PROTAC GPX4 degrader-1

MedChemExpress

Cat. No.:	HY-149236	
CAS No.:	2916433-81-1	
Molecular Formula:	C ₅₀ H ₅₇ ClN ₁₀ O ₁₀	
Molecular Weight:	993.5	
Target:	Glutathione Peroxidase; Ferroptosis; PROTACs	
Pathway:	Apoptosis; Metabolic Enzyme/Protease; PROTAC	
Storage:	4°C, stored under nitrogen * In solvent : -80°C, 6 months; -20°C, 1 month (stored under nitrogen)	

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (100.65 mM; Need ultrasonic)						
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg		
		1 mM	1.0065 mL	5.0327 mL	10.0654 mL		
		5 mM	0.2013 mL	1.0065 mL	2.0131 mL		
		10 mM	0.1007 mL	0.5033 mL	1.0065 mL		
	Please refer to the solubility information to select the appropriate solvent.						
In Vivo	 Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (2.52 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (2.52 mM); Clear solution 						

BIOLOGICAL ACTIVITY						
BIOLOGICAL ACTIVITY						
Description	PROTAC GPX4 degrader-1 (DC-2) is a PROTAC-based GPX4 degrader, with a DC $_{50}$ of 0.03 μ M in HT1080 cells ^[1] .					
In Vitro	PROTAC GPX4 degrader-1 (DC-2) has been found to induce GPX4 degradation and subsequent ferroptosis ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Western Blot Analysis ^[1] .					
	Cell Line: Concentration:	HT1080 cells. 0.003-1 μM.				
	Incubation Time:	24 h.				

Product Data Sheet

Result:

REFERENCES

[1]. Han Wang, et al. Discovery of ML210-Based glutathione peroxidase 4 (GPX4) degrader inducing ferroptosis of human cancer cells. Eur J Med Chem. 2023 Jun 5;254:115343.

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

 Tel: 609-228-6898
 Fax: 609-228-5909
 E-mail: tech@MedChemExpress.com

 Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA