

Factor I Polyclonal Antibody

Catalog No :	YT1652
Reactivity :	Human;Rat;Mouse;
Applications :	WB;ELISA
Target :	Factor I
Fields :	>>Complement and coagulation cascades;>>Staphylococcus aureus infection
Gene Name :	CFI
Protein Name :	Complement factor I
Human Gene Id :	3426
Human Swiss Prot No :	P05156
Mouse Swiss Prot No :	Q61129
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.
Source :	Polyclonal, Rabbit,IgG
Dilution :	WB 1:500-2000. ELISA: 1:10000. Not yet tested in other applications.
Purification :	The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.
Concentration :	1 mg/ml
Storage Stability :	-15°C to -25°C/1 year(Do not lower than -25°C)

Observed Band : Full length:66kD, heavy chain: 50-58kD

Cell Pathway : Complement and coagulation cascades;

Background : This gene encodes a serine proteinase that is essential for regulating the complement cascade. The encoded preprotein 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 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

Function : catalytic activity:Inactivates complement subcomponents C3b, iC3b and C4b by proteolytic cleavage.,disease:Defects in CFI are the cause of complement factor I deficiency (CFI deficiency) [MIM:610984]. CFI deficiency is an autosomal recessive condition associated with a propensity to pyogenic infections.,disease:Defects in CFI are the cause of component I deficiency (CFI deficiency) [MIM:217030]. CFI deficiency is an autosomal recessive condition associated with a propensity to pyogenic infections.,disease:Defects in CFI may be associated with or predispose to hemolytic uraemic syndrome (HUS) [MIM:235400]. HUS, the most frequent cause of acute renal failure in childhood, is characterized by the association of acute renal failure, microangiopathic hemolytic anemia, and thrombocytopenia. The majority of HUS cases occur after an episode of infectious diarrhea, and are associated with E.coli

Subcellular Location : Secreted, extracellular space. Secreted .

Expression : Expressed in the liver by hepatocytes (PubMed:6327681). Also present in other cells such as monocytes, fibroblasts or keratinocytes (PubMed:6444659, PubMed:17320177).

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