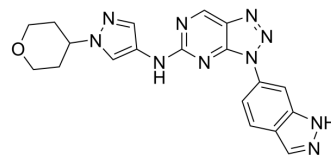


GCN2-IN-1

Cat. No.:	HY-100877		
CAS No.:	1448693-69-3		
Molecular Formula:	C ₁₉ H ₁₈ N ₁₀ O		
Molecular Weight:	402		
Target:	Eukaryotic Initiation Factor (eIF)		
Pathway:	Cell Cycle/DNA Damage		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



SOLVENT & SOLUBILITY

In Vitro	DMSO : 60 mg/mL (149.25 mM; Need ultrasonic and warming)					
		Solvent Concentration	Mass	1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM		2.4876 mL	12.4378 mL	24.8756 mL
		5 mM		0.4975 mL	2.4876 mL	4.9751 mL
10 mM			0.2488 mL	1.2438 mL	2.4876 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: ≥ 2.5 mg/mL (6.22 mM); Clear solution					

BIOLOGICAL ACTIVITY

Description	GCN2-IN-1 (A-92) is a potent general control nonderepressible 2 kinase (GCN2) inhibitor with an IC ₅₀ of <0.3 μM in the enzyme assay and an IC ₅₀ of 0.3-3 μM in the cell assay ^[1] .
IC ₅₀ & Target	eIF2
In Vitro	GCN2-IN-1 (Compound A-92) is a GCN2 inhibitor, which may be useful as a chemotherapeutic drug for the treatment of cancer ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Cell Discov. 2021 Oct 26;7(1):98.
- Nat Commun. 2023 Oct 25;14(1):6777.
- Nat Commun. 2023 Sep 8;14(1):5535.
- Mol Cell. 2023 Nov 2:S1097-2765(23)00849-3.
- Nucleic Acids Res. 2024 Feb 16:gkae119.

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REFERENCES

[1]. Dieter Dorsch, et al. Triazolo[4,5-d]pyrimidine derivatives. WO2013110309A1.

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

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