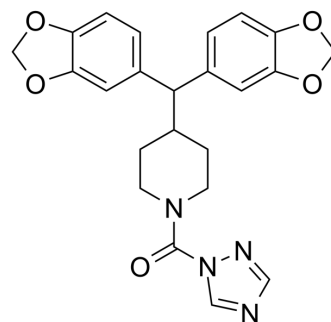


## JJKK 048

Cat. No.:	HY-108613		
CAS No.:	1515855-97-6		
Molecular Formula:	C <sub>23</sub> H <sub>22</sub> N <sub>4</sub> O <sub>5</sub>		
Molecular Weight:	434.44		
Target:	MAGL		
Pathway:	Metabolic Enzyme/Protease		
Storage:	Powder	-20°C	3 years
	In solvent	-80°C	6 months
		-20°C	1 month



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 33.33 mg/mL (76.72 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent	Mass	1 mg	5 mg	10 mg
	Concentration				
	1 mM		2.3018 mL	11.5091 mL	23.0181 mL
	5 mM		0.4604 mL	2.3018 mL	4.6036 mL
	10 mM		0.2302 mL	1.1509 mL	2.3018 mL

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

#### In Vivo

- Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline  
Solubility: ≥ 2.5 mg/mL (5.75 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline)  
Solubility: ≥ 2.5 mg/mL (5.75 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil  
Solubility: ≥ 2.5 mg/mL (5.75 mM); Clear solution

### BIOLOGICAL ACTIVITY

#### Description

JJKK 048 is an ultrapotent and highly selective inhibitor of Monoacylglycerol lipase (MAGL).

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

MAGL<sup>[1]</sup>

### REFERENCES

[1]. Aaltonen N, et al. In Vivo Characterization of the Ultrapotent Monoacylglycerol Lipase Inhibitor {4-[bis-(benzo[d][1,3]dioxol-5-yl)methyl]piperidin-1-yl}(1H-1,2,4-triazol-

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1-yl)methanone (JJKK-048). J Pharmacol Exp Ther. 2016 Oct;359(1):62-72.

[2]. Aaltonen N, et al. Piperazine and piperidine triazole ureas as ultrapotent and highly selective inhibitors of monoacylglycerol lipase. Chem Biol. 2013 Mar 21;20(3):379-90.

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

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