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# **DATASHEET**

LIH383

### **Product overview**

Name LIH383 Cat No HB9242

Biological description Novel, potent and highly selective atypical chemokine receptor ACKR3 (CXCR7) agonist. Competes

directly with CXCL12-AF647 for ACKR3 binding at low nanomolar concentrations and is more potent in inducing beta-arrestin recruitment to ACKR3 (EC $_{50}$  = 0.61 nM) than the CXCL12 and CXCL11 chemokine ligands (EC $_{50}$  = 1.2 and 2.2 nM respectively). Restrains the negative regulatory function of

ACKR3 on opioid peptides to potentiate their activity in rat brain.

**Biological action** Peptide Purity >95%

**Description** Potent and selective ACKR3 (CXCR7) agonist. Potentiates opioid peptide activity.

## **Solubility & Handling**

Storage instructions

Solubility overview

Soluble in aqueous buffer

-20°C

Important This product is for RESE

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not

for human or veterinary use

### **Chemical Data**

 $\begin{tabular}{lll} \textbf{Molecular Weight} & 997.23 \\ \textbf{Molecular Formula} & $C_{45}H_{72}N_{16}O_8S$ \\ \textbf{Sequence (one letter)} & FGGFMRRK-NH2 \\ \end{tabular}$ 

Sequence (three letter) H-Phe-Gly-Gly-Phe-Met-Arg-Arg-Lys-NH2

**Modifications** Amide at C terminal lysine

### References

The atypical chemokine receptor ACKR3/CXCR7 is a broad-spectrum scavenger for opioid peptides

Meyrath M *et al* (2020) Nat Commun 11(1) **PubMedID**32561830