**Proteins** 

# **Product** Data Sheet

## L-Glutamic acid-d<sub>5</sub>

Cat. No.: HY-14608S7 CAS No.: 2784-50-1 Molecular Formula:  $C_5H_4D_5NO_4$ Molecular Weight: 152.16

Target: Apoptosis; iGluR; Ferroptosis; Endogenous Metabolite

Pathway: Apoptosis; Membrane Transporter/Ion Channel; Neuronal Signaling; Metabolic

Enzyme/Protease

Storage: Powder -20°C 3 years

4°C 2 years -80°C 6 months

In solvent -20°C 1 month

### **SOLVENT & SOLUBILITY**

In Vitro

H<sub>2</sub>O: 14.29 mg/mL (93.91 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	6.5720 mL	32.8601 mL	65.7203 mL
	5 mM	1.3144 mL	6.5720 mL	13.1441 mL
	10 mM	0.6572 mL	3.2860 mL	6.5720 mL

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

In Vivo

1. Add each solvent one by one: PBS

Solubility: 7.14 mg/mL (46.92 mM); Clear solution; Need ultrasonic and warming and heat to 60°C

### **BIOLOGICAL ACTIVITY**

Description L-Glutamic acid-d<sub>5</sub> is the deuterium labeled L-Glutamic acid. L-Glutamic acid acts as an excitatory transmitter and an

agonist at all subtypes of glutamate receptors (metabotropic, kainate, NMDA, and AMPA). L-Glutamic acid shows a direct

activating effect on the release of DA from dopaminergic terminals.

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[1].

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

#### **REFERENCES**

	uticals. Ann Pharmacother. 2019;53(2):211-216.	
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