SANTA CRUZ BIOTECHNOLOGY, INC.

C4BPα (N-20): sc-17206



BACKGROUND

The complement component proteins C3, C4 and C5 are potent anaphylatoxins that are released during classical complement activation, a system of ligandsurface protein interactions that aid in the elimination of pathogens. These proteins belong to the α 2-Macroglobulin family, but retain distinctive features including an anaphylatoxin domain and a netrin (NTR) domain. They are also expressed as single-chain precursors, which are cleaved into α , β and γ subunits that are linked by disulfide bonds. Complement C4 is an essential component for the activation of the complement pathway, which acts through the receptor CR1 (CD35). Complement C4 is predominately expressed in liver and its precursor contains C4a anaphylatoxin and C4b. The full length C4 protein is cleaved into an α chain, a β chain and a γ chain. C4 exists as two functionally distinct isotypes, C4A and C4B, which react preferentially with amino groups and hydroxyl groups, respectively. Excessive complement activation by C4 is negatively regulated by C4BP (C4 binding protein), a fluid-phase complement inhibitor that protects against complement-induced cell apoptosis. The C4BP complex contains α and γ chains which act together to accelerate inactivation of C4, thereby controlling the classical pathway of complement activation.

REFERENCES

- Scharfstein, J., et al. 1978. Human C4-binding protein. I. Isolation and characterization. J. Exp. Med. 148: 207-222.
- Chung, L.P., et al. 1985. Molecular cloning and characterization of the cDNA coding for C4β-binding protein, a regulatory protein of the classical pathway of the human complement system. Biochem. J. 230: 133-141.
- 3. Blom, A.M., et al. 1999. A cluster of positively charged amino acids in the C4BP α chain is crucial for C4 β binding and factor I cofactor function. J. Biol. Chem. 274: 19237-19245.
- 4. Blom, A.M., et al. 2000. Positively charged amino acids at the interface between α chain CCP1 and CCP2 of C4BP are required for regulation of the classical C3-convertase. Mol. Immunol. 37: 445-453.
- 5. Blom, A.M., et al. 2000. Human C4 β -binding protein has overlapping, but not identical, binding sites for C4 β and streptococcal M proteins. J. Immunol. 164: 5328-5336.
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CHROMOSOMAL LOCATION

Genetic locus: C4BPA (human) mapping to 1q32.2.

SOURCE

C4BP α (N-20) is an affinity purified goat polyclonal antibody raised against a peptide mapping within an internal region of C4BP α of human origin.

STORAGE

Store at 4° C, **DO NOT FREEZE**. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PRODUCT

Each vial contains 200 μg lgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-17206 P, (100 μ g peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

C4BP α (N-20) is recommended for detection of C4BP α of human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for C4BP α siRNA (h): sc-42739, C4BP α shRNA Plasmid (h): sc-42739-SH and C4BP α shRNA (h) Lentiviral Particles: sc-42739-V.

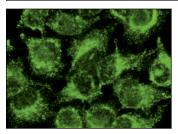
Molecular Weight of C4BPa: 70 kDa.

Positive Controls: HeLa whole cell lysate: sc-2200.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluo-rescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

DATA



C4BPa (N-20): sc-17206. Immunofluorescence staining of methanol-fixed HeLa cells showing cytoplasmic localization.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.