milieu. *vp1* is highly upregulated in
717 the presence of host cells and is induced by an upstream transcription factor, Rgg/Shp. In
718 addition, previous work shows that levels of *vp1* respond to nutrients and to the master
719 regulator CodY. VP1 induces changes in the expression of multiple operons, including two
720 implicated in the processing and acquisition of hyaluronic acid. We propose that VP1 promotes
721 attachment via its influence on the products of genes encoded in the PTS-HAL operon.

722

723 **Fig S1. *vp1* enhances pneumococcal attachment to epithelial cells.**

724 (A) A549 cells were inoculated with the wild-type PMEN1 strain PN4595-T23. After washing
725 the unbound bacteria, the total cell-bound bacteria were recovered. For visualization, five
726 microliters of ten-fold dilutions were used for spot assays on TSA plates (left panel). For
727 quantification, the total number of bacteria was determined by enumeration on TSA plates
728 (right panel). (B and C). Attachment assay were performed on CMEE cells (B) and NIH-3T3
729 cells (C) inoculated with the strain wild-type PN4595-T23, the *vp1* mutant and *vp1*
730 overexpressor strain ($\Delta vp1$ and the $\Delta vp1:vp1$, respectively). As in A, cell-bound bacteria were
731 recovered and enumerated on TSA plates (bottom panels). (D) A549 cells were inoculated with
732 the wild-type TIGR4 strain and its derivative *vpo* mutant and *vpo* overexpressor strains (Δvpo
733 and the $\Delta vpo::vpo$, respectively). The total cell-bound bacteria were recovered and
734 enumerated as in A. (E) Representative orthogonal view image of A549 cells inoculated with
735 wild-type PN4595-T23. Note the extracellular *foci* indicating surface binding of the bacteria.
736 Bacterial cells were stained with BODYPI-FL vancomycin (green), actin was visualized with
737 TRITC-phalloidin (red), and bacterial and human DNA was visualized with DAPI (blue). Data
738 on A, B, C, and D represents the mean \pm S.E.M of at least four independent experiments.
739 Statistical significance was calculated using unpaired one-way ANOVA analysis with
740 Bonferroni correction, and multiple comparison between samples were performed. * $p < 0.005$,
741 ** $p < 0.001$, *** $p < 0.0001$, ns = not significant.

742

743 **Fig S2. The *vp1* product promotes degradation of the hyaluronan polymer via its**
744 **controls of the operon encoding the PTS-EII system in TIGR4.**

745 (A) Schematic of the genomic region implicated in the metabolism of hyaluronic acid. Arrows
746 correspond to predicted open reading frames in TIGR4. Black and white denote predicted
747 operons; PTS-HAL has two additional open reading frames at the end of the predicted operon

748 (represented in white). Grey corresponds to genes involved in HA metabolism, but not highly
749 controlled by *vp1*. The IDs refer to gene names in the TIGR4 genome. (B and C)
750 Measurements of breakdown of hyaluronan by strain TIGR4 and isogenic mutants. Both the
751 supernatants (B) and the pellets (C) of bacteria grown in liquid cultures were collected and
752 assessed for hyaluronidase activity relative to the wild-type strain. Data on B and C represents
753 the mean \pm S.E.M of 8 independent experiments. Statistical significance was calculated using
754 unpaired one-way ANOVA analysis with Bonferroni correction, and multiple comparison
755 between samples were performed. * $p < 0.05$, ** $p < 0.05$, *** $p < 0.0001$ ns = not significant
756 relative to the wild-type strain. Note that on C, the $\Delta vpo:vpo$ rescues the hyaluronidase
757 deficiency of the Δvpo mutant ($p < 0.0001$) and the overexpressor Δvpo -OE PTS-HAL rescues
758 the hyaluronidase activity of the mutants Δvpo ($p < 0.0001$). Further, Δvpo -OE PTS-HAL is
759 also significantly different from the double-mutant Δvpo - Δ PTS-HAL ($p < 0.0001$).

