

Rabbit anti Retinoblastoma Polyclonal Antibody

Alternative Name(s): retinoblastoma 1; RB

Order Information

- Description: Retinoblastoma (RB)
- Catalogue: 500-4314
- Lot: See label
- Size: 100ug/200ul
- Host: Rabbit
- Clone: nan
- Application: IHC(P), WB
- Reactivity: Hu, Ms, Rt

ANTIGEN PREPARATION

A synthetic peptide corrsponding to the internal segment of human retinoblastoma RB

BACKGROUND

RB, retinoblastoma, is a negative regulator of the cell cycle and was the first tumor suppressor gene found. It is a key regulator of the cell cycle during the transition from the G1 to S phases. RB also stabilizes constitutive heterochromatin to maintain the overall chromatin structure. The active, hypophosphorylated form of the protein binds transcription factor E2F1. Defects in this gene are a cause of childhood cancer retinoblastoma (RB), bladder cancer, and osteogenic sarcoma.

PURIFICATION

The Rabbit IgG is purified by Epitope Affinity Purification

FORMULATION

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

SPECIFICITY

This antbody recognizes Retinoblastoma (RB) protein. It reacts to human, mice and rat. The other species not tested.

STORAGE

The antibodies are stable for 24 months from date of receipt when stored at -200C to -700C. The antibodies can be stored at 20C-80C for three month without detectable loss of activity. Avoid repeated freezing-thawing cycles.

APPLICATIONS/SUGGESTED WORKING DILUTIONS*

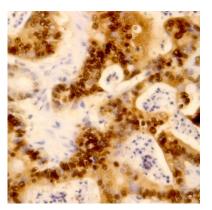
- Western Blot: 0.1-1 µg/ml
- ELISA: 0.01-0.1 µg/ml
- Immunoprecipitation: 2-5 µg/ml
- IHC: 2-10 µg/ml
- Flow cytometry: Not tested
- Molecular Weight: 110.0
- Positive Control: Kidney Tissue
- Cellular Location: Cell Membrane

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

FOR RESEARCH USE ONLY.

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Immunohistochemistry: Human Breast carcinoma (FFPE) stained with Rabbit anti-Retinoblastoma (SB) (Cat# 500-4314) 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