Genotype replacement within serotype 23F *Streptococcus pneumoniae* in Beijing, China: characterization of serotype 23F


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**SUMMARY**

We investigated the genetic structure of 99 isolates of serotype 23F *Streptococcus pneumoniae* from children with acute respiratory infections collected over two periods from 1997 to 2006, and 2010. All isolates were susceptible to vancomycin and amoxicillin–clavulanic acid; 97 were resistant to erythromycin, 95 of which carried the *ermB* gene and two carried both *mefA/E* and *ermB* genes. Multidrug resistance to three or more classes of antibiotics was exhibited by 90 isolates. Sequence types ST342 and ST81 were the most frequent in 1997–2006 and 2010, respectively. All CC81 isolates were non-susceptible to β-lactam antibiotics and had higher minimum inhibitory concentration values for penicillin than other clone complexes and sequence types. The increased β-lactam antibiotic resistance may have resulted from the replacement of multidrug-resistant clones related to ST81. Long-term studies on *S. pneumoniae* serotype 23F, especially the ST81 clone, should be conducted to better understand the epidemiological picture of this pathogen in China.

**Key words**: Multidrug resistance, serotype 23F, *Streptococcus pneumoniae*.

**INTRODUCTION**

*Streptococcus pneumoniae* is the leading cause of bacterial infection in infants and young children. In 2005, the World Health Organization (WHO) estimated that 1·6 million people die of pneumococcal disease annually, including 0·7–1 million children (aged < 5 years) mostly from developing countries [1]. This situation has been worsened by the rapid spread of antimicrobial-resistant *S. pneumoniae* and is now a global problem.

To date, more than 93 serotypes have been identified in *S. pneumoniae*. Before the first pneumococcal conjugate vaccine was introduced, the antibiotic-resistant serotype 23F was one of the most common types worldwide [2, 3]. Spain23F-1 clone or sequence type (ST) 81 first emerged in 1970 and became the first antibiotic-resistant *S. pneumoniae* clone whose standard nomenclature was given by the Pneumococcal Molecular Epidemiology Network (PMEN) (http://www.sph.emory.edu/PMEN/index.html). This clone was one of the first pandemic penicillin (PEN)-resistant isolates from Spain in the 1980s [4] and in the late 1990s, it constituted about 40% of PEN-resistant isolates in the USA [5]. Previous studies have reported this clone to be resistant to tetracycline (TCY) and
S. pneumoniae serotype 23F in China

METHODS

Clinical isolates

The study protocol was in accordance with the ethical standards of the responsible regional committee on human experimentation and the Helsinki Declaration of 1975 (revised 1983); it was approved by the ethics committee of the Beijing Children’s Hospital. Nasopharyngeal swabs were taken with written informed consent from the parents or legal guardians of the children. A total of 1132 isolates were recovered in Beijing [10, 11] and almost all isolates collected in 2010 were ST81 [12]. However, whether ST81 was predominant before 2010 is unknown. In the present study, we analysed serotype 23F S. pneumoniae isolates collected in Beijing from 1997 to 2006 and 2010 and report our findings on their antibiotic susceptibility, genotype distribution, and temporal trends.

Antimicrobial susceptibility testing

The minimum inhibitory concentrations (MICs) for PEN, amoxicillin–clavulanic acid (AMC), ceftriaxone (CRO), cefuroxime (CXM), erythromycin (ERY), and vancomycin (VAN) were determined using E-test strips (AB Biodisk, Sweden) [13], and for TCY, sulfamethoxazole–trimethoprim (SXT), and CHL by the disk diffusion method. Breakpoints were determined in accordance with the Clinical and Laboratory Standards Institute 2010 criteria [14] using S. pneumoniae ATCC49619 as the quality control. Isolates were considered multidrug resistant (MDR) if they were not susceptible to three or more classes of antimicrobials.

Detection of macrolide resistance genes

The ermB and mefA/E resistance genes were amplified by polymerase chain reaction (PCR) for all ERY non-susceptible strains using the primers and PCR conditions as previously described [15]. Each PCR reaction contained 500 ng template DNA, 50 mm KCl, 10 mm Tris–HCl (pH 8.3), 200 μM of each deoxynucleotide triphosphate, 2-5 U Taq DNA polymerase (Dalian, Takara Bio, China), 1-5 mm MgCl2, and 1-5 μM of each primer. PCR products were visualized by 1-5% agarose gel electrophoresis and gold-view staining.