760

761 **Fig S3. Pneumococcus attaches to host HA in a process controlled by VP1 and**
762 **mediated by molecules involved in hyaluronic acid processing.**

763 (A) Spot assay to assess attachment to host cells. The wild-type PN4595-T23, and its
764 derivative strains $\Delta vp1$, $\Delta vp1:vp1$, Δ PTS-HAL, $\Delta vp1$ - Δ PTS-HAL, $\Delta vp1$ -OE PTS and $\Delta hysA$,
765 were assessed for attachment to A549 cells using a spot assay. After washing the unbound
766 bacteria, the total cell-bound bacteria were recovered and plated on TSA plates. (B)
767 Quantification of binding, where the total number of the $\Delta vp1$ - Δ PTS, $\Delta regR$ and $\Delta hysA$, bound
768 to epithelial cells was enumerated on TSA plates. (C) Measurement of the expression levels of
769 hyaluronic acid synthases-encoding genes *HAS1*, *HAS2*, and *HAS3* in A549 cells at two time
770 points where levels are compared to treatment with a scrambled control. (D) Measurement of
771 the expression levels of hyaluronic acid synthases-encoding genes *HAS2* and *HAS3* after

772 treatment with specific siRNA. (E) Quantification of bacterial binding to A549 pre-incubated
773 with varying concentration of HA. Data on B to E represents the mean \pm S.E.M of at least four
774 independent experiments. Statistical significance was calculated using unpaired one-way
775 ANOVA (B, D, and E) analysis with Bonferroni correction, and multiple comparison between
776 samples were performed. * $p < 0.005$, ** $p < 0.0001$, ns = not significant.

777

778 **Fig S4. *vp1* mediates attachment to hyaluronic acid of *Streptococcus equis* via genes in**
779 **the PTS-HAL operon.**

780 (A) hyaluronic acid (HA) coating of 6 well plates under listed treatments, visualized by staining
781 with Alcian Blue 8GX. (B) The PMEN1 strain PN4595-T23 (top panel) was assessed for
782 attachment to hyaluronic acid-coated surfaces in FBS-supplemented DMEM media (A549
783 growth media) or Columbia broth (bacterial growth media). The total cell-bound bacteria were
784 recovered and tested for growth on TSA plates.

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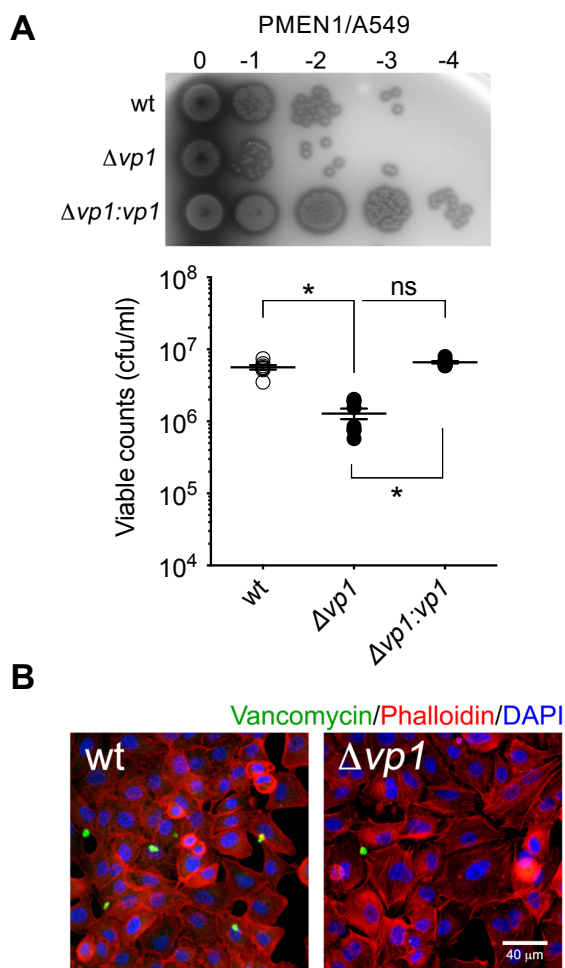


Fig 1. *vp1* enhances pneumococcal attachment to epithelial cells.

