

CaSR

Calcium-sensing receptor

HDAC Inhibitor:
Vorinostat (SAHA)

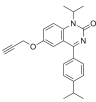
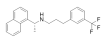

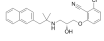

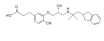


HDAC (Histone deacetylase)

of parathyroid hormone. It also inhibits the cAMP dependent pathway.

CaSR (calcium-sensing receptor) is a Class C G-protein coupled receptor which senses extracellular levels of calcium ion. In the parathyroid gland, the calcium-sensing receptor controls calcium homeostasis by regulating the release of parathyroid hormone (PTH). The release of PTH is inhibited in response to elevations in plasma calcium concentrations and activation of the calcium receptor. Increased calcium binding on the extracellular side gives a conformational change in the receptor, which, on the intracellular side, initiates the phospholipase C pathway, presumably through a Gq α type of G protein, which ultimately increases intracellular concentration of calcium, which inhibits vesicle fusion and exocytosis

CaSR Inhibitors & Modulators

<p>Calcium-Sensing Receptor Antagonists I Cat. No.: HY-50713</p> <p>Bioactivity: Calcium-Sensing Receptor Antagonists I is an antagonist of calcium-sensing parathyroid hormone receptors.</p> <p>Purity: 99.02% Clinical Data: No Development Reported Size: 10mM x 1mL in DMSO, 10 mg, 100 mg</p> 	<p>Cinacalcet (AMG 073) Cat. No.: HY-70037</p> <p>Bioactivity: Cinacalcet is an orally active, allosteric agonist of Ca receptor (CaR), used for cardiovascular disease treatment.</p> <p>Purity: 99.65% Clinical Data: Launched Size: 10mM x 1mL in DMSO, 10 mg, 50 mg</p> 
<p>Cinacalcet hydrochloride (AMG-073 hydrochloride; Cinacalcet) Cat. No.: HY-70037A</p> <p>Bioactivity: Cinacalcet hydrochloride is an orally active, allosteric agonist of Ca receptor (CaR), used for cardiovascular disease treatment.</p> <p>Purity: 99.97% Clinical Data: Launched Size: 10mM x 1mL in DMSO, 10 mg, 50 mg, 100 mg</p> 	<p>Evocalcet (KHK7580) Cat. No.: HY-17613</p> <p>Bioactivity: Evocalcet has an activating effect on calcium sensing receptor (CaSR) extracted from patent WO 2017061621 A1, compound A.</p> <p>Purity: 98.52% Clinical Data: No Development Reported Size: 10mM x 1mL in DMSO, 1 mg, 5 mg, 10 mg, 50 mg, 100 mg</p> 
<p>GSK3004774 Cat. No.: HY-107773</p> <p>Bioactivity: GSK3004774 is a potent, nonabsorbable agonist of CaSR, with an pEC₅₀ of 7.3, 6.6 and 6.5 for human, mouse and rat CaSR, respectively. GSK3004774 shows an EC₅₀ of 50 nM for human CaSR ^[1].</p> <p>Purity: >98% Clinical Data: No Development Reported Size:</p> 	<p>NPS-2143 (SB 262470A) Cat. No.: HY-10007</p> <p>Bioactivity: NPS-2143 is a selective antagonist of calcium-sensing receptor (CaSR) with an IC₅₀ of 43 nM.</p> <p>Purity: 99.42% Clinical Data: No Development Reported Size: 10mM x 1mL in DMSO, 5 mg, 10 mg, 50 mg, 100 mg</p> 
<p>NPS-2143 hydrochloride Cat. No.: HY-10171</p> <p>Bioactivity: NPS-2143(SB 262470A) is a selective potent calcium ion-sensing receptor antagonist with IC50 of 43 and 41 nM for cytoplasmic Ca2+ concentrations and parathyroid hormone secretion, respectively.</p> <p>Purity: 99.96% Clinical Data: No Development Reported Size: 10mM x 1mL in DMSO, 5 mg, 10 mg, 50 mg</p> 	<p>SB-423562 Cat. No.: HY-15105</p> <p>Bioactivity: SB-423562 is a short-acting calcium-sensing receptor (CaR) antagonist.</p> <p>Purity: >98% Clinical Data: No Development Reported Size: 1 mg, 5 mg, 10 mg</p> 
<p>Strontium Ranelate (Distronium renelate; S12911) Cat. No.: HY-17397</p> <p>Bioactivity: Strontium ranelate(S12911) stimulates the calcium sensing receptors (CaSR) and leads to the differentiation of pre-osteoblast to osteoblast which increases the bone formation.</p> <p>Purity: 99.16% Clinical Data: Launched Size: 100 mg, 500 mg</p> 