Cleaved PARP (Asp214) Rabbit mAb

Catalog No.: A27147 Recombinant 1 Publications



Basic Information

Observed MW 89kDa

Calculated MW 113kDa

Category

SMab Recombinant Monoclonal Antibody

Applications WB, IF/ICC, ELISA

Cross-Reactivity Human

Background

This gene encodes a chromatin-associated enzyme, poly(ADP-ribosyl)transferase, which modifies various nuclear proteins by poly(ADP-ribosyl)ation. The modification is dependent on DNA and is involved in the regulation of various important cellular processes such as differentiation, proliferation, and tumor transformation and also in the regulation of the molecular events involved in the recovery of cell from DNA damage. In addition, this enzyme may be the site of mutation in Fanconi anemia, and may participate in the pathophysiology of type I diabetes.

Recommended Dilutions

WB	1:1000 - 1:10000
IF/ICC	1:2000 - 1:6000
ELISA	Recommended starting concentration is 1 µg/mL. Please optimize the concentration based on your specific assay requirements.

Immunogen Information

Gene ID 142

Swiss Prot P09874

Immunogen

Synthetic peptide. This information is considered to be commercially sensitive.

Synonyms

PARP; PARS; PPOL; ADPRT; ARTD1; ADPRT1; PARP-1; ADPRT 1; pADPRT-1; Poly-PARP

Contact

Product Information

G www.abclonal.com Source Rabbit

Isotype IgG

Purification Affinity purification

Storage

Store at -20°C. Avoid freeze / thaw cycles. Buffer: PBS containing 50% glycerol and 0.05% BSA, preserved with proclin300 or sodium azide (as specified on the Certificate of Analysis), pH 7.3.

Validation Data



Western blot analysis of lysates from Jurkat cells using Cleaved PARP (Asp214) Rabbit mAb (A27147) at 1:1000 dilution incubated overnight at 4°C. Jurkat cells were treated with Etoposide (25 μ M) at 37°C for 5 hours. Secondary antibody: HRP-conjugated Goat anti-Rabbit IgG (H+L) (AS014) at 1:10000 dilution. Lysates/proteins: 30 μ g per lane. Blocking buffer: 3 % nonfat dry milk in TBST. Detection: ECL Basic Kit (RM00020). Exposure time: 0.5s.



Confocal imaging of HeLa cells (treated with Staurosporine) and HeLa cells (untreated) cells using Cleaved PARP (Asp214) Rabbit mAb (A27147, dilution 1:4200) 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). Objective: 100x.