AMG-337

Cat. No.: HY-18696
CAS No.: 1173699-31-4
Molecular Formula: C₂₃H₂₂FN₇O₃
Molecular Weight: 463.46
Target: c-Met/HGFR
Pathway: Protein Tyrosine Kinase/RTK
Storage: Powder
-20°C: 3 years
4°C: 2 years
In solvent
-80°C: 6 months
-20°C: 1 month

SOLVENT & SOLUBILITY

**In Vitro**
DMSO: 100 mg/mL (215.77 mM; Need ultrasonic)

<table>
<thead>
<tr>
<th>Preparing Stock Solutions</th>
<th>Solvent Concentration</th>
<th>Mass (1 mg)</th>
<th>Mass (5 mg)</th>
<th>Mass (10 mg)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 mM</td>
<td>2.1577 mL</td>
<td>10.7884 mL</td>
<td>21.5768 mL</td>
<td></td>
</tr>
<tr>
<td>5 mM</td>
<td>0.4315 mL</td>
<td>2.1577 mL</td>
<td>4.3154 mL</td>
<td></td>
</tr>
<tr>
<td>10 mM</td>
<td>0.2158 mL</td>
<td>1.0788 mL</td>
<td>2.1577 mL</td>
<td></td>
</tr>
</tbody>
</table>

Please refer to the solubility information to select the appropriate solvent.

**In Vivo**
1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline
   Solubility: ≥ 2.5 mg/mL (5.39 mM); Clear solution
2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)
   Solubility: ≥ 2.5 mg/mL (5.39 mM); Clear solution
3. Add each solvent one by one: 10% DMSO >> 90% corn oil
   Solubility: ≥ 2.5 mg/mL (5.39 mM); Clear solution

BIOLOGICAL ACTIVITY

**Description**
AMG-337 is a potent and highly selective small molecule ATP-competitive MET kinase inhibitor. AMG 337 inhibits MET kinase activity with an IC50 of < 5nM in enzymatic assays. IC50 value: < 5nM [1] Target: MET in vitro: AMG-337 demonstrates exquisite selectivity for MET when profiled against a diverse panel of over 400 protein and lipid kinases in a competitive binding assay. In cellular assays, AMG 337 inhibits HGF-dependent MET phosphorylation with an IC50 of < 10 nM. [1] AMG 337 is a selective inhibitor of Met, which inhibits multiple mechanisms of Met activation. [2] In vivo: AMG-337 demonstrates robust activity in MET-dependent cancer models. Oral administration of AMG 337 results in robust dose-dependent anti-tumor efficacy in MET amplified gastric cancer xenograft models, with inhibition of tumor growth consistent with the pharmacodynamic
modulation of MET signaling.[1]

REFERENCES
