

For Research Use Only

Cleaved PARP1 Monoclonal antibody

Catalog Number: 60555-1-Ig



Basic Information

Catalog Number: 60555-1-Ig	GenBank Accession Number: BC037545	Purification Method: Protein G purification
Size: 150ul , Concentration: 1000 ug/ml by Nanodrop;	GeneID (NCBI): 142	CloneNo.: 4G4C8
Source: Mouse	UNIPROT ID: P09874	Recommended Dilutions: WB 1:5000-1:50000 IHC 1:1000-1:4000 IF/ICC 1:500-1:2000
Isotype: IgG1	Full Name: poly (ADP-ribose) polymerase 1	
	Calculated MW: 1014 aa, 113 kDa	
	Observed MW: 89 kDa	

Applications

Tested Applications: WB, IHC, IF/ICC, FC (Intra), ELISA	Positive Controls:
Species Specificity: human, mouse, rat	WB : A2780 cells, HSC-T6 cells, mouse splenocytes, Staurosporine treated A2780 cells, Staurosporine treated HSC-T6 cells, Staurosporine treated mouse splenocytes
Note-IHC: suggested antigen retrieval with TE buffer pH 9.0; (*) Alternatively, antigen retrieval may be performed with citrate buffer pH 6.0	IHC : Jurkat cells,
	IF/ICC : 1 μM Staurosporine (3 hours) treated HSC-T6 cells, 1 μM Staurosporine (3 hours) treated HeLa cells

Background Information

PARP1 (poly(ADP-ribose) polymerase 1) is a nuclear enzyme catalyzing the poly(ADP-ribosyl)ation of many key proteins in vivo. The normal function of PARP1 is the routine repair of DNA damage. Activated by DNA strand breaks, the PARP1 is cleaved into an 85 to 89-kDa COOH-terminal fragment and a 24 kDa NH2-terminal peptide by caspases during the apoptotic process. The appearance of PARP fragments is commonly considered an important biomarker of apoptosis. In addition to caspases, other proteases like calpains, cathepsins, granzymes, and matrix metalloproteinases (MMPs) have also been reported to cleave PARP1 and give rise to fragments ranging from 42-89 kDa.

This antibody only recognizes the cleaved form of PARP1 but not full-length PARP1.

Storage

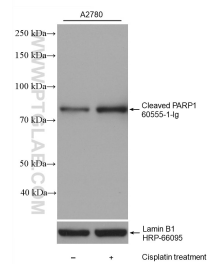
Storage:
Store at -20°C. Stable for one year after shipment.
Storage Buffer:
PBS with 0.02% sodium azide and 50% glycerol pH 7.3.
Aliquoting is unnecessary for -20°C storage

*** 20ul sizes contain 0.1% BSA

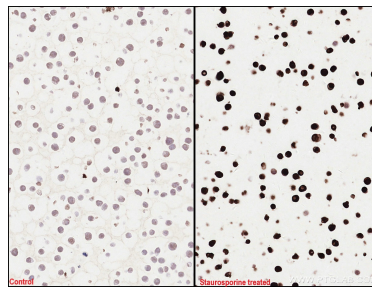
For technical support and original validation data for this product please contact:
T: 1 (888) 4PTGLAB (1-888-478-4522) (toll free in USA), or 1(312) 455-8498 (outside USA)
E: proteintech@ptglab.com
W: ptglab.com

This product is exclusively available under Proteintech Group brand and is not available to purchase from any other manufacturer.

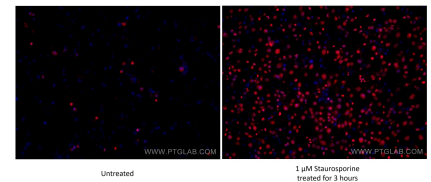
Selected Validation Data



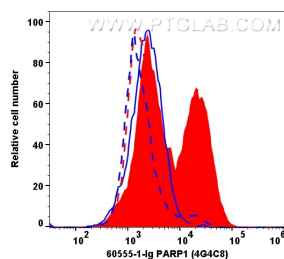
Staurosporine treated and untreated A2780 cells were subjected to SDS PAGE followed by western blot with 60555-1-Ig (Cleaved PARP1 antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with HRP-conjugated Lamin B1 (HRP-66095) antibody as a loading control.



