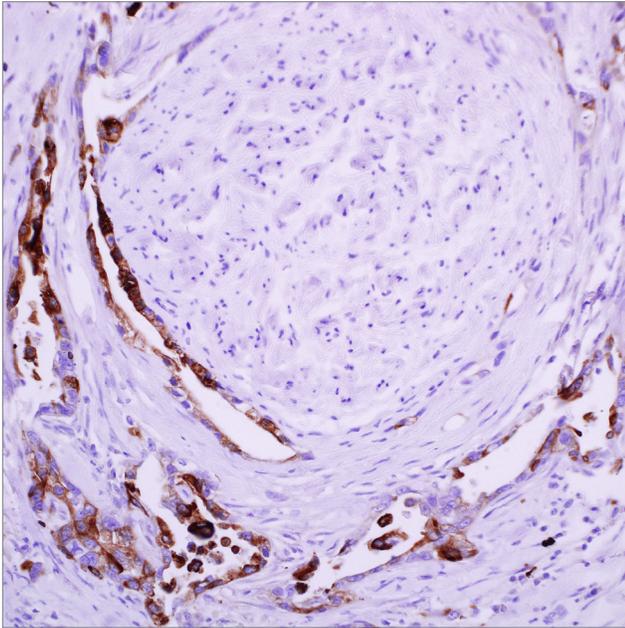


# MUC4 (8G7) Mouse Monoclonal Antibody

**For In Vitro Diagnostic Use (IVD)**



## Product Identification

REF	Description
406S	Positive control slides, 5 slides/pack
406M-14	0.1 ml concentrate
406M-15	0.5 ml concentrate
406M-16	1.0 ml concentrate
406M-17	1.0 ml predilute ready-to-use
406M-18	7.0 ml predilute ready-to-use

## Symbol Definitions

KEY-CODE	keycode
P	predilute
C	concentrate
A	ascites
E	serum
S	supernatant
DIL	concentrate dilution range

## Intended Use

This antibody is intended for *in vitro* diagnostic (IVD) use.

The MUC4 (8G7) antibody is intended for qualified laboratories to qualitatively identify by light microscopy the presence of associated antigens in sections of formalin-fixed, paraffin-embedded tissue sections using IHC test methods. Use of this antibody is indicated, subsequent to clinical differential diagnosis, as an aid in the identification of pancreatic ductal adenocarcinoma within the context of an antibody panel, the patient's clinical history, and other diagnostic tests evaluated by a qualified pathologist.

## Summary and Explanation

Mucin 4 (MUC4) is a transmembranous glycoprotein. MUC4 is expressed in the cytoplasm of certain epithelial surfaces, such as colonic, breast, and lung epithelia.<sup>1</sup> An abnormal expression of MUC4 has been reported in various carcinomas of the colon, pancreas, breast, and ovaries.<sup>2</sup> MUC4 is highly expressed in the vast majority of human pancreatic neoplasms, such as intraductal papillary mucinous neoplasia of the pancreas, pancreatic intraepithelial neoplasms and pancreatic ductal adenocarcinoma; however, it is not detected in the normal pancreas or in chronic pancreatitis.<sup>1-3</sup> MUC4 is not expressed in normal bile ducts, but it can be overexpressed in intrahepatic cholangiocarcinoma and bile duct cholangiocarcinoma.<sup>3</sup> MUC4 overexpression has been reported in low-grade fibromyxoid sarcoma (LGFMS).<sup>4</sup> In this study a large cohort of LGFMS, all with FUS gene rearrangement confirmed by FISH, showed cytoplasmic staining for MUC4, usually in a strong and diffuse manner.<sup>4</sup> Similarly, strong, diffuse cytoplasmic staining for MUC4 was identified in 78% (32/41) of cases of sclerosing epithelioid fibrosarcoma.<sup>5</sup> MUC4 expression is also detected in the glandular component of biphasic synovial sarcomas (90%).<sup>5</sup> Focal staining, usually interpreted as only scattered cells, was also seen in a subset of ossifying fibromyxoid tumors (29%), epithelioid GISTs (20%) and myoepithelial carcinomas (10%); whereas all other epithelioid soft tissue tumors, including clear cell sarcoma, epithelioid sarcoma, epithelioid hemangiosarcoma, PEComa and melanoma, were negative. It should be noted that various carcinomas may express MUC4, and therefore additional keratins or lineage-specific markers may be needed to exclude this possibility in some cases.<sup>5</sup>

## Principles and Procedures

The stated primary antibody may be used as the primary antibody for immunohistochemical staining of formalin-fixed, paraffin-embedded tissue sections. In general, immunohistochemical staining in conjunction with a HRP or Alk Phos linked detection system allows the visualization of antigens via the sequential application of a specific antibody (primary antibody) to the antigen, a secondary antibody (link antibody) to the primary antibody, an enzyme complex and a chromogenic substrate with interposed washing steps. The enzymatic activation of the chromogen results in a visible reaction product at the antigen site. The specimen may then be counterstained and a coverslip applied. Results are interpreted using a light microscope.