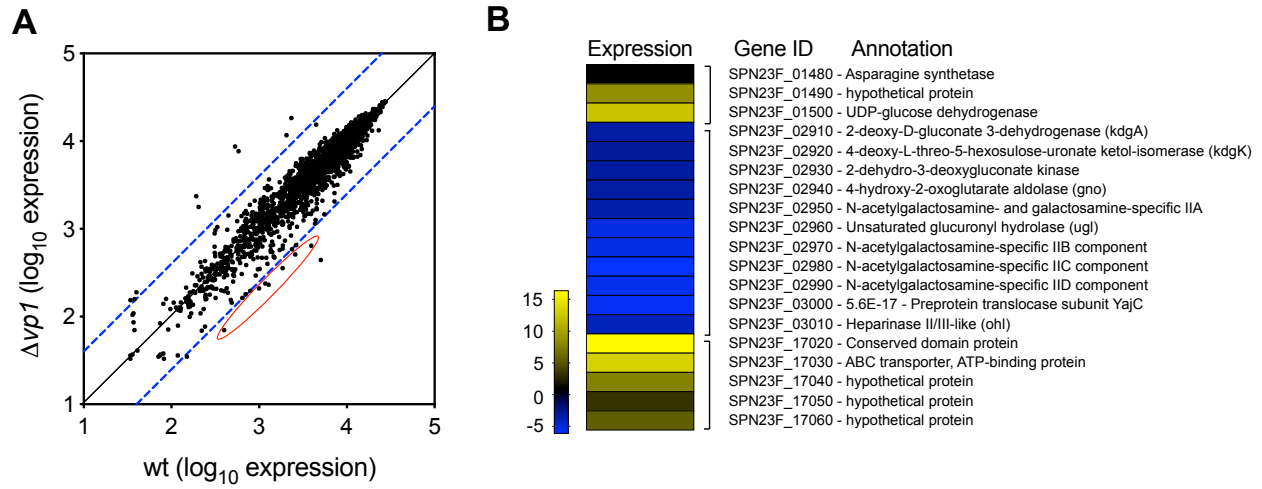


Fig 2. The *vp1* product positively regulates the expression of an operon critical to processing of hyaluronic acid.

