

Product Name: Fostamatinib (R788)
Revision Date: 07/23/2024

## **Product Data Sheet**

# Fostamatinib (R788)

Cat. No.: B2284

CAS No.: 901119-35-5

Formula: C23H26FN6O9P

M.Wt: 580.46

Synonyms:

In Vitro

Target: Tyrosine Kinase

**Pathway:** Spleen Tyrosine Kinase (Syk)

Storage: Store at -20°C



insoluble in H2O; ≥100.4 mg/mL in DMSO; ≥3.05 mg/mL in EtOH with gentle warming and ultrasonic

Preparing Stock Solutions	Solvent Concentration	1mg	5mg	10mg
	1 mM	1.7228 mL	8.6139 mL	17.2277 mL
	5 mM	0.3446 mL	1.7228 mL	3.4455 mL
	10 mM	0.1723 mL	0.8614 mL	1.7228 mL

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

# **Biological Activity**

Shortsummary	Spleen tyrosine kinase (Sy	Spleen tyrosine kinase (Syk) inhibitor		
IC <sub>50</sub> & Target		SIQ.		
	Cell Viability Assay	Esperine W.		
In Vitro	Cell Line;	Diffuse large B-cell lymphoma (DLBCL) cell lines		
	Preparation method:	The solubility of this compound in DMSO is > 100.4mg/mL. 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:	4 μM, 96 h		

	Applications:	R788 was a prodrug of the active metabolite. R406 induced apoptosis of the
		majority of examined DLBCL cell lines. In R406-sensitive DLBCL cell lines,
		R406 specifically inhibited both tonic- and ligand-induced BCR signaling
		(autophosphorylation of SYK525/526 and SYK-dependent phosphorylation of
	Introven	the B-cell linker protein [BLNK]).
	Animal experiment	
	Animal models:	Eμ- TCL1 transgenic mouse model of CLL, B6/C3H F1 mice
	Dosage form:	Intraperitoneal administration, 18 consecutive days at a daily dose of 80 mg/kg
	Applications:	R788 (80 mg/kg/d) inhibited the growth of adoptively transferred TCL1
		leukemias in vivo. R788 treatment administered from days 4 to 25 after
		adoptive transfer significantly prevented the outgrowth of leukemias. Treatment
		with R788 significantly prolonged the survival of the animals. In B6/C3H mice,
In Vivo		after 4 days of treatment animals receiving R788 showed a greater rise in the
		number of circulating malignant lymphocytes than controls. In B6/C3H F1 mice,
	The Unstrown	treatment with R788 for 7 days blocked BCR signaling and inhibited leukemic
	Ton Endore	cell survival and proliferation in vivo. In Eµ- TCL1 transgenic mouse model of
	Achiere 2 orte	CLL, R788 inhibited the growth of spo <mark>ntan</mark> eously developing TCL1 leukemias.
	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**

See more customer validations on www.apexbt.com.

### References

[1]. Chen L, Monti S, Juszczynski P, et al. SYK-dependent tonic B-cell receptor signaling is a rational treatment target in diffuse large B-cell lymphoma[J]. Blood, 2008, 111(4): 2230-2237.

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[2]. Suljagic M, Longo P G, Bennardo S, et al. The Syk inhibitor fostamatinib disodium (R788) inhibits tumor growth in the Eµ-TCL1 transgenic mouse model of CLL by blocking antigen-dependent B-cell receptor signaling[J]. Blood, 2010, 116(23): 4894-4905.

#### 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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