Multilocus sequence typing (MLST)

Internal fragments ~450 bp long from the aroE, gdh, gki, recP, spi, xpt, and ddx genes were amplified by PCR as previously described [16]. Alleles that were not present in the pneumococcal MLST database were verified by resequencing the gene fragments on both strands. These alleles were submitted to the MLST S. pneumoniae database for designation. The eBURST algorithm (http://eburst.mlst.net) was used to estimate the relationships among isolates. Various STs were subdivided into one group as a clonal complex (CC) only if they shared six identical alleles of the seven MLST loci with another ST in the group.

Pulsed-field gel electrophoresis (PFGE)

Pneumococci from overnight cultures were suspended in low-melting-point agarose at a concentration of 1-2 in a Dade turbidity meter to form plugs. The bacteria were incubated in 500 mm EDTA/1% N-lauroylsarcosine overnight at 55 °C and DNA was digested with Smal (New England Biolabs, USA) at 30 °C overnight according to the protocol described by Lefevre et al. [17]. PFGE was performed in 1-1% agarose (SeaKem GTG agarose, Switzerland) and 0-5× Tris-borate-EDTA buffer using an electrophoresis system (CHEF-DR III, Bio-Rad Laboratories, USA) at 6 V/cm (3-40 s switch time) for 23 h. Gels were stained with ethidium bromide for 30 min and visualized under UV light. The image was captured using the Gel Doc system (Cell-Bio Sciences, USA), and saved dendrograms of unweighted-pair group method with arithmetic means were constructed using Bionumerics software (Applied Maths, Belgium). PFGE patterns were clustered for isolates with ≥80% genetic relatedness on the dendrogram.
An optimization value of 1% and a position tolerance of 2% were used for the analysis.

Statistical analysis

Drug susceptibility data were analysed using WHONET v. 5.6 software, recommended by the WHO. The $\chi^2$ test, calculated using SPSS software v. 10.0 (SPSS Inc., USA), was used for statistical comparisons. A two-tailed cut-off of $P<0.05$ indicated statistical significance.

RESULTS

Frequency of serotype 23F

In the entire study period, serotype 23F was identified in 8.9% (101/1132) S. pneumoniae isolates and their frequencies for different years were 10.7% ($n=25$), 6.7% ($n=12$), 10.4% ($n=21$), 6.6% ($n=16$), 10.2% ($n=14$), and 9.3% ($n=13$), respectively. No temporal trend was found ($\chi^2=30.00, P>0.05$).

Antimicrobial susceptibility testing

All 99 strains were susceptible to VAN and AMC. The number of intermediate and resistant strains, as well as their frequency against seven other antimicrobials, is shown in Table 1. Only one isolate was resistant to PEN (MIC 16 $\mu$g/ml). Non-susceptibility to CXM significantly increased from 4% (1997) to 28.5% (2006) to 92.4% (2010). A total of 97 (98.0%) isolates were resistant to ERY with high MICs (94 isolates had a MIC $\geq$256 $\mu$g/ml); 95 carried only the $ermB$ gene and two carried both $ermB$ and $mefA/E$ genes. No strain was positive for the $mefA/E$ gene alone. Moreover, 90 isolates had multidrug resistance to ERY, SXT, TCY, and 25 isolates in addition were resistant to $\beta$-lactam antibiotics.

MLST and PFGE

MLST analysis revealed 16 STs, six of which were novel (ST6825, ST7909–913). The most common STs were ST342 (45.5%), ST81 (17.2%), ST802 (8.1%), ST2624 (6.1%), ST242 (5.1%), and ST6321 (5.1%). The predominant STs varied during the study period. The first ST81 was isolated in 2001, and its frequency increased thereafter, reaching 84.6% in 2010. In 1997–2006, ST342 was predominant and fluctuated over the study with a decreasing trend; none was identified in 2010 (Fig. 1).

