

Rabbit Anti-Human CD20 Monoclonal Antibody (Clone SP32)

CATALOG #:

M3320 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.

M3322 0.5 m

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.

M3324

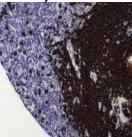
1.0 ml rabbit monoclonal antibody purified by protein A/G in PBS/1% BSA buffer pH 7.6 with less than 0.1%

sodium azide.

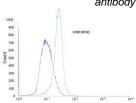
M3321

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. (For IHC

only)



Human B cell lymphoma stained with anti-CD20 antibody



Western Blot analysis of Ramos cell lysate with CD20 antibody

Tel: 1-925-474-8440

Toll Free: 1-800-787-6896 Fax: 1-925-474-8469

Flow cytometric analysis of rabbit anti-CD20 (SP32) antibody in Jurkat (green) compare to negative control of rabbit IgG (blue)

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

CLONE: SP32

IMMUNOGEN: Synthetic peptide from C-terminus of human CD20 protein.

IG ISOTYPE: Rabbit IgG
EPITOPE: Not determined

MOLECULAR WEIGHT 35kDa

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

sequence homology.)

DESCRIPTION: CD20 is a 35kDa leukocyte surface antigen consisting of four transmembrane regions and

cytoplasmic N- and C-termini. CD20 is expressed primarily on B cells but has also been detected on both normal and neoplastic T cells. Activation of CD20 significantly increases DNA synthesis and is

thought to involve basic helix-loop-helix leucine zipper transcription factors.

APPLICATIONS: Immunohistochemistry (IHC), Western Blotting 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 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 10mM citrate buffer, pH 6.0 for 10 min followed by cooling

at room temperature for 20 min.

Primary Antibody Incubation: 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.

IHC POSITIVE CONTROL: Tonsil, B cell lymphoma

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:

Ramos cell lysate

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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 POSITIVE CONTROL:

Jurkat cell line Membrane

STORAGE & STABILITY

CELLULAR LOCALIZATION:

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:

 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.