## **Product** Data Sheet

## **GB-110**

 Cat. No.:
 HY-120528

 CAS No.:
 1252806-70-4

 Molecular Formula:
 C<sub>33</sub>H<sub>48</sub>N<sub>6</sub>O<sub>5</sub>

Molecular Weight: 608.77

Target: Protease-Activated Receptor (PAR)

Pathway: GPCR/G Protein

Storage: Please store the product under the recommended conditions in the Certificate of

Analysis.

## **BIOLOGICAL ACTIVITY**

Description	GB-110 is a potent, orally active, and nonpeptidic protease activated receptor 2 (PAR2) agonist. GB-110 selectively induces PAR2-mediated intracellular $Ca^{2+}$ release in HT29 cells with an $EC_{50}$ of 0.28 $\mu$ M <sup>[1]</sup> .
IC <sub>50</sub> & Target	Protease activated receptor $2^{[1]}$
In Vitro	In an intracellular $Ca^{2+}$ (i $Ca^{2+}$ ) mobilization assay using HT29 colon cancer cells, GB110 (E $C_{50}$ 240±20 nM; pE $C_{50}$ 6.7±0.05) is equipotent with the peptide agonist 2f-LIGRLO-NH2 (E $C_{50}$ 210±30 nM; pE $C_{50}$ 6.6±0.05), 10-fold more potent than SLIGRL-NH2, but ~35-fold less potent than trypsin (E $C_{50}$ 6±0.5 nM; pE $C_{50}$ 8.2±0.8) <sup>[2]</sup> . MCE has not independently confirmed the accuracy of these methods. They are for reference only.

## **REFERENCES**

[1]. Barry GD, et al. Novel agonists and antagonists for human protease activated receptor 2. J Med Chem. 2010 Oct 28;53(20):7428-40.

[2]. Suen JY, et al. Modulating human proteinase activated receptor 2 with a novel antagonist (GB88) and agonist (GB110). Br J Pharmacol. 2012 Mar;165(5):1413-23.

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

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