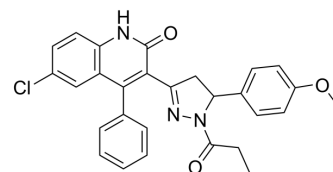


## Homologous recombination-IN-1

Cat. No.:	HY-148183
CAS No.:	391889-85-3
Molecular Formula:	C <sub>28</sub> H <sub>24</sub> ClN <sub>3</sub> O <sub>3</sub>
Molecular Weight:	485.96
Target:	RAD51
Pathway:	Cell Cycle/DNA Damage
Storage:	Powder    -20°C    3 years 4°C    2 years In solvent   -80°C    6 months -20°C    1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 50 mg/mL (102.89 mM; ultrasonic and warming and heat to 60°C)

	Solvent Concentration	Mass	1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM		2.0578 mL	10.2889 mL	20.5778 mL
	5 mM		0.4116 mL	2.0578 mL	4.1156 mL
	10 mM		0.2058 mL	1.0289 mL	2.0578 mL

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

### BIOLOGICAL ACTIVITY

#### Description

Homologous recombination-IN-1 is a novel RAD51-BRCA2 protein-protein interaction inhibitor (EC<sub>50</sub>=19 μM). Homologous recombination-IN-1 can interfere with homologous recombination<sup>[1]</sup>.

#### In Vitro

Homologous recombination-IN-1 (10-40 μM; 5 h) reduces cell homologous recombination in a dose-dependent manner<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Western Blot Analysis<sup>[1]</sup>

Cell Line:	BxPC3 cells
Concentration:	10-40 μM
Incubation Time:	5 hours
Result:	Led to homologous recombination inhibition with an EC <sub>50</sub> value of 18.4 μM.

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## REFERENCES

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[1]. Bagnolini G, et al. Synthetic Lethality in Pancreatic Cancer: Discovery of a New RAD51-BRCA2 Small Molecule Disruptor That Inhibits Homologous Recombination and Synergizes with Olaparib. J Med Chem. 2020 Mar 12;63(5):2588-2619.

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

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