

Anti- *Vibrio parahaemolyticus* TDH / TRH Toxin antibody, mouse monoclonal (vp-01)

64-013 100 µg

Shipping and Storage: Shipped at 4°C or -20°C. Store at -20°C. Do not freeze.

Immunogen: Culture supernatant of *V. parahemolyticus*

Form: 0.5 mg/ml in PBS- with 50% glycerol, filter sterilized.

Purity: IgG, affinity-purified with Protein A/G mix

Isotype: mouse IgG1

Reactivity: Reacts with *V. parahemolyticus* TDH and TRH toxins

Applications:

1. Western blotting (1/500~1/1,000)
2. ELISA (assay dependent)

This antibody is useful for detecting food-poisoning *V. hemolyticus* strains.

Background: Many *Vibrio parahaemolyticus* strains isolated as a cause of food poisoning, produce toxin called hemolysin, and this is the main cause of illness. Two kinds of hemolysins, **T**hermo-resistant **D**irect **H**emolysin (TDH) and **T**DH **R**elated **H**emolysin (TRH), are known. TDH is the heat labile toxin protein of molecular weight 21.3 kDa (189 aa). Homology of TRH (21.1 kDa, 189 aa) with TDH is about 60%, and shows partial antigenic similarities.

Data Link: UniProtKB: [P19249](#) (Thermostable direct hemolysin1),
[Q769J9](#) (TDH related hemolysin)

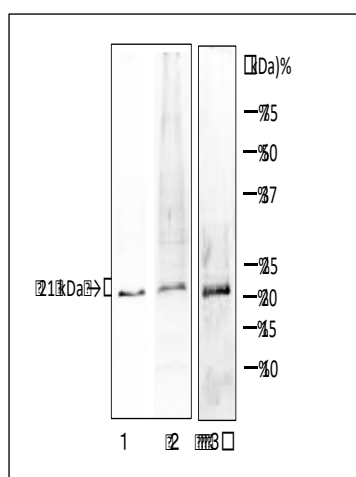


Fig.1. Detection of *V. parahaemolyticus* TDH and TRH by Western blotting with MAb (vp-01)

1. Culture medium of *V. parahaemolyticus* (trh⁺)
2. Culture medium of *V. parahaemolyticus* (tdh⁺)
3. Culture medium of *V. parahaemolyticus* (trh⁺)

MAb (vp-01) was used at 1/1,000 dilution in lanes 1 and 2.

Polyclonal antiTRH antibody (BioAcademia 64-015) was used at 1/1,000 dilution in lane 3.

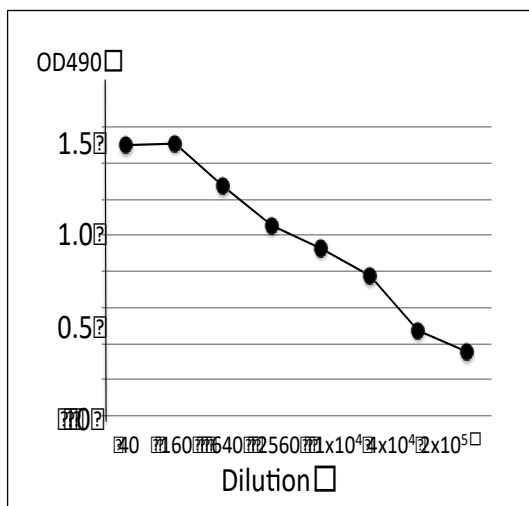


Fig.2. Titration of antibody reactivity of MAb (vp-1) by indirect ELISA, using culture medium of *V. parahaemolyticus* trh⁺

The wells of plate were coated with culture medium of *V. parahaemophilus* trh⁺(100 μ l, 1 μ g/ml). After blocking with 5% skim milk, 100 μ l of antibody at the indicated dilution was added to the each well. HRP-conjugate goat anti-mouse IgG (100 μ l, x2000 dilution) was added. Color was developed with orthophenylenediamine as substrate. Optical densities (OD) measured at 490nm.

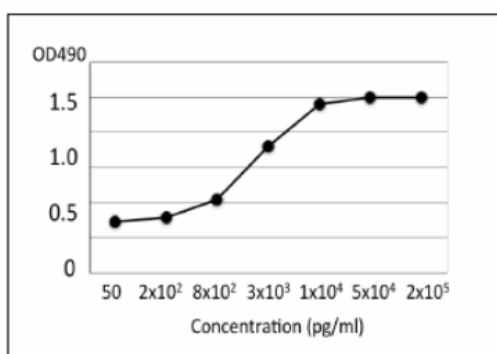


Fig.3. Indirect ELISA of TDH in extract of *V. parahaemolyticus* trh⁺ with MAb (vp-01)

ELISA plate was coated with indicated amounts of the extract of *V. parahaemolyticus* trh⁺. MAb (vp-01) was used at 1/500 dilution. ELISA was performed as in Fig.2.

Table 1. Reactivity of MAb (vp-01) with various food poisoning bacteria.

	ELISA	WB
<i>Vibrio parahaemolyticus</i> (NBRC12711)	+	21K
Other 3 isolated strains	+	21K
<i>Salmonella</i> Enteritidis	-	-
<i>E. coli</i> 0157:H7	-	
<i>Staphylococcus aureus</i>	-	
<i>Bacillus cereus</i>	-	
Partially purified TDH	+	
TDH: Thermostable direct hemolysin		

Reference: There has been no publication using this antibody.

Please let us know when your research using this antibody is published. We will offer one vial of our antibody as compliment.