Increase in Pilus Islet 2–encoded Pili among Streptococcus pneumoniae Isolates, Atlanta, Georgia, USA

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Journal Title: Emerging Infectious Diseases
Volume: Volume 16, Number 6
Publisher: U.S. National Center for Infectious Diseases | 2010-06, Pages 955-962
Type of Work: Article | Final Publisher PDF
Publisher DOI: 10.3201/eid1606.091820
Permanent URL: http://pid.emory.edu/ark:/25593/ffdbz

Final published version:
http://wwwnc.cdc.gov/eid/article/16/6/09-1820_article.htm

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Accessed April 30, 2019 7:36 PM EDT
To define the prevalence of pilus islet 2 (PI-2)–encoded pili in *Streptococcus pneumoniae* in a geographically defined area, we examined 590 *S. pneumoniae* isolates from population-based surveillance of invasive pneumococcal disease in Atlanta, Georgia, USA, 1994–2006. In 2006, PI-2 was present in 21% of all invasive isolates, including serotypes 1 (100%), 7F (89%), 11A (21%), 19A (40%), and 19F (75%). Only serotype 19F is included in the 7-valent pneumococcal conjugate vaccine that is in use worldwide. In 1999, PI-2-containing isolates were of the same serotypes but accounted for only 3.6% of all invasive isolates. The increase of PI-2 in 2006 resulted predominantly from the emergence of serotype 19A isolates of sequence type 320 and the expansion of serotype 7F isolates. The increase in PI-2-containing isolates and the finding that isolates of all identified serotypes expressed highly conserved PI-2 pili supports their potential as a vaccine candidate.

*Streptococcus pneumoniae* is a major human pathogen, which causes pneumonia, sinusitis, otitis, bacterial meningitis, and septicemia. Introduction of the 7-valent pneumococcal conjugate vaccine (PCV7, Prevnar; Wyeth, Madison, NJ, USA) in 2000 has dramatically decreased invasive pneumococcal disease in the United States in children and in adults (1) and is now in use worldwide. PCV7 includes capsule polysaccharide serotypes 4, 6B, 9V, 14, 18C, 19F, and 23F, serotypes that were responsible for >80% of cases of invasive pneumococcal disease in the United States before the introduction of the conjugate vaccine (1). In recent years other non-PCV7 serotypes, including 7F and 19A, have been emerging in the United States, (2–4).

Pili have recently been identified on several gram-positive bacteria (5–7). To date, 2 different pilus islets (PIs) have been described in *S. pneumoniae* that encode the structural and biosynthetic genes for 2 antigenically different types of pili, PI-1 (8) and recently described PI-2 (9). The so-called *rlrA* pathogenicity islet (10) or PI-1 is a 14-kb genetic region present in 21%–27% of clinical isolates, depending on the geographic region analyzed. It is most prevalent, but not restricted to, the PCV7-vaccine serotypes 4, 6B, 9V, 14, and 19F (11–13).

PI-2 is a 7-kb region located between the genes that encode peptidase T (PepT) and ferrochelatase (HemH) (9). The PI-2 pilus is composed of polymers of the major pilus protein PitB (9).
positive bacteria that contain accessory pilus proteins, that serve as adhesins (15–17) or adaptors for pilus attachment to the cell wall (18, 19), PI-2 pili appear to consist solely of PitB polymers, and these polymers themselves have been shown to mediate adhesion of _S. pneumoniae_ to eukaryotic cells (9).

Bagnoli et al. identified PI-2–containing isolates in a random, worldwide collection of pneumococcal isolates (9). We describe the distribution of PI-2 pili in a well-documented, comprehensive collection from a defined geographic region, i.e., in invasive pneumococcal isolates collected as part of a population-based surveillance program conducted in the metropolitan area of Atlanta, Georgia, USA.

