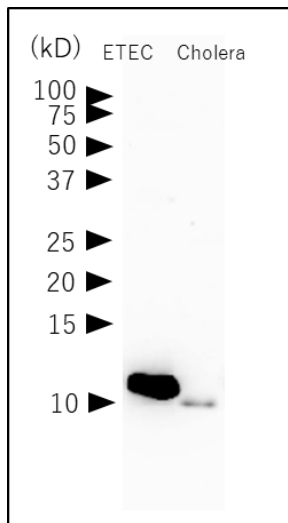


抗 LT サブユニット B (E.coli) 抗体, マウスモノクローナル (ec-01)

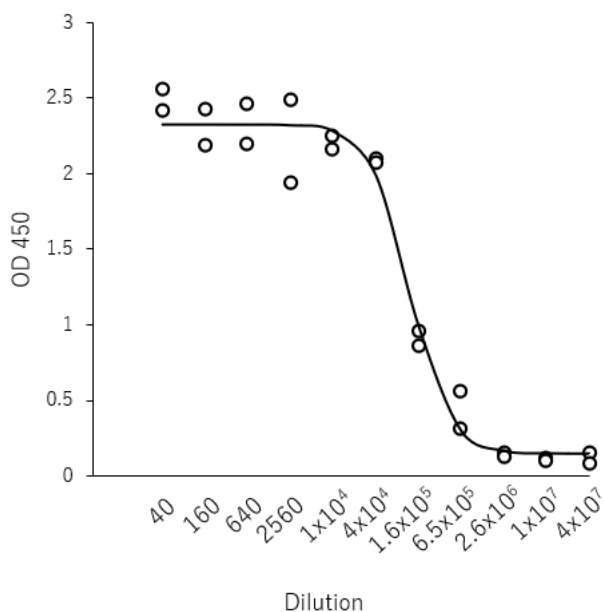
<b>Product code</b>	64-023
<b>Size</b>	100 µg
<b>Storage</b>	-20°C
<b>Concentration</b>	0.5 mg/ml
<b>Buffer</b>	PBS- with 50% glycerol
<b>Purity</b>	Purified IgG fraction with protein A from hybridoma cell culture medium
<b>Immunogen</b>	Crude extract of <i>Escherichia coli</i> ( ETEC LT <sup>+</sup> ) cells
<b>Isotype</b>	Mouse IgG2ak
<b>Reactivity</b>	subunit B of <i>E. coli</i> LT and <i>V. Cholera</i> CT.
<b>Special notes</b>	N/A
<b>Application</b>	1. Western blotting (1/500~1/5000 ) 2. ELISA (assay dependent) This antibody is useful for detecting food poisoning Enterotoxigenic <i>E. coli</i> (ETEC)
<b>Background</b>	Pathogenic <i>Escherichia coli</i> is one of the major causative agents of food poisoning. One group of them, enterotoxigenic <i>E. coli</i> (ETEC) produces some toxins. Heat labile enterotoxin (LT) produced by ETEC is similar to cholera toxin (CT). The identity of the amino acid sequences of LT and CT is about 80% and both toxins are consisted of one subunit A and five subunit B. LT continuously activates adenylate cyclase and elevated level of cAMP inhibits absorption of Na <sup>+</sup> by intestinal villi cells, and stimulates secretion of Cl <sup>-</sup> by villi and crypt cells, thus causing diarrhea. Subunit A possesses signal peptide of the amino acids 1-18, and the mature form consists of 19-258 amino acids (MW: 28.8 kDa). Subunit B has signal peptide of 1-21, and the mature form consists of 22-124 amino acids (MW: 11.8 kDa). The holotoxin MW is 86.4 kDa.
<b>Data Link</b>	UniProtKB: <a href="#">P0CK94</a> (Heat-labile enterotoxin B chain)
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.	

**Data Images:** 抗 LT サブユニット B (*E.coli*) 抗体, マウスモノクローナル (ec-01)



**Fig.1. Detection of LT in crude extract of *E. coli* ETEC strain and Cholera toxin (#01-511) by Western blot.**

The anti- LT toxin subunit B antibody was used at 1/1,000 dilution.



**Fig.2. Titration of antibody reactivity of MAb by indirect ELISA using crude extract of ETEC cells.**

The wells of plate were coated with crude extract of *E. coli*. After blocking with 5% skim milk, 100  $\mu$ l of antibody at the indicated dilutions was added to the each well. HRP-conjugated goat anti-mouse IgG (100 $\mu$ l, x 2000 dilution) was added. Color was developed with orthophenylenediamine as substrate. Optical densities (OD) measured at 450nm.

**Reference:** There has been no publication using this antibody.

**Related Product:**

64-020 抗 LT (*E.coli*) 抗体, ウサギ抗血清