

**(** Tel: 400-999-8863 ■ Email:Upingbio.163.com



# C8 α Polyclonal Antibody

of the membrane attack complex. C8 binds to the C5b-7 complex, forming the C5b-8 complex. C5-b8 binds C9 and acts as a catalyst in the polymerization of C9.,similarity:Belongs to the complement C6/C7/C8/C9 family.,similarity:Contair 1 EGF-like domain.,similarity:Contains 1 LDL-receptor class A domain.,similarity:Contains 1 MACPF domain.,similarity:Contains 2 TSP type-1	Catalog No	YP-Ab-10777
Applications WB;ELISA  Gene Name C8 α  Protein Name C8 α  Immunogen Synthesized peptide derived from human C8 α. at AA range: 201-250  Specificity C8 α Polyclonal Antibody detects endogenous levels of C8 α  Formulation Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.  Source Polyclonal, Rabbit,IgG  Purification The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.  Dilution WB 1:500-2000, ELISA 1:10000-20000  Concentration 1 mg/ml  Purity ≥90%  Storage Stability -20°C/1 year  Synonyms Complement component C8 alpha chain (Complement component 8 subunit alpha)  Observed Band 60kD  Cell Pathway Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.  Tissue Specificity Blood, Liver, Plasma,  disease: Defects in C8A are a cause of complement C8 deficiency type I [MIM:120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningtidics. function: 8 is a constitue of the membrane attack complex. C8 binds to a catalyst in the C5B-8 complex. C5-9 binds C9 and acts as a catalyst in polymerization of C9 s. similarity. Belongs to the complement C8/C7/C8/C9 family, similarity. Contains 1 LD1-peoppro class. a constitue of the membrane attack complex. 1 MACPF domain, similarity. Contains 2 MACPF domain, similarity. Contains 1 LD1-peoppro class. a constitue of the membrane and gamma chains are disulfide bonded.,  C8 is a component of the complement system and contains three polypeptides alpha, beta and gamma. This gene encodes the alpha subunit of C8. C8	Isotype	IgG
Gene Name         C8 α           Protein Name         C8 α           Immunogen         Synthesized peptide derived from human C8 α. at AA range: 201-250           Specificity         C8 α Polyclonal Antibody detects endogenous levels of C8 α           Formulation         Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.           Source         Polyclonal, Rabbit, IgG           Purification         The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.           Dilution         WB 1:500-2000, ELISA 1:10000-20000           Concentration         1 mg/ml           Purity         ≥90%           Storage Stability         -20°C/1 year           Synonyms         Complement component C8 alpha chain (Complement component 8 subunit alpha)           Observed Band         60kD           Cell Pathway         Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.           Tissue Specificity         Blood, Liver, Plasma,           Function         disease: Defects in C8A are a cause of complement C8 deficiency type I [MiM: 120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitidis, function: C8 is a constitue of the membrane attack complex. C8 binds C9 and acts as a catalyst in the polymerization of C9, similarity. Contains 1 tD1-receptor class A domai	Reactivity	Human;Rat;Mouse;
Protein Name         C8 α           Immunogen         Synthesized peptide derived from human C8 α. at AA range: 201-250           Specificity         C8 α Polyclonal Antibody detects endogenous levels of C8 α           Formulation         Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.           Source         Polyclonal, Rabbit, IgG           Purification         The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.           Dilution         WB 1:500-2000, ELISA 1:10000-20000           Concentration         1 mg/ml           Purity         ≥90%           Storage Stability         -20°C/1 year           Synonyms         Complement component C8 alpha chain (Complement component 8 subunit alpha)           Observed Band         60kD           Cell Pathway         Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.           Tissue Specificity         Blood, Liver, Plasma,           Function         disease: Defects in C8A are a cause of complement C8 deficiency type I [MiM: 120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitidis, function: C8 is a constitue of the membrane attack complex. C8 binds to the C5b-7 complex, forming the C5b-8 complex. C9-50 binds complex as a catalyst in the polymerization of C9, similarity. Belongs to the complement C6/C7/C8/C9 family, sim	Applications	WB;ELISA
Immunogen         Synthesized peptide derived from human C8 α. at AA range: 201-250           Specificity         C8 α Polyclonal Antibody detects endogenous levels of C8 α           Formulation         Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.           Source         Polyclonal, Rabbit, IgG           Purification         The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.           Dilution         WB 1:500-2000, ELISA 1:10000-20000           Concentration         1 mg/ml           Purity         ≥90%           Storage Stability         -20°C/1 year           Synonyms         Complement component C8 alpha chain (Complement component 8 subunit alpha)           Observed Band         60kD           Cell Pathway         Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.           Tissue Specificity         Blood, Liver, Plasma,           Function         disease: Defects in C8A are a cause of complement C8 deficiency type I [MIM: 120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitids, function: C8 is a constitue of the membrane attack complex. C8 binds to the C5b-7 complex, forming the c5b-8 complex. C5-0 so and acts as a catalyst in the polymerization of C9, similarity. Belongs to the complement C6/C7/C8/C9 family, similarity. Contains 1 LDL-receptor class A domain, similarity. Contains 1 MA	Gene Name	C8A