**Materials and Methods**

**Strain Collection**

A set of 590 strains of _S. pneumoniae_ from population-based surveillance of invasive pneumococcal disease isolated between 1994 and 2006 in the 8-county metropolitan Atlanta area, Georgia Health District 3 formed the basis for this study. The isolates were obtained as a part of the Centers for Disease Control and Prevention–sponsored Active Bacterial Core surveillance of the Georgia Emerging Infections Program (1,20). Information about serotype and antimicrobial drug susceptibility was determined for all isolates as described (21). Only viable strains with documented serotype and available antibiogram were included in the study. Laboratory strain _S. pneumoniae_ R6 is a non-encapsulated derivative of the serotype 2 strain D39 (22).

**PCR-based Screening for PI-2**

To ascertain whether PI-2 was absent from the described insertion site (9), we designed primers pepT_F and hemH_R (Table 1) against the flanking genes _pepT_ and _hemH_. To determine whether PI-2 was present, we used primers sipA_up and sipA_dn (Table 1) designed against the PI-2–specific gene _sipA_. We tested for PI-1 using primers Rlr_up_F and Rlr_do_R for its absence and Rlr_SrtC_F and Rlr_srtD_R for its presence, respectively (Table 1).

**Multilocus Sequence Typing and Clonal Complexes**

Multilocus sequence typing (MLST) was performed as described (23). CCs were assigned by using the eBURST program (24) to partition the MLST database (http://spneumoniae.mlst.net). The predicted founder of a CC was assigned by eBURST to the ST with the highest number of single-locus variants.

**Construction of a pitB-Deletion Mutant**

The 3′- and 5′-flanking regions of _pitB_ were amplified by using primer pairs sipA_up_F/PilA_del_3 and PilA_del_2/SrtC_do_R (Table 1). Both PCR products were used in a PCR-based overlap-extension reaction, which introduced an _MfeI_ restriction site between the 2 fused fragments. The PCR product was cloned into pCR-Blunt II (Invitrogen, Carlsbad, CA, USA), resulting in pCR-pitB. The _erm_-cassette of pSK-erm (25) was amplified by using primers _erm_F_ and _erm_R_ (Table 1) and ligated into _MfeI_-digested pCR-pitB. The resulting plasmid, pCR-pitB:erm, was used to transform GA41070 according to established protocols (26), which resulted in mutant GA41070ΔpitB.

**Production of Recombinant PitB and Antiserum**

The region of _pitB_ that encoded the mature protein (bp 126–bp 1128) was amplified from chromosomal DNA of serotype 1 strain GA19686 and cloned into the _Escherichia coli_ expression vector pET-32a+ (Stratagene, La Jolla, CA, USA). The thioredoxin/6xHis-PitB fusion protein was purified by using the B-PER 6xHis Fusion Purification Kit (Pierce, Rockford, IL, USA). The N-terminal thioredoxin/6xhis-tag was removed by cleavage with recombinant enterokinase (Novagen, Madison, WI, USA).

<table>
<thead>
<tr>
<th>Region</th>
<th>Primer</th>
<th>Sequence (5′ → 3′)†‡</th>
</tr>
</thead>
<tbody>
<tr>
<td>Erm cassette</td>
<td><em>erm_F</em></td>
<td>gctctagaCGTTAGATTAATTCTACCAGTGAC</td>
</tr>
<tr>
<td></td>
<td><em>erm_R</em></td>
<td>gctctagaCTCCATTCCTCCCTTTATGAAGGTGTAAC</td>
</tr>
<tr>
<td>PI-2</td>
<td><em>pepT_F</em></td>
<td>TAAGAACGGGTCCAAGAGATTTGG</td>
</tr>
<tr>
<td></td>
<td><em>hemH_R</em></td>
<td>AATAATGGGCTCCAAAATACAAAGC</td>
</tr>
<tr>
<td></td>
<td><em>sipA_up_F</em></td>
<td>CTCTAGGAGGGATCTTCTTTATCATC</td>
</tr>
<tr>
<td></td>
<td><em>sipA_do_R</em></td>
<td>CTACAGCCGTTGGTTGATGTGTC</td>
</tr>
<tr>
<td></td>
<td>PilA_del_3</td>
<td>gcaattgcgggctgtagCTGTATAGGGATGTTCTCCAAAG</td>
</tr>
<tr>
<td></td>
<td>PilA_del_2</td>
<td>ctagccggcgcaattgGCTTGGGGGCAGATGTG</td>
</tr>
<tr>
<td></td>
<td>Rlr_up_F_</td>
<td>CTCCACGGAATCTCTTCAATGG</td>
</tr>
<tr>
<td>PI-1</td>
<td>Rlr_do_R_</td>
<td>GTCTTAGATATCATGTTGTATCAGTGAC</td>
</tr>
<tr>
<td></td>
<td>Rlr_SrtC_F</td>
<td>GGGGAAGATTATGGCGACCTT</td>
</tr>
<tr>
<td></td>
<td>Rlr_srtD_R_</td>
<td>GCTTG GCCCTGACGGTGCC</td>
</tr>
</tbody>
</table>

