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Mouse anti Myc-Tag mAb

Catalog No.: AE010 102 Publications

Basic Information

Observed MW

62kDa,60kDa

Calculated MW

Category

SMab Recombinant Monoclonal Antibody

Applications

WB,IP,ELISA,IF

Cross-Reactivity

Species independent

CloneNo number

AMC0504

Background

Protein tags are peptide sequences genetically grafted onto a recombinant protein. Often these tags are removable by chemical agents or by enzymatic means, such as proteolysis or intein splicing. Tags are attached to proteins for various purposes. Epitope tags are short peptide sequences which are chosen because high-affinity antibodies can be reliably produced in many different species. These are usually derived from viral genes, which explain their high immunoreactivity. Epitope tags include V5-tag, Myc-tag, HA-tag and NE-tag. These tags are particularly useful for western blotting, immunofluorescence and immunoprecipitation experiments, although they also find use in antibody purification.

Recommended Dilutions

WB 1:2000 - 1:6000

IF 1:1000-1:5000

IP 0.5μg-4μg antibody for

200µg-400µg 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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Immunogen Information

Gene ID Swiss Prot

Immunogen

A synthetic peptide corresponding to Myc tag.

Synonyms

Myc;Myc tag;Myc-tag

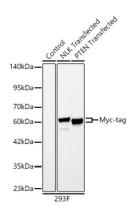
Product Information

SourceIsotypePurificationMouseIgG1Affinity purification

Storage

Store at -20℃. Avoid freeze / thaw cycles.

Buffer: PBS with 0.05% proclin300,50% glycerol,pH7.3.



Western blot analysis of lysates from wild type (WT) and 293F cells transfected with NLK-Myc (C-terminal) or PTEN-Myc (C-terminal) using Mouse anti Myc-Tag mAb (AE010) at 1:10000 dilution incubated overnight at 4°C.

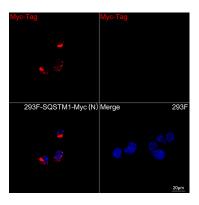
Secondary antibody: HRP-conjugated Goat anti-Mouse IgG (H+L) (AS003) at 1:10000 dilution.

Lysates/proteins: 30 µg per lane.

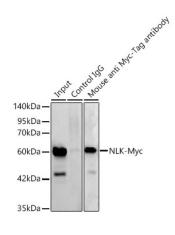
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020)

.Exposure time: 30s.



Confocal imaging of 293F cells transfected with SQSTM1-Myc(N) cells using Mouse anti Myc-Tag mAb (AE010, dilution 1:2000) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). Objective: 100x.



Immunoprecipitation of NLK-Myc from 300 μ g extracts of 293F cells transfected with a NLK expression vector containing a single C-terminal Myc-Tag was performed using 3 μ g of Mouse anti Myc-Tag mAb (AE010). Mouse IgG isotype control (AC011) was used to precipitate the Control IgG sample. IP samples were eluted with 1X Laemmli Buffer. The Input lane represents 10 % of the total input. Western blot analysis of immunoprecipitates was conducted using Mouse anti Myc-Tag mAb (AE102) at a dilution of 1:3000.