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Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

F8 (Human) Recombinant Protein (P01)

Catalog Number: H00002157-P01

Regulation Status: For research use only (RUO)

Product Description: Human F8 full-length ORF (NP_063916.1, 1 a.a. - 216 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence:

```
MRIQDPGKVFVFGNVDSGSIKHNFNPPIIARYIRLHPTH  
YSIRSTLRMELMGCDLNSCSMPLGMESKAISDAQITAS  
SYFTNMFATWSPSKARLHLQGRSNAWRPQVNNPKE  
WLQVDFQKTMKVTGVTQGVKSLTSMYVKEFLISS  
QDGHQWTLFFQNGKVKVFQGNQDSFTPVVNSLDPPL  
LTRYLRHPQSWVHQIALRMEVLGCEAQDLY
```

Host: Wheat Germ (in vitro)

Theoretical MW (kDa): 51

Applications: AP, Array, ELISA, WB-Re
(See our web site product page for detailed applications information)

Protocols: See our web site at
<http://www.abnova.com/support/protocols.asp> or product page for detailed protocols

Preparation Method: [in vitro wheat germ expression system](#)

Purification: Glutathione Sepharose 4 Fast Flow

Storage Buffer: 50 mM Tris-HCl, 10 mM reduced Glutathione, pH=8.0 in the elution buffer.

Storage Instruction: Store at -80°C. Aliquot to avoid repeated freezing and thawing.

Entrez GeneID: 2157

Gene Symbol: F8

Gene Alias: AHF, DXS1253E, F8B, F8C, FVIII, HEMA

Gene Summary: 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 Ca²⁺ 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]