

Mouse Monoclonal anti-Desmin, Clone GM007

60-0077; 60-0077-7 61-0077; 61-0077-2; 61-0077-5

6 mL; 7 mL predilute Antibody, Ready-To-Use 1 mL; 0.2 mL; 0.5 mL Concentrate Antibody lgG1

Isotype

Concentration See container label

Intended Use For In Vitro Diagnostic Use.

> This product is intended for laboratory use to qualitatively detect Desmin by light microscopy in normal and neoplastic formalin fixed paraffin embedded (FFPE) tissue sections using immunohistochemical (IHC) methodology. Interpretation of any positive or negative staining shall be supported by a proper control and must be made within the context of the patient's clinical

history and other diagnostic test by a qualified pathologist.

Desmin is a characteristic intermediate filament of all three types of muscle cells (skeletal, cardiac, Description

and smooth muscle) and neoplasms associated with them. In general, desmin is a specific marker for myogenic differentiation among soft tissue tumors. It is seen in the majority of rhadbomyomas, leiomyomas, rhadbomyosarcoma, and leiomyosarcomas. Desmin is also seen in myofibroblasts.

Myoepithelial cells typically lack desmin.

The antibody labels smooth and striated muscle cells as well as mesothelial cells. It allows the subtyping of many undifferentiated and pleomorphic tumors through intermediate filament analysis. With selected panels of antibodies, it is a useful tool to separate the different pleomorphic spindle cell tumors and round cell tumors in soft tissues and skin. The antibody labels strongly reactive mesothelial cells, but not malignant mesothelioma and adenocarcinoma.

This antibody is purified antibody diluted in 10 mM phosphate buffered saline (PBS), pH 7.2 Reagent provided

containing 1% bovine serum albumin (BSA) and 0.09% sodium azide (NaN₃) as antimicrobial

agent.

Precautions For professional users.

Proper handling of this product as with any product derived from biological sources shall be

followed according to local and applicable regulations.

Sodium azide is a toxic chemical. The concentration in this product is not classified as hazardous, however, the build-ups of NaN₃ may react with lead and copper plumbing to form highly explosive metal azides. Flush the disposed reagent with large volume of water to prevent azide build-up.

Usage

Dilution 60-0077; 60-0077-7: Ready-To-Use

> 61-0077; 61-0077-2; 61-0077-5: Dilute 1:50 to 1:100 before use when using Acu-Stain[™] detection system. Optimum dilution factor may vary depending on the specimen and preparation process

and should be determined by each individual investigator.

Staining procedure Incubate this antibody with tissue section for 30-60 minutes at room temperature. Follow the

instructions from the selected detection system.

Positive control tissue Uterus

Epitope retrieval Not Required. HIER, Citrate pH 6 may enhance staining in some tissues.

Staining pattern Cytoplasmic

Storage Store at 2-8°C.

1. Miettinen M, et al. Am J Surg Pathol. 2000 Feb;24(2):211-22. References

2. Hurlimann J. Hum Pathol. 1994 Aug;25(8):753-7.

Symbols REF LOT IVD Catalog No. Batch No. In Vitro Diagnostic Use Temperature Range Use By

30331 Rev.03







