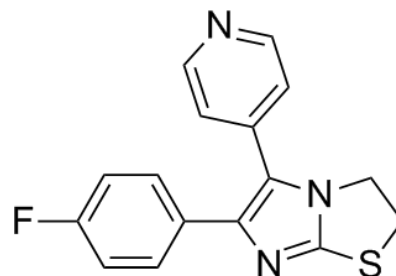


Product Name : SKF-86002
Cat. No. : PC-43080
CAS No. : 72873-74-6
Molecular Formula : C₁₆H₁₂FN₃S
Molecular Weight : 297.35
Target : p38 MAPK
Solubility : 10 mM in DMSO



Biological Activity

SKF-86002 is a potent inhibitor of **p38 MAPK** with IC₅₀ of 0.5-1 uM, inhibits LPS-induced IL-1 and TNF-α production in human monocyte with IC₅₀ of 1 uM.

SKF-86002 inhibits prostaglandin H₂ (PGH₂) synthase activity (IC₅₀=120 uM) as well as prostanoid production by rat basophilic leukemia (RBL-1) cells (IC₅₀=70 uM).

SKF-86002 blocks superoxide anion production in response to FMLP and reduces adhesion and chemotaxis in response to PAF or FMLP in human neutrophils.

SKF-86002 also inhibits 5-lipoxygenase- and cyclooxygenase-mediated arachidonic acid metabolism in RBL-1 cells (IC₅₀=10 and 100 uM respectively), shows anti-inflammatory in vivo.

References

Griswold DE, et al. *Biochem Pharmacol.* 1987 Oct 15;36(20):3463-70.

Nick JA, et al. *J Clin Invest.* 1997 Mar 1;99(5):975-86.

Lee JC, et al. *Nature.* 1994 Dec 22-29;372(6508):739-46.

Perregaux DG, et al. *Mol Pharmacol.* 1995 Sep;48(3):433-42.

Caution: Product has not been fully validated for medical applications. Lab Use Only!

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