

Product Name: 5-R-Rivaroxaban Revision Date: 01/10/2021

Product Data Sheet

5-R-Rivaroxaban

Cat. No.:	A3126
CAS No.:	8 <mark>6547</mark> 9-71-6
Formula:	C19H18CIN3O5S
M.Wt:	435.88
Synonyms:	Xarelto; BAY 59-7939
Target:	Proteases
Pathway:	Thrombin
Storage:	Store at -20°C
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Solvent & Solubility

	≥20.85 mg/mL in DN	\geq 20.85 mg/mL in DMSO; insoluble in H2O; insoluble in EtOH			
In Vitro	Preparing Stock Solutions	Mass Solvent Concentration	1mg	5mg	10mg
		1 mM	2.2942 mL	11.4710 mL	22.9421 mL
		5 mM	0.4588 mL	2.2942 mL	4.5884 mL
		10 mM	0.2294 mL	1.1471 mL	2.2942 mL

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

Biological Activity

Shortsummary

Factor Xa (FXa) inhibitor

IC₅₀ & Target

In Vitro

Cell Viability Assay	Part and
Cell Line:	Human, rabbit and rat plasma
Preparation method:	The solubility of this compound in DMSO is >20.85mg/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:	IC50: 21 nM (human and rabbit plasma)

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	Applications:	Rivaroxaban competitively inhibited human FXa with the Ki value of 0.4 nM.		
		Rivaroxaban inhibited prothrombinase activity with the IC50 of 2.1 nM.		
		Rivaroxaban inhibited endogenous FXa more potently in human and rabbit		
		plasma (IC50: 21 nM) than rat plasma (IC50:290 nM). Rivaroxaban		
		demonstrated anticoagulant effects in human plasma, doubling prothrombin		
	E BIO	time (PT) and activated partial thromboplastin time at 0.23 and 0.69 μ M, respectively.		
	Animal experiment			
	Animal models:	Rat venous stasis model, Anaesthetised rat model		
	Dosage form:	Intravenous inection, Oral administration, 2 mg/kg		
	Applications:	In a rat venous stasis model, Rivaroxaban (i.v.) dose-dependently reduced		
		venous thrombosis with the ED50 of 0.1 mg/kg. Rivaroxaban (p.o.) reduced		
In Vivo		arterial (fibrin- and platelet-rich) thrombus formation in an arteriovenous (AV)		
		shunt in rats (ED50: 5 mg/kg) and rabbits (ED50: 0.6 mg/kg). In anaesthetised		
	810	rat model, pretreatment with 5-R-Rivaroxaban (i.v., 2 mg/kg) shortened		
	OF	bleeding time and clotting time.		
	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]. Perzborn, E., Strassburger, J., Wilmen, A., Pohlmann, J., Roehrig, S., SCHLEMMER, K. H., & amp; Straub, A. (2005). In vitro and in vivo studies of the novel antithrombotic agent BAY 59 - 7939—an oral, direct Factor Xa inhibitor. Journal of Thrombosis and Haemostasis, 3(3), 514-521.

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[2]. Perzborn, E., et al., Reversal of rivaroxaban anticoagulation by haemostatic agents in rats and primates. Thromb Haemost, 2013. 110(1): p. 162-72.

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 **2** | www.apexbt.com

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

www.apexbt.com 7505 Fannin street, Suite 410, Houston, TX 77054. Tel: +1-832-696-8203 | Fax: +1-832-641-3177 | Email: info@apexbt.com





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