

**ihcDirect® SMMS-1 Ab**  
**Anti-Human Smooth Muscle Myosin Heavy Chain**  
**(Clone: SMMS-1)**

Ab: K31024-015, 150 tissue stains\*  
Ab: K31024-005, 50 tissue stains\*

**Intended Use:** For *In Vitro* Diagnostic Use

Polymerized horseradish peroxidase (pHRP)-labeled anti-human smooth muscle myosin heavy chain antibody (SMMS-1 pHRP) is intended for laboratory use to qualitatively identify by light microscopy the presence of smooth muscle myosin heavy chain in sections of formalin-fixed, paraffin-embedded tissue sections or frozen tissues using immunohistochemistry (IHC) test methods. The clinical interpretation of any staining or its absence should be complemented by morphological studies using proper controls and should be evaluated within the context of the patient's clinical history and other diagnostic tests by a qualified pathologist/physician. This conjugate has been pre-diluted and optimized for IHC use without further dilution.

**Summary and Explanation:**

Smooth muscle myosin heavy chain is a major component of the contractile apparatus of the smooth muscle cells, and also is a myoepithelium-associated protein. SMMS-1 antibody reacts with human visceral and vascular smooth muscle cells. The antibody also recognizes human myoepithelial cells. IHC staining with SMMS-1 antibody is very helpful in distinguishing benign sclerosing breast lesions and carcinoma in situ from infiltrating breast carcinoma. (Kalof et al., 2004; Hill CB, Yeh IT, 2005). The key component in this product is horseradish peroxidase (HRP) polymer (PolyHRP) labeled anti human smooth muscle myosin heavy chain antibody (clone SMMS-1). The ihc Blocker is used prior to applying SMMS-1 pHRP antibody conjugate to help reduce background and nonspecific staining. A chromogen such as 3,3'-diaminobenzidine (DAB) is then used to develop color at the reaction site.

**Principle of Procedure:**

The ready-to-use ihcDirect SMMS-1 pHRP conjugate is directly applied to pretreated tissue sections, where it binds to smooth muscle myosin heavy chain. A Working Solution (WS) of a chromogen such as ihc DAB 1:1 is then applied to the tissue. The SMMS-1 antibody-linked pHRP reacts with the chromogen to form a visible colored product at the location of smooth muscle myosin heavy chain. The specimen may then be counterstained and a coverslip applied. Results are viewed and interpreted using a light microscope. Volumes are based upon 100µl antibody per tissue. This product may be used to perform IHC either manually or on an open automatic IHC staining system.

**Reagents Provided:**

Part No.	Σ	Description
K31024-015*	150*	15ml size ihcDirect SMMS-1 ready-to-use antibody conjugate. See Ancillary Reagents section.
K31024-005*	50*	5ml size ihcDirect SMMS-1 ready-to-use antibody conjugate. See Ancillary Reagents section.

\* At estimated volume of 100µl of antibody conjugate per tissue

Immunogen	Clone	Species	Ig Class	Total Protein Conc.
Human smooth muscle myosin heavy chain	SMMS-1	Mouse	IgG	10 mg/ml

SMMS-1 antibody is a mouse monoclonal antibody to human smooth muscle myosin heavy chain purified from ascites. HRP is extracted from horseradish plant.

Novodiox ihc Blocker and chromogens such as ihc DAB 1:1 Kit, ihc Magenta 1:1 Kit, and DAB Kit are recommended for use with the SMMS-1 antibody.

