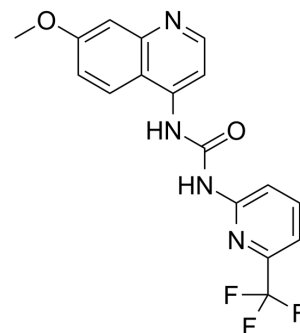


## A 1070722

<b>Cat. No.:</b>	HY-107531		
<b>CAS No.:</b>	1384424-80-9		
<b>Molecular Formula:</b>	C <sub>17</sub> H <sub>13</sub> F <sub>3</sub> N <sub>4</sub> O <sub>2</sub>		
<b>Molecular Weight:</b>	362.31		
<b>Target:</b>	GSK-3		
<b>Pathway:</b>	PI3K/Akt/mTOR; Stem Cell/Wnt		
<b>Storage:</b>	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year



### SOLVENT & SOLUBILITY

#### In Vitro

DMSO : 83.33 mg/mL (230.00 mM; Need ultrasonic)

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.7601 mL	13.8003 mL	27.6007 mL
	5 mM	0.5520 mL	2.7601 mL	5.5201 mL
	10 mM	0.2760 mL	1.3800 mL	2.7601 mL

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

### BIOLOGICAL ACTIVITY

#### Description

A 1070722 is a potent and selective glycogen synthase kinase 3 (GSK-3) inhibitor, with a K<sub>i</sub> of 0.6 nM for both GSK-3α and GSK-3β. A 1070722 can penetrate the blood-brain barrier (BBB) and accumulates in brain regions, thus potential for PET radiotracer for the quantification of GSK-3 in brain<sup>[1]</sup>.

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

GSK-3α	GSK-3β
0.6 nM (K <sub>i</sub> )	0.6 nM (K <sub>i</sub> )

### REFERENCES

[1]. Prabhakaran J, et al. Radiosynthesis and in Vivo Evaluation of [11C]A1070722, a High Affinity GSK-3 PET Tracer in Primate Brain. ACS Chem Neurosci. 2017 Aug 16;8(8):1697-1703.

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

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