# DB008

Cat. No.:	HY-150221	
Molecular Formula:	$C_{25}H_{21}FN_{4}O_{3}$	0
Molecular Weight:	444.46	NH
Target:	PARP	Ň O
Pathway:	Cell Cycle/DNA Damage; Epigenetics	
Storage:	4°C, sealed storage, away from moisture and light * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)	F C

## SOLVENT & SOLUBILITY

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In Vitro	DMSO : 100 mg/mL (2	DMSO : 100 mg/mL (224.99 mM; Need ultrasonic)					
		Mass Solvent Concentration	1 mg	5 mg	10 mg		
	Preparing Stock Solutions	1 mM	2.2499 mL	11.2496 mL	22.4992 mL		
		5 mM	0.4500 mL	2.2499 mL	4.4998 mL		
		10 mM	0.2250 mL	1.1250 mL	2.2499 mL		
	Please refer to the so	lubility information to select the app	propriate solvent.				
In Vivo		1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 5 mg/mL (11.25 mM); Clear solution					
		2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 5 mg/mL (11.25 mM); Clear solution					
		3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 5 mg/mL (11.25 mM); Clear solution					

BIOLOGICAL ACTIVITY			
Description	DB008 is potent and selective PARP16 inhibitor with an IC <sub>50</sub> value of 0.27 μM, containing an acrylamide electrophilic reagent. DB008 is membrane-permeable and marks PARP16 selectively <sup>[1]</sup> . DB008 is a click chemistry reagent, it contains an Alkyne group and can undergo copper-catalyzed azide-alkyne cycloaddition (CuAAc) with molecules containing Azide groups.		
IC₅₀ & Target	PARP16 0.27 μM (IC <sub>50</sub> )		
In Vitro	DB008 (0-1µM; 0-120 min; HEK 293T cells) covalently modifies Cys169 of PARP16 and exhibits excellent proteome-wide selectivity in the irreversible binding mode <sup>[1]</sup> .		

Product Data Sheet

DB008 (100 nM; 16 h; HAP1 WT and HAP1 PARP16 KO cells) rescues nutrient starvation-induced loss of soluble PARP16<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## REFERENCES

[1]. Bejan DS, et, al. Structure-guided design and characterization of a clickable, covalent PARP16 inhibitor. Chem Sci. 2022 Nov 16;13(46):13898-13906.

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

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