

Product Name: Imatinib Mesylate (STI571)

Revision Date: 02/24/2023

Product Data Sheet

Imatinib Mesylate (STI571)

Cat. No.: A1805

CAS No.: 220127-57-1

Formula: C29H31N7O·CH4SO3

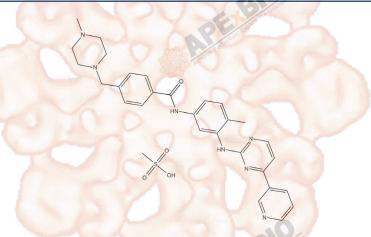
M.Wt: 589.71

Synonyms:

Target: TGF-β / Smad Signaling

Pathway: Bcr-Abl

Storage: Store at -20°C



Solvent & Solubility

≥29.5mg/mL in DMSO

In Vitro

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	1.6957 mL	8.4787 mL	16.9575 mL
	5 mM	0.3391 mL	1.6957 mL	3.3915 mL
-10	10 mM	0.1696 mL	0.8479 mL	1.6957 mL

Please refer to the solubility information to select the appropriate solvent.

Biological Activity

Shortsummary	Abl/c-kit/PDGFR inhibitor			
IC ₅₀ & Target	600 nM (v-Abl), 100 nM (PDGFR), 100 nM (c-Kit)			
In Vitro	Cell Viability Assay			
	Cell Line:	T cells		
	Preparation method:	The solubility of this compound in DMSO is >10 mM. General tips for obtaining		
		a higher concentration: Please warm the tube at 37 °C for 10 minutes and/or		
		shake it in the ultrasonic bath for a while.Stock solution can be stored below		
		-20°C for several months.		
	Reacting conditions:	IC50: 3.9 μM for inhibiting DCs-stimulated T-cell proliferation 2.9 μM for		
		inhibiting PHA-stimulated T-cell proliferation 4 days		

	Applications:	Cells were stimulated with allogeneic mature DCs or PHA in the presence of
		imatinib mesylate. The drug inhibited T-cell proliferation as a function of
		concentration. The effects were significant at 0.5 μM imatinib mesylate for the
		cells stimulated by DCs and at 1.0 µM imatinib mesylate for the cells stimulated
		with PHA. The IC50 values for imatinib mesylate-inhibited T-cell proliferation
	SIQ.	stimulated by DCs and PHA were 3.9 µM and 2.9 µM, respectively.
	Animal experiment	and the state of t
	Animal models:	Female C57BL/6 mice
	Dosage form:	Intraperitoneal injection, 25 or 50mg/kg/day
	Applications:	Administration of imatinib alone did not generate any changes in lung
		morphology. However, when imatinib was administered in bleomycin-treated
		mice, a reduction of fibrotic lesions in the subpleural areas of lung was
		observed at doses of 25 and 50 mg/kg/day. The quantitative histologic analysis
In Vivo		demonstrated that the fibrotic score in mice treated with bleomycin and 50
	-0	mg/kg/day of imatinib was significantly lower than that treated with bleomycin
	B & Jungann	alone. The collagen content of the lung was also significantly lower in mice
	Too typoe in	treated with bleomycin and imatinib (50 mg/kg/day) as compared with those
		treated with bleomycin alone.
	Other notes:	Please test the solubility of all compounds indoor, and the actual solubility may
		slightly differ with the theoretical value. This is caused by an experimental
		system error and it is normal.

Product Citations

1. Ivey MJ, Kuwabara JT, et al. "Platelet-derived growth factor receptor-α is essential for cardiac fibroblast survival." Am J Physiol Heart Circ Physiol. 2019 Aug 1;317(2):H330-H344.PMID:31125253

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References

[1] Dietz A B, Souan L, Knutson G J, et al. Imatinib mesylate inhibits T-cell proliferation in vitro and delayed-type hypersensitivity in vivo. Blood, 2004, 104(4): 1094-1099.

[2] Aono Y, Nishioka Y, Inayama M, et al. Imatinib as a novel antifibrotic agent in bleomycin-induced pulmonary fibrosis in mice. American journal of respiratory and critical care medicine, 2005, 171(11): 1279-1285.

Caution

FOR RESEARCH PURPOSES ONLY.

NOT FOR HUMAN, VETERINARY DIAGNOSTIC OR THERAPEUTIC USE.

Specific storage and handling information for each product is indicated on the product datasheet. Most APExBIO products are stable

under the recommended conditions. Products are sometimes shipped at a temperature that differs from the recommended storage temperature. Shortterm storage of many products are stable in the short-term at temperatures that differ from that required for long-term storage. We ensure that the product is shipped under conditions that will maintain the quality of the reagents. Upon receipt of the product, follow the storage recommendations on the product data sheet.





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