

Pain Research

The development of novel pain therapeutics relies on understanding the highly complex nociceptive signaling pathways. Cayman offers a select group of tools to study TRP ion channels, purinergic receptors, voltage-gated ion channels, and various G protein-coupled receptors (GPCRs) that are known to be expressed by nociceptive primary afferent nerves. Modulators of glutamate, GABA, and nicotinic acetylcholine receptors are also available.



Pain Transmission

Primary nociceptor afferents release neurotransmitters such as glutamate and prostaglandins, which bind to specific receptors on the postsynaptic membrane to transmit a pain signal.

The antinociceptive response includes inhibitory interneurons, signaling via GABA receptors, that activate μ -opioid receptors, among others.

Inhibitory pathways descending from the brainstem release neurotransmitters such as serotonin (5-HT) or activate small opioid-containing interneurons in the spinal dorsal horn to release opioid peptides.

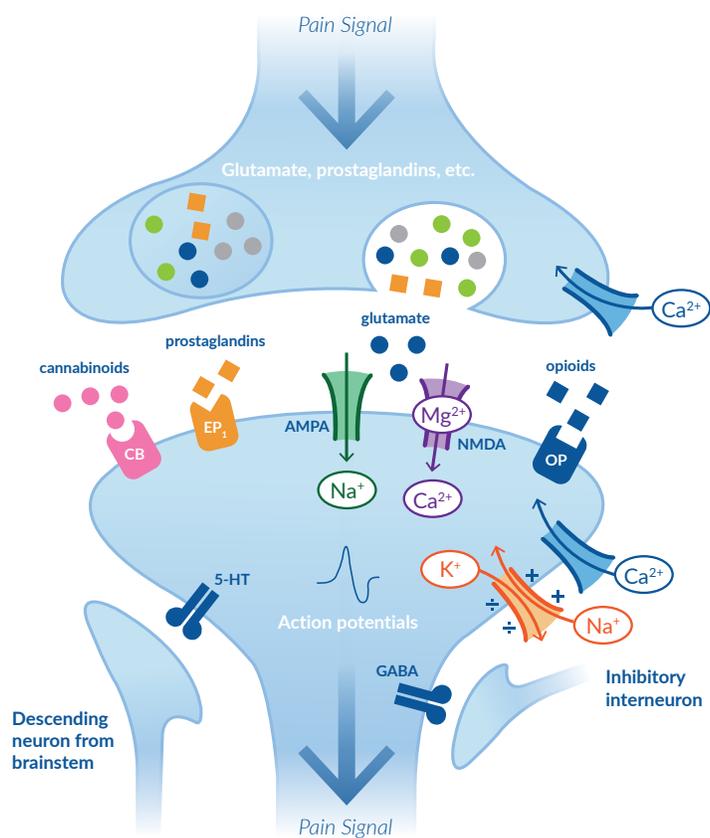


Image adapted from: Olesen, A.E., Andresen, T., Staahl, C., et al. *Pharmacol. Rev.* 64(3), 722-779 (2012).

Ligand-Gated Ion Channels

TRP Ion Channel Agonists and Antagonists

Item No.	Product Name	Summary
14715	AMG 9810	A competitive antagonist of capsaicin activation of TRPV1 (IC_{50} s = 24.5 and 85.6 nM for human and rat, respectively)
92350	Capsaicin	The primary heat and pain-eliciting compound of the <i>Capsicum</i> pepper
10007518	Capsazepine	A competitive TRPV1 antagonist that blocks the capsaicin-induced uptake of Ca^{2+} (IC_{50} = 0.42 μ M in neonatal rat dorsal root ganglia and IC_{50} = 17 nM in Chinese hamster ovary cells)
11923	HC-030031	A selective TRPA1 channel blocker that antagonizes TRPA1-mediated calcium influx induced by AITC and formalin (IC_{50} s = 6.2 and 5.3 μ M, respectively)
10137	Icilin	A CMR1/TRPM8 super agonist that serves as the reference cold nociceptive agonist for TRP-type ion channels
20458	JYL1421	A TRPV1 receptor antagonist
25328	Nonivamide	A TRPV1 agonist
21019	Supercinnamaldehyde	A TRPA1 activator (EC_{50} = 0.8 μ M)

Glutamate/GABA Receptor Agonists and Antagonists

Item No.	Product Name	Summary
18600	(±)-Baclofen	A selective agonist of GABA _B (IC ₅₀ = 200 nM)
16355	Gaboxadol (hydrochloride)	A GABA _A receptor agonist whose potency varies depending on the receptor subunit composition
20919	Ganaxolone	A positive modulator of GABA _A receptors containing α_1 , α_2 , α_3 , β_2 , and γ_{2L} subunits
15351	LY379268	A brain permeable, selective agonist of mGluR2 and mGluR3
10009019	(+)-MK 801 (hydrogen maleate)	A selective, non-competitive NMDA receptor antagonist (K _i = 30.5 nM)