Table 1. Susceptibility against seven antimicrobials of the 99 serotype 23F strains isolated in Beijing, n (%)

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<tr>
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<td>9 (69.3)</td>
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<td>3 (23.1)</td>
<td>9 (9.1)</td>
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<tr>
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<td>1 (6.3)</td>
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<td>1 (7.7)</td>
<td>5 (5.1)</td>
</tr>
<tr>
<td>Res</td>
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<td>0</td>
<td>0</td>
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<td>0</td>
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<tr>
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<tr>
<td>Res</td>
<td>24 (96.0)</td>
<td>9 (90)</td>
<td>21 (100)</td>
<td>16 (100)</td>
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<td>97 (98.0)</td>
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<td>0</td>
<td>2 (9.5)</td>
<td>0</td>
<td>1 (7.1)</td>
<td>3 (23.1)</td>
<td>6 (6.1)</td>
</tr>
<tr>
<td>Res</td>
<td>21 (84.0)</td>
<td>10 (100)</td>
<td>18 (85.7)</td>
<td>16 (100)</td>
<td>13 (92.9)</td>
<td>10 (76.9)</td>
<td>88 (88.9)</td>
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<td>Sulfamethoxazole–trimethoprim</td>
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<tr>
<td>Int</td>
<td>1 (4.0)</td>
<td>0</td>
<td>1 (4.7)</td>
<td>2 (12.5)</td>
<td>1 (7.1)</td>
<td>1 (7.7)</td>
<td>6 (6.1)</td>
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<tr>
<td>Res</td>
<td>21 (84.0)</td>
<td>10 (100)</td>
<td>20 (95.3)</td>
<td>14 (87.5)</td>
<td>12 (85.8)</td>
<td>12 (92.3)</td>
<td>89 (89.9)</td>
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<tr>
<td>Chloramphenicol</td>
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<td>0</td>
</tr>
<tr>
<td>Res</td>
<td>4 (16.0)</td>
<td>2 (20.0)</td>
<td>4 (19.0)</td>
<td>5 (31.3)</td>
<td>2 (14.3)</td>
<td>0</td>
<td>17 (17.2)</td>
</tr>
</tbody>
</table>

Int, Intermediate; Res, resistant.
eBURST analysis found four CCs and five singletons. CC342 comprising five STs was the most common group and accounted for 59.6% of all the isolates and CC81 for 18.2%; only one of this complex was not identified as ST81. CC802 (nine isolates), CC242 (seven isolates), and singletons are shown in the population snapshot in Figure 2. Seven different PFGE patterns were found in all isolates and the four major patterns (A–D) corresponded respectively to CC242, CC81, CC802, and CC342 (Supplementary Fig. S1, available online).

Except for one isolate, all isolates identified as CC81 and CC242 were non-susceptible to CXM and showed higher PEN MICs than the other CC/STs (Table 2, Supplementary Fig. S1). Most of the strains (96%, 24/25) in these two CCs were consistently resistant to β-lactam, ERY, SXT, and TCY. By contrast, all CC342 strains were susceptible to the four tested β-lactam antimicrobials, except for one isolate that was classified as intermediate to CRO (MIC 2 μg/ml). The CC342 strains were frequently resistant to ERY, SXT, and TCY.

**DISCUSSION**

The present study showed that the frequency of serotype 23F in all the *S. pneumoniae* isolates was 8.9%, which is similar to previously reported data from...
No temporal trend was found. A significant decrease in serotype 23F was reported in Western countries after the global introduction of the 7-valent conjugate pneumococcal vaccine (PCV7) [2, 20, 21]. Since September 2008, PCV7 has become available in the private sector in China but to date, only a few parents have sought immunization for their children because of the very high cost (860 RMB or US$136 per dose). Thus, PCV7 immunization is not yet ‘universally administered’ in China and hence, no decrease in the frequency of serotype 23F was evident in our study.

ST81, in the present study, was found to be the predominant ST in 2010. Zhang et al. [9] also collected ST81 isolates from 2009 to 2010 in southern China but this clone proved to be rather uncommon during the period from 1997 to 2006. ST81 was first identified in 1978 by Spanish researchers [22] and since then has been found to have spread globally. This ST was very common in the USA and Europe before the introduction of PCV [8, 23]. Some earlier surveys indicated that ST81 had emerged and remained stable in some Asian countries or regions from the 1990s [24–28] and thus the clone may have spread into the hinterland of China from other countries in the region, and may have the potential to spread further. This epidemic spread may be a result of the increasing movements of populations, which contributes to the emergence of antimicrobial resistance worldwide. This situation emphasizes the need for vaccination to interrupt the transmission of resistant clones.

