

## **German Research Products - GRP GmbH**

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## **Product Datasheet**

## F8 - Coagulation factor VIII **GRP13213**

Species/Host Chicken

Reactivity Human

**Predicted Reactivity** Human

purified full length native human Coagulation Factor VIII-VWF-F **Immunogen** 

VIII:c complex

Form/Appearance Liquid in 0.9% NaCl, 0.1% sodium azide

Storage Store at 4°C; make aliquots to avoid working with a stock. Please,

remember to spin tubes briefly prior to opening them to avoid any losses that might occur from liquid material adhering to the

cap or sides of the tubes.

Note For research use only.

Clonality Polyclonal

**Purity** Total IgY fraction

MW 267 kDa

**Dilution Range** 1:200 (IHC)

**Application Notes** Additional Information: Immunohistochemistry: antibody was tested on

bonin-fixed rat blood vessel. IgG concentration is 22.2 µg/µl Background: The factor VIII complex, with a molecular weight in excess of 1 million, has 2 components: (1) factor VIII P00451 (molecular weight of 267 kDa) called factor VIII C, when measured by procoagulant activity or factor VIII Ag, when measured immunologically; (2) factor VIII R P04275 (the von Willebrand factor or vWF) has a molecular weight of 225 kDa. Factor VIII, along with calcium and

phospholipids, acts as a cofactor for factor IXa when it converts factor X to the activated form, factor Xa. Domain F5/8 type C 2 Is responsible for phospholipid binding and essential for factor VIII activity. Defects in F8 are the cause of hemophilia A (HEMA). HEMA is a common recessive X-linked coagulation

disorder.