

Mouse anti His-Tag mAb

Catalog No.: AE003 224 Publications

Basic Information

Observed MW 27kDa,35kDa/

Calculated MW

Category **Monoclonal Antibody**

Applications WB, IF/ICC, IP, ELISA

Cross-Reactivity Species independent

CloneNo number AMC0149

Recommended Dilutions

WB	1:5000 - 1:30000
IF/ICC	1:5000-1:10000
IP	0.5ug-4ug antibody for 200ug-400ug extracts of whole cells
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Contact

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www.abclonal.com

Background

Consecutive histidine residues (usually 6 to 10 in length) are often inserted into the amino acid sequences of recombinant proteins. The resulting His-tagged proteins can be detected by using His antibodies.

Immunogen Information

0000	Gene ID	Swi	iss Prot
.0000	Immunogen		
ly for	Synthetic peptide.	This information is considered	d to be commercially sensitive.
tracts	Synonyms		
cells	His;His tag;His-tag	I	
arting			
n is 1			
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ecific nents.	Product Information		
	Source	Isotype	Purification

Affinity purification

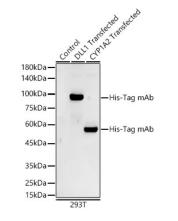
Storage

Mouse

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS with 0.09% Sodium azide, 50% glycerol, pH7.3.

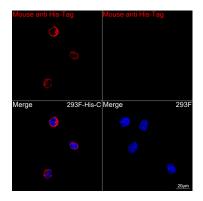
IgG1

Validation Data

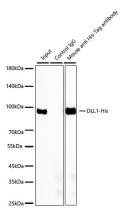


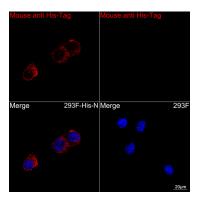
Western blot analysis of various lysates, using His-Tag mAb (AE003) at 1:5000 dilution. Secondary antibody: HRP-conjugated Goat anti-Mouse IgG (H+L) (AS003) at 1:10000 dilution.

Lysates/proteins: 25µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 90s.



Confocal imaging of 293F cells transfected with His-C using Mouse anti His-Tag mAb (AE003, dilution 1:10000) followed by a further incubation with Cy3 Goat Anti-Mouse IgG (H+L) (AS008, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.





Confocal imaging of 293F cells transfected with His-N using Mouse anti His-Tag mAb (AE003, dilution 1:10000) followed by a further incubation with Cy3 Goat Anti-Mouse IgG (H+L) (AS008, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.

Immunoprecipitation of DLL1-His from 300 µg extracts of 293F cells transfected with a DLL1 expression vector containing a single N-terminal His-Tag was performed using 1 µg of Mouse anti His-Tag mAb (AE003). Mouse IgG isotype control (AC011) was used to precipitate the Control IgG sample. The IP sample was eluted with 1X reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using Rabbit anti His-tag mAb (AE086) at a dilution of 1:10000.