

## Product datasheet for TP720638XL

## OriGene Technologies, Inc.

9620 Medical Center Drive, Ste 200 Rockville, MD 20850, US Phone: +1-888-267-4436 https://www.origene.com techsupport@origene.com EU: info-de@origene.com CN: techsupport@origene.cn

## Complement factor B (CFB) (NM\_001710) Human Recombinant Protein

**Product data:** 

**Product Type:** Recombinant Proteins

**Description:** Purified recombinant protein of Human complement factor B (CFB)

Species: Human Expression Host: HEK293

**Expression cDNA Clone** 

Thr26-Leu764

or AA Sequence:

Tag: C-His

Predicted MW: 84.07 kDa

**Purity:** >95% as determined by SDS-PAGE and Coomassie blue staining

**Buffer:** Provided lyophilized from a 0.2 μm filtered solution of 20 mM Tris-HCl, 150 mM NaCl

**Endotoxin:** Endotoxin level is < 0.1 ng/µg of protein (< 1 EU/µg)

**Storage:** Store at -80°C.

Stability: Stable for at least 6 months from date of receipt under proper storage and handling

conditions.

**RefSeq:** NP 001701

Locus ID: 629

UniProt ID: <u>P00751</u>, <u>A0A1U9X7H8</u>

RefSeq Size: 2646
Cytogenetics: 6p21.33
RefSeq ORF: 2292

Synonyms: AHUS4; ARMD14; BF; BFD; CFAB; CFBD; FB; FBI12; GBG; H2-Bf; PBF2





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Summary: This gene encodes complement factor B, a component of the alternative pathway of

complement activation. Factor B circulates in the blood as a single chain polypeptide. Upon activation of the alternative pathway, it is cleaved by complement factor D yielding the noncatalytic chain Ba and the catalytic subunit Bb. The active subunit Bb is a serine protease which associates with C3b to form the alternative pathway C3 convertase. Bb is involved in the proliferation of preactivated B lymphocytes, while Ba inhibits their proliferation. This gene localizes to the major histocompatibility complex (MHC) class III region on chromosome 6.

This cluster includes several genes involved in regulation of the immune reaction. Polymorphisms in this gene are associated with a reduced risk of age-related macular degeneration. The polyadenylation site of this gene is 421 bp from the 5' end of the gene for

complement component 2. [provided by RefSeq, Jul 2008]

**Protein Families:** Druggable Genome, Protease, Secreted Protein

**Protein Pathways:** Complement and coagulation cascades