

## CSD-CH2(1,8)-NH2

<b>Cat. No.:</b>	HY-P5756
<b>CAS No.:</b>	3032600-19-1
<b>Molecular Formula:</b>	C <sub>76</sub> H <sub>125</sub> N <sub>25</sub> O <sub>15</sub> S <sub>2</sub>
<b>Molecular Weight:</b>	1693.09
<b>Target:</b>	Opioid Receptor
<b>Pathway:</b>	GPCR/G Protein; Neuronal Signaling
<b>Storage:</b>	Please store the product under the recommended conditions in the Certificate of Analysis.

### BIOLOGICAL ACTIVITY

<b>Description</b>	CSD-CH2(1,8)-NH2 is a selective and competitive KOR antagonist (K <sub>i</sub> : 6.8 nM). CSD-CH2(1,8)-NH2 inhibits calcium mobilization in DRG neurons. CH2(1,8)-NH2 antagonizes the antinociceptive effect of U50,488. CSD-CH2(1,8)-NH2 can be used for research of neuropsychiatric disorders <sup>[1]</sup> .
<b>IC<sub>50</sub> &amp; Target</b>	K <sub>i</sub> : 6.8 nM (KOR) <sup>[1]</sup>

### REFERENCES

[1]. Muratspahić E, et al. Development of a Selective Peptide κ-Opioid Receptor Antagonist by Late-Stage Functionalization with Cysteine Staples. *J Med Chem.* 2023 Sep 14;66(17):11843-11854.

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

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