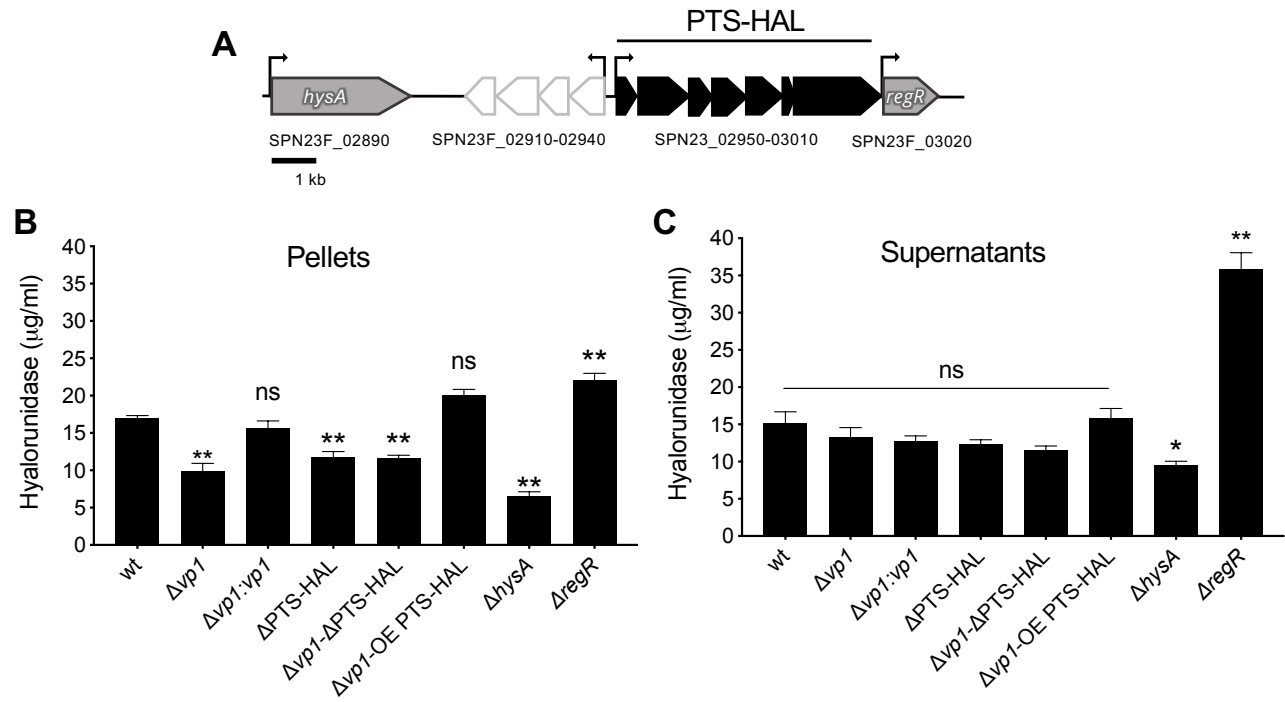


Fig 3. The *vp1* product promotes degradation of the hyaluronan polymer via its controls of the operon encoding the PTS-EII system.

