CC-401 . HCl

Potent inhibitor of JNK

CC-401 is a competitive inhibitor of the ATP binding site in the active, phosphorylated, form of JNK. This prevents JNK from phosphorylating its various target molecules, including the amino terminus of c-Jun. It is a potent inhibitor of all three forms of JNK (K_i of 25-50 nM), and has at least 40-fold selectivity for JNK compared with other related kinases, including: p38, ERK, IKK2, PKC, Lck, and ZAP70. CC-401 acts to inhibit JNK signaling by competitive binding to the adenosine triphosphate-binding site in the active, phosphorylated, form of JNK, resulting in inhibition of the phosphorylation of JNK targets, such as the amino-terminal activation domain of the transcription factor, c-Jun. In cell-based assays, 1–5 µmol/l CC-401 provides specific JNK inhibition.

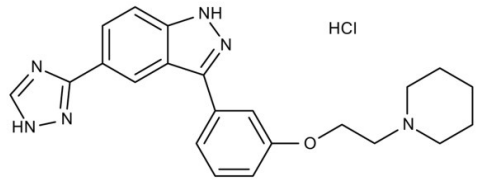
Ordering Information

Order Online »

ENZ-CHM194-0001	1mg

Manuals, SDS & CofA

View Online »


NH
N
HCl
N
HN
N
O
N

Handling & Storage

Use/Stability	As indicated on product label or CoA when stored as recommended.
Short Term Storage	+4°C
Long Term Storage	-20°C
Shipping	Ambient Temperature

Regulatory Status

RUO - Research Use Only

Product Details

Alternative Name	3-[3-(2-piperidin-1-ylethoxy)phenyl]-5-(1H-1,2,4-triazol-5-yl)-1H-indazole hydrochloride
Appearance	White powder.
CAS	1438391-30-0
Couple Target	Jnk
Couple Type	Inhibitor
Formula	$C_{22}H_{24}N_6O$. HCl
Identity	Determined by EM-MS, NMR
MW	424.9
Purity	≥99% (HPLC)
Solubility	Soluble in DMSO (15mM).

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