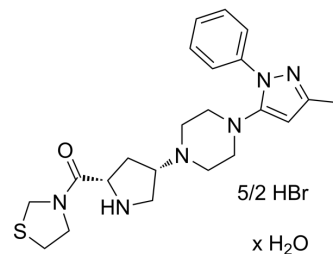


## Teneligliptin hydrobromide hydrate

<b>Cat. No.:</b>	HY-14806B
<b>CAS No.:</b>	1572583-29-9
<b>Molecular Formula:</b>	C <sub>22</sub> H <sub>30</sub> N <sub>6</sub> OS
<b>Molecular Weight:</b>	426.58
<b>Target:</b>	Dipeptidyl Peptidase
<b>Pathway:</b>	Metabolic Enzyme/Protease
<b>Storage:</b>	4°C, sealed storage, away from moisture * In solvent : -80°C, 6 months; -20°C, 1 month (sealed storage, away from moisture)



### SOLVENT & SOLUBILITY

#### In Vitro

H<sub>2</sub>O : ≥ 100 mg/mL (234.42 mM)  
\* "≥" means soluble, but saturation unknown.

Preparing Stock Solutions	Solvent		Mass		
	Concentration		1 mg	5 mg	10 mg
	1 mM		2.3442 mL	11.7211 mL	23.4423 mL
	5 mM		0.4688 mL	2.3442 mL	4.6885 mL
	10 mM		0.2344 mL	1.1721 mL	2.3442 mL

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

### BIOLOGICAL ACTIVITY

#### Description

Teneligliptin hydrobromide hydrate is a potent chemotype prolylthiazolidine-based DPP-4 inhibitor, which competitively inhibits human plasma, rat plasma, and human recombinant DPP-4 in vitro, with IC<sub>50</sub>s of approximately 1 nM<sup>[1]</sup>.

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

DPP-4

### CUSTOMER VALIDATION

- Antioxidants (Basel). 2023 Jul 24;12(7):1478.
- Antioxidants (Basel). 2021 Sep 9;10(9):1438.
- Eur J Med Chem. 2021 Feb 15;212:113030.
- Nephrol Dial Transplant. 2019 Oct 1;34(10):1669-1680.
- Chem Res Toxicol. 2020 Aug 17;33(8):2164-2171.

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## REFERENCES

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[1]. Fukuda-Tsuru S, et al. A novel, potent, and long-lasting dipeptidyl peptidase-4 inhibitor, teneligliptin, improves postprandial hyperglycemia and dyslipidemia after single and repeated administrations. *Eur J Pharmacol.* 2012 Dec 5;696(1-3):194-202.

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

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