

# **Product datasheet**

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# ARG22573 anti-Factor VIII antibody [RFF-VIIIC/8]

Package: 250 μg Store at: -20°C

## **Summary**

Product Description Mouse Monoclonal antibody [RFF-VIIIC/8] recognizes Factor VIII

Tested Reactivity Hu, Pig

Species Does Not React With Ms, Rat, Dog

Tested Application ELISA, RIA, WB

Host Mouse

Clone RFF-VIIIC/8

Isotype IgG1

Target Name Factor VIII
Species Human

Immunogen Affinity purified human Factor VIII.

Conjugation Un-conjugated

Alternate Names AHF; Antihemophilic factor; Coagulation factor VIII; HEMA; F8B; F8C; Procoagulant component; FVIII;

DXS1253E

### **Application Instructions**

Application table	Application	Dilution
	ELISA	1:500 - 1:15000
	RIA	Assay-dependent
	WB	Assay-dependent
Application Note	* The dilutions indicate recommended starting dilutions and the optimal dilutions or concentrations should be determined by the scientist.	

#### **Properties**

Form	Liquid	
Purification	Purification with Protein G.	
Buffer	PBS and 0.09% Sodium azide	
Preservative	0.09% Sodium azide	
Concentration	1 mg/ml	
Storage instruction	For continuous use, store undiluted antibody at 2-8°C for up to a week. For long-term storage, aliquot and store at -20°C or below. Storage in frost free freezers is not recommended. Avoid repeated freeze/thaw cycles. Suggest spin the vial prior to opening. The antibody solution should be gently mixed before use.	

#### Bioinformation

Gene Symbol F8

Gene Full Name coagulation factor VIII, procoagulant component

Background This gene encodes coagulation factor VIII, which participates in the intrinsic pathway of blood

coagulation; factor VIII is a cofactor for factor IXa which, in the presence of Ca+2 and phospholipids, converts factor X to the activated form Xa. This gene produces two alternatively spliced transcripts. Transcript variant 1 encodes a large glycoprotein, isoform a, which circulates in plasma and associates with von Willebrand factor in a noncovalent complex. This protein undergoes multiple cleavage events. Transcript variant 2 encodes a putative small protein, isoform b, which consists primarily of the phospholipid binding domain of factor VIIIc. This binding domain is essential for coagulant activity. Defects in this gene results in hemophilia A, a common recessive X-linked coagulation disorder.

[provided by RefSeq, Jul 2008]

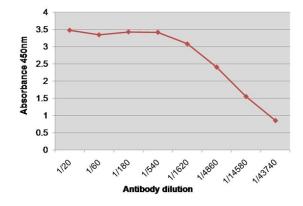
Function Factor VIII, along with calcium and phospholipid, acts as a cofactor for factor IXa when it converts factor

X to the activated form, factor Xa. [UniProt]

Calculated Mw 267 kDa

PTM Sulfation on Tyr-1699 is essential for binding vWF.

#### **Images**



#### ARG22573 anti-Factor VIII antibody [RFF-VIIIC/8] ELISA image

ELISA: Titration curve of ARG22573 anti-Factor VIII antibody [RFF-VIIIC/8]. Antigen: Recombinant Factor VIII.