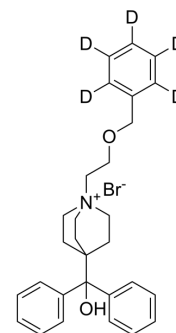


## Umeclidinium-d<sub>5</sub> bromide

<b>Cat. No.:</b>	HY-12100S
<b>Molecular Formula:</b>	C <sub>29</sub> H <sub>29</sub> D <sub>5</sub> BrNO <sub>2</sub>
<b>Molecular Weight:</b>	513.52
<b>Target:</b>	mAChR; Isotope-Labeled Compounds
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling; Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Umeclidinium-d <sub>5</sub> (bromide) is the deuterium labeled Umeclidinium bromide. Umeclidinium bromide is a novel mAChR antagonist. The affinity (K <sub>i</sub> ) of Umeclidinium bromide for the cloned human M1-M5 mAChRs ranges from 0.05 to 0.16 nM.	
<b>IC<sub>50</sub> &amp; Target</b>	mAChR1	mAChR5
<b>In Vitro</b>	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;53(2):211-216.
- [2]. Salmon M, et al. Pharmacological characterization of GSK573719 (umeclidinium): a novel, long-acting, inhaled antagonist of the muscarinic cholinergic receptors for treatment of pulmonary diseases. *J Pharmacol Exp Ther.* 2013 May;345(2):260-70.
- [3]. Cazzola M, et al. Pharmacology and therapeutics of bronchodilators. *Pharmacol Rev.* 2012 Jul;64(3):450-504.
- [4]. Calzetta L, et al. Pharmacological characterization of the interaction between umeclidinium and vilanterol in human bronchi. *Eur J Pharmacol.* 2017 Jul 14. pii: S0014-2999(17)30470-3.

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

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