

Product Name: ST 2825 Revision Date: 01/10/2020 Product Data Sheet

ST 2825

Cat. No.:	A3840
CAS No.:	894787-30-5
Formula:	C27H28Cl2N4O5S
M.Wt:	591.51
Synonyms:	ST2825;ST-2825
Target:	Others
Pathway:	MyD88
Storage:	Store at -20°C

Solvent & Solubility

Soluble in DMSO

In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg
		1 mM	1.6906 mL	8.4529 mL	16.9059 mL
		5 mM	0.3381 mL	1.6906 mL	3.3812 mL
		10 mM	0.1691 mL	0.8453 mL	1.6906 mL

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

Biological Activity

Shortsummary

Inhibitor of MyD88 dimerization

IC₅₀ & Target

In Vitro

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Cell Viability Assay		
Cell Line:	HEK 293T and HeLa cell [1], B cells and plasmacytoid dendritic cells [1]	
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:	6-7 h (HEK 293T cell), 15 min (HeLa cell), 5 days (B cell culture) [1]	
Applications:	In HEK293T cells, ST2825 specifically inhibited homodimerization of MyD88	

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TIR domains with 40% inhibition of dimerization at 5 μM and 80% inhibition
at 10 μM. This effect was specific for homodimerization of the TIR domains
and did not affect homodimerization of the death domains. Moreover, ST2825
interfered with recruitment of IRAK1 and IRAK4 by MyD88, causing inhibition of
IL-1-mediated activation of NF-κB transcriptional activity [1]. B cell
proliferation and differentiation into plasma cells in response to CpG-induced
activation of TLR9 were also suppressed by ST2825 (> 8 μM). These
results showed that ST2825 blocked IL-1R/TLR signaling by interfering with
MyD88 homodimerization and suggested that it may have therapeutic potential
in treatment of chronic inflammatory diseases [1]. TLR9-induced plasma cell
(PC) generation was blocked by ST2825 in Peripheral blood mononuclear cells
from SLE patients [3].

Animal experiment

Animal models:	C57BL female mice [1]		
Dosage form:	Orally taken at 100 or 200 mg/kg/day [1]; injection at 25 mg/kg/day [2]		
Applications:	IL-1 beta-induced production of IL-6 was dose-dependently inhibited by		
	ST2825 (100 or 200mg/kg daily) in treated mice [1]. In a murine model of		
	non-reperfused acute myocardial infarction, ST2825 (25 mg/kg) protected left		
	ventricular from dilatation and hypertrophy. No measurable reduction in infarct		
	size was found [2].		
Preparation method:	ST2825 dissolved in 0.5% carboxymethylcellulose [1]		
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

In Vivo

1. Pantaleão L, Rocha GHO, et al. "Connections of annexin A1 and translocator protein-18 kDa on tolllike receptor stimulated BV-2 cells." Exp Cell Res. 2018 Jun 15;367(2):282-290.PMID:29649428

2. Janda J, Burkett NB, et al. "Resatorvid-based Pharmacological Antagonism of Cutaneous TLR4Blocks UV-induced NF-κB and AP-1 Signaling in Keratinocytes and Mouse Skin."Photochem Photobiol. 2016 Nov;92(6):816-825.PMID:27859308

3. Wang SH, Wang SC, et al. "Induction of cyclooxygenase-2 gene by Candida albicans through EGFR, ERK, and p38 pathways in human urinaryepithelium." Med Mycol. 2016 Sep 23.PMID:27664170

4. Hu LT, et al. "Role of TREM-1 in response to Aspergillus fumigatus infection in corneal epithelial cells.." Int Immunopharmacol.2014 Sep 18. pii:S1567-5769(14)00363-4.PMID:25242387

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References

[1]. Loiarro M, Capolunghi F, Fantò N et al. Pivotal advance: inhibition of MyD88 dimerization and recruitment of IRAK1 and IRAK4 by a novel peptidomimetic compound. Journal of Leukocyte Biology (2007), 82(4), 801-810.

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[2]. Van Tassell BW, Seropian IM, Toldo S et al. Pharmacologic Inhibition of Myeloid Differentiation Factor 88 (MyD88) Prevents Left Ventricular Dilation and Hypertrophy After Experimental Acute Myocardial Infarction in the Mouse. Journal of Cardiovascular Pharmacology (2010), 55(4), 385-390.

[3]. Capolunghi F1, Rosado MM, Cascioli S et al., Pharmacological inhibition of TLR9 activation blocks autoantibody production in human B cells from SLE patients. Rheumatology (Oxford). 2010 Dec;49(12):2281-9.

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.

APExBIO Technology

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