UK-78282 hydrochloride

Cat. No.:	HY-W013727	
CAS No.:	136647-02-4	
Molecular Formula:	C ₂₉ H ₃₆ CINO ₂	
Molecular Weight:	466.05	
Target:	Potassium Channel	
Pathway:	Membrane Transporter/Ion Channel	H-CI
Storage:	-20°C, sealed storage, away from moisture	
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)	

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (214.57 mM; Need ultrasonic)					
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg	
		1 mM	2.1457 mL	10.7285 mL	21.4569 mL	
		5 mM	0.4291 mL	2.1457 mL	4.2914 mL	
		10 mM	0.2146 mL	1.0728 mL	2.1457 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: ≥ 2.5 mg/mL (5.36 mM); Clear solution					
	2. Add each solvent one by one: 10% DMSO >> 90% (20% SBE-β-CD in saline) Solubility: ≥ 2.5 mg/mL (5.36 mM); Clear solution					
	3. Add each solvent o Solubility: ≥ 2.5 mg	one by one: 10% DMSO >> 90% cor g/mL (5.36 mM); Clear solution	n oil			

Description	UK-78282, a novel piperidine, potent and selective Kv1.3 blocker with an IC ₅₀ of 200 nM. UK-78,282 effectively suppresses human T-lymphocyte activation in vitro. UK-78,282 binds to residues at the inner surface of the channel overlapping the site of action of verapamil ^[1] .			
IC ₅₀ & Target	IC50: 200 nM (Kv1.3) ^[1]			

REFERENCES



[1]. D C Hanson, et al. UK-78,282, a novel piperidine compound that potently blocks the Kv1.3 voltage-gated potassium channel and inhibits human T cell activation. Br J Pharmacol. 1999 Apr;126(8):1707-16

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

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