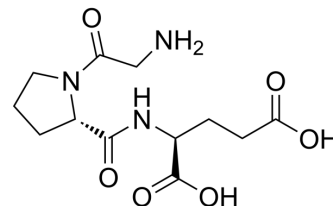


## Gly-Pro-Glu

<b>Cat. No.:</b>	HY-117483
<b>CAS No.:</b>	32302-76-4
<b>Molecular Formula:</b>	C <sub>12</sub> H <sub>19</sub> N <sub>3</sub> O <sub>6</sub>
<b>Molecular Weight:</b>	301.3
<b>Target:</b>	Others
<b>Pathway:</b>	Others
<b>Storage:</b>	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 : 66.67 mg/mL (221.27 mM; ultrasonic and adjust pH to 2 with 1M HCl)

Concentration	Mass		
	1 mg	5 mg	10 mg
1 mM	3.3190 mL	16.5948 mL	33.1895 mL
5 mM	0.6638 mL	3.3190 mL	6.6379 mL
10 mM	0.3319 mL	1.6595 mL	3.3190 mL

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

### BIOLOGICAL ACTIVITY

#### Description

Gly-Pro-Glu is a neuroactive peptide with a potent action on acetylcholine release. Gly-Pro-Glu is the N-terminal tripeptide of insulin-like growth factor-I. Gly-Pro-Glu inhibits glutamate binds to N-methyl-D-aspartate (NMDA) receptor with an IC<sub>50</sub> value of 14.7 μM. Gly-Pro-Glu can be used for the research of neuroprotection <sup>[1][2]</sup>.

#### IC<sub>50</sub> & Target

IC<sub>50</sub>: 14.7 μM (glutamate binds to NMDA receptor)<sup>[1]</sup>

#### In Vitro

Gly-Pro-Glu (0-100 μM) potentiates the potassium evoked release of both acetylcholine and dopamine, increases K<sup>+</sup> evoked acetylcholine release even at concentrations of 0.1 nM and significantly enhances evoked dopamine release<sup>[1]</sup>. Gly-Pro-Glu (1-1000 μM) shows an inhibition of L-[<sup>3</sup>H]glutamate binding with an IC<sub>50</sub> value of 14.7 μM<sup>[1]</sup>. MCE has not independently confirmed the accuracy of these methods. They are for reference only.

#### In Vivo

Gly-Pro-Glu (300 mg; i.p. once per day; on day 0, 6 and 12) shows an in vivo effect protecting the temporal cortical somatostatinergic system from Abeta insult.<sup>[2]</sup> MCE has not independently confirmed the accuracy of these methods. They are for reference only.

Animal Model:	Ovariectomized rats with Abeta25-35 injection <sup>[2]</sup>
Dosage:	300 mg
Administration:	Intraperitoneal injection; 300 mg per day; on day 0, 6 and 12
Result:	Recovered Abeta25-35-induced the reduction of somatostatin (SRIF) content and SRIF receptor density, and reduced the inhibitory effect of SRIF on adenylyl cyclase activity.

## REFERENCES

[1]. Sara VR, et al. Identification of Gly-Pro-Glu (GPE), the aminoterminal tripeptide of insulin-like growth factor 1 which is truncated in brain, as a novel neuroactive peptide. *Biochem Biophys Res Commun.* 1989 Dec 15;165(2):766-71.

[2]. Aguado-Llera D, et al. Gly-Pro-Glu protects beta-amyloid-induced somatostatin depletion in the rat cortex. *Neuroreport.* 2004 Aug 26;15(12):1979-82.

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

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