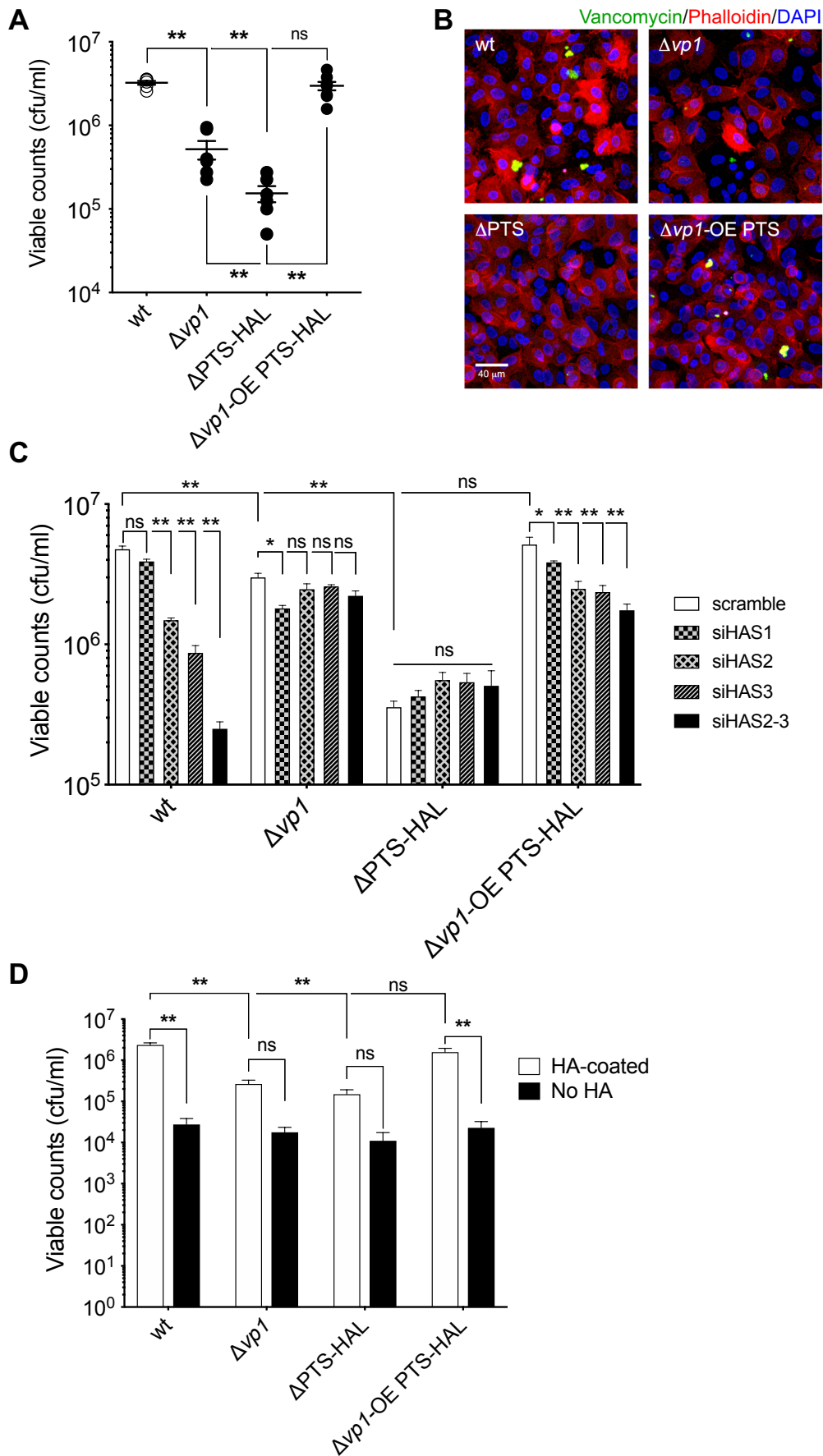


Fig 4. Pneumococcus attaches to host HA in a process controlled by VP1 and mediated by molecules involved in HA processing.

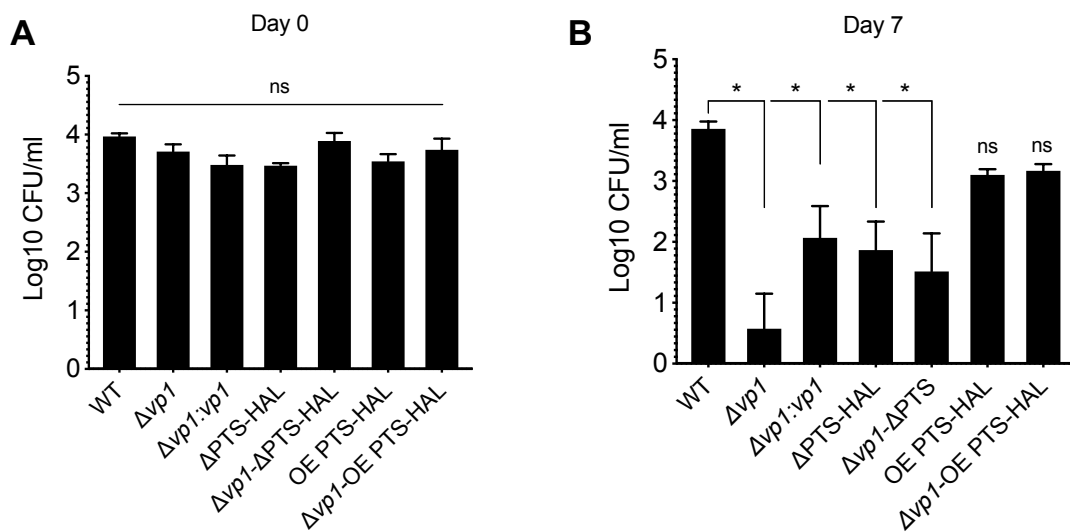


Fig 5. HA processing region is required for pneumococcal colonization.

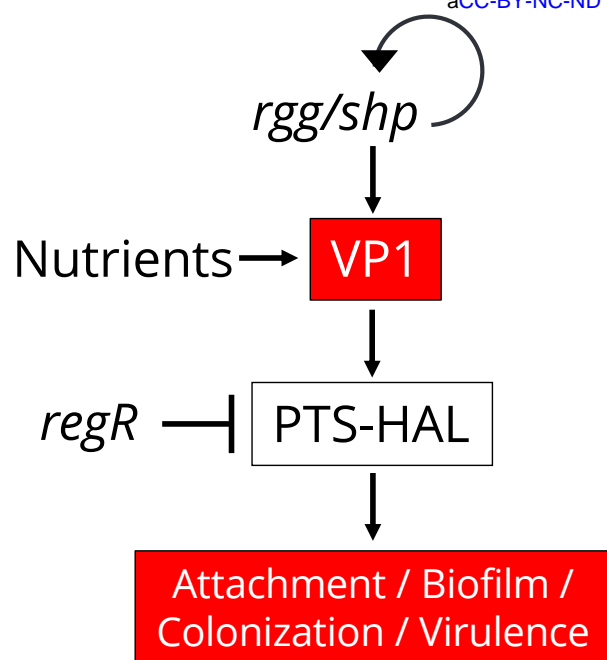


Fig 6. Working model for VP1 regulation and its role in cell-cell communication in *Streptococcus pneumoniae*.

