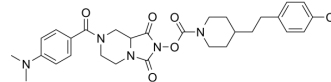


## JJH260

<b>Cat. No.:</b>	HY-116895		
<b>CAS No.:</b>	1831135-30-8		
<b>Molecular Formula:</b>	C <sub>29</sub> H <sub>34</sub> ClN <sub>5</sub> O <sub>5</sub>		
<b>Molecular Weight:</b>	568.06		
<b>Target:</b>	Androgen Receptor; MAGL		
<b>Pathway:</b>	Vitamin D Related/Nuclear Receptor; Metabolic Enzyme/Protease		
<b>Storage:</b>	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 6.25 mg/mL (11.00 mM; Need ultrasonic)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	1.7604 mL	8.8019 mL	17.6038 mL
5 mM	0.3521 mL	1.7604 mL	3.5208 mL
10 mM	0.1760 mL	0.8802 mL	1.7604 mL

Please refer to the solubility information to select the appropriate solvent.

### BIOLOGICAL ACTIVITY

#### Description

JJH260 is AIG1 inhibitor, and inhibit the fluorophosphonate reactivity and fatty acid esters of hydroxy fatty acid (FAHFA) hydrolysis activity of AIG1 in HEK293T cells, with IC<sub>50</sub> values of 0.50 μM and 0.57 μM, respectively<sup>[1]</sup>.

#### IC<sub>50</sub> & Target

0.50 μM (AIG1) and 0.57 μM (FAHFA) in HEK293T cells<sup>[1]</sup>

#### In Vitro

JJH260 (30 min) inhibits ADTRP, FP-Rh labeling of hAIG1, and the 9-PAHSA hydrolysis activity of hAIG1, in HEK293T, with the IC<sub>50</sub> of 0.5 and 0.57 μM<sup>[1]</sup>.

JJH260 (5 μM, 4 h) inhibits FAHFA hydrolysis in LNCaP cells and human T-cells<sup>[1]</sup>.

MCE has not independently confirmed the accuracy of these methods. They are for reference only.

### REFERENCES

[1]. William H Parsons, et al. AIG1 and ADTRP are atypical integral membrane hydrolases that degrade bioactive FAHFAs. Nat Chem Biol. 2016 May;12(5):367-372.

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

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