## ARUK2001607

Cat. No.: HY-152155 CAS No.: 2924824-56-4 Molecular Formula:  $C_{14}H_{13}N_3O_2S_2$ 

Molecular Weight: 319.4 PI5P4K Target:

Pathway: Metabolic Enzyme/Protease

Storage: Powder -20°C 3 years  $4^{\circ}C$ 2 years

> -80°C 6 months In solvent

> > -20°C 1 month

## **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 25 mg/mL (78.27 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	3.1309 mL	15.6544 mL	31.3087 mL
	5 mM	0.6262 mL	3.1309 mL	6.2617 mL
	10 mM	0.3131 mL	1.5654 mL	3.1309 mL

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

## **BIOLOGICAL ACTIVITY**

Description ARUK2001607 is a selective phosphatidylinositol 5-phosphate 4-kinase  $\gamma$  (PI5P4K $\gamma$ ) inhibitor with a K<sub>d</sub> of 7.1 nM.

ARUK2001607 shows high selectivity over other >150 kinases<sup>[1]</sup>.

IC<sub>50</sub> & Target Kd: 7.1 nM (PI5P4Kγ)<sup>[1]</sup>

In Vivo Mouse Pharmacokinetics Parameters for Selected ARUK2001607 (15) (5 mg/kg; ip Administration)  $^{[1]}$ In Vivo

	T <sub>1/2</sub> (h)	AUC (hr·ng/mL)	[plasma] at 0.5h (ng/mL)	[brain] at 0.5h (ng/g)	K <sub>p</sub>
ARUK2001607 (15)	0.74	4520	2773	1590	0.58

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

REFERENCES					
[1]. Timothy P C Rooney, et al. The Identification of Potent, Selective, and Brain Penetrant PI5P4Ky Inhibitors as In Vivo-Ready Tool Molecules. J Med Chem. 2022 Dec 14.					
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