

PARP-8 (C-17): sc-82985

BACKGROUND

Poly(ADP-ribosylation) is a method of DNA damage-dependent posttranslational modification that helps to rescue injured proliferating cells from cell death. The PARP (poly(ADP-ribose) polymerase) proteins comprise a superfamily of enzymes that functionally modify histones and other nuclear proteins, thereby preventing cell death. PARPs use NAD⁺ as a substrate to catalytically transfer ADP-ribose residues onto protein acceptors; a process that, when repeated multiple times, leads to the formation of poly(ADP-ribose) chains on the protein. The presence of these chains alters the function of the target protein and promotes cell survival. PARP proteins are implicated in a variety of diseases, including cancer, neurodegenerative and inflammatory disorders. PARP-8 (poly(ADP-ribose) polymerase family, member 8), also designated pART16, is a 854 amino acid protein containing a single PARP catalytic domain and may become phosphorylated upon DNA damage by ATM or ATR. PARP-8 exists as two alternatively spliced isoforms.

REFERENCES

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- Aguiar, R.C., Takeyama, K., He, C., Kreinbrink, K. and Shipp, M.A. 2005. B-aggressive lymphoma family proteins have unique domains that modulate transcription and exhibit poly(ADP-ribose) polymerase activity. *J. Biol. Chem.* 280: 33756-33765.
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- Elser, M., Borsig, L., Hassa, P.O., Erener, S., Messner, S., Valovka, T., Keller, S., Gassmann, M. and Hottiger, M.O. 2008. Poly(ADP-ribose) polymerase 1 promotes tumor cell survival by coactivating hypoxia-inducible factor-1-dependent gene expression. *Mol. Cancer Res.* 6: 282-290.
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CHROMOSOMAL LOCATION

Genetic locus: PARP8 (human) mapping to 5q11.1; Parp8 (mouse) mapping to 13 D2.3.

SOURCE

PARP-8 (C-17) is an affinity purified goat polyclonal antibody raised against a peptide mapping near the C-terminus of PARP-8 of human origin.

STORAGE

Store at 4° C, ****DO NOT FREEZE****. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

PRODUCT

Each vial contains 200 µg IgG in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

Blocking peptide available for competition studies, sc-82985 P, (100 µg peptide in 0.5 ml PBS containing < 0.1% sodium azide and 0.2% BSA).

APPLICATIONS

PARP-8 (C-17) is recommended for detection of PARP-8 of mouse, rat and human origin by Western Blotting (starting dilution 1:200, dilution range 1:100-1:1000), immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000); non cross-reactive with other PARP family members.

PARP-8 (C-17) is also recommended for detection of PARP-8 in additional species, including equine, canine, bovine, porcine and avian.

Suitable for use as control antibody for PARP-8 siRNA (h): sc-76066, PARP-8 siRNA (m): sc-76067, PARP-8 shRNA Plasmid (h): sc-76066-SH, PARP-8 shRNA Plasmid (m): sc-76067-SH, PARP-8 shRNA (h) Lentiviral Particles: sc-76066-V and PARP-8 shRNA (m) Lentiviral Particles: sc-76067-V.

PARP-8 (C-17) X TransCruz antibody is recommended for Gel Supershift and ChIP applications.

Molecular Weight of PARP-8: 96 kDa.

RECOMMENDED SECONDARY REAGENTS

To ensure optimal results, the following support (secondary) reagents are recommended: 1) Western Blotting: use donkey anti-goat IgG-HRP: sc-2020 (dilution range: 1:2000-1:100,000) or Cruz Marker™ compatible donkey anti-goat IgG-HRP: sc-2033 (dilution range: 1:2000-1:5000), Cruz Marker™ Molecular Weight Standards: sc-2035, TBS Blotto A Blocking Reagent: sc-2333 and Western Blotting Luminol Reagent: sc-2048. 2) Immunofluorescence: use donkey anti-goat IgG-FITC: sc-2024 (dilution range: 1:100-1:400) or donkey anti-goat IgG-TR: sc-2783 (dilution range: 1:100-1:400) with UltraCruz™ Mounting Medium: sc-24941.

RESEARCH USE

For research use only, not for use in diagnostic procedures.

PROTOCOLS

See our web site at www.scbt.com or our catalog for detailed protocols and support products.