

MART-1 + Tyrosinase + SOX10

Prediluted Monoclonal Antibody Cocktail
901-3165-091025



Available Product Formats

Format	Catalog Number	Description	Dilution
NeoPATH PRO	NPAI 3165 T40	40 tests	Ready-to-use

Intended Use:

For *in vitro* Diagnostic Use

The Biocare MART-1 + Tyrosinase + SOX10 mouse monoclonal antibody cocktail is intended for laboratory professional use in the qualitative identification of MART-1, Tyrosinase, and SOX10 proteins in sections of formalin-fixed, paraffin-embedded (FFPE) human tissue by an immunohistochemistry (IHC) procedure performed manually or on an automated slide stainer and visualized by light microscopy. The clinical interpretation of any staining or its absence should be complemented by morphological studies and proper controls and should be evaluated within the context of the patient's clinical history and other diagnostic tests by a qualified pathologist.

Summary and Explanation:

MART-1 recognizes an 18 kDa protein (Melanoma Antigen Recognized by T cells 1) and is specific for melanocytic lesions, making it a valuable component of melanoma panels.¹⁵⁻¹⁷ MART-1 has been shown to be more sensitive than HMB45 for labeling metastatic melanomas. Tyrosinase, a key enzyme in early melanin biosynthesis, is more sensitive than HMB45 and MART-1 and labels a higher percentage of desmoplastic melanomas.¹⁵ The combination of MART-1 and Tyrosinase aids in identifying metastatic melanoma in sentinel lymph nodes.¹⁶

SOX10, a transcription factor in the SRY-related HMG-Box family, plays an important role in neural crest, peripheral nervous system, and melanocytic cell development.¹⁸⁻²² It is widely expressed in normal human tissues including melanocytes and breast tissue and is a marker for malignant tumors such as melanoma, breast carcinoma, gliomas, and benign tumors including schwannomas.¹⁸⁻²⁰ SOX10 is expressed in the vast majority of desmoplastic and spindle cell melanomas and in 100% of nevi.^{18,19} Compared to S100, SOX10 is less likely to stain background fibroblasts or histiocytes within scars, making it a more specific marker in these contexts.²⁰

The combination of SOX10 with MART-1 and/or Tyrosinase increases detection sensitivity in lymph node and metastatic melanoma specimens compared to S100 and provides greater specificity, as S100 stains dendritic processes in lymph nodes while SOX10, MART-1, and Tyrosinase remain negative.^{21,22} This antibody cocktail is suitable for tumors of unknown origin or for differential diagnosis of melanoma and its mimics.

Principle of Procedure:

This antibody cocktail product may be used as the primary antibody in immunohistochemistry testing of formalin-fixed, paraffin-embedded tissue sections. In general, immunohistochemical (IHC) staining techniques allow for the visualization of antigens via the sequential application of a specific antibody cocktail to the antigen (primary antibody), a secondary antibody cocktail containing anti-rabbit and anti-mouse enzyme complexes to the primary antibody and the chromogenic substrates with interposed washing steps. The enzymatic activation of the chromogens results in a visible reaction product at the antigen sites. The specimen may then be counterstained, and cover slipped. Results are interpreted using a light microscope and aid in the differential diagnosis of pathophysiological processes, which may or may not be associated with a particular antigen.

Materials and Methods:

Reagents Provided:

Antibody:	Anti-MART-1	Anti-Tyrosinase	Anti-SOX10
Host Source:	Mouse monoclonal	Mouse monoclonal	Mouse monoclonal
Species Reactivity:	Human. Other species not tested		
Clone:	M2-7C10 + M2-9E3	T311	BC34
Isotype:	IgG2b/kappa + IgG2b	IgG2a/kappa	IgG1/kappa
Protein Concentration:	Contact Biocare's Technical Support for specific Ig concentration.		
Specificity:	MART-1	Tyrosinase	SOX10
Cellular Localization:	Cytoplasmic	Cytoplasmic	Nuclear
Staining:	Brown (DAB)	Brown (DAB)	Brown (DAB)

Method: Affinity purified mouse monoclonal antibodies.

