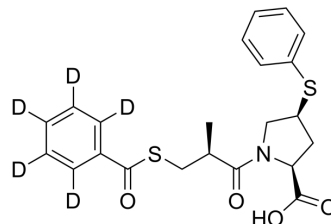


Zofenopril-d₅

Cat. No.:	HY-108321S		
Molecular Formula:	C ₂₂ H ₁₈ D ₅ NO ₄ S ₂		
Molecular Weight:	434.58		
Target:	Angiotensin-converting Enzyme (ACE); Isotope-Labeled Compounds		
Pathway:	Metabolic Enzyme/Protease; Others		
Storage:	Pure form	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	6 months
		-20°C	1 month



BIOLOGICAL ACTIVITY

Description	Zofenopril-d ₅ is deuterium labeled Zofenopril. Zofenopril is an angiotensin-converting enzyme (ACE) inhibitor with an IC ₅₀ of 81 μM.
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

- [1]. Lin CJ, et al. Competitive inhibition of glycylsarcosine transport by enalapril in rabbit renal brush border membrane vesicles: interaction of ACE inhibitors with high-affinity H⁺/peptide symporter. *Pharm Res.* 1999 May;16(5):609-15.
- [2]. Sarro GD, et al. Fosinopril and zofenopril, two angiotensin-converting enzyme (ACE) inhibitors, potentiate the anticonvulsant activity of antiepileptic drugs against audiogenic seizures in DBA/2 mice. *Pharmacol Res.* 2012 Mar;65(3):285-96.
- [3]. Russak EM, et al. Impact of Deuterium Substitution on the Pharmacokinetics of Pharmaceuticals. *Ann Pharmacother.* 2019;53(2):211-216.

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

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