

Catalog # 10-5124 WH-4-023

CAS# 837422-57-8

N-(2,4-dimethoxyphenyl)-N-[2-[[4-(4-methyl-1-piperazinyl)phenyl]amino]-4-pyrimidinyl]-carbamic acid 2,6-dimethylphenyl ester

Lot # X109247

A potent and selective inhibitor of Src family kinases Lck and Src, $IC_{50}=2$ and 6 nM respectively.^{1,2} Also inhibits SIK, $IC_{50}=10$, 22 and 60 nM for SIK 1, 2 and 3 respectively.³ May be used along with PD-325901 and CHIR-99021 to support self-renewal of naïve human embryonic stem cells.⁴

- 1) Martin et al. (2006), Novel 2-aminopyrimidine carbamates as potent and orally active inhibitors of Lck: synthesis, SAR, and in vivo anti-inflammatory activity; J. Med. Chem., **49** 4981
- 2) Moroco et al. (2015), A discovery Strategy for Selective Inhibitors of c-Src in Complex with the Focal Adhesion Kinase SH3/SH2-binding Region; Chem. Biol. Drug. Des., **86** 144
- 3) Clark et al. (2012), Phosphorylation of CRTC3 by the salt-inducible kinases controls the interconversion of classically activated and regulatory macrophages; Proc. Natl. Acad. Sci. USA, **109** 16986
- 4) Theunissen et al. (2014), Systemic identification of culture conditions for introduction and maintenance of naïve human pluripotency; Cell Stem Cell, **15** 471

PHYSICAL DATA

Molecular Weight: 568.67

Molecular Formula: C₃₂H₃₆N₆O₄

Purity: 98% by HPLC

NMR: (Conforms)

Solubility: DMSO (up to 60 mg/ml)
Physical Description: White or off-white solid

Storage and Stability: Store as supplied desiccated at -20°C for up to 2 years from the date of purchase.

Solutions in DMSO may be stored at -20°C for up to 2 months.

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