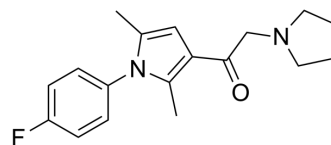


IU1

Cat. No.:	HY-13817		
CAS No.:	314245-33-5		
Molecular Formula:	C ₁₈ H ₂₁ FN ₂ O		
Molecular Weight:	300.37		
Target:	Deubiquitinase; Autophagy		
Pathway:	Cell Cycle/DNA Damage; Autophagy		
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 : 41.67 mg/mL (138.73 mM; Need ultrasonic)				
		Solvent Concentration	Mass 1 mg	5 mg	10 mg
	Preparing Stock Solutions	1 mM	3.3292 mL	16.6461 mL	33.2923 mL
		5 mM	0.6658 mL	3.3292 mL	6.6585 mL
10 mM		0.3329 mL	1.6646 mL	3.3292 mL	
Please refer to the solubility information to select the appropriate solvent.					
In Vivo	<ol style="list-style-type: none"> Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 1.67 mg/mL (5.56 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 1.67 mg/mL (5.56 mM); Clear solution Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1.67 mg/mL (5.56 mM); Clear solution 				

BIOLOGICAL ACTIVITY

Description	IU1 is a special Usp14 inhibitor with an IC ₅₀ of 4-5 μM.
IC₅₀ & Target	IC ₅₀ : 4-5 μM (Usp14) ^[1]
In Vitro	Usp14, a proteasome-associated deubiquitinating enzyme, can inhibit the degradation of ubiquitin-protein conjugates, in vivo and in vitro. IU1 inhibits the catalytic activity of proteasome associated Usp14 in vitro. The IC ₅₀ of IU1 for Usp14 is 4-5 μM. IU1 binds specifically to the activated form of Usp14. IU1 can potentially inhibit Usp14 by preventing its docking on the

proteasome, but direct tests of this scenario proved negative. Usp14 inhibition is rapidly established upon addition of IU1 and rapidly reversed upon its removal. IU1 also promotes degradation of Sic1, a CDK inhibitor from *S. cerevisiae*^[1]. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

CUSTOMER VALIDATION

- Nat Commun. 2022 Mar 31;13(1):1700.
- Cell Death Differ. 2022 Jul 29.
- EMBO J. 2022 Jul 11;e108791.
- Pharmacol Res. 8 October 2021, 105933.
- Front Oncol. 2021 Feb 23;11:615568.

See more customer validations on www.MedChemExpress.com

REFERENCES

[1]. Lee BH, et al. Enhancement of proteasome activity by a small-molecule inhibitor of USP14. Nature. 2010 Sep 9;467(7312):179-84.

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

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