*PI, pilus islet.
†Not included are primers used for sequencing and multilocus sequence typing.
‡Lowercase letters represent bases added for mutagenesis; uppercase letters represent bases complementary to _S. pneumoniae_.

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PI-2 Pili of Streptococcus pneumoniae

Preparation of Cell Wall Extracts from S. pneumoniae

Cultures were grown to mid-exponential phase (optical density, 0.5–0.6) in Todd-Hewitt medium containing 0.5% yeast extract. After centrifugation, the cell pellets were resuspended in 1/20 of the culture volume in extraction buffer (50 mmol/L Tris-HCl; pH 8.0) containing 200 U/mL mutanolysin (Sigma-Aldrich, St. Louis, MO, USA) and 30% raffinose. After 2 h of incubation at 30°C, cell wall extracts were collected by centrifugation at 12,000 × g for 15 min; 20 μL cell wall extract were mixed with sodium dodecyl sulfate–polyacrylamide gel electrophoresis sample buffer and boiled for 10 min immediately before separation on a 4%–12% Tris-HCl polyacrylamide gel (NuPAGE; Invitrogen). For immunostaining, proteins were transferred to a nitrocellulose membrane and detected with polyclonal anti-PitB antiserum at a dilution of 1:20,000.

Statistical Analysis

Proportions were compared by using the Fisher exact test. p values <0.005 were considered significant.

Results

PI-2 in Strains of Serotypes 1, 7F, 11A, 19A, and 19F

To determine the post PCV-7 prevalence of PI-2 in S. pneumoniae, we analyzed all 381 viable and documented invasive isolates collected in 2006 in metropolitan Atlanta (381/409) in a PCR-based screen. Two PCRs were performed with each strain; 1 resulted in a PCR product in strains that lacked PI-2, and a second resulted in a product only in strains that contained PI-2. Strains that produced a PCR product for PI-2 were further characterized by amplification of pitB, which encoded the pilus backbone protein. For all strains, 1 of the 2 PCRs resulted in a PCR product. No strain gave a PCR product for both PCRs, which could have indicated the presence of the PI in an alternative integration site. These results strongly suggest that the pepT-hemH intergenic region is the only integration site of PI-2 in S. pneumoniae. Overall, 21% of all viable isolates collected in 2006 contained PI-2. The PI-2 of S. pneumoniae strain GA47784 was assigned the GenBank accession no. GU256423.

PI-2 was identified only in isolates of serotypes 1, 7F, 11A, 19A, and 19F (Figure 1). Serotypes 19A and 7F were the most frequent PI-2–containing serotypes isolated in 2006, with a PI-2 prevalence of 40% and 89%, respectively. In serotype 11A, PI-2 was identified sporadically, whereas for serotype 19F, 75% of the isolates contained PI-2. Serotype 1 was represented by a single isolate in 2006. To determine the prevalence of PI-2 in serotype 1 isolates, we analyzed an additional 20 invasive isolates, collected in Atlanta from 1994 through 2005; all contained PI-2. Serotype 1 had the highest presence of PI-2 of all analyzed serotypes. Except for serotype 19F, the PI-2–containing serotypes are not included in PCV7.