## Materials and Methods

### Reagents Provided

Product Composition	
Predilute: diluted in	Tris Buffer, pH 7.3-7.7, with 1% BSA and <0.1% Sodium Azide
Concentrate: diluted in	Tris Buffer, pH 7.3-7.7, with 1% BSA and <0.1% Sodium Azide
Isotype	IgG <sub>1</sub> /k
Recommended working dilution range	1:10-1:50

See product label for lot specific information for the following:

1. Antibody immunoglobulin concentration
2. Source details

### Reconstitution, Mixing, Dilution, Titration

Prediluted antibody is ready-to-use and optimized for staining. No reconstitution, mixing, dilution, or titration is required. The concentrated antibody is optimized to be diluted to within the dilution range using Cell Marque Diamond Diluent.

### Materials and Reagents Needed But Not Provided

The following reagents and materials may be required for staining but are not provided with the primary antibody:

1. Positive and negative control tissue
  2. Microscope slides, positively charged
  3. Drying oven capable of maintaining a temperature of 53-65°C
  4. Staining jars or baths
  5. Timer
  6. Xylene or xylene substitute
  7. Ethanol or reagent alcohol
- Note: Cell Marque's one-step pretreatment, Trilogy™ (cat. #920P-06), can replace both 6 and 7 above.*
8. Deionized or distilled water
  9. Heating Equipment, such as Electric Pressure Cooker (cat. #976L), for tissue pretreatment step
  10. Detection system, such as HiDef Detection™ HRP Polymer System (cat. #954D-20) or HiDef Detection™ Alk Phos Polymer System (cat. #962D-20)
  11. Chromogen, such as DAB Substrate Kit (cat. #957D-20) or Permanent Red Chromogen Kit (cat. #960D-10)
  12. TBS IHC Wash Buffer + Tween®\* 20 (cat. #935B-09)
  13. Hematoxylin (cat. #930B-05) or other counterstain
  14. Antibody diluents, such as Diamond: Antibody Diluent (cat. #938B-05) or Emerald: Antibody Diluent (cat. #936B-08)
  15. Peroxide Block (cat. #925B-05) for use with HRP
  16. Avidin-Biotin Blocking Reagents (cat. # 928B-02 for use with streptavidin-biotin detection)
  17. Negative Control Reagent (cat. #932B-02 for mouse; cat. #933B-02 for rabbit, cat. #939B-02 for universal)
  18. Permanent Aqueous Mounting Medium (cat. #931B-03)

19. Cover glass
20. Light microscope (40-400x)

### Storage and Handling

Store at 2-8°C. Do not freeze.

To ensure proper reagent delivery and stability of the antibody after every run, the cap must be replaced and the bottle must be immediately placed in the refrigerator in an upright position.

Every antibody reagent is expiration dated. When properly stored, the reagent is stable to the date indicated on the label. Do not use reagent beyond the expiration date for the prescribed storage method.

There are no definitive signs to indicate instability of this product; therefore, positive and negative controls should be run simultaneously with unknown specimens. Contact Cell Marque customer service if there is a suspected indication of reagent instability.

### Specimen Collection and Preparation for Analysis

Routinely processed, neutral-buffered formalin-fixed, paraffin-embedded, tissues are suitable for use with this primary antibody when used with Cell Marque detection kits (see Materials, Reagents, and Equipment Needed But Not Provided section). Note: Cell Marque evaluates performance only on human tissues. The recommended tissue fixative is 10% neutral-buffered formalin. Variable results may occur as a result of prolonged fixation or special processes such as decalcification of bone marrow preparations.

Each section should be cut to the appropriate thickness (approximately 3 µm) and placed on a positively charged glass slide. Slides containing the tissue section may be baked for at least 2 hours (but not longer than 24 hours) in a 53-65°C oven.

