

Produktinformation



Forschungsprodukte & Biochemikalien



Zellkultur & Verbrauchsmaterial



Diagnostik & molekulare Diagnostik



Laborgeräte & Service

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siehe unsere Liefer- und Versandbedingungen

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Proteins

Product Data Sheet

RI-2

Cat. No.: HY-16904

CAS No.: 1417162-36-7 Molecular Formula: $C_{21}H_{18}Cl_2N_2O_4$

Molecular Weight: 433 Target: RAD51

Pathway: Cell Cycle/DNA Damage

Powder -20°C Storage: 3 years

2 years

-80°C In solvent 2 years

> -20°C 1 year

SOLVENT & SOLUBILITY

In Vitro

DMSO: 130 mg/mL (300.23 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.3095 mL	11.5473 mL	23.0947 mL
	5 mM	0.4619 mL	2.3095 mL	4.6189 mL
	10 mM	0.2309 mL	1.1547 mL	2.3095 mL

Please refer to the solubility information to select the appropriate solvent.

BIOLOGICAL ACTIVITY

Description RI-2 is a reversible RAD51 inhibitor, with an IC $_{50}$ of 44.17 μ M, and specifically inhibits homologous recombination repair in human cells.

IC50: 44.17 μM (RAD51)^[1] IC₅₀ & Target

> RI-2 (7a) is a reversible RAD51 inhibitor, with an IC₅₀ of 44.17 µM. RI-2 specifically inhibits homologous recombination repair in human cells. RI-2 (150 μ M) induces a significant sensitization of cells [1].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

PROTOCOL

In Vitro

Cell Assay [1]

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% at 30°C, 5% CO_2 .

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].

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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.

Caution: Product has not been fully validated for medical applications. For research use only.

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