

Mouse anti Melan-A Monoclonal Antibody

Alternative Name(s): MART-1

Order Information

• Description: Melan-A/MART-1

• Catalogue: 603-920 • Lot: See label • Size: 100ug/200ul • Host: Mouse • Clone: A103 • Application: IHC(P)

• Reactivity: Hu

ANTIGEN PREPARATION

A synthetic peptide from C-term of human MART-1 protein

BACKGROUND

Melanocyte lineage-specific protein (MART-1; melanoma antigen recognized by T cells 1) is a widely shared melanoma antigen recognized by the T lymphocytes of patients with established malignancy. This 18 kD transmembrane protein is expressed in most melanoma tumor samples and, among normal cells in melanocytes. It is a useful marker for melanocytic tumors however it is normally found in benign nevi as well. Antibodies against the antigen are used to recognize cells of melanocytic differentiation. MART-1 antibody is a superior immunohistochemical marker for the diagnosis of malignant melanoma.

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 human Melan-A/MART-1 protein. The other species are not tested.

STORAGE

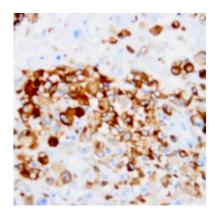
The antibodies are stable for 24 months from date of receipt when stored at -20oC to -70oC. The antibodies can be stored at 2oC-8oC for three month without detectable loss of activity. Avoid repeated freezing-thawing cycles.

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: 18.0
- Positive Control: Kidney Tissue
- Cellular Location: Cell Membrane

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





Immunohistochemistry: Human Melanoma (FFPE) stained with Mouse anti-Melan A (Cat# 603-920) 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