### Warnings and Precautions

1. Take reasonable precautions when handling reagents. Use disposable gloves and lab coats when handling suspected carcinogens or toxic materials (example: xylene).
2. Avoid contact of reagents with eyes and mucous membranes. If reagents come in contact with sensitive areas, wash with copious amounts of water.
3. Patient specimens and all materials contacting them should be handled as biohazardous materials and disposed of with proper precautions. Never pipette by mouth.
4. Avoid microbial contamination of reagents, as this could produce incorrect results.
5. The user must validate incubation times and temperatures.
6. The prediluted, ready-to-use reagents are optimally diluted, and further dilution may result in loss of antigen staining.
7. The concentrated reagents may be diluted optimally based on validation by user. Any diluent used that is not specifically recommended herein must likewise be validated by the user for both its compatibility and effect on stability.
8. When used according to instructions, this product is not classified as a hazardous substance. The preservative in the reagent is less than 0.1% sodium azide and does not meet the OSHA (USA) criteria for hazardous substance at the stated concentration. See SDS.
9. The user must validate any storage conditions other than those specified in the package insert.
10. Diluent may contain bovine serum albumin and supernatant may contain bovine serum. The products containing fetal bovine serum and products containing bovine serum albumin are purchased from commercial suppliers. Certificates of Origin for the animal

source used in these products are on file at Cell Marque. The certificates support that the bovine sources are from countries with negligible BSE risk and state sources of bovine from USA and Canada.

- As with any product derived from biological sources, proper handling procedures should be used.

## Instructions For Use

### Recommended Staining Protocols for the stated primary antibody:

#### HiDef Detection™ HRP:

HiDef Detection™ HRP Polymer System (cat. #954D-20)

- Epitope Retrieval Technique: HIER, Epitope Retrieval Reagent: Trilogy
- Antibody Incubation Time (Minutes): 5
- HiDef Detection Amplifier Incubation Time (Minutes): 10
- HiDef Detection Polymer Detector Incubation Time (Minutes): 10
- DAB Incubation Time (Minutes): 1-10
- Dehydrate and Coverslip.

#### HiDef Detection™ Alk Phos:

HiDef Detection™ Alk Phos Polymer System (cat. #962D-20)

- Epitope Retrieval Technique: HIER, Epitope Retrieval Reagent: Trilogy
- Antibody Incubation Time (Minutes): 5
- HiDef Detection Amplifier Incubation Time (Minutes): 10
- HiDef Detection Polymer Detector Incubation Time (Minutes): 10
- DAB Incubation Time (Minutes): 15-30
- Dehydrate and Coverslip.

## Quality Control Procedures

### Positive Tissue Control

A positive tissue control must be run with every staining procedure performed. This tissue may contain both positive and negative staining cells or tissue components and serve as both the positive and negative control tissue. Control tissues should be fresh autopsy, biopsy or surgical specimens prepared or fixed as soon as possible in a manner identical to the test sections. Use of a tissue section fixed or processed differently from the test specimen will serve to provide control for all reagents and method steps except fixation and tissue processing.

A tissue with weak positive staining is more suitable for optimal quality control and for detecting minor levels of reagent degradation. Positive tissue control for the stated primary antibody may include the following:

Positive Tissue Control	
Tissue	Visualization
Pancreatic Ductal Adenocarcinoma	Cytoplasmic
Colon	Cytoplasmic
Colorectal Adenocarcinoma	Cytoplasmic

Known positive tissue controls should be utilized only for monitoring the correct performance of processed tissues and test reagents, not as an aid in determining a specific diagnosis of patient samples. If the positive tissue controls fail to demonstrate appropriate positive staining, results with the test specimens must be considered invalid.

### Negative Tissue Control

The same tissue used for the positive tissue control may be used as the negative tissue control. The variety of cell types present 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.

### Unexplained Discrepancies

Unexplained discrepancies in controls should be referred to Cell Marque Customer Service immediately. If quality control results do not meet specifications, patient results are invalid. See the Troubleshooting section of this insert. Identify and correct the problem, then repeat the entire procedure with the patient samples.

### Negative Control Reagent

A negative control reagent must be run for every specimen to aid in the interpretation of results. A negative control reagent is used in place of the primary antibody to evaluate nonspecific staining. The slide should be treated with negative control reagent, matching the host species of the primary antibody, and ideally having the same IgG concentration. The incubation period for the negative control reagent should equal the primary antibody incubation period.

## Interpretation of Results

The immunostaining procedure causes a colored reaction product to precipitate at the antigen sites localized by the primary antibody. Refer to the appropriate detection system package insert for expected color reactions. A qualified pathologist experienced in immunohistochemistry procedures must evaluate positive and negative tissue controls before interpreting results.

