## **Product** Data Sheet

## Valproic acid-d<sub>15</sub>

Cat. No.: HY-10585S2 CAS No.: 362049-65-8 Molecular Formula:  $C_8HD_{15}O_2$  Molecular Weight: 159.3

Target: Notch; Autophagy; HDAC; HIV; Mitophagy; Endogenous Metabolite

Pathway: Neuronal Signaling; Stem Cell/Wnt; Autophagy; Cell Cycle/DNA Damage; Epigenetics;

Anti-infection; Metabolic Enzyme/Protease

Storage: Pure form -20°C 3 years

In solvent

4°C 2 years -80°C 6 months

-20°C 1 month

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## **BIOLOGICAL ACTIVITY**

Description	Valproic acid-d <sub>15</sub> is the deuterium labeled Valproic acid. Valproic acid (VPA; 2-Propylpentanoic Acid) is an HDAC inhibitor, with IC50 in the range of 0.5 and 2 mM, also inhibits HDAC1 (IC50, 400 μM), and induces proteasomal degradation of HDAC2. Valproic acid activates Notch1 signaling and inhibits proliferation in small cell lung cancer (SCLC) cells. Valproic acid sodium salt is used in the treatment of epilepsy, bipolar disorder and prevention of migraine headaches[1][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**

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- [3]. Zhang ZH, et al. Valproic acid inhibits tumor angiogenesis in mice transplanted with Kasumi 1 leukemia cells. Mol Med Rep. 2013 Nov 28.
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 $\label{lem:caution:Product} \textbf{Caution: Product has not been fully validated for medical applications. For research use only.}$ 

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