



上海源叶生物科技有限公司
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产品名称: LXS196

产品别名: LXS196

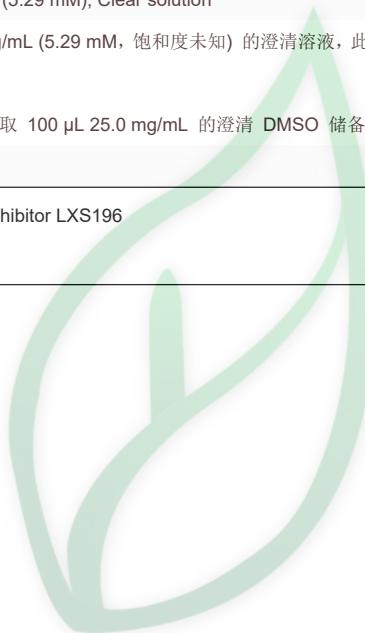
生物活性:

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|---|---|---|----------------------------|------------|------------|
| Description | LXS196 is a potent, selective and orally active protein kinase C (PKC) inhibitor, with IC ₅₀ values of 1.9 nM, 0.4 nM and 3.1 μM for PKCα, PKCθ and GSK3β, respectively. It can be used for the treatment of uveal melanoma[1][2]. | | | | |
| IC ₅₀ & Target | PKCα | PKCθ | GSK3β | | |
| | 1.9 nM (IC ₅₀) | 0.4 nM (IC ₅₀) | 3.1 μM (IC ₅₀) | | |
| In Vitro | Upon oral administration, protein kinase C inhibitor LXS196 binds to and inhibits PKC, which prevents the activation of PKC-mediated signaling pathways. This may lead to the induction of cell cycle arrest and apoptosis in susceptible tumor cells. PKC, a serine/threonine protein kinase overexpressed in certain types of cancer cells, is involved in tumor cell differentiation, proliferation, invasion and survival[1]. | | | | |
| In Vivo | LXS196 (compound 9) (15, 30, 75, 150 mg/kg, P.O., mice) shows improved efficacy (regression) in a 92.1 GNAQ uveal melanoma xenograft model in a dose-dependently manner[2]. | | | | |
| | Animal Model: | Mice implanted with 92.1 GNAQ mutant uveal melanoma cells[2]. | | | |
| | Dosage: | 15, 30, 75, 150 mg/kg | | | |
| | Administration: | P.O. (bid) for 35 days | | | |
| | Result: | Dose-dependently suppressed the tumor growth. | | | |
| Solvent&Solubility | In Vitro: DMSO : ≥ 100 mg/mL (211.65 mM) * "≥" means soluble, but saturation unknown. | | | | |
| | Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg | 10 mg |
| | | 1 mM | 2.1165 mL | 10.5827 mL | 21.1654 mL |
| | | 5 mM | 0.4233 mL | 2.1165 mL | 4.2331 mL |
| | | 10 mM | 0.2117 mL | 1.0583 mL | 2.1165 mL |
| *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。 | | | | | |
| 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。 | | | | | |
| In Vivo: 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: ——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 | | | | | |
| 1.请依序添加每种溶剂: 10% DMSO→40% PEG300 →5% Tween-80 → 45% saline Solubility: ≥ 2.5 mg/mL (5.29 mM); Clear solution 此方案可获得 ≥ 2.5 mg/mL (5.29 mM, 饱和度未知) 的澄清溶液。 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 | | | | | |



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| | <p>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: $\geq 2.5 \text{ mg/mL}$ (5.29 mM); Clear solution</p> <p>此方案可获得 $\geq 2.5 \text{ mg/mL}$ (5.29 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: $\geq 2.5 \text{ mg/mL}$ (5.29 mM); Clear solution</p> <p>此方案可获得 $\geq 2.5 \text{ mg/mL}$ (5.29 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> |
| References | [1]. Protein Kinase C Inhibitor LXS196 [2]. US20180179181. |



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