

# **Product datasheet for AP06114PU-M**

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OriGene Technologies, Inc.

### **Factor VIII (F8) Rabbit Polyclonal Antibody**

**Product data:** 

**Product Type:** Primary Antibodies

**Applications:** IHC, WB

Recommended Dilution: Western blot: 1/500-1/1000.

Immunohistochemistry on paraffin sections: 1/50-1/200.

Reactivity: Human, Mouse

**Host:** Rabbit

Clonality: Polyclonal

**Immunogen:** Synthetic peptide, corresponding to amino acids 2161-2210 of Human F8.

Specificity: This antibody detects endogenous levels of Coagulation factor VIII (F8) protein (region

surronding Ser2194).

**Formulation:** Phosphate buffered saline (PBS), pH 7.2.

State: Aff - Purified

State: Liquid purified lg fraction Preservative: 0.05% Sodium azide

**Concentration:** 1.0 mg/ml

**Purification:** Affinity chromatography using epitope-specific immunogen (> 95% pure by SDS-PAGE).

Conjugation: Unconjugated

Storage: Store undiluted at 2-8°C for one month or (in aliquots) at -20°C for longer.

Avoid repeated freezing and thawing.

**Stability:** Shelf life: one year from despatch.

**Predicted Protein Size:** ~267 kDa

Gene Name: coagulation factor VIII

Database Link: Entrez Gene 2157 Human

P00451





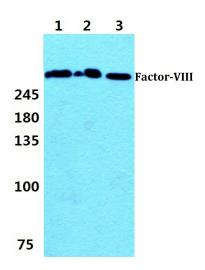
#### Background:

Hemostasis following tissue injury involves the deployment of essential plasma procoagulants (prothrombin, and Factors X, IX, V, and VIII), which are involved in a blood coagulation cascade that leads to the formation of insoluble fibrin clots and the promotion of platelet aggregation. Coagulation Factor VII (serum prothrombin conversion accelerator, proconvertin, F7, Factor VII) is a 406 amino acid, vitamin K-dependent, single chain serine protease that is synthesized in the liver and circulates as an inactive precursor. Factor IX A, Factor X A, Factor XII A, or thrombin mediated proteolytic cleavage of Factor VII at Arg152-Ile153 generates Factor VII A, an active serine protease composed of a catalytic heavy chain disulfide linked to a light chain, containing 2 EGF-like domains. Mutations at the F7 locus that lead to Factor VII deficiencies are generally asymptomatic or phenotypically uncharacterized, with hemorrhagic diathesis occurring at extremely low levels.

Synonyms:

Procoagulant component, Antihemophilic factor, F8C, AHF

## **Product images:**



Western blot (WB) analysis of F8 antibody at 1/500 dilution Lane 1:HepG2 whole cell lysate Lane 2:Mouse liver tissue lysate Lane 3:Rat liver tissue lysate