

Product datasheet for AP31080HR-N

OriGene Technologies, Inc.

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C3 Goat Polyclonal Antibody

Product data:

Product Type: Primary Antibodies

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

Recommended Dilution: Suitable for use in Enzyme-Immunocytochemical and Histochemical staining of Mouse C3c in

appropriately prepared substrates at the cellular and subcellular level. Locally deposited immune complexes in tissue usually contain complement, pointing to activation of the

classical pathway. Complement activation in vivo implies active disease and may contribute to the elicitation of the pathogenesis and he extent of tissue destruction.

In ELISA and Western blotting to identify Mouse C3c in serum or other body fluids.

This immunoconjugate is not pre-diluted. The optimum working dilution of each conjugate should be established by titration before being used. Excess labelled antibody must be avoided because it may cause high unspecific background staining and interfere with the

specific signal.

Recommended Working Dilutions:

Histochemical and Cytochemical Use: 1/100-1/500.

ELISA and comparable non-precipitating antibody-binding assays: 1/1,000-1/5,000.

Reactivity: Mouse
Host: Goat
Isotype: IgG

Clonality: Polyclonal

Immunogen: C3c isolated and purified from pooled normal Mouse serum.

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

C3 Goat Polyclonal Antibody - AP31080HR-N

Specificity:

In Immunoelectrophoresis against fresh Mouse serum, a single precipitin line is obtained in

the beta-1 region representing native C3.

Against serum containing partly activated C3, a precipitin line is obtained which extends from the beta-1 into the alpha-2 region, demonstrating a gradient. In old serum containing totally activated C3 a single precipitin line in the alpha-2 region is obtained. Antisera to C3c cab also react with the fragments C3b, C3bi and smaller fragments, since they all carry antigenic determinants of the C3c domain.

The product does not react with any other protein components of Mouse serum or plasma.

Cross-reactivity:

The antiserum does not cross-react with any other component of Mouse plasma. Interspecies cross-reactivity is a normal feature of antibodies to plasma proteins since they frequently share antigenic determinants. Cross-reactivity of this antiserum has not been tested in detail.

Formulation: PBS, pH 7.2 without preservatives

Label: HRP

State: Lyophilized hyperimmune IgG fraction

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

Reconstitution Method: Restore by adding 1 ml of sterile distilled water.

Concentration: lot specific

Purification: The IgG fraction is isolated and purified from the antiserum and contains the bulk of the

defined antibody specificity. It is free of other serum proteins as tested by

immunoelectrophoresis and double radial immunodiffusion.

Conjugation: HRP

Storage: Prior to reconstitution store at 2-8°C.

Following reconstitution store undiluted at 2-8°C for one week

or (in aliquots) at -20°C for longer. Avoid repeated freezing and thawing.

Stability: Shelf life: one year from despatch.

Gene Name: complement component 3

Database Link: Entrez Gene 24232 Rat

P01026



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Background:

C3 is the most abundant complement protein in rat serum. Its biological function strongly resembles that of C3 in man and other laboratory animal species. It has a central role in the activation system being common to both pathways.

Activation of C3 is achieved by very specific limited proteolysis resulting in the release of a number of degradation fragments. The anaphylotoxin C3a promotes smooth muscle contraction and increases vascular permeability: the large C3b fragment is involved in binding to the complement activator and can be interact with specific receptors to allow efficient clearance of the activating cell or particle; degradation fragments of C3b (C3bi, C3c, C3dg C3d) are important in receptor binding and clearance mechanisms, in virus neutralization and possibly in the immune response.

Synonyms:

CPAMD1, Complement component 3

Note:

Adsorption: Immunoaffinity adsorbed using insolubilized antigens as required, to eliminate antibodies cross-reacting with other with other plasma proteins. The use of insolubilized adsorption antigens prevents the presence of excess adsorbent protein or immune complexes in the antiserum.