



Rabbit Anti-Human Cleaved PARP-1 (Asp214) Monoclonal Antibody (Clone SP276)

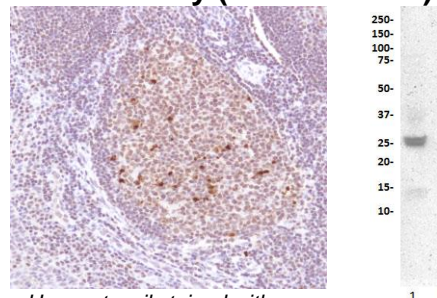
CATALOG #:

M5760 0.1 ml rabbit monoclonal antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.

M5762 0.5 ml rabbit monoclonal antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.

M5764 1.0 ml rabbit monoclonal antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.

M5761 7.0 ml pre-diluted rabbit monoclonal antibody purified by protein A/G in TBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.



Human tonsil stained with anti-cleaved PARP-1 (SP276) antibody

Western Blot analysis of NIH3T3 cell lysate (lane 1 cell treated with staurosporine and lane 2 untreated cell) with anti-cleaved PARP-1 antibody (SP276)

INTENDED USE:

For Research Use Only. Not for use in diagnostic procedures.

CLONE:

SP276

IMMUNOGEN:

Synthetic peptide surrounding Asp 214 of human PARP-1 protein.

IG ISOTYPE:

Rabbit IgG

EPITOPE:

Not determined

MOLECULAR WEIGHT:

24 kDa

SPECIES REACTIVITY:

Human (tested). (See www.springbio.com for information on species reactivity predicted by sequence homology.)

DESCRIPTION:

Poly [ADP-ribose] polymerase 1 (PARP-1) is a nuclear DNA binding protein for repairing single-strand DNA break and for maintaining cell viabilities. Human PARP are cleaved by some caspases such as caspase-3 and caspase-7 between Asp214 and Gly215 during apoptosis and can be used as an apoptosis marker.

APPLICATIONS:

Immunohistochemistry (IHC) and Western Blotting

IHC POSITIVE CONTROL:

Tonsil, NIH3T3 cells treated with staurosporine

IHC PROCEDURE:

Specimen Preparation: Formalin-fixed, paraffin-embedded tissues are suitable for use with this primary antibody.

Deparaffinization: Deparaffinize slides using xylene or xylene alternative and graded alcohols.

Antibody Dilution: If using the concentrate format of this product, dilute the antibody 1:100. The dilutions are estimates; actual results may differ because of variability in methods and protocols.

Antigen Retrieval: Boil tissue section in 10 mM citrate buffer, pH 6.0 for 10 min followed by cooling at room temperature for 20 min.

Primary Antibody Incubation: Incubate for 10 minutes at room temperature.

Slide Washing: Slides must be washed in between steps. Rinse slides with PBS/0.05% Tween.

Visualization: Detect the antibody as instructed by the instructions provided with the visualization system.

WESTERN BLOTTING:

Recommended starting protocol: Dilute the antibody 1:100. Incubate for 1 hour at room temperature.

The dilution is an estimate; actual results may differ because of variability in methods and protocols. Optimal dilution and procedure should be determined by the end user.

WESTERN BLOTTING

POSITIVE CONTROL:

NIH3T3 cells treated with staurosporine

CELLULAR LOCALIZATION:

Nucleus

STORAGE & STABILITY:

Store at 2-8°C. Do not freeze. The user must validate any other storage conditions. When properly stored, the reagent is stable to the date indicated on the label. Do not use the reagent beyond the expiration date.

There are no definitive signs to indicate instability of this product; therefore, positive and negative controls should be tested simultaneously with unknown specimens.

If unexpected results are observed which cannot be explained by variations in laboratory procedures and a problem with the reagent is suspected, contact Technical Support at spring.tech@roche.com.

**WARNINGS &
PRECAUTIONS:**

1. Avoid contact of reagents with eyes and mucous membranes. If reagents come into contact with sensitive areas, wash with copious amounts of water.
2. This product is harmful if swallowed.
3. Consult local or state authorities with regard to recommended method of disposal.
4. Avoid microbial contamination of reagents.