

**Mouse Anti-MART-1/Melan A [A103]: MC0189, MC0189RTU7**

**Intended Use:** For Research Use Only

**Description:** MART-1, also known as Melan-A, is a melanocyte lineage-specific protein (MART-1; melanoma antigen recognized by T cells 1) recognized by the T lymphocytes of patients with established malignancy. MART-1 labels both normal melanocyte and diseased cell with melanocyte differentiation. It is useful for diagnosis of tumors with melanocyte differentiation, especially metastatic melanoma. Identification of MART-1 also opens possibilities for the development of immunotherapies for patients with melanoma.

**Specifications**

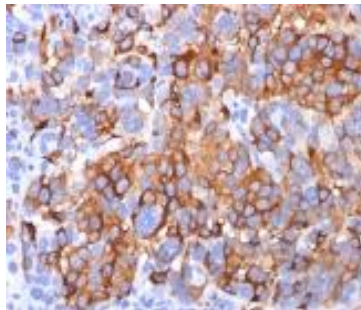
Clone:	A103
Source:	Mouse
Isotype:	IgG1k
Reactivity:	Human, mouse, rat, dog
Immunogen:	Recombinant human MART-1 protein
Localization:	Cytoplasm
Formulation:	Antibody in PBS pH7.4, containing BSA and $\leq 0.09\%$ sodium azide (NaN <sub>3</sub> )
Storage:	Store at 2°- 8°C
Applications:	IHC, Flow Cyt., IF, WB
Package:	

Description	Catalog No.	Size
MART-1/Melan A Concentrated	MC0189	1 ml
MART-1/Melan A Prediluted	MC0189RTU7	7 ml

**IHC Procedure\***

Positive Control Tissue:	Skin, melanoma
Concentrated Dilution:	50-200
Pretreatment:	Tris EDTA pH9.0, 15 minutes Pressure Cooker or 30-60 minutes water bath at 95°-99°C
Incubation Time and Temp:	30-60 minutes @ RT
Detection:	Refer to the detection system manual

\* Result should be confirmed by an established diagnostic procedure.



FFPE human Melanoma stained with MART-1/Melan-A using DAB

**References:**

1. Reconstitution of full-thickness skin by microcolumn grafting. Tam J, et al. J Tissue Eng Regen Med N/A:N/A, 2016.
2. Quantitative measurement of melanoma spread in sentinel lymph nodes and survival. Ulmer A, et al. Med 11:e1001604, 2014.
3. Localisation of epithelial cells capable of holoclone formation in vitro and direct interaction with stromal cells in the native human limbal crypt. Dziasko MA, et al. PLoS One 9:e94283, 2014.
4. Direct chemosensitivity monitoring ex vivo on undissociated melanoma tumor tissue by impedance spectroscopy. Jahnke HG, et al. Cancer Res 74:6408-18, 2014.