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SQSTM1/p62 Rabbit mAb

Catalog No.: A19700 Recombinant 98 Publications

Basic Information

Observed MW

62kDa

Calculated MW

48kDa

Category

SMab Recombinant Monoclonal Antibody

Applications

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

Cross-Reactivity

Human, Mouse, Rat

CloneNo number

ARC0180

Background

This gene encodes a multifunctional protein that binds ubiquitin and regulates activation of the nuclear factor kappa-B (NF-kB) signaling pathway. The protein functions as a scaffolding/adaptor protein in concert with TNF receptor-associated factor 6 to mediate activation of NF-kB in response to upstream signals. Alternatively spliced transcript variants encoding either the same or different isoforms have been identified for this gene. Mutations in this gene result in sporadic and familial Paget disease of bone.

Recommended Dilutions

WB 1:20000 - 1:80000

IHC-P 1:500 - 1:5000

IF/ICC 1:50 - 1:200

IP0.5μg-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 ID8878

Swiss Prot
Q13501

Immunogen

A synthetic peptide corresponding to a sequence within amino acids 341-440 of human SQSTM1/p62 (Q13501).

Synonyms

p60; p62; A170; DMRV; OSIL; PDB3; ZIP3; p62B; NADGP; FTDALS3; SQSTM1/p62

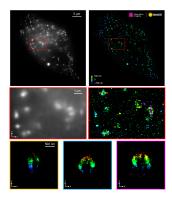
Product Information

SourceIsotypePurificationRabbitIgGAffinity purification

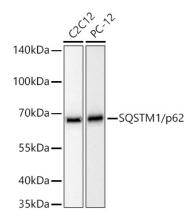
Storage

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

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



The STORM super-resolution (SR) imaging of HeLa cells using SQSTM1/p62 Rabbit mAb (A19700, ABclonal) at dilution of 1:200 with 3% paraformaldehyde (PFA) +0.1% glutaraldehyde (GA) fixation. The immunostaining was performed by Full Automatic Immunofluorescence Workflow System (Workflow Ultra300, Nano-Micro imaging, China). Image was performed with Single-Molecule Localization Super-Resolution Microscopy (STORM Ultra300, Nano-Micro imaging, China). We acknowledge Nano-Micro imaging Biotechnology Co., Ltd. in SR image processing and kindly providing this image.



Western blot analysis of various lysates using SQSTM1/p62 Rabbit mAb (A19700) at 1:20000 dilution incubated overnight at 4° C.

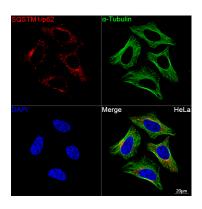
Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution.

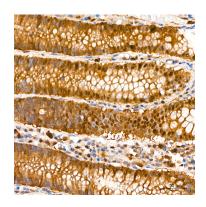
Lysates/proteins: 25 µg per lane.

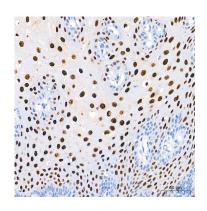
Blocking buffer: 3% nonfat dry milk in TBST.

Detection: ECL Basic Kit (RM00020).

Exposure time: 10s.





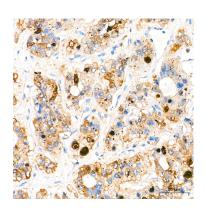


Validation Data

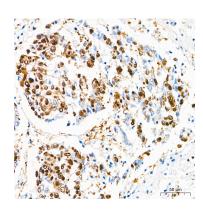
Confocal imaging of HeLa cells using SQSTM1/p62 Rabbit mAb (A19700, dilution 1:200) followed by a further incubation with Cy3 Goat Anti-Rabbit IgG (H+L) (AS007, dilution 1:500) (Red). The cells were counterstained with α -Tubulin Mouse mAb (AC012, dilution 1:400) followed by incubation with ABflo® 488-conjugated Goat Anti-Mouse IgG (H+L) Ab (AS076, dilution 1:500) (Green). DAPI was used for nuclear staining (Blue). Objective: 100x.

Immunohistochemistry analysis of paraffin-embedded Human colon tissue using SQSTM1/p62 Rabbit mAb (A19700) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer(pH 6.0) prior to IHC staining.

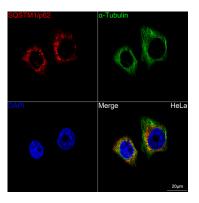
Immunohistochemistry analysis of paraffin-embedded Human esophagus tissue using SQSTM1/p62 Rabbit mAb (A19700) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer(pH 6.0) prior to IHC staining.



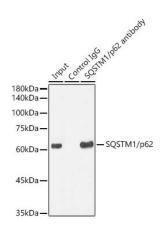
Immunohistochemistry analysis of paraffin-embedded Human liver cancer tissue using SQSTM1/p62 Rabbit mAb (A19700) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer(pH 6.0) prior to IHC staining.



Immunohistochemistry analysis of paraffin-embedded Human lung squamous carcinoma tissue using SQSTM1/p62 Rabbit mAb (A19700) at a dilution of 1:500 (40x lens). High pressure antigen retrieval performed with 0.01M Citrate Buffer(pH 6.0) prior to IHC staining.



Confocal imaging of HeLa cells using SQSTM1/p62 Rabbit mAb (A19700, dilution 1:200) (Red). The cells were counterstained with Alphatubulin (ubiquitous) chain Rabbit mAb (AC049, dilution 1:100) (Green). DAPI was used for nuclear staining (Blue). Objective: 60x.



Immunoprecipitation analysis of 300 μ g extracts from HeLa cells using 3 μ g SQSTM1/p62 Rabbit mAb (A19700). Western blot was performed from the immunoprecipitate using SQSTM1/p62 Rabbit mAb (A19700) at a dilition of 1:1000.