

## Product Data Sheet

## Orexin A (human, rat, mouse) (TFA)

Cat. No.:	HY-106224A				
Molecular Formula:	C <sub>152</sub> H <sub>243</sub> N <sub>47</sub> O <sub>44</sub> S <sub>4</sub> .xC <sub>2</sub> HF <sub>3</sub> O <sub>2</sub>				
Sequence Shortening:	{Glp}-PLPDCCRQKTCSCRLYELLHGAGNHAAGILTL-NH2 (Disulfide bridge: Cys6-Cys12, C ys7-Cys14)				
Target:	Orexin Receptor (OX Receptor)				
Pathway:	GPCR/G Protein; Neuronal Signaling				
Storage:	Sealed stora Powder	ge, away -80°C -20°C	from moisture 2 years 1 year		
	* In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)				

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In Vitro

H<sub>2</sub>O:100 mg/mL (Need ultrasonic)

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DIOLOGICAL ACTIV					
Description	Orexin A (human, rat, mouse) (Hypocretin-1 (human, rat, mouse)) TFA, a 33 amino acid excitatory neuropeptide, orchestrates diverse central and peripheral processes. Orexin A (human, rat, mouse) TFA is a specific, high-affinity agonist for G-protein-coupled receptor OX1R. Orexin A (human, rat, mouse) TFA has a role in the regulation of feeding behavior. Orexin A (human, rat, mouse) TFA is an effective anti-nociceptive and anti-hyperalgesic agent in mice and rats <sup>[1][2]</sup> .				
IC <sub>50</sub> & Target	OX <sub>1</sub> Receptor				
In Vitro	Orexin A (human, rat, mouse) TFA has high affinity for OX1R, with 38 nM IC <sub>50</sub> and 34 nM EC <sub>50</sub> values in the the [Ca <sup>2⊠</sup> ]i transient assay <sup>[1]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.				
In Vivo	Orexin A (human, rat, mouse) TFA (3-30 mg/kg; i.v.; 5 min pre-test) significantly increases the latency to response at 10 and 30 mg/kg i.v. when given 5 min pre-test from 24.8±2.0 s in vehicle-treated mice to 35.0±3.7 s and 45.7±4.5 s, respectively <sup>[2]</sup> . Orexin A (human, rat, mouse) TFA (3, 10 and 30 mg/kg; i.v.) is given immediately before phenylp-quinone (PPQ) and increases the latency to the first PPQ-induced constriction from 357.4±35.2 s in vehicle-treated mice to 500.3±31.2 s at 10 mg/kg and 594.5±5.5 s at 30 mg/kg <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.Animal Model:Female mice (mouse carrageenan-induced thermal hyperalgesia test <sup>[2]</sup>				
	Dosage:	3, 10 and 30 mg/kg			
	Administration:	i.v.; 5 min pre-test			
	Result:	Significantly increased the latency to response at 10 and 30 mg/kg.			

Cym-Cymu) (TFA self

## CUSTOMER VALIDATION

- J Inflamm Res. 2021 May 18;14:2007-2017.
- Brain Res Bull. 2021 Apr;169:81-93.
- Med Sci Monit. 2019 Apr 19;25:2886-2895.

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## REFERENCES

[1]. Sakurai T, et al. Orexins and orexin receptors: a family of hypothalamic neuropeptides and G protein-coupled receptors that regulate feeding behavior. Cell. 1998 Feb 20;92(4):573-85.

[2]. Bingham S, et al. Orexin-A, an hypothalamic peptide with analgesic properties. Pain. 2001 May;92(1-2):81-90.

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

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