P2X Receptor Antagonists

Item No.	Product Name	Summary
14580	A-438079 (hydrochloride)	A competitive antagonist of the P2X ₇ receptor (pIC ₅₀ = 6.9)
21256	A-740003	A competitive antagonist of the P2X ₇ receptor (IC ₅₀ s = 40 and 18 nM for human and rat receptors, respectively)
21895	JNJ-47965567	A selective P2X ₇ antagonist

Nicotinic Acetylcholine Receptor Modulators

Item No.	Product Name	Summary
21018	NS 1738	A positive allosteric modulator of $\alpha 7$ -containing nAChRs
14304	PNU-120596	A positive allosteric modulator of the $\alpha 7$ nAChR, increasing agonist-evoked calcium flux (EC ₅₀ = 216 nM)
15030	Varenicline (tartrate)	A partial agonist of the $\alpha 4\beta 2$ neuronal nAChR (EC ₅₀ = 2.3 μ M)

Additional modulators of ligand-gated ion channels available at www.caymanchem.com

Voltage-Gated Ion Channels

Sodium Channel Blockers

Item No.	Product Name	Summary
10012588	A-803467	A Na _v 1.8 sodium channel blocker (IC ₅₀ = 8 nM when stimulated at half-maximal inactivation and IC ₅₀ = 79 nM at a resting state)
10011032	QX-314 (bromide)	A selective blocker of sodium channels on nociceptive neurons
21422	Ropivacaine (hydrochloride hydrate)	A reversible blocker of sodium channels in nerve fibers
14964	Tetrodotoxin (citrate)	A neurotoxic blocker of Na _v channels

Calcium Channels

Item No.	Product Name	Summary
10008346	Gabapentin	Penetrates the central nervous system and blocks the $\alpha_2\delta$ -type voltage-gated calcium channels
22475	Mirogabalin	Blocks $\alpha_2\delta$ -1 and $\alpha_2\delta$ -2 subunits of voltage-dependent calcium channels
9002652	Phenibut (hydrochloride)	Blocks $\alpha_2\delta$ subunit-containing voltage-gated calcium channels

Potassium Blockers

Item No.	Product Name	Summary
15185	ML-213	A selective activator of KCNQ2 and KCNQ4 channels
15191	ML-252 (hydrochloride)	A selective inhibitor of the voltage-gated potassium channel subtype K _v 7.2 (IC ₅₀ = 69 nM in a patch clamp assay)
15193	ML-277	An activator of KCNQ1 channels (EC ₅₀ = 260 nM)
21449	Retigabine	An activator of KCNQ2/3, KCNQ3/5, KCNQ2, KCNQ3, KCNQ4, and KCNQ5 channels (EC ₅₀ s = 1.6, 1.4, 2.5, 0.6, 5.2, and 6.4 μ M, respectively)

Additional modulators of voltage-gated ion channels available at www.caymanchem.com

GPCR Receptors

Prostanoid Signaling

Cayman is the gold standard for manufacturing compounds that target the inflammatory mediators of the arachidonic acid cascade. This includes selective cyclooxygenase (COX) inhibitors, NSAIDs, blockers of microsomal prostaglandin E synthase-1 (mPGES-1), and antagonists of prostaglandin receptors.

COX Inhibitors

Item No.	Product Name	Summary
10024	Acetaminophen	An analgesic and antipyretic compound that inhibits COX-2 activity
10008672	Celecoxib	A selective COX-2 inhibitor (IC ₅₀ = 50 nM)
70280	(±)-Ibuprofen	A non-selective, reversible COX-1 and COX-2 inhibitor (IC ₅₀ s = 2.6 and 1.53 μM, respectively)
70220	Lornoxicam	An inhibitor of both COX-1 and COX-2 (IC ₅₀ s = 5 and 8 nM, respectively)
20304	Tenoxicam	A COX-2 inhibitor (IC ₅₀ s = 20 and 322 nM for COX-1 and COX-2, respectively)

A wide range of selective and non-selective COX-1 and -2 inhibitors is available at www.caymanchem.com

mPGES-1 Inhibitors

Item No.	Product Name	Summary
13164	CAY10589	A dual mPGES-1 and 5-lipoxygenase inhibitor
13217	MF63	A selective, orally active inhibitor of human mPGES-1 (IC ₅₀ = 1.3 nM)
18904	PF-9184	An inhibitor of mPGES-1 (IC ₅₀ = 16.5 nM for recombinant human enzyme)

Additional selective and non-selective mPGES-1 inhibitors as well dual mPGES-1/lipoxygenase inhibitors available at www.caymanchem.com

Prostaglandin EP₁ Receptor Antagonists

Item No.	Product Name	Summary
14070	ONO-8711	A selective EP ₁ receptor antagonist (K _s = 0.6 and 1.7 nM for human and mouse EP ₁ , respectively)
14060	SC-19220	A selective EP ₁ antagonist (IC ₅₀ = 6.7 μM for displacing radiolabeled PGE ₂ from hEP ₁)
10010744	SC-51322	A selective EP ₁ antagonist with analgesic activity in mouse pain models (ED ₅₀ = 0.9 mg/kg)

Additional EP₁ selective antagonists available at www.caymanchem.com

Cannabinoid Receptors

The cannabinoid receptors CB₁ and CB₂ can be activated by endogenous cannabinoids, plant cannabinoids, and synthetic cannabinoids to modulate pain perception. Cayman offers a collection of incredibly potent cannabinoid receptor ligands, many of which are highly selective for either the central or peripheral receptor, as well as inhibitors of endocannabinoid transport and enzyme degradation.

Item No.	Product Name	Summary
71670	AM251	A CB ₁ receptor antagonist (K _i = 7.5 nM) and GPR55 agonist (EC ₅₀ = 39 nM)
90060	AM404	A blocker of AEA reuptake with an IC ₅₀ of 1 μM in rat neurons and 5 μM in rat astrocytes
91054	Arachidonoyl 2'-Chloroethylamide	A stable, selective CB ₁ receptor agonist (K _i = 1.4 nM)
90050	Arachidonoyl Ethanolamide	An endogenous cannabinoid (K _s = 61 and 1,940 nM for CB ₁ and CB ₂ , respectively)
90067	N-Arachidonoyl-γ-Aminobutyric Acid	A suppressor of formalin-induced pain in rats
14497	JNJ-1661010	A selective FAAH inhibitor (IC ₅₀ s = 34 and 33 nM in rat and human, respectively)
10005428	JWH 133	A selective CB ₂ receptor agonist (K _i = 3.4 and 677 nM for CB ₂ and CB ₁ , respectively)
13158	JZL 184	A selective MAGL inhibitor (IC ₅₀ s = 8 nM and 4 μM for MAGL and FAAH in mouse brain membranes, respectively)

A complete list of cannabinoid receptor ligands and inhibitors of endocannabinoid reuptake and degradation is available at www.caymanchem.com

Opioid Signaling

Cayman offers specific opioid receptor agonists as well as related analgesics to study pain relief by modulating the signaling of various receptors involved in nociception.

Opioid Receptor Agonists

Item No.	Product Name	Summary
19410	Carfentanil	An Analytical Reference Material
22801	Crotonyl fentanyl	An Analytical Reference Standard
19567	<i>ortho</i> -Fluorobutyryl fentanyl (hydrochloride)	An Analytical Reference Standard
21553	DAMGO	A selective peptide agonist of the μ -opioid receptor ($K_{1/2}$ = 1.18, 1,430, and 213 nM for human μ -, δ -, and κ -opioid receptors, respectively)
23184	DPDPE (trifluoroacetate salt)	A synthetic peptide δ -opioid receptor agonist (K_i = 2.7 nM)
18169	Dynorphin A	A κ -, μ -, and δ - opioid receptor agonist ($K_{1/2}$ = 0.5-1 nM); activates human ORL1 (K_i = 386 nM)
18178	Dynorphin B	A κ_{1b} -opioid receptor agonist (K_i = 1.1 nM)
20782	Methoxyacetyl fentanyl (hydrochloride)	An Analytical Reference Standard
22245	Salvinorin A Carbamate	A potent κ -opioid receptor full agonist (EC_{50} = 6.2 nM)
22290	Salvinorin A Propionate	A selective, partial κ_1 -opioid receptor agonist (K_i = 32.6 nM)
22244	Salvinorin B Mesylate	A selective κ -opioid receptor agonist (EC_{50} = 30 nM)
15599	SNC 80	A nonpeptide δ -opioid receptor agonist (K_i = 0.18 nM; IC_{50} = 2.73 nM)
24480	SR 17018	A functionally selective μ -opioid receptor agonist

Additional opioid receptor agonists available at www.caymanchem.com

Non-Opioid Analgesics

Item No.	Product Name	Summary
22928	BD 1047 (hydrobromide)	A selective σ_1 receptor antagonist (K_i = 0.9 nM in a radioligand binding assay)
21254	JTC 801	An antagonist of the nociceptin receptor (ORL1; K_i = 44.5 nM; IC_{50} = 94 nM)
15196	ML-335	An agonist for OPRM1-OPRD1 heterodimerization (EC_{50} = 403 nM); may alleviate the unwanted effects associated with prolonged opiate use
12037	MCOPPB (hydrochloride)	An agonist for the NOP receptor (pK_i = 10.1, EC_{50} = 0.39 nM)
15070	Nociceptin	A NOP1 (ORL1) receptor ligand (K_i = 88 μ M in CHO cells)
16279	S1RA	A selective σ_1 receptor antagonist (K_i = 17 nM); enhances peripheral μ -opioid analgesia without affecting opioid-induced constipation
20390	Racecadotril	An enkephalinase inhibitor that prevents the degradation of endogenous opioids
15600	Thiophan	A potent inhibitor of neprilysin produced <i>in vivo</i> via the metabolism of racecadotril

Additional non-opioid analgesics available at www.caymanchem.com



To view a complete list of our Pain Research products, visit us online at www.caymanchem.com

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