## **Product** Data Sheet

# 3-Methyladenine-d<sub>3</sub>

Cat. No.: HY-19312S CAS No.: 110953-39-4 Molecular Formula:  $C_6H_4D_3N_5$ 

Molecular Weight: 152.17

Target: PI3K; Autophagy; Mitophagy; Endogenous Metabolite

Pathway: PI3K/Akt/mTOR; Autophagy; Metabolic Enzyme/Protease

Storage: 4°C, sealed storage, away from moisture and light

\* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)

### **SOLVENT & SOLUBILITY**

In Vitro

DMSO: 41.67 mg/mL (273.84 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	6.5716 mL	32.8580 mL	65.7160 mL
	5 mM	1.3143 mL	6.5716 mL	13.1432 mL
	10 mM	0.6572 mL	3.2858 mL	6.5716 mL

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

### **BIOLOGICAL ACTIVITY**

Description	3-Methyladenine-d <sub>3</sub> is the deuterium labeled 3-Methyladenine[1]. 3-Methyladenine (3-MA) is a PI3K inhibitor. 3-Methyladenine is a widely used inhibitor of autophagy via its inhibitory effect on class III PI3K[2].
In Vitro	Stable heavy isotopes of hydrogen, carbon, and other elements have been incorporated into drug molecules, largely as tracers for quantitation during the drug development process. Deuteration has gained attention because of its potential to affect the pharmacokinetic and metabolic profiles of drugs <sup>[1]</sup> .  MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### **REFERENCES**

- $[1]. \ Russak\ EM, et\ al.\ Impact\ of\ Deuterium\ Substitution\ on\ the\ Pharmacokinetics\ of\ Pharmaceuticals.\ Ann\ Pharmacother.\ 2019\ Feb; 53(2): 211-216.$
- [2]. Miller S, et al. Finding a fitting shoe for Cinderella: searching for an autophagy inhibitor. Autophagy. 2010 Aug;6(6):805-7.
- [3]. Hou H, et al. Inhibitors of phosphatidylinositol 3'-kinases promote mitotic cell death in HeLa cells. PLoS One. 20127(4):e35665.

4]. Wang X, et al. Acanthopana Rats.Mediators Inflamm. 20162		eliorates Sodium Taurocholate-I	nduced Severe Acute Pancreatitis by Inhibiti	ng the Autophagic Pathway in
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Page 2 of 2 www.MedChemExpress.com