Simian Immunoglobulin G4 (IgG4) ELISA Kit

Catalog No: #EK11559



Package Size: #EK11559-1 48T #EK11559-2 96T

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| Description | |
|--------------------|--|
| Product Name | Simian Immunoglobulin G4 (IgG4) ELISA Kit |
| Brief Description | ELISA Kit |
| Applications | ELISA |
| Species Reactivity | Simian |
| Storage | The stability of ELISA kit is determined by the loss rate of activity. The loss rate of this kit is less than 5% |
| | within the expiration date under appropriate storage condition. |
| | The loss rate was determined by accelerated thermal degradation test. Keep the kit at 37C for 4 and 7 days, |
| | and compare O.D.values of the kit kept at 37C with that of at recommended temperature. (referring from China |
| | Biological Products Standard, which was calculated by the Arrhenius equation. For ELISA kit, 4 days storage |
| | at 37C can be considered as 6 months at 2 - 8C, which means 7 days at 37C equaling 12 months at 2 - 8C). |

Application Details

| Detect Range:1.23-100 μg/mL | |
|--|--|
| Sensitivity:0.48 μg/mL | |
| Sample Type:Serum, Plasma, Other biological fluids | |
| Sample Volume: 1-200 μL | |
| Assay Time:1-4.5h | |
| Detection wavelength:450 nm | |

Product Description

Detection Method:SandwichTest principle:This assay employs a two-site sandwich ELISA to quantitate IgG4 in samples. An antibody specific for IgG4 has been pre-coated onto a microplate. Standards and samples are pipetted into the wells and anyIgG4 present is bound by the immobilized antibody. After removing any unbound substances, a biotin-conjugated antibody specific for IgG4 is added to the wells. After washing, Streptavidin conjugated Horseradish Peroxidase (HRP) is added to the wells. Following a wash to remove any unbound avidin-enzyme reagent, a substrate solution is added to the wells and color develops in proportion to the amount of IgG4 bound in the initial step. The color development is stopped and the intensity of the color is measured. Product Overview: Three closely linked loci (IgG1, IgG2 and IgG3) were thought to be responsible for the Gm specificities. Van Loghem et al. (1970) presented evidence on the linkage relationship of immunoglobulin markers (gamma 1, 2, 3, Am). A fourth IgG locus (gamma-G4) was identifiable in the cluster. Using a gamma-4 probe, Kirsch et al. (1982) assigned the IGH cluster to 14q32 by in situ hybridization. Bech-Hansen et al. (1983) found that an RFLP produced by BamHI is a marker for the heavy chain genes G2 and G4 and a gamma-pseudogene. Considerable linkage disequilibrium was found. Quantitative assessment of the degree of association between C-gamma RFLPs, Gm markers, and switch region RFLPs adjacent to C-mu and C-alpha-1 showed that the gamma-pseudogene is most tightly associated with G2, suggesting that it lies between A1 and G2.

Note: This product is for in vitro research use only