**SMMS-1 Antibody (K31024-###):**

Kit Components	Component Part Numbers	Sizes (ml)
SMMS-1 pHRP	H31024-(R###) (005, 015)	5, 15

**Ancillary Reagents for Use with SMMS-1 Antibody:**

Reagent Description	Part Numbers	Sizes (ml)
ihc Blocker Intl. USA	K51001-(###) (015) (OR)	15
	K51002-(###) (015, 030)	15, 30
ihc DAB 1:1 Kit	K50002-### (015, 030)	15, 30
ihc Magenta 1:1 Kit USA	K50011-### (015, 030)	15, 30
DAB Kit	K50001-### (015, 030)	15, 30

**Materials Needed but Not Provided:**

The following reagents/supplies may be required in staining but are not provided:

- |   |  |
|---|--|
| 1. Frozen section fixative (Acetone or NBF§)          | 9. Peroxide blocker (optional)   |
| 2. Positive and negative control tissues              | 10. Instruments used for tissue pretreatment, such as water bath, or pressure cooker or microwave oven (when using FFPE tissues) |
| 3. Microscope slides, positively charged (required)   | 11. Hematoxylin  |
| 4. Staining jars, baths or processing tools           | 12. Xylene or Xylene substitute  |
| 5. ihc Wash Buffer (PBS-T)                            | 13. Ethanol  |
| 6. Pipettor and pipet tips                            | 14. Mounting medium  |
| 7. Timer  | 15. Cover slips  |
| 8. Antigen retrieval buffer (when using FFPE tissues) | 16. Light microscope (40 - 400x)   |
- §NBF – neutral buffered formalin

**Novodiox Bulk Reagent Formulations:**

- ihc Wash Buffer (PBS-T), (10 mM phosphate buffer, pH7.2, 150 mM NaCl, 0.05% Tween-20).
- ihc Antigen Retrieval Buffer (10mM Citric buffer, pH 6.0, 0.02% Tween 20).

**Storage and Handling:**

This product should be stored at 2-8°C and is suitable for use until expiry date when stored at this temperature. Do not freeze. Do not use the product after expiration date unless dating extension information is provided by Novodiox. If reagents are stored under any conditions other than those specified in the package insert, they must be verified by the user.

**Specimen Preparation:**

**Paraffin Sections:** Tissues routinely processed with 10% NBF are suitable for use prior to paraffin embedding. Consult references (Kiernan, 1981; Sheehan & Hrapchak, 1980). Variable results may occur as a result of prolonged fixation. Each section should be cut to the appropriate thickness (approximately 4-5 µm) and placed on a positively charged glass slide. Slides containing the tissue section may be baked for at least one hour but not exceeding 24 hours in a 58-60°C±5°C oven. Osseous tissues should be decalcified prior to tissue processing to facilitate tissue cutting and prevent damage to microtome blades (Kiernan, 1981; Sheehan & Hrapchak, 1980).



**Frozen Tissue Sections:** Frozen tissue is sectioned to the appropriate thickness (approximately 5 µm) and placed on a positively charged glass slide. Tissues should be fixed in either a reagent grade acetone or 10% NBF for 1-2 minutes immediately after sectioning. A short time to air dry may help tissues adhere to the slide. Other fixatives should be validated prior to use. Reagent grade acetone may be kept cold, e.g. at cryostat temperatures or at room temperature. Following fixation, tissues should be processed within a few minutes or may be stored in a PBS buffer for as long as a day.

**Treatment of Tissues Prior to Staining:** Pretreatment is tissue dependent and should be performed as suggested in the staining procedure sections.