### Positive Tissue Control

The stained positive tissue control should be examined first to ascertain that all reagents are functioning properly. The presence of an appropriately colored reaction product within the target cells is indicative of positive reactivity. Refer to the package insert of the detection system used for expected color reactions. Depending on the incubation length and potency of the hematoxylin used, counterstaining will result in a pale to dark blue coloration of cell nuclei. Excessive or incomplete counterstaining may compromise proper interpretation of results. If the positive tissue control fails to demonstrate appropriate positive staining, any results with the test specimens are considered invalid.

### Negative Tissue Control

The negative tissue control should be examined after the positive tissue control to verify the specific 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 or cellular components. If specific staining occurs in the negative tissue control, results with the patient specimen are considered invalid. Nonspecific staining, if present, will have a diffuse appearance. Sporadic light staining of connective tissue may also be observed in sections

from tissues that are not optimally fixed. Intact cells should be used for interpretation of staining results. Necrotic or degenerated cells show non-specific staining.

### Patient Tissue

Patient specimens should be examined last. Positive staining intensity should be assessed within the context of any background staining of the negative reagent control. As with any immunohistochemical test, a negative result means that the antigen in question was not detected, not that the antigen is absent in the cells or tissue assayed. A panel of antibodies may aid in the identification of false negative reactions (see Summary of Expected Results section). The morphology of each tissue sample should also be examined utilizing a hematoxylin and eosin stained section when interpreting any immunohistochemical result. The patient's morphologic findings and pertinent clinical data must be interpreted by a qualified pathologist.

### Limitations

- Color does not affect performance.
- This reagent is "for professional use only" as immunohistochemistry is a multiple step process that requires specialized training in the selection of the appropriate reagents, tissues, fixation, processing; preparation of the immunohistochemistry slide; and interpretation of the staining results.
- For laboratory use only.
- For *in vitro* diagnostic use.
- 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 result from variations in fixation and embedding methods, as well as from inherent irregularities within the tissue.
- Excessive or incomplete counterstaining may compromise proper interpretation of results.
- The clinical interpretation of any positive staining, or its absence, must be evaluated within the context of clinical history, morphology, other histopathological criteria as well as other diagnostic tests. This antibody is intended to be used in a panel of antibodies if applicable. It is the responsibility of a qualified pathologist to be familiar with the antibodies, reagents, diagnostic panels, and methods used to produce the stained preparation. Staining must be performed in a certified, licensed laboratory under the supervision of a pathologist who is responsible for reviewing the stained slides and assuring the adequacy of positive and negative controls.
- Cell Marque provides antibodies and reagents at optimal dilution for use as instructed. Any deviation from recommended test procedures may invalidate expected results. Appropriate controls must be employed and documented. Users in any circumstance must accept responsibility for interpretation of patient results.
- Cell Marque provides primary antibodies in concentrated format so that the user may subsequently optimally dilute for use subject to the user's determination of and adherence to suitable validation techniques. Users must validate the use of any diluents other than what is recommended herein. Once the primary is validated to be suitable for use, any deviation from recommended test procedures may invalidate expected results. Appropriate controls must be employed and documented. Users in any circumstance must accept responsibility for interpretation of patient results.
- This product is not intended for use in flow cytometry.
- Reagents may demonstrate unexpected reactions in previously untested tissues. The possibility of unexpected reactions even in tested tissue groups cannot be completely

eliminated because of biological variability of antigen expression in neoplasms, or other pathological tissues. Contact Cell Marque customer service with any suspected, documented unexpected reactions.

- Tissues from persons infected with hepatitis B virus and containing hepatitis B surface antigen (HBsAg) may exhibit nonspecific staining with horseradish peroxidase.
- When used in blocking steps, normal sera from the same animal source as the secondary antisera may cause false negative or false positive results because of the effect of autoantibodies or natural antibodies.
- False positive results may be seen because of non immunological binding of proteins or substrate reaction products. They may also be caused by pseudoperoxidase activity (erythrocytes), endogenous peroxidase activity (cytochrome C), or endogenous biotin (example: liver, brain, breast, kidney) subject to the type of immunostaining technique used.
- As with any immunohistochemistry test, a negative result means that the antigen was not detected, not that the antigen was absent in the cells or tissue assayed.
- The prediluted antibody products are optimized as a ready-to-use product. Because of the possibility of variation in tissue fixation and processing, it may be necessary to increase or decrease the primary antibody incubation time on individual specimens.
- The antibody, in combination with detection systems and accessories, detects antigen(s) that survive routine formalin fixation, tissue processing and sectioning. Users who deviate from recommended test procedures remain, as they would in any circumstance, responsible for interpretation and validation of patient results.