Reconstitution, Mixing, Dilution, Titration:

Prediluted antibody reagent is optimally diluted for use with the below listed staining system. Further dilution may result in loss of antigen staining. The user must validate any such change. Differences in tissue processing and technical procedures in the user's laboratory may produce significant variability in results necessitating regular performance of in-house controls (see Quality Control section).

Known Applications:

Immunohistochemistry (formalin-fixed paraffin-embedded tissues)

Supplied As:

Buffered saline solution, pH 5.9-6.0, contains a protein carrier and less than 0.1% sodium azide preservative. See Safety Data Sheet for additional details.

Materials and Reagents Needed but Not Provided:

Microscope slides positively charged.
Positive and negative tissue controls
Desert Chamber (or similar Drying oven)
Xylene or xylene substitute
Ethanol or reagent alcohol
Decloaking Chamber (Pressure cooker)
Deionized or distilled water
Wash buffer
Pretreatment reagents
Peroxidase block
Protein block (optional) Remove (Optional) if protein block is required.
Detection probe and polymer
Negative control reagents
Chromogens
Hematoxylin (counterstain)
Bluing reagent
Mounting medium
Coverglass
Light Microscope (40-400X magnification)
Automated Staining Platform

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Configurations of the antibody product are available for use on the instruments indicated in the table above.

Storage and Stability:

Store at 2°C to 8°C. The product is stable to the expiration date printed on the vial label, when stored under these conditions. Do not use after expiration date. Storage under any condition other than those specified must be verified. Diluted reagents should be used promptly; store any remaining reagent at 2°C to 8°C. The stability of user diluted reagents has not been established by Biocare.

Positive and negative controls should be run simultaneously with all patient specimens. If unexpected staining is observed, which cannot be explained by variations in laboratory procedures and a problem with the antibody is suspected, contact Biocare's Technical Support at 1-800-542-2002 or via the technical support information provided on biocare.net.

Specimen Preparation:

Tissues fixed in formalin are suitable for use prior to paraffin embedding. Osseous tissues should be decalcified prior to tissue processing to facilitate tissue cutting and prevent damage to microtome blades.^{1,2}

Properly fixed and embedded tissues expressing the specified antigen target should be stored in a cool place. The Clinical Laboratory Improvement Act (CLIA) of 1988 requires in 42 CFR §493.1259(b) that "The laboratory must retain stained slides at least ten years from the date of examination and retain specimen blocks at least two years from the date of examination."³

Treatment of Tissues Prior to Staining:

Perform Heat Induced Epitope Retrieval (HIER) per recommended protocol below. The routine use of HIER prior to IHC has been shown to minimize inconsistency and standardize staining.^{4,5}

Warning and Precautions:

1. This antibody contains less than 0.1% sodium azide. Concentrations less than 0.1% are not reportable hazardous materials according to U.S. 29 CFR 1910.1200, OSHA Hazard communication and EC Directive 91/155/EC. Sodium azide (NaN₃) used as a preservative is toxic if ingested. Sodium azide may react with lead and copper plumbing to form highly explosive metal azides. Upon disposal, flush with large volumes of water to prevent azide build-up in plumbing. (Center for Disease Control, 1976, National Institute of Occupational Safety and Health, 1976)⁶
2. Specimens, before and after fixation, and all materials exposed to them should be handled as if capable of transmitting infection and disposed of with proper precautions. Never pipette reagents by mouth and avoid contacting the skin and mucous membranes with reagents and specimens. If reagents or specimens come into contact with sensitive areas, wash with copious amounts of water.⁷
3. Microbial contamination of reagents may result in an increase in nonspecific staining.
4. Incubation times or temperatures other than those specified may give erroneous results. The user must validate any such change.
5. Do not use reagent after the expiration date printed on the vial.
6. Prediluted antibody reagent is optimally diluted for use. Further dilution may result in loss of antigen staining.
7. To prevent evaporation and ensure maximum test capacity, promptly cap and remove reagents from automated instruments after each run. Leaving reagents exposed can reduce their effectiveness and the number of tests they can provide. Always store reagents as directed to maintain their integrity.
8. Dispose of all used reagents and any other contaminated disposable materials following procedures for infectious or potentially infectious waste.

