

PRODUCT SPECIFICATION



HYB 333-02B Anti Pertussis Toxin, Biotinylated

Mouse monoclonal antibody

Article No.	59284															
Product Name	HYB 333-02B Anti Pertussis Toxin, Biotinylated															
Clone	21.3. D11A2G5															
Subclass	IgG1 / kappa															
Description	<p>Preparation: Protein-A purified Biotinylated</p> <p>Concentration: 1 mg/mL ± 10%, based on A₂₈₀. See Certificate of Analysis for details.</p> <p>Solvent: PBS, pH 7.2 – 7.4</p> <p>Storage: 2–8 °C</p>															
Antigen	<p>Pertussis toxin (islet-activating protein) is the major protein toxin produced by virulent strains of <i>Bordetella pertussis</i>, the organism that causes whooping cough (1). As revealed by polyacrylamide gel electrophoresis, the purified protein consists of five dissimilar subunits: S1 (MW 28,000), S2 (MW 23,000), S3 (MW 22,000), S4 (MW 11,700) and S5 (MW 9,300), in a molar ratio of 1:1:1:2:1. The A-protomer, S1 is responsible for the enzymatic activity of the toxin. Together, S2, S3, S4 and S5 comprise the B-oligomer, responsible for binding the toxin to the cell surface (2).</p>															
Immunogen	Pertussis toxin.															
Specificity	HYB 333-02 (3) reacts with pertussis toxin. Some reactivity towards the toxoid is also present.															
Epitope Specificity	HYB 333-02 has a different epitope specificity compared with HYB 333-01, HYB 333-03, HYB 333-05, and HYB 333-09..															
Reactivity	<p>HYB 333-02 (MAb clone 21.3 D11) (3) is well suited for ELISA based measurement of pertussis toxin.</p> <p>We recommend using HYB 333-02 as detection antibody (conjugated with signal molecules) in combination with HYB 333-01 as catching antibody.</p> <p>HYB 333-02 can neutralize pertussis toxin when measured by Chinese Hamster Ovary cell assays, Leucocytosis promoting activity and <i>in-vivo</i> experiments (4,6).</p>															
Culture Medium	Dulbecco's modified Eagle's medium with 10% fetal calf serum..															
Fusion Partner	X63-Ag8.653..															
Immunization	Female CF1xBalb/c F1 hybrid mice were immunized i.p. with immunogen.															
Application	<table><thead><tr><th>Method</th><th>Usability</th><th>References</th></tr></thead><tbody><tr><td>ELISA</td><td>yes</td><td>4-6</td></tr><tr><td>Immunoblotting</td><td>nd.</td><td>4</td></tr><tr><td>Immunofluorescence</td><td>nd.</td><td></td></tr><tr><td>Neutralization</td><td>yes</td><td>4,6</td></tr></tbody></table>	Method	Usability	References	ELISA	yes	4-6	Immunoblotting	nd.	4	Immunofluorescence	nd.		Neutralization	yes	4,6
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See next page

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References

- 1) **Pittman, M.** (1979) Rev. Infect. Dis. 1, 401-412.
- 2) **Tamura, M., Nogomori, K., Murai, S., Yajima, M., Ito, K., Katada, T., Ui, M. and Ishi, S.** (1982) Biochem. 21, 5516-5522.
- 3) HYB 333-02 is known from the literature as "21.3 D11".
- 4) **Schou C, Au-Jensen M, Heron I.** The interaction between pertussis toxin and 10 monoclonal antibodies. Acta Pathol Microbiol Immunol Scand C. 1987 Oct;95(5):177-87.
- 5) **Ibsen, P.H. and Heron, I.** (1990) Quantification of pertussis toxin in an enzyme linked immunosorbent assay with improved specificity. Biologicals, 18, 123-126.
- 6) **Ibsen, P.H.** (1996). The effect of formaldehyde, hydrogen peroxide and genetic detoxification of pertussis toxin on epitope recognition by murine monoclonal antibodies. Vaccine, 14, 359-368.

Conditions

For research use only. Not for use in diagnostic procedures. Not for therapeutic use or applications.

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