To further characterize PI-2 of a serotype 11A strain, we sequenced the 7-kb region between the flanking genes pepT and hemH of strain GA47784. The sequence was 99.9% identical to the published PI-2 sequences, with the highest sequence identity to serotype 1 strain PN110, with only 1-bp difference over the entire PI-2 region. The mutation was located downstream of the stop-codon in pitA, which results in premature termination of PitA, as has been reported (9).

Effects of PCV7 Introduction on PI-2 Distribution

To assess whether PI-2 was associated with the emergence of serotypes in the aftermath of PCV7 introduction (2000), the PI-2 distribution was determined in invasive isolates from 1999 that belonged to the PI-2–containing serotypes (Figure 2). The numbers of isolates for the different serotypes in the 2 periods reflected the expansion of isolates of the nonvaccine serotypes 7F and 19A and the decline in 19F isolates (Figure 2) that has been described in
the United States since the introduction of PCV7 (4). PI-2 was present in isolates of all 5 serotypes before the introduction of PCV7; hence, PI-2 was not recently acquired. For serotype 7F, the ratio between PI-2–containing and PI-2–lacking isolates in 2006 compared to the ratio in 1999 remained essentially the same (Figure 2), which suggests that PI-2 pili may not provide a major selective advantage to invasive 7F isolates. Serotype 19A showed a significant increase in PI-2–containing isolates. To assess the presence of PI-2 in other serotypes in 1999, we analyzed isolates of all represented serotypes of PI-2: all isolates of infrequent serotypes (1, 5, 7C, 7F, 8, 9A, 9N, 10A, 15B, 15C, 16, 16F, 18F, 20, 23A, 24B, 31, 33F, 34, 35B, 35F, 37) and at least 10% of the isolates of serotypes with >10 isolates (3, 4, 6A, 6B, 9V, 11A, 12F, 14, 18C, 19A, 19F, 22F, 23F). No additional PI-2–containing isolates were detected, which suggests that serotypes 1, 7F, 11A, 19A, and 19F were the only serotypes that contained substantial numbers of PI-2–containing isolates. These PI-2–containing isolates accounted for ≈3.6% of all viable isolates collected in 1999. The estimation of total viable isolates in 1999 was based on the known proportion of viable isolates to total collected isolates of the PI-2–containing serotypes of 1999 (93%).

**Genetic Relationships Between PI-2–containing Strains**

To analyze the clonality of the PI-2–containing isolates, we performed MLST for all PI-2–containing isolates from 1999 and 2006 (Table 2). For serotype 1, the isolates from 1999 and 2006 belonged to ST304 and ST227, respectively (Table 2). Both STs have been described as the most prevalent serotype 1 STs in the United States (27,28). PI-2–containing serotype 7F isolates of 2006 belonged to either ST191 or ST1176, and ST191 isolates were responsible for the strong expansion of this serotype in 2006. 7F isolates lacking PI-2 were tested and found to be of ST191 (data not shown), which indicated that PI-2 is present in only a subpopulation of this ST. ST1176 is unrelated to ST191, and 7F isolates of this ST were not present in 1999. All PI-2–containing isolates of serotype 11A belonged to ST62. Analysis of additional serotype 11A isolates that lacked PI-2 showed that they all belonged to ST62 (data not shown). This finding suggests that the structure of this serotype is clonally homogeneous and that PI-2 is present in a subset of this ST.

For serotype 19A, 27 of the 39 PI-2–containing 19A isolates in 2006 were ST320 (Table 2). This ST was absent in 1999 (29); thus these isolates were the major contributor to the increase of PI-2–containing isolates of serotype 19A in 2006. Additionally, PI-2 was identified in 19A isolates of ST1339 and ST2268, and both sequence types belonged to CC2090. One ST1339 isolate was present in 1999, which indicated that PI-2 was present in this ST before PCV7 was introduced. 19A isolates of 2006 that lacked PI-2 were tested and found to belong to other STs, e.g., ST199 and ST695, STs consistent with the described population structure of 19A isolates in the United States (4).