Specificity         C8 α Polyclonal Antibody detects endogenous levels of C8 α           Formulation         Liquid in PBS containing 50% glycerol, 0.5% BSA and 0.02% sodium azide.           Source         Polyclonal, Rabbit, IgG           Purification         The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.           Dilution         WB 1:500-2000, ELISA 1:10000-20000           Concentration         1 mg/ml           Purity         ≥90%           Storage Stability         -20°C/1 year           Synonyms         Complement component C8 alpha chain (Complement component 8 subunit alpha)           Observed Band         60kD           Cell Pathway         Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.           Tissue Specificity         Blood, Liver, Plasma,           Function         disease: Defects in C8A are a cause of complement C8 deficiency type I MIM: 120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningities, function: C8 is a constitute of the membrane attack complex. C8 binds to the C5b-7 complex, forming the C5b-8 complex. C5-b8 binds C9 and acts as a catalyst in the polymerization of C9. s.similarity:Belongs to the complement C6/C7/C8/C9 family, similarity:Contain 1 EGF-like domain., similarity:Contains 1 MACPF domain., similarity:Contains 2 TSP type-1 domain., subunit:C8 is composed of three chains: alpha, beta and gamma. This gene encodes	Protein Name	C8 α
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Purification The antibody was affinity-purified from rabbit antiserum by affinity-chromatography using epitope-specific immunogen.  Dilution WB 1:500-2000, ELISA 1:10000-20000 Concentration 1 mg/ml  Purity ≥90% Storage Stability -20°C/1 year  Synonyms Complement component C8 alpha chain (Complement component 8 subunit alpha) Observed Band 60kD  Cell Pathway Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.  Tissue Specificity Blood,Liver,Plasma,  disease:Defects in C8A are a cause of complement C8 deficiency type I MIM:120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitidis., function:C8 is a constitue of the membrane attack complex. C6 binds to the C5b-7 complex, forming the C5b-8 complex. C5-b8 binds C9 and acts as a catalyst in the polymerization of C9. similarity Eelongs to the complement C6/C7/C8/C9 family, similarity:Contain 1 EGF-like domain., similarity:Contains 1 LDL-receptor class A domain., similarity:Contains 1 MACPF domain., similarity:Contains 2 TSP type-1 domains, subunit:C8 is composed of three chains: alpha, beta and gamma. The alpha and gamma chains are disulfide bonded.,  C8 is a component of the complement system and contains three polypeptides alpha, beta and gamma. This gene encodes the alpha subunit of C8. C8	Specificity	C8 $\alpha$ Polyclonal Antibody detects endogenous levels of C8 $\alpha$
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Concentration       1 mg/ml         Purity       ≥90%         Storage Stability       -20°C/1 year         Synonyms       Complement component C8 alpha chain (Complement component 8 subunit alpha)         Observed Band       60kD         Cell Pathway       Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.         Tissue Specificity       Blood,Liver,Plasma,         Function       disease:Defects in C8A are a cause of complement C8 deficiency type I [MIM:120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitidis, function:C8 is a constitue of the membrane attack complex. C8 binds to the C5b-7 complex, forming the C5b-8 complex. C5-b8 binds C9 and acts as a catalyst in the polymerization of C9. similarity:Belongs to the complement C6/C7/C8/C9 family, similarity:Contains 1 LDL-receptor class A domain., similarity:Contains 1 MACPF domain., similarity:Contains 2 TSP type-1 domains., subunit:C8 is composed of three chains: alpha, beta and gamma. The alpha and gamma chains are disulfide bonded.         Background       C8 is a component of the complement system and contains three polypeptides alpha, beta and gamma. This gene encodes the alpha subunit of C8. C8	Purification	
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alpha, beta and gamma. This gene encodes the alpha subunit of C8. C8	Observed Band Cell Pathway	alpha)  60kD  Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.
assembles on bacterial membranes to form a pore, permitting disruption of	Observed Band Cell Pathway Tissue Specificity	alpha)  60kD  Secreted. Cell membrane; Multi-pass membrane protein. Secreted as soluble protein. Inserts into the cell membrane of target cells.  Blood,Liver,Plasma,  disease:Defects in C8A are a cause of complement C8 deficiency type I [MIM:120950]. Patients with deficiency of C8 suffer from recurrent bacterial infections, predominantly from Neisseria meningitidis.,function:C8 is a constituent of the membrane attack complex. C8 binds to the C5b-7 complex, forming the C5b-8 complex. C5-b8 binds C9 and acts as a catalyst in the polymerization of C9.,similarity:Belongs to the complement C6/C7/C8/C9 family.,similarity:Contains 1 EGF-like domain.,similarity:Contains 1 LDL-receptor class A domain.,similarity:Contains 1 MACPF domain.,similarity:Contains 2 TSP type-1 domains.,subunit:C8 is composed of three chains: alpha, beta and gamma. The



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bacterial membrane organization. Mutations in this gene cause complement C8 alpha-gamma deficiency. [provided by RefSeq, Nov 2008],

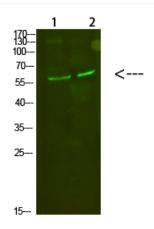
## matters needing attention

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**



Western Blot analysis of 1,mouse-kidney 2,mouse-heart cells using primary antibody diluted at 1:500(4°C overnight). Secondary antibody:Goat Anti-rabbit IgG IRDye 800( diluted at 1:5000, 25°C, 1 hour)