

**Cleaved PARP (Asp214) Antibody**  
**Purified Rabbit Polyclonal Antibody (Pab)**  
**Catalog # AP22119a**

## Specification

### Cleaved PARP (Asp214) Antibody - Product Information

Application	WB,E
Primary Accession	<a href="#">P09874</a>
Host	Rabbit
Clonality	polyclonal
Isotype	Rabbit Ig

### Cleaved PARP (Asp214) Antibody - Additional Information

Gene ID 142

### Target/Specificity

This Cleaved PARP (Asp214) antibody is generated from a rabbit immunized with a KLH conjugated synthetic peptide between 205-225 amino acids from the human region of human Cleaved PARP (Asp214).

### Dilution

WB~~1:500

### Format

Purified polyclonal antibody supplied in PBS with 0.09% (W/V) sodium azide. This antibody is purified through a protein A column, followed by peptide affinity purification.

### Storage

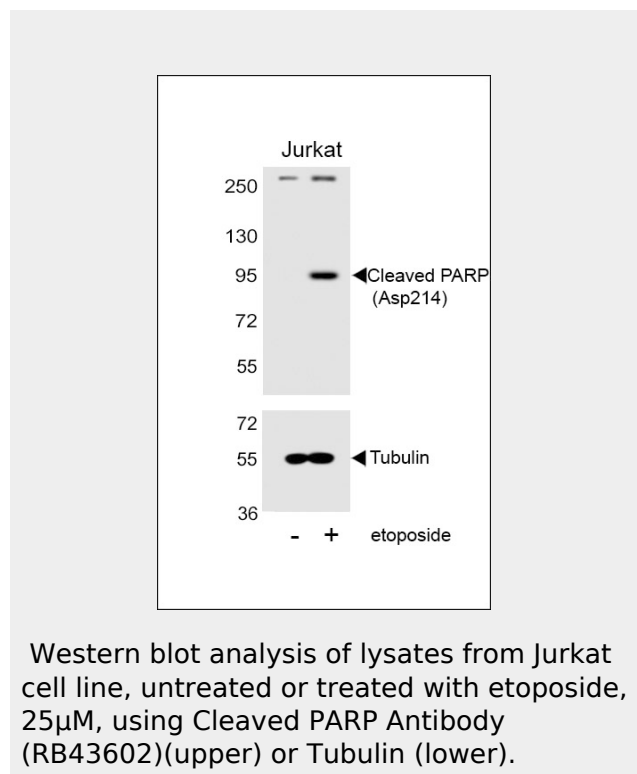
Maintain refrigerated at 2-8°C for up to 2 weeks. For long term storage store at -20°C in small aliquots to prevent freeze-thaw cycles.

### Precautions

Cleaved PARP (Asp214) Antibody is for research use only and not for use in diagnostic or therapeutic procedures.

### Cleaved PARP (Asp214) Antibody - Protein Information

Name PARP1 ([HGNC:270](#))



**Function**

Poly-ADP-ribosyltransferase that mediates poly-ADP- ribosylation of proteins and plays a key role in DNA repair (PubMed:<a href="http://www.uniprot.org/citations/17177976" target="\_blank">17177976</a>, PubMed:<a href="http://www.uniprot.org/citations/18172500" target="\_blank">18172500</a>, PubMed:<a href="http://www.uniprot.org/citations/19344625" target="\_blank">19344625</a>, PubMed:<a href="http://www.uniprot.org/citations/19661379" target="\_blank">19661379</a>, PubMed:<a href="http://www.uniprot.org/citations/23230272" target="\_blank">23230272</a>, PubMed:<a href="http://www.uniprot.org/citations/25043379" target="\_blank">25043379</a>, PubMed:<a href="http://www.uniprot.org/citations/33186521" target="\_blank">33186521</a>, PubMed:<a href="http://www.uniprot.org/citations/32028527" target="\_blank">32028527</a>, PubMed:<a href="http://www.uniprot.org/citations/26344098" target="\_blank">26344098</a>). Mediates glutamate, aspartate, serine or tyrosine ADP-ribosylation of proteins: the ADP-D-ribosyl group of NAD(+) is transferred to the acceptor carboxyl group of target residues and further ADP-ribosyl groups are transferred to the 2'-position of the terminal adenosine moiety, building up a polymer with an average chain length of 20-30 units (PubMed:<a href="http://www.uniprot.org/citations/7852410" target="\_blank">7852410</a>, PubMed:<a href="http://www.uniprot.org/citations/9315851" target="\_blank">9315851</a>, PubMed:<a href="http://www.uniprot.org/citations/19764761" target="\_blank">19764761</a>, PubMed:<a href="http://www.uniprot.org/citations/25043379" target="\_blank">25043379</a>, PubMed:<a href="http://www.uniprot.org/citations/28190768" target="\_blank">28190768</a>, PubMed:<a href="http://www.uniprot.org/citations/29954836" target="\_blank">29954836</a>). Serine ADP- ribosylation of proteins constitutes the