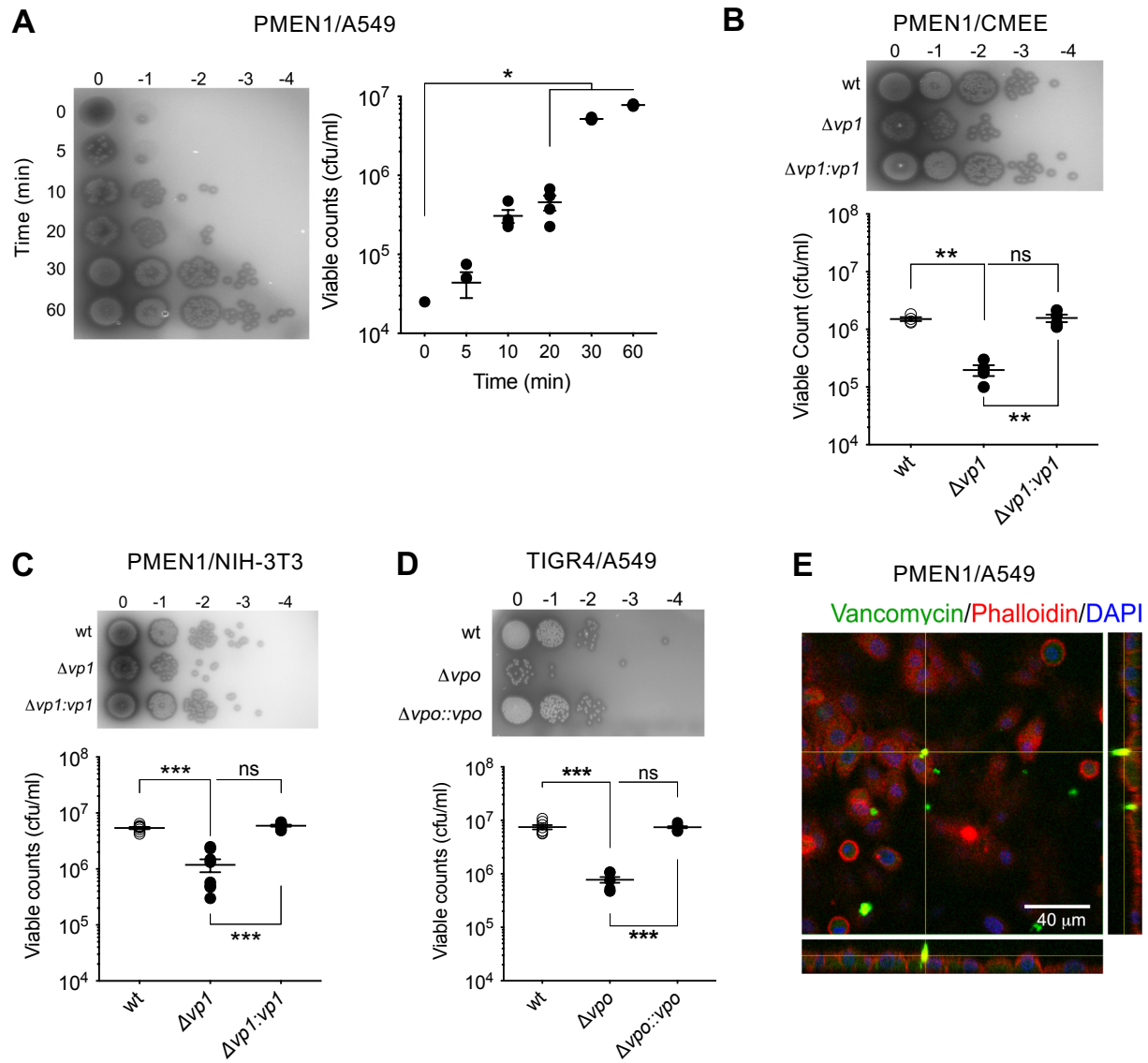


Fig S1. *vp1* enhances pneumococcal attachment to epithelial cells.

