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## DATASHEET

SLIGRL-NH2

### Product overview

**Name** SLIGRL-NH2  
**Cat No** HB2920  
**Biological description** Overview

SLIGRL-NH2 is a PAR<sub>2</sub> peptide agonist which is specific for activation of PAR<sub>2</sub> over PAR<sub>1</sub> and PAR<sub>4</sub> (EC<sub>50</sub> = 0.5-2.0µM).

The peptide is derived from the N-terminus of the rat PAR<sub>2</sub> receptor sequence.

#### Uses

SLIGRL-NH2 is an itch-inducing agent which is commonly used to study histamine-independent itch. It evokes dose dependent scratching behaviour in mice. Interestingly, in pruritogen-responsive neurons that transmit the itch signal induced by SLIGRL-NH2, sex-related differences may exist.

The peptide has also been shown to enhance gastrointestinal transit in mice and rats

**Biological action** Active *in vivo*.  
**Purity** Agonist  
**Description** >95%  
PAR<sub>2</sub> peptide agonist

### Solubility & Handling

**Storage instructions** -20 °C  
**Solubility overview** Soluble in water (1 mg/ml)  
**Important** This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not for human or veterinary use

### Chemical Data

**Chemical name** SLIGRL (modifications: C-terminal amide)  
**Molecular Weight** 656.82  
**Chemical structure**   
**Molecular Formula** C<sub>29</sub>H<sub>56</sub>N<sub>10</sub>O<sub>7</sub>  
**Sequence (one letter)** SLIGRL  
**Modifications** C-terminal amide  
**CAS Number** 171436-38-7  
**PubChem identifier** 9831050  
**SMILES** CC[C@H](C)[C@@H](C(=O)NCC(=O)N[C@@H](CCCN=C(N)N)C(=O)N[C@@H](CC(C)C)C(=O)N)NC(=O)[C@H](CC(C)C)NC(=O)[C@H](CO)N  
SGPMJRPYYIJZPC-JYAZKYGWSA-N  
**InChiKey** MFCD03093421  
**MDL number**

### References

**Protease-activated receptor-1 (PAR1) and PAR2 but not PAR4 mediate relaxations in lower esophageal sphincter.**

Huang et al (2007) Regul Pept. 142(1-2)

**PubMedID** [17335921](#)

**The protease-activated receptor-2-specific agonists 2-aminothiazol-4-yl-LIGRL-NH<sub>2</sub> and 6-aminonicotiny-LIGRL-NH<sub>2</sub> stimulate multiple signaling pathways to induce physiological responses in vitro and in vivo.**

Flynn et al (2011) J Biol Chem 286(21)

**PubMedID** [21467041](#)

**The distinct roles of two GPCRs, MrgprC11 and PAR2, in itch and hyperalgesia.**

Liu et al (2011) Sci Signal. 4(181)

**PubMedID** [21775281](#)

**Sex-related differences in SLIGRL-induced pruritus in mice.**

Yamaura et al (2014) Life Sci. 94(1)

**PubMedID** [24239643](#)

**Therapeutic effect of protease-activated receptor 2 agonist SLIGRL-NH<sub>2</sub> on loperamide-induced Sprague-Dawley rat constipation model and the related mechanism.**

Zhang et al (30122898) Drug Des Devel Ther. 12

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