(S)-Terazosin

Cat. No.:	HY-B0371D		
CAS No.:	109351-33-9	9	
Molecular Formula:	C ₁₉ H ₂₅ N ₅ O	4	
Molecular Weight:	387.43		
Target:	Adrenergic Receptor		
Pathway:	GPCR/G Protein; Neuronal Signaling		
Storage:	Powder	-20°C	3 years
		4°C	2 years
	In solvent	-80°C	2 years
		-20°C	1 year

®

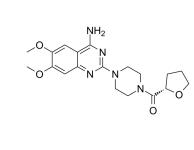
MedChemExpress

SOLVENT & SOLUBILITY

	Mass Solvent Concentration	1 mg	5 mg	10 mg
Preparing Stock Soluti	1 mM	2.5811 mL	12.9056 mL	25.8111 mL
	5 mM	0.5162 mL	2.5811 mL	5.1622 mL
	10 mM	0.2581 mL	1.2906 mL	2.5811 mL

BIOLOGICAL ACTIVITY			
Description	(S)-Terazosin is an active S-enantiomer of Terazosin. (S)-Terazosin is a potent and high-affinity α-adrenoceptor antagonist with K _i values of 3.91 nM, 0.79 nM and 1.16 nM for α1a, α1b and α1d-adrenoceptor, respectively. (S)-Terazosin also has high-affinity for α2a, α2B and α2c-adrenoceptor with K _i values of 729 nM, 3.5 nM and 46.4 nM, respectively ^[1] .		
IC ₅₀ & Target	Ki: 3.91 nM (α1a-adrenoceptor), 0.79 nM (α1b-adrenoceptor) and 1.16 nM (α1d-adrenoceptor); 729 nM (α2a-adrenoceptor), 3.5 nM (α2Ba-adrenoceptor) and 46.4 nM (α2c-adrenoceptor) ^[1]		
In Vitro	The racemic compound and its enantiomers show high and apparently equal affinity for subtypes of α 1-adrenoceptors with K_i values in the low nanomolar range, and showed potent antagonism of α 1-adrenoceptors in isolated tissues, with the enantiomers approximately equipotent to the racemate at each α 1-adrenoceptor subtype. At α 2b sites, (R)-Terazosin binds less potently than either the (S)-Terazosin or racemate ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.		
In Vivo	(S)-Terazosin shows antagonism of at rat atrial α2B receptor with a pEC30 of 6.93. (s)-Terazosin shows antagonism of at rat vas deferens α1A and α2A receptor with pA2 values of 8.3 and 6.12, respectively ^[1] .		

Product Data Sheet



MCE has not independently confirmed the accuracy of these methods. They are for reference only.

REFERENCES

[1]. Hancock AA, et al. Actions of terazosin and its enantiomers at subtypes of alpha 1- and alpha 2-adrenoceptors in vitro. J Recept Signal Transduct Res. 1995 Sep-Dec;15(7-8):863-85.

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