primary form of ADP- ribosylation of proteins in response to DNA damage (PubMed:<a href="http://www.uniprot.org/citations/33186521" target="\_blank">33186521</a>). Mainly mediates glutamate and aspartate ADP-ribosylation of target proteins in absence of HPF1 (PubMed:<a href="http://www.uniprot.org/citations/19764761" target="\_blank">19764761</a>, PubMed:<a href="http://www.uniprot.org/citations/25043379" target="\_blank">25043379</a>). Following interaction with HPF1, catalyzes serine ADP-ribosylation of target proteins; HPF1 conferring serine specificity by completing the PARP1 active site (PubMed:<a href="http://www.uniprot.org/citations/28190768" target="\_blank">28190768</a>, PubMed:<a href="http://www.uniprot.org/citations/29954836" target="\_blank">29954836</a>, PubMed:<a href="http://www.uniprot.org/citations/33186521" target="\_blank">33186521</a>, PubMed:<a href="http://www.uniprot.org/citations/32028527" target="\_blank">32028527</a>). Also catalyzes tyrosine ADP-ribosylation of target proteins following interaction with HPF1 (PubMed:<a href="http://www.uniprot.org/citations/30257210" target="\_blank">30257210</a>, PubMed:<a href="http://www.uniprot.org/citations/29954836" target="\_blank">29954836</a>). PARP1 initiates the repair of DNA breaks: recognizes and binds DNA breaks within chromatin and recruits HPF1, licensing serine ADP-ribosylation of target proteins, such as histones, thereby promoting decompaction of chromatin and the recruitment of repair factors leading to the reparation of DNA strand breaks (PubMed:<a href="http://www.uniprot.org/citations/17177976" target="\_blank">17177976</a>, PubMed:<a href="http://www.uniprot.org/citations/18172500" target="\_blank">18172500</a>, PubMed:<a href="http://www.uniprot.org/citations/19344625" target="\_blank">19344625</a>, PubMed:<a href="http://www.uniprot.org/citations/19661379" target="\_blank">19661379</a>, PubMed:<a href="http://www.uniprot.org/citations/19661379" target="\_blank">19661379</a>).

PubMed:<a href="http://www.uniprot.org/citations/23230272" target="\_blank">23230272</a>, PubMed:<a href="http://www.uniprot.org/citations/27067600" target="\_blank">27067600</a>). In addition to base excision repair (BER) pathway, also involved in double-strand breaks (DSBs) repair: together with TIMELESS, accumulates at DNA damage sites and promotes homologous recombination repair by mediating poly-ADP-ribosylation (PubMed:<a href="http://www.uniprot.org/citations/26344098" target="\_blank">26344098</a>, PubMed:<a href="http://www.uniprot.org/citations/30356214" target="\_blank">30356214</a>). Mediates the poly(ADP-ribosyl)ation of a number of proteins, including itself, APLF and CHFR (PubMed:<a href="http://www.uniprot.org/citations/17396150" target="\_blank">17396150</a>, PubMed:<a href="http://www.uniprot.org/citations/19764761" target="\_blank">19764761</a>). In addition to proteins, also able to ADP-ribosylate DNA: catalyzes ADP-ribosylation of DNA strand break termini containing terminal phosphates and a 2'-OH group in single- and double-stranded DNA, respectively (PubMed:<a href="http://www.uniprot.org/citations/27471034" target="\_blank">27471034</a>). Required for PARP9 and DTX3L recruitment to DNA damage sites (PubMed:<a href="http://www.uniprot.org/citations/23230272" target="\_blank">23230272</a>). PARP1-dependent PARP9-DTX3L-mediated ubiquitination promotes the rapid and specific recruitment of 53BP1/TP53BP1, UIMC1/RAP80, and BRCA1 to DNA damage sites (PubMed:<a href="http://www.uniprot.org/citations/23230272" target="\_blank">23230272</a>). Acts as a regulator of transcription: positively regulates the transcription of MTUS1 and negatively regulates the transcription of MTUS2/TIP150 (PubMed:<a href="http://www.uniprot.org/citations/19344625" target="\_blank">19344625</a>). Plays a role in the positive regulation of IFNG transcription in T-helper 1 cells as part of an IFNG promoter-binding complex with TXK and EEF1A1 (PubMed:<a href="http://www.uniprot.org/citations/17177976" target="\_blank">17177976</a>

target="\_blank">17177976</a>). Involved in the synthesis of ATP in the nucleus, together with NMNAT1, PARG and NUDT5 (PubMed:<a href="http://www.uniprot.org/citations/27257257"

target="\_blank">27257257</a>). Nuclear ATP generation is required for extensive chromatin remodeling events that are energy-consuming (PubMed:<a href="http://www.uniprot.org/citations/27257257" target="\_blank">27257257</a>).

#### **Cellular Location**

Nucleus. Nucleus, nucleolus. Chromosome  
Note=Localizes to sites of DNA damage.

#### **Cleaved PARP (Asp214) Antibody - Protocols**

Provided below are standard protocols that you may find useful for product applications.

- [Western Blot](#)
- [Blocking Peptides](#)
- [Dot Blot](#)
- [Immunohistochemistry](#)
- [Immunofluorescence](#)
- [Immunoprecipitation](#)
- [Flow Cytometry](#)
- [Cell Culture](#)

#### **Cleaved PARP (Asp214) Antibody - Citations**

- [Isochamaejasmin induces toxic effects on Helicoverpa zea via DNA damage and mitochondria-associated apoptosis.](#)
- [PSMD7 downregulation induces apoptosis and suppresses tumorigenesis of esophageal squamous cell carcinoma the mTOR/p70S6K pathway.](#)