

Anti-p62/SQSTM1-Alexa Fluor[®] 488

CODE No. M162-A48
CLONALITY Monoclonal
CLONE 5F2
ISOTYPE Mouse IgG1 κ
QUANTITY 100 μ L, 1 mg/mL

SOURCE Purified IgG from hybridoma supernatant
IMMUNOGEN Human p62, 120-440 aa (recombinant)
FORMURATION PBS containing 1% BSA and 0.09% NaN₃.

*Azide may react with copper or lead in plumbing system to form explosive metal azides. Therefore, always flush plenty of water when disposing materials containing azide into drain.

STORAGE This antibody solution is stable for one year from the date of purchase when stored at 4°C.

APPLICATIONS-CONFIRMED

Immunocytochemistry 2 μ g/mL
Flow cytometry 1 μ g/mL

SPECIES CROSS REACTIVITY on IC

Species	Human	Mouse	Rat	Hamster
Cells	Transfectant	Not Tested	Not Tested	Not Tested
Reactivity	+			

Entrez Gene ID 8878 (Human)

REFERENCES
1) Ichimura, Y., *et al.*, *J. Biol. Chem.* **283**, 22847-22857 (2008)
2) Komatsu, M., *et al.*, *Cell* **131**, 1149-1163 (2007)

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RELATED PRODUCTS

Antibodies

PD014	anti-LC3 (polyclonal)	[WB]
PD015	anti-LC3 (polyclonal)	[IC]
PM036	anti-LC3 (polyclonal)	[WB, IP, IC, IHC, FCM]
PM046	anti-LC3 (polyclonal)	[WB, IC]
M115-3	anti-LC3 (51-11)	[WB]
M152-3	anti-LC3 (4E12)	[WB, IP, IC, FCM]
M186-3	anti-LC3 (8E10)	[WB]
M135-3	anti-GABARAP (1F4)	
PM037	anti-GABARAP (polyclonal)	
PM038	anti-GATE-16 (polyclonal)	
PM034	anti-Atg3 (polyclonal)	
M133-3	anti-Atg3 (3E8)	
M134-3	anti-Atg4B (9H5)	
M153-3	anti-Atg5 (4D3)	
PM050	anti-Atg5 (polyclonal)	
PM039	anti-Atg7 (polyclonal)	
M151-3	anti-Atg10 (5A7)	
M154-3	anti-Atg12 (6E5)	
PD036	anti-Atg13 (polyclonal)	
M183-3	anti-Atg13 (5G4)	
PD026	anti-Atg14 (polyclonal)	
M184-3	anti-Atg14 (4H8)	
PM040	anti-Atg16L (polyclonal)	
M150-3	anti-Atg16L (1F12)	
M162-3	anti-p62/SQSTM1 (5F2)	
M162-A48	anti-p62/SQSTM1-Alexa Fluor [®] 488 (5F2)	
M162-A59	anti-p62/SQSTM1-Alexa Fluor [®] 594 (5F2)	
M162-A64	anti-p62/SQSTM1-Alexa Fluor [®] 647 (5F2)	
PM045	anti-p62/SQSTM1 (polyclonal)	
PM066	anti-p62 C-terminal (polyclonal)	
M160-3	anti-UVRAG (1H4)	
PD017	anti-Beclin 1 (polyclonal)	
PD027	anti-Rubicon (polyclonal)	
M170-3	anti-Rubicon (1H6)	
PM036-P	Positive control for anti-LC3 antibody	
M175-3	anti- α -Tubulin (2F9)	
M175-A48	anti- α -Tubulin-Alexa Fluor [®] 488 (2F9)	
M175-A59	anti- α -Tubulin-Alexa Fluor [®] 594 (2F9)	
M175-A64	anti- α -Tubulin-Alexa Fluor [®] 647 (2F9)	
PM054	anti- α -Tubulin (polyclonal)	
M176-3	anti-EEA1 (3C10)	
M176-A48	anti-EEA1-Alexa Fluor [®] 488 (3C10)	
M176-A59	anti-EEA1-Alexa Fluor [®] 594 (3C10)	
M176-A64	anti-EEA1-Alexa Fluor [®] 647 (3C10)	
PM062	anti-EEA1 (polyclonal)	
M178-3	anti-Calnexin (4F10)	
M178-A48	anti-Calnexin-Alexa Fluor [®] 488 (4F10)	
M178-A59	anti-Calnexin-Alexa Fluor [®] 594 (4F10)	
M178-A64	anti-Calnexin-Alexa Fluor [®] 647 (4F10)	
PM060	anti-Calnexin (polyclonal)	
M181-3	anti-KDEL (1D5)	
PM059	anti-KDEL (polyclonal)	
M179-3	anti-GM130 (5G8)	
M179-A48	anti-GM130-Alexa Fluor [®] 488 (5G8)	
M179-A59	anti-GM130-Alexa Fluor [®] 594 (5G8)	
M179-A64	anti-GM130-Alexa Fluor [®] 647 (5G8)	
PM061	anti-GM130 (polyclonal)	

PM063	anti-COX4 (polyclonal)
PM064	anti-Lamin B1
PM067	Normal Guinea pig IgG (polyclonal)

