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产品名称: **KN-93 (phosphate)**
产品别名: **KN-93 phosphate**

| 生物活性: | | | | |
|---------------------------|--|----------------------------|-----------|-----------|
| Description | KN-93 phosphate is a novel membrane-permeant synthetic inhibitor of purified neuronal CaMK-II, with K_i of 370 nM. | | | |
| IC ₅₀ & Target | K _i : 370 nM (CaMK-II) | | | |
| In Vitro | After 2 days of KN-93 treatment, 95% of cells are arrested in G1. G1 arrest is reversible; 1 day after KN-93 release, a peak of cells had progressed into S and G2-M. KN-93 also blocks cell growth stimulated by basic fibroblast growth factor, platelet-derived growth factor-BB, and epidermal growth factor in NIH 3T3 fibroblasts ^[1] . KN-93 inhibits the H ⁺ , K ⁺ -ATPase activity but strongly dissipates the proton gradient formed in the gastric membrane vesicles and reduces the volume of luminal space ^[2] . KN-93 (0.5 μM) prevents increased LV developed pressure during action potential prolongation and early afterdepolarizations. Ca ²⁺ -independent CaM kinase activity is increased during early afterdepolarizations and this increase is prevented by KN-93 ^[3] . | | | |
| Solvent&Solubility | In Vitro: DMSO : 100 mg/mL (166.94 mM; Need ultrasonic) H₂O : 50 mg/mL (83.47 mM; Need ultrasonic) | | | |
| | Preparing Stock Solutions | Solvent Mass Concentration | 1 mg | 5 mg |
| | | 1 mM | 1.6694 mL | 8.3468 mL |
| | | 5 mM | 0.3339 mL | 1.6694 mL |
| | | 10 mM | 0.1669 mL | 0.8347 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: ≥ 10 mg/mL (16.69 mM); Clear solution 此方案可获得 ≥ 10 mg/mL (16.69 mM, 饱和度未知) 的澄清溶液。 以 1 mL 工作液为例, 取 100 μL 100.0 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。 2.请依序添加每种溶剂: 10% DMSO→ 90% (20% SBE-β-CD in saline) Solubility: ≥ 10 mg/mL (16.69 mM); Clear solution | | | |
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| | <p>此方案可获得 ≥ 10 mg/mL (16.69 mM, 饱和度未知) 的澄清溶液。</p> <p>以 1 mL 工作液为例, 取 100 μL 100.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 20% 的 SBE-β-CD 生理盐水水溶液中, 混合均匀。</p> <p>3.请依序添加每种溶剂: 10% DMSO \rightarrow90% corn oil</p> <p>Solubility: ≥ 10 mg/mL (16.69 mM); Clear solution</p> <p>此方案可获得 ≥ 10 mg/mL (16.69 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 100.0 mg/mL 的澄清 DMSO 储备液加到 900 μL 玉米油中, 混合均匀。</p> |
| References | <p>[1]. Tombes RM, et al. G1 cell cycle arrest and apoptosis are induced in NIH 3T3 cells by KN-93, an inhibitor of CaMK-II (the multifunctional Ca²⁺/CaM kinase). Cell Growth Differ. 1995 Sep;6(9):1063-70.</p> <p>[2]. Li J, et al. Curcumin Attenuates Retinal Vascular Leakage by Inhibiting Calcium/Calmodulin-Dependent Protein Kinase II Activity in Streptozotocin-Induced Diabetes. Cell Physiol Biochem. 2016;39(3):1196-208.</p> <p>[3]. Mamiya N, et al. Inhibition of acid secretion in gastric parietal cells by the Ca²⁺/calmodulin-dependent protein kinase II inhibitor KN-93. Biochem Biophys Res Commun. 1993 Sep 15;195(2):608-15.</p> <p>[4]. Anderson ME, et al. KN-93, an inhibitor of multifunctional Ca⁺⁺/calmodulin-dependent protein kinase, decreases early afterdepolarizations in rabbit heart. J Pharmacol Exp Ther. 1998 Dec;287(3):996-1006.</p> |

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