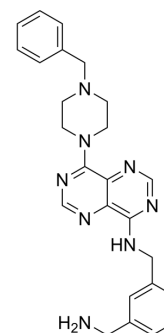


BI8626

Cat. No.:	HY-120204		
CAS No.:	1875036-75-1		
Molecular Formula:	C ₂₅ H ₂₈ N ₈		
Molecular Weight:	440.54		
Target:	E1/E2/E3 Enzyme		
Pathway:	Metabolic Enzyme/Protease		
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 : 83.33 mg/mL (189.15 mM; Need ultrasonic)					
		Solvent Concentration	Mass	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM		2.2699 mL	11.3497 mL	22.6994 mL
		5 mM		0.4540 mL	2.2699 mL	4.5399 mL
10 mM			0.2270 mL	1.1350 mL	2.2699 mL	
Please refer to the solubility information to select the appropriate solvent.						
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.72 mM); Clear solution 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.72 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	BI8626 is a specific inhibitor of the ubiquitin ligase HUWE1 with an IC ₅₀ of 0.9 μM ^[1] .
IC₅₀ & Target	IC ₅₀ : 0.9 μM (HUWE1) ^[1]
In Vitro	BI8626 induces HUWE1 ectopically expresses to abolish ubiquitination of MCL1 in HeLa cells ^[1] . ?BI8626 suppresses colony formation of Ls174T cells with estimated IC ₅₀ value of 0.7 μM, and BI8622 (1-4 days) treatment retards passage of Ls174T cells through all phases of the cell cycle, with the effect being strongest for G1 ^[1] . ?BI8626 (0-50 μM; 0-6 hours) retards the degradation of MCL1 in response to UV irradiation to the same extent as depletion of HUWE1 in U2OS cells ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Cell Cycle Analysis^[1]

Cell Line:	Ls174T cells
Concentration:	0 μ M, 5 μ M, 10 μ M, 15 μ M, 20 μ M
Incubation Time:	0-4 days
Result:	Retarded passage of Ls174T cells through all phases of the cell cycle, with the effect being strongest for G1.

Western Blot Analysis^[1]

Cell Line:	U2OS cells
Concentration:	0 μ M, 20 μ M, 50 μ M
Incubation Time:	0 hour, 1 hour, 2 hours, 4 hours, 6 hours
Result:	Retarded the degradation of MCL1 in response to UV irradiation in HeLa cells by inhibiting HUWE1 in U2OS cells.

REFERENCES

[1]. Peter S, et al. Tumor cell-specific inhibition of MYC function using small molecule inhibitors of the HUWE1 ubiquitin ligase. *EMBO Mol Med.* 2014 Dec;6(12):1525-41.

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

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