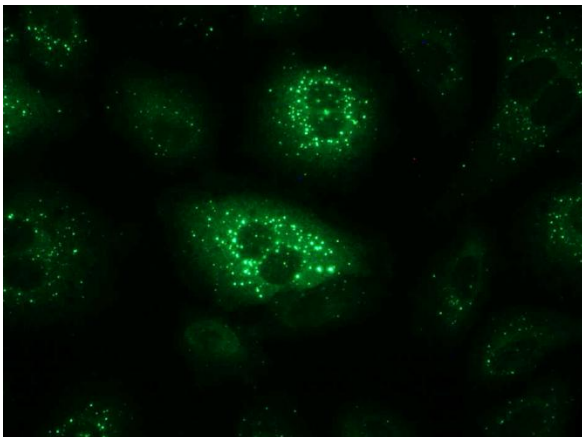
WB: Western blotting
IP: Immunoprecipitation
IC: Immunocytochemistry
IHC: Immunohistochemistry
FCM: Flow cytometry

Other related antibodies and kits are also available.
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Immunocytochemistry

- 1) Spread the cells in the nutrient condition on a glass slide, then incubate in a CO₂ incubator for one night.
- 2) Remove the culture supernatant by careful aspiration.
- 3) Fix the cells by immersing the slide in 4% paraformaldehyde (PFA)/PBS for 10 minutes at room temperature (20~25°C).
- 4) Prepare a wash container such as a 500 mL beaker with a magnetic stirrer. Then wash the fixed cells on the glass slide by soaking the slide with a plenty of PBS in the wash container for 5 minutes. Take care not to touch the cells. Repeat another wash once more.
- 5) Immerse the slide in 100 µg/mL digitonin in PBS for 10 minutes at room temperature.
- 6) Wash the slide in a plenty of PBS as in the step 4).
- 7) Add 200 µL of Clear Back (human Fc receptor blocking reagent, MBL; code no. MTG-001) onto the cells and incubate for 5 minutes at room temperature.
- 8) Add 200 µL of the primary antibody diluted with 2% fetal calf serum (FCS)/PBS as suggested in the **APPLICATIONS** onto the cells and incubate for 60 minutes at room temperature. (Optimization of antibody concentration or incubation condition is recommended if necessary.)
- 9) Wash the slide in a plenty of PBS as in the step 4).
- 10) Promptly add mounting medium onto the slide, then put a cover slip on it.

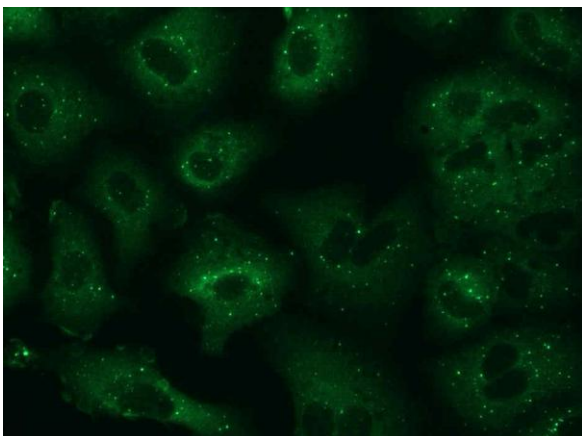
(Positive control for Immunocytochemistry; A549)



Immunocytochemical detection of p62 in A549

Upper: Starved A549

Lower: Nutrient A549



Flow cytometric analysis for adherent cells

- 1) Detach the cells from culture dish.
- 2) Wash the cells 1 time with 1 mL of washing buffer [PBS containing 2% fetal calf serum (FCS)].
- 3) Add 200 μ L of 4% paraformaldehyde (PFA) to the cell pellet after tapping. Mix well, then fix the cells for 10 minutes at room temperature.
- 4) Wash the cells 2 times with 1 mL of washing buffer.
- 5) Add 200 μ L of 100 μ g/mL digitonin in PBS to the cell pellet after tapping. Mix well, then permeabilize the cells for 10 minutes at room temperature.
- 6) Wash the cells 1 time with 1 mL of washing buffer.
- 7) Resuspend the cells with washing buffer (5×10^6 cells/mL).
- 8) Add 100 μ L of the cell suspension into each tube, and centrifuge at 500 x g for 1 minute at room temperature (20~25°C). Remove supernatant by careful aspiration.
- 9) Add 20 μ L of Clear Back (human Fc receptor blocking reagent, MBL; code no. MTG-001) to the cell pellet after tapping. Mix well and incubate for 5 minutes at room temperature.
- 10) Add 40 μ L of the primary antibody at the concentration as suggested in the **APPLICATIONS** diluted in the washing buffer. Mix well and incubate for 30 minutes at room temperature.
- 11) Add 1 mL of the washing buffer followed by centrifugation at 500 x g for 1 minute at room temperature. Remove supernatant by careful aspiration. Repeat another wash once more.
- 12) Resuspend the cells with 500 μ L of the washing buffer and analyze by a flow cytometer.

(Positive control for Flow cytometry; A549)

