

Product Data Sheet

Product Name: AZD1208 hydrochloride

Cat. No.: GC35447

Chemical Properties

Cas. No. 1621866-96-3

SMILES O=C(NC(=O)SC1=CC=C(N3C[C@H](N)CCC3)C(C4=CC=CC=C4)=CC=C2.[H]Cl

Formula $C_{21}H_{22}ClN_3O_2S$ M.Wt 415.94

Solubility DMSO: 83.3 mg/mL (200.27 mM and warming) Storage Store at -20°C

General tips For obtaining a higher solubility, please warm the tube at 37 °C and shake it in the ultrasonic bath for a while. Stock solution can be stored below -20°C for several months.

Shipping Condition Evaluation sample solution: ship with blue ice All other available size: ship with RT, or blue ice upon request.

Structure

Background

AZD 1208 is a potent, selective, and orally available inhibitor of all three forms of the proto-oncogene Pim kinase (IC_{50} s = 0.4, 5.0, and 1.9 nM for Pim-1, Pim-2, and Pim-3, respectively).¹ It causes cell cycle arrest and apoptosis in MOLM-16 megakaryoblastic leukemia cells and inhibits the growth of MOLM-16 xenograft tumors in mice.¹ It has efficacy against acute myeloid leukemia and glioblastoma cells when combined with mTOR and p110 α inhibitors, respectively.^{2,3}

1. Keeton, E.K., McEachern, K., Dillman, K.S., et al. AZD1208, a potent and selective pan-Pim kinase inhibitor, demonstrates efficacy in preclinical models of acute myeloid leukemia. *Blood* 123(6):905-913 (2014)
 2. Harada, M., Benito, J., Yamamoto, S., et al. The novel combination of dual mTOR inhibitor AZD2014 and pan-PIM inhibitor AZD1208 inhibits growth in acute myeloid leukemia via HSF pathway suppression. *Oncotarget* 6(35):37930-37947 (2015)
 3. Iqbal, A., Eckerdt, F., Bell, J., et al. Targeting of glioblastoma cell lines and glioma stem cells by combined PIM kinase and PI3K-p110 α inhibition. *Oncotarget* 7(22):33192-33201 (2016)

Caution: Product has not been fully validated for medical applications. For research use only.

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