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产品名称: RI-2  
 产品别名: RI-2

生物活性:				
<b>Description</b>	RI-2 is a reversible RAD51 inhibitor, with an IC <sub>50</sub> of 44.17 μM, and specifically inhibits homologous recombination repair in human cells.			
<b>IC<sub>50</sub> &amp; Target</b>	IC50: 44.17 μM (RAD51) <sup>[1]</sup>			
<b>In Vitro</b>	RI-2 (7a) is a reversible RAD51 inhibitor, with an IC <sub>50</sub> of 44.17 μM. RI-2 specifically inhibits homologous recombination repair in human cells. RI-2 (150 μM) induces a significant sensitization of cells <sup>[1]</sup> .			
<b>Solvent&amp;Solubility</b>	<b>In Vitro:</b> DMSO : 15.5 mg/mL (35.77 mM; Need ultrasonic and warming)			
		<b>Solvent</b>	<b>Mass</b>	
		<b>Concentration</b>		
	<b>Preparing</b>	1 mM	2.3080 mL	11.5399 mL
	<b>Stock Solutions</b>	5 mM	0.4616 mL	2.3080 mL
		10 mM	0.2308 mL	1.1540 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。			
<b>References</b>	[1]. Budke B, et al. An optimized RAD51 inhibitor that disrupts homologous recombination without requiring Michael acceptor reactivity. J Med Chem. 2013 Jan 10;56(1):254-63.			
实验参考:				
<b>Cell Assay</b>	HEK293 cells are plated into 96-well tissue culture plates at a density of 300 cells per well in the presence or absence of 50 nM mitomycin C (MMC) for 24 hours at 37°C, 5% CO <sub>2</sub> . Media is subsequently replaced with fresh media containing 0.5% DMSO plus RI-2 for an additional 24 hours. RI-2 is then removed, and cultures are allowed to grow to a 50-70% confluence. Average survival from at least three replicates is measured using CellGlo reagent. RI-2 is deemed successful in sensitizing cells to MMC if they generate significantly greater toxicity in the presence of MMC relative to the absence of MMC. Specifically, sensitization is scored as a "+" when non-overlapping standard errors are observed for at least two pairs of compound doses <sup>[1]</sup> .			
<b>References</b>	[1]. Budke B, et al. An optimized RAD51 inhibitor that disrupts homologous recombination without requiring Michael acceptor reactivity. J Med Chem. 2013 Jan 10;56(1):254-63.			