**Table 2. Clonal distribution of PI-2-containing Streptococcus pneumoniae isolates, Atlanta, Georgia, USA, 1994–2006**

<table>
<thead>
<tr>
<th>Serotype</th>
<th>CC</th>
<th>ST</th>
<th>1999</th>
<th>2006</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>304</td>
<td>304</td>
<td>1</td>
<td>1</td>
</tr>
<tr>
<td>1</td>
<td>306</td>
<td>227(DLV)</td>
<td>3</td>
<td>–</td>
</tr>
<tr>
<td>7F</td>
<td>191</td>
<td>191</td>
<td>3</td>
<td>28</td>
</tr>
<tr>
<td>7F</td>
<td>218</td>
<td>1176(DLV)</td>
<td>–</td>
<td>3</td>
</tr>
<tr>
<td>11A</td>
<td>53</td>
<td>62</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>19A</td>
<td>2090</td>
<td>1339(SLV)</td>
<td>1</td>
<td>9</td>
</tr>
<tr>
<td>19A</td>
<td>2268(DLV)</td>
<td>–</td>
<td>3</td>
<td></td>
</tr>
<tr>
<td>19A</td>
<td>271</td>
<td>320(SLV)</td>
<td>–</td>
<td>27</td>
</tr>
<tr>
<td>19F</td>
<td>271</td>
<td>271</td>
<td>1</td>
<td>–</td>
</tr>
<tr>
<td>19F</td>
<td>236(SLV)</td>
<td>3</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>19F</td>
<td>3039(SLV)</td>
<td>–</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>19F</td>
<td>651(DLV)</td>
<td>3</td>
<td>2</td>
<td></td>
</tr>
<tr>
<td>19F</td>
<td>654(SLV)</td>
<td>4</td>
<td>1</td>
<td></td>
</tr>
<tr>
<td>19F</td>
<td>1258(DLV)</td>
<td>2</td>
<td>–</td>
<td></td>
</tr>
</tbody>
</table>

*CC, clonal complex; ST, sequence type; PI, pilus islet; DLV, double-locus variant; SLV, single-locus variant.*
population confirmed PI-1 in isolates of CC271 (ST320, ST271, ST3039, ST236, ST651). Additionally, PCR results showed that PI-1 was in ST1339 isolates but not in isolates of the single locus variant ST2268. Isolates of all other PI-2–containing serotypes or STs did not contain PI-1.

**Antimicrobial Drug Resistance and PI-2–containing Isolates**

19F and 19A isolates of CC271 are closely related to the globally distributed multidrug-resistant clone Taiwan19F-14, and all were resistant to penicillin, erythromycin, cotrimoxazole, tetracycline, and chloramphenicol. In addition, serotype 19A isolates of CC2090 (ST1339, ST2268) were resistant to penicillin, erythromycin, and cotrimoxazole. All PI-2–containing 19A isolates in 2006 were multidrug resistant. In contrast, PI-2–containing isolates of serotypes 1, 7F, 11A, and 19F (ST654, ST251) were susceptible to all tested antimicrobial drugs. Hence, antimicrobial drug resistance was not a consistent feature of PI-2–containing isolates.

**Expression of High Molecular Weight PitB on the Surface of PI-2–containing Isolates**

The first report of PI-2 pili described mutations in several genes encoded by PI-2 (namely, pitA and srtG2), which resulted in their inactivation (9). To confirm that PI-2–containing isolates indeed express PitB polymers on their surface, we produced an anti-PitB antiserum to detect PitB polymers. The specificity of the PitB antiserum was tested against cell wall extracts of strain GA41070 and its isogenic pitB deletion mutant, GA41070ΔpitB (Figure 3). In the wild type, the antibody reacted with several bands that produced the high molecular weight (HMW) banding pattern characteristic of gram-positive pili (30, 31) (Figure 3). As expected, the PitB monomer and the HMW banding pattern were absent in the pitB-deletion mutant (Figure 3).

