For Research Use Only

## CoraLite®594-conjugated PARP1 Monoclonal antibody



Catalog Number: CL594-66520

**Basic Information** 

Catalog Number: CL594-66520

CL594-66520 BC037545 Size: GeneID (NCBI): 1000 μ g/ml 142

Source: UNIPROT ID:
Mouse P09874

Isotype: Full Name:

IgG1 poly (ADP-ribose) polymerase 1

Immunogen Catalog Number: Calculated MW: AG19173 1014 aa, 113 kDa

Observed MW:

113-116 kDa, 85-89 kDa

GenBank Accession Number:

Purification Method:

Protein G purification CloneNo.:

1D7D4
Recommended Dilutions:
IF/ICC 1:50-1:500

Excitation/Emission maxima

wavelengths: 588 nm / 604 nm

**Applications** 

Tested Applications: IF/ICC, FC (Intra) Species Specificity:

human, mouse, rat

(Intra)

**Positive Controls:** 

IF/ICC: HeLa cells, Neuro-2a 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 as 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 gave rise to fragments ranging from 42-89-kD. This antibody was generated against the N-terminal region of human PARP1 and it recognizes the full-length as well as the cleavage of the PARP1.

Storage

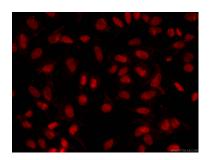
Storage:

Store at -20°C. Avoid exposure to light. Stable for one year after shipment. Storage Buffer.

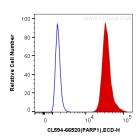
PBS with 50% Glycerol, 0.05% Proclin300, 0.5% BSA, pH 7.3.

Aliquoting is unnecessary for -20°C storage

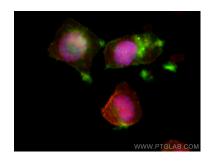
## Selected Validation Data



Immunofluorescent analysis of (4% PFA) fixed HeLa cells using CL594-66520 (PARP1 antibody) at dilution of 1:100.



1X10^6 HeLa cells were intracellularly stained with 0.4 ug CoraLite®594 Anti-Human PARP1 (CL594-66520, Clone:1D7D4) (red), or 0.4 ug Control Antibody. Cells were fixed and permeabilized with Transcription Factor Staining Buffer Kit (PF00011).



Immunofluorescent analysis of (4% PFA) fixed Neuro-2a cells using Coralite®594 PARP1 antibody (CL594-66520, Clone: 1D7D4) at dilution of 1:2000, CL488-Phalloidin (green).