

## Rabbit Anti-Human Bax Monoclonal Antibody (Clone SP47)

M3470 0.1 ml rabbit monoclonal CATALOG #:

antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1%

sodium azide.

0.5 ml rabbit monoclonal M3472

antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1%

sodium azide.

M3474 1.0 ml rabbit monoclonal

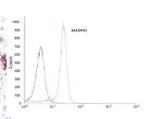
antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1 sodium azide.

7.0 ml pre-diluted rabbit M3471

> monoclonal antibody purified by protein A/G in TBS/1% BSA buffer pH 7.6 with less than 0.1% sodium azide.

Human normal breast stained with anti-Bax

antibody



Flow cytometric analysis of rabbit anti-BAX (SP47) antibody in HeLa (green) compare to negative control of rabbit IgG (blue)

INTENDED USE: For Research Use Only. Not for use in diagnostic procedures.

CLONE:

**IMMUNOGEN:** Synthetic peptide corresponding to internal region of human Bax protein.

Rabbit IgG IG ISOTYPE: Not determined EPITOPE: **MOLECULAR WEIGHT** 21 kDa

**SPECIES REACTIVITY:** Human (tested). (See www.springbio.com for information on species reactivity predicted by

sequence homology.)

**DESCRIPTION:** Bax is a member of the Bcl-2 family of proteins, which function as regulators of apoptosis.

Overexpression of Bax functions to promote cell death. Bax can form homodimers and is also able

to heterodimerize with other Bcl-2 related proteins.

APPLICATIONS: Immunohistochemistry (IHC) and Flow Cytometry

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 Dilutions: If using the concentrate format of this product, dilute the antibody 1:50 prior to use. The dilutions are estimates; actual results may differ because of variability in methods and protocols.

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

**Primary Antibody:** Incubate for 30 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.

**POSITIVE CONTROL:** 

FLOW CYTOMETRY: Recommended starting protocol: Dilute the antibody 1:100. Incubate for 30 minutes at 4°C.

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.

FLOW CYTOMETRY

HeLa Cell Line POSITIVE CONTROL: CELLULAR LOCALIZATION: Cytoplasm, membrane

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## **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@ventana.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.

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