(Glu2)-TRH

Cat. No.:	HY-P3960			
CAS No.:	85541-78-2			
Molecular Formula:	$C_{15}H_{22}N_4O_6$			
Molecular Weight:	354.36			
Sequence Shortening:	Glp-EP-NH2			H
Target:	Endogenous	Metabol	ite; Thyroid Hormone Receptor	°⇒(")
Pathway:	Metabolic Er	nzyme/Pr	rotease; Vitamin D Related/Nuclear Receptor	
Storage:	Sealed stora Powder * In solvent : and light)	ge, away -80°C -20°C -80°C, 6	r from moisture and light 2 years 1 year months; -20°C, 1 month (sealed storage, away from moisture	

SOLVENT & SOLUBILITY

		Solvent Mass Concentration	1 mg	5 mg	10 mg
Pr	Preparing Stock Solutions	1 mM	2.8220 mL	14.1099 mL	28.2199 mL
		5 mM	0.5644 mL	2.8220 mL	5.6440 mL
	10 mM	0.2822 mL	1.4110 mL	2.8220 mL	

BIOLOGICAL ACTIV		
Description	(Glu2)-TRH, a metabolically stable analogue of Thyrotropin-releasing hormone (TRH; HY-P0002), is a negative modulator for the cholinergic effect of TRH in the mouse brain. (Glu2)-TRH significantly attenuates TRH-induced hippocampal extracellul acetylcholine release. (Glu2)-TRH is not metabolized by thyroliberinase. (Glu2)-TRH manifests neuroprotective, antidepressant, anticonvulsant in the CNS ^[1] .)r ar
In Vivo	(Glu2)-TRH (1-100 μmol/kg; IV) dose-dependently attenuated TRH-induced analeptic action ^[1] . (Glu2)-TRH (10 μmol/kg; IV) 10 min before Pentobarbital (60 mg/kg; i.p.) decreases the sleeping time by approximately 20% compared to the vehicle treated group ^[1] . MCE has not independently confirmed the accuracy of these methods. They are for reference only. Animal Model: Male Swiss-Webster mice (30 g) ^[1])

 H_2N

¥ ОН



Dosage:	1, 2, 5, 10, 15, 25, 50, 100 μmol/kg
Administration:	IV; 10 min before Pentobarbital (60 mg/kg, i.p.)
Result:	Dose-dependently attenuated TRH-induced (10 μ mol/kg; iv) analeptic action.

REFERENCES

[1]. Vien Nguyen, et al. [Glu2]TRH dose-dependently attenuates TRH-evoked analeptic effect in mice. Brain Res Bull. 2010 Apr 29;82(1-2):83-6.

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

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