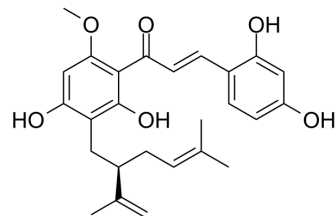


Kuraridine

Cat. No.:	HY-121381	
CAS No.:	34981-25-4	
Molecular Formula:	C ₂₆ H ₃₀ O ₆	
Molecular Weight:	438.51	
Target:	Phosphodiesterase (PDE)	
Pathway:	Metabolic Enzyme/Protease	
Storage:	Powder	-20°C 3 years
	In solvent	-80°C 6 months
		-20°C 1 month



SOLVENT & SOLUBILITY

In Vitro

DMSO : ≥ 100 mg/mL (228.04 mM)
 * "≥" means soluble, but saturation unknown.

Concentration	Solvent	Mass		
		1 mg	5 mg	10 mg
Preparing Stock Solutions	1 mM	2.2804 mL	11.4022 mL	22.8045 mL
	5 mM	0.4561 mL	2.2804 mL	4.5609 mL
	10 mM	0.2280 mL	1.1402 mL	2.2804 mL

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

BIOLOGICAL ACTIVITY

Description

Kuraridine is a prenylated flavonol extract from the roots of *Sophora flavescens*. Kuraridine has an inhibitory effect on cGMP specific phosphodiesterase type 5 (PDE5) (IC₅₀=0.64 μM)^[1].

IC₅₀ & Target

PDE5
 0.64 μM (IC₅₀)

In Vitro

Kuraridine shows potent inhibitory activity (IC₅₀=0.64 μM) against cGMP PDE5 with 2.0- and 12.9-fold selectivity over PDE3 and PDE4, respectively^[1].

Kuraridine induces pronounced chloride inward currents in the absence of GABA. The currents does not exceed 10% of the maximal I_{GABA} induced by a saturating GABA concentration (1 mM)^[2].

Kuraridine induces a concentration-response curves for I_{GABA} enhancement (EC₅₀=4.0±2.4 μM) in *Xenopus* oocytes expressing GABA_A receptors composed of α1, β2, and γ2S subunits^[2].

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

REFERENCES

- [1]. Hye Joo Shin, et al. A prenylated flavonol, sophoflavescenol: a potent and selective inhibitor of cGMP phosphodiesterase 5. *Bioorg Med Chem Lett*. 2002 Sep 2;12(17):2313-6.
- [2]. Xinzhou Yang, et al. HPLC-based activity profiling for GABAA receptor modulators from the traditional Chinese herbal drug Kushen (*Sophora flavescens* root). *Mol Divers*. 2011 May;15(2):361-72.
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Caution: Product has not been fully validated for medical applications. For research use only.

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