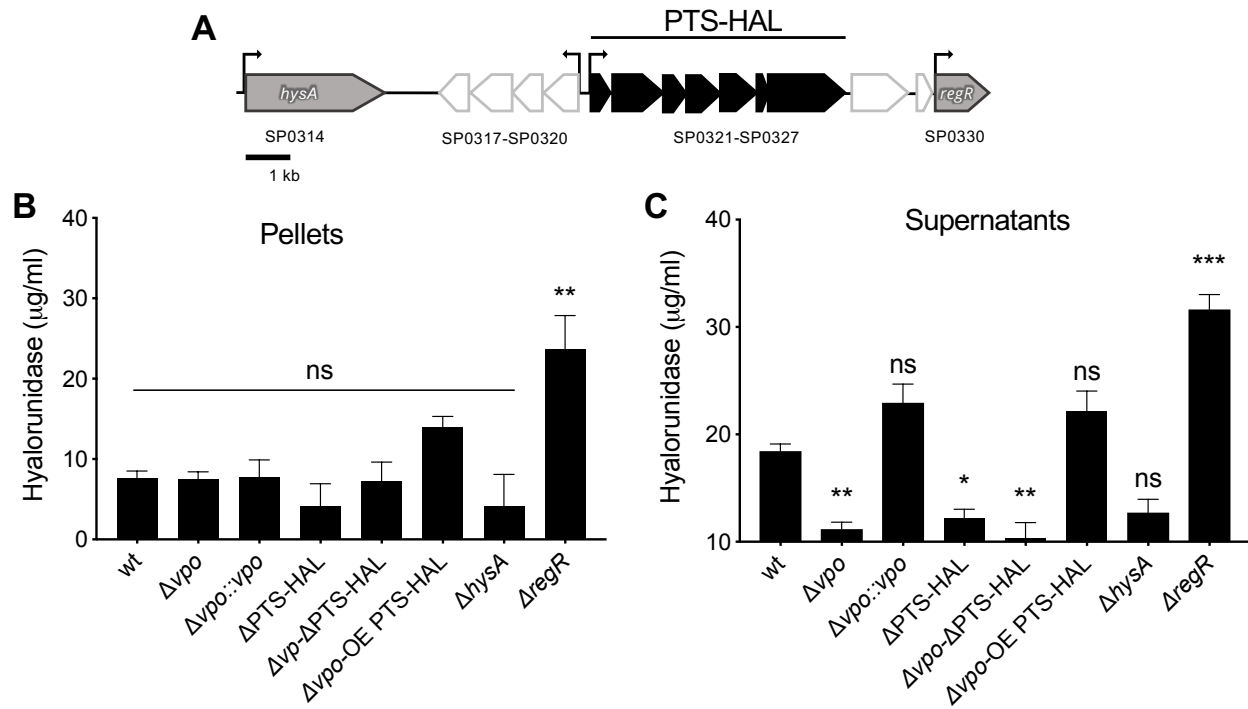


Fig S2. The *vp1* product promotes degradation of the hyaluronan polymer via its controls of the operon encoding the PTS-EII system in TIGR4.

