

Mouse anti CD20 Monoclonal Antibody

Alternative Name(s): MS4A1, CD20, CD20 antigen, CD20 receptor, B-Lymphocyte cell-surface antigen B1

Order Information

Description: CD20
Catalogue: 500-11214
Lot: See label
Size: 100ug/200ul
Host: Mouse
Clone: C273

• Application: IHC(P), WB

• Reactivity: Hu

ANTIGEN PREPARATION

A synthetic peptide corresponding to C-terminus of human CD20 protein.

BACKGROUND

CD20, a non-glycosylated protein, belongs to a member of the membrane-spanning 4A gene family. The molecular weight is at a range of ~34-37kDa depending on the degree of the protein phosphorylation. The CD20 antigen is present on human pre B lymphocytes and B lymphocytes at all stages of maturation. Thus, it plays an important role in the development and differentiation of B-cells into plasma cells. Low level of CD20 expression appears on normal T lymphocytes. Combination of chemo and humanized CD20 antibodies has been reported in clinical study of lymphoma diseases.

PURIFICATION

The mouse IgG is purified by Protein A-Affinity Chromatography according to Isotyping

FORMULATION

This affinity purified antibody is supplied in sterile Phosphatebuffered saline (pH7.2) containing antibody stabilizer

SPECIFICITY

This antibody recognizes ~34 kDa of human CD20. The other species are not tested

STORAGE

The antibodies are stable for 24 months from date of receipt when stored at -20oC to -70oC. The antibodies can be stored at 2oC-8oC for three month without detectable loss of activity. Avoid repeated freezing-thawing cycles.

APPLICATIONS/SUGGESTED WORKING DILUTIONS*

• Western Blot: 0.1-1 μg/ml

• ELISA: 0.01-0.1 µg/ml

• Immunoprecipitation: 2-5 µg/ml

• IHC: 2-10 µg/ml

· Flow cytometry: Not tested

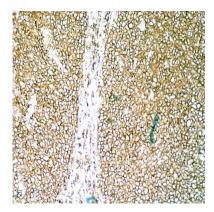
• Molecular Weight: 35.0

• Positive Control: Kidney Tissue

• Cellular Location: Cell Membrane

^{*}Optimal dilutions should be determined by researchers for the specific applications.





Immunohistochemistry: Human Tonsil (FFPE) stained with Mouse anti-CD20 (Cat# 500-11214) at 1:200 for 10 min @ RT. Staining of formalin-fixed tissue requires boiling tissue sections in 10 mM Citrate Buffer, pH 6.0 for 10 min followed by cooling at RT for 20 min.

REFERENCES