R&D SYSTEMS
a bio-techne brand

Human/Mouse PARP Antibody

Antigen Affinity-purified Polyclonal Goat IgG
Catalog Number: AF-600-NA

DESCRIPTION

Species Reactivity	Human/Mouse
Specificity	Detects human and mouse PARP in Western blots.
Source	Polyclonal Goat IgG
Purification	Antigen Affinity-purified
Immunogen	*E. coli*-derived recombinant mouse PARP Val71-Pro329 Accession # NP_031441
Formulation	Lyophilized from a 0.2 μm filtered solution in PBS with Trehalose. See Certificate of Analysis for details. *Small pack size (-SP) is supplied as a 0.2 μm filtered solution in PBS.

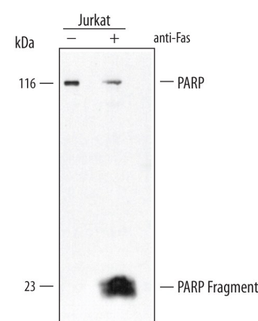
APPLICATIONS

Please Note: *Optimal dilutions should be determined by each laboratory for each application. General Protocols are available in the Technical Information section on our website.*

	Recommended Concentration	Sample
Western Blot	0.4 μg/mL	See Below
Immunocytochemistry	1-25 μg/mL	See Below
Immunoprecipitation	5 μg/10^6 cells	See Below
Simple Western	5 μg/mL	See Below

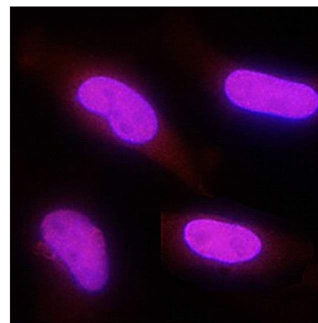
DATA

Western Blot



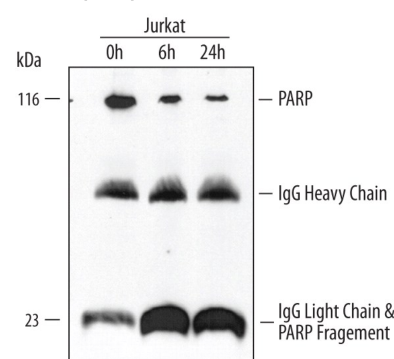
Detection of Human PARP by Western Blot. Western blot shows lysates of Jurkat human acute T cell leukemia cell line untreated (-) or treated (+) with 200 ng/mL anti-Fas for 24 hours. PVDF membrane was probed with 0.4 µg/mL of Goat Anti-Human/Mouse PARP Affinity-purified Polyclonal Antibody (Catalog # AF-600-NA) followed by HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF109). A specific band was detected for PARP at approximately 116 kDa (as indicated). This experiment was conducted under reducing conditions and using Immunoblot Buffer Group 2.

Immunocytochemistry



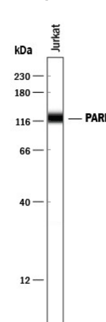
PARP in HeLa Human Cell Line. PARP was detected in immersion fixed HeLa human cervical epithelial carcinoma cell line using Goat Anti-Human/Mouse PARP Antigen Affinity-purified Polyclonal Antibody (Catalog # AF-600-NA) at 1 µg/mL for 3 hours at room temperature. Cells were stained using the NorthernLights™ 557-conjugated Anti-Goat IgG Secondary Antibody (red; Catalog # NL001) and counterstained with DAPI (blue). Specific staining was localized to nuclei. View our protocol for Fluorescent ICC Staining of Cells on Coverslips.

Immunoprecipitation



Immunoprecipitation of Human PARP. Jurkat human acute T cell leukemia cell line was treated with apoptosis inducer anti-Fas for the indicated times. PARP was immunoprecipitated from cell lysates (1 - 2 x 10^6 cells) following incubation with 5 µg Goat Anti-Human/Mouse PARP Antigen Affinity-purified Polyclonal Antibody (Catalog # AF-600-NA) for overnight at 4 °C. PARP-antibody complexes were absorbed using Protein G expressing Staph cells (Sigma). Immunoprecipitated PARP was detected by Western blot using 0.4 µg/mL Goat Anti-Human/Mouse PARP Antigen Affinity-purified Polyclonal Antibody (Catalog # AF-600-NA). View our recommended buffer recipes for immunoprecipitation.

Simple Western



Detection of Human PARP by Simple Western™. Simple Western lane view shows lysates of Jurkat human acute T cell leukemia cell line, loaded at 0.2 mg/mL. A specific band was detected for PARP at approximately 122 kDa (as indicated) using 5 µg/mL of Goat Anti-Human/Mouse PARP Antigen Affinity-purified Polyclonal Antibody (Catalog # AF-600-NA) followed by 1:50 dilution of HRP-conjugated Anti-Goat IgG Secondary Antibody (Catalog # HAF109). This experiment was conducted under reducing conditions and using the 12-230 kDa separation system.

bio-techne
bio-techne.com
Global info@bio-techne.com techsupport@bio-techne.com
North America TEL 800 343 7475 Europe | Middle East | Africa TEL +44 (0)1235 529449
China info.cn@bio-techne.com TEL +86 (21) 52380373
Rest of World bio-techne.com/find-us/distributors TEL +1 612 379 2956

PREPARATION AND STORAGE

Reconstitution	Reconstitute at 0.2 mg/mL in sterile PBS.
Shipping	The product is shipped at ambient temperature. Upon receipt, store it immediately at the temperature recommended below. *Small pack size (-SP) is shipped with polar packs. Upon receipt, store it immediately at -20 to -70 °C
Stability & Storage	**Use a manual defrost freezer and avoid repeated freeze-thaw cycles.** ● 12 months from date of receipt, -20 to -70 °C as supplied. ● 1 month, 2 to 8 °C under sterile conditions after reconstitution. ● 6 months, -20 to -70 °C under sterile conditions after reconstitution.

BACKGROUND

PARP, Poly [ADP-ribose] polymerase 1 (PARP1), is a component of a base excision repair (BER) complex, containing at least XRCC1, PARP2, POLB and LRIG3. Widely expressed. Expression is correlated with proliferation, with higher levels occurring during early fetal development and organogenesis and in the highly proliferative cell compartments of adult. Expressed in B-cells that have been induced to switch to various Ig isotypes. PARP interacts with the DNA polymerase alpha catalytic subunit POLA1; this interaction functions as part of the control of replication fork progression.

bio-techne
bio-techne.com
Global info@bio-techne.com techsupport@bio-techne.com
North America TEL 800 343 7475 **Europe | Middle East | Africa** TEL +44 (0)1235 529449
China info.cn@bio-techne.com TEL +86 (21) 52380373
Rest of World bio-techne.com/find-us/distributors TEL +1 612 379 2956