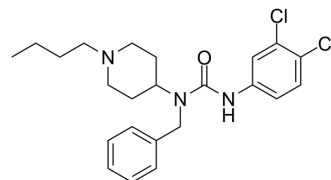


## NACM-OPT

<b>Cat. No.:</b>	HY-111505		
<b>CAS No.:</b>	2089293-61-6		
<b>Molecular Formula:</b>	C <sub>23</sub> H <sub>29</sub> Cl <sub>2</sub> N <sub>3</sub> O		
<b>Molecular Weight:</b>	434.4		
<b>Target:</b>	E1/E2/E3 Enzyme; NEDD8-activating Enzyme		
<b>Pathway:</b>	Metabolic Enzyme/Protease		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



## SOLVENT & SOLUBILITY

<b>In Vitro</b>	DMSO : 125 mg/mL (287.75 mM; Need ultrasonic)			
		Solvent Concentration	Mass	
			1 mg	5 mg
			10 mg	
	<b>Preparing Stock Solutions</b>	1 mM	2.3020 mL	11.5101 mL
	5 mM	0.4604 mL	2.3020 mL	4.6041 mL
	10 mM	0.2302 mL	1.1510 mL	2.3020 mL
Please refer to the solubility information to select the appropriate solvent.				
<b>In Vivo</b>	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.08 mg/mL (4.79 mM); Clear solution  2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.08 mg/mL (4.79 mM); Clear solution			

## BIOLOGICAL ACTIVITY

<b>Description</b>	NACM-OPT is an orally bioavailable cullin neddylation 1 (DCN1) inhibitor, which potently inhibits the DCN1-UBE2M interaction <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	DCN1 <sup>[1]</sup>
<b>In Vitro</b>	NACM-OPT (Compound 67) is orally bioavailable, well tolerated in mice, and currently used to study the effects of acute pharmacologic inhibition of the DCN1-UBE2M interaction on the NEDD8/CUL pathway <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

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## REFERENCES

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[1]. Hammill JT, et al. Discovery of an Orally Bioavailable Inhibitor of Defective in Cullin Neddylation 1 (DCN1)-Mediated Cullin Neddylation. J Med Chem. 2018 Apr 12;61(7):2694-2706.

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

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