



上海源叶生物科技有限公司
Shanghai yuanye Bio-Technology Co., Ltd
电话: 021-61312973 传真: 021-55068248
网址: www.shyuanye.com
邮箱: shyysw@sina.com

产品名称: RI-2
产品别名: RI-2

生物活性:					
Description	RI-2 is a reversible RAD51 inhibitor, with an IC ₅₀ of 44.17 μM, and specifically inhibits homologous recombination repair in human cells.				
IC ₅₀ & Target	IC50: 44.17 μM (RAD51) ^[1]				
In Vitro	RI-2 (7a) is a reversible RAD51 inhibitor, with an IC ₅₀ of 44.17 μM. RI-2 specifically inhibits homologous recombination repair in human cells. RI-2 (150 μM) induces a significant sensitization of cells ^[1] .				
Solvent&Solubility	In Vitro: DMSO : 15.5 mg/mL (35.77 mM; Need ultrasonic and warming)				
	<div>Preparing Stock Solutions</div>	<div>Solvent / Mass Concentration</div>	1 mg	5 mg	10 mg
		1 mM	2.3080 mL	11.5399 mL	23.0798 mL
		5 mM	0.4616 mL	2.3080 mL	4.6160 mL
		10 mM	0.2308 mL	1.1540 mL	2.3080 mL
	<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>				
References	[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:					
Cell Assay	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 ₂ . 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 reagentor. 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 ^[1] .				
References	[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.				