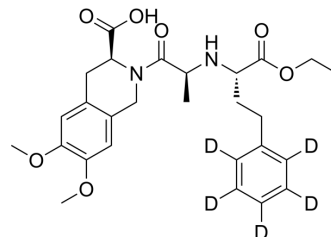


## Moexipril-d<sub>5</sub>

<b>Cat. No.:</b>	HY-117281S
<b>CAS No.:</b>	1356929-49-1
<b>Molecular Formula:</b>	C <sub>27</sub> H <sub>29</sub> D <sub>5</sub> N <sub>2</sub> O <sub>7</sub>
<b>Molecular Weight:</b>	503.6
<b>Target:</b>	Angiotensin-converting Enzyme (ACE); Apoptosis; Isotope-Labeled Compounds
<b>Pathway:</b>	Metabolic Enzyme/Protease; Apoptosis; Others
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.



### BIOLOGICAL ACTIVITY

<b>Description</b>	Moexipril-d <sub>5</sub> is the deuterium labeled Moexipril. Moexipril hydrochloride is a potent orally active non-sulfhydryl angiotensin converting enzyme(ACE) inhibitor, which is used for the treatment of hypertension and congestive heart failure[1][2].
<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]. Chrysant, S.G. and G.S. Chrysant, Pharmacological and clinical profile of moexipril: a concise review. *J Clin Pharmacol*, 2004. 44(8): p. 827-36.
- [3]. Edling, O., et al., Moexipril, a new angiotensin-converting enzyme (ACE) inhibitor: pharmacological characterization and comparison with enalapril. *J Pharmacol Exp Ther*, 1995. 275(2): p. 854-63.
- [4]. Song, J.C. and C.M. White, Clinical pharmacokinetics and selective pharmacodynamics of new angiotensin converting enzyme inhibitors: an update. *Clin Pharmacokinet*, 2002. 41(3): p. 207-24.

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

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