

Factor I rabbit pAb

Cat No.: ES5825

For research use only

Overview

Product Name Factor I rabbit pAb

Host species Rabbit
Applications WB;ELISA

Species Cross-Reactivity Human; Rat; Mouse;

Recommended dilutions IHC: 1/500 - 1/2000. ELISA: 1/10000. Not yet tested

in other applications.

Immunogen The antiserum was produced against synthesized

peptide derived from human CFI. AA range:441-490

Specificity Factor I Polyclonal Antibody detects endogenous

levels of Factor I protein.

Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and

0.02% sodium azide.

Storage Store at -20° C. Avoid repeated freeze-thaw cycles.

Protein Name Complement factor I

Gene Name CFI

Cellular localization Secreted, extracellular space. Secreted.

Purification The antibody was affinity-purified from rabbit

antiserum by affinity-chromatography using

epitope-specific immunogen.

ClonalityPolyclonalConcentration1 mg/ml

Observed band Full lenth:66kD, heavy chain: 50-58kD

Human Gene ID 3426 Human Swiss-Prot Number P05156

Background

Alternative Names CFI; IF; Complement factor I; C3B/C4B inactivator

This gene encodes a serine proteinase that is essential for regulating the complement cascade. The encoded preproprotein is cleaved to produce both heavy and light chains, which are linked by

disulfide bonds to form a heterodimeric

glycoprotein. This heterodimer can cleave and inactivate the complement components C4b and C3b, and it prevents the assembly of the C3 and C5



+86-27-59760950 ELKbio@ELKbiotech.com

www.elkbiotech.com



convertase enzymes. Defects in this gene cause complement factor I deficiency, an autosomal recessive disease associated with a susceptibility to pyogenic infections. Mutations in this gene have been associated with a predisposition to atypical hemolytic uremic syndrome, a disease characterized by acute renal failure, microangiopathic hemolytic anemia and thrombocytopenia. Primary glomerulonephritis with immune deposits and age-related macular degeneration are other conditions associated with mutations of this gene. [provided by Ref



+86-27-59760950