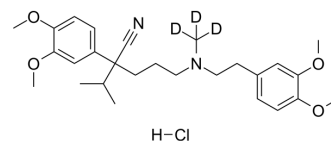


Verapamil-d₃ hydrochloride

Cat. No.:	HY-A0064S
CAS No.:	2714485-49-9
Molecular Formula:	C ₂₇ H ₃₆ D ₃ ClN ₂ O ₄
Molecular Weight:	494.08
Target:	P-glycoprotein; Calcium Channel; Cytochrome P450; Isotope-Labeled Compounds
Pathway:	Membrane Transporter/Ion Channel; Neuronal Signaling; Metabolic Enzyme/Protease; Others
Storage:	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



BIOLOGICAL ACTIVITY

Description	Verapamil-d ₃ (hydrochloride) is the deuterium labeled Verapamil hydrochloride. Verapamil hydrochloride ((±)-Verapamil hydrochloride) is a calcium channel blocker and a potent and orally active first-generation P-glycoprotein (P-gp) inhibitor. Verapamil hydrochloride also inhibits CYP3A4. Verapamil hydrochloride has the potential for high blood pressure, heart arrhythmias and angina research[1][2][3].
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

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- [2]. Krikler DM. Verapamil in arrhythmia. *Br J Clin Pharmacol.* 1986;21 Suppl 2:183S-189S.
- [3]. Rehnqvist N, et al. Effects of metoprolol vs verapamil in patients with stable angina pectoris. The Angina Prognosis Study in Stockholm (APSIS). *Eur Heart J.* 1996 Jan;17(1):76-81.
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Caution: Product has not been fully validated for medical applications. For research use only.

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