abcam

Product datasheet

Anti-C9 antibody ab118902

4 Images

Overview

Product name Anti-C9 antibody

Description Rabbit polyclonal to C9

Host species Rabbit

Tested applications Suitable for: WB, IHC-P, ICC/IF

Species reactivity Reacts with: Human

Immunogen Recombinant Human C9 protein.

Positive control Human Small Intestine, Vessels Human Placenta methanol-fixed HepG2 Raji cell lysate

General notes

The Life Science industry has been in the grips of a reproducibility crisis for a number of years.

Abcam is leading the way in addressing this with our range of recombinant monoclonal antibodies and knockout edited cell lines for gold-standard validation. Please check that this product meets

your needs before purchasing.

If you have any questions, special requirements or concerns, please send us an inquiry and/or contact our Support team ahead of purchase. Recommended alternatives for this product can be

found below, along with publications, customer reviews and Q&As

Properties

Form Liquid

Storage instructions Shipped at 4°C. Upon delivery aliquot and store at -20°C. Avoid freeze / thaw cycles.

Storage buffer pH: 7.00

Preservative: 0.01% Thimerosal (merthiolate)

Constituents: 10% Glycerol (glycerin, glycerine), Tris glycine

Purity Immunogen affinity purified

Clonality Polyclonal

Isotype IgG

Applications

The Abpromise guarantee Our Abpromise guarantee covers the use of ab118902 in the following tested applications.

The application notes include recommended starting dilutions; optimal dilutions/concentrations should be determined by the end user.

Application	Abreviews	Notes
WB		1/500 - 1/3000. Predicted molecular weight: 62 kDa.
IHC-P		Use a concentration of 10 µg/ml.
ICC/IF		1/100 - 1/200.

Target

Function Constituent of the membrane attack complex (MAC) that plays a key role in the innate and

adaptive immune response by forming pores in the plasma membrane of target cells. C9 is the

pore-forming subunit of the MAC.

Tissue specificity Plasma.

Involvement in disease Defects in C9 are a cause of complement component 9 deficiency (C9D) [MIM:613825]. A rare

defect of the complement classical pathway associated with susceptibility to severe recurrent

infections, predominantly by Neisseria gonorrhoeae or Neisseria meningitidis.

Sequence similarities Belongs to the complement C6/C7/C8/C9 family.

Contains 1 EGF-like domain.

Contains 1 LDL-receptor class A domain.

Contains 1 MACPF domain.
Contains 1 TSP type-1 domain.

Post-translational Thrombin cleaves factor C9 to produce C9a and C9b.

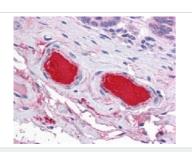
modifications Phosphorylation sites are present in the extracelllular medium.

Cellular localization Secreted. Cell membrane. Secreted as soluble monomer. Oligomerizes at target membranes,

forming a pre-pore. A conformation change then leads to the formation of a 100 Angstrom

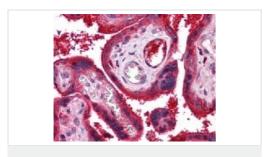
diameter pore.

Images



Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-C9 antibody (ab118902)

Immunohistochemistry analysis (Formalin-Fixed, Paraffin-Embedded) of Human Small Intestine, Vessels stained with ab118902 at 10 μ g/ml



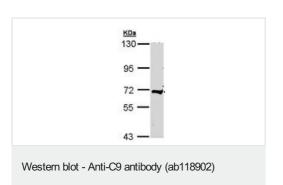
Immunohistochemistry (Formalin/PFA-fixed paraffinembedded sections) - Anti-C9 antibody (ab118902)

Immunohistochemistry analysis (Formalin-Fixed, Paraffin-Embedded) of Human Placenta stained with ab118902 at 10 μ g/ml



C9 antibody (ab118902)

Immunofluorescence analysis of methanol-fixed HepG2, using ab118902 at 1:200 dilution.



Anti-C9 antibody (ab118902) at 1/500 dilution + Raji cell lysate at 30 μg

Predicted band size: 62 kDa

Please note: All products are "FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES"

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