Web: www.taiclone.com
Tel: +886-2-2735-9682
Email: order@taiclone.com

Falnidamol

Catalog No: tcsc0691

Available Sizes

Size: 5mg

Size: 10mg

Size: 50mg

Specifications

CAS No:
196612-93-8

Formula:
$C_{18}H_{19}ClFN_7$

Pathway:
JAK/STAT Signaling;Protein Tyrosine Kinase/RTK

Target:
EGFR;EGFR

Purity / Grade:
>98%

Solubility:
DMSO : ≥ 41 mg/mL (105.71 mM)

Alternative Names:
BIBX 1382

Observed Molecular Weight:
387.84

Product Description

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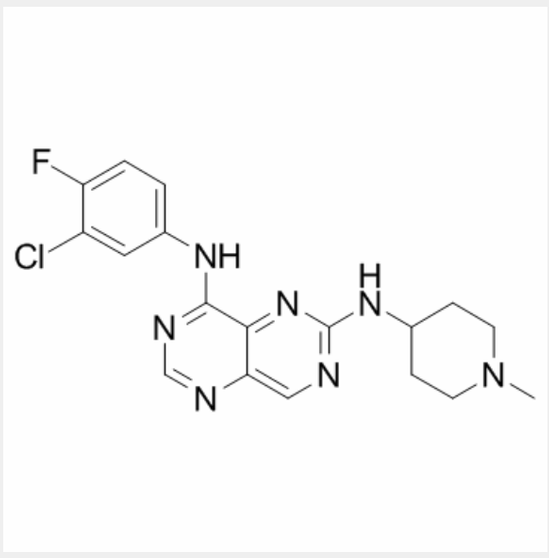
Falnidamol (BIBX 1382) is a potent, selective inhibitor of EGFR tyrosine kinase (IC_{50} = 3 nM); displays > 1000-fold lower potency against ErbB2 (IC_{50} = 3.4 μM) and a range of other related tyrosine kinases (IC_{50} > 10 μM).

IC50 & Target: IC50: 3 nM (EGFR)[1].

In Vitro: Falnidamol (BIBX 1382) and BIBU1361 are both potent and selective submicromolar inhibitors of the EGFR kinase activity. An IC_{50} value of 3 nM was determined for both compounds. The potency of these two compounds compares with the one obtained with Iressa, which is a leading EGFR inhibitor in the field. Inhibition of the closest family member, HER2, was 100- to 1000-fold less potent. Furthermore, Falnidamol (BIBX 1382) and BIBU1361 did not inhibit a number of other related tyrosine kinases[1].

In Vivo: In nude mice, oral once daily dosing at 10 mg/kg with either Falnidamol (BIBX 1382) or BIBU1361 completely suppressed tumor growth of human A431 xenografts with respective T/C values of 15 and 6% after 2 weeks of treatment[1].



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