In the present study, ST342 was found to have been the predominant strain before 2006 but its frequency decreased slightly from 1997 to 2006 and unexpectedly was found in 2010, but was largely displaced by ST81. Considering the high β-lactam resistance of ST81 (or CC81), the low resistance of ST342 (or CC342), and the common use of PEN and cephalosporin in China [29], the genotype replacement in serotype 23F may be driven by antibiotic use. In our previous reports [30–32], we emphasized that antibiotic abuse increased serogroup 19 (especially 19A) and subtype 6B-II before the introduction of PCV7 in China. The present study reconfirms this phenomenon because the genotype replacement described here cannot be associated with vaccine administration. A similar genotype replacement was discovered in other serotypes and in other countries where serotype replacement was identified before the introduction of PCV7 [33].

<p>| Table 2. Susceptibility of the 97 serotype 23F strains to seven antimicrobial in different clonal complexes (CCs) and sequence types (STs) |</p>
<table>
<thead>
<tr>
<th>CCs</th>
<th>CRO</th>
<th>PEN</th>
<th>ERY</th>
<th>AMC</th>
<th>CXM</th>
<th>CHL</th>
<th>TCY</th>
<th>SXT</th>
<th>TCY</th>
<th>Int</th>
<th>Res</th>
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<tbody>
<tr>
<td>ST81</td>
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<td>0</td>
<td>1 (100)</td>
<td>0</td>
<td>1 (100)</td>
<td>0</td>
<td>1 (100)</td>
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<td>1 (100)</td>
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<tr>
<td>ST83</td>
<td>5</td>
<td>0</td>
<td>1 (20)</td>
<td>0</td>
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<td>1 (20)</td>
<td>0</td>
<td>1 (20)</td>
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<td>1 (20)</td>
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<tr>
<td>Others</td>
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<td>0</td>
<td>0</td>
<td>0</td>
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<td>0</td>
<td>14 (100)</td>
<td>0</td>
<td>13 (92.9)</td>
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<tr>
<td>Total</td>
<td>97</td>
<td>1 (1.0)</td>
<td>4 (4.0)</td>
<td>2 (2.0)</td>
<td>0</td>
<td>24 (24.2)</td>
<td>0</td>
<td>97 (97.9)</td>
<td>5 (5.1)</td>
<td>90 (90.9)</td>
<td>6 (6.1)</td>
</tr>
</tbody>
</table>

PEN, Penicillin; AMC, amoxicillin–clavulanic acid; CXM, cefuroxime; CHL, chloramphenicol; ERY, erythromycin; TCY, tetracycline; Int, Intermediate; Res, resistant.

Beijing [11, 18, 19].
In Asia, most ERY-resistant strains that carry both \(ermB\) and \(mefA/E\) are serotypes 19F and 14 [34]. The prevalence of macrolide-resistant genes varies in different countries and regions. For example, the dominant gene in the hinterland of China and Taiwan is \(ermB\), whereas in Hong Kong it is \(mefA/E\) [33–36]. Therefore, almost all ERY-resistant strains expressing 23F (97.9%) carried only the \(ermB\) gene. However, the very low carriage of \(mefA/E\) in serotype 23F (2%) should be considered. Ip et al. [24] found that the \(mefA/E\) gene was expressed in serotype 23F isolates with ST81 in Hong Kong in 1994. Recently, Byung et al. [28] reported for the first time in Korea three isolates belonging to CC81 (with serotypes 23F, 15B/C, and 6A) that were positive for both \(ermB\) and \(mefA/E\) genes. In the present study, two ST81 isolates (CC81) harbouring both genes were identified in 2004 and 2010. These findings suggest that the \(mefA/E\) gene may have been transferred from another clone to ST81 when it spread into Asia.

A limitation of the present study is the disruption of survey continuity from 2007 to 2009 and further that all strains were isolated from a single hospital. Considering the high number of patient visits to the largest paediatric hospital in China (5000–6000 visits per day on average) and the total number of pneumococcal isolates, we believe that the present results objectively demonstrate the epidemiological changes of serotype 23F \(S.~pneumoniae\) in Beijing.

In summary, serotype 23F remains the most common serotype in children with respiratory infections in Beijing. However, the genotype component has changed because of antibiotic misuse. The MDR PMEN1 clone (ST81) replaced another MDR ST342 when it spread into Asia.

### SUPPLEMENTARY MATERIAL

For supplementary material accompanying this paper visit http://dx.doi.org/10.1017/S0950268812002269.

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### DECLARATION OF INTEREST

None.

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