

A25919

Leader in Biomolecular Solutions for Life Science



TRAP1 Rabbit mAb

Catalog No.: A25919

Recombinant

Basic Information

Observed MW

80kDa

Calculated MW

80kDa

Category

SMab Recombinant Monoclonal
Antibody

Applications

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

Cross-Reactivity

Human,Mouse

CloneNo number

ARC67266

Background

This gene encodes a mitochondrial chaperone protein that is member of the heat shock protein 90 (HSP90) family. The encoded protein has ATPase activity and interacts with tumor necrosis factor type I. This protein may function in regulating cellular stress responses. Alternate splicing results in multiple transcript variants.

Recommended Dilutions

WB	1:2000 - 1:8000
IHC-P	1:200 - 1:800
IF/ICC	1:200 - 1:800
IP	0.5µg-4µg antibody for 400µg-600µg extracts of whole cells
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID	Swiss Prot
10131	Q12931

Immunogen

Recombinant protein (or fragment).This information is considered to be commercially sensitive.

Synonyms

HSP75; HSP 75; HSP90L; TRAP-1

Product Information

Source	Isotype	Purification
Rabbit	IgG	Affinity purification

Storage

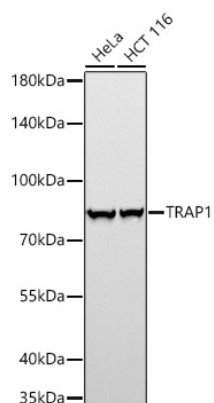
Store at -20°C. Avoid freeze / thaw cycles.

Buffer: PBS with 0.09% Sodium azide,0.05% BSA,50% glycerol,pH7.3.

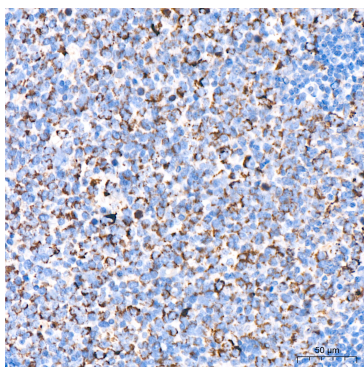
Contact

 www.abclonal.com

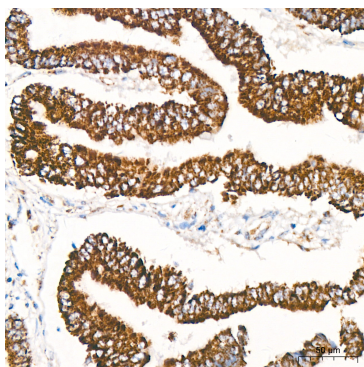
Validation Data



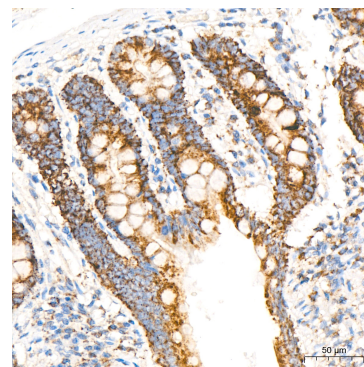
Western blot analysis of various lysates using TRAP1 Rabbit mAb (A25919) at 1:2000 dilution incubated overnight at 4°C.
 Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.
 Lysates/proteins: 25 µg per lane.
 Blocking buffer: 3% nonfat dry milk in TBST.
 Detection: ECL Basic Kit (RM00020).
 Exposure time: 20s.



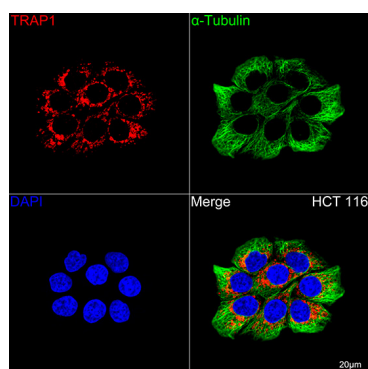
Immunohistochemistry analysis of paraffin-embedded Human tonsil tissue using TRAP1 Rabbit mAb (A25919) at a dilution of 1:300 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



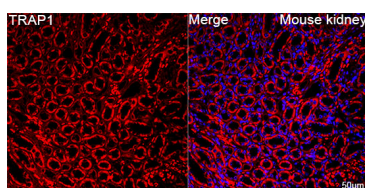
Immunohistochemistry analysis of paraffin-embedded Human colon carcinoma tissue using TRAP1 Rabbit mAb (A25919) at a dilution of 1:300 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Human colon tissue using TRAP1 Rabbit mAb (A25919) at a dilution of 1:300 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate buffer (pH 6.0) prior to IHC staining.



Confocal imaging of HCT 116 cells using TRAP1 Rabbit mAb (A25919, dilution 1:200) followed by a further incubation with Cy3 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

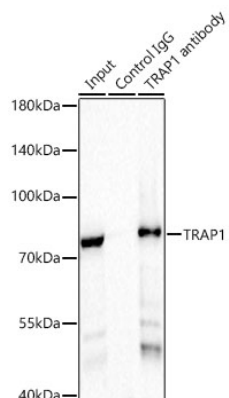


Confocal imaging of paraffin-embedded Mouse kidney tissue using TRAP1 Rabbit mAb (A25919, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). DAPI was used for nuclear staining (Blue). High pressure antigen retrieval performed with 0.01M Citrate Buffer (pH 6.0) prior to IF

Validation Data

(H+L) Ab (AS076, dilution 1:500)
(Green). DAPI was used for nuclear
staining (Blue). Objective: 100x.

staining. Objective: 40x.



Immunoprecipitation of TRAP1 from 500 µg extracts of HeLa cells was performed using 2 µg of TRAP1 Rabbit mAb (A25919). Rabbit IgG isotype control (AC042) was used to precipitate the Control IgG sample. IP samples were eluted with 1X non-reducing Laemmli Buffer. The Input lane represents 10% of the total input. Western blot analysis of immunoprecipitates was conducted using TRAP1 Rabbit mAb (A25919) at a dilution of 1:2000.