abcam

Product datasheet

Human Complement C9 ELISA Kit ab137972

3 References 1 Image

Overview

Product name Human Complement C9 ELISA Kit

Detection methodColorimetric

Precision

Sample	n	Mean	SD	CV%	
Overall				3.2%	

Inter-assay

Intra-assav

Sample	n	Mean	SD	CV%
Overall				8.4%

Sample type Cell culture supernatant, Saliva, Milk, Urine, Serum, Plasma, Tissue, Cell Lysate, Cerebral Spinal

Fluid

Assay type Sandwich (quantitative)

Sensitivity = 0.059 ng/ml

Range 0.234 ng/ml - 15 ng/ml

Recovery = 101 %

Assay time 4h 0m

Assay duration Multiple steps standard assay

Species reactivity Reacts with: Human

Product overview Abcam's Complement C9 Human in vitro ELISA (Enzyme-Linked Immunosorbent Assay) kit is

designed for the quantitative measurement of C9 in Human plasma, serum, saliva, milk, urine,

cerebrospinal fluid and cell culture supernatants.

A Complement C9 specific antibody has been precoated onto 96-well plates and blocked. Standards or test samples are added to the wells and subsequently a Complement C9 specific biotinylated detection antibody is added and then followed by washing with wash buffer. Streptavidin-Peroxidase Conjugate is added and unbound conjugates are washed away with wash buffer. TMB is then used to visualize Streptavidin-Peroxidase enzymatic reaction. TMB is catalyzed by Streptavidin-Peroxidase to produce a blue color product that changes into yellow after adding acidic stop solution. The density of yellow coloration is directly proportional to the

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amount of Complement C9 captured in plate.

The entire kit may be stored at -20°C for long term storage before reconstitution - Avoid repeated freeze-thaw cycles.

Platform Microplate

Properties

Storage instructions Store at -20°C. Please refer to protocols.

Components	1 x 96 tests
100X Streptavidin-Peroxidase Conjugate	1 x 80µl
10X Diluent M Concentrate	1 x 30ml
20X Wash Buffer Concentrate	2 x 30ml
50X Biotinylated Human Complement C9 Antibody	1 x 120µl
Chromogen Substrate	1 x 7ml
Complement C9 Microplate (12 x 8 well strips)	1 unit
Complement C9 Standard	2 vials
Sealing Tapes	3 units
Stop Solution	1 x 11ml

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 modifications

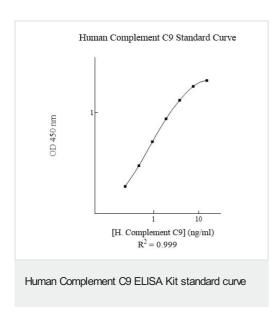
Thrombin cleaves factor C9 to produce C9a and C9b.

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.



Representative Standard Curve using ab137972

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