

产品名称: Verubecestat

产品别名: MK-8931

| 生物活性: | | | | | | | | | | | | | | | | | | | | | | | | | | |
|------------------------------|---|-----------|-----------|------------|------------|------------------------------|--------------------------|------|------|------|-------|------|--|-----------|------------|------------|------|--|-----------|-----------|-----------|-------|--|-----------|-----------|-----------|
| Description | Verubecestat (MK-8931) is a beta-secretase 1 (BACE1) inhibitor under investigation for the treatment of Alzheimer's Disease. | | | | | | | | | | | | | | | | | | | | | | | | | |
| | In Vitro: DMSO : $\geq 35 \text{ mg/mL}$ (85.49 mM) * " \geq " means soluble, but saturation unknown. | | | | | | | | | | | | | | | | | | | | | | | | | |
| | <table border="1"><thead><tr><th rowspan="2">Preparing Stock Solutions</th><th>Solvent Concentration</th><th>Mass</th><th>1 mg</th><th>5 mg</th><th>10 mg</th></tr></thead><tbody><tr><th>1 mM</th><td></td><td>2.4425 mL</td><td>12.2127 mL</td><td>24.4254 mL</td></tr><tr><th>5 mM</th><td></td><td>0.4885 mL</td><td>2.4425 mL</td><td>4.8851 mL</td></tr><tr><th>10 mM</th><td></td><td>0.2443 mL</td><td>1.2213 mL</td><td>2.4425 mL</td></tr></tbody></table> | | | | | Preparing Stock Solutions | Solvent Concentration | Mass | 1 mg | 5 mg | 10 mg | 1 mM | | 2.4425 mL | 12.2127 mL | 24.4254 mL | 5 mM | | 0.4885 mL | 2.4425 mL | 4.8851 mL | 10 mM | | 0.2443 mL | 1.2213 mL | 2.4425 mL |
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| Solvent&Solubility | <p>*请根据产品在不同溶剂中的溶解度选择合适的溶剂配制储备液。一旦配成溶液, 请分装保存, 避免反复冻融造成的产品失效。</p> <p>储备液的保存方式和期限 -80°C, 6 months; -20°C, 1 month。 -80°C 储存时, 请在 6 个月内使用, -20°C 储存时, 请在 1 个月内使用。</p> <p>In Vivo:</p> <p>请根据您的实验动物和给药方式选择适当的溶解方案。以下溶解方案都请先按照 In Vitro 方式配制澄清的储备液, 再依次添加助溶剂:</p> <p>——为保证实验结果的可靠性, 澄清的储备液可以根据储存条件, 适当保存; 体内实验的工作液, 建议您现用现配, 当天使用; 以下溶剂前显示的百分比是指该溶剂在您配制终溶液中的体积占比; 如在配制过程中出现沉淀、析出现象, 可以通过加热和/或超声的方式助溶</p> <p>1.请依序添加每种溶剂: 10% DMSO → 40% PEG300 → 5% Tween-80 → 45% saline</p> <p>Solubility: $\geq 2.5 \text{ mg/mL}$ (6.11 mM); Clear solution</p> <p>此方案可获得 $\geq 2.5 \text{ mg/mL}$ (6.11 mM, 饱和度未知) 的澄清溶液。</p> <p>以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 400 μL PEG300 中, 混合均匀。向上述体系中加入 50 μL Tween-80, 混合均匀; 然后继续加入 450 μL 生理盐水定容至 1 mL。</p> <p>2.请依序添加每种溶剂: 10% DMSO → 90% (20% SBE-β-CD in saline)</p> <p>Solubility: $\geq 2.5 \text{ mg/mL}$ (6.11 mM); Clear solution</p> <p>此方案可获得 $\geq 2.5 \text{ mg/mL}$ (6.11 mM, 饱和度未知) 的澄清溶液。</p> <p>以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 20% 的 SBE-β-CD 生理盐水溶液中, 混合均匀。</p> <p>3.请依序添加每种溶剂: 10% DMSO → 90% corn oil</p> <p>Solubility: $\geq 2.5 \text{ mg/mL}$ (6.11 mM); Clear solution</p> <p>此方案可获得 $\geq 2.5 \text{ mg/mL}$ (6.11 mM, 饱和度未知) 的澄清溶液, 此方案不适用于实验周期在半个月以上的实验。</p> <p>以 1 mL 工作液为例, 取 100 μL 25.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 玉米油中, 混合均匀。</p> | | | | | | | | | | | | | | | | | | | | | | | | | |
| | [1]. Yan R Stepping closer to treating Alzheimer's disease patients with BACE1 inhibitor drugs. <i>Transl Neurodegener.</i> 2016 Jul 14;5:13. | | | | | | | | | | | | | | | | | | | | | | | | | |

References

[2]. Nisha CM et al. Molecular Docking and In Silico ADMET Study Reveals Acylguanidine 7a as a Potential Inhibitor of β -Secretase. Adv Bioinformatics. 2016 Apr 10.



源叶生物