It is the responsibility of each laboratory to handle solid and liquid waste according to their nature and degree of hazard and to treat and dispose of it (or have them treated and disposed of) in accordance with any applicable regulations.

9. Follow local disposal regulations for your location along with recommendations in the Safety Data Sheet to determine the safe disposal of this product

10. The SDS is available upon request and is located at <http://biocare.net>.

11. To report suspected serious incidents related to this device, contact the local Biocare representative and the competent authority of the Member State or Country in which the user is established.

Instructions for Use:

Recommended Staining Protocols for MART-1 + Tyrosinase + SOX10:

NeoPATH PRO:

NPAI3165 is intended for use with the NeoPATH PRO. Refer to the User Manual for specific instructions for use. Recommended protocol parameters are as follows:	
Antibody Protocol:	MART + Tyrosinase + SOX10, 20 min at RT
Template:	HRP_HIGH_105C_20MINAB_STD
Dewax:	Dewax STD; 20 min at 75°C
Antigen Retrieval (HIER Option):	HIGH_105C_30MIN
Enzyme:	N/A
Block Option:	N/A
Detection:	HRP_20AB_STD (Amplifier; 10 min at RT, Polymer; 25 min at RT)
Chromogen:	7 min DAB + 2 min DAB Enhancer at RT.
Hematoxylin:	7 min at RT

Quality Control:

Refer to CLSI Quality Standards for Design and Implementation of Immunohistochemistry Assays; Approved Guideline-Second edition (I/LA28-A2) CLSI Wayne, PA USA (www.clsi.org). 2011⁸

Positive Tissue Control: Melanoma

External Positive control materials should be fresh specimens fixed, processed, and embedded as soon as possible in the same manner as the patient sample(s). Positive tissue controls are indicative of correctly prepared tissues and proper staining techniques. One positive external tissue control for each set of test conditions should be included in each staining run.

The tissues used for the external positive control materials should be selected from patient specimens with well-characterized low levels of the positive target activity that gives weak positive staining. The low level of positivity for external positive controls is designed to ensure detection of subtle changes in the primary antibody sensitivity from instability or problems with the IHC methodology. Commercially available tissue control slides or specimens processed differently from the patient sample(s) validate reagent performance only and do not verify tissue preparation.

Known positive tissue controls should only be utilized for monitoring the correct performance of processed tissues and test reagents, rather than as an aid in formulating a specific diagnosis of patient samples. If the positive tissue controls fail to demonstrate positive staining, results with the test specimens should be considered invalid.

Negative Tissue Control:

Use a negative tissue control (known to be MART-1 + Tyrosinase + SOX10 negative) fixed, processed, and embedded in a manner identical to the patient

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sample(s) with each staining run to verify the specificity of the IHC primary antibody for demonstration of the target antigen, and to provide an indication of specific background staining (false positive staining). Also, the variety of different cell types present in most tissue sections can be used by the laboratorian as internal negative control sites to verify the IHC's performance specifications. The types and sources of specimens that may be used for negative tissue controls are listed in the Performance Characteristics section.

If specific staining (false positive staining) occurs in the negative tissue control, results with the patient specimens should be considered invalid.

Nonspecific Negative Reagent Control:

Use a nonspecific negative reagent control in place of the primary antibody with a section of each patient specimen to evaluate nonspecific staining and allow better interpretation of specific staining at the antigen site. Ideally, a negative reagent control contains a MART-1 / IgG2b, kappa + IgG2b, Tyrosinase / IgG2a, kappa, and SOX10 / IgG1, kappa mouse monoclonal antibody produced from tissue culture supernatant in the same way as the primary antibody but exhibits no specific reactivity with human tissues in the same matrix/solution as the Biocare antibody. Dilute a negative control antibody to the same immunoglobulin or protein concentration as the diluted primary antibody using the identical diluent. If fetal calf serum is retained in the neat antibody after processing, fetal calf serum at a protein concentration equivalent to the diluted primary antibody in the same diluent is also suitable for use. (Refer to reagent provided). Diluent alone may be used as a less desirable alternative to the previously described negative reagent controls. The incubation period for the negative reagent control should correspond to that of the primary antibody.

