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CGRP Receptor

Calcitonin gene-related peptide receptor

CGRP receptor is a heterodimer formed by calcitonin-receptor-like receptor (CRLR), a type II (family B) G-protein-coupled receptor, and receptor-activity-modifying protein 1 (RAMP1), a single-membrane-pass protein. RAMP1 is needed for CGRP binding and also cell-surface expression of CLR. CLR is an example of a family B GPCR.

CGRP is a neuropeptide abundant in the trigeminal system and widely expressed in both the peripheral and central nervous systems. CGRP has several functions including vasodilation, the perception of painful stimuli, and inflammation. CGRP exerts its biological action by interacting with its receptors. There are two types of CGRP receptors, CGRP-A and CGRP-B.

<p>CGRP antagonist 1</p> <p>Cat. No.: HY-112262</p>	<p>HCGRP-(8-37) (Human α-CGRP (8-37))</p> <p>Cat. No.: HY-P1014</p>
<p>CGRP antagonist 1 is a highly potent CGRP receptor antagonist with a K_i and IC_{50} of 35 and 57 nM, respectively.</p> <p>Purity: >98%</p> <p>Clinical Data: No Development Reported</p> <p>Size: 1 mg, 5 mg</p>	<p>HCGRP-(8-37) is a human calcitonin gene-related peptide (hCGRP) fragment and also an antagonist of CGRP receptor.</p> <p>Purity: 98.0%</p> <p>Clinical Data: No Development Reported</p> <p>Size: 500 μg, 1 mg, 5 mg</p>
<p>Kendomycin (-)-TAN2162</p> <p>Cat. No.: HY-121300</p>	<p>MK-3207</p> <p>Cat. No.: HY-10301</p>
<p>Kendomycin ((-)-TAN 2162) is a polyketide antibiotic with remarkable antibacterial and cancer cells cytotoxic activities. Kendomycin tends to be bacteriostatic rather than bactericidal and inhibits the growth of the.</p> <p>Purity: >98%</p> <p>Clinical Data: No Development Reported</p> <p>Size: 1 mg, 5 mg</p>	<p>MK-3207 is a potent and orally bioavailable CGRP receptor antagonist (IC_{50}= 0.12 nM; K_i value= 0.024 nM); highly selective versus human AM1, AM2, CTR, and AMY3.</p> <p>Purity: 98.76%</p> <p>Clinical Data: Phase 2</p> <p>Size: 10 mM \times 1 mL, 5 mg, 10 mg</p>
<p>MK-3207 Hydrochloride</p> <p>Cat. No.: HY-10302</p>	<p>Olcegepant (BIBN-4096; BIBN 4096BS)</p> <p>Cat. No.: HY-10095</p>
<p>MK-3207 (Hydrochloride) is a potent and orally bioavailable CGRP receptor antagonist with IC_{50} of 0.12 nM and K_i of 0.024 nM, and is highly selective versus human AM1, AM2, CTR, and AMY3.</p> <p>Purity: 99.06%</p> <p>Clinical Data: Phase 1</p> <p>Size: 10 mM \times 1 mL, 5 mg, 10 mg</p>	<p>Olcegepant (BIBN-4096) is a potent and selective non-peptide antagonist of the calcitonin gene-related peptide 1 (CGRP1) receptor with IC_{50} of 0.03 nM and K_i of 14.4 pM for human CGRP.</p> <p>Purity: 99.65%</p> <p>Clinical Data: Phase 2</p> <p>Size: 10 mM \times 1 mL, 2 mg, 5 mg, 10 mg, 50 mg</p>
<p>Olcegepant hydrochloride (BIBN-4096 hydrochloride; BIBN4096BS hydrochloride)</p> <p>Cat. No.: HY-10095A</p>	<p>PHM-27 (human)</p> <p>Cat. No.: HY-P1072</p>
<p>Olcegepant hydrochloride (BIBN-4096 hydrochloride) is a potent and selective non-peptide antagonist of the calcitonin gene-related peptide 1 (CGRP1) receptor with IC_{50} of 0.03 nM and with a K_i of 14.4 pM for human CGRP.</p> <p>Purity: 99.31%</p> <p>Clinical Data: Phase 2</p> <p>Size: 10 mM \times 1 mL, 5 mg, 10 mg, 50 mg</p>	<p>PHM-27 (human) is a human prepro-vasoactive intestinal polypeptide (27 amino acid). PHM-27 (human) is a potent the human calcitonin receptor agonist with an EC_{50} of 11 nM.</p> <p>Purity: >98%</p> <p>Clinical Data: No Development Reported</p> <p>Size: 1 mg, 5 mg</p>
<p>Rat CGRP-(8-37)</p> <p>Cat. No.: HY-P0209</p>	<p>Rimegepant (BMS-927711)</p> <p>Cat. No.: HY-15498</p>
<p>Rat CGRP-(8-37) (VTHRLAGLLSRSGGVVKNFVPTNVGSEAF-NH₂) is a highly selective CGRP receptor antagonist.</p> <p>Purity: 98.54%</p> <p>Clinical Data: No Development Reported</p> <p>Size: 500 μg, 1 mg, 5 mg</p>	<p>Rimegepant (BMS-927711) is a highly potent, oral calcitonin gene-related peptide (CGRP) receptor antagonist with a K_i of 0.027 nM and an IC_{50} of 0.14 nM for hCGRP receptor.</p> <p>Purity: 99.08%</p> <p>Clinical Data: Launched</p> <p>Size: 5 mg, 10 mg, 50 mg, 100 mg</p>

