Recommendations for the Use of Antiretroviral Drugs in Pregnant Women with HIV Infection and Interventions to Reduce Perinatal HIV Transmission in the United States

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**Tipranavir (Aptivus, TPV)**

(Last reviewed December 7, 2018; last updated December 7, 2018)

Tipranavir is classified as Food and Drug Administration Pregnancy Category C. **Tipranavir should not be used during pregnancy.**

**Animal Studies**

**Carcinogenicity**

Tipranavir was neither mutagenic nor clastogenic in a battery of five screening tests, both in vitro and, in animals, in vivo. Long-term carcinogenicity studies of tipranavir have been conducted in mice and rats. Mice were administered tipranavir doses ranging from 30 to 300 mg/kg/day, with or without ritonavir 40 mg/kg/day; all doses resulted in systemic exposures below those seen in humans receiving the recommended dose. Incidence of benign hepatocellular adenomas, combined adenomas/carcinomas, and hepatocellular carcinoma was increased in both male and female mice receiving tipranavir/ritonavir (TPV/r). The clinical relevance of the carcinogenic findings in mice is unknown. Rats were administered doses ranging from 30 to 300 mg/kg/day tipranavir, with or without ritonavir. No drug-related findings were observed in male rats. At the highest dose of tipranavir (approximately equivalent to exposure in humans at the recommended therapeutic dose), an increased incidence of benign follicular cell adenomas of the thyroid gland was observed in female rats. This finding is probably not relevant to humans, because thyroid follicular cell adenomas are considered a rodent-specific effect secondary to enzyme induction.\(^1\)

**Reproduction/Fertility**

Tipranavir had no effect on fertility or early embryonic development in rats at exposure levels that are similar to human exposure levels at the recommended clinical dose (TPV/r 500 mg/200 mg administered twice daily).\(^1\)

**Teratogenicity/Adverse Pregnancy Outcomes**

No teratogenicity was detected in studies of pregnant rats and rabbits with exposure levels that were approximately 1.1-fold and 0.1-fold human exposure levels. Fetal toxicity (decreased ossification and body weights) was observed in rats exposed to 400 mg/kg/day or more of tipranavir (~0.8-fold human exposure). Fetal toxicity was not seen in rats and rabbits at levels of 0.2-fold and 0.1-fold human exposures. In rats, no adverse effects on development occurred at exposure levels of 40 mg/kg/day (~0.2-fold human exposure), but growth inhibition in pups and maternal toxicity were observed at 400 mg/kg/day (~0.8-fold human exposure).\(^1\)

**Placental and Breast Milk Passage**

No animal studies of placental or breast milk passage of tipranavir have been reported.

**Human Studies in Pregnancy**

**Pharmacokinetics**

No studies of tipranavir have been completed in pregnant women or neonates.

**Placental and Breast Milk Passage**

It is unknown if tipranavir passes through the placenta or breast milk in humans. A single case report described relatively high levels of tipranavir in the third trimester and relatively high placental transfer (0.41), as measured by cord blood.\(^2\)

**Teratogenicity/Adverse Pregnancy Outcomes**

The five first-trimester exposures to tipranavir that have been monitored to date in the Antiretroviral Pregnancy Registry are insufficient to draw conclusions regarding the risk of birth defects.\(^3\)
### Excerpt from Table 10\textsuperscript{a}

<table>
<thead>
<tr>
<th>Generic Name (Abbreviation)</th>
<th>Formulation</th>
<th>Dosing Recommendations</th>
<th>Use in Pregnancy</th>
</tr>
</thead>
<tbody>
<tr>
<td>Tipranavir (TPV)</td>
<td>Capsules: 250 mg</td>
<td>Standard Adult Dose: • TPV/r 500 mg/200 mg twice daily With RTV Tablets: • Take with food. With RTV Capsules or Solution: • Take without regard to food; however, administering with food may help make the dose more tolerable.</td>
<td><strong>TPV should not be used</strong> during pregnancy. Moderate placental transfer to fetus reported in 1 patient.\textsuperscript{b} Insufficient data to assess teratogenicity in humans. No evidence of teratogenicity in rats or rabbits. Must be given as low-dose, RTV-boosted regimen.</td>
</tr>
<tr>
<td><strong>Note:</strong> Must be combined with RTV for PK boosting</td>
<td>Oral Solution: 100 mg/mL</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Tipranavir (TPV) Capsules:</strong> 250 mg</td>
<td><strong>Standard Adult Dose:</strong> • TPV/r 500 mg/200 mg twice daily</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Oral Solution:</strong> 100 mg/mL</td>
<td><strong>With RTV Tablets:</strong> • Take with food.</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>With RTV Capsules or Solution:</strong> • Take without regard to food; however, administering with food may help make the dose more tolerable.</td>
<td></td>
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</tr>
<tr>
<td><strong>Dosing in Pregnancy:</strong> • Insufficient data to make dosing recommendation PK in Pregnancy: • Limited PK data in human pregnancy</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

\textsuperscript{a} Individual ARV drug dosages may need to be adjusted in patients with renal or hepatic insufficiency (for details, see the *Adult and Adolescent Guidelines, Appendix B, Table 8.*

\textsuperscript{b} Placental transfer categories are determined by mean or median cord blood/maternal delivery plasma drug ratio:

- **High:** >0.6
- **Moderate:** 0.3–0.6
- **Low:** <0.3

**Key to Acronyms:** PK = pharmacokinetic; RTV = ritonavir; TPV = tipranavir; TPV/r = tipranavir/ritonavir

### References

