



Mouse anti-Melansome (HMB45)

Cat. No.: AIB-30132 (1 ml Concentrate); AIB-30133 (0.5 ml Concentrate); AIB-30131 (6 ml Ready-to-use)

Instructions for use

Intended use

This antibody is designed for the specific localisation of melanocytic differentiation antigen gp100 in formalin-fixed, paraffin-embedded tissue sections. Anti-Melanosome antibody is intended for in vitro diagnostic use.

Specifications

Specificity: Melansome

Immunogen: Metastatic malignant melanoma cells

Clone: HMB45

Isotype: Mouse IgG1 kappa

Species reactivity: Human +, rat -, dog-, others not tested

Summary and Description

Diagnosis of metastatic amelanotic melanomas can be difficult. They can easily be mistaken for poorly differentiated carcinomas, sarcomas, or large-cell lymphomas. Primary spindle-cell melanomas can be mistaken as spindle-cell carcinomas or different types of mesenchymal neoplasms.

In the literature HMB45 is described as melanocytic differentiating antigen gp100. It is expressed in normal fetal and neonatal melanocytes and some nevus cells, but not in adult melanocytes or intradermal nevi. Hence, the marker is suitable for detection of melanocytic tumours. Other markers for diagnosis of melanomas are: S-100, Tyrosinase, Melan-A/MART1, and MiTF.

Reagent provided

Monoclonal mouse antibody from cell culture supernatant in buffer solution with carrier protein and preservative for stabilisation in the following formats:

 Concentrate:
 1 ml
 (Cat. No. AIB-30132)

 Concentrate:
 0.5 ml
 (Cat. No. AIB-30133)

 Ready-to-use:
 6 ml
 (Cat. No. AIB-30130)

Dilution of primary antibody

Dilution of Nordic Biosite' concentrated antibody depends on the detection system used. The final working dilution must always be determined by the user. The elaboration of staining protocol should be done by an experienced specialist. For Nordic Biosite' recommendations see chapter 'Staining procedure'.

Storage and handling

The antibody should be stored at 2-8°C without furt her dilution.

Dilutions of the concentrated antibody should be done with a suitable antibody dilution buffer (e.g. BCB-20005/BCB-20006from Nordic Biosite). The diluted antibody should be stored at 2-8°C after use. The stability of this work ing solution depends on various parameters and has to be confirmed by appropriate controls.

The antibody provided is suitable for use until the expiry date indicated on the label, if stored at 2-8°C. Do not use product after the expiry date. Positive and negative controls should be run simultaneously with all specimens. If unexpected staining is observed which cannot be explained by variations in laboratory procedures and a problem with the antibody is suspected, contact Nordic Biosite' technical support or your local distributor.

Explanations of the symbols on the product label:

REF	Catalog Number Bestellnummer Reference du catalogue	LOT	Batch Code Chargenbezeichnung Code du lot	Manufacturer Nordic BioSite AB
	Use By Verwendbar bis Utiliser jusque	IVD	In Vitro Diagnostic Medical Device In vitro Diagnostikum Dispositif médical de diagnostic in vitro	Propellervägen 4A S-183 62 Täby Sweden Tel: +46 (0)8 5444 33 40
	Consult Instructions for use Gebrauchsanweisung beachten Consulter les instructions d'utilisation		Temperature Limitation Lagerungstemperatur Limites de température	Fax: +46 (0)8 756 94 90 info@nordicbiosite.com www.nordicbiosite.com

Precautions

Use through qualified personnel only.

Wear protective clothing to avoid contact of reagents and specimens with eye, skin and mucous membranes. If reagents or specimens come in contact with sensitive area, wash with large amounts of water.

Microbial contamination of the reagent must be avoided, since otherwise non-specific staining may occur. Sodium azide (NaN3), used for stabilisation, is not considered hazardous material in the concentration used.

Reaction of sodium azide with lead or copper in drainage pipes can result in the formation of highly explosive metallic azides. Sodium azide should be discarded in a large volume of running water to avoid formation of deposits. A material safety data sheet (MSDS) is available upon request.

Staining procedure

Refer to the following table for conditions specifically recommended for this antibody. Also refer to detection system data sheets for guidance on specific staining protocols or other requirements.

<u>Parameters</u> <u>Nordic BioSites recommendations</u>

*Pre-treatment Heat Induced Epitope Retrieval (for example in Citrate Buffer pH 6.0 (BCB-

20015/-20016)

*Control tissue Melanoma

*Working dilution 1:50-1:100 (for concentrated antibodies only)

*Incubation time 30-60 minutes

Quality control

The recommended positive control tissue for this antibody is a melanoma. We recommend carrying out a positive and a negative control with every staining run. Please refer to the instructions of the detection system for guidance on general quality control procedures.

Troubleshooting

If you observe unusual staining or other deviations from the expected results please read these instructions carefully, refer to the instructions of the detection system for relevant information or contact your local distributor.

Expected results

This antibody stains positive in the cytoplasm of malignant melanoma cells in formalin-fixed, paraffin-embedded tissue sections. Interpretation of the staining results is solely the responsibility of the user. Any experimental result should be confirmed by a medically established diagnostic procedure.

Limitations of the Procedure

Immunohistochemistry is a complex technique involving both histological and immunological detection methods. Tissue processing and handling prior to immunostaining, for example variations in fixation and embedding or the inherent nature of the tissue can cause inconsistent results (Nadji and Morales, 1983). Endogenous peroxidase, alkaline phosphatase or biotin may cause non-specific staining depending on the detection system used. Tissues containing Hepatitis B Surface Antigen (HBsAg) may give false positive results with HRP (horse radish peroxidase) detection systems (Omata et al, 1980). Inadequate counterstaining and mounting can influence the interpretation of the results.

Nordic Biosite warrants that the product will meet all requirements described from its shipping date until the expiry date is reached, if the product is stored and utilised as recommended. No additional guarantees can be given. Under no circumstances shall Nordic Biosite be liable for any damages arising out of the use of the reagent provided.

Performance characteristics

Nordic Biosite has conducted studies to evaluate the performance of the antibody for use with a standard detection system. The product has been found to be sensitive and specific to the antigen of interest with minimal or no cross-reactivity.

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Bibliography

Bruggen J, et al. Cancer Immunol Immunother 15:200-205, 1983 DeLellis, R., et al. Semin Oncol 14:173-192, 1987 Gown AM, et al. Am J Pathol 123:195-203, 1986 Omata M et al. Am J Clin Pathol 73(5): 626-32, 1980 Ordonez NG, et al. Am J Clin Pathol 90:385:390,1988 Wick MR, et al. J Cutan Pathol 15:201-207, 1988 Yaziji H, Gown AM, Int J Surg Pathol, 11:11-15, 2003 Nadji M and Morales AR Ann N.Y. Acad Sci 420:134-9, 1983

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