

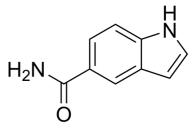
# **Data Sheet**

Product Name: SD-169

Target: p38 MAPK

Pathway: MAPK/ERK Pathway

**Solubility:** DMSO: 100 mg/mL (624.30 mM; Need ultrasonic)



## **BIOLOGICAL ACTIVITY:**

SD-169 is an orally active ATP-competitive inhibitor of **p38\alpha MAPK**, with an **IC**<sub>50</sub> of 3.2 nM. SD-169 also weakly inhibits **p38\beta MAPK** with an **IC**<sub>50</sub> of 122 nM. SD-169 prevents the development and progression of diabetes by inhibiting T cell infiltration and activation<sup>[1]</sup>. IC50 & Target: IC50: 3.2 nM (p38 $\alpha$  MAPK), 122 nM (p38 $\beta$  MAPK)<sup>[1]</sup> *In Vitro:* SD-169 significantly reduces p38 and HSP60 expression in T cells of the pancreatic beta islets<sup>[1]</sup>.

SD-169 demonstrates 38-fold potency against p38 $\alpha$  MAP kinase (IC<sub>50</sub>=3.2 nM) than p38 $\beta$  MAP kinase (IC<sub>50</sub>=122 nM)<sup>[1]</sup>.

#### References:

[1]. Medicherla S, et al. Preventive and therapeutic potential of p38 alpha-selective mitogen-activated protein kinase inhibitor in nonobese diabetic mice with type 1 diabetes. J Pharmacol Exp Ther. 2006 Jul;318(1):99-107.

### **CAIndexNames:**

1H-Indole-5-carboxamide

## SMILES:

O=C(C1=CC2=C(NC=C2)C=C1)N

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

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