

## Product datasheet

# anti-6-His-tag mouse monoclonal, 6His, lyophilized, purified

#### Short overview

 Cat. No.
 910HIS

 Quantity
 25 μg

Concentration 0.25 mg/ml after reconstitution with 100 µl PBS

#### Product description

HostMouseAntibody TypeMonoclonalIsotypeIgG1Clone6His

ImmunogenRecombinant protein containing the sequence HHHHHHFormulationLyophilized; reconstitute in 100 μl sterile PBS, pH 7.4

Synomym ChR2

Conjugate Unconjugated

Purification Affinity chromatography

**Storage before** 2-8°C until indicated expiry date

reconstitution

Storage after -20°C (avoid freeze/thaw cycles)

reconstitution

Intended useResearch use onlyApplicationICC/IF, IP, WB

**Reactivity** 6-His

### **Applications**

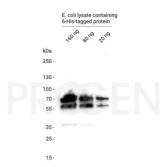
Immunocytochemistry (ICC)Assay dependentImmunoprecipitation (IP)Assay dependent

Western Blot (WB) 1:2,000-1:5,000 (0.125-0.05 μg/ml)

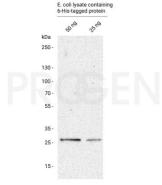
#### Background

The monoclonal 6-His antibody recognizes polyhistidine (6-His). The 6-His tag is commonly added to recombinant proteins and can be used for detection or purification of the tagged protein.

#### **Product images**



Western blot analysis of E. coli lysate containing 6-His-tagged protein with anti-6-His-tag antibody. Western blot analysis was performed on 160 ng, 80 ng or 20 ng of E. coli lysate containing 6-His-tagged protein. Cells were lysed with SDS sample buffer. The PVDF membrane was blocked with 5% dry milk in PBST for 1 h at RT. The primary antibody anti-6-His-tag mouse monoclonal, 6His (Cat. No. 910HISL) was diluted in blocking buffer (antibody concentration 0.125  $\mu$ g/ml) and incubated for 1 h at RT. The secondary antibody goat anti-mouse IgG polyclonal, HRP conjugate was also diluted in blocking buffer (antibody concentration 0.2  $\mu$ g/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using PierceTM ECL Western Blotting Substrate.



Western blot analysis of E. coli lysate containing 6-His-tagged protein with anti-6-His-tag antibody. Western blot analysis was performed on 50 ng or 25 ng of E. coli lysate containing 6-His-tagged protein. Cells were lysed with SDS sample buffer. The PVDF membrane was blocked with 5% dry milk in PBST for 1 h at RT. The primary antibody anti-6-His-tag mouse monoclonal, 6His (Cat. No. 910HISL) was diluted in blocking buffer (antibody concentration 0.05  $\mu$ g/ml) and incubated for 1 h at RT. The secondary antibody goat anti-mouse IgG polyclonal, HRP conjugate was also diluted in blocking buffer (antibody concentration 0.2  $\mu$ g/ml) and incubated for 1 h at RT. The bands were visualized by chemiluminescent detection using PierceTM ECL Western Blotting Substrate.