Inhibitors

Proteins

Telmisartan-13C,d₃

Cat. No.: HY-13955S2 CAS No.: 1261396-33-1 Molecular Formula: $C_{32}^{13}CH_{27}D_3N_4O_2$

Molecular Weight: 518.63

Target: Autophagy; Angiotensin Receptor; Isotope-Labeled Compounds

Pathway: Autophagy; GPCR/G Protein; Others

Please store the product under the recommended conditions in the Certificate of Storage:

Analysis.

BIOLOGICAL ACTIVITY

Description	Telmisartan- 13 C,d ₃ is the 13 C- and deuterium labeled Telmisartan. Telmisartan is a potent, long lasting antagonist of angiotensin II type 1 receptor (AT1), selectively inhibiting the binding of 125I-AngII to AT1 receptors with IC50 of 9.2 nM.
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 ^[55] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

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- [3]. Fujihara S, et al. The angiotensin II type 1 receptor antagonist telmisartan inhibits cell proliferation and tumor growth of esophageal adenocarcinoma via the AMPKq/ mTOR pathway in vitro and in vivo. Oncotarget. 2017 Jan 31;8(5):8536-8549.
- [4]. Maillard MP, et al. In vitro and in vivo characterization of the activity of telmisartan: an insurmountable angiotensin II receptor antagonist. J Pharmacol Exp Ther. 2002 Sep;302(3):1089-95.
- [5]. Torika N, et al. Intranasal telmisartan ameliorates brain pathology in five familial Alzheimer's disease mice. Brain Behav Immun. 2017 Apr 3.
- [6]. Xuan H, et al. Inhibition or deletion of angiotensin II type 1 receptor suppresses elastase-induced experimental abdominal aortic aneurysms. J Vasc Surg. 2017 Apr 20. pii: S0741-5214(17)30100-3.

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

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