

NEDD8 Rabbit mAb

Catalog No.: A29552 **Recombinant**

Basic Information

Observed MW

9 kDa/90 kDa

Calculated MW

9 kDa

Category

Primary antibody

Applications

WB, IP, IF/ICC, IHC-P, ELISA

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC4057

Background

Enables ubiquitin protein ligase binding activity. Acts upstream of or within protein neddylation. Located in nucleoplasm. Biomarker of Parkinson's disease; alcohol-associated liver disease; and malignant astrocytoma.

Recommended Dilutions

WB 1:3000 - 1:15000

IP 0.5 µg - 4 µg antibody for
200 µg - 400 µg extracts of
whole cells

IF/ICC 1:100 - 1:400

IHC-P 1:800 - 1:3200

ELISA Recommended starting
concentration is 1 µg/mL.
Please optimize the
concentration based on your
specific assay requirements.
For high-ratio antibody
dilutions (≥1:10)

Contact

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✉ | cn.market@abclonal.com.cn

Immunogen Information

Gene ID

4738

Swiss Prot

Q15843

Immunogen

This information is considered to be commercially sensitive.

Synonyms

NEDD-8

Product Information

Source

Rabbit

Isotype

IgG

Purification

Affinity purification

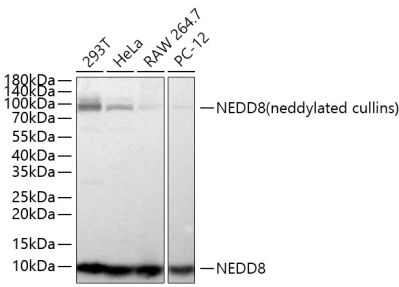
Storage

Store at -20°C

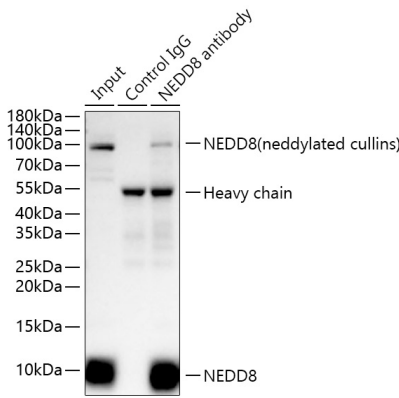


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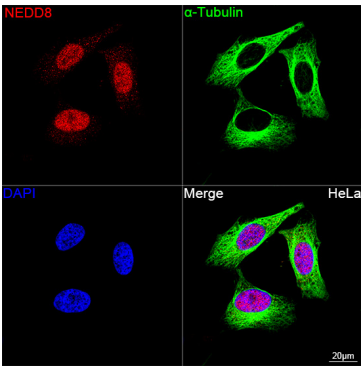
Validation Data



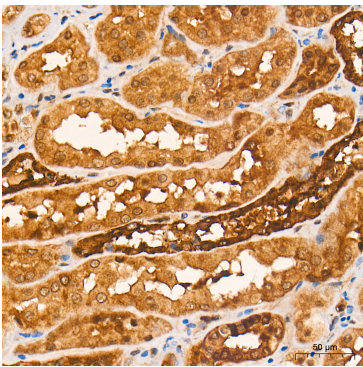
Western blot analysis of various lysates using NEDD8 Rabbit mAb (A29552) at 1:5000 dilution incubated overnight at 4°C. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 25 µg per lane. Blocking buffer: 3% nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 1 s.



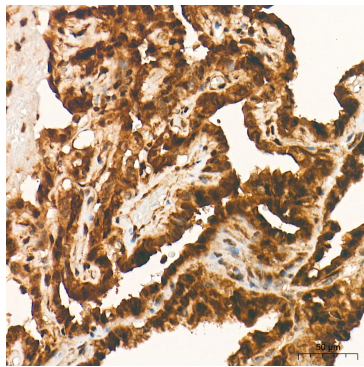
Immunoprecipitation of NEDD8 from 300 µg extracts of HeLa cells was performed using 2 µg of NEDD8 Rabbit mAb (A29552). Rabbit Control IgG (AC005) was used to precipitate the Control IgG sample. IP samples were eluted with 1x Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using NEDD8 Rabbit mAb (A29552) at a dilution of 1:5000.



Confocal imaging of HeLa cells using NEDD8 Rabbit mAb (A29552, dilution 1:200) followed by a further incubation with Cy3-conjugated Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α-Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) (AS037, dilution 1:800) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

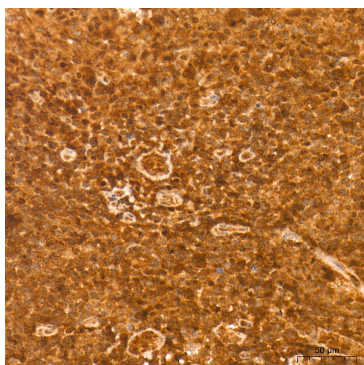


Immunohistochemistry analysis of paraffin-embedded Human kidney tissue using NEDD8 Rabbit mAb (A29552) at a dilution of 1:800 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

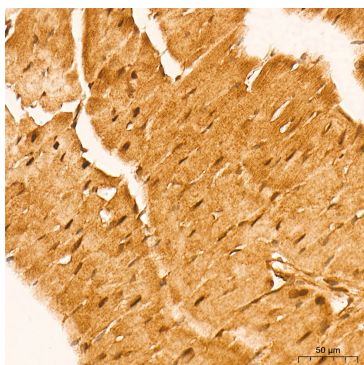


Immunohistochemistry analysis of paraffin-embedded Human thyroid cancer tissue using NEDD8 Rabbit mAb (A29552) at a dilution of 1:800 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.

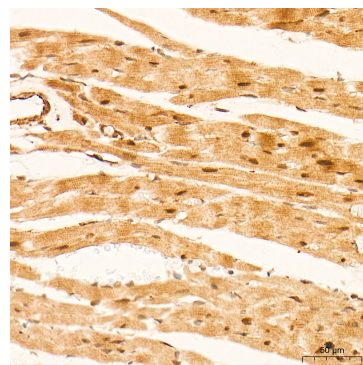
Validation Data



Immunohistochemistry analysis of paraffin-embedded Human tonsil tissue using NEDD8 Rabbit mAb (A29552) at a dilution of 1:800 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Mouse heart tissue using NEDD8 Rabbit mAb (A29552) at a dilution of 1:800 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Rat heart tissue using NEDD8 Rabbit mAb (A29552) at a dilution of 1:800 (40x lens). High pressure antigen retrieval performed with 0.01M Tris-EDTA Buffer (pH 9.0) prior to IHC staining.