Product Data Sheet



(R)-Terazosin

Cat. No.: HY-B0371B CAS No.: 109351-34-0 Molecular Formula: $C_{19}H_{25}N_5O_4$ Molecular Weight: 387.43

Target: Adrenergic Receptor

Pathway: GPCR/G Protein; Neuronal Signaling

Powder

Storage: 3 years 4°C

2 years -80°C 2 years

In solvent

-20°C

-20°C 1 year

SOLVENT & SOLUBILITY

In Vitro

DMSO: 75 mg/mL (193.58 mM; Need ultrasonic)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	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

Please refer to the solubility information to select the appropriate solvent.

In Vivo

- 1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 7.5 mg/mL (19.36 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 7.5 mg/mL (19.36 mM); Clear solution
- 3. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 7.5 mg/mL (19.36 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	(R)-Terazosin is an active R-enantiomer of Terazosin. (R)-Terazosin is a potent $\alpha 1$ -adrenoceptor antagonist with K_i values of 6.51 nM, 1.01 nM and 1.97 nM for $\alpha 1a$, $\alpha 1b$ and $\alpha 1d$ -adrenoceptor, respectively ^[1] .
IC ₅₀ & Target	Ki: 6.51 nM (α 1a-adrenoceptor), 1.01 nM (α 1b-adrenoceptor) and 1.97 nM (α 1d-adrenoceptor) $^{[1]}$
In Vitro	(R)-Terazosin is low affinity for α 2a, α 2B and α 2c-adrenoceptor with K_i values of 3.85 μ M, 0.33 μ M and 0.37 μ M, respectively $^{[1]}$.

	(R)-Terazosin may prove to be a useful probe in understanding the functional role of subtypes of adrenoceptors in various tissues. Because it is a weaker antagonist at $\alpha 2B$ sites than its enantiomer, it may be possible to use (R)-Terazosin to differentiate between pharmacological effects mediated by subtypes of $\alpha 2$ -adrenoceptors in animal studies ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only.
In Vivo	(R)-Terazosin shows antagonism of at rat atrial α 2B receptor with a pEC ₃₀ of 5.69. (R)-Terazosin shows antagonism of at rat vas deferens α 1A and α 2A receptor with pA ₂ values of 7.5 and 5.31, respectively ^[1] . 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

Page 2 of 2 www.MedChemExpress.com