To analyze whether the PI-2–containing isolates of the different STs produced PitB polymers, we used Western blots to examine their cell wall extracts (Figure 4). All PI-2–containing strains showed the same HMW banding pattern (Figure 4, lanes 2–15). In addition, all 81 PI-2–containing isolates from 2006 were analyzed by whole cell dot blots that confirmed that all isolates expressed PitB on their surface (data not shown).

**High Conservation of PitB among Different STs**

To determine the degree of variation in PitB, we sequenced pitB in isolates representing the different STs (Table 2). The pitB sequences were 100% identical for all strains, except the 7F isolates GA47077 (ST191) and GA47340 (ST1176), which have a single point mutation that has been observed in other serotype 7F strains (9). The high conservation of PitB was consistent with the reported PitB sequences by Bagnoli et al. (9) and is an essential feature of PI-2 as a potential vaccine candidate. Additionally, we analyzed the gene for the second surface protein encoded in PI-2, pitA, and confirmed the presence of the described stop-codon in all strains representing the different STs (Table 2).

**Discussion**

PI-1 pili on *S. pneumoniae* have been reported as promising vaccine candidates for this important human pathogen (13,32). One limitation to this approach is that PI-1 has been found in only ≈25% of strains; it is most frequently found in pneumococcal isolates of serotypes covered by PCV7, which are decreasing in countries that have introduced the vaccine. The recent identification of PI-2 in serotypes not covered by PCV7 has raised the possibility that broad coverage of strains can be achieved by a bivalent pilus-based vaccine.

The first report on PI-2 pili described the serotypes and sequence types associated with PI-2 (9). In the current study, we defined the prevalence of PI-2–expressing isolates in a defined geographic region and population and assessed whether the introduction of PCV7 had an effect on the distribution of PI-2–containing isolates. Twenty-one
percent of the 381 viable invasive pneumococcal isolates collected in 2006 in metropolitan Atlanta expressed PI-2. This percentage is higher than the PI-2 incidence reported from a global strain collection (9) and is similar to the reported percentage for PI-1 (11–13). The higher incidence rate of PI-2 in our study refers to a post-PCV7-era population, whereas the earlier report was based on a random strain collection, including pre- and post-PCV7 isolates. Therefore, our higher results are consistent with the assessment of Bagnoli et al. (9) that PI-2 is most prevalent in emerging, non-PCV7 serotypes.

Notably, all serotype 1 strains had PI-2. This serotype does not play a large role in pneumococcal invasive disease in the United States (27), a conclusion reflected by the low number of serotype 1 isolates in our dataset. However, serotype 1 isolates have been identified in several studies as particularly invasive, in contrast to other serotypes that are more frequently associated with carriage (33–35) and associated with epidemic outbreaks (27,36,37). Further study of PI-2 prevalence in serotype 1 requires a geographically different dataset with a higher prevalence of PI-2-containing isolates (serotypes 1, 7F, 19A) are included in the 13-valent pneumococcal conjugate vaccine now being introduced in Europe, Canada, and the United States. Use of this vaccine is expected to greatly reduce the frequency of all included serotypes and may thus reduce the proportion of S. pneumoniae isolates that contain PI-2. In summary, the prevalence of PI-2 increased from ≈3.6% of the invasive pneumococcal isolates in 1999 to 21% in 2006 in metropolitan Atlanta, Georgia, especially in the emerging serotypes 7F and 19A. PI-2-containing isolates of all identified sequence types expressed polymers of the highly conserved pilus protein PitB on their surface. These findings support the potential of PI-2 pili as a vaccine candidate.

**Acknowledgments**

We are grateful to Monica M. Farley, Wendy Baughman, Sarah W. Satola, and the Georgia Emerging Infections Program.
for providing the clinical isolates and the corresponding documentation. We thank Leslie McGee and Scott T. Chaneey for sequence type data of serotype 19A isolates.

This work was supported by funds from National Institutes of Health grant #R01AI 070829 (D.S.S.) and a Veterans Administration Merit Award from the Department of Veterans Affairs (D.S.S.).

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References


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