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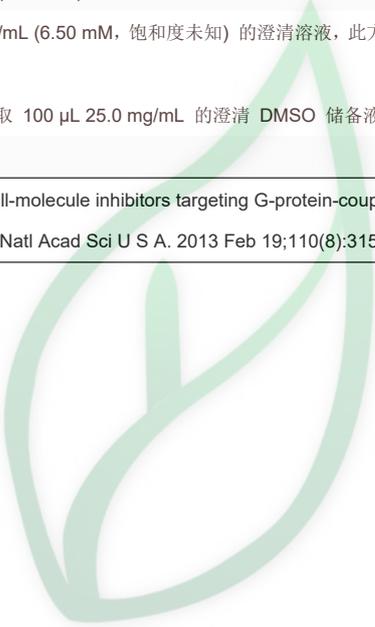
产品名称: Y16  
 产品别名: Y16

生物活性:																											
<b>Description</b>	Y16 is a specific inhibitor of Leukemia-associated Rho guanine nucleotide exchange factor (LARG) with a $K_d$ value of 76 nM. Y16 is active in blocking the interaction of LARG and related G-protein-coupled Rho GEFs with RhoA. Y16 shows no detectable effect on other diffuse B-cell lymphoma (Dbl) family Rho GEFs, Rho effectors, or a RhoGAP[1].																										
<b>IC<sub>50</sub> &amp; Target</b>	Kd: 76 nM (LARG)[1]																										
<b>In Vitro</b>	Y16 (10-30 $\mu$ M; 24 hours; NIH 3T3 cells) could inhibit RhoA-GTP formation induced by serum dose dependently and is specific for RhoA[1]. Y16 (10-30 $\mu$ M; 24 hours; NIH 3T3 cells) efficiently inhibits serum or SDF-1 $\alpha$ -induced phospho-MLC and phospho-FAK formation, which are downstream of RhoA[1].																										
	<b>Cell Viability Assay[1]</b>																										
	Cell Line: NIH 3T3 cells																										
	Concentration: 10 $\mu$ M, 30 $\mu$ M																										
	Incubation Time: 24 hours																										
	Result: Inhibited RhoA-GTP formation induced by serum dose dependently and was specific for RhoA.																										
	<b>Western Blot Analysis[1]</b>																										
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<b>Solvent&amp;Solubility</b>	<b>In Vitro:</b> DMSO : 25 mg/mL (65.03 mM; Need ultrasonic) H <sub>2</sub> O : < 0.1 mg/mL (insoluble)																										
	<table border="1"> <thead> <tr> <th rowspan="2">Preparing Stock Solutions</th> <th>Solvent</th> <th>Mass</th> <th rowspan="2">1 mg</th> <th rowspan="2">5 mg</th> <th rowspan="2">10 mg</th> </tr> <tr> <th>Concentration</th> <th></th> </tr> </thead> <tbody> <tr> <td></td> <td>1 mM</td> <td></td> <td>2.6013 mL</td> <td>13.0063 mL</td> <td>26.0125 mL</td> </tr> <tr> <td></td> <td>5 mM</td> <td></td> <td>0.5203 mL</td> <td>2.6013 mL</td> <td>5.2025 mL</td> </tr> <tr> <td></td> <td>10 mM</td> <td></td> <td>0.2601 mL</td> <td>1.3006 mL</td> <td>2.6013 mL</td> </tr> </tbody> </table>	Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg	Concentration			1 mM		2.6013 mL	13.0063 mL	26.0125 mL		5 mM		0.5203 mL	2.6013 mL	5.2025 mL		10 mM		0.2601 mL	1.3006 mL	2.6013 mL
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*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。																											
<b>In Vivo:</b> 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																											



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	<p>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: 2.5 mg/mL (6.50 mM); Suspended solution; Need ultrasonic</p> <p>此方案可获得 2.5 mg/mL (6.50 mM)的均匀悬浊液, 悬浊液可用于口服和腹腔注射。</p> <p>以 1 mL 工作液为例, 取 100 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 400 <math>\mu</math>L PEG300 中, 混合均匀向上述体系中加入 50 <math>\mu</math>L Tween-80, 混合均匀; 然后继续加入 450 <math>\mu</math>L 生理盐水定容至 1 mL。</p> <p>2.请依序添加每种溶剂: 10% DMSO →90% corn oil</p> <p>Solubility: <math>\geq</math> 2.5 mg/mL (6.50 mM); Clear solution</p> <p>此方案可获得 <math>\geq</math> 2.5 mg/mL (6.50 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 <math>\mu</math>L 25.0 mg/mL 的澄清 DMSO 储备液加到 900 <math>\mu</math>L 玉米油中, 混合均匀。</p>
<b>References</b>	<p>[1]. Shang X, et al. Small-molecule inhibitors targeting G-protein-coupled Rho guanine nucleotide exchange factors. Proc Natl Acad Sci U S A. 2013 Feb 19;110(8):3155-60.</p>



# 源叶生物