Selection of *Streptococcus pneumoniae* Mutants Having Reduced Susceptibility to Moxifloxacin and Levofloxacin

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With *Streptococcus pneumoniae*, moxifloxacin was 4- and 10-fold more effective than levofloxacin at restricting selection of resistant mutants and at killing resistant mutants, respectively. The selection frequency for first-step topoisomerase mutants was 1,000 times lower for moxifloxacin than for levofloxacin; this difference was lost when second-step mutants were selected.

Resistance to penicillin and macrolides is widespread among isolates of *Streptococcus pneumoniae* (6), and use of fluoroquinolones for treatment of pneumonia has increased (3). Decreased fluoroquinolone susceptibility is being reported (3), and resistant isolates have been recovered from patients after levofloxacin has failed to effect a cure (15, 16, 19; R. J. Davidson, J. DeAzavedo, D. Bast, J. Arbique, R. Bethune, C. Duncan, A. McGeer, and D. E. Low, Abstr. 40th Intersci. Conf. Antimicrob. Agents Chemother., abstr. 825, p. 127, 2000; N. O. Fishman, B. Suh, L. M. Weigel, B. Lorber, S. Gelone, A. L. Truant, and T. D. Gootz, 39th Intersci. Conf. Antimicrob. Agents Chemother., abstr. 825, p. 111, 1999). While the prevalence of resistance to levofloxacin is still low in the United States, a five- to sixfold increase in resistance was recorded between surveys of 1997 to 1998 and 1998 to 1999 (18). With several bacteria, addition of a C-8-methoxy group to N-1 cyclopropyl fluoroquinolones improves potency against gyrase and topoisomerase IV resistance mutants (7, 12–14, 20–22), and work with clinical isolates of *S. pneumoniae* suggests that the C-8-methoxy compound moxifloxacin might restrict the selection of resistant mutants more effectively than levofloxacin (1). The present study compared moxifloxacin and levofloxacin for selection of resistant mutants of *S. pneumoniae* in vitro.

*S. pneumoniae* strain ATCC 49619 was grown as liquid cultures in Todd-Hewitt broth (THB; Difco, Detroit, Mich.) containing 10% sheep blood (Hemostat, Dixon, Calif.). For selection of resistant mutants, late-exponential-stage cultures were applied to brain heart infusion (BHI; Difco) agar plates (≤10⁹ CFU/150-mm-diameter plate) containing 10% sheep blood and either moxifloxacin (Bayer Corporation, West Haven, Conn.) or levofloxacin (R. W. Johnson Pharmaceutical Research Institute, Spring House, Pa.). Colonies were recovered and retested for growth on the selecting concentration of fluoroquinolone. DNA was isolated, quinolone-resistance-determining regions (QRDRs) were amplified, and nucleotide sequences were determined as described previously (1, 19). Lethal activity was determined by incubating bacteria grown as liquid cultures (about 3 × 10⁷ CFU/ml) with fluoroquinolone at 37°C for 16 h. Aliquots from samples diluted into cold THB were spotted in triplicate onto drug-free BHI agar containing 10% sheep blood. Colonies were counted after overnight incubation at 37°C in the presence of 5% CO₂.

Fluoroquinolone concentration dramatically affected the recovery of resistant mutants. As the concentration increased, the fraction of input cells recovered as mutant colonies dropped sharply, passed through an inflection point, and then continued to drop sharply (Fig. 1A). When the QRDRs of genes encoding DNA gyrase (*gyrA* and *gyrB*) and DNA topoisomerase IV (*parE* and *parC*) were determined, GyrA variants (Ser-81 to Tyr and Ser-81 to His) were found, but only when 10⁹ or more cells were tested. No mutation was found in *gyrB, parC, or parE*. With levofloxacin, ParC variants (Ser-79 to Tyr and Asp-83 to His) were recovered, but only when at least 1.8 × 10⁶ cells were tested. No mutation was detected in *gyrA, gyrB, or parE*. These data confirm that DNA gyrase is the primary target for moxifloxacin, while topoisomerase IV is the primary target for levofloxacin (15, 16). The 1,000-fold difference in selection of target mutants has been observed with other pairs of fluoroquinolones (9). It may reflect intrinsic differences between the targets. (With *Escherichia coli*, parC resistance mutations are codominant with the wild type, while *gyrA* mutations are recessive [10, 11].) Non-topoisomerase mutants were recovered at low concentration: for some, reserpine lowered the fluoroquinolone MIC (not shown), indicating involvement of active efflux in reduced fluoroquinolone susceptibility (2). The concentration at which no mutant was recovered when more than 10¹⁰ cells were tested (mutant prevention concentration [MPC]) (8) was fourfold lower for moxifloxacin (Fig. 1A, arrowheads, and Table 1).

When second-step mutants were selected from a first-step *parC* resistance strain (KD2138) by using moxifloxacin or levofloxacin, the inflection region was more pronounced than with wild-type cells, and the MPC was higher (Fig. 1B). Determination of QRDR nucleotide sequences in mutant DNA showed that *gyrA* and *parC* double mutants were recovered (labeled arrows, Fig. 1B). Similar phenomena were observed with a first-step *gyrA* resistance mutant (strain KD2139; Fig. 1C). For both mutants, the MPC was higher for levofloxacin. This
may reflect the inability of levofloxacin to kill resistant mutants at the MPC (Fig. 2 and Table 1).

