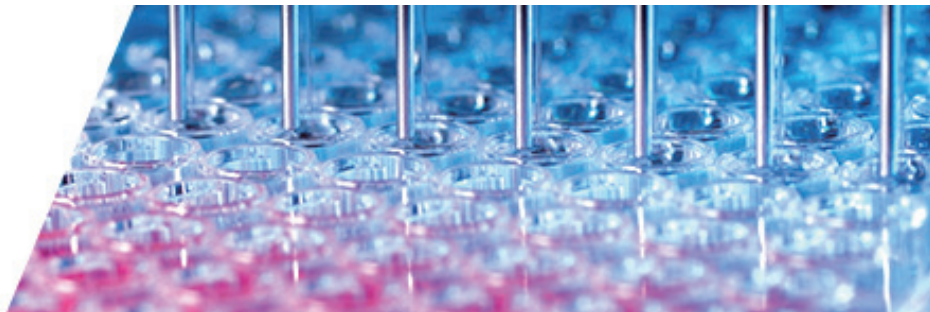


## New Products

September 2019



MedChemExpress (MCE) offers a wide range of high quality research chemicals and biochemicals including novel bioactive compounds, dye reagents, peptides and natural compounds for laboratory and scientific use.

- Over 10,000 bioactive molecules in stock
- Target 375 key proteins in 20 signaling pathways
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- Release more than 100 of the newest biochemicals per month
- High purity & competitive prices
- Data sheet with detailed biological information
- Delivery within 24 hours

### • JAK Inhibitor

**BMS-986165**

CAS No.: 1609392-27-9

BMS-986165 is a highly selective, orally bioavailable allosteric TYK2 inhibitor for the treatment of autoimmune diseases, which selectively binds to TYK2 pseudokinase (JH2) domain ( $IC_{50}=1.0$  nM) and blocks receptor-mediated Tyk2 activation by stabilizing the regulatory JH2 domain. BMS-986165 inhibits IL-12/23 and type I IFN pathways.

### • Epigenetic Reader Domain Inhibitor

**SGC-iMLLT**

CAS No.: 2255338-25-9

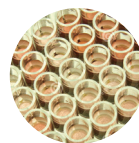
SGC-iMLLT is a first-in-class chemical probe and a potent, selective inhibitor of MLLT1/3-histone interactions with an  $IC_{50}$  of 0.26  $\mu$ M. SGC-iMLLT shows high binding activity towards MLLT1 YEATS domain (YD) and MLLT3 YD (AF9/YEATS3) with  $K_d$ s of 0.129 and 0.077  $\mu$ M, respectively.

### • LRRK2 Inhibitor

**PFE-360**

CAS No.: 1527475-61-1

PFE-360 (PF-06685360) is a potent, selective, brain penetrated and orally active leucine-rich repeat kinase 2 (LRRK2) inhibitor with a mean  $IC_{50}$  of 2.3 nM in vivo.



### Compound Screening Libraries

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

Optimized for drug screening and new indication research.

### Free Sample

Get your free inhibitor sample from MCE!

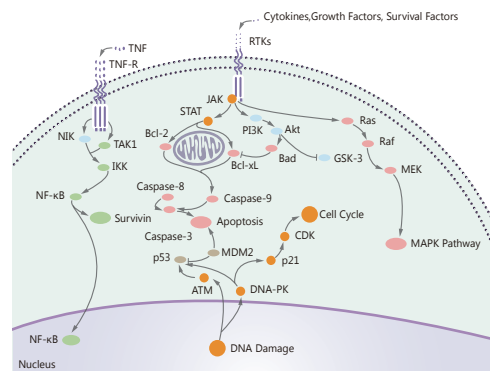


### Cell Signaling Pathway

## • TRP Channel Antagonist

### BI-749327

BI-749327 is a potent, high selectivity and orally bioavailable TRPC6 antagonist, with  $IC_{50}$ s of 13 nM, 19 nM and 15 nM for mouse, human and guinea pig TRPC6, respectively. BI-749327 is 85-fold more selective for mouse TRPC6 than TRPC3 and 42-fold versus TRPC7.



## • Adenosine Receptor Antagonist

### Tonapofylline

CAS No.: 340021-17-2

Tonapofylline (BG 9928) is an orally active and selective adenosine A1 receptor antagonist with a  $K_i$  of 7.4 nM for human adenosine A1 receptor (hA1), which displays 915-fold selectivity versus human adenosine A2A receptor and 12-fold selectivity versus human adenosine A2B receptor and is used in development for the treatment of heart failure.

## Other New Products This Month

KT185 (MAGL)		LY2922470 (GPR40)		L-690330 (Phosphatase)	
AZD8329 (HSD)		NCC007 (Casein Kinase)		Indibulin (Microtubule/Tubulin)	
SC75741 (NF-κB)		UNC2541 (TAM Receptor)		Derenofylline (Adenosine Receptor)	
dBET57 (PROTAC) ...					

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