**Warnings and Precautions:**

1. Read and understand all of the Novodiox Instructions for Use (IFUs) before product use.
2. The SMMS-1 antibody-pHRP conjugate is pre-diluted. Further dilution may reduce signal intensity or increase false-negative staining. These recommendations are for guidance only. Laboratory managers should determine their own procedures and quality policies.
3. To obtain best results when working with frozen tissues, it is desirable to freeze tissues as quickly as possible following extraction.
4. Use caution and shorten incubation times when utilizing intense hematoxylin counterstains such as Gills as these stains may tend to overstain and obscure antibody staining.
5. Take reasonable precautions when handling reagents. Use protective equipment such as disposable gloves and lab coats when handling suspected carcinogens or toxic materials. Read Safety Data Sheets (SDS) prior to use.
6. Avoid contact of reagents with eyes and mucous membranes. If reagents come in contact with sensitive areas, wash with copious amounts of water.
7. Use charged slides to secure tissue adhesion.
8. Patient specimens and all materials that come into contact with patient specimens should be handled as bio-hazardous materials and disposed of appropriately.
9. Consult local or state authorities with regard to recommended methods of disposal of bio-hazardous and hazardous chemical waste materials.
10. Incubation time and temperature other than those specified may give erroneous results. The user must validate any such changes.
11. Use lab grade quality chemicals such as acetone, ethanol and water when preparing solutions and buffers. Users should validate performance including stability for laboratory prepared reagents (at 1X).
12. Avoid microbial contamination of reagents.
13. Fixation is a vital part of the protocol and fixation times may vary with the fixative chosen, tissue type, e.g. containing fat and other parameters. Generally, an acetone or NBF fixation of 1-2 minutes is recommended. Place frozen tissue sections into fixation solution shortly after sectioning.
14. **Prolonged exposure to room or freezing temperatures may alter targeted epitopes.** It is best to prevent slides from drying out during the staining process to avoid unwanted background staining.

**Staining Procedures:**

General Operating Notes:

1. Equilibrate all reagents to room temperature prior to use. Swirl or shake the ihc Blocker and pHRP-labeled antibody solution before use. **Do not vortex.** Calculate the amount of chromogen WS needed (100µl per tissue) and **freshly** prepare chromogen WS. See instructions for use.
2. Gently and thoroughly wash tissues during manual wash steps. Avoid direct high velocity streams of wash that might tend to damage or cut delicate tissues.
3. Following each manual assay step, remove excess fluids on tissue slides with tissue paper. Excessive residual solution may dilute subsequent reagents, causing negative or uneven staining.
4. To reduce background signal, wash thoroughly following antibody step.
5. For the tissues with high levels of oxidase activity, e.g. gastrointestinal or renal tissues, an additional blocking step with H<sub>2</sub>O<sub>2</sub> is required to minimize background.
6. The following protocol for frozen tissues has been validated at temperatures between 21-30°C (70-86°F) for incubating ihc Blocker, SMMS-1 pHRP and chromogen WS. If room temperature is less than 21°C, incubate labeled antibody for a longer period of time (≤5 minutes depending upon temperature). Consistent results have been obtained using a slide warmer set to 30°C at the surface of the slide.

**Frozen Tissue Sections:**

1. Following fixation, rinse slides with 1X ihc Wash buffer and then wipe away any excess fluid with a Kimwipe® or paper towel.
2. Dispense 100µl of ihc Blocker, covering the entire tissue, and incubate at room temperature for 1 minute. Then firmly tap off any excess ihc Blocker onto an absorbent surface but do not rinse the slides.
3. Dispense 100µl of pHRP antibody, covering the entire tissue, and incubate for 3 minutes at room temperature. To obtain a darker stain, users may extend incubation up to a total of 5 minutes. Then, rinse slides with 1X ihc Wash buffer and wipe away any excess fluid.
4. Dispense 100µl of a chromogen WS, such as DAB covering the entire tissue, and incubate for 1-3 minutes at room temperature. Users should determine the optimal incubation time for their particular chromogen or lab environment. Then rinse slides with either 1X ihc Wash buffer or lab grade water and wipe away any excess fluid.
5. Add a counterstain. Incubation times will vary according to the counterstain formulation. Then, rinse slides with water and wipe away any excess fluid.
6. Apply aqueous media or dehydrate slides with the user's typical dehydration protocol and then add coverslip.

