CH 5450

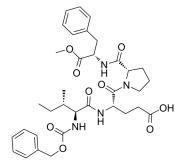
Cat. No.: HY-16707 CAS No.: 252557-97-4 Molecular Formula: $C_{34}H_{44}N_4O_9$ Molecular Weight: 652.73

Sequence: Z-Ile-Glu-Pro-Phe-Ome

Sequence Shortening: ZIEPF-Ome Others Target: Others Pathway:

Storage: Sealed storage, away from moisture

> -80°C Powder 2 years -20°C 1 year



Product Data Sheet

SOLVENT & SOLUBILITY

In Vitro

DMSO: ≥ 43 mg/mL (65.88 mM)

* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent Mass Concentration	1 mg	5 mg	10 mg
	1 mM	1.5320 mL	7.6601 mL	15.3203 mL
	5 mM	0.3064 mL	1.5320 mL	3.0641 mL
	10 mM	0.1532 mL	0.7660 mL	1.5320 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 (3.83 mM); Clear solution
- 2. Add each solvent one by one: 10% DMSO >> 90% corn oil Solubility: ≥ 2.5 mg/mL (3.83 mM); Clear solution

BIOLOGICAL ACTIVITY

Description

CH 5450 (Z-Ile-Glu-Pro-Phe-Ome) is a selective short peptide human cardiac chymase inhibitor. CH-5450 inhibits the action of rat MAB elastase 2 on substrate Ang I with an IC $_{50}$ value of 49 μM and N-succinyl-Ala-Ala-Pro-Phe-p-nitroanilide with an IC $_{50}$ value of 4.8 μ M^[1].

IC₅₀ & Target

Chymase

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

REFERENCES				
[1]. Carlos F Santos, et al. Kinetic characterization and inhibition of the rat MAB elastase-2, an angiotensin I-converting serine protease. Can J Physiol Pharmacol. 2002 Jan;80(1):42-7.				
	Caution: Product has not been fully validated for medical applications. For research use only.			
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