Several parameters obtained from the experiments described above allow quantitative comparison of moxifloxacin and levofloxacin (Table 1). First, moxifloxacin MICs required to block colony formation by 99% (MIC99, unlabeled arrows in Fig. 1) were 21, 62, and 17% of that of levofloxacin with wild-type, gyrA-resistant, and parC-resistant cells, respectively. The difference between the two mutants reflects the target preference of the two compounds (16, 17). When the MPC is taken as the ability to restrict the growth of the most resistant mutant, moxifloxacin was four to five times more effective with wild-type cells and resistant mutants. With respect to lethal activity, moxifloxacin concentrations that allowed only 1% survival (LD99) were 18, 8, and 9% that of levofloxacin with wild-type, gyrA-resistant, and parC-resistant cells, respectively. Collectively, these data show that moxifloxacin is intrinsically more active against wild-type and resistant S. pneumoniae.

From published pharmacokinetic measurements (4, 5, 18a), we calculated that drug concentrations in serum were above the MPC longer for moxifloxacin (38 h at the 400-mg dose) than for levofloxacin (8 h at the 500-mg dose; 18 h at the 750-mg dose). The area under the time-concentration curve was greater for moxifloxacin than that of levofloxacin (500-mg dose). Collectively, these data support the previous conclusion (1) that drug concentrations in serum are more likely to exceed the MPC of moxifloxacin than that of levofloxacin with S. pneumoniae. Additional work is required to provide estimates of available drug at the relevant sites of infection.

The acquisition of a first-step parC mutation by levofloxacin may have only a small effect on the ability of moxifloxacin to prevent growth (compare MIC99s in Table 1); however, it increases by several orders of magnitude the probability for acquiring a second mutation (compare Fig. 1A with B). Thus, preservation of the effectiveness of new fluoroquinolones, such as moxifloxacin and gemifloxacin, may require avoiding the selective enrichment of first-step mutants by older compounds such as ciprofloxacin and levofloxacin.

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TABLE 1. Effect of gyrA and parC resistance mutations on fluoroquinolone activity

<table>
<thead>
<tr>
<th>S. pneumoniae strain</th>
<th>MIC99 (µg/ml)a</th>
<th>MPC (µg/ml)b</th>
<th>LD99 (µg/ml)c</th>
<th>% Survival at wild-type MPCd</th>
</tr>
</thead>
<tbody>
<tr>
<td>ATCC 49619 (wild type)</td>
<td>0.15</td>
<td>0.7</td>
<td>0.5</td>
<td>2.3</td>
</tr>
<tr>
<td>KD2139 (gyrA)c</td>
<td>0.56</td>
<td>0.9</td>
<td>6.0</td>
<td>30</td>
</tr>
<tr>
<td>KD2138 (parC)c</td>
<td>0.29</td>
<td>1.7</td>
<td>10</td>
<td>50</td>
</tr>
</tbody>
</table>

a Data obtained from Fig. 1. A replicate experiment in each case gave similar values.
b Data obtained from Fig. 2. A replicate experiment in each case gave similar values.
c Survival was determined relative to that of cells present immediately before addition of fluoroquinolone. Survival >100% indicates cell growth.

d Survival was determined relative to the number of input CFU. Unlabeled arrows indicate MIC99. The triangle indicates no colony recovered at that drug concentration and bacterial load. The dashed line indicates one colony recovered per 10^10 cells tested. Similar results were obtained in two independent experiments. Three strains were tested: wild-type strain ATCC 49619 (A); strain KD2138, a ParC (Ser-79 to Tyr) variant (B); and strain KD2139, a GyrA (Ser-81 to Phe) variant (C). Two double mutants were recovered from each point indicated by an arrow, and the nucleotide sequence of the QRDRs of gyrA and parC was determined for each mutant. Letters indicate recovery of the following mutants: a (GyrA Ser-81 to Phe), b (GyrA Ser-81 to Tyr), c (GyrA Glu-85 to Lys), and d (Par C Ser-79 to Tyr). A single letter with an arrow indicates that both mutants examined had the same change.

FIG. 1. Effect of fluoroquinolone concentration on the recovery of single-step, resistant mutants. S. pneumoniae was grown to late log phase and then distributed at various dilutions to agar plates containing the indicated concentrations of moxifloxacin (open circles) or levofloxacin (solid circles). After incubation, colonies were counted, and the fraction recovered was determined relative to the number of input CFU. Unlabeled arrows indicate MIC99. The triangle indicates no colony recovered at that drug concentration and bacterial load. The dashed line indicates one colony recovered per 10^10 cells tested. Similar results were obtained in two independent experiments. Three strains were tested: wild-type strain ATCC 49619 (A); strain KD2138, a ParC (Ser-79 to Tyr) variant (B); and strain KD2139, a GyrA (Ser-81 to Phe) variant (C). Two double mutants were recovered from each point indicated by an arrow, and the nucleotide sequence of the QRDRs of gyrA and parC was determined for each mutant. Letters indicate recovery of the following mutants: a (GyrA Ser-81 to Phe), b (GyrA Ser-81 to Tyr), c (GyrA Glu-85 to Lys), and d (Par C Ser-79 to Tyr). A single letter with an arrow indicates that both mutants examined had the same change.
FIG. 2. Survival of S. pneumoniae in the presence of fluoroquinolone. Cells were incubated in the presence of the indicated concentrations of moxifloxacin (open circles) or levofloxacin (solid circles), after which they were plated on drug-free agar. Percent survival was calculated relative to CFU at the time of drug addition. (A) Wild-type strain ATCC 49619. (B) gyrA resistance mutant strain KD2139. (C) parC resistance mutant KD2138. MPCs, determined as described in the legend to Fig. 1, are indicated by arrows. Similar results were obtained in a replicate experiment.

REFERENCES