**Test Timing Est. (10-minute IHC protocol for frozen tissue sections):**

ihcDirect Frozen Tissue Procedure	Time in minutes
Fix with Acetone or Neutral Buffered Formalin	Start
- Wash with <input type="text" value="ihc Wash"/> remove excess fluid	
Block with <input type="text" value="ihc Blocker"/>	1
- Tap+Absorb to remove excess blocker	- - -
Novodiox ihcDirect pHRP Antibody	3
- Wash thoroughly with <input type="text" value="ihc Wash"/>	
- Tap+Absorb to remove excess wash buffer	- - -
Novodiox Chromogen Working Solution	1-3
- Wash with <input type="text" value="ihc Wash"/> or DI water	
Hematoxylin counterstain (Conc. Dependent)	2-45 sec
Wash with water	
Dehydrate/Mounting Medium and Coverslip	User Det.
<b>Total</b>	<b>~10</b>

**Paraffin Tissues:**

1. Deparaffinization: Soak slides in Xylene 3 times for 5 minutes each. Next, 3 minutes each in 100%, 95% and 75% ethanol. Then wash slides with tap water in slide tank for two times, 2 minutes each time.
2. Antigen retrieval: Using a water bath, incubate slides in antigen retrieval buffer in a slide tank at 96°C for 30 minutes, then cool the slides down to room temperature for 30 minutes. Rinse the slides twice with tap water, 2 minutes each time.
3. (Optional) Block tissue with H<sub>2</sub>O<sub>2</sub>: Soak the slides in 3% H<sub>2</sub>O<sub>2</sub> in slide tank, stand for 10 minutes. Rinse the slides with tap water twice and then wash once with PBS-T in slide tank for 2 minutes.
4. Dispense 100µl of ihc Blocker covering the entire tissue and incubate at room temperature for 15 minutes. Remove ihc Blocker as much as possible but do not rinse the slides with PBS-T or water.
5. Dispense 100 µl of pHRP labeled SMMS-1 antibody on slides covering the entire tissue and incubate for 30 minutes at room temperature. Rinse the slides three times with PBS-T in slide tank, 2 minutes each time. Note: Place slides in a wet chamber to prevent evaporation if longer incubation times are used.
6. Dispense 100µl of chromogen WS covering the entire tissue, incubate for 3-10 minutes at room temperature. Rinse the slides twice with tap water in slide tank, 2 minutes each time.



- Counterstaining: Add hematoxylin and incubate for 1 minute at room temperature. Rinse twice with tap water for 2 minutes, each time.
- Dehydration: Soak slides in the following order: 75% ethanol for 3 minutes, 95% ethanol for 3 minutes, 100% ethanol for 3 minutes and Xylene twice at 5 minutes each time.
- Applying Coverslip: Add one drop of permanent mounting medium on both the slide and the coverslip, then apply coverslip.

#### Quality Control Procedures:

Positive and negative controls should be run simultaneously with patient specimens.

**Positive Tissue Control:** The recommended positive control tissues for this antibody are properly processed normal breast or colon tissues. The staining is cytoplasmic. One positive tissue control for each set of test conditions should be included in each staining run. Previous tissue specimens that have been frozen and freshly cut or in some cases, an individual's own tissue may be used as a control.

The tissues used for the positive control should be selected from patient specimens with well-characterized low levels of positive target activity that give weak positive staining. 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 patient specimens should be considered invalid.

**Negative Tissue Control:** The same tissue used for the positive control may be used as the negative tissue control. The variety of cell types in most tissue sections offers internal negative control sites, but this should be verified by the user. The components that do not stain should demonstrate the absence of specific staining, and provide an indication of non-specific background staining. If specific staining occurs in the negative tissue control sites, results with the patient specimens must be considered invalid.

#### Troubleshooting:

If an unexpected staining pattern occurs on control tissues or patient samples, consider the following:

