## Anti-His-tag mAb-Biotin

CODE No. D291-6

| CLONALITY | Monoclonal |
| :--- | :--- |
| CLONE | OGHis |
| ISOTYPE | Mouse IgG2a $\kappa$ |
| QUANTITY | $50 \mu \mathrm{~L}, 1 \mathrm{mg} / \mathrm{mL}$ |

SOURCE Purified IgG from hybridoma supernatant
IMMUNOGEN 6xHis tagged protein
FORMULATION $\quad 1 \mathrm{mg} / \mathrm{mL}$ in PBS ( pH 7.2 ) containing $1 \%$ BSA and $0.09 \% \mathrm{NaN}_{3}$
*Azide may react with copper or lead in plumbing system to form explosive metal azides. Therefore, always flush plenty of water when disposing materials containing azide into drain.
STORAGE This antibody solution is stable for one year from the date of purchase when stored at $4^{\circ} \mathrm{C}$.

## APPLICATION-CONFIRMED <br> ELISA <br> $0.25-1 \mu \mathrm{~g} / \mathrm{mL}$

For more information, please visit our web site https://ruo.mbl.co.jp/.

The descriptions of the following protocols are examples. Each user should determine the appropriate condition.

## ELISA

1) Add $100 \mu \mathrm{~L} /$ well of antigen solution in PBS to the 96 -well plate. Incubate overnight at $4^{\circ} \mathrm{C}$.
2) Add $200 \mu \mathrm{~L} /$ well of $5 \%$ BSA in PBS. Incubate overnight at $4^{\circ} \mathrm{C}$.
3) Add $100 \mu \mathrm{~L} /$ well of primary antibody diluted with PBS as suggested in the APPLICATION for 1 hour at room temperature. (The concentration of antibody will depend on the conditions.)
4) Wash the plate with PBS-T [ $0.05 \%$ Tween- 20 in PBS] (4 times).
5) Add $100 \mu \mathrm{~L} /$ well of $1: 50,000$ diluted HRP-conjugated streptavidin in PBS. Incubate for 1 hour at room temperature.
6) Wash the plate with PBS-T (4 times).
7) Add $100 \mu \mathrm{~L} /$ well of substrate solution (ex. TMB). Incubate for appropriate time at room temperature.
8) Add $100 \mu \mathrm{~L} /$ well of stop solution (ex. $1 \mathrm{M} \mathrm{H}_{2} \mathrm{SO}_{4}$ ).
9) Read absorbance at 450 nm .


ELISA for measurement of His-tagged protein

