Pharmacokinetics of WCK 2349 in Healthy Indian Male Adults under Fasting Conditions

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The global spread of methicillin resistant Staphylococcus aureus (MRSA) in hospital and, more recently, in community is an unmet medical need1,2. WCK 2349, a novel fluoroquinolone, is being developed by Wockhardt as an oral anti-MRSA agent.

The primary objective of this analysis was to characterize the pharmacokinetics (PK) of oral WCK 2349 in healthy adult Indian male volunteers under fasting conditions using Phase 1 data that included one single dose study with 7 dose levels (200 mg, 400 mg, 600 mg, 800 mg, 1000 mg, 1200 mg and 1500 mg) and one multiple dose study with 4 dose levels (600 mg, 800 mg, 1000 mg and 1200 mg twice daily (BID) for five days).

Methods

Intensive venous blood samples were collected per protocol and WCK 2349 plasma concentrations were measured with LC-MS/MS.

1322 observations from 70 subjects (10 subjects per dose level) after single dose administration and 528 observations form 24 subjects (6 subjects per dose level) in steady state after multiple dose administration were analyzed using a non-compartmental approach with Phoenix WinNonlin.

Results

Table 1: Mean Pharmacokinetic Parameters of WCK 2349 for all dose levels

<table>
<thead>
<tr>
<th>STUDY</th>
<th>DOSE (mg)</th>
<th>N</th>
<th>Mean</th>
<th>SD</th>
<th>Mean</th>
<th>SD</th>
<th>Mean</th>
<th>SD</th>
<th>Mean</th>
<th>SD</th>
<th>Mean</th>
<th>SD</th>
</tr>
</thead>
<tbody>
<tr>
<td>S.D.</td>
<td>200</td>
<td>10</td>
<td>31.58</td>
<td>5.92</td>
<td>6.11</td>
<td>1.07</td>
<td>5.11</td>
<td>1.31</td>
<td>6.12</td>
<td>1.26</td>
<td>57.08</td>
<td>14.26</td>
</tr>
<tr>
<td>S.D.</td>
<td>400</td>
<td>10</td>
<td>66.84</td>
<td>10.92</td>
<td>6.12</td>
<td>0.92</td>
<td>9.12</td>
<td>2.21</td>
<td>8.38</td>
<td>1.43</td>
<td>73.71</td>
<td>15.06</td>
</tr>
<tr>
<td>S.D.</td>
<td>600</td>
<td>10</td>
<td>81.48</td>
<td>21.67</td>
<td>7.84</td>
<td>2.04</td>
<td>12.16</td>
<td>1.57</td>
<td>7.67</td>
<td>1.45</td>
<td>85.93</td>
<td>25.31</td>
</tr>
<tr>
<td>S.D.</td>
<td>800</td>
<td>10</td>
<td>138.97</td>
<td>26.16</td>
<td>5.99</td>
<td>1.38</td>
<td>17.84</td>
<td>3.86</td>
<td>7.29</td>
<td>0.97</td>
<td>62.50</td>
<td>14.99</td>
</tr>
<tr>
<td>S.D.</td>
<td>1000</td>
<td>10</td>
<td>157.36</td>
<td>32.82</td>
<td>6.60</td>
<td>1.32</td>
<td>19.15</td>
<td>4.73</td>
<td>7.57</td>
<td>1.68</td>
<td>71.95</td>
<td>21.38</td>
</tr>
<tr>
<td>S.D.</td>
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<td>10</td>
<td>203.39</td>
<td>47.94</td>
<td>6.21</td>
<td>1.45</td>
<td>23.44</td>
<td>3.87</td>
<td>8.09</td>
<td>1.08</td>
<td>71.41</td>
<td>14.26</td>
</tr>
<tr>
<td>S.D.</td>
<td>1500</td>
<td>10</td>
<td>226.06</td>
<td>51.22</td>
<td>7.03</td>
<td>2.00</td>
<td>25.64</td>
<td>4.42</td>
<td>8.39</td>
<td>1.29</td>
<td>83.32</td>
<td>18.30</td>
</tr>
<tr>
<td>BID=SD</td>
<td>600</td>
<td>6</td>
<td>91.11</td>
<td>20.25</td>
<td>125.61</td>
<td>43.33</td>
<td>6.85</td>
<td>1.41</td>
<td>14.81</td>
<td>2.25</td>
<td>6.86</td>
<td>1.60</td>
</tr>
<tr>
<td>BID=SD</td>
<td>800</td>
<td>6</td>
<td>120.10</td>
<td>21.12</td>
<td>156.06</td>
<td>31.21</td>
<td>6.82</td>
<td>1.14</td>
<td>20.56</td>
<td>2.67</td>
<td>6.85</td>
<td>0.97</td>
</tr>
<tr>
<td>BID=SD</td>
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<td>186.36</td>
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<td>254.65</td>
<td>65.24</td>
<td>5.56</td>
<td>1.16</td>
<td>29.68</td>
<td>4.09</td>
<td>7.46</td>
<td>1.11</td>
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<tr>
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<td>6</td>
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<td>321.76</td>
<td>80.34</td>
<td>5.45</td>
<td>0.85</td>
<td>33.78</td>
<td>4.21</td>
<td>8.68</td>
<td>1.54</td>
</tr>
</tbody>
</table>

• Linear increase in Cmax and AUC([0, infinity]) was observed from 200 – 1500 mg single doses.
• For BID for 5 days, the AUC([0, tau]) at steady state was not significantly different from the AUC([0, infinity]) after respective single doses, and the accumulation was minimal.

Conclusions

Conclusions: Single and multiple dose PK studies for WCK 2349 show that the drug was well tolerated and the PK are linear across all dose levels.

References