When panels of several antibodies are used on serial sections, the negatively staining areas of one slide may serve as a negative/nonspecific binding background control for other antibodies. To differentiate endogenous enzyme activity or nonspecific binding of enzymes from specific immunoreactivity, additional patient tissues may be stained exclusively with substrate-chromogen or enzyme complexes (PAP, avidin-biotin, streptavidin) and substrate-chromogen, respectively.

Assay Verification:

Prior to initial use of an antibody or staining system in a diagnostic procedure, the user should verify the antibody's specificity by testing it on a series of in-house tissues with known immunohistochemical performance characteristics representing known positive and negative tissues. Refer to the quality control procedures previously outlined in this section of the product insert and to the quality control recommendations of the CAP Certification Program⁹ for Immunohistochemistry and/or the NCCLS IHC guideline¹⁰). These quality control procedures should be repeated for each new antibody lot, or whenever there is a change in assay parameters.

Troubleshooting:

Follow the antibody specific protocol recommendations according to the data sheet provided. If atypical results occur, contact Biocare's Technical Support at 1-800-542-2002.

Interpretation of Staining:

Positive Tissue Control:

The positive tissue control stained with indicated antibody should be examined first to ascertain that all reagents are functioning properly. The appropriate staining of target cells (as indicated above) is indicative of positive reactivity. If the positive tissue controls fail to demonstrate positive staining, any results with the test specimens should be considered invalid.

The color of the reaction product may vary depending on substrate chromogens used. Refer to substrate package inserts for expected color reactions. Further, metachromasia may be observed in variations of the method of staining.¹¹

When a counterstain is used, depending on the incubation length and potency of the counterstain used, counterstaining will result in a coloration of the cell nuclei. Excessive or incomplete counterstaining may compromise proper interpretation of results. Refer to protocol(s) for recommended counterstain.

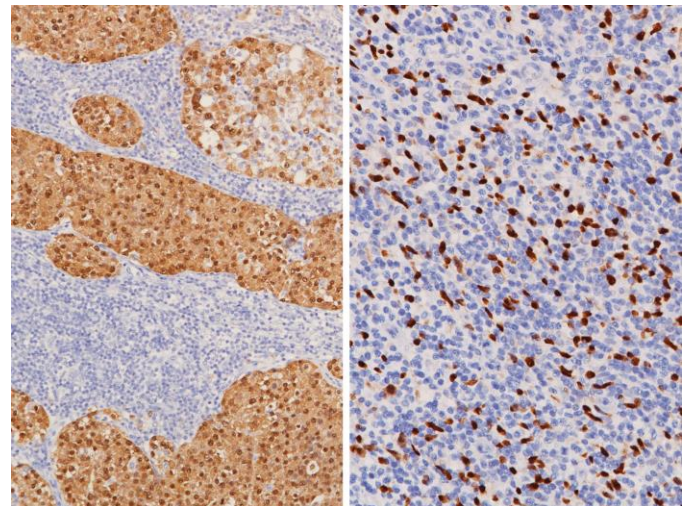
Negative Tissue Control:

The negative tissue control should be examined after the positive tissue control to verify the specificity of the labeling of the target antigen by the primary antibody. The absence of specific staining in the negative tissue control confirms the lack of antibody cross reactivity to cells/cellular components. If specific staining (false positive staining) occurs in the negative external tissue control, results with the patient specimen should be considered invalid.

Nonspecific staining, if present, usually has a diffuse appearance. Sporadic staining of connective tissue may also be observed in sections from excessively formalin-fixed tissues. Use intact cells for interpretation of staining results. Necrotic or degenerated cells often stain nonspecifically.

Patient Tissue:

Examine patient specimens stained with indicated antibody last. Positive staining intensity should be assessed within the context of any nonspecific background staining of the negative reagent control. As with any immunohistochemical test, a negative result means that the antigen was not detected, not that the antigen was absent in the cells/tissue assayed. If necessary, use a panel of antibodies to identify false-negative reactions.



