

SB202190 (FHP1)

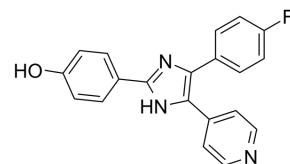
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OrganRegen, INC.
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DESCRIPTION

Background	SB202190 is a selective p38 MAP kinase inhibitor with IC50s of 50 nM and 100 nM for p38 α and p38 β 2, respectively. SB 202190 binds to the ATP pocket of the active recombinant human p38 kinase with a Kd of 38 nM. SB 202190 has anti-cancer activity and rescued memory deficits ^{[1][2]} . SB202190 induces autophagy ^[3] .		
M. W t	331.34		
Formula	C ₂₀ H ₁₄ FN ₃ O		
CAS No	152121-30-7		
Storage	Powder	- 20°C	3 years
		4°C	2 year
	In solvent	-80°C	6 months
		-20°C	1 month
Solubility	DMSO	100 mg/mL(301.80 mM; Need ultrasonic)	
	Ethanol	12 mg/mL(36.22 mM)	
	H ₂ O	< 0.1 mg/mL(insoluble)	



C₂₀H₁₄FN₃O

BIOLOGICAL ACTIVITY

In Vitro

SB202190 (0-10 μ M; 0-72 hours) attenuates growth of a subgroup of CRC cell lines such as RKO, CACO2 and SW480 in a dose- and time-dependent manner^[1].

SB 202190 strongly inhibited colony formation and anchorage-independent growth (10 μ M for 7–10 days) and elevated apoptotic cell death (10 μ M for 72 h) in this same subset of CRC lines (RKO, CACO2 and SW480)^[2].

In RKO, CACO2 and SW480 cells, SB202190 (10 μ M; 2 hours) abrogates phosphorylation of S6K1(T389) and S6(S235/236), but not AKT(S473), indicating that p38i selectively blocks mTORC1 signaling^[2].

In Vivo

SB202190 (5 mg/kg; intraperitoneal injection; daily for 10-12 days) shows inhibition of tumor cell survival and tumor growth^[2].

REFERENCES

- [1]. Davies SP, et al. Specificity and mechanism of action of some commonly used protein kinase inhibitors. *Biochem J.* 2000 Oct 1;351(Pt 1):95-105.
- [2]. Nemoto S, et al. Induction of apoptosis by SB202190 through inhibition of p38beta mitogen-activated protein kinase. *J Biol Chem.* 1998 Jun 26;273(26):16415-20.
- [3]. Grossi V, et al. Bay 43-9006 inhibits p38 α activity in colorectal cancer cells and synergizes with the DFG-in inhibitor SB202190 to increase apoptotic response. *Cancer Biol Ther.* 2012 Dec;13(14):1471-81.