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See the following pages for more information!



Lieferung & Zahlungsart

siehe unsere [Liefer- und Versandbedingungen](#)

Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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Datasheet

CFP (Human) Recombinant Protein (P01)

Catalog Number: H00005199-P01

Regulation Status: For research use only (RUO)

Product Description: Human CFP full-length ORF (AAH15756.1, 1 a.a. - 469 a.a.) recombinant protein with GST-tag at N-terminal.

Sequence:

MITEGAQAPRLLLPLLLLLLTPATGSDPVLCFTQYEES
SGKCKGLLGGVSVEDCCLNTAFAYQKRSGGLCQPC
RSPRWLWSTWAPCSVTCSEGSQLRYYRRCVGNWNGQ
CSGKVAPGTLEWQLQACEDQQCCPEMGGWSGWGP
WEPCSVTCSKGTRRRRACNHPAPKCGGHCPGQAQ
ESEACDTQQVCPHGWATWGPWTPCSASCHGGPH
EPKETRSRKCSAPEPSQKPPGKPCPGLAYEQRRCTG
LPPCPVAGGWGPWGPVSPCPVTCGLGQTMEQRTC
HPVPQHGGPFCAGDATRTHICNTAVPCPVDGEWDSW
GEWSPCIRRNMKSSISCQEIPGQQSRGRTCRGRKFDG
HRCAGQQDIRHCYSIQHCPLKGSWSEWSTWGLCM
PPCGPNPTRARQLCTPLLPKYPTVSMVEGQGEKN
VTFWGRPLPRCEELQGQKLVVEEKRPCLHVPACKDP
EEEEEL

Host: Wheat Germ (in vitro)

Theoretical MW (kDa): 77.33

Interspecies Antigen Sequence: Mouse (77); Rat (79)

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: 5199

Gene Symbol: CFP

Gene Alias: BFD, PFC, PFD, PROPERDIN

Gene Summary: This gene encodes a plasma glycoprotein that positively regulates the alternative complement pathway of the innate immune system. This protein binds to many microbial surfaces and apoptotic cells and stabilizes the C3- and C5-convertase enzyme complexes in a feedback loop that ultimately leads to formation of the membrane attack complex and lysis of the target cell. Mutations in this gene result in two forms of properdin deficiency, which results in high susceptibility to meningococcal infections. Multiple alternatively spliced variants, encoding the same protein, have been identified]