### Summary of Expected Results

See the following tables of reactivity:

Normal Study			
Tissue	# Stained (+)	Total #	Notes
Brain	0	2	
Adrenal Cortex	0	2	
Ovary	0	1	
Pancreas	0	5	Occasional large ducts stained
Parathyroid	0	1	
Pituitary	0	1	
Testis	0	2	
Thyroid	0	1	
Breast	1	1	Ducts weakly stained
Spleen	0	1	
Tonsil	0	1	Squamous stained
Thymus	2	2	Epithelium stained
Bone Marrow	0	2	
Lung	3	3	Bronchial epithelium stained
Heart	0	1	
Esophagus	1	1	
Stomach	1	1	Deep glands stained
Small Intestine	3	3	
Colon	2	2	
Liver	0	1	
Salivary Gland	0	1	

Normal Study (continued)			
Tissue	# Stained (+)	Total #	Notes
Gall Bladder	2	3	
Kidney	0	1	
Bladder	1	1	
Prostate	0	1	Occasional acini stained
Uterus	0	2	
Fallopian Tube	0	1	
Ureter	0	1	
Cervix	1	2	Endocervix stained
Skeletal Muscle	0	1	
Smooth Muscle	0	1	
Skin	0	1	
Peripheral Nerve	0	1	
Mesothelium	0	1	
Fat	0	1	
Placenta	0	1	

This antibody performs as indicated in the literature.

Disease Tissue Study			
Tissue	# Stained (+)	Total #	Notes
Renal cell carcinoma	22	23	
Colorectal adenocarcinoma	13	19	
Pancreatic ductal adenocarcinoma	10	10	
Lung adenocarcinoma	9	10	
Stomach carcinoma intestinal type	9	9	
Stomach carcinoma diffuse type	8	10	
Lung squamous cell carcinoma	7	11	
Transitional cell carcinoma	5	9	
Esophageal Adenocarcinoma	4	5	
Duodenal Adenocarcinoma	1	1	
Gall Bladder Adenocarcinoma	1	4	
Hepatocellular carcinoma	1	9	
Rhabdomyosarcoma	1	4	
Synovial Sarcoma	1	8	
Breast invasive ductal carcinoma	0	23	
Diffuse large B-cell lymphoma	0	4	
Fibrosarcoma	0	6	
Follicular lymphoma	0	3	
Leiomyosarcoma	0	4	
Prostate carcinoma	0	9	
Small lymphocytic lymphoma	0	3	

This antibody performs as indicated in the literature.

## Troubleshooting

1. If the positive control exhibits weaker staining than expected, other positive controls run during the same staining run should be checked to determine if it is because of the primary antibody or one of the common secondary reagents.

2. If the positive control is negative, other positive controls used on the same run should be checked to determine if the underlying cause relates to the primary antibody or one of the common secondary reagents. Tissues may have been improperly collected, fixed or deparaffinized. The proper procedure should be followed for collection, storage, and fixation.
3. If excessive background staining occurs, high levels of endogenous biotin may be present. A biotin blocking step should be included unless a biotin-free detection system is being used in which case any biotin present would not be a contributing factor to background staining.
4. If all of the paraffin has not been removed, the deparaffinization procedure should be repeated.
5. If tissue sections wash off the slide, slides should be checked to ensure that they are positively charged. Other possibilities that could have adverse affect on tissue adhesion include insufficient drying of the tissue section on the slide prior to staining or fixation in formalin that was not properly neutral-buffered. Tissue thickness may also be a contributing factor.

For corrective action, refer to the Instructions for Use section or contact Cell Marque customer service.

## References

1. Moniaux N, et al. Generation and characterization of anti-muc4 monoclonal antibodies reactive with normal and cancer cells in humans. *J Histochem Cytochem.* 2004; 52:253-61.
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3. Yonezawa S, et al. Significance of mucin expression in pancreatobiliary neoplasms. *J Hepatobiliary Pancreat Sci.* 2010; 17:108-24.
4. Doyle LA, et al. MUC4 is a highly sensitive and specific marker for low-grade fibromyxoid sarcoma. *Am J Surg Pathol.* 2011; 35:733-41.
5. Doyle LA, et al. MUC4 is a sensitive and extremely useful marker for sclerosing epithelioid fibrosarcoma: association with fus gene rearrangement. *Am J Surg Pathol.* 2012; 36:1444-51.

## Disclaimers

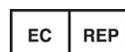
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