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产品名称: **M-110**
 产品别名: **M-110**

生物活性:	
Description	M-110 is a highly selective, ATP-competitive inhibitor of PIM kinases with a preference for PIM-3 (IC ₅₀ =47 nM). M-110 inhibits PIM-1 and PIM-2 with similar IC ₅₀ s of 2.5 μM. M-110 inhibits the proliferation of prostate cancer cell lines with IC ₅₀ s of 0.6 to 0.9 μM ^[1] .
In Vitro	M-110 (0.01-10 μM; 72 hours) inhibiting the growth of DU-145 cells with an IC ₅₀ value of 0.9 μM ^[1] . M-110 has no activity on normal human peripheral blood mononuclear cells up to 40 μM ^[1] . M-110 (10 μM; 18 hours) inhibits STAT3 Tyr705 phosphorylation ^[1] . M-110 inhibits the expression of active STAT3 through inhibition of PIM-3. M-110 also inhibits the proliferation of 22Rv1, PC3, and SW480 cells, with IC ₅₀ values of 0.6 to 0.8 μM ^[1] .
	Cell Viability Assay^[1]
	Cell Line: DU-145 cells
	Concentration: 0.01, 0.1, 1, 10 μM
	Incubation Time: 72 hours
	Result: Inhibiting the growth of DU-145 cells with an IC ₅₀ value of 0.9 μM.
	Western Blot Analysis^[1]
	Cell Line: DU-145 cells
	Concentration: 10 μM
	Incubation Time: 18 hours
Result: Reduced the expression of p-STAT3 Tyr705 to 23.5%, compared with untreated cells without affecting the expression of STAT3.	
References	<p>[1]. Ther. 2010 Sep;9(9):2478-87. Cancer Chang M, et al. PIM kinase inhibitors downregulate STAT3(Tyr705) phosphorylation. Mol</p> <p>[2]. He Y, et al. Schisantherin A suppresses osteoclast formation and wear particle-induced osteolysis via modulating RANKL signaling pathways. Biochem Biophys Res Commun. 2014 Jul 4;449(3):344-50.</p> <p>[3]. Zhou E, et al. Schisantherin A protects lipopolysaccharide-induced acute respiratory distress syndrome in mice through inhibiting NF-κB and MAPKs signaling pathways. Int Immunopharmacol. 2014 Sep;22(1):133-40.</p> <p>[4]. Sa F, et al. Discovery of novel anti-parkinsonian effect of schisantherin A in in vitro and in vivo. Neurosci Lett. 2015 Apr 23;593:7-12.</p> <p>[5]. Zhang LQ, et al. Schisantherin A protects against 6-OHDA-induced dopaminergic neuron damage in zebrafish and cytotoxicity in SH-SY5Y cells through the ROS/NO and AKT/GSK3β pathways. J Ethnopharmacol. 2015 Apr 29. pii: S0378-8741(15)00306-2.</p>