

Product datasheet for AP21480HR-N

OriGene Technologies, Inc.

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Human IgE (Fc specific) Goat Polyclonal Antibody

Product data:

Product Type: Secondary Antibodies

Product Name: Human IgE (Fc specific) Goat Polyclonal Antibody

Applications: ELISA, ID, IF, IHC, IP, WB

Recommended Dilution: Enzyme-Immunocytochemical and Immunohistochemical staining for the detection of IgE at

the cellular and subcellular level by staining of appropriately treated cell and tissue

substrates.

To demonstrate circulating IgE antibodies in serodiagnostic microbiology and autoimmune

diseases.

To identify a specific antigen using an reference antibody of Human origin known to be of the

IgE isotype in the middle layer of the indirect test procedure.

In non-isotopic assay methodology (e.g. ELISA) to measure IgE in Human serum or other body

fluids.

Recommneded Dilutions:

Histochemistry and Cytochemistry: 1/50-1/250.

ELISA and comparable non-precipitating antibody-binding assays: 1/1,000-1/25,000.

Reactivity: Human

Host: Goat

Immunogen: Purified monoclonal IgE isolated from Human serum.

Feund's complete adjuvant is used in the first step of the immunization procedure.

Isotype: IgG

Formulation: PBS, pH 7.2 without preservatives and foreign proteins

Label: HRP

State: Lyophilized hyperimmune IgG fraction

Label: Horseradish Peroxidase Molar radio: Peroxidase/IgG ~ 1.7

Reconstitution Method: Restore by adding 1.0 ml of sterile distilled water

Concentration: 7.5 mg/ml

Purification: Hyperimmune antisera with strong precipitating activity are selected for Fractionation by Salt-

Precipitation and purification of the IgG fraction by DEAE-Chromatography.

Conjugation: HRP





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Storage: Store lyophilized at 2-8°C and reconstituted at 2-8°C for one week or (in aliquots) at -20°C for

longer.

Avoid Repeated thawing and freezing.

Note: Adsorption: Immunoaffinity adsorbed using insolubilized antigens as required to eliminate

antibodies cross-reacting with other components of the immunoglobulin system or reacting with other serum proteins. Special attention is given to the removal of antibodies to common

Ig/Fab. The use of insolubilized adsorption antigens prevents the presence of excess

adsorbent protein or immune complexes in the antiserum.