- High background:** Possible causes include (1) insufficient washing, (2) ihc Blocker not being applied or washed out after application, (3) specimens drying out, (4) prolonged chromogen incubation, (5) prolonged pHRP-labeled antibody incubation, and (6) specimens containing high level of endogenous peroxidase, which necessitates an additional blocking step (refer to the Staining Procedures for Paraffin Tissues). Perform any required corrective actions and repeat the procedure.
- Low signal or faint staining:** Please check whether (1) the reagents are not expired, (2) temperature of the testing environment was at least 21°C or a 30°C slide warmer was used, (3) chromogen WS was prepared freshly and correctly, (4) excess ihc Wash solution was not left on the slide, causing subsequent reagents to be diluted, and (5) for FFPE tissue, dewaxing and antigen retrieval were performed adequately. Perform any required corrective actions and repeat the procedure. Alternatively, if using a DAB chromogen, consider using another stain, e.g. ihc Magenta 1:1 to obtain more vibrant staining. In addition, some individuals may naturally have low expression of certain antigens. In these cases, users may extend the antibody incubation times by 1-2 minutes.
- High background:** Possible causes include (1) insufficient washing, (2) ihc Blocker not being applied or washed out after application, (3) specimens drying out, (4) prolonged chromogen incubation, (5) prolonged pHRP-labeled antibody incubation, and (6) specimens containing high level of endogenous peroxidase, which necessitates an additional blocking step (refer to the Staining Procedures for Paraffin Tissues). Perform any required corrective actions and repeat the procedure.

If an unexpected staining pattern is observed on control tissues or patient samples which cannot be explained by variations in laboratory procedures or a problem with the antibody is suspected, contact NovodiAx Technical Support or your local distributor immediately. Within the US and Canada call 1 (888) 439-2716 ext. 2 or 1 (510) 342-3043 ext. 2.

#### Expected Results:

Intense color stains with a clean background if smooth muscle myosin heavy chain-expression cells exist. There will be no color staining if no smooth muscle myosin heavy chain-expression cells exist in the tissue. Interpretation of the staining result is solely the responsibility of the user.


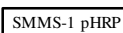



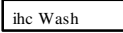







#### General Limitations:

Immunohistochemistry is a multistep diagnostic process that requires 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. 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 (Nadji M, Morales AR. 1983).

The manufacturer provides these antibodies/reagents at optimal dilution for use following the provided instructions for IHC on prepared tissue sections. Any deviation from recommended test procedures may invalidate declared expected results; appropriate controls must be employed and documented. Users who deviate from recommended test procedures must accept responsibility for interpretation of patient results under these circumstances.

#### Performance Characteristics:

The ihcDirect SMMS-1 pHRP test performance has been determined using both frozen and FFPE tissue sections. NovodiAx has conducted studies to evaluate the performance of the antibody conjugates and recommended ancillary reagents and supplies. The antibodies and systems have been found to be sensitive and show specific binding to the antigen of interest with minimal to no binding of non-specific tissues or cells. NovodiAx antibodies and ancillary reagents have shown reproducible and consistent results when used within a single run, between runs and between lots. These products have been determined to be stable for the periods of time specified on the labels either by standard real-time and/or accelerated methods. NovodiAx ensures product quality by testing each lot of material and by testing materials at regular intervals and via surveillance programs.

ihcDirect SMMS-1 Key to Symbols			
	In vitro diagnostic medical device		pHRP SMMS-1 antibody conjugate
	Catalog Number		Blocking reagent
	Use by: YYYY-MM-DD		ihc Wash Buffer
	Consult Instruction for Use		Manufacturer
	Batch Code		Authorized European Representative
	Contains sufficient for < n > tests		CE Mark
	Temperature Limitation		

#### Instructions for Use (IFU) Access:

To obtain a translation or the latest electronic version of an IFU document, visit our website at <https://www.novodiAx.com/literature/instructions-for-use-ifu/>. Printed copies of an IFU document may be obtained by contacting NovodiAx Technical Support or your local distributor.

#### Bibliography:

- Kalof AN et al. Immunostaining patterns of myoepithelial cells in breast lesions: a comparison of CD10 and smooth muscle myosin heavy chain. J Clin Pathol. 2004; 57: 625-629.
- Hill CB, Yeh IT. Myoepithelial cell staining patterns of papillary breast lesions: from intraductal papillomas to invasive papillary carcinomas. AM J Clin Pathol. 2005 123(1) 36-44.
- Kiernan JA. Histological and Histochemical Methods: Theory and Practice. New York: Pergamon Press 1981.
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