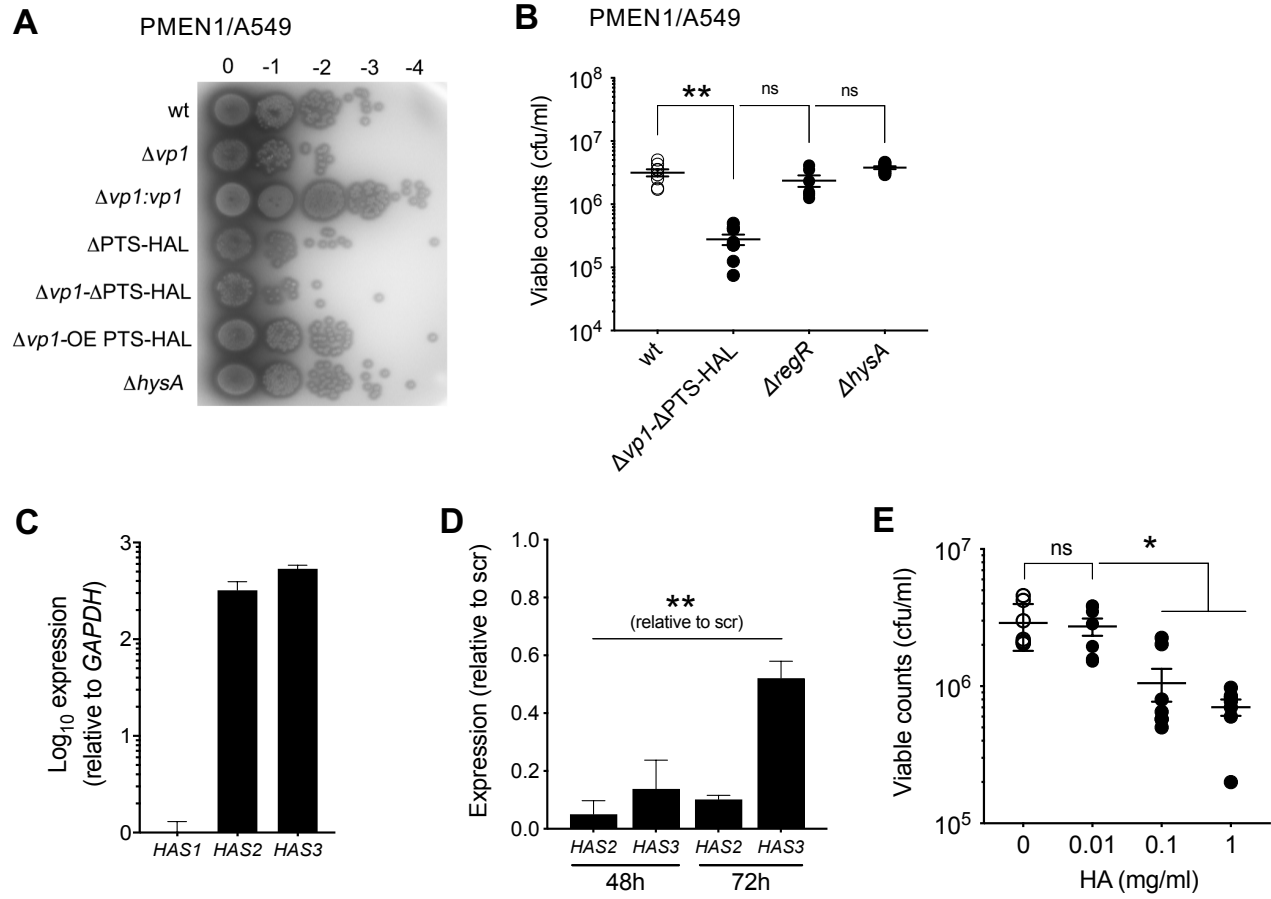


Fig S3. Pneumococcus attaches to host HA in a process controlled by VP1 and mediated by molecules involved in hyaluronic acid processing.

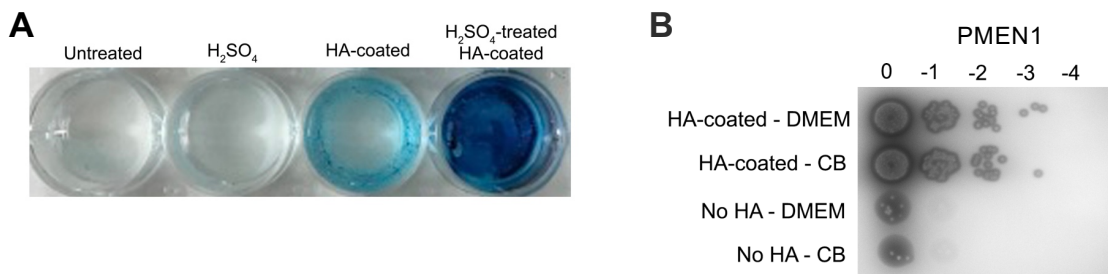


Fig S4. *vp1* mediates attachment to hyaluronic acid of *Streptococcus equis* via genes in the PTS-HAL operon.