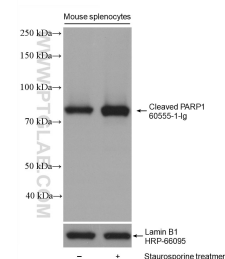
Immunohistochemical analysis of paraffin-embedded Jurkat (left) and Staurosporine treated Jurkat (right) cells slide using 60555-1-Ig (Cleaved PARP1 antibody) at dilution of 1:2000 (under 40x lens). Heat mediated antigen retrieval with Tris-EDTA buffer (pH 9.0).



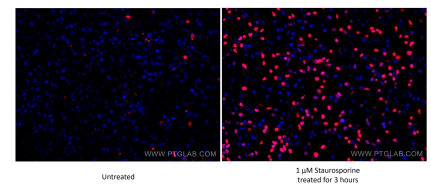
Immunofluorescent analysis of (4% PFA) fixed untreated and 1 μ M Staurosporine (3 hours) treated HSC-T6 cells using Cleaved PARP1 antibody (60555-1-Ig, Clone: 4G4C8) at dilution of 1:1000 and Multi-rAb Coralite® Plus 594-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO. RGAM004).



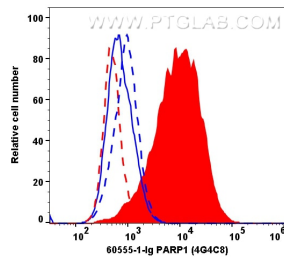
1x10⁶ HSC-T6 cell (dash lines) and 1 μ M Staurosporine (3 hours) treated HSC-T6 cells (full lines) were intracellularly stained with 0.4 μ g Cleaved PARP1 Monoclonal Antibody (60555-1-Ig, Clone:4G4C8, red) and Coralite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA.



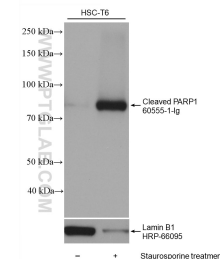
Staurosporine treated and untreated mouse splenocytes were subjected to SDS PAGE followed by western blot with 60555-1-Ig (Cleaved PARP1 antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with HRP-conjugated Lamin B1 (HRP-66095) antibody as a loading control.



Immunofluorescent analysis of (4% PFA) fixed untreated and 1 μ M Staurosporine (3 hours) treated HeLa cells using Cleaved PARP1 antibody (60555-1-Ig, Clone: 4G4C8) at dilution of 1:366 and Anti-rAb Coralite® Plus 594-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO. RGAM004).



1x10⁶ HeLa cell (dash lines) and 1 μ M Staurosporine (3 hours) treated HeLa cells (full lines) were intracellularly stained with 0.1 μ g Cleaved PARP1 Monoclonal Antibody (60555-1-Ig, Clone:4G4C8, red) and Coralite® Plus 647-Goat Anti-Mouse Recombinant Secondary Antibody (H+L) (Cat.NO.RGAM005). Mouse IgG1 isotype control (66360-1-Ig, Clone: 1F8D3, blue) was parallel stained as control. Cells were fixed with 4% PFA .



Staurosporine treated and untreated HSC-T6 cells were subjected to SDS PAGE followed by western blot with 60555-1-Ig (Cleaved PARP1 antibody) at dilution of 1:20000 incubated at room temperature for 1.5 hours. The membrane was stripped and re-blotted with HRP-conjugated Lamin B1 (HRP-66095) antibody as a loading control.