

Myeloid-Associated Differentiation Marker (MYADM); Clone BM-2 (Concentrate)

Availability/Contents:

<u>Item #</u>	<u>Volume</u>
RA0366-C.5	0.5 ml

Description:

Species: Mouse

Immunogen: Nuclei from pokeweed mitogen stimulated human peripheral blood lymphocytes

Clone: BM-2

Isotype: IgG1

Entrez Gene ID: 91663 (Human)

Hu Chromosome Loc.: 19q13.42

Synonyms: MYADM; myeloid associated differentiation marker; Myeloid upregulated protein; Protein SB135

Mol. Weight of Antigen: 35kDa

Format: 200µg/ml of Ab purified from Bioreactor Concentrate by Protein A/G. Prepared in 10mM PBS with 0.05% BSA & 0.05% azide.

Specificity: Recognizes a myeloid associated differentiation antigen in the cytoplasm of mature granulocytes. It shows no reactivity with any other cell type in human tissues.

Background: Markers of myeloid cells are useful in the identification of different levels of cellular differentiation. It reacts with early precursor and mature forms of human and monkey myeloid cells. This antibody is useful for the detection of myeloid leukemias and granulocytic sarcomas. It can be used as a marker of granulocytes in normal tissues or inflammatory processes.

Species Reactivity: Human and Macaque Monkey. Others not known.

Positive Control: HL60 cells. Tonsil or lymph node.

Cellular Localization: Cytoplasmic

Titer/ Working Dilution: Immunohistochemistry (Frozen and Formalin-fixed): 0.5-1 µg/ml
Flow Cytometry: 0.5-1 µg/million cells
Immunofluorescence: 1-2 µg/ml

Microbiological State: This product is not sterile.

Storage: 2° C  8° C

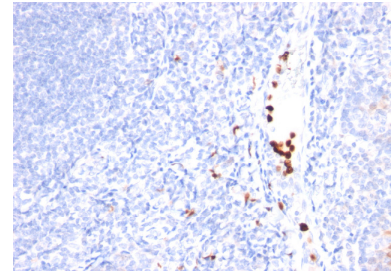


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Uses/Limitations: Not to be taken internally.
 For Research Use Only.
 This product is intended for qualitative immunohistochemistry with normal and neoplastic formalin-fixed, paraffin-embedded tissue sections, to be viewed by light microscopy.
 Do not use if reagent becomes cloudy.
 Do not use past expiration date.
 Non-Sterile.



Formalin-fixed, paraffin-embedded human tonsil stained with MYDAM; Clone BM-2. Note specific cytoplasmic staining of granulocytes.

Ordering Information and Current Pricing at www.scytek.com

Procedure:

1. **Tissue Section Pretreatment (Required):** Staining of formalin fixed, paraffin embedded tissue sections is significantly enhanced by pretreatment with Citrate Plus (ScyTek catalog# CPL500).
2. **Primary Antibody Incubation Time:** We suggest an incubation period of 30 minutes at room temperature. However, depending upon the fixation conditions and the staining system employed, optimal incubation should be determined by the user.
3. **Visualization:** For maximum staining intensity we recommend the “UltraTek HRP Anti-Polyvalent Lab Pack” (ScyTek catalog# UHP125, see IFU for instructions) combined with the “DAB Chromogen/Substrate Bulk Pack (High Contrast)” (ScyTek catalog# ACV500, see IFU for instructions).

Precautions:


Contains Sodium Azide as a preservative (0.09% w/v).
 Do not pipette by mouth.
 Avoid contact of reagents and specimens with skin and mucous membranes.
 Avoid microbial contamination of reagents or increased nonspecific staining may occur.
 This product contains no hazardous material at a reportable concentration according to U.S. 29 CFR 1910.1200, OSHA Hazardous Communication Standard and EC Directive 91/155/EC.


References:

1. Membrane protein hMYADM preferentially expressed in myeloid cells is up-regulated during differentiation of stem cells and myeloid leukemia cells. Wang Q, *et al.* Life Sci, 2007 Jan 9. PMID 17097684.
2. Cloning of human myeloid-associated differentiation marker (MYADM) gene whose expression was up-regulated in NB4 cells induced by all-trans retinoic acid. Cui W, *et al.* Mol Biol Rep, 2001. PMID 12075932.
3. MYADM controls endothelial barrier function through ERM-dependent regulation of ICAM-1 expression. Aranda JF, *et al.* Mol Biol Cell, 2013 Feb. PMID 23264465.
4. MYADM regulates Rac1 targeting to ordered membranes required for cell spreading and migration. Aranda JF, *et al.* Mol Biol Cell, 2011 Apr 15. PMID 21325632.
5. Isolation of MYADM, a novel hematopoietic-associated marker gene expressed in multi-potent progenitor cells and up-regulated during myeloid differentiation. Pettersson M, *et al.* J Leukoc Biol, 2000 Mar. PMID 10733104.


Warranty:

No products or “Instructions For Use (IFU)” are to be construed as a recommendation for use in violation of any patents. We make no representations, warranties or assurances as to the accuracy or completeness of information provided on our IFU or website. Our warranty is limited to the actual price paid for the product. ScyTek Laboratories, Inc. is not liable for any property damage, personal injury, time or effort or economic loss caused by our products. Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining can cause inconsistent results. Variations in fixation and embedding or the inherent nature of the tissue specimen may cause variations in results. Endogenous peroxidase activity or pseudoperoxidase activity in erythrocytes and endogenous biotin may cause non-specific staining depending on detection system used.

Storage: 2° C  8° C

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