Cya A (bN-13): sc-17899



The Power to Question

BACKGROUND

Bordetella pertussis, the causative agent of whooping cough, secretes several toxins implicated in this disease. One of these putative virulence factors is the adenylate cyclase toxin (Cya A or ACT) that elevates intracellular cAMP in eukaryotic cells to cytotoxic levels upon activation by endogenous calmodulin. The Bordetella pertussis Cya toxin-encoding locus (Cya) is composed of five genes. The CyaA gene encodes a virulence factor Cya A, exhibiting adenylate cyclase, hemolytic and invasive activities. Cya A is related to the RTX (repeats in toxin) family of pore-forming toxins. Like all RTX toxins, Cya A is synthesized as a protoxin (proCya A), encoded by the CyaA gene. Activation to the mature cell-invasive toxin involves palmitoylation of lysine 983 and is dependent on co-expression of Cya C. The CyaB, D and E gene products are necessary for Cya A transport, and the CyaC gene product is required to activate Cya A. Additionally, Cya A uses the α M β 2 integrin (CD11b/CD18) as a cell receptor. Thus, the cellular distribution of CD11b, mostly on neutrophils, macrophages, and dendritic and natural killer cells, supports a role for Cya A in disrupting the early, innate antibacterial immune response.

REFERENCES

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- Gross, M.K., Au, D.C., Smith, A.L., and Storm, D.R. 1992. Targeted mutations that ablate either the adenylate cyclase or hemolysin function of the bifunctional Cya A toxin of *Bordetella pertussis* abolish virulence. Proc. Natl. Acad. Sci. USA 89: 4898-4902.
- 3. Ehrmann, I.E., Weiss, A.A., Goodwin, M.S., Gray, M.C., Barry, E., and Hewlett, E.L. 1992. Enzymatic activity of adenylate cyclase toxin from *Bordetella pertussis* is not required for hemolysis. FEBS Lett. 304: 51-56.
- Westrop, G.D., Hormozi, E.K., Da Costa, N.A., Parton, R., and Coote, J.G. 1996. Bordetella pertussis adenylate cyclase toxin: proCya A and Cya C proteins synthesised separately in Escherichia coli produce active toxin in vitro. Gene 180: 91-99.
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SOURCE

Cya A (bN-13) is an affinity purified goat polyclonal antibody raised against a peptide mapping at the N-terminus of Cya A of B. pertussis origin.

PRODUCT

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

Blocking peptide available for competition studies, sc-17899 P, (100 μg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

Cya A (bN-13) is recommended for detection of Cya A of *B. pertussis* origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Molecular Weight of Cya A: 233 kDa.

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.

STORAGE

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

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.

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