MCE MedChemExpress

Product Data Sheet

Radequinil

 Cat. No.:
 HY-106025

 CAS No.:
 219846-31-8

 Molecular Formula:
 C₁₈H₁₄N₄O₃

 Molecular Weight:
 334.33

Target: GABA Receptor

Pathway: Membrane Transporter/Ion Channel; Neuronal Signaling

Storage: 4°C, sealed storage, away from moisture and light

* In solvent: -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture

and light)

SOLVENT & SOLUBILITY

In Vitro

DMSO: 10 mg/mL (29.91 mM; ultrasonic and warming and heat to 60°C)

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	2.9911 mL	14.9553 mL	29.9106 mL
	5 mM	0.5982 mL	2.9911 mL	5.9821 mL
	10 mM	0.2991 mL	1.4955 mL	2.9911 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: ≥ 1 mg/mL (2.99 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE- β -CD in saline) Solubility: \geq 1 mg/mL (2.99 mM); Clear solution
- Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 1 mg/mL (2.99 mM); Clear solution

BIOLOGICAL ACTIVITY

Description	Radequinil (AC-3933) is a benzodiazepine receptor (BzR) partial inverse agonist. AC-3933 binds to GABA(-) and GABA(+) ligand with K_i s of 5.15 and 6.11 nM, respectively ^[1] .
IC ₅₀ & Target	Ki: 5.15±0.39 nM (GABA(-) ligand), 6.11±0.26 nM (GABA(+) ligand) ^[1]

REFERENCES

1]. Hashimoto T, et al. Pharmad	cological properties of AC-3933, a novel benz	odiazepine receptor partial inverse aş	gonist. Neuroscience. 2014 Jan 3;256:352-9.	
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