



C1s Monoclonal Antibody

Catalog No	YP-mAb-13874
Isotype	IgG
Reactivity	Human;Mouse;Rat
Applications	WB
Gene Name	C1S
Immunogen	The antiserum was produced against synthesized peptide derived from human C1S. AA range:541-590
Specificity	C1s Monoclonal Antibody detects endogenous levels of C1s protein.
Formulation	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.
Source	Monoclonal, Mouse,IgG
Purification	The antibody was affinity-purified from mouse antiserum by affinity-chromatography using epitope-specific immunogen.
Dilution	WB 1:500-1:2000
Concentration	1 mg/ml
Purity	≥90%
Storage Stability	-20°C/1 year
Synonyms	C1 esterase;C1S;Complement C1s subcomponent;Complement component 1 subcomponent s
Observed Band	77kD
Calculated Molecular Weight	77kD
Cell Pathway	extracellular region,extracellular exosome,blood microparticle,
Tissue Specificity	Liver,Peripheral blood leukocyte,Plasma,PNS,
Function	catalytic activity:Cleavage of Arg- -Ala bond in complement component C4 to form C4a and C4b, and Lys(or Arg)- -Lys bond in complement component C2 to form C2a and C2b: the 'classical' pathway C3 convertase.,disease:Defects in C1S are the cause of selective C1s deficiency [MIM:120580]; that is associated with early onset multiple autoimmune diseases.,enzyme regulation:Inhibited by SERPING1.,function:C1s B chain is a serine protease that combines with C1q and C1s to form C1, the first component of the classical pathway of the complement system. C1r activates C1s so that it can, in turn, activate C2 and C4.,online information:C1S mutation db,PTM:The iron and 2-oxoglutarate dependent 3-hydroxylation of aspartate and asparagine is (R) stereospecific within EGF domains.,similarity:Belongs to the peptidase S1 family.,



similarity:Contains 1 EGF-like domain.,similarity:Contains 1 peptidase S1 dom

Background

This gene encodes a serine protease, which is a major constituent of the human complement subcomponent C1. C1s associates with two other complement components C1r and C1q in order to yield the first component of the serum complement system. Defects in this gene are the cause of selective C1s deficiency. [provided by RefSeq, Mar 2009],

matters needing attention

Avoid repeated freezing and thawing!

Usage suggestions

This product can be used in immunological reaction related experiments. For more information, please consult technical personnel.

Products Images

