# SANTA CRUZ BIOTECHNOLOGY, INC.

# MICA/B (F-6): sc-137242



## BACKGROUND

MICA and MICB are stress-induced antigens that are related to major histocompatibility complex (MHC) class I molecules. MICA and MICB are frequently expressed in epithelial tumors. These highly glycosylated cell surface proteins are stably expressed without conventional class I peptide ligands or association with  $\beta$ -2-Microglobulin. The expression is induced on proliferating or heat shock-stressed epithelial cells. MICA and MICB are broadly recognized by intestinal epithelial V $\delta$ 1  $\gamma\delta$ T cells expressing variable TCRs, suggesting that these antigens may play a central role in the signaling of cellular distress to evoke immune responses in the intestinal epithelium.

#### **CHROMOSOMAL LOCATION**

Genetic locus: MICA/MICB (human) mapping to 6p21.33.

## SOURCE

MICA/B (F-6) is a mouse monoclonal antibody raised against amino acids 1-300 mapping at the N-terminus of MICA of human origin.

## PRODUCT

Each vial contains 200  $\mu g$  IgG\_1 kappa light chain in 1.0 ml of PBS with < 0.1% sodium azide and 0.1% gelatin.

MICA/B (F-6) is available conjugated to agarose (sc-137242 AC), 500 µg/ 0.25 ml agarose in 1 ml, for IP; to HRP (sc-137242 HRP), 200 µg/ml, for WB, IHC(P) and ELISA; to either phycoerythrin (sc-137242 PE), fluorescein (sc-137242 FITC), Alexa Fluor<sup>®</sup> 488 (sc-137242 AF488), Alexa Fluor<sup>®</sup> 546 (sc-137242 AF546), Alexa Fluor<sup>®</sup> 594 (sc-137242 AF594) or Alexa Fluor<sup>®</sup> 647 (sc-137242 AF647), 200 µg/ml, for WB (RGB), IF, IHC(P) and FCM; and to either Alexa Fluor<sup>®</sup> 680 (sc-137242 AF680) or Alexa Fluor<sup>®</sup> 790 (sc-137242 AF790), 200 µg/ml, for Near-Infrared (NIR) WB, IF and FCM.

# **APPLICATIONS**

MICA/B (F-6) is recommended for detection of MICA and MICB of human origin by Western Blotting (starting dilution 1:100, dilution range 1:100-1:1000), immunoprecipitation [1-2  $\mu$ g per 100-500  $\mu$ g of total protein (1 ml of cell lysate)], immunofluorescence (starting dilution 1:50, dilution range 1:50-1:500), immunohistochemistry (including paraffin-embedded sections) (starting dilution 1:50, dilution range 1:50-1:500) and solid phase ELISA (starting dilution 1:30, dilution range 1:30-1:3000).

Suitable for use as control antibody for MICA/B siRNA (h): sc-43931, MICA/B shRNA Plasmid (h): sc-43931-SH and MICA/B shRNA (h) Lentiviral Particles: sc-43931-V.

Molecular Weight of truncated MICA/B: 38 kDa.

Molecular Weight of glycosylated MICA/B: 62 kDa.

Positive Controls: MICA (h2): 293T Lysate: sc-113460, U-87 MG cell lysate: sc-2411 or Jurkat whole cell lysate: sc-2204.

## **STORAGE**

Store at 4° C, \*\*DO NOT FREEZE\*\*. Stable for one year from the date of shipment. Non-hazardous. No MSDS required.

## **RESEARCH USE**

For research use only, not for use in diagnostic procedures.

## DATA





of formalin fixed, paraffin-embedded human esophagus

tissue showing cytoplasmic staining of squamous

MICA/B (F-6): sc-137242. Western blot analysis of MICA expression in non-transfected: sc-117752 (A) and human MICA transfected: sc-113460 (B) 293T whole cell lysates.

# SELECT PRODUCT CITATIONS

 Del Toro-Arreola, S., et al. 2011. MHC class I-related chain A and B ligands are differentially expressed in human cervical cancer cell lines. Cancer Cell Int. 11: 15.

epithelial cells

- Tsukagoshi, M., et al. 2016. Overexpression of natural killer group 2 member D ligands predicts favorable prognosis in cholangiocarcinoma. Cancer Sci. 107: 116-122.
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- 4. Murakami, T., et al. 2017. Neoadjuvant chemoradiotherapy of pancreatic cancer induces a favorable immunogenic tumor microenvironment associated with increased major histocompatibility complex class I-related chain A/B expression. J. Surg. Oncol. 116: 416-426.
- 5. Ou, Z.L., et al. 2019. Hypoxia-induced shedding of MICA and HIF1Amediated immune escape of pancreatic cancer cells from NK cells: role of circ\_0000977/miR-153 axis. RNA Biol. 16: 1592-1603.
- Ding, H., et al. 2021. MICA-G129R: a bifunctional fusion protein increases PRLR-positive breast cancer cell death in co-culture with natural killer cells. PLoS ONE 16: e0252662.
- Sasagawa, S., et al. 2024. Improvement of histone deacetylase inhibitor efficacy by SN38 through TWIST1 suppression in synovial sarcoma. Cancer Innov. 3: e113.
- Kim, J.E., et al. 2025. Isoxazole-based molecules restore NK cell immune surveillance in hepatocarcinogenesis by targeting TM4SF5 and SLAMF7 linkage. Signal Transduct. Target. Ther. 10: 15.

## **PROTOCOLS**

See our web site at www.scbt.com for detailed protocols and support products.

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