

• Inhibitors • Agonists • Screening Libraries

New Products

December 2019



MedChemExpress (MCE) provides a wide range of lifescience biochemicals, including more than 15,000 **bioactive compounds**, **dye reagents**, **peptides and natural compounds** for laboratory and scientific use. If you need these products, please do not hesitate to contact us via **sales@MedChemExpress.com**.

• p38 MAPK Inhibitor

AMG-548

AMG-548 dihydrochloride, a potent, orally active and selective **p38** α inhibitor (K_i=0.5 nM), shows slightly selective over **p38** β (K_i=36 nM) and >1000 fold selective against p38 γ and p38 δ . AMG-548 dihydrochloride is also extremely potent in the inhibition of whole blood LPS stimulated **TNF** α (IC₅₀=3 nM). AMG-548 dihydrochloride inhibits Wnt signaling by directly inhibiting **Casein kinase 1** isoforms δ and ϵ .



Compound Screening Libraries

50+ specific libraries for high throughput screening (HTS) and high content screening (HCS).

Optimized for drug screening and new indication research.

Thrombin Inhibitor

Ximelagatran

CAS No.: 192939-46-1

Ximelagatran (H 376/95) is an orally active **thrombin** inhibitor that selectively and competitively inhibits both free and clot-bound **thrombin**. Ximelagatran is an anticoagulant agent with a rapid onset of anticoagulant effect, predictable, dose-dependent pharmcokinetics and pharmacodynamics.

• PI3K Inhibitor

PI-828

CAS No.: 942289-87-4

PI-828 is a dual PI3K and casein kinase 2 (CK2) inhibitor with IC₅₀s of 173 nM, 149 nM, and 1127 nM for p110 α , CK2, and CK2 α 2 in lipid kinase assay, respectively.

Free Sample

from MCE!

Get your free inhibitor sample



Cell Signaling Pathway

• Protease-Activated Receptor (PAR) Agonist

GB-110 hydrochloride

GB-110 hydrochloride is a potent, orally active, and nonpeptidic **protease activated receptor 2 (PAR2)** agonist. GB-110 hydrochloride selectively induces PAR2-mediated intracellular Ca²⁺ release in HT29 cells with an EC₅₀ of 0.28 μ M.



Btk Inhibitor

Remibrutinib

CAS No.: 1787294-07-8

Remibrutinib is an orally active **bruton tyrosine kinase (BTK)** inhibitor with an IC_{50} value of 1 nM. Remibrutinib inhibits BTK activity with an IC_{50} value of 0.023 μ M in blood. Remibrutinib has the potential for Chronic urticaria (CU) treatment.

Other New Products This Month

Mavacoxib (COX)		WAY-213613 (EAAT2)		AT-007 (Aldose Reductase)	
AS-604850 (PI3K)		Nosiheptide (Bacterial)		NS13001 (Potassium Channel)	
BT2 (Bcl-2 Family)		GSK205 (TRP Channel)		Lenalidomide-Br (Ligand for E3 Ligase)	
Splitomicin (HDAC)					

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