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Lieferung & Zahlungsart

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Zuschläge

- Mindermengenzuschlag
- Trockeneiszuschlag
- Gefahrgutzuschlag
- Expressversand

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CD96 (Human) Matched Antibody Pair

Catalog # : H00010225-AP51

規格 : [1 Set]

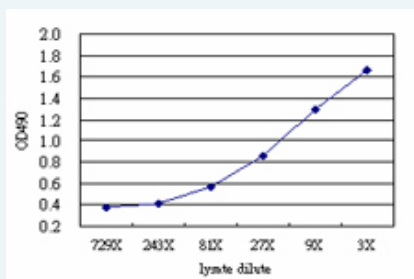
[List All](#)

Specification

Product Description: This antibody pair set comes with matched antibody pair to detect and quantify protein level of human CD96.

Reactivity: Human

Quality Control Testing: Standard curve using CD96 293T overexpression lysate (non-denatured) as an analyte.



Sandwich ELISA detection sensitivity ranging from approximately 81x to 3x dilution of the CD96 293T overexpression lysate (non-denatured).

Supplied Product: Antibody pair set content:
 1. Capture antibody: mouse monoclonal anti-CD96 (100 ug)
 2. Detection antibody: rabbit purified polyclonal anti-CD96 (50 ug)
 *Reagents are sufficient for at least 3-5 x 96 well plates using recommended protocols.

Storage Instruction: Store reagents of the antibody pair set at -20°C or lower. Please aliquot to avoid repeated freeze thaw cycle. Reagents should be returned to -20°C storage immediately after use.

MSDS:  [Download](#)

Applications

ELISA Pair (Transfected lysate)

 [Protocol Download](#)

Gene Information

Entrez GeneID: [10225](#)

Gene Name: CD96

Gene Alias: DKFZp667E2122, MGC22596, TACTILE

Gene Description: CD96 molecule

Omim ID: [211750](#), [605039](#), [606037](#)

Gene Ontology: [Hyperlink](#)

Application Image

ELISA Pair (Transfected lysate)

Gene Summary: The protein encoded by this gene belongs to the immunoglobulin superfamily. It is a type I membrane protein. The protein may play a role in the adhesive interactions of activated T and NK cells during the late phase of the immune response. It may also function in antigen presentation. Alternative splicing occurs at this locus and two transcript variants encoding distinct isoforms have been identified. [provided by RefSeq]

Other Designations: CD96 antigen, T cell activation, increased late expression, T-cell surface protein tactile

Related Disease

Abnormalities, Multiple Craniofacial Abnormalities Syndrome Translocation, Genetic

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