MEN 11270

Cat. No.:	HY-103289			
CAS No.:	235082-52-7	HO		
Molecular Formula:	$C_{60}H_{90}N_{20}O_{11}S$			
Molecular Weight:	1299.55			
Sequence:	{d-Arg}-Arg-Pro-{Hyp}-Gly-{Thi}-c({Dab}-{d-Tic}-{Oic}-Arg)c(7γ-10α)			
Sequence Shortening:	$\label{eq:constraint} $$ d-Arg}-RP-{Hyp}-G-{Thi}-c({Dab}-{d-Tic}-{Oic}-R)c(7\gamma-10\alpha)$$$	NH HN=		
Target:	Bradykinin Receptor			
Pathway:	GPCR/G Protein			
Storage:	Sealed storage, away from moisture and light Powder -80°C 2 years -20°C 1 year * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture and light)			

SOLVENT & SOLUBILITY

In Vitro	DMSO : 100 mg/mL (76.95 mM; Need ultrasonic)				
	Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
		1 mM	0.7695 mL	3.8475 mL	7.6950 mL
		5 mM	0.1539 mL	0.7695 mL	1.5390 mL
		10 mM	0.0769 mL	0.3847 mL	0.7695 mL
	Please refer to the so	lubility information to select the app	propriate solvent.		
In Vivo	1. Add each solvent one by one: 10% DMSO >> 40% PEG300 >> 5% Tween-80 >> 45% saline Solubility: ≥ 2.5 mg/mL (1.92 mM); Clear solution				
	 Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (1.92 mM); Clear solution 				

BIOLOGICAL ACTIV	
Description	MEN 11270, a cyclic decapeptide, is a B2 kinin receptor antagonist. MEN 11270 bound with high-affinity to the B2 kinin receptor constitutively expressed by WI38 human fibroblasts, inhibiting 3H-bradykinin (BK) with a pK _i value of 10.3 ^[1] .

REFERENCES



[1]. Meini S, et, al. MEN 11270, A novel selective constrained peptide antagonist with high affinity at the human B2 kinin receptor. J Pharmacol Exp Ther. 1999 Jun;289(3):1250-6.

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

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