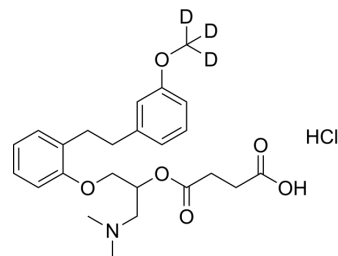


Sarpogrelate-d₃ hydrochloride

Cat. No.:	HY-10564S
Molecular Formula:	C ₂₄ H ₂₉ D ₃ ClNO ₆
Molecular Weight:	468.99
Target:	5-HT Receptor; Isotope-Labeled Compounds
Pathway:	GPCR/G Protein; Neuronal Signaling; Others
Storage:	Please store the product under the recommended conditions in the Certificate of Analysis.



BIOLOGICAL ACTIVITY

Description	Sarpogrelate-d ₃ (hydrochloride) is the deuterium labeled Sarpogrelate hydrochloride. Sarpogrelate hydrochloride (MCI-9042) is a selective 5-HT _{2R} antagonist, with pK _i s of 8.52, 6.57, and 7.43 for 5-HT _{2A} , 5-HT _{2B} , and 5-HT _{2C} receptors, respectively. Sarpogrelate hydrochloride displays selectivity over 5-HT ₁ , 5-HT ₃ , 5-HT ₄ , α ₁ -, α ₂ - and β-adrenoreceptor, histamine H ₁ , H ₂ and muscarinic M ₃ receptors. Sarpogrelate hydrochloride can be used for the research of vascular disease associated with thrombosis ^{[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

- [1]. Rashid M, et, al. Identification of the binding sites and selectivity of sarpogrelate, a novel 5-HT₂ antagonist, to human 5-HT_{2A}, 5-HT_{2B} and 5-HT_{2C} receptor subtypes by molecular modeling. *Life Sci.* 2003 May 30;73(2):193-207.
- [2]. Maruyama K, et, al. MCI-9042: high affinity for serotonergic receptors as assessed by radioligand binding assay. *J Pharmacobiodyn.* 1991 Apr;14(4):177-81.

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

Tel: 609-228-6898

Fax: 609-228-5909

E-mail: tech@MedChemExpress.com

Address: 1 Deer Park Dr, Suite Q, Monmouth Junction, NJ 08852, USA