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DATASHEET

SB 202190

Product overview

Name SB 202190 Cat No HB1301 **Alternative names** SB-202190 **Biological action** Inhibitor **Purity** >99%

Description Potent ATP competitive p38 MAPK inhibitor. 3D growth matrix component, promotes long-term

organoid growth and can be used in production of gastric organoids. Promotes stability of human

pluripotent stem cells.

Images



Biological Data

Biological description

Potent ATP competitive p38 MAPK inhibitor (IC $_{50}$ values are 50 and 100 nM for p38 α and p38 β respectively).

It inhibits TNF, IL-1 and CK1 (approx IC $_{50}$ = 0.6 μ M) and therefore inhibits CREB phosphorylation. It exhibits little or no activity at CK2.

It displays anti-inflammatory properties and induces apoptosis.

It can also be used as a 3D growth matrix component, promotes long-term organoid growth and can be used in production of gastric organoids.

Also promotes stability of human pluripotent stem cells.

Solubility & Handling

Solubility overview Storage instructions Storage of solutions Soluble in DMSO (100mM)

Prepare and use solutions on the same day if possible. Store solutions at -20 °C for up to one month if storage is required. Equilibrate to RT and ensure the solution is precipitate free before use.

Stable for ambient temperature shipping. Follow storage instructions on receipt.

This product is for RESEARCH USE ONLY and is not intended for therapeutic or diagnostic use. Not

Shipping Conditions Important

for human or veterinary use.

Chemical Data

Chemical name 4-[4-(4-Fluorophenyl)-5-(4-pyridiny l)-1*H*-imidazol-2-yl]phenol

Molecular Weight 331.35

Chemical structure

м Н — ОН

InChiKey YKRNPHOBDOUQTG-UHFFFAOYSA-N

References

Non-specific in vivo inhibition of CK1 by the pyridinyl imidazole p38 inhibitors SB 203580 and SB 202190.

Shanware NP *et al* (2009) BMB Rep 42(3) **PubMedID** 19336000

Specificity and mechanism of action of some commonly used protein kinase inhibitors.

Davies SP *et al* (2000) Biochem J 351(Pt 1) **PubMedID**10998351

Induction of apoptosis by SB202190 through inhibition of p38beta mitogen-activated protein kinase.

Nemoto S *et al* (1998) J Biol Chem 273(26) **PubMedID**9632706