(L) Metastatic melanoma stained with MART-1 + Tyrosinase (cytoplasmic, positive) and SOX10 (nuclear, positive), (R) Melanoma stained with MART-1 + Tyrosinase (cytoplasmic, negative) and SOX10 (nuclear, positive).

Refer to Summary and Explanation and Limitations for specific information regarding indicated antibody immunoreactivity.

Limitations:

General Limitations:

1. For *in vitro* diagnostic Use

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2. This product is for professional use only: Immunohistochemistry is a multistep diagnostic process that consists of specialized training in the selection of the appropriate reagents; tissue selection, fixation, and processing; preparation of the IHC slide; and interpretation of the staining results.
3. Tissue staining is dependent on the handling and processing of the tissue prior to staining. Improper fixation, freezing, thawing, washing, drying, heating, sectioning or contamination with other tissues or fluids may produce artifacts, antibody trapping, or false negative results. Inconsistent results may be due to variations in fixation and embedding methods, or to inherent irregularities within the tissue.¹²
4. Excessive or incomplete counterstaining may compromise proper interpretation of results.
5. The clinical interpretation of any positive or negative staining should be evaluated within the context of clinical presentation, morphology, and other histopathological criteria. The clinical interpretation of any positive or negative staining should be complemented by morphological studies using proper positive and negative internal and external controls as well as other diagnostic tests. It is the responsibility of a qualified pathologist who is familiar with the proper use of IHC antibodies, reagents, and methods to interpret all the steps used to prepare and interpret the final IHC preparation.
6. The optimum antibody dilution and protocols for a specific application can vary. These include, but are not limited to fixation, heat-retrieval method, incubation times, tissue section thickness and detection kit used. Due to the superior sensitivity of these unique reagents, the recommended incubation times and titers listed are not applicable to other detection systems, as results may vary. The data sheet recommendations and protocols are based on exclusive use of Biocare products. Ultimately, it is the responsibility of the investigator to determine optimal conditions.
7. This product is not intended for use in flow cytometry. Performance characteristics have not been determined for flow cytometry.
8. Tissues from persons infected with hepatitis B virus and containing hepatitis B surface antigen (HBsAg) may exhibit nonspecific staining with horseradish peroxidase.¹³
9. Reagents may demonstrate unexpected reactions in previously untested tissues. The possibility of unexpected reactions even in tested tissue groups cannot be completely eliminated due to biological variability of antigen expression in neoplasms, or other pathological tissues.¹⁴ Contact Biocare's Technical Support at 1-800-542-2002 or via the technical support information provided on biocare.net, with documented unexpected reaction(s).
10. Normal/nonimmune sera from the same animal source as secondary antisera used in blocking steps may cause false-negative or false-positive results due to autoantibodies or natural antibodies.
11. False-positive results may be seen due to non-immunological binding of proteins or substrate reaction products. They may also be caused by pseudo peroxidase activity (erythrocytes), endogenous peroxidase activity (cytochrome C), or endogenous biotin (e.g., liver, breast, brain, kidney) depending on the type of immunostain used.¹²

Product Specific Limitations:

No additional product specific limitations noted.

Troubleshooting:

1. No staining of any slides – Check to determine appropriate positive control tissue, antibody, and detection products have been used.
2. Weak staining of all slides – Check to determine appropriate positive control tissue, antibody, and detection products have been used.
3. Excessive background of all slides – There may be high levels of endogenous biotin (if using biotin-based detection products), endogenous HRP activity converting chromogen to colored end product (use

peroxidase block), or excess non-specific protein interaction (use a protein block, such as serum- or casein-based blocking solution).

4. Tissue sections wash off slides during incubation – Check slides to ensure they are positively charged.
5. Specific staining too dark – Check protocol to determine if proper antibody titer was applied to slide, as well as proper incubation times for all reagents. Additionally, ensure the protocol has enough washing steps to remove excess reagents after incubation steps are completed.

References:

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