

**Product Name: Cleaved-C1s HC (R437) Rabbit Polyclonal Antibody**  
**Catalog #: APRab08951**

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## Summary

<b>Production Name</b>	Cleaved-C1s HC (R437) Rabbit Polyclonal Antibody
<b>Description</b>	Rabbit Polyclonal Antibody
<b>Host</b>	Rabbit
<b>Application</b>	WB,ELISA
<b>Reactivity</b>	Human,Rat,Mouse

## Performance

<b>Conjugation</b>	Unconjugated
<b>Modification</b>	Unmodified
<b>Isotype</b>	IgG
<b>Clonality</b>	Polyclonal
<b>Form</b>	Liquid
<b>Storage</b>	Store at 4°C short term. Aliquot and store at -20°C long term. Avoid freeze/thaw cycles.
<b>Buffer</b>	Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% New type preservative N.
<b>Purification</b>	Affinity purification

## Immunogen

<b>Gene Name</b>	C1S
<b>Alternative Names</b>	C1S; Complement C1s subcomponent; C1 esterase; Complement component 1 subcomponent s
<b>Gene ID</b>	716.0
<b>SwissProt ID</b>	P09871.The antiserum was produced against synthesized peptide derived from human C1S. AA range:388-437

## Application

<b>Dilution Ratio</b>	WB 1:500 - 1:2000. ELISA: 1:20000
<b>Molecular Weight</b>	47+76kD

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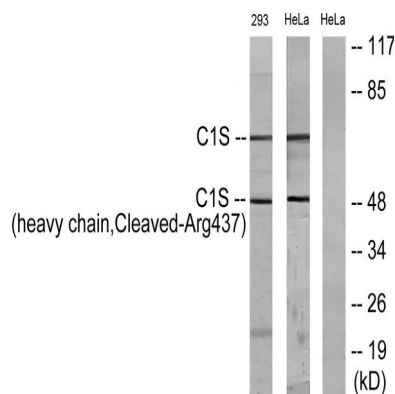
## Background

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],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 C1r 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 domain.,similarity: Contains 2 CUB domains.,similarity: Contains 2 Sushi (CCP/SCR) domains.,subunit: C1 is a calcium-dependent trimolecular complex of C1q, C1r and C1s in the molar ratio of 1:2:2. Activated C1s is a disulfide-linked heterodimer of a heavy chain and a light chain.,

## Research Area

Complement and coagulation cascades; Systemic lupus erythematosus;

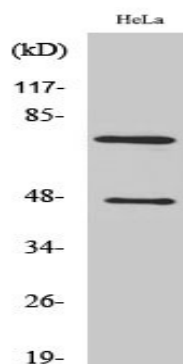
## Image Data



Western blot analysis of lysates from 293 and HeLa cells, treated with etoposide 25uM 1h, using C1S (heavy chain, Cleaved-Arg437) Antibody. The lane on the right is blocked with the synthesized peptide.

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Western Blot analysis of various cells using Cleaved-C1s HC (R